MEETING ABSTRACTS

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Recurrent fevers

O1
Efficacy and safety of Canakinumab in patients with periodic fever syndromes (colchicine-resistant fmf, hids/mkd and traps): results from a phase 3, pivotal, umbrella trial
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Pediatric Rheumatology 2017, 15(Suppl 1):O1

Introduction: Periodic fever syndromes (PFS) are rare auto-inflammatory conditions that include, among others, cryopyrin-associated periodic syndromes (CAPS), familial Mediterranean fever (FMF), hyper-IGD syndrome/mevalonate kinase deficiency (HIDS/MKD), TNF receptor-associated periodic syndrome (TRAPS) [1]. Canakinumab (CAN), a fully human, highly specific anti-IL-1β neutralising monoclonal antibody, is effective in CAPS [2]. IL-1β has been shown to be involved in the pathogenesis of FMF, HIDS/MKD and TRAPS, for which no or limited treatment options exist [1]. Open-label studies have suggested the efficacy of CAN in colchicine-resistant/intolerant FMF (crFMF), HIDS/MKD and TRAPS [3-5]. We report the efficacy and safety of CAN from a randomised treatment epoch of a Phase 3 pivotal study in patients (pts) with crFMF, HIDS/MKD or TRAPS.

Objectives: Primary objective was to demonstrate that CAN 150 mg (or 2 mg/kg for pts <40 kg) q 2 wks is superior to placebo (PBO) in achieving a clinically meaningful response defined as resolution of the index flare at Day 15 and no new disease flares over 16 weeks (wks) of treatment. Secondary objectives included proportion of pts who achieved a physician global assessment (PGA) of disease activity <2 (minimal/none) and proportions of pts with C-reactive protein (CRP) ≤10 mg/L and serum amyloid A (SAA) ≤10 mg/L at Wk 16.

Methods: The study consists of 3 disease cohorts (crFMF, HIDS/MKD and TRAPS) and 4 study epochs: a screening epoch (E1) of up to 12 wks, a randomised treatment epoch (E2) of 16 wks, a randomised withdrawal epoch (E3) of 24 wks and an open-label treatment epoch (E4) of 72 wks. Pts (aged ≥2 years) with a flare during E1 were randomised (1:1) in E2 to receive CAN or PBO. Safety assessments included adverse events (AEs) and serious AEs (SAEs).

Results: Of 181 pts (crFMF, n = 63; HIDS/MKD, n = 72; TRAPS, n = 46) randomised in E2, 6 discontinued (5 PBO; 1 CAN). In all 3 disease cohorts, the proportion of responders for the primary outcome at Wk 16 was significantly higher with CAN vs PBO (Table). At Wk 16, a significantly higher proportion of pts achieved a PGA score <2, CRP ≤10 mg/L and SAA ≤10 mg/L in the CAN group vs PBO in all 3 cohorts. The most frequently affected system organ class across 3 cohorts was infections and infestations typically involving the upper respiratory tract. The incidence of SAEs was 8.6%, 4.7% and 11.8% in crFMF, TRAPS and HIDS/MKD cohort, respectively.

Conclusion: These results demonstrated superior efficacy of canakinumab after a 16-week treatment period compared with placebo. The overall safety profile was not distinct from those reported in previous controlled studies.

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Trial registration identifying number: NCT02059291

Disclosure of Interest
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Table 1 (abstract O1). Proportion of responders at Week 16 by cohort

| Cohort | Canakinumab (150 mg q4w), n (%) | Placebo n/m (%) | p-value |
|--------|---------------------------------|-----------------|---------|
| crFMF  | 19/31 (61.3)                    | 2/52 (3.8)      | <0.001* |
| HIDS/MKD| 13/37 (35.1)                    | 2/35 (5.7)      | 0.0020* |
| TRAPS  | 10/22 (45.5)                    | 2/24 (8.3)      | 0.0050* |

*n number of patients who responded, m number of patients evaluated for response

*Statistical significance (1-sided) at the 0.025 level based on Fisher exact test/logistic regression model

O2
An update on the Italian cohort of DADA2 patients and an assessment of a novel functional screening test
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This abstract is not included here as it has already been published.

Mechanisms of disease

O3
Inhibition of interferon production driving tmem173-associated auto-inflammation
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Pediatric Rheumatology 2017, 15(Suppl 1):O3

Introduction: Gain-of-function mutations in TMEM173 encoding STING (Stimulator of Interferon Genes) underlie a novel type I interferonopathy, minimally responsive to conventional immunosuppressive therapies and associated with high childhood morbidity and mortality. A newly emerging treatment strategy in STING-related inflammation aims to control interferon (IFN) signalling post-binding of the IFN receptor, by targeting JAK1/2. We hypothesized that inhibition of IFN production itself might represent an alternative therapeutic approach in this disease.

Objectives: To evaluate the effect of BX795, a TBK1 inhibitor, on constitutive production of type I IFN in TMEM173-associated vasculopathy.

Methods: Human embryonic kidney 293 cells (HEKs) transfected with wild-type (WT) or mutant STING, and peripheral blood mononuclear cells (PBMCs) from three patients were treated with BX795 2 µM. The effect of BX795 on TBK1 signalling cascading was assessed in HEKs by western blotting and IFNβ promoter activity induced by cGAMP or mutant STING. In patient-derived PBMCs, exposure to BX795 inhibited IFNα production measured by digital ELISA, and was associated with a decrease of STAT1 phosphorylation and reduced ISG transcription. In addition, BX795 decreased the positive feedback loop of STING on the type I IFN pathway.

Results: In HEKs, BX795 inhibited the phosphorylation of IRF3 and IFNβ promoter activity induced by cGAMP or mutant STING. In patient-derived PBMCs, exposure to BX795 inhibited IFNα production measured by digital ELISA, and was associated with a decrease of STAT1 phosphorylation and reduced ISG transcription. In addition, BX795 decreased the positive feedback loop of STING on the type I IFN pathway.

Conclusion: Our findings demonstrate that inhibition of IFNα secretion is a potential approach to control disease-associated inflammation in patients with gain-of-function mutations in TMEM173, and may also be relevant in other monogenic type I interferonopathies.

Disclosure of Interest
None Declared

O4
NLRP3 inflammasome activity in monocytes is regulated by 12/15-lipoxygenase
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Pediatric Rheumatology 2017, 15(Suppl 1):O4

This abstract is not included here as it has already been published.

Other autoinflammatory diseases

O5
PLAS: a novel autoinflammatory and lymphoproliferative syndrome associated with PIM-1 mutations
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Introduction: We present a novel syndrome characterised by both lymphoproliferative and autoinflammatory features, due to a PIM-1 kinase gene mutation (PIM-1 associated Lymphoproliferative Autoinflammatory Syndrome, PLAS).

Objectives: To describe clinical and genetic features in two cases with PLAS.

Methods: Whole exome sequencing (WES) analysis with trio based strategy in the first case and direct sequencing of candidate gene in the second one.

Results: A 35-year-old man was referred to ophthalmologist’s evaluation for blurry vision in his left eye. The fundus examination showed confluent, yellowish choroidal lesions in both eyes. During childhood, his past medical history was relevant for a diagnosis of celiac disease, recurrent episodes of protracted unexplained fever and skin rashes. He also referred unexplained inflammatory lesions of the osteoarticular and muscular system, one episode of aseptic meningitis, an intra-crinal granuloma with features of vascular proliferation and two episodes of anterior uveitis. In adulthood, his medical history included episodes of fever with urticaria and subcutaneous nodules. The histological examination of two skin lesions showed a reactive angioendotheliomatosis and a leukocytoclastic vasculitis, respectively.

Since childhood its laboratory data ever showed persistent elevated reactive protein (60–80 mg/l) and polyclonal gammopathy (IgG
3600 mg/dl; IgM 126 mg/dl), elevated serum amyloid A levels (160 mg/ml) and decreased CD4/CD8 T cell ratio. Autoantibody (ANA, ENA, anti-DNA) and HLA B 27 were negative. WES analysis showed the de novo heterozygous missense variation c.C1132A (p.H378N) in PIM-1 gene (NM_001243186). This variation was never described in on-line database (HGMD, Exac, 1000 g, ESP6000). The second patient was an adult female with a clinical history of persistent microcytic anemia, splenomegaly, striking hypergammaglobulinemia (IgG > 3000 mg/dL) and recurrent protracted fever episodes during childhood. At 13 years, she underwent splenectomy because of hypersplenism. At 28 years she presented a cerebral oligoastrocytoma that was removed by surgery. No mutations were found in genes associated with autoimmune lymphoproliferative syndromes and histiocytosis (FAS, FASLG, XIAP, TNFRSF13B, UNC13D, CASP9, CASP10).

The direct sequencing of PIM-1 revealed again the above described de novo mutation in PIM-1.

Conclusion: We propose that these patients represent two cases of a novel syndrome associated with a specific mutation in PIM-1 gene (PILAS). First, the two cases share significant overlap of autoinflammatory and proliferative features involving the endothelial and the immune systems. Second, the gene mutation identified was never described in healthy people or in patients with other disorders and is predicted to be pathogenic based on all the bioinformatic tools. Finally, previous functional studies showed that PIM-1 pathway is relevant to immune activation and indeed somatic mutations in this gene have been reported in association with lymphomas, and it is thought to contribute to survival of cancerous cells.

Disclosure of Interest
None Declared

Diagnosis of Disease

O6 Biallelic hypomorphic mutations in a linear deubiquitinase define otulipenia, an early-onset systemic autoinflammatory disease

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Pediatric Rheumatology 2017, 15(Suppl 1):O6

Introduction: Autoinflammatory diseases are caused by mutations in genes regulating innate immune responses. More than 20 genes have been associated with various monogenic autoinflammatory disorders; however there is a significant number of undiagnosed patients including familial cases.

Objectives: To identify a disease-causing gene in two inbred families presenting with a neonatal-onset systemic inflammatory disease. We expected to find homozygous pathogenic mutations and subsequently to screen other undiagnosed patients who present with similar disease.

Methods: We performed whole-exome and candidate gene sequencing in the patients and their unaffected family members. NFkB luciferase assay and overexpression experiments in HEK 293 cells were used to confirm the causality of the mutations. Patient samples were analyzed using immunoprecipitation, immunoblotting, gene expression and cytokine profiling.

Results: We studied 3 unrelated patients, one of Pakistani and two of Turkish ancestry, with early-onset systemic inflammation and generalized erythematous rash diagnosed as neutrophilic dermatitis/panniculitis, lipoedema, failure to thrive, and without obvious primary immunodeficiency.

We identified three novel homozygous mutations in the FAM105B gene, which encodes OTULIN, the deubiquitinase that specifically hydrolyzes Met1-linked ubiquitin chains (Ub) from target proteins. Post-translational modifications by ubiquitination are important for the regulation of many signaling complexes including the NF-kB pathway. Met1-Ub chains are generated by the linear ubiquitin assembly complex (LUBAC). OTULIN is an evolutionarily highly conserved protein and in mice complete deficiency is embryonically lethal.

The p.Leu272Pro and p.Tyr244Cys mutations are located near the linear ubiquitination binding region, while the p.Gly174Aspfs12 mutation truncates the protein. Cells transfected with mutated versus wildtype OTULIN had decreased enzyme activity and showed a substantial defect in the linear ubiquitination of target molecules NEMO, RIPK1, TNFR1 and ASC. Stimulated patients’ cells, fibroblasts and PBMCs, showed increased phosphorylation of IKKα/β, IκBα, JNK, P38 and a higher linear-ubiquitination level of the target proteins. These results indicate that inefficient deubiquitination of OTULIN target proteins might explain increased NF-kB activity in mutant cells. Levels of many proinflammatory cytokines including IL-1β, TNF, IL-6, IL-18, IL-12, and IFN-γ were substantially increased in patient serum samples and stimulated cells. Cytokine profiling in serum samples was consistent with disease activity. Transcriptome profiling of patient whole blood samples and stimulated fibroblasts showed similar results. One patient had excellent response to treatment with the TNF inhibitor, infliximab. Two other patients have been treated with either etanercept or anakinra, but they still have an active disease requiring steroids.

Conclusion: Loss-of-function mutations in OTULIN result in increased linear ubiquitination of signaling molecules and lead to enhanced TNFR1, NF-kB, and ASC-dependent inflammatory activity. This new disease, which we named otulipenia, adds to the emerging spectrum of human diseases caused by defects in the ubiquitin pathway and suggests a role for targeted cytokine therapies.

Disclosure of Interest
None Declared

Differential Diagnosis

O7 Discrimination among monogenic and acquired disease with inflammatory and autoimmune component through the in vitro study of signal transducers of type I interferon

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This abstract is not included here as it has already been published.

O8 A targeted next-generation sequencing gene panel for autoinflammation

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Pediatric Rheumatology 2017, 15(Suppl 1):O8
Introduction: Monogenic autoinflammatory diseases (AID) are severe lifelong systemic inflammatory disorders with dysregulated innate immunity, causing significant morbidity, mortality, and economic burden. The number of AID is rapidly expanding. Securing a molecular diagnosis is of major importance for treatment, prognosis, and genetic counselling. Routine genetic screening is time-consuming, costly, and lacks sensitivity since only common disease harbouring exons of a minority of the known AID genes are currently tested using Sanger sequencing. Next-generation sequencing (NGS) offers the ability to rapidly and cost-effectively screen all exons of gene panels containing hundreds of genes. This approach has not yet been routinely introduced in the UK for AID.

Objectives: To develop and evaluate the performance of a NGS gene panel for AID.

Methods: The Agilent SureDesign tool was used to design an NGS panel targeting 113 genes, grouped into 9 broad clinical phenotypes: AID; monogenic vasculitis/vasculopathy; complement defects; monogenic lupus; hemophagocytic lymphohistiocytosis (HLH); early-onset inflammatory bowel disease; autoimmune lymphoproliferative syndromes; monogenic stroke; and hereditary amyloidosis. The targeted region includes coding exons, conserved non-coding exons, upstream promoter regions, and splice sites. Captured and indexed libraries (QXT Target Enrichment System) were sequenced as a multiplex of 16 samples on an Illumina MiSeq sequencer in paired-end mode. Positive controls for panel validation comprised 21 DNA samples from patients with confirmed mutations in a variety of genes on the panel. We then applied the panel to test 27 prospective samples with suspected but unconfirmed monogenic inflammation. Read alignment, variant calling, and annotation were performed using 2 pipelines: the Galaxy online bioinformatics tool (https://galaxyproject.org/) and Agilent SureCall v3.0 software.

Results: In the validation stage, our targeted panel detected all known mutations in the 19 control samples with an average read depth of 244X (+/- 29X) across the captured genes. The panel was effective at detecting different types of variants, including rare and common single nucleotide variants (SNVs), insertion/deletions, splice-junction variants, upstream promoter region variants, and somatic mosaicism. The Agilent SureCall software was more sensitive and able to call mosaic mutation with allele frequency as low as 3%. Pro-mosaicism. The Agilent SureCall software was more sensitive and able to call mosaic mutation with allele frequency as low as 3%. Pro-

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mation of hospital stay between the two groups (p: 0.375) The difference in baseline characteristics between the two groups. IVIG resistance Secondary outcomes: 1. Duration of hospital stay 2. Incidence of coronary artery abnormalities

Methods: Retrospective chart review of patients admitted with a diagnosis of Kawasaki disease at 2 Canadian centres. 150 charts from centre 1 and 178 charts from centre 2 were reviewed. Inclusion criteria: 1. Definitive diagnosis of KD as per AHA guidelines. 2. Received IVIG at AHA recommended dose 3. Received ASA (3–5 mg/kg/day) at centre 1 and 80–100 mg/kg in initial phase at centre 2. Charts with incomplete information were excluded. Data from both centres was entered into the REDCAP database and exported to SPSS version 22.0 for analysis. Data analysis: Baseline characteristics were compared with descriptive statistics. Chi square test was used to compute p values for the primary outcome. Multiple logistic regression analysis was used to adjust for potential confounders. Duration of hospital stay was compared with ANOVA to adjust for confounders. p value of <0.05 was considered significant.

Results: 122/150 and 127/178 patients from centre 1 and 2 respectively were included for the final analysis. There were no significant differences in baseline characteristics between the two groups. IVIG resistance was seen in 28/122 patients in Centre 1 vs 11/127 in Centre 2 (p = 0.030). There was no significant difference in the duration of hospital stay between the two groups (p: 0.375) The differences in coronary artery abnormalities between the two groups did not reach a significant p value after adjusting for confounders (p = 0.05).

Conclusion: In this retrospective study, a lower dose of ASA led to an increased risk of IVIG resistance without a significant difference in coronary artery outcomes. The 154% increase in the use of IVIG translates to potential increase in health care costs. Given that IVIG is a scarce resource worldwide, this justifies conducting larger multicentric trials to explore cost benefit ratio of high versus low dose aspirin.

Disclosure of Interest None Declared

Table 2 (abstract O9). See text for description

| Variable names | Unadjusted | | | | | Adjusted | | | |
|----------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|-------------|
|                | ODDS | 95% CI lower | 95% CI upper | P-value | ODDS | 95% CI lower | 95% CI upper | P-value |
| Aspirin dose (HIGH vs LOW) | 0.318 | 0.151 | 0.673 | 0.003 | 0.313 | 0.110 | 0.892 | 0.030 |
| Platelet count | 1.585 | 0.889 | 2.827 | 0.119 | 1.734 | 0.889 | 3.384 | 0.107 |
| CRP value | 1.852 | 0.305 | 11.258 | 0.503 | 1.003 | 0.998 | 1.008 | 0.267 |
| Age categories (<12 MONTHS, >12 MONTHS) | 1.138 | 0.444 | 2.916 | 0.788 | 1.096 | 0.379 | 3.173 | 0.865 |
| Sodiuim <133 mmol/L | 1.368 | 0.711 | 2.631 | 0.348 | 1.752 | 0.842 | 3.646 | 0.134 |

Vasculitis

09 Optimal aspirin dose in Kawasaki disease - the ongoing battle

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Pediatric Rheumatology 2017, 15(Suppl 1):O9

Introduction: Kawasaki disease (KD) is an acute systemic vasculitis of childhood with potential cardiac morbidity in the form of coronary aneurysms. Pediatric Rheumatologists play a key role in the diagnosis and management of difficult cases of KD. Guidelines for KD have been published by American Heart Association (AHA). Standard use of 2 g/kg intravenous immunoglobulin (IVIG) is well supported by randomized trials, however, there is controversy about the best dose of acetyl salicylic acid (ASA). Consequently, there is worldwide difference in the current ASA dose used. Concerns regarding gastrointestinal side effects of ASA have prompted use of moderate doses (30-50 mg/kg) in some parts of the world, supported by evidence from meta analyses. The use of high dose ASA (80-100 mg/kg/day) in the initial phase of KD, at one Canadian Centre, as opposed to low dose ASA (3-5 mg/kg) at another centre with a similar patient population, led to this retrospective study

Objectives: To determine if low dose ASA in initial phases of KD at one Canadian centre leads to worse short term outcomes compared to higher doses of aspirin used at another centre. Primary Outcome: IVIG resistance Secondary outcomes: 1. Duration of hospital stay 2. Incidence of coronary artery abnormalities

Methods: Retrospective chart review of patients admitted with a diagnosis of Kawasaki disease at 2 Canadian centres. 150 charts from centre 1 and 178 charts from centre 2 were reviewed. Inclusion criteria: 1. Definitive diagnosis of KD as per AHA guidelines. 2. Received IVIG at AHA recommended dose. 3. Received ASA (3–5 mg/kg/day) at centre 1 and 80–100 mg/kg in initial phase at centre 2. Charts with incomplete information were excluded. Data from both centres was entered into the REDCAP database and exported to SPSS version 22.0 for analysis. Data analysis: Baseline characteristics were compared with descriptive statistics. Chi square test was used to compute p values for the primary outcome. Multiple logistic regression analysis was used to adjust for potential confounders. Duration of hospital stay was compared with ANOVA to adjust for confounders. p value of <0.05 was considered significant.

Results: 122/150 and 127/178 patients from centre 1 and 2 respectively were included for the final analysis. There were no significant differences in baseline characteristics between the two groups. IVIG resistance was seen in 28/122 patients in Centre 1 vs 11/127 in Centre 2 (p = 0.030). There was no significant difference in the duration of hospital stay between the two groups (p: 0.375) The differences in coronary artery abnormalities between the two groups did not reach a significant p value after adjusting for confounders (p = 0.05).

Conclusion: In this retrospective study, a lower dose of ASA led to an increased risk of IVIG resistance without a significant difference in coronary artery outcomes. The 154% increase in the use of IVIG translates to potential increase in health care costs. Given that IVIG is a scarce resource worldwide, this justifies conducting larger multicentric trials to explore cost benefit ratio of high versus low dose aspirin.

Disclosure of Interest None Declared
### O10

**Childhood Takayasu Arteritis: disease course and response to therapy**

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**Pediatric Rheumatology** 2017, 15(Suppl 1):O10

**Introduction:** Takayasu Arteritis (TA) is a devastating large vessel vasculitis commonly associated with systemic inflammation and resulting in severe ischemic end organ dysfunction. To date, treatment efficacy, disease course and flare risk in children with TA are widely unknown.

**Objectives:** To describe clinical presentation, treatment regimens and outcome of childhood TA and to identify risk factors for adverse outcome.

**Methods:** A single-center cohort study of consecutive children fulfilling the EULAR/PRINTO/PReS criteria for childhood TA between 1986 and 2015 was performed. Clinical, laboratory and imaging features, treatment, disease course and outcome were documented. Disease activity was assessed by the Pediatric Vasculitis Disease Activity Score (PVAS), damage by the Pediatric Vasculitis Damage Index (PVDI). Active disease was defined as PVAS ≥ 1 and/or increased inflammatory markers not explained by other causes and/or active disease on imaging (new lesions and/or evidence of vessel wall inflammation). Inactive disease was defined as PVAS = 0 and normal inflammatory markers or inactive disease on imaging. Adverse outcome was defined as disease flare (new TA symptoms and/or increased inflammatory markers necessitating therapy escalation and/or new angiographic lesions) or death.

**Analyses:** Kaplan-Meier statistics compared treatment regimens. Logistic regression analyses tested putative predictors for adverse outcome.

**Results:** Twenty-seven children (74% female) were included; median age at diagnosis was 12.4 years (range 4.2–17.7); median follow-up duration was 2.1 years. The most common presenting symptoms were malaise (48%), headaches (33%), weight loss (30%) and claudication of the extremities (22%), the most common clinical findings blood pressure discrepancy between limbs (67%), decreased/absent pulse (59%), hypertension and bruits (56%). Median ESR was 35 mm/h (range 1–67%), decreased/absent pulse (59%), hypertension and bruits (56%). Median follow-up duration was 2.1 years. The most common presenting symptoms were malaise (48%), headaches (33%), weight loss (30%) and claudication of the extremities (22%), the most common clinical findings blood pressure discrepancy between limbs (67%), decreased/absent pulse (59%), hypertension and bruits (56%). Median ESR was 35 mm/h (range 1–109), CRP 31.9 mg/dl (range 0.5–140). Twenty-two (81%) children presented with active disease at diagnosis and received immunosuppressive treatment (Table 3). Adverse outcome was seen in 14/27 (52%) children: 13 (48%) experienced flares after a median of 9.2 months after diagnosis and 2 (7%) died within months of diagnosis. The two-year flare-free survival was 80% with biologic therapies compared to 43% in conventional therapies (p = 0.03). At last follow-up, biologic therapies resulted in significantly higher rates of inactive disease (p = 0.02). None of the other tested variables were associated with an increased flare risk.

**Conclusion:** Childhood TA carries a dramatic burden: half of the children experienced flares and 7% died. Biologic treatment regimens were associated with significantly better control of disease activity.

**Disclosure of Interest** None Declared

| Induction therapy | No. of patients (focal N = 27) | Median PVAS at diagnosis (range) | Inactive disease at 6 months | Therapy for maintenance of remission during disease course | No. of patients with flares | Death | Inactive disease at last follow-up |
|-------------------|-------------------------------|---------------------------------|-------------------------------|----------------------------------------------------------|----------------------------|-------|----------------------------------|
| CS only           | 4 (15)                        | 16 (9–15)                       | 1/5                           | CS, Cyclo, Aza, MMF, MTX                                   | 3/4                       | 1/5   | 1/3                              |
| CS + MTX          | 10 (37)                       | 15.5 (8–21)                     | 2/10                          | CS, Cyclo, Aza, Lef, MTX, Biologics                       | 7/10                      | 0/10  | 5/10                            |
| CS + Cyclo        | 5 (19)                        | 7 (6–35)                        | 2/4                           | CS, Cyclo, Aza, Lef, MTX, Biologics                       | 3/5                       | 1/5   | 3/4                              |
| CS + MTX + Biologic | 3 (11)                      | 10 (6–25)                       | 3/3                           | CS, Biologic = Lef + MTX                                  | 0/3                       | 0/3   | 3/3                              |
| No treatment      | 5 (19)                        | 10 (6–17)                       | 5/5                           | None                                                      | 0/5                       | 0/5   | 5/5                              |

Aza azathioprine, CS corticosteroids, Cyclo cyclophosphamide, Lef leflunomide, MTX methotrexate, MMF mycophenolate mofetil

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**Immune aspects of JIA**

### O11

**Dysbalanced serum factor environments can drive IL-17 over-expression by gamma delta T cells in systemic juvenile idiopathic arthritis**

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**Pediatric Rheumatology** 2017, 15(Suppl 1):O11

**Introduction:** Systemic-onset juvenile idiopathic arthritis (sJIA) is considered to follow a biphasic course with an initial systemic disease phase driven by innate immune mechanisms and interleukin (IL)-1β as a key cytokine, while a second chronic arthritic phase may be dominated by adaptive immunity and cytokines such as IL-17A.

**Objectives:** A recent sJIA mouse model points to a critical implication of IL-17A expressing γδ T cells (Tyβ17) in disease pathology [1]. In this model, IL-17 expressing γδ T cells drive a phenotype that resembles arthritis-prominent sJIA in an environment of low interferon (IFN)γ. However, human data on the role of IL-17 expressing γδ T cells in sJIA are lacking.

**Methods:** Paired serum samples of sJIA patients (n = 12) during active and inactive disease as well as sera from healthy children (n = 10) were analyzed for expression of IL-1β, IL-6, IL-17A, IL-23, IFNγ and S100A12 by bead array assay. Paired serum samples of sJIA patients (n = 12) during active and inactive disease as well as sera from healthy children (n = 10) were analyzed for expression of IL-1β, IL-6, IL-17A, IL-23, IFNγ and S100A12 by bead array assay. Ex vivo T cellular cytokine expression was assayed in whole peripheral blood samples collected from active and inactive sJIA (n = 12) as well as healthy children (HC, n = 13). Cells in whole peripheral blood were stimulated with phorbol 12-myristate 13-acetate (PMA) and ionomycin and were stained for expression of CD3, CD4, γδ TCR, IL-17A and IFNγ before analysis by flow cytometry. Finally, CD4+ and γδ TCR+ T cells were isolated from sJIA patients’ as well as healthy control PBMCs (both n = 5) and co-cultured with healthy control monocytes under different stimulating conditions. IL-17A and IFNγ expression was quantified.

**Results:** Particularly in active sJIA patients’ serum IL-17A expression was significantly increased. Ex vivo T cellular cytokine expression analysis revealed sJIA γδ T cells to over-express IL-17A compared to cells in whole blood obtained from healthy donors. This was not observed...
with CD4+ T cells. During active disease peripheral Tγδ17 cell frequencies were decreased and appeared to recover in patients with no clinical disease activity. IL-17A over-expression by sJIA γδT cells was confirmed in in vitro studies on isolated cell level and found to occur particularly upon stimulation of monocyteγδT cell co-cultures with TLR4 ligands such as LPS or S100A12. On the contrary to γδT cellular IL-17A over-expression, IFNγ expression by sJIA CD3+CD4+ cells was strikingly low, both in ex vivo analysis of patients’ whole blood as well as in vitro T cell cultures. Finally, in our assays particularly therapeutic IL-1 blockade appeared to expand peripheral IFNγ+CD4+ cells and restore intracellular IFNγ levels, while γδT cellular IL-17A expression and release normalized to that of healthy donors.

Conclusion: An sJIA serum factor environment with elevated levels of endogenous TLR4 ligands such as S100 proteins and increased expression of cytokines like IL-1β can result in IL-17 over-expression by sJIA γδT cells. As already suggested from the sJIA mouse model, this may be promoted by impaired IFNγ expression as we observed with CD4+ T cells. During active disease Tγδ17 cells may migrate from the periphery to inflamed tissues to drive local inflammation and promote downstream differentiation and expansion of Th17 cells. As a cell-type bridging innate and adaptive immunity, γδT cells may thus play a key pathological role in tilting systemic to arthritis-prone disease in men and mice.

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Disclosure of Interest

None Declared

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**O12**

The composition of the gut microbiota differs between children with JIA and healthy controls

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Pediatric Rheumatology 2017, 15(Suppl 1):O12

Introduction: The human body is home to a myriad of non-pathogenic bacteria, the so-called microbiota. It has been hypothesised that the gut microbiota plays a role in the pathogenesis of autoimmune diseases, such as RA. Its role in the pathogenesis of JIA needs further investigation.

Objectives: To study differences in the composition of the gut microbiota in patients with JIA compared to healthy controls.

Methods: A prospective follow up study was initiated in October 2013, in which untreated patients were enrolled with a new onset of JIA according to ILAR criteria and a duration of symptoms of at most 6 months. Patients were followed for 2 years. At baseline, and if ever a patient presented with inactive disease according to Wallace criteria or a disease flare, a faecal sample was collected and stored at -80 °C. Operational taxonomic units were pyrosequenced targeting 16S ribosomal RNA. Additionally, faecal samples were collected from age-, gender- and geography-matched healthy controls and processed as above.

The microbial composition of the faecal samples was analysed using compositional data analysis. The relative abundance was visualised using ternary plots. Differences between healthy controls and patients, as well as between Italian and Dutch patients were tested using MANOVA. Paired baseline, inactive disease and flare samples were analysed using Hotelling’s T2 paired sample test. Association with various clinical parameters was tested using MANOVA. Finally, a random forest algorithm was used to distinguish Italian patients and healthy controls based on their microbiota composition.

Results: Overall, 307 samples were available for analysis. Of these, 100 belonged to patients at baseline (72 Italian samples and 28 Dutch samples). These were matched 1:1 with 107 healthy control samples. Moreover, 100 samples were obtained during follow-up: 70 in inactive disease and 30 in persistent activity or disease flare.

There was strong evidence the composition of the microbiota at the phyla level was different in patients at baseline with respect to healthy controls, when analysing Italian and Dutch patients both separately and jointly, taking into account the different age at onset of Italian and Dutch patients (p ≤ 0.003). There was no evidence of differences between patients at baseline and in inactive disease, or between patients at baseline and in persistent activity (p 0.048). There was evidence that the composition of the microbiota at the phyla level was different in ANA positive patients with respect to ANA negative patients, even when taking the nationality of patients into account (p = 0.02). There was no evidence for an association with uveitis at baseline (p = 0.18) and only weak evidence for an association with the number of active joints (p = 0.07).

Visualisation of the results, as well as compositional Principal Components Analysis showed that the relative abundance of Proteobacteria and Tenericutes contributed the most to the differences between the groups in all comparisons. The random forest algorithm separated patients and healthy controls well, using only the microbiota composition at the species level. The AUC of the model was 0.99 (95% CI 0.98–0.99).

Conclusion: This study provides strong evidence for differences in the composition of the microbiota, and especially the relative abundance of Proteobacteria and Tenericutes, between patients with JIA and healthy controls. Future research should be directed to elucidate the effects of the microbiota on JIA.

Disclosure of Interest

None Declared

**Vasculitis and Behcet disease**

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**O13**

A new approach to assessment of juvenile vasculitis: the juvenile vasculitis multidimensional assessment report (J-VAMAR)

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Pediatric Rheumatology 2017, 15(Suppl 1):O13
Introduction: There are many effects of vasculitis (fatigue, reduced energy level, school absenteeism, physical appearance) that were quite important to patients but not measured with the outcome instruments currently included in clinical trials of vasculitis.

Objectives: To develop and test a new multidimensional questionnaire for assessment of children with juvenile vasculitis in standard clinical care.

Methods: The Juvenile Vasculitis Multidimensional Assessment Report (J-VAMAR) includes 15 parent or patient-centered measures or items that assess well-being, pain, functional status, health-related quality of life, problems about school, disease activity, disease status and course, symptoms of vasculitis, use of medications, side effects of medications, therapeutic compliance. The J-VAMAR is proposed for use as both a proxy-report and a patient self-report, with the suggested age range of 7-18 years for use as a self-report. The Outcome Measures in Rheumatology Clinical Trials (OMERACT) filter for outcome measures in rheumatology was applied to validate the J-VAMAR.

Results: The analysis data set was collected between 2013 to 2016, the questionnaire was completed by 164 children with vasculitis in 429 visits. The diagnoses of the patients were as follows: 46.6% HSP, 19% Behçet vasculitis, 9.2% CPAN, 9.2% Takayasu, 6.6% Kawasaki, 5.5% Wegener, 3.2% Unclassified and 0.5% Churg-Strauss. The mean disease onset age of children with vasculitis was 8.36 ± 4.61. The J-VAMAR was found to be feasible and to possess face, content, criterion and construct validity. All parents and children reported that the questionnaire was simple and easy to understand. Completion and scoring appeared to be quick, requiring less than 15 minutes. The Cronbach’s alpha coefficient for internal consistency for the J-VAMAR dimensions was between 0.735–0.963. Between the test-retest scale scores, there is a significant and a positive correlation (0.326–1.000). When the criterion validity is considered, we would say that the correlation level between the each subscale and the related scale spanned from medium (r = 0.282, p < 0.05) to large (r = 0.694, p < 0.001). Parents’ proxy-reported and children’s self-reported data were remarkably concordant.

Conclusion: The J-VAMAR provided thorough information for the study patients about recent medical history and current health status. It performed similarly across different children’s ages and characterized the level of disease activity and disability well. The development of the J-VAMAR introduces a new and a multidimensional approach in pediatric rheumatology practice. This new questionnaire is a valid tool may help enhance the quality of care of children with vasculitis.

Disclosure of Interest: None Declared

O14
Behçet’s disease in children: Eastern Mediterranean experience
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Pediatric Rheumatology 2017, 15(Suppl 1):O14

Introduction: Behçet’s disease (BD) is a variable vessel vasculitis characterized by recurrent oral and/or genital aphthous ulcers accompanied by cutaneous, ocular, articular, vascular, gastrointestinal and/or central nervous system inflammatory lesions. The most widely used diagnostic criteria for adult onset disease is the International Behçet’s Study Group (ISG) criteria. An international expert consensus group (the pediatric BD [PEDBD] group) has recently proposed a new set of criteria for the classification of Behçet’s disease (BD) in children. To evaluate the disease activity in the follow-up; there are mainly two severity scores Behçet’s disease dynamic activity measure (IBDDAM) and Behçet’s disease current activity form (BDCAF).

Objectives: Our aim was to test the performance of the PEDBD criteria compared to the ISG criteria. Our second aim was to check the correlation between the severity score systems and physician global assessment (PGA) in pediatric BD patients.

Methods: Two centers from Turkey and one center from Israel participated in this study. The disease onset was ≤16 years of age. As controls, pediatric patients with other diseases such as primary vasculitides, periodic fever syndrome, or systemic lupus erythematosus were included.

Results: Sixty eight BD (44.1% male) patients and 93 control patients were included. The median age at symptom onset was 132 (14–191) months and the median time to diagnosis from symptom onset was 12 (0–108) months. Acneiform lesions were significantly associated with female gender (18.4% in females versus 0% in males; p = 0.015). The sensitivity and specificity of the PEDBD and the ISG criteria and revised ICBD criteria were 73.5%/52.9% and 98.9%/100% respectively. Thirty two (47%) patients with BD failed to fulfill the ISG BD criteria. However, almost all of these patients met the PEDBD criteria. For systemic therapy, 62 (91.2%) patients received oral colchicine, 15 (22.1%) prednisolone, 16 (23.5%) azathioprine, 6 (8.8%) methotrexate, 5 (7.4%) anti-tumor necrosis factor alpha treatment, and one patient received cyclophosphamide. The median (min-max) IBDDAM and BDCAF scores at diagnosis were 6 (1–23) and 4 (1–7) and significantly decreased to 1 (0–8) and 1 (0–4) respectively at latest follow-up (p < 0.001 for both). The median (min-max) physician global assessment score at diagnosis was 5 (2–9) and significantly decreased to 1 (0–7) at latest follow-up (p < 0.001). IBDDAM score had a strong positive correlation with BDCAF score (r = 0.637; p < 0.001). PGA positively correlated with BDCAF and IBDDAM scores (r = 0.502; p < 0.001 and r = 0.624; p < 0.001, respectively).

Conclusion: In our pediatric series, the PEDBD criteria showed better sensitivity (73.5%) than the ISG criteria and the specificity was close. The higher sensitivity of the PEDBD criteria set is a big advantage for pediatric patients since early diagnosis and timely treatment is very important. Currently, BD patients are treated according to the physician’s judgement of disease activity. Our study demonstrated that the severity scores were positively correlated with each other and PGA and thus may be used in clinical practice.

Disclosure of Interest: None Declared

O15
Macrophage activation syndrome

Immune abnormalities leading to exaggerated production of IFN-gamma and the therapeutic response to an anti-IFN-gamma antibody in a patient with NLRC4 mediated disease
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Pediatric Rheumatology 2017, 15(Suppl 1):O15

Introduction: Animal and human data suggest that IFNγ plays a pathogenic role in HLH. A phase 2 trial with the anti-IFNγ monoclonal
antibody Ni-0501 in primary HLH provides encouraging preliminary data [1]. Gain-of-function mutations in NLR4 are associated with a distinct autoinflammatory syndrome, with recurrent HLH, fever episodes, and enterocolitis [2].

**Objectives:** To report a patient with severe early onset progressive HLH carrying a de novo missense mutation in NLR4 (T337N), the immune abnormalities leading to abnormal production of IFNγ and the response to treatment with Ni-0501.

**Methods:** We evaluated cytokine levels by multiplex assay and by specific ELISAs and expression of IFNγ in freshly isolated PBMCs by cytometry.

**Results:** LR, caucasian male, presented, at 20 days of age, fever and rash and progressively developed clinical and laboratory features of HLH leading rapidly to liver failure and subsequent multi-organ failure. Genes causing primary-HLH and functional tests were negative. High-dose glucocorticoids and cyclosporine-A led to partial improvement. Development of sepsis triggered HLH reactivation. Measurable IFNγ levels (6 pg/ml) and high levels of the IFNγ inducible chemokines CXCL9 (5670 pg/ml) and CXCL10 (4400 pg/ml) were found, the latter demonstrating activation of the IFNγ pathway. Ni-0501 was started (compassionate use) on background of dexamethasone (13.6 mg/m2) and cyclosporine-A. After 3 months, the child was discharged in excellent conditions (prednisone 0.3 mg/kg). Markedly elevated production of IFNγ was revealed by the administration of Ni-0501 through measurement of total IFNγ bound to circulating Ni-0501. This IFNγ was fully neutralized, as shown by rapidly undetectable levels of CXCL9 and CXCL10. The percentage of CD4+, CD8+ and CD56+ cells expressing IFNγ was significantly increased in the NLR4 patient without (0.95%, 0.88%, 0.80%, respectively) and with stimulation with PMA/ionomycin (6.44%, 10.20%, 25.00%). Serum levels of IL-18 were markedly higher (as expected in NLR4-mediated activation) with other autoinflammatory diseases (0.59 ± 0.47 ng/ml). This increase in IFNγ production was neutralized by incubation with an anti-IL18 receptor antibody. After 7 months of Ni-0501 treatment, all therapies, including glucocorticoid and cyclosporine-A. No safety concern emerged.

**Conclusion:** Our data suggest that in NLR4-related disease overproduction of IL-18, induced by a hyperfunctional NLR4 inflammasome, may be one of the contributors to the up-regulation of IFNγ production that appears to be driving HLH. In this patient, neutralization of IFNγ allowed control of all disease features, enabling withdrawal of all treatments, including glucocorticoid and cyclosporine-A. No safety concern emerged.

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**Disclosure of Interest**
C. Bracaglia: None Declared, G. Prencipe: None Declared, M. Pardeo: None Declared, G. Lapeyre Employee of: Novimmune, E. Haddad 2, E. Nasonov 1, M. Trachana 1, O. Vougiouka 1, K. Leon, E. Vritzali, K. Lheritier, A. Martinis, D. Lovell, and PRINTO/PRTC
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**Pediatric Rheumatology** 2017, 15(Suppl 1):O16

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**O16**

**Long-term efficacy and safety of Canakinumab in patients with active systemic juvenile idiopathic arthritis (SJIA): results from a phase III extension study**

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**Pediatric Rheumatology** 2017, 15(Suppl 1):O16

**Introduction:** The management of SJIA with biological therapies is aimed to achieve and maintain clinical remission (CR), and accordingly taper corticosteroids (CS). Canakinumab (CAN) demonstrated high inactive disease (ID) rates in about 33% of patients (pts) in Day 15 and 30 respectively in previous studies [1]. However, little is known about high level response rates in SJIA pts using CAN long-term.

**Objectives:** To evaluate the long-term treatment response in terms of safety and efficacy in CAN treated pts with active SJIA.

**Methods:** This was an open-label, non-comparative study of CAN-naive SJIA pts (≥2 to <20 years) receiving subcutaneous CAN 4 mg/kg every 4 weeks. Efficacy was assessed every 3 months by the aACR 30/50/70/90/100 responses compared to baseline (BL), ID or CR (ID for >6 months) and changes in JADAS10-CRP scores over time. Safety was assessed by adverse events (AEs) and serious AEs (SAEs). The results are based on the observed data with imputations to carry the last observation forward.

**Results:** Of 123 pts with active SJIA, 70 (57%) had fever and 71 (57.7%) used corticosteroids at BL. Mean CRP was 117.8 mg/L (normal: 0-10 mg/L), and, on average, pts had 9.9 active joints and 8.9 joints with limited motion. A rapid response was observed at Day 15: 59 (51%) and 27 (26%) pts had aACR >70 and aACR 100 responses, respectively. These responses were maintained at subsequent time points (Table). At Month 6, CR was achieved in 52 (42.3%) pts. Overall, 33 (26.8%) pts had CR for at least 12 months. At BL, the median CRP score was 22.3, with median changes from BL of –12.0 at Day 15 and –16.8 at last assessment, respectively. At the last assessment, 59 (48.4%) pts had ID (JADAS10≤11); 14 (11.5%) had low disease activity (JADAS10>1 and ≤3.8), while 14 (11.5%) had moderate and 35 (28.7%) had high disease activity. Overall, 24 (33.8%) pts were steroid-free at last assessment. In total, 108 (87.8%) pts had at least 1 AE. Overall, exposure adjusted AE and SAE rate was 8.22 and 5.48 events/pt-years (pyr) respectively, with 183.56 pyr exposure and 40 (32.5%) pts had SAEs; most commonly reported SAEs were disease flares or worsening of SJIA in 13 (10.6%) pts; macrophage activation syndrome in 6 (4.9%) pts, and fever in 4 (3.3%) pts. No deaths occurred in this study.

**Conclusion:** CAN treatment was associated with rapid response and sustained therapeutic effect over the long-term in the pts with active SJIA. The safety profile is consistent with other CAN studies.

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**Trial registration identifying number:** NCT00891046

**Disclosure of Interest**
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Consultant for: Novartis, Roche, Pfizer, UCB, Celgene, Regeneron, Amgen, Astrazeneca, GSK and BMS, Speaker Bureau of: Novartis and Roche, P. Quartiier Grant / Research Support from: Abbvie, BMS, Chugai-Roche, Novartis, Pfizer, Consultant for: Abbvie, Novartis, Servier and SOBI, Speaker Bureau of: Abbvie, BMS, Chugai-Roche, Medimmune, Novartis, Pfizer and SOBI, T. Constantin: None Declared, E. Alexeeva Grant / Research Support from: Roche, Abbott, Novartis, Pfizer, Bristol-Myers Squibb and Centocor, Speaker Bureau of: Roche, Novartis, Merck Sharp Dohme, Bristol-Myers Squibb, Medac and Pfizer, I. Kone-Paut Grant / Research Support from: SOBI, Novartis and Roche, Consultant for: Novartis, SOBI, Pfizer, Abbott and Roche/Chugai, K. Marzan Grant / Research Support from: Novartis, Abbvie, N. Wuifnaat Grant / Research Support from: Novartis, R. Schneider: None Declared, S. Padhe: None Declared, V. Chasnuk: None Declared, C. Wouters Grant / Research Support from: GSK, Novartis and Roche, J. Kuemmel-Deschiner Grant / Research Support from: Novartis, Consultant for: Novartis, SOBI and Baxalta, T. Kallinich Grant / Research Support from: Novartis, Speaker Bureau of: Sobi, Novartis, BMS and Roche, B. Lauwerys: None Declared, E. Haddad: None Declared, E. Nasonov: None Declared, M. Deschner Grant / Research Support from: Novartis, Consultant for: Novartis, SOBI, Grant / Research Support from: Novartis, Abbvie, Amgen, Biogen and Novartis, Consultant for: Genentech, Glaxo Smith Kline, Boehringer Ingelheim, Celgene and Bristol Meyers Squibb, Abbvie, Medimmune, Novartis, Pfizer, Sanofi, Roche, Servier and Takeda, D. Lovell Grant / Research Support from: National Institutes of Health, Consultant for: Hoffman-La Roche, Italfarmaco, Janssen, Novartis, Pfizer, Sanofi, Aventis, Schwarz Biosciences, Sobi, Xoma and Wyeth, Speaker Bureau of: Abbott, Abbvie, Amgen, Biogen and Novartis, Consultant for: Genentech, Glaxo Smith Kline, Boehringer Ingelheim, Celgene and Janssen, Speaker Bureau of: Genentech, Roche and Novartis

Table 4 (abstract O16). ACR responses achieved in the cohort by time point

| Time point | CAN N = 123 |
|------------|-------------|
|            | Minimum adapted ACR pediatric response | n (n/m) (%) | Patients with inactive disease (n/m) (%) |
| Month 12   | m            | 85 (52/88) | 59.1 |
| Non-Responders | aACR ≥30 | 4 (4.7) | 81 (95.3) |
|             | aACR ≥50 | 77 (90.6) |
|             | aACR ≥70 | 78 (85.9) |
|             | aACR 100 | 49 (57.6) |
| Month 21   | m            | 65 (48/65) | 73.8 |
| Non-Responders | aACR ≥30 | 3 (4.6) | 62 (95.4) |
|             | aACR ≥50 | 58 (89.2) |
|             | aACR ≥70 | 54 (83.1) |
|             | aACR 100 | 39 (60.0) |
| Last assessment | m | 121 (62/122) | 50.8 |
| Non-Responders | aACR ≥30 | 28 (23.1) | 93 (76.9) |
|             | aACR ≥50 | 89 (73.6) |
|             | aACR ≥70 | 81 (66.9) |
|             | aACR 100 | 62 (51.2) |

n number of patients who satisfy the criteria, m number of patients with an assessment in the time period
O18
Previously undescribed SAVI-associated mutations identify a novel region in human sting necessary for the control of type I interferon signalling

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Presenting author: Isabelle Melki

Pediatric Rheumatology 2017, 15(Suppl 1):O18

Introduction: Gain-of-function mutations in TMEM173 encoding STING (stimulator of interferon genes) underlie a very recently described type I interferonopathy referred to as SAVI (STING Associated Vasculopathy in Infancy). Thirteen patients belonging to ten different families have been published so far, all carrying heterozygous substitutions involving one of three residues at positions 147, 154 and 155, located in a small linker region connecting the N-terminal transmembrane domain of STING to the C-terminal cyclic dinucleotide-binding domain. It has been shown that the constitutive activation driven by these mutants is mediated by exit of STING from the endoplasmic reticulum (ER) and stable sequestration within the ER-Golgi intermediate compartment (ERGIC/Golgi compartment). This triggers the downstream protein kinase TBK1, which then phosphorylates IF3 and induces the transcription of interferon stimulated genes (ISGs).

Objectives: To report three new individuals variably exhibiting the core features of SAVI including systemic inflammation, destructive skin lesions and interstitial lung disease.

Methods: One patient was studied by exome sequencing and all three by Sanger sequencing of the candidate gene TMEM173. Interferon alpha titres were assessed by Single Molecule Array (Simoa®) technology and ISGs were measured by qPCR on PBMCs. Functional studies were performed using site-directed mutagenesis, studying the response to cGAMP in a HEK 293 T transfection system. TBK1, IRF3 and STAT1 phosphorylation were also studied.

Results: Although all three patients demonstrated characteristic features of SAVI, in two individuals the disease was essentially limited to the skin. Furthermore, none of the three experienced recurrent fevers, and bacterial infections were seen, thus perhaps broadening the phenotypic spectrum.

All three patients demonstrated high interferon alpha titres in serum and / or increased expression of ISGs. Molecular and in vitro data revealed that the pathology in these patients is due to missense substitutions at positions 206, 281 and 284 of the human STING protein. These mutations confer cGAMP independent constitutive activation of type I interferon signalling through TBK1, and independent from the alternative STING pathway triggered by membrane fusion of enveloped RNA viruses. Structural analysis indicates that these three amino acids lie in a discrete region of the protein thereby providing a physical explanation for the human genetic findings, and implicating a novel cluster of amino acids in STING as functionally important in the regulation of type I interferon signalling.

Conclusion: Our data clearly implicate a novel region of STING as important in type I interferon signalling, leading us to speculate that the surface-exposed arginine residues at 281 and 284 may be involved in ER retention or in binding to a negative regulator of STING signalling. Thus, the findings presented here suggest a previously unappreciated aspect of the control of STING.

Disclosure of Interest
None Declared

O19
Efficacy of the Janus kinase 1/2 inhibitor ruxolitinib in the treatment of vasculopathy associated with TMEM173-activating mutations in three children

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Presenting author: Marie-Louise Frémond

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Introduction: Gain-of-function mutations in TMEM173 encoding STING underlie a novel type I interferonopathy, minimally responsive to conventional immunosuppressive therapies and associated with high childhood morbidity and mortality.

Objectives: We hypothesized that inhibition of interferon (IFN) signalling could be used as a therapeutic approach in this disease. We describe here the medical and molecular consequences of treatment with ruxolitinib, an oral JAK1/2 inhibitor.

Methods: During follow-up of three children over a period of between 12 to 24 months, clinical data were collected and disease activity scores were determined for systemic, skin and lung manifestations. The effect of ruxolitinib on JAK-STAT signaling was assessed in vitro and ex vivo by transcriptomic analysis and STAT1 phosphorylation status.

Results: We observed marked clinical improvement in all three children in each of the central aspects of the phenotype – namely, destructive skin lesions, interstitial lung disease and systemic inflammation. This clinical effect was mirrored by in vitro and ex vivo data, where we recorded decreased STAT1 phosphorylation and reduced expression of both IFN-stimulated genes and a subset of genes associated with fever and vasculopathy. Tolerance of treatment was good without an obvious infectious risk.

Conclusion: These results suggest that JAK1 inhibition represents a promising therapeutic approach in TMEM173-mutated patients and may be also relevant to the treatment of other type I interferonopathies.

Disclosure of Interest
None Declared
O20 Development of the autoinflammatory disease damage index (ADDI)
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Introduction: Autoinflammatory diseases cause systemic inflammation which can result in damage to multiple organs. A validated instrument to measure damage is essential to quantify damage in individual patients, and to compare disease outcomes in clinical studies. At this moment, there is no instrument to qualify damage in autoimmune inflammatory diseases.

Objectives: Our objective was to develop a common autoinflammatory disease damage index (ADDI) for Familial Mediterranean Fever (FMF), Cryopyrin Associated Periodic Syndromes (CAPS), Tumor Necrosis Factor Receptor Associated Periodic Syndrome (TRAPS) and Mevalonate Kinase Deficiency (MKD).

Methods: We have developed the ADDI by consensus building based on the Delphi method. The top 40 enrollers of patients in the Eurofever registry and nine important experts from the Americas participated in multiple rounds of online surveys to suggest and select items and definitions. Patients fulfilled a role in this project in order to develop a damage index that represents what is important for patients. Further, 22 patients and parents of patients rated damage items and suggested new items.

Lastly, we decided the scoring of each item based on the results of a decision making survey (1000minds program), which was completed by experts and (parents) of patients.

Results: More than 80% of the 49 experts and 22 patients completed the online surveys. At the consensus meeting, 31 experts and a patient representative reached consensus on the inclusion and definition of the items. The scoring of each item was based on the 1000minds program, which was completed by 37 experts and fourteen patients. The preliminary ADDI contains eighteen relevant damage items, categorised per organ system. The damage items are considered relevant by the experts and all cause an important burden for patients according to the experts and (parents) of patients. Examples of damage items of the preliminary ADDI are hearing loss, amyloidosis, infertility and serosal scarring. It will take approximately five to ten minutes to complete the ADDI.

Conclusion: An instrument to measure damage caused by autoinflammatory diseases is developed based on consensus building with multiple rounds of online surveys and a consensus meeting. Patients fulfilled a significant role in this process.

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O21 The phenotype and genotype of mevalonate kinase deficiency: a series of 114 cases from the Eurofever registry
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Pediatric Rheumatology 2017, 15(Suppl 1):O21

This abstract is not included here as it has already been published.

O22 Autoinflammatory diseases: role of next-generation sequencing
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Presenting author: Chiara Passarelli
Pediatric Rheumatology 2017, 15(Suppl 1):O22

Introduction: Autoinflammatory disorders (AIDs) are an expanding group of complex diseases marked by periodic or chronic systemic inflammation. Mutations in approximately 25 genes have been associated with autoinflammatory disorders (AIDs). Genetic analysis of based on the candidate gene has a low efficiency and it is expensive and time consuming. Next Generation Sequencing (NGS) has emerged as a new diagnostic tool in this field, with the goal of allowing for faster genetic diagnosis and of providing a better comprehension of the phenotype/genotype correlation in AID patients.

Objectives: To evaluate the results obtained by the application of NGS in a cohort of patients affected by undefined AID evaluated at our center.

Methods: In this study we enrolled 242 patients evaluated at our center from 2010 to march 2016 affected by undefined AIDs. In the first 4 years of the study, we performed Targeted resequencing on 179 patients using a panel including 11 AIDs related genes (MVK, MEFV, NLRP12, NLRP3, NOD2, TNFRSF1A, PSTPIP1, IL1RN, LPIN2, IL36RN, PSM8). The customized panel was subsequently implemented, adding 14 genes more recently demonstrated to be involved in AIDs, used to analyse 63 additional patients. Sequencing analysis were performed on MiSeq® sequencing platform (Illumina, San Diego, CA) and all variants identified have been confirmed by Sanger sequencing. Validated variants were studied by in silico analysis using SIFT and PolyPhen software in order to predict their possible functional impact.

Results: NGS analysis led to the identification of 80 patients with different variants in the typically AIDs-associated genes: MVK, MEFV, NLRP12, NLRP3, NOD2, TNFRSF1A, PSTPIP1, with a detection rate of 31%. We took into account only variants with a frequency in the
global population <1%, as reported in dbSNP and Ensembl databases. 25% of patients showed monogenic variants, both novel or already known, while variants in two or more genes were found in 5% of patients. In details, 33% of total variants was found in NOD2 gene, 19% in MEFV, 16% in NLPR12, 12% in NLPR3, 5% in P53 and 5% in both TNFRSF1A and MVK. Several of these variants do not have a clear functional significance yet, while some of them are described as risk factors. We performed 19 familial study to unravel the segregation of some variants. 

**Conclusion:** NGS allowed the identification of a large amount of genetic variants in different genes. The major challenge is represented by the interpretation of the clinical relevance of the identified variants, particularly of variants that are found at low, but >1% frequency in various populations. These may indeed function as susceptibility alleles to inflammation rather than disease associated mutations.

Some patients show variants in different genes that may cooperate to cause a clinical phenotype or, on the other hand, may be irrelevant. Therefore large-scale population studies, in vitro functional assays and careful correlation between genetic information and phenotypic data are needed. Cooperations with clinicians as well as collaborations with different laboratories using NGS is crucial in order to generate information that are scientifically rigorous and relevant for patient management. 

**Disclosure of Interest**

C. Passarelli: None Declared, E. Pisanesci: None Declared, V. Messia: None Declared, M. Pardeo: None Declared, A. Novelli: None Declared, C. Passarelli: None Declared, E. Pisaneschi: None Declared, V. Messia: None Declared.

**O23** Effectiveness of childhood vaccinations in CAPS patients treated with Canakinumab: results from an open-label phase 3 extension study

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**Presenting author:** P.A. Brogan

**Pediatric Rheumatology 2017, 15(Suppl 1):O23**

**Introduction:** Canakinumab (CAN) has been shown not to impair antibody production following vaccination in children in an open-label phase 3 study (NCT01302860) [1]. Here we present the results from the extension of this study.

**Objectives:** To evaluate the presence of protective antibody levels following immunisation with inactivated vaccines in CAPS patients during extension study.

**Methods:** Patients who completed the core study were allowed to continue into the extension study on the standard dosing regimen of 2 mg/kg subcutaneous CAN every 8 weeks or on last dose/dosing regimen received in the core study. Vaccination response was evaluated using post-vaccination antibody titres at 4 and 8 weeks after immunisation. Patients were considered assessable for an antibody response to a specific vaccination if they had a measurement of antibody titre 0-14 days post-vaccination (pre-vaccination assessment) and at least 1 subsequent measurement of antibody titre at 4 weeks and/or 8 weeks post-vaccination. However, for patients with adequate pre-dose antibody titres and maintained during the trial, the specific patient vaccination was deemed non-assessable.

**Results:** During the extension phase, of 17 patients (≤5 years), 4 received 8 types of vaccinations against Corynebacterium diphtheria, Bordetella pertussis, Neisseria meningitidis, Clostridium tetani, influenza type A and type B, Haemophilus influenza B, Streptococcus pneumoniae, or hepatitis B. Of 20 unique patient-vaccination cases, 17 were assessable for a vaccination response, whereas for the remaining 3, pre-dose antibody titre was not available. For 16 (94.1%) assessable cases, post-vaccination antibody titres increased above protective levels. For one patient who received Tetravac formulation (diphtheria, tetanus and acellular pertussis combination), the response observed for 1 (vaccination against *Clostridium tetani*) of the 3 vaccines included in Tetravac represented optical density rather than antibody concentrations and hence considered non-evaluable. For 19/20 patient-vaccinations, including those without pre-dose antibody titres, protective levels were observed during the study, which were maintained throughout the extension.

**Conclusion:** Canakinumab appeared to have no effect on post-vaccination antibody production following the administration of non-live vaccines in CAPS patients.

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**Disclosure of Interest**

P. Brogan Consultant for: Roche Pharmaceuticals and SOBI, M. Hofer Consultant for: Abbvie and Novartis, Speaker Bureau of: Abbvie, J. Kuemmerle-Deschner Grant / Research Support from: Novartis, Consultant for: Novartis, Sobi and Baxalta, B. Lauwerys: None Declared, A. Speziale Employee of: Novartis, X. Wei Employee of: Novartis, R. Laxer Grant / Research Support from: Novartis and Abbvie, Consultant for: Novartis and Abbvie

**O24** Baseline characteristics of 116 patients with chronic non-bacterial osteomyelitis (CNO) included in a national longitudinal cohort study

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**Presenting author:** Antonella Insalaco

**Pediatric Rheumatology 2017, 15(Suppl 1):O24**

**Introduction:** Chronic nonbacterial osteomyelitis (CNO) is one of the most common autoinflammatory bone disorders in childhood. Until now few data are available about the outcome and treatment of these patients. Several studies, all uncontrolled, have suggested the efficacy of non-steroidal anti-inflammatory drugs (NSAIDs), glucocorticoids, methotrexa, bisphosphonates and tumour necrosis factor α (TNF-α) inhibitors. These drugs are differently used with variable degree of response.

**Objectives:** To evaluate demographics, clinical features, laboratory and radiological assessment, therapeutic response in a large cohort of Italian CNO patients

**Methods:** Since 2015 CNO patients from 8 Italian centers were included in the new longitudinal section of the Eurofever registry. Demographic data, clinical manifestations and response to treatment were analysed.

**Results:** Baseline data from 116 patients are so far available. 72 (61%) females, 111 (95.7%) Caucasian with a median age at disease onset of
Patients presented with musculoskeletal involvement at the onset of disease (bone pain 91.4%, arthralgia 84.5%, bone alteration 55.2%, osteitis 55%, osteolytic lesions 48.3%, myalgia and arthritis were found in a minority of the patients). Other features include constitutional symptoms in 64 patients (55.2%), muco-cutaneous manifestations in 17 (14.7%), gastrointestinal involvement in 10 (8.6%) and neurological manifestations in 4 (3.4%). At diagnosis erythrocyte sedimentation rate (ESR) and C-reactive protein (CRP) were elevated respectively in 84 (72.4%) and 62 (53.4%), with 26 (22.4%) patients having ESR and CRP both normal. Serum amyloid A was elevated in 34 out of the 50 patients tested (68%). MRI/CT was performed in 107 patients (56 Whole Body MRI, 52 local MRI, 19 CT), while bone scintigraphy in 90 patients, leading to the detection of a median of 4 lesions (range 1-21). A bone biopsy was performed in 52 patients. At the baseline visit, thirty-one (26.7%) patients had received NSAIDs, 8 (7%) were treated with paminodronate, 29 (25%) with glucocorticoids, 8 with anakinra (7%), 7 (6%) with sulfasalazine. Etanercept was used in 5 patients, Methotrexate was used in 4, adalimumab, colchicine in 1 patient.

**Conclusion:** In this study we describe the baseline demographic, clinical, laboratory and radiological findings and therapy in a cohort of patients affected by CNO. Treatment used at baseline is extremely variable. Analysis of the outcome of bone lesions and of the efficacy and safety of the various treatments is ongoing on the data available from follow-up visits.

**Disclosure of Interest**
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**O25**
Analyses of 247 patients with recurrent inflammation of unknown cause

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**Pediatric Rheumatology 2017, 15(Suppl 1):O25**

**Introduction:** Auto-inflammatory diseases (AIDs) are a heterogeneous group of disorders caused by dysfunction of the innate immune system. Although many genetic mutations causing AIDs have been discovered, there is still a large group of patients with recurrent inflammation of unknown cause.

**Objectives:** Our objective is to describe the clinical features and available genetic data of patients with an undefined AID.

**Methods:** Clinical and genetic data from patients with an undefined AID were retrospectively collected in the Eurofever registry. Clinical classification criteria for FMF, MKD, TRAPS and CAPS as defined by Federici et al. were applied on the dataset [1].

**Results:** Out of the 268 patients in the Eurofever database registered with an undefined AID, 21 were excluded from further analyses because of incomplete clinical data (20) or a positive mutation for TRAPS (1). The median age of disease onset was 4.2 years (interquartile range 1.3–12.8). Patients had a median of 12 (5–15) episodes per year with a median duration of 4 (3–7) days. In 47 patients relatives were affected as well. In 204 patients analysis of one or multiple genes was performed. In 31 patients genetic variants, excluding benign polymorphisms, were found in the MEFV (12), TNFRSF1A (8), MND2 (9), NLRP3 (6), MEFV (3) and NLRP12 gene (3). None of these variants were confirmative for diagnosis of a known AID. Patients with positive genetic screening more often had relatives affected (p < 0.001, Chi-squared) and had more episodes per year (p = 0.035, Mann-Whitney U).

**Clinical manifestations are shown in the table below.** NSAI Ds were beneficial in 108/128 patients, but rarely completely effective. Corticosteroids were beneficial in 109/118 patients (51 complete, 58 partial response). Of 50 patients treated with colchicine, 10 had a complete response and 22 had a partial response. Of 15 patients treated with anakinra, 7 had a complete response and 6 a partial response. There was no correlation between the response to medication and presence of genetic variants (Fisher’s exact).

**Conclusion:** This study describes the clinical features of a group of patients with undefined periodic inflammation. Almost one third of the patients (32%) had a positive clinical classification score for a hereditary AID.

**References**
1. Federici S, Sormani MP, Ozen S, Lachmann HJ, Amaryan G, Woo P, et al. Evidence-based provisional clinical classification criteria for autoinflammatory periodic fevers. Ann Rheum Dis. England; 2015 May;74(5):799–805.

**Disclosure of Interest**
None Declared

**Table 5 (abstract O25). Clinical manifestations during episodes**

| Variable                  | Always* | Sometimes/often* | Most common complaints* |
|---------------------------|---------|------------------|------------------------|
| Musculo-cutaneous         | 22      | 47               | Aphthous stomatitis (27) |
| Gastro-intestinal         | 14      | 41               | Abdominal pain (47) |
| Musculoskeletal           | 21      | 46               | Arthralgia (61), myalgia (44) |
| Ocular                    | 2       | 12               | Conjunctivitis (8) |
| Lymphoid                  | 33      | 14               | Enlarged cervical Inn (38) |
| Cardio-respiratory        | 3       | 15               | Chest pain (13) |
| Neurological              | 15      | 26               | Headache (32) |
| Genito-urinary            | 1       | 5                | Urethritis/cystitis (3) |
| Constitution              | 25      | 40               | Fatigue (55), malaise (49) |

*% of cohort
The multifaceted presentation of chronic recurrent multifocal osteomyelitis: a series of 466 cases from the Eurofever international registry

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Introduction: Chronic Recurrent Multifocal Osteomyelitis (CRMO) is a rare autoinflammatory disorder featured by sterile bone osteolytic lesions. Objectives: To evaluate the demographic data and the clinical, instrumental and therapeutic features at baseline in a large series of CRMO patients enrolled in the Eurofever registry. Methods: A web-based registry collected retrospective data on patients affected by CRMO. Participating hospitals included paediatric and adult centers with a specific interest in autoinflammatory diseases. Results: Complete baseline information on 466 patients were available (M:F = 171/295, paediatrics/adults = 439/27). 441 patients were Caucasian, 2 Hispanic, 4 Afro-American, 4 Asian and 7 of unknown ethnicity. Mean onset age was 11 years (range 3 months-62 years). An adult onset (after the 18 years) was observed in 27 (5.8%) patients. The mean time from disease onset to final diagnosis was 1 year (range 0–15 years). Disease course was described as continuous in 197 (42%), recurrent in 244 (52%), continuous and recurrent in 25 patients (5%). At baseline, 464 patients (99.5%) displayed musculoskeletal symptoms (431 bone pain, 301 myalgia, 72 monoarthritis, 54 oligoarthritis); 87 patients (19%) mucocutaneous manifestations (25 acne, 24 palmo-plantar pustulosis, 21 psoriasis, 16 papulo-pustular lesions, 9 urticarial rash); 37 patients (8%) displayed gastrointestinal symptoms (27 abdominal pain, 15 diarrhea). Among imaging techniques, whole body magnetic resonance was performed at baseline in 141 patients (30%), revealing a mean number of 7 lesions (range 1-26); Overall, 174 patients (37%) displayed metaphyseal, 105 (23%) epiphyseal, 72 (15%) diaphyseal, 116 (25%) pelvic, 105 (23%) vertebral, 45 (10%) thoracic, 72 (15%) tarsal, 12 (3%) carpal, 13 (3%) cranial and 89 (19%) clavicular lesions. Biopsy was performed in 281 patients (60%). Complete information concerning the treatment in the period between disease onset and the enrollment in the registry were available for 386 patients. 361 (94%) patients had been treated with NSAIDs (142 complete response); 111 (29%) with steroids (42 complete response), 63 (16%) with Bisphosphonates (31 complete response), 57 (15%) with Methotrexate (13 complete response), 46 (12%) with Sulfasalazine (18 complete response), 17 (4%) with Etanercept (7 with complete response), 8 (2%) with infliximab (3 complete response) and 4 (1%) with Anakinra (2 with complete response). Eighty-one patients have recently entered in the longitudinal phase of the registry that is specifically devoted to the evaluation of long-term efficacy and safety of the different treatments performed during their follow-up. Conclusion: In conclusion, this is the largest reported case series of CRMO patients. The study shows that the disease can present with a late onset and that the range of clinical manifestations is heterogeneous. At least in the early phases of the disease NSAIDs are the most widely used drugs with a complete response in almost 40% of the patients.

Disclosure of Interest

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Clinical features distinguishing patients with Chronic Recurrent Multifocal Osteomyelitis (CRMO) from patients with tumoral osteolytic lesions: a single center experience

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Pediatric Rheumatology 2017, 15(Suppl 1):O27

Introduction: Chronic Recurrent Multifocal Osteomyelitis (CRMO) is a rare autoinflammatory disorder featured by sterile bone osteolytic lesions. The differential diagnosis includes bone malignancies, benign tumors, bacterial subacute or chronic osteomyelitis.

Objectives: The aim of our study is to analyze the possible differences in the clinical presentation, laboratory and instrumental features distinguishing CRMO from patients with tumoral osteolytic lesions.
examinations between patients with CRMO and patients presenting osteolytic lesions secondary to malignancies.

**Methods:** 48 pediatric patients affected by CRMO (M:F = 33/15, mean onset age 9.6 years, range 4-14 years), 21 patients affected by Lang- erhans cell histiocytosis (LCH, M:F = 12/9, mean onset age 4.3 years, range 3 months -12 years) and of 10 patients affected by hemat-oncological disorder (lymphoma, sarcoma and neuroblastoma, M:F = 4/6, mean onset age 7 years, range 2-11 years) presenting with osteolytic lesions and followed by a single centre since 1999 to 2015 were enrolled in the study. Clinical features, acute phase reactants values and radiological investigations (whole-body magnetic reson- ance) were obtained at diagnosis and at last available follow-up. On-set features of CRMO patients have been compared to those of LCH and hemato-oncological patients. A univariate analysis was performed comparing first CRMO, LCH and hemato-oncological patients and then only CRMO and LCH patients. Nominal variables were ana- lyzed using the χ² test and Fisher’s exact test. Continuous variables were analyzed with the nonparametric Mann- Whitney U test and the Wilcoxon test.

**Results:** Comparison among the three categories of patients (CRMO, LCH and hemato-oncological), showed that patients with bone malignancies (but not Langerhans cell histiocytosis patients) displayed a significant higher incidence of systemic symptoms: el- evated fever (>38 °C) and malaise in fact resulted prevalent in hemato-oncological cohort than in the other two categories (P re- spectively: <0.0001 and <0.01). CRP and LDH levels were signifi- cantly higher and hemoglobin levels lower in patients with bone malignancies than in patient with CRMO or LCH, whilst arthritis signs (pain, swelling and functional limitation) were more fre- quent in CRMO patients (P: <0.007). Comparing only CRMO with LCH patients, some differences in the onset variables were ob- served: onset age was higher in CRMO than in LCH patients (P <0.001). Some differences have also been detected in the radio- logical distribution of bone localizations: clavicle, distal fibula and foot were exclusive for CRMO. Moreover, patients with CRMO dis- played a greater number of lesions than LCH, (Table). (Table)

**Conclusion:** Our study shows that some clinical manifestations and laboratory data could be useful for differential diagnosis between CRMO and bone malignancies. However, LCH represents the most difficult disease implicated in differential diagnosis with CRMO. Bi- opsy still represents the gold standard examination for a correct diagnosis.

**Disclosure of Interest**
None Declared

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**Table 6 (abstract O27). Comparison between CRMO and LCH cohort of patients**

| Variables                  | CRMO (n=48) | LCH (n=21) | P       | Odds ratio (OR) |
|---------------------------|-------------|------------|---------|-----------------|
| Demographic Manifestations|             |            |         |                 |
| Mean onset age (years)    | 9.6         | 4.3        | <0.001  |                 |
| Clinical Manifestations   |             |            |         |                 |
| Musculoskeletal pain      | 48/48 (100%)| 10/21 (48%)| <0.0001 | 0.06 (0.01,0.26) |
| Musculoskeletal limitation| 31/48 (65%) | 7/21 (33%) | <0.02   | 0.27 (0.09,0.81) |
| MR lesions site           |             |            |         |                 |
| Clavicle                  | 11/48 (22.9%)| 0/21 (0%) | <0.01   |                 |
| Distal Fibula             | 9/48 (19%)  | 0/21 (0%)  | <0.05   |                 |
| Foot                      | 10/48 (20.8%)| 0/21 (0%) | <0.03   |                 |
| Mean number of MR lesions| 6.5         | 2          | <0.03   |                 |

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**O28**

Cryopyrin associated periodic syndromes (CAPS): immunological characterization of knock-in mouse model to exploit novel approaches for the modulation of the NLRP3 inflammasome

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This abstract is not included here as it has already been published.

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**Oral presentations 2**

**O29**

Predicting which children with juvenile idiopathic arthritis will have a severe disease course: results from the REACCh-Out cohort

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**Introduction:** We previously reported that the disease course of chil- dren with juvenile idiopathic arthritis (JIA) could be described in four groups, based on five variables prioritized by patients, parents and clinicians (Ann Rheum Dis 2015;74:Suppl 2 609 doi:10.1136/ annrheumdis-2015-eular.2396). The four groups were labeled as Mild (43.8% of children), Moderate (35.6%), Severe Controlled (9%) and Severe Persisting (11.5%).

**Objectives:** To develop a risk stratification method to estimate the probability of a severe disease course for each individual child with JIA at diagnosis.

**Methods:** We used data from 609 children followed in the Research in Arthritis in Canadian Children emphasizing Outcomes (REACCh- Out) prospective inception cohort who attended at least 6 of 8 planned visits in the five years after diagnosis. We used information available at study enrollment to predict disease course in the subse- quent 5 years. We divided the cohort at random into a training sam- ple (75%) to develop candidate risk stratification methods and a testing sample (25%) to test the accuracy of stratification. This process was repeated 50 times for each of 100 candidate models de- rived from logistic regression, random forest, neural net or nearest neighbor methods. The 50 random splits allowed estimation of the stability and accuracy of the models in subjects that were not used in the development of the model, to select the most reliable, accur- ate and parsimonious method. Model performance was measured with c-index, Maximum Likelihood and Pearson statistics (higher values reflect better performance).

**Results:** For prediction of a severe disease course (Severe Controlled and Severe Persisting combined) the best performing model was de- rived using logistic regression. It had a c-index of 0.85 (95% CI 0.80, 0.90) and 91% of children assigned to the highest decile of risk actu- ally had a severe disease course in the test sample (Table 7),
compared to 5% of those assigned to the lowest decile of risk. The model used a combination of JIA category, active joint count, pattern of joint involvement and routine tests available at the first study visit. The model’s performance was better than using JIA category alone (c-index of 0.71; 95% CI 0.65, 0.79). For reference, the widely used Framingham’s Cardiovascular Risk Score has c-index values of 0.75 to 0.80 depending on the cohort.

Conclusion:
In this cohort, a risk stratification method that used variables usually available at diagnosis accurately estimated the risk of a severe JIA course during the following 5 years. Knowing the probability of a severe disease course at diagnosis will help parents and physicians better tailor the optimal aggressiveness of treatment for each child with JIA.

Disclosure of Interest
None Declared

Table 7 (abstract O29). Proportion of children with a severe disease course (severe controlled and severe persisting combined) in the test data across 50 re-samples, according to their decile of risk

| Decile of risk | 1) Logistic Random forest 2) K-nearest neighbor 3) Neural network 4) JIA category alone |
|---------------|--------------------------------------------------------------------------------------------|
| First (low)   | 0.05 0.04 0.16 0.11 0.11 0.05                                                             |
| Third         | 0.05 0.05 0.05 0.09 0.09 0.06                                                             |
| Sixth         | 0.10 0.13 0.17 0.10 0.10 0.32                                                             |
| Ninth         | 0.43 0.45 0.29 0.41 0.36                                                                 |
| Tenth (high)  | 0.91 0.88 0.51 0.74 0.37                                                                 |

C-Index (95% CI) 0.85 (0.80, 0.90) 0.85 (0.82, 0.88) 0.67 (0.59, 0.75) 0.75 (0.71, 0.79) 0.71 (0.65, 0.79)

Maximum likelihood (95% CI) -51 (-53, -53) -58 (-49, -66) -81 (-61, -100) -107 (-99, -116) -69 (-60, -77)

Pearson (95% CI) 47 (34, 61) 45 (37, 53) 12 (3, 20) 26 (18, 33) 13 (7, 19)

O30
The vasculopathy of JDM: evidence of persistent endothelial injury, hypercoagulability, subclinical inflammation and increased arterial stiffness
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Introduction: Vasculopathy is considered central to the pathogenesis of Juvenile Dermatomyositis (JDM). Moreover, the interplay between persistent JDM-vasculopathy, traditional cardiovascular risk factors, exposure to corticosteroids, and chronic inflammation could create a perfect storm for early atherosclerosis. One major hurdle to the study and detection of the vasculopathy of JDM, monitoring of its trajectory over time and contribution to excess cardiovascular disease has been a lack of non-invasive biomarkers. Recently, we described a number of methods for detecting endothelial cell components in blood which allow non-invasive assessment of vascular injury: circulating endothelial cells (CEC), and endothelial microparticles (EMP).

Objectives: To explore the vasculopathy of JDM by assessing: (i) biomarkers of endothelial injury (CEC and EMP), subclinical inflammation (cytokines) and hypercoagulability (MP-mediated thrombin generation); and (ii) structural arterial changes (indictative of premature atherosclerosis), and their relation to disease activity and treatment in children with JDM.

Methods: 64 patients recruited to the UK JDM Cohort & Biomarker Study were included; median age 10.5 (range 6.9–13.7) years with median disease duration of 1.5 (0.3–4.7) years. 40 (62.5%) were females. Inactive disease was defined as per modified PRINTO criteria: no skin rashes, CK ≤150, CMAS >48, MMTB ≥78. Physician’s global assessment ≤0.2 on a visual analogue scale. CECs and MPs were identified with immunomagnetic bead extraction and flow cytometry, respectively. MP function as assessed by thrombin generation was determined using a fluorogenic assay. Cytokines and chemokines were measured by electrochemiluminescence. Arterial stiffness was assessed using pulse wave velocity (PWV). Results are expressed as median and range.

Results: CECs were higher in JDM patients at 68 (32-128) cell/ml compared to 12 (8-21) cells/ml in 66 age-sex matched healthy controls, p < 0.0001. Patients with active JDM had higher CEC than those with inactive JDM, p = 0.02. Patients with calcinosis had higher CEC compared to patients without calcinosis, p = 0.03. CEC counts significantly correlated with levels of inflammatory cytokines/chemokines implicated previously in JDM disease pathogenesis: interferon regulated Monocyte Chemoattractant Protein-1 (MCP-1; r = 0.63, p = 0.02) and interleukin-8 (IL-8; r = 0.65 and p = 0.01). Total circulating MP counts were also significantly higher in active JDM, 1781 (981–2616) x10³/ml compared to inactive JDM, 1116 (263–1393) x10³/ml, p = 0.02; and healthy controls 89 (25–236) x10³/ml, p = 0.0001. These circulating MPs were predominantly of platelet and endothelial origin. Enhanced MP mediated thrombin generation was demonstrated in active compared to inactive JDM (p = 0.03) and controls (p = 0.001). Lastly, children with JDM had increased carotid-radial PWV adjusted for age compared to healthy controls (p = 0.005).

Conclusion: Our data demonstrate: 1. Increased endothelial damage in children with active JDM, possibly driven by pro-inflammatory cytokines; 2. High levels of circulating MP with propensity to drive thrombin generation and hence occlusive vasculopathy; and 3. Increased arterial stiffness, suggestive of accelerated atherosclerosis in patients with JDM. Validation of these biomarkers in multicentre prospective studies will provide data regarding their prognostic relevance.

Disclosure of Interest
None Declared

O31
Online peer mentoring for adolescents with juvenile idiopathic arthritis: the iPeer2Peer Program
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Presenting author: Lynn R. Spiegel
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O32
Adjudication of infections in the pharmacovigilance in juvenile idiopathic arthritis patients (Pharmachild) treated with biologic agents and/or methotrexate

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Introduction: The Pharmachild study is aimed at observing children with JIA undergoing treatment primarily with biologics ± methotrexate (MTX). It is conducted by the participating centres of over 60 countries belonging to PRINTO and PReS.

Objectives: Primary endpoint of the study is safety in JIA patients undergoing immunosuppressive therapy. A preliminary analysis of infections has been performed by an independent Safety Adjudication Committee of 6 experts (3 pediatric rheumatologists and 3 pediatric infectious disease specialists).

Methods: The centres were asked to report all infections encountered by the patients followed for JIA. Data were checked by PRINTO and the medical monitor based on MedDRA dictionary. Upon completion of these steps, all eligible cases for adjudication, represented by serious/severe/very severe infections, were submitted to the expert Committee. The experts were asked to answer 6 questions. The events with consensus of at least 4/6 experts on more than 3/6 questions were considered. An additional analysis was performed on opportunistic infections, with referral to the recommendations by Withrop et al.[1], and on the correlation between infections and 3 treatment phases: MTX only, first biologic ± MTX and at least 2 sequential biologics ± MTX after failure of the first biologic.

Results: 7817 patients were enrolled in Pharmachild at the time of preliminary analysis, with 27% of total safety events reported as infections. 360 events were submitted to the Safety Adjudication Committee. 94 infections (26.1%) achieved maximum consensus among the experts (6/6 questions), 237 (65.8%) consensus on 4/6 or 5/6 questions, the remaining events were discarded. 330 infections were finally retained for the analysis, mostly represented by serious infections of moderate/severe intensity. With regard to the other questions, 74.2% infections were adjudicated as common, appropriately treated, with no clear correlation to the immunosuppressive therapy. The experts achieved consensus on 18 infections adjudicated as opportunistic. Referring to the list of opportunistic infections provided by Winthrop et al.[1], we identified 89/330 (27%) opportunistic infections, mostly represented by EBV and herpes zoster infections. 98.8% of events were not recoded. A doubled risk of infections was found in the group of patients who were started on their first biologic ± MTX or who had been treated with at least 2 sequential biologics ± MTX (1.3 and 1.9 infectious events per 100 py, respectively), compared to the MTX only group (0.8 infectious events per 100 py). The same was found for serious and opportunistic infections.

Conclusions: Our preliminary analysis showed a significant number of infections in JIA patients on immunosuppressive therapy, with one third classified as opportunistic infection. The introduction of biologics in therapy increased the risk of opportunistic and serious infection.

Disclosure of Interest
None Declared

O33
The influence of early achievement of clinically inactive disease on long-term disability outcomes in juvenile idiopathic arthritis

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Introduction: Published definitions for clinically inactive disease (CID) in juvenile idiopathic arthritis (JIA) have recently been shown to identify different groups of children. Wallace’s preliminary criteria captures children with no objective inflammatory signs, whereas the clinical Juvenile Arthritis Disease Activity Score (cJADAS) cut-off identifies children with little objective inflammation and, since a parent global assessment is included, little chronic pain and fatigue. It is unclear whether long-term outcomes, such as functional ability, differ between those who, early in disease, achieve inactive inflammatory disease but have ongoing subjective symptomatology versus those who achieve CID on both aspects.

Objectives: To assess whether five-year functional outcome using the Childhood Health Assessment Questionnaire (CHAQ) differs according to disease activity state at one year following JIA diagnosis.

Methods: Children were recruited from the Childhood Arthritis Prospective Study (CAPS), a multicentre UK inception cohort. Children with a physician’s diagnosis of JIA (limited to oligoarticular and polyarticular subtypes) and recruitment prior to 1st January 2011 were selected. Children were categorised into three disease states at one year following presentation: i) CID on both tools, ii) CID on Wallace’s preliminary criteria only and iii) No CID. Mean CHAQ score and percent of children with CHAQ ≤0.5 were assessed at five years and compared descriptively between each disease state group. Multiple imputation accounted for missing data.

Results: Of 832 children included, 70% were female and median symptom duration at diagnosis was five months (IQR 3 to 11 months). The most common subtype was oligoarticular (68%), followed by 27% RF-negative (27%) and 5% RF-positive polyarticular JIA. At one year, 22% had achieved CID according to both Wallace’s preliminary criteria and cJADAS, 7% according to only Wallace’s preliminary criteria and 56% to neither. Compared to those who had not achieved CID, children who had achieved CID on both tools at one year showed a trend to improved CHAQ scores at five years. However, those who had achieved CID only according to Wallace’s preliminary criteria developed similar CHAQ scores to the no CID group (Table 8). This difference may have been driven by the parental global assessment, with mean one year scores of 0.4 cm for the children in ID on both tools, but a similar scores for children in CID only according to Wallace’s preliminary criteria and those not in CID, (2.8 cm and 2.5 cm respectively).

Conclusion: Absence of clinically evident inflammation in JIA does not in itself predict better long-term functional ability. Better long-term functional ability is only observed to increase when both objective measures of inflammation and subjective measures of the global picture of disease are taken into account. Children in CID according to Wallace’s preliminary criteria but not the cJADAS may require alternative interventions to manage their ongoing non-inflammatory symptoms.

Disclosure of Interest
None Declared
Table 8 (abstract O34). Functional ability at five years in children who achieved different disease states at one year following diagnosis

| CID state at one year following diagnosis | Mean CHAQ at five years following diagnosis (95% CI) | Percent with CHAQ ≤ 0.5 at five years following diagnosis (95% CI) |
|----------------------------------------|---------------------------------------------------|------------------------------------------------------------------|
| CID on Wallace's preliminary criteria and cJADAS | 0.7 (0.5, 0.8) | 51 (40, 63) |
| CID only on Wallace's preliminary criteria | 1.0 (0.8, 1.3) | 33 (16, 50) |
| No CID | 0.9 (0.8, 1.0) | 37 (29, 44) |

O34
Cost-effectiveness of first year treatment for juvenile idiopathic arthritis (JIA) in the era of biosimilars
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Introduction: Biologics are costly, but effective in JIA. Little is known about the cost-effectiveness of non-biologic treatment of JIA.

Objectives: We aimed to compare the cost-effectiveness of three treatment strategies of JIA during the first year after onset.

Methods: In Acute-JA study1, 60 patients were randomized to receive either infliximab (IFX) plus methotrexate (MTX); combination (COMBO) of MTX, hydroxychloroquine, and sulphasalazine; or MTX alone. Efficacy was measured at eight study visits, and data on both health care and non-health care costs were collected. Costs were assessed at 2012 level, despite for biologics, which were assessed at 2016 level. Societal perspective was used. Health utility was produced from CHAQ using the NICE quadratic mapping algorithm.

Results: COMBO seemed more cost-effective over aTNF and MTX. (Table 9). Cost per additional months spent in inactive disease state when using prices for original molecules were 729, 446, and 1372, respectively. When using the prices for biosimilars, this cost was 543, 436, and 1372, respectively.

Conclusion: COMBO seemed a dominant treatment for JIA, over MTX and aTNF.

Disclosure of Interest
None Declared

Table 9 (abstract O34). Incremental cost-effectiveness ratio (ICER) using prices of biosimilars in treatment of JIA

| Treatment | Incremental Costs, € | Incremental QALYs | ICER estimate 6/Qaly |
|-----------|----------------------|-------------------|----------------------|
| COMBO vs MTX | -2918 | 0.072 | dominant* |
| aTNF vs combo | 7627 | 0.023 | 331 609 |
| aTNF vs MTX | 4926 | 0.095 | 51 853 |

QALY Quality-Adjusted Life-Year
*less costly and more effective treatment option

O35
Peripheral blood B cells are expanded and their cytokine expression is dysregulated in juvenile dermatomyositis
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Introduction: Juvenile dermatomyositis (JDM) is a rare form of childhood autoimmune myositis that presents with proximal muscle weakness and heliotrope rash. B cells are strongly implicated in the pathogenesis of the disease and are responsible for the generation of myositis specific antibodies, that are closely associated with specific clinical subgroups. B cells have been detected in inflamed muscle suggesting another pathological function. We propose that B cells may also contribute to the disease by producing pro-inflammatory cytokines, triggered by type I interferons.

Objectives: To characterise B cell subsets and B cell production of interleukin 6 (IL-6) and interleukin 10 (IL-10) in JDM patients at presentation and after treatment with corticosteroids and methotrexate.

Methods: Samples and data were collected from children with JDM recruited to the UK JDM Cohort and Biomarker Study. Peripheral blood mononuclear cells (PBMC) samples from JDM patients before treatment and on treatment were selected for analysis. PBMC from child healthy controls (CHC) (n = 25) and JDM patients (n = 61) were incubated in the presence of phorbol 12-myristate 13-acetate (PMA), Ionomycin and Golgi Plug for 4 hrs and cytokine expression was analysed by flow cytometry. PBMC and sorted B cells were also stimulated for 72 hr with CD40L or 48 hr with R848 (TLR7/8) and assessed for intracellular IL-6 and IL-10 by flow cytometry and protein levels were measured by ELISA. PBMC were also stained ex-vivo to detect B cell subsets and Ki67 expression. Differences between groups were analysed using paired/ unpaired t tests or one-way ANOVA (Prism 6 software).

Results: The B cell frequency was significantly higher in pre-treatment JDM patient compared to CHC samples (25.72% of total B cells Vs. 22.69% (rs) (<6 months treatment) or 16.26% in CHC; p < 0.01). This difference was most marked in the immature B cell subset (36.05% of total B cells Vs. 3.66% (<6 months on treatment) or 12.4% in CHC; p < 0.01). The immature B cell subset was more proliferative before treatment (mean Ki-67% of 17.63% Vs. 8.86% (>6 months on treatment) or 7.18% in CHC; p < 0.01). This difference was specific to the immature B cell subset. Whole transcriptome RNA sequencing of B cells from paired pre and on treatment patient samples showed an upregulation of the type I interferon signature. Of particular interest Toll-like receptor 7 (TLR7) which was up-regulated in B cells from patients' pre-treatment. When B cells were stimulated with TLR7 agonist (R848) there was reduced expression of IL-10 detected by ELISA from JDM compared to CHC samples (486.18 pg/ml Vs. 414.29 pg/ml). When B cells were stimulated with CD40L there was equivalent expression of IL-10 from JDM pre-treatment and CHC samples (25.49% of total B cells Vs. 13.26% (<6 months on treatment)); p < 0.01) or 20.07% in CHC).

Conclusion: We have identified a subset of immature B cells that are expanded, more proliferative and express both IL-6 and IL-10 in pre-treatment patients. The up-regulated type I interferon signature in JDM could contribute to the expansion of immature B cells with an imbalance of IL-6 and IL-10 production.

Disclosure of Interest
None Declared
**O36**

**Uveitis associated to polyarticular juvenile idiopathic arthritis (pJIA): data from strive registry**

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**Introduction:** Juvenile idiopathic arthritis (JIA) is one of the most common of all systemic diseases associated with childhood uveitis. Approximately 10–15% of JIA patients (pts) experience comorbid uveitis¹.

**Objectives:** To explore events of uveitis and associated safety in pts with moderately to severely active polyarticular or polyarticular-course JIA (pJIA) who were prescribed and treated with adalimumab (ADA) and/or methotrexate (MTX) in routine clinical practice.

**Methods:** STRIVE is an ongoing, multicenter, non-interventional, observational registry of up to 10 years duration in pts with moderately to severely active pJIA who are treated with either ADA ± MTX or MTX alone as part of their routine clinical care. Pts could initiate ADA and/or MTX within 24 months prior to registry entry. Pts that completed ADA studies (DE038, M10-444) had an option to roll-over into this registry. Ophthalmologists performed slit-lamp examination for uveitis at registry entry and specified visits in 3-6 month intervals through 5 yrs. Beyond 5 yrs, uveitis events were collected solely through adverse event (AE) reporting. Observational ocular AEs (e.g. cataract, glaucoma) were recorded from registry entry through yr 6.

**Results:** As of 1 June 2015, a total of 21/303 (6.9%) and 68/543 (12.5%) enrolled pts reported at least 1 case of JIA-associated uveitis at any visit in the MTX and ADA ± MTX groups, respectively. In the JIA-associated uveitis population, 10 (47.6%) in the MTX and 42 (61.8%) in the ADA ± MTX group presented with documented uveitis at registry entry. In the population without uveitis at registry entry, 11/293 (3.8%) pts and 26/501 (5.2%) pts in MTX and ADA ± MTX arms, respectively had first documentation of uveitis post-enrollment. Most pts in the JIA-associated uveitis sub-population were female (73%), white (96%), with a mean age of 8.1 yrs; mean pJIA disease duration was 1.8 and 4.8 yrs for the MTX and ADA ± MTX groups at registry entry, respectively. Nineteen (42.9%) MTX and 48 ADA ± MTX (72.7%) pts were positive for antinuclear antibodies at registry enrollment. Uveitis in the vast majority of pts was localized to the anterior layer. In the ADA group, 45 (66.2%) pts with documented uveitis received concomitant MTX during the course of the registry. Through Month 42, the majority of the JIA-associated uveitis sub-population had either no new manifestation of uveitis or stabilized uveitis. A higher proportion of MTX vs ADA ± MTX pts discontinued the registry drug (15/21 [71.4%] vs. 21/68 [30.9%]), but continued to be monitored for safety follow-up. Of these, 2 (9.5%) and 1 (1.5%) in the MTX and ADA ± MTX group, respectively, discontinued the registry drug due to an AE, and 4 of the 15 pts in the MTX group discontinued MTX group and switched to the ADA ± MTX registry group. Two (0.4%) pts with glaucoma and 1 (0.2%) pt with cataract were reported in the ADA ± MTX group and none in the MTX group; 2 of these patients had documented uveitis at registry enrollment.

**Conclusion:** Among pJIA pts with uveitis documented at registry entry, a higher percentage of pts were enrolled in the ADA ± MTX group as per investigator judgment. No new safety signals for adalimumab were observed in the JIA-associated uveitis sub-population treated per standard of care. Based on this interim analysis, JIA-associated uveitis appeared well-controlled during the course of this registry for pJIA pts with existing uveitis.

**Trial registration identifying number:** NCT00783510

**Reference**

1) Rabinovich CE, 2007; 19(5): 482–486.

**Disclosure of Interest**

I. Foeldvari Consultant for: AbbVie and Novartis, N. Ruperto Grant / Research Support from: AbbVie Inc., AstraZeneca, Bristol-Myers Squibb, Janssen Biologics B.V., Eli Lilly and Co., “Francesco Angelini”, GiaxomithKline, Italfarmacopo, Novartis, Pfizer, Roche, Sanofi Aventis, Schwarz Biosciences GmbH, Xoma, and Wyeth Phgroupaceuticals, Employee of: GASNLI Hospital, Speaker Bureau of: Astellas, AstraZeneca, Bristol-Myers Squibb, Italfarmaco, Janssen Biologics B.V., MedImmune, Roche, and Wyeth/Pfizer, D. Lovell Consultant for: AbbVie Inc., AstraZeneca, Centocor, Bristol-Myers Squibb, Boehringer-Ingelheim, Pfizer, Regeneron, Hoffman La-Roche, Novartis, UCB, and Genentech, G. Horneff Grant / Research Support from: AbbVie, Pfizer, and Roche, H.-I. Hupperetz Consultant for: Wyeth Pharmaceuticals, Essex Pharmaceuticals, and Abbott Laboratories, Speaker Bureau of: Wyeth Pharmaceuticals, Essex Pharmaceuticals, and Abbott Laboratories, P. Quarter Consultant for: AbbVie, Novartis, Pfizer, ADM, Chugai-Roche, MedImmune, Servier, and Swedish Orphan Biovitrum, Speaker Bureau of: Member of a data monitoring board for a Sanofi trial, G. Simonini: None Declared, M. Bereswill Shareholder of: AbbVie Inc, Employee of: AbbVie Deutschland GmbH & Co. KG, J. Kalabic Shareholder of: AbbVie Inc, Employee of: AbbVie Deutschland GmbH & Co. KG, A. Martini Grant / Research Support from: Abbott, BMS, Francesco Angelini S.P.A., GSK, Janssen Biotech, Novartis, Pfizer, Roche, Sanofi-Aventis, Schwarz Biosciences GmbH, Consultant for: BMS, Janssen Biotech, GSK, Novartis, Pfizer, Roche, Sanofi-Aventis, Schwarz Biosciences GmbH, Employee of: GASNLI Hospital, Speaker Bureau of: Abbott, AbbVie, Amgen, Baxalta Biosimilaris, Biogenidec, BMS, Astellas, Boehringer, Italfarmaco, Janssen, MedImmune, Novartis, NovoNordisk, Pfizer, Sanofi, Roche, Servier, Takeda, UCB Biosciences GmbH, H. Brunner Consultant for: AbbVie Inc., AstraZeneca, Centocor, Bristol-Myers Squibb, Boehringer-Ingelheim, Pfizer, Regeneron, Hoffman La-Roche, Novartis, UCB, and Genentech, Speaker Bureau of: Genentech Pharmaceuticals.

**O37**

**Changes in quality of life over time in children with juvenile idiopathic arthritis: results from the ReACCh-Out cohort**

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**Introduction:** Health-related quality of life (HRQOL) measures provide a more holistic view of the impact of health conditions than single outcome measures. Recent studies suggest a dissociation of HRQOL and traditional Juvenile Idiopathic Arthritis (JIA) disease outcomes, with suboptimal HRQOL reported in children whose disease was inactive or minimally active.

**Changes in quality of life over time in children with juvenile idiopathic arthritis:**

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**Presenting author:** Jaime Guzman

**Pediatric Rheumatology 2017, 15(Suppl 1):O37**
Objectives: To describe changes in HRQOL from diagnosis in a large prospective inception cohort of children with JIA.

Methods: We analyzed data from the Research in Arthritis in Canadian Children Emphasizing Outcomes cohort (REACCh-Out). Families (the child and a parent) of children newly diagnosed with JIA between January 2005 and December 2010 were asked to complete the Juvenile Arthritis Quality of Life Questionnaire (JAQQ) and the Quality of My Life visual analogue scale (QoML) at 0, 6, 12, 18, 24, 36, 48 and 60 months after enrolment; data to complete the Juvenile Arthritis Quality of Life Questionnaire with JIA between January 2005 and December 2010 were asked.

Families (the child and a parent) of children newly diagnosed with JIA shortly after diagnosis, and it improves over time at different rates across JIA categories. Achievement of best possible HRQOL scores lagged behind improvements in other outcome measures. This may explain the apparent dissociation of HRQOL from these measures in cross-sectional studies.

Disclosure of Interest
None Declared

Table 10 (abstract O37). See text for description

| Total | Gross motor | Fine motor | Psycho-social | Systemic Symptoms | QoML | Active joint pain | Physician's | CHAQ-
|-------|-------------|------------|---------------|-------------------|------|------------------|-------------|-------|
| Median score first visit (IQR) | 2.7 (2.4) | 2.3 (2.5) | 2.4 (1.3) | 2.4 (1.3) | 3.6 (2.5) | 7.8 (5.9) | 2 (1.6) | 2.8 (1.5) | 0.4 (0.1, 1) |
| Median months to best score | 59.3 | 21.2 | 7.2 | 21.8 | 40.1 | 34.5 | 11.4 | 14.9 | 14 |
| Probability of best score within 1 year | 0.10 | 0.32 | 0.61 | 0.33 | 0.15 | 0.25 | 0.53 | 0.38 | 0.44 |
| Probability of best score within 2 years | 0.24 | 0.52 | 0.79 | 0.53 | 0.32 | 0.42 | 0.81 | 0.68 | 0.68 |

Best possible scores were a JAQQ = 1, a QoML = 10
a an active joint count = 0
b a physician global = 0 in a 100 mm visual analogue scale
c a Childhood Health Assessment Questionnaire (CHAQ) = 0

O38
Subcutaneous abatacept (ABA) 10 mg/kg every 4 weeks is well tolerated and effective in reducing the signs and symptoms of polyarticular-course juvenile idiopathic arthritis (pJIA). SC ABA 125 mg weekly has equivalent efficacy and comparable safety to IV ABA in adult RA.

Objectives: To evaluate SC ABA treatment in patients with active pJIA.

Methods: Patients with pJIA and an inadequate response/intolerance to ≥1 DMARD were enrolled in two age cohorts (2–5 years and 6–17 years) in this single-arm, open-label (OL), Phase III study and received SC ABA weekly OL for 4 months based on body weight tier (10–<25 kg [50 mg ABA]; 25–50 kg [87.5 mg ABA]; >50 kg [125 mg ABA]). JIA-ACR criteria 30 (JIA-ACR30; ACR Pediatric 30) responders at 4 months could enter a 20-month OL extension. Primary endpoint was ABA steady-state blood trough concentration (Cminss) at 4 months in the 6–17-year cohort.

Results: For the 2–5-year (n = 32; interim analysis) and 6–17-year (n = 173; complete analysis) cohorts, respectively: median (min, max) age: 5.0 (2.0, 5.0) and 13.0 (6.0, 17.0) years; number of active joints: 7.0 (2.0, 27.0) and 10.0 (2.0, 42.0). The target therapeutic Cminss was achieved in both cohorts (mean [SD] Cminss at 4 months: 2–5 years: 50.1 [14.2] μg/mL; 6–17 years: 42.1 [14.7] μg/mL). Robust JIA-ACR30, 70 and inactive disease (no active joints, physician’s global assessment of disease activity <10 mm, CRP <0.6 mg/dL) responses were seen at 4 months in the 2–5-year cohort: 86.7, 70.0 and 51.7%, respectively, and in the 6–17-year cohort: 80.9, 52.6 and 29.5%, respectively. No new or unexpected safety concerns were reported (Table).

Conclusion: Target therapeutic exposure for SC abatacept was achieved in patients with pJIA aged 2–17 years; exposures were within the range observed for IV abatacept in pJIA. Robust efficacy was seen after 4 months of treatment with no new safety concerns.

1. Ruperto N, et al. Lancet 2008;372:383–91.
2. Ruperto N, et al. EULAR Annual European Congress of Rheumatology 2016: Abstract OP0215.

Trial registration identifying number: ClinicalTrials.gov NCT01844518

Disclosure of Interest
None Declared
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### Table 11 (abstract O38), Summary of AEs during the combined initial 4-month and 20-month extension period (All Treated Patients)

| Event Type | 2-5-year cohort (n = 32) | 6-17-year cohort (n = 173) |
|------------|------------------------|---------------------------|
| Deaths     | 0                      | 0                         |
| All AEs    | 26 (81.3)              | 127 (73.4)                |
| Related AEs| 11 (34.4)              | 45 (26.0)                 |
| AEs leading to discontinuation | 0  (2.3) | 4 (2.3)*                  |
| SAEs       | 0                      | 8 (4.6)                   |
| Related SAEs | 0 (0.6)               | 1 (0.6)                   |
| SAEs leading to discontinuation | 0 (1.2)*               | 2 (1.2)*                   |
| AEs of special interest | | | |
| Malignancies | 0 (1.0)               | 0 (0.6)                   |
| Autoimmune disorders | 0 (3.7)               | 0 (3.7)                   |
| Local injection-site reactions | 0 (10.5)              | 0 (10.5)                  |
| Infections | 22 (68.8)              | 90 (52.0)                 |

Data are n (%)

SAE serious adverse event

*Exanthema (n = 1) and fatigue (n = 1), both related to study drug (as well as the two SAEs leading to discontinuation)

*Sepsis (n = 1), related to study drug; stage III ovarian germ cell teratoma (n = 1), not related to study drug

### O39 Expression of type I and type II interferons is increased in muscle biopsies of juvenile dermatomyositis patients and related to clinical and histological features

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**Introduction:** Juvenile dermatomyositis (JDM) is the most common juvenile inflammatory myopathy, with a still not fully clarified immunopathogenesis. However, there is relevant evidence for an involvement of interferons (IFNs) in the chronic inflammation that characterizes JDM, and a better characterization of their role may provide promising targets for new therapies.

**Objectives:** The aim of this study was to investigate muscle expression of type I (IFNα/β) and type II (IFNγ) IFN inducible genes in muscle biopsies of JDM patients and their correlations with clinical and histological aspects of the disease.

**Methods:** A retrospective cohort of patients diagnosed with JDM (n = 22), expression of specific genes induced by IFNα/β (IFIT2, IFI44L, IFIT1, ISG15, RSAD2, SIGLEC1), the so called “type I IFN signature”, by IFNγ (CXCL9, CXCL10, CXCL11, CIITA), and IFNγ itself, were analysed by real-time PCR on snap frozen muscle biopsies and compared with samples from Duchenne muscular dystrophy (DMD) patients (n = 24). We also analysed mRNA expression of pro-inflammatory cytokines such as interleukin-1β (IL-1β), tumor necrosis factor-α (TNFa) and interleukin-6 (IL-6). For each patient charts were reviewed to record clinical features at diagnosis, physician’s global assessment of the patient’s overall disease activity, serum levels of muscle enzymes (CK, ALT, AST, LDH), erythrocyte sedimentation rate (ESR), C-reactive protein level, antinuclear antibodies status, time to inactive disease, number of immunosuppressants used over disease course and relapses. We also evaluated typical histological aspects of JDM (inflammatory infiltrate, necrosis, perifascicular atrophy and fibrosis) on tissue sections of the muscle biopsies.

**Results:** Since glucocorticoid therapy strongly reduced muscle expression of cytokines, JDM patients treated before biopsy were excluded from analysis. The mRNA expression of type I IFN signature genes (type I-IFN score) was significantly higher in untreated JDM patients (n = 16) compared with DMD patients (p < 0.0001). Expression of IFNγ and IFNγ related genes (CXCL9, CXCL10, CXCL11, CIITA) were significantly higher in biopsies of untreated JDM patients compared with those of DMD patients (p < 0.01, p < 0.01, p < 0.0001, p < 0.0001, p < 0.0001, respectively). Expression of TNFa, but not of IL-1β and IL-6, was significantly higher in untreated JDM muscles compared with those of DMD patients (p < 0.01). IFNγ expression significantly correlated with CIITA, CXCL9, CXCL10 and CXCL11 mRNA levels (p < 0.05, p < 0.0001, p < 0.01, respectively). Type I-IFN score significantly correlated with ERS, CK, time to inactive disease and number of immunosuppressants (p < 0.05, p < 0.05, p < 0.01, respectively). IFNγ mRNA levels significantly correlated with time to inactive disease and relapse after remission (p < 0.05, p < 0.01, respectively). We also found that type I-IFN score correlated with inflammatory infiltrate and necrosis (p < 0.05), while IFNγ correlated with inflammatory infiltrate, perifascicular atrophy and fibrosis (p < 0.01, p < 0.05, p < 0.05 respectively).

**Methods:** The increased expression of IFN related genes in muscle biopsies of JDM patients and their association with clinical and histological features suggest a pathogenic role of IFNs in muscle damage and inflammation in JDM. Thus, both type I and type II IFNs pathways may represent therapeutic targets in JDM.

**Disclosure of Interest:** None Declared

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**Table 11 (abstract O38), Summary of AEs during the combined initial 4-month and 20-month extension period (All Treated Patients)**

- Deaths: 0
- All AEs: 26 (81.3) vs. 127 (73.4)
- Related AEs: 11 (34.4) vs. 45 (26.0)
- AEs leading to discontinuation: 0 vs. 4 (2.3)*
- SAEs: 0 vs. 8 (4.6)
- Related SAEs: 0 vs. 1 (0.6)
- SAEs leading to discontinuation: 0 vs. 2 (1.2)*
- AEs of special interest:
  - Malignancies: 0 vs. 0 (1.0)
  - Autoimmune disorders: 0 vs. 3 (1.7)
  - Local injection-site reactions: 0 vs. 10 (5.8)
  - Infections: 22 (68.8) vs. 90 (52.0)

*Data are n (%)
O40

Next generation sequencing reveals complement deficiencies are the most frequent causes of monogenic lupus in children
Alexandre Belot, Gillion Rice, Anne-Laure Mathieu, Sulliman O. Omarjee, Brigitte Bader-Meunier, Thierry Walzer, Tracy A. Briggs, James O'Sullivan, Simon Williams, Rolando Cimaz, Eve Smith, Michael W. Beresford, Yanick J. Crow and GENIAL Investigators, UK JSLE Study Group
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Presenting author: Alexandre Belot
Pediatric Rheumatology 2017, 15(Suppl 1):O40

This abstract is not included here as it has already been published.

Various topics

O41
Prevention and treatment of steroid induced osteopenia in children and adolescents with rheumatic diseases: the pops study
Madeleine Rooney, Nick Bishop, Joyce Davidson, Clarissa Pilkington, Michael Beresford, Jacqui Clinch, Rangara Satyapal, Helen Foster, Janet Gardner Medwin, Janet McDonagh, Sue Wyatt,1 and On Behalf of the British Society for Paediatric and Adolescent Rheumatology
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Presenting author: Madeleine Rooney
Pediatric Rheumatology 2017, 15(Suppl 1):O41

Introduction: Children with rheumatic diseases have reduced bone density and increased vertebral fracture rate compared to controls and that children on high dose steroids have twice the fracture risk of those on low dose. Despite the widespread use of biologics some 30% of children with JIA for example remain on steroids. There is ample evidence base for the treatment of adults with low bone density (BMD); there is none for children and adolescents.

Objectives: To investigate whether the bisphosphonate risedronate was superior to 1-alfacalcidol or placebo, in increasing BMD as a surrogate for reducing fracture risk

Methods: Children and adolescents with JIA, JDM, JSLE or vasculitis, commencing or receiving steroids were eligible. Patients were stratified using a minimisation technique according to: disease type; commencing or receiving steroids were eligible. Patients were stratified for reducing fracture risk was superior to 1-alfacalcidol or placebo, in increasing BMD as a surrogate for reducing fracture risk

Table 12 (abstract O41).

| Variables          | Placebo | One alpha | Risedronate | All patients |
|--------------------|---------|-----------|-------------|--------------|
| N                  | 77      | 71        | 69          | 217          |
| Age (years), Mean (SD) | 12.1 (3.5) | 12.1 (3.7) | 12.0 (3.4) | 12.1 (3.5) |
| Female, n (%        | 55 (71.4) | 48 (67.6) | 53 (76.8) | 156 (71.9) |
| Male, n (%)         | 22 (28.6) | 23 (32.4) | 16 (23.2) | 61 (28.1) |
| Tanner Score, Median (IQR) | 2 (1 to 4) | 2 (1 to 4) | 2 (1 to 3) | 2 (1 to 4) |
| Steroid Dose, n (%) | 37 (48.1) | 30 (42.3) | 32 (46.4) | 99 (45.6) |
| Steroid Dose, n (%) | 40 (52.0) | 41 (57.8) | 37 (53.6) | 118 (54.4) |
| Etanercept; Infliximab; Anakinra; Tolctizumab | 8 (10.5%) | 17 (23.9%) | 7 (10.1%) | 32 (14.8%) |
| Prior fracture History (Yes, n (%) | 13 (17.1) | 9 (12.7) | 8 (11.6) | 30 (13.8%) |

O42
Reduced joint count for ultrasound assessment in juvenile idiopathic arthritis
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Presenting author: Valentina Litta Modignani
Pediatric Rheumatology 2017, 15(Suppl 1):O42

Introduction: Juvenile idiopathic arthritis (JIA) is a heterogeneous group of arthritides with different prevalence and distribution of joint involvement. Ultrasound (US) is a powerful tool for the assessment of joint disease and has been shown to be more accurate than clinical examination in detecting synovitis. However, assessment of a large number of joints is tedious and time consuming in daily practice, making the feasibility of US questionable.

Objectives: To evaluate the prevalence of clinically active joints in different JIA categories, to propose a core set for reduced joint assessment for US (rUS), and to provide preliminary evidence of validity of rUS.

Methods: The data collected in a large multinational study (the EPOCA study) were analyzed to evaluate the prevalence of active
joints in different ILAR categories of JIA. For each patient, the visit with the highest number of active joints was selected. The 10 most frequently affected joints in the different JIA subsets, and the Juvenile Arthritis Disease Activity Score (JADAS) were calculated. A core set of 6, 8 and 10 joints was considered for inclusion in the rUS and the construct validity of the three rUS was assessed by calculating their Spearman’s correlations with the clinical (3-item) JADAS and with the other JIA outcome measures.

Results: 8,269 JIA patients seen in pediatric rheumatology centers worldwide were included in the study. The table shows the Spearman’s correlations of the three versions of the rUS with the clinical JADAS, the physician’s global assessment of overall disease activity, and the active joint count.

Conclusion: The reduced joint count revealed the ability to serve as surrogate for the whole joint count. The version that includes 6 joints performed similarly to the 8- and 10-joint counts in the assessment of severity of joint disease in all ILAR categories and is proposed as a core set for standard screening in US assessment.

Disclosed Interest
None Declared

Table 13 (abstract O42). See text for description

| Clinical JADAS | Systemic arthritis | RF + polyarthritis | RF arthritis | Oligoarthritis | Psoriatic arthritis | PsA | Undifereniated arthritis |
|----------------|-------------------|--------------------|-------------|----------------|---------------------|-----|-------------------------|
| N 880          | N 346             | N 1014             | N 3516      | N 282          | N 871               | N 457 |
| rUS 6          | 0.77              | 0.82               | 0.77        | 0.73           | 0.70                | 0.65 | 0.69                    |
| rUS 10         | 0.79              | 0.86               | 0.82        | 0.77           | 0.78                | 0.70 | 0.76                    |
| Physician’s global assessment | rUS 6            | 0.74              | 0.73        | 0.72           | 0.75                | 0.67 | 0.62                    | 0.69 |
|                | rUS 10           | 0.75              | 0.76        | 0.76           | 0.77                | 0.74 | 0.65                    | 0.75 |
| Active joint count | rUS 6          | 0.95              | 0.95        | 0.92           | 0.92                | 0.89 | 0.89                    | 0.89 |
|                | rUS 10           | 1                 | 1           | 1              | 1                   | 1    | 1                       | 1    |

Poster Session: Autoinflammatory diseases I

P1

The impact of gastrointestinal clinical manifestations in autoinflammatory diseases (AIDS): lessons from the international Eurofever registry

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Presenting author: Alessia Omenetti
Pediatric Rheumatology 2017, 15(Suppl 1):P1

Introduction: The spectrum of autoinflammatory diseases (AIDs) is expanding and delay in diagnosis may occur if those rare entities are not promptly identified.

Objectives: To assess the impact and significance of gastrointestinal (GI) clinical manifestations in a broad spectrum of patients with primary diagnosis of AIDs.

Methods: The Eurofever registry containing data retrospectively collected about patients with primary diagnosis of AIDs enrolled by 108 participating centres (Nov 2009-Apr 2016) was evaluated. Demographic, genetic and clinical features were analyzed by filtering for GI symptoms including vomiting, abdominal pain, constipation, diarrhea, GI ulcers, anal/perianal ulcers, GI bleeding and aseptic peritonitis. When present, each manifestation was ranked by its clinical significance and classified as grade 1 or grade 2 (i.e. occasionally or persistently present during flares, respectively). Development of complications such as gut perforation, peritoneal adhesions, occlusions/sub-occlusions or any, if any, were also assessed.

Results: Total 3322 patients affected by AIDs including Bechet (n = 208, 6.26%), Blau (n = 49, 1.47%), CANDLE (n = 1, 0.03%), CAPS (n = 263, 7.91%), CRMO (n = 486, 14.62%), DADADA2 (n = 2, 0.06%), DIRA (n = 3, 0.09%), FMF (n = 905, 27.24%), Majeed (N = 3, 0.09%) MKD (n = 192, 5.77%), NLRP12 (n = 13, 0.39%), PAPA (n = 28, 0.84%), PFPA (n = 631, 18.99%), Majeed (n = 3, 0.09%), SAVI (n = 1, 0.03%), TRAPS (n = 261, 7.85%), undefined (n = 276, 8.30%) were evaluated. At least 1 GI manifestation was reported in 1570 patients (47.2%)(MF = 1:1) including 83 Bechet, 7 Blau, 1 CANDLE, 61 CAPS, 38 CRMO, 1 DADADA2, 2 DIRA, 676 FMF, 1 Majeed, 168 MKD, 2 NLRP12, 4 PAPA, 229 PFPA, 160 TRAPS, and 137 undefined AIDs. Among patients displaying GI manifestations, symptoms were reported as grade 2 in 731 cases (46.5%) and severe GI complications occurred in 107 cases (6.81%). Namely, 20 patients (1.27%) suffered from gut perforation (i.e. 1 CAPS, 7 FMF, 3 MKV, 1 PFTA, 1 DIRA, 1 CRMO, 3 Bechet, 1 PAPA, 2 undefined), while 49 subjects (3.1%) were affected by peritoneal adhesions (i.e. 3 CAPS, 27 FMF, 2 TRAPS, 6 MKV, 1 PFTA, 1 DIRA, 1 CRMO, 3 Bechet, 1 PAPA, 4 undefined), and 38 patients (2.4%) developed occlusions/sub-occlusions (i.e. 1 CAPS, 17 FMF, 4 TRAPS, 5 MKV, 1 PFTA, 1 DIRA, 1 CRMO, 3 Bechet, 1 PAPA, 4 undefined). Finally, secondary diagnosis of GI disease was found in 5 subjects (0.3%) (i.e. gastroodventitis, IBD, coeliac disease, eosinophilic colitis) and hepatomegaly was reported in 154 patients (9.8%).

Conclusion: Although descriptive, these data unveiled the unforeseen significance of GI involvement in AIDs. These findings have threefold implications in the daily clinical management of AIDs. 1) GI specialists should be aware of these rare diseases in order to avoid delayed AID diagnosis in case patients come to their attention first; 2) pediatric rheumatologists and GI specialists should develop standardized work-up protocols in order to monitor these signs and prevent potential complications; 3) these data strongly address the need of a prospective analysis in order to delineate the disease history and potential evolutions (if any) into more defined GI diseases in the context of autoinflammation.

Disclosed Interest
A. Omenetti: None Declared, J. Frenkel Grant / Research Support from: unrestricted grant from Novartis and SOBI, H. Lachmann Grant / Research Support from: unrestricted grant from Novartis and SOBI, S. Ozen Grant / Research Support from: unrestricted grant from Novartis and SOBI, N. Ruperto Grant / Research Support from: unrestricted grant from Novartis and SOBI, M. Gattorno Grant / Research Support from: unrestricted grant from Novartis and SOBI

P2

Variable clinical phenotypes and relation of interferon signature with disease activity in ADA 2 deficiency

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Presenting author: Antonella Insalaco
Pediatric Rheumatology 2017, 15(Suppl 1):P2

Introduction: The deficiency of adenosine deaminase 2 (ADA2) is a recently described autosomal recessive autoinflammatory disease, caused by mutations of CECR1 and characterized by early onset vasculopathy with livedoid skin associated to systemic inflammation. In some patients, the disease is mild and skin-limited, in others is severe, with multi-organ involvement including ischemic or hemorrhagic strokes. In some ADA2 patients a mild immunodeficiency was detected involving adaptive immunity. TNF inhibitors are very efficacious. Recently, an upregulation of type I interferon-stimulated gene transcripts, so-called interferon signature, was described also in two ADA2 patients.
**P3**

**Association between familial Mediterranean fever and spondyloarthritis. Clinical characteristics of the French ceremai Pediatric center**

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**Presenting author:** Bilade Cherqaoui

Pediatric Rheumatology 2017, 15(Suppl 1):P3

**Introduction:** Spondyloarthritis (SpA) is more frequently observed in familial Mediterranean fever (FMF), and possibly related to MEFV mutations in these patients. No studies to date report this association in FMF-FSPA patients. We screened 16 FMF patients with SpA (FMF-SpA) and compared them to 20 FMF patients without SpA (FMF-ASAS) in a French pediatric center.

**Methods:** We screened FMF patients seen at Bicêtre hospital, a tertiary referral center. FMF was defined according to either Tel-Hashomer or Turkish pediatric criteria, with at least 1 exacerbation of MEFV gene. J-SpA was defined according to ILAR criteria of enthesitis-related psoriatic juvenile arthritis; we also assessed the performance of ASAS criteria. FMF patients with SpA (FMF-SpA) were then compared to those with typical FMF and to 20 patients with J-SpA.

**Results:** Among 125 screened files, 99 fulfilled criteria of FMF. Among FMF patients, 16 fulfilled criteria of J-SpA (16%). The demographic and genetic features of FMF-SpA patients were: 1.6 girls/1 boy, median age 7.5 years, frequency of abdominal pain and arthralgia/plantalgia (OR = 34.3 and 44.4, p < 0.001) in FMF-SpA group. No significant biological or MEFV genotypes differences were observed. FMF-SpA patients received more frequently anti-IL1 drugs after failure of colchicine (OR = 42, p = 0.04). Articular features seemed to respond at least partially to anti-IL1 drugs. When comparing the 16 FMF-SpA patients with 20 typical J-SpA patients, we observed: lower age at onset of articular signs (4.5±1.1 vs 9±0.6 years respectively, p = 0.001), higher frequency of plantar pain (OR = 31.6, p < 0.001), less frequent synovitis on ultrasound (p = 0.03) in FMF-SpA group. Non-steroidal anti-inflammatory and anti-TNF drugs were less frequently used in FMF-SpA patients (p = 0.008, <0.001 respectively). ILAR criteria were more sensitive and less specific than ASASs in FMF-SpA and J-SpA groups (92.3/57.1% and 100/60% respectively).

**Conclusion:** A non- incidental association of FMF and J-SpA was observed in this French cohort, in accordance with data observed in Middle East (Barut K, et al. Articular involvement in childhood FMF. Pediatr Rheumatol 2015). Frequency of this association reached 16%, a bit higher than previously defined. Compared to typical FMF, SpA-FMF patients present a predominant articular/enthesitis pattern, less febrile crises, and more frequent resistance to colchicine. Except earlier age at onset, more frequent plantar involvement and less frequent ultrasound synovitis, SpA-FMF features were virtually similar to J-SpA. Thus FMF-FMF related arthritis should be followed and treated as J-SpA, including anti-TNF drug use.

**Disclosure of Interest** None Declared

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**P4**

Health-related quality of life (HRQOL) of children with PFAPA syndrome

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**Presenting author:** Claire Grimwood

Pediatric Rheumatology 2017, 15(Suppl 1):P4

**Introduction:** PFAPA syndrome (Periodic Fever, Aphthous stomatitis, Pharyngitis and Adenitis) is generally considered to be a benign disease compared to other autoinflammatory syndromes because it normally disappears before adulthood. However fever episodes may...
have a huge impact on daily activities, schooling and family functioning.

Objectives: To describe and compare the physical and psychosocial HRQOL of children with PFAPA compared to FMF peers using a multidimensional, well validated, and reliable HRQOL instrument.

Methods: Thirty three voluntary PFAPA patients attending the French reference centre for autoinflammatory diseases (CeRéMAI) during the year 2015 and having an active disease were included in the study. The control group consisted of 27 FMF patients age matched, attending the autoinflammatory clinic during the same period. All the subjects and/or their parents of the study were asked to complete the age appropriate questionnaire of the Pediatric Quality of Life Inventory TM 4.0 (PedsQL™ 4.0) Generic Core Scale. Patients and controls were grouped according to their ages as follows: pre-school age children (2–7 years) and school-age children and youth (8–18 years).

Results: PedsQL™ self-report scores of pre-school age children (2–7 years) with PFAPA were significantly lower than FMF peers for general quality of life, physical and psychosocial functioning (Table). The parent proxy-report did not find a significant difference even though scores were systematically lower in PFAPA patients. The parent proxy-report and child self-reported PedsQL™ scores of school age children and adolescent patients (8–18 years) with PFAPA were lower than the FMF group for general quality of life, physical and psychosocial functioning; however the difference was not significant.

Conclusion: HRQOL in PFAPA children seems to be lower than in FMF peers while the latter are known to have impaired QOL if compared to the general population.

Disclosure of Interest
None Declared

Table 14 (abstract P4). Parent proxy-reported and child self-reported PedsQL™ scores of pediatric patients with

| Scale                          | PFAPA patients | FMF controls | P       |
|-------------------------------|----------------|--------------|---------|
|                               | Number | Mean         | Number | Mean         |       |
| Pre-school age children (2–7 years) |       |              |        |              |       |
| Self report: Total score       | 15     | 66.6         | 8      | 80.3         | 0.01   |
| Proxi-report: Total score      | 23     | 68.9         | 14     | 76.9         | 0.12   |
| School age children and youth (8–18 years) |       |              |        |              |       |
| Self report: Total score       | 8      | 58           | 13     | 74           | 0.06   |
| Proxi-report: Total score      | 10     | 57.4         | 13     | 72.3         | 0.08   |

P5
Cryopyrin associated periodic syndromes in Italian patients: evaluation of the rate of NLRP3 mosaicism and search for novel genes
Denise Lasiglié1, Denise Ferrera1, Giulia Amico1, Marco Di Duca1, Roberta Caorsi1, Loredana Lepore2, Antonella Insalaco3, Marco Cattalini4, Laura Obici5, Rita Consolini6, Roberto Ravazzolo6, Alberto Martini6, Isabella Ceccherini6, Ryuta Nishikomori7, Juan Arostegui8, Marco Gattorno1, Silvia Borghini1, Denise Ferrera1, Giulia Amico1, Marco Di Duca1, Roberta Caorsi1, Loredana Lepore2, Antonella Insalaco3, Marco Cattalini4, Laura Obici5, Rita Consolini6, Roberto Ravazzolo6, Alberto Martini6, Isabella Ceccherini6, Ryuta Nishikomori7, Juan Arostegui8, Marco Gattorno1, Silvia Borghini1

Introduction: Most patients affected with Cryopyrin associated periodic syndromes (CAPS) bear a heterozygous mutation in NLRP3 gene, a little proportion of which in mosaic status at level not evaluable by Sanger sequencing. Nevertheless, a proportion of patients with a clinical phenotype of CAPS, particularly Muckle Wells Syndrome and Familial Cold Autoinflammatory Syndrome, remain without a genetic diagnosis.

Objectives: (1) Evaluation of the rate of NLRP3 mosaicism in Italian patients previously described as mutation negative (2) Identification of novel genes responsible for CAPS symptoms (3) Providing patients with a correct genetic diagnosis and a risk recurrence

Methods: Patients with a clinical and immunological phenotype resembling CAPS phenotypes and resulted negative to the screening of the total coding sequence of NLRP3 gene by Sanger screening have been enrolled in this study. Patients DNA have been subjected to (i) Next generation sequencing at high coverage of NLRP3 gene to identify mutations in mosaic status and, in negative cases, to (ii) whole exome sequencing (WES) for novel genes discovery.

Results: Massively parallel DNA sequencing revealed somatic NLRP3 mosaicism in four patients: three affected with CINCA (100%) and one with MWS (12.5%). Identified nucleotide substitutions are all located in the exon three of NLRP3 and encode for four different amino-acid changes, two of them being novel (p.G564S and p.Y563C) and remaining already reported (p.R260P as full mutation and p.T433I in mosaic state). None of the detected variants was reported in about 63000 genomes at Exac Browser Consortium. Compared to wild type NLRP3, mutant proteins rapidly induce necrotic cell death when transiently transfected in human monocyte THP-1 cells and cause enhanced NF-kB reporter activity after co-expression with its partner ASC in HEK293FT. In almost all cases, mutated alleles have been identified not only in all the cell population from the blood but also in ectodermal tissues, suggesting that the event has been established very early in development.

DNA from the remaining patients have been subjected to WES in search of novel candidate genes. Unfortunately, same SNPs or different variants at the same genes have not been identified in at least two different patients. Subsequent analysis has been conducted considering single family or trio. A de novo mutation has been identified in a transcription factor belonging to the FOX gene family. Preliminary functional tests confirm a causative role of this amino acid substitution. In a family, the disease segregates with a novel variant in a protein sensor of viral RNA. Finally, in a patient an immune receptor displays impaired membrane expression in white blood cells, due to the coupling of an allele with low frequency from the mother and a non-functional one from the father.

Conclusion: CAPS are widely associated with mutations in the NLRP3 gene, being mutated in mosaic status in the most severe patients, and only a small percentage to private mutations that arise in individual patients. Although these changes have apparently a very rare frequency, only an analysis of the genes identified in this study in more patients with CAPS resembling symptoms can definitively assess their actual impact.

Disclosure of Interest
None Declared

P6
Monocytes proteomic profile of patients with different autoinflammatory diseases: an approach to identify new biomarkers
Federica Penco1, Andrea Petretto2, Chiara Lavarello2, Etriva Inglese2, Alessia Omenetti1, Martina Finetti1, Claudia Pastorino1, Arina Bertoni1, Marco Gattorno1, Marco Mattiaccia1,2,3,4

Presenting author: Federica Penco

Introduction: Autoinflammatory diseases are a group of inherited disorders characterized by early onset and systemic inflammation, often manifesting with unexplained fevers. These pathologies are usually caused by mutations in genes involved in the regulation of innate immune response with consequent inflammatory phenotype driven by activation of monocytes, macrophages and granulocytes. Part of these pathologies are however genetically undefined.
Objectives: Our aim is to evaluate the differences in the expression of proteins or pathway in monocytes, untreated or treated with LPS, of patients with autoinflammatory diseases and healthy subjects in order to clusterize the different diseases and better characterize the genetically undefined pathologies.

Methods: Monocytes, purified from peripheral blood of patients and healthy subjects with CD14 microbeads positive selection, were collected and incubated for 4 hours with or without LPS stimulation. Cells were lysed and prepared to be analyzed at MS. The samples were processed by iST protocol and protein expression evaluated by an High Resolution/Mass Accuracy Liquid Chromatography Tandem Mass Spectrometry (HR/MA LC MS/MS). Each sample was run at least in triplicate. PCA analysis and Person’s correlation were used as quality control of experimental design, while the statistical analysis was performed with the Perseus software.

Results: We analyze the monocytes of patients with CAPS, TRAPS and FMF (each group compared with healthy subjects). We have identified about 4000 proteins of each 3500 are quantified by LFQ approach. PCA analysis and Person’s correlation show a good reproducibility of data and a good separation between the different groups. In synchronous way using a cluster analysis and heatmap, based on a machine learning protein selection, we observe a protein signature specific for each group of pathology and for each condition of monocytes treatment.

Conclusion: Here, we addressed how an high resolution proteomics approach could be used to better understand the biology of autoinflammatory diseases. The characterization of a broad spectrum of proteins and their interaction network will allow us to identify new biomarkers for the different pathologies and to better comprehend and recognize the genetically undefined disorders.

Disclosure of Interest
None Declared

P7
Results of the Eurofever Delphi survey for the classification criteria of PFAPA syndrome
Federica Vanoni1, Silvia Federici2, Seza Ozen3, Joost Frenkel5, Helen Lachmann2, Alberto Martini1, Nicola Ruperto1, Marco Gattorno3, Michæll Hofer1,6, and on behalf of EUROFEVER PROJECT
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Presenting author: Federica Vanoni
Pediatric Rheumatology 2017, 15(Suppl 1):P7

Introduction: Diagnosis of Periodic fever, aphthous stomatitis, pharyngitis and cervical adenitis (PFAPA) is currently based on a set of criteria proposed in 1999 modified from Marshall’s criteria. Nevertheless no validated evidence based set of classification criteria for PFAPA has been established so far.

Objectives: To identify evidence based candidate classification criteria for PFAPA syndrome using the consensus formation techniques (Delphi and Nominal Group Techniques).

Methods: Two following Delphi surveys were sent to health professionals working in the field of autoinflammation. In the first open survey 124 experts belonging to the PRINTO Network were asked to list as many variables they wanted they consider as important for the diagnosis of patients with PFAPA. The variables may be of any type (clinical, laboratory, instrumental variables, genetic tests etc.). In the second survey 162 experts (also American experts were involved in this phase) were asked to select, from a list of items coming from the first survey, the 10 top variables and to rank them by assigning a score from 10 to 1 in order of importance. There was no possibility to assign the same rank to multiple variables. This process has been conducted in parallel with an analogous process for the monogenic periodic fevers.

Results: The overall rate of response to the first and second Delphi, including monogenic periodic fevers, was respectively 86% (107/124 experts) and 87% (141/162 experts). For PFAPA, the rate of response to the first and second Delphi was respectively 55% (68/124 experts) and 70% (101/162 experts). We obtained 92 variables from the first survey among which 62 were selected in the second survey. Table 15 shows the total score obtained and the rate of selection of the top variables (3rd quartile).

Conclusion: Our process led to the identification of those features that are considered to be the most important as candidate variable to be included in a new set of evidence based classification criteria for PFAPA. The next step of this project (now ongoing) is the evaluation by a panel of experts of 360 real patients affected by AIDs (PFAPA, monogenic periodic fever and undefined periodic fever) randomly selected from EUROFEVER Registry. Those patients reaching the 80% of consensus among experts in the evaluation process will be used for the subsequent statistical analysis aimed at identified the best evidence-based classification criteria (in term of sensitivity and specificity) for PFAPA. The process will end with a consensus conference that will be held in Genoa on March 16-18, 2017.

Disclosure of Interest
None Declared

Table 15 (abstract P7). See text for description

| Rank | Variable | Global Score | Votes (n) |
|------|----------|--------------|-----------|
| 1    | Regular periodicity                  | 436          | 56        |
| 2    | Aphthous stomatitis                | 431          | 77        |
| 3    | Response to steroid                | 401          | 66        |
| 4    | Cervical adenitis                  | 368          | 72        |
| 5    | Well being between flares          | 299          | 57        |
| 6    | Pharyngitis (exudative or not)     | 288          | 47        |
| 7    | Increase of acute phase reactants and serum amyloid A during fever episodes | 271 | 44 |
| 8    | Normal growth/development          | 236          | 51        |
| 9    | Pharingotonsillitis                | 228          | 35        |
| 10   | Periodic fever 3–5 days            | 202          | 24        |
| 11   | Periodic fever 3–6 days            | 202          | 23        |
| 12   | Self limiting episodes             | 183          | 35        |
| 13   | Response to tonsillectomy          | 182          | 33        |
| 14   | Improvement with age               | 160          | 40        |
| 15   | Exclusion cyclic neutropenia/immunodeficiency | 150 | 34 |
| 16   | Normalization of acute phase reactants in well-being | 146 | 33 |
| 17   | Recurrence every 3–6 weeks         | 145          | 21        |

P8
Safety and efficacy of long-term canakinumab therapy in patients with cap: final results from β-confident registry
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Presenting author: J.B. Kuenmerle-Deschner
Pediatric Rheumatology 2017, 15(Suppl 1):P8

This abstract is not included here as it has already been published.
P9
Development and validation of diagnostic criteria for cryopyrin associated periodic syndromes
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Pediatric Rheumatology 2017, 15(Suppl 1)P9
This abstract is not included here as it has already been published.

P10
Application of Gaslini Diagnostic Score in Japanese children with periodic fever
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Pediatric Rheumatology 2017, 15(Suppl 1)P10
Introduction: The Gaslini Diagnostic Score (GDS) is a differential diagnostic score intended to determine a patient’s risk of having genetic factors contributing to PFAPA syndrome (periodic fevers with aphthous stomatitis, pharyngitis, and adenitis). The GDS is calculated based on the following criteria: age at onset, positive family history of periodic fever, chest pain, abdominal pain, and diarrhea. While the presence of any of the clinical symptoms is associated with an increased likelihood of a genetic basis, a child with a GDS of 1.32 or higher is considered likely to be at high risk of having a hereditary periodic fever syndrome. When applied in European countries, the GDS showed very high sensitivity (>90%) and a specificity of 59% to 82%.
Objectives: To investigate the utility of GDS in Japanese children diagnosed with PFAPA.
Methods: Children 5 years old or younger who had been diagnosed with periodic fever were enrolled in this study. The observed clinical symptoms for each subject were used in determining his or her GDS, in order to compare the score to the presence or absence of polymorphisms in 10 genes that are frequently implicated in autoinflammatory diseases. Each patient’s genomic DNA was isolated from peripheral blood and subjected to polymerase chain reaction (PCR) to amplify selected exons of the genes of interest. The three genes most frequently reported as being associated with PFAPA (and the exons amplified for each in this study) include: pyrin/marenostrin (MEFV, exons 1 to 10), tumor necrosis factor receptor superfamily member 1A (TNFRSF1A, exons 2, 3, 4), and mevalonate kinase (MKV, exons 9, 10, 11). The PCR products were purified, sequenced using the BigDye® Terminator v3.1 Cycle Sequencing Kit (ThermoFisher Scientific, Waltham, Mass, USA), and analyzed using an ABI 3130xl Prism Genetic Analyzer (Applied Biosystems, Inc., Foster City, Calif, USA).
Results: Of the 35 children enrolled (23 boys and 12 girls, with an average age at onset of 36.2 ± 18.4 months), 42.8% developed aphthous stomatitis, 34.3% reported stomachache, 11.4% had diarrhea, and 11.4% reported pectoralgia. Thirteen (37.1%) had a family history of periodic fever. A GDS score higher than 1.32 was seen in only 5 patients. However, 2 of those 5 had MEFV E148Q hetero, a fairly common polymorphism, found in 20% to 30% of healthy Japanese people. Of the 5, 2 other patients had MEFV polymorphisms (E148Q/P369S/R408Q hetero, E148Q homo). Of the 30 subjects with a GDS lower than 1.32, 13 showed no polymorphisms, while the other 17 patients showed MEFV polymorphisms (6 E148Q heteros, 4 E148Q/P369S/R408Q heteros, 3 E148Q/L110P heteros, 2 R202 heteros, 1 P369S/R408Q homo, 1 E84K hetero). Two patients with high GDS improved after administration of colchicine or prednisolone during an acute attack. Four patients with low GDS were clinically diagnosed with Familial Mediterranean fever, based on efficacy of colchicine. Conclusion: Application of the GDS in the Japanese children with PFAPA revealed low sensitivity (15%) and comparatively high specificity (86%), excluding E148Q heteros. Therefore, we concluded that the GDS is not an informative diagnostic tool for Japanese PFAPA patients.
Disclosure of Interest: None Declared

P11
Our experience in the treatment of PAPA syndrome
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Pediatric Rheumatology 2017, 15(Suppl 1)P11
Introduction: PAPA syndrome, is a rare autosomal dominant autoinflammatory disorder and its management still represents a challenge.
Objectives: Our goal is to present our only experience in the treatment of PAPA syndrome.
Methods: We report the case of a 13 year old boy who was referred to our hospital with two weeks history of swollen and painful left knee and right ankle. Since the ages of 2 he was treated in another hospital for recurrent sterile arthritis and skin ulcer; last two years he was treated with Etanercept. His first relative suffers from recurrent sterile arthritis, skin ulcers and acnae. Although genetic analysis not made, arthritis of boy was suspected associated with PAPA syndrome. On examination, left knee, right ankle were swollen, painful with limitation of movement, right elbow with the flexion contracture. A rough scars on the skin above elbows, ankles, right knee, lower legs and on the back were present. Laboratory findings were abnormal with the ESR of 110 mm/h, CRP of 90.0 mg/l, mild anemia, and sterile culture of an aspirate from left knee.
Results: The boy responded well to start treatment with Infliximab, pulses and intra-arterial steroids, in terms that after a few days the swelling and pain diminished, movements was better also; in addition, laboratory results returned to normal. In the following months the disease was partially controlled considering that after 7 months of treatment the boy had skin manifestations (ulcer and milde acne), so tetracycline were added. After 16 months achieved complete response but he developed generalized urticaria during of administration of Infliximab. In coming weeks he had flares of arthritis and skin manifestations, elevated values of ESR, CRP, along with sterile cultures of an aspirate from left knee.

P12
Methotrexate and Adalimumab 40 mg every 14 days. After administration of Infliximab. In coming weeks he had flares of arthritis and skin manifestations, elevated values of ESR, CRP, along with sterile culture of an aspirate from left knee.
treatment the boy was without skin manifestations but with occasional manifestations of arthritis who were treated by intra-articular steroids.

Conclusion: Our observations indicate that treatment with infliximab, if tolerated, may be more effective than adalimumab, although adalimumab had better effects on the skin manifestations.

Disclosure of Interest
None Declared

P12

Enterococcalis and recurrent macrophage activation syndrome in a patient with G786V mutation of NLR4C gene
Anna Kozlova, Irina Mersijanova, Mariya Panina, Lily Hachtryan, Vasily Burlakov, Elena Raikina, Alexey Maschan, Anna Shcherbina

Introduction: Autoinflammatory disorders comprise heterogeneous group of diseases characterized by chronic or recurrent systemic sterile inflammation as a result of genetic defects of regulators of inflammation. The NLR4C is a member of the NOD-like receptor family of intracellular sensors that has a pivotal role in antimicrobial defense via intracellular sensing of a wide range of pathogens. NLR4C is activated by NALP-mediated recognition of flagellin or components of bacterial type 3 secretion systems (TTSS) in the host cell cytosol, leading to the assembly of inflammasome that mediates proteolytic cleavage of immature pro-IL-1β and pro-IL-18 and release of their bioactive forms into the extracellular space. Heterozygous mutations of NLR4C gene lead to autoinflammation with infantile enterocolitis (AIEFEC) syndrome. AIEFEC symptoms include early onset growth retardation, enterocolitis, splenomegaly and fatal or near fatal episodes of systemic inflammation with macrophage activation.

Objectives: We report a patient with clinical picture typical of NLR4C defect.

Methods: Case report.

Results: An index case is a 3 years old female with symptoms similar to AIEFEC syndrome: episodes of fever, rash, enterocolitis, arthritis, hepatosplenomegaly, lymphadenopathy and laboratory changes (elevated ESR, CRP, hypergammaglobulinemia, anemia, thrombocytosis/ thrombocytopenia, leukocytosis/leukopenia, high level of ferritin and LDG) since infancy. Mutations in the genes typical for some autoinflammatory disorders (MEFV, MVK, TNFRSF1A, NLRP3), familiar hemoglobinocytosis (SH2D1A, PRF1, STX11, STXB2), as well as FAS, KRAS, Nras, PTPN11, CBL were not found. The mutation c. 2357G > C (G786V) of NLR4C gene, which has been described as polymorphism, but also described in an AIEFEC-like patient, was found. Healthy father is a carrier of the same mutation. Prednisone therapy didn't alleviate symptoms. Treatment according to HLH-2004 protocol (dexamethasone, etoposide, cyclosporin A) and rituximab led to a temporary remission, with relapse of symptoms upon protocol completion. IL1 inhibitor (Anakinra, 5 mg/kg) therapy led to complete resolution of the symptoms.

Conclusion: In the light of patient's clinical phenotype consistent with AIEFEC the pathogenic role the NLR4C c. 2357G>C variant requires further investigation. The patient is currently submitted to WES to rule out other genetic mechanisms of her disease.

Disclosure of Interest
None Declared

P13

Decreased vitamin B12 levels in children with familial Mediterranean fever
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Pediatric Rheumatology 2017, 15(Suppl 1):P13

Introduction: Familial Mediterranean fever (FMF), the most common autoinflammatory hereditary fever disease, is characterized by paroxysmal attacks of serosal inflammation. Colchicine is highly effective in preventing these attacks but it may also disrupt the intestinal absorption of vitamin B12.

Objectives: We hypothesized that patients treated with colchicine for a prolonged period could develop deficiency of the vitamin. Symptoms of B12 deficiency are confused with some findings of FMF if the elusiveness to investigate vitamin B12 deficiency.

Methods: Twenty FMF patients on regular colchicine treatment for at least 1 year and age and sex-matched 20 healthy controls were enrolled. Weakness, tiredness, a loss of appetite, nerve problems (numbness or tingling, muscle weakness, and problems walking) mental problems like memory loss, or behavioral changes were recorded. Complete blood count, peripheral blood smear, vitamin B12, homocysteine and folic acid were measured in each child. FMF patients diagnosed with vitamin B12 deficiency were given regular vitamin B12 treatment for a month. At the end of treatment complaints of patients were re-evaluated.

Results: Demographic, clinical, and laboratory features of the patients are shown in Table 16. The median hemoglobin, folic acid levels were not significantly different between the groups. Prevalence of B12 deficiency among patients with FMF was 35% compared with 15% in healthy controls. The median vitamin B12 values were not significantly different between the groups 310.41 (SD = 158.22) pg/mL vs. 351.83 (SD = 170.41) pg/mL, p = 0.67). At the beginning of vitamin B12 treatment, the median hemoglobin level was 11.3 g/dL, which increased to 12.4 g/dL at the end of the first month of vitamin B12 treatment. After the treatment of vitamin B12, arthralgia (25%), tiredness (10%) and memory loss (10%) declined in patients with FMF.

Conclusion: Vitamin B12 deficiency is common in patients with FMF. Although chronic arthralgia and fatigue are classic symptoms of FMF, they can also be the result of a vitamin B12 deficiency. The decision on an inadequate dosage of colchicine should not be drawn too hastily, reasons such as vitamin B12 deficiency, which can cause the patient's symptoms should be considered.

Disclosure of Interest
None Declared

Table 16 (abstract P13). Demographic, clinical, and laboratory features of the patients

| Characteristics                  | FMF              | Control group |
|----------------------------------|------------------|---------------|
| Age (years)                      | 8.2 ± 2.72       | 8.5 ± 2.85    |
| Arthralgia                       | 8 (40)           | 3 (15)        |
| Limb pain with exercise / Heel pain| 1 (5)/2 (10)   | 0 /0          |
| Myalgia / Weakness               | 0/0              | 1 (5)/0       |
| Tiredness                        | 4 (20)           | 2 (10)        |
| A loss of appetite               | 3 (15)           | 1 (5)         |
| Mental problems like memory loss | 4 (20)           | 1 (5)         |
| or behavioral changes            | 351.83           |               |
| Vitamin B12 level (median-pg/ml) | 310.41           |               |
| Folic acid (median-ng/ml)        | 8.9 ± 7.2        | 10.3 ± 5.6    |

P14

Canakinumab-treatment in autoinflammatory disease patients younger than 2 years old
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Pediatric Rheumatology 2017, 15(Suppl 1):P14

Introduction: Twenty FMF patients on regular colchicine treatment for at least 1 year and age and sex-matched 20 healthy controls were enrolled. Weakness, tiredness, a loss of appetite, nerve problems (numbness or tingling, muscle weakness, and problems walking) mental problems like memory loss, or behavioral changes were recorded. Complete blood count, peripheral blood smear, vitamin B12, homocysteine and folic acid were measured in each child. FMF patients diagnosed with vitamin B12 deficiency were given regular vitamin B12 treatment for a month. At the end of treatment complaints of patients were re-evaluated.

Results: Demographic, clinical, and laboratory features of the patients are shown in Table 16. The median hemoglobin, folic acid levels were not significantly different between the groups. Prevalence of B12 deficiency among patients with FMF was 35% compared with 15% in healthy controls. The median vitamin B12 values were not significantly different between the groups 310.41 (SD = 158.22) pg/mL vs. 351.83 (SD = 170.41) pg/mL, p = 0.67). At the beginning of vitamin B12 treatment, the median hemoglobin level was 11.3 g/dL, which increased to 12.4 g/dL at the end of the first month of vitamin B12 treatment. After the treatment of vitamin B12, arthralgia (25%), tiredness (10%) and memory loss (10%) declined in patients with FMF.

Conclusion: Vitamin B12 deficiency is common in patients with FMF. Although chronic arthralgia and fatigue are classic symptoms of FMF, they can also be the result of a vitamin B12 deficiency. The decision on an inadequate dosage of colchicine should not be drawn too hastily, reasons such as vitamin B12 deficiency, which can cause the patient's symptoms should be considered.

Disclosure of Interest
None Declared
Introduction: The proinflammatory cytokine IL-1β has significantly increased the understanding of the pathogenesis of many autoinflammatory diseases (AID), including cryopyrin-associated periodic syndromes (CAPS), hyper IgD syndrome (HIDS), familial Mediterranean fever (FMF), and systemic onset juvenile idiopathic arthritis (sJIA). Canakinumab is a fully humanized monoclonal antibody (mAb) specific for IL-1β that has been indicated for a wide range of inflammatory disorders. Based on data from the treatment-withdrawal phase III trial, canakinumab was approved for use for the treatment of children aged >4 years for CAPS and >2 years for SJA indications.

Objectives: The aim of this case report is to verify the efficacy and safety of canakinumab in patients younger than 2 years old.

Methods: Eight unrelated AID patients (4 sJIA, 2 NOMID/CINCA, 1 HIDS and 1 FMF and inflammatory bowel disease) are reported from two pediatric rheumatology centers. All patients were younger 24 months old and obtained their drugs off-label by approval of Ministry of Health.

Results: The results of eight patients (5 female, 3 male) are analyzed in this study. All patients were younger 24 months old (9 to 23 months; mean age = 18.6 months), and had canakinumab (3 mg/kg up to 6 mg/kg) s.c. every 4-8 weeks. Six out of eight patients were previously treated with anakinra for a median period of 55 days (range 1 to 150 days). The median follow up time was 20 months. Seven patients achieved a complete response while one patient was switched due to adverse event. Of the seven patients with the sJIA, HIDS and NOMID/CINCA, four patients (1,2,4 and 5) did not require any adjustment of the therapy due to a persistent optimal control of both clinical and laboratory parameters during the whole follow-up period. Three others required at least one modification of the treatment schedule due to a persistent elevation of acute phase reactants. Compliance with treatment was good, and no reactions at the site of the injection were recorded. A patient (number 2) had pneumonia at month 20 who treated ambulatory with antibiotic. The administration of canakinumab was postponed until the complete resolution of the infection, which occurred without complications.

Conclusion: In this case series, we report the response to canakinumab in a AID pediatric patients. Canakinumab was an effective treatment for patients with ADIs younger than 24 months old and well tolerated.

Disclosure of Interest
None Declared

P15 Response of pamidronate treatment assessed by whole body magnetic resonance imaging in chronic non-bacterial osteomyelitis
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Pediatric Rheumatology 2017, 15(Suppl 1):P15

Introduction: Pamidronate (PAM) may be effective to diminish pain and bone changes in chronic non-bacterial osteomyelitis (CNO). Whole body magnetic resonance imaging (WB MRI) can be used to assess bone involvement.

Objectives: To evaluate clinical and radiographic responses to PAM treatment as assessed by WB MRI and clinical response one and two year post-treatment.

Methods: We did a retrospective evaluation of children diagnosed with CNO according to Bristol criteria (ref.) and who had an insufficient response to NSAID treatment. Children were treated with i.v. PAM 1 mg/kg (max 60 mg/day) on 3 consecutive days every 3 months. Seventeen children were treated for one year (8 girls and 9 boys) median age 11 years (range 5-13). All children were scheduled for two years of treatment. Data were available in 17 children after one year of treatment and nine children after two years of treatment.

WB MRI (1,5 Tesla) short tau inversion recovery (STIR) and T1-weighted images were performed at baseline (median -1 month, range -6-0), after one year (median 12 months, range 9-19) and two years (median 24 months, range 22-26). Clinical assessments, medical history and inflammatory biochemistry were obtained. For data not following a normal distribution Wilcoxon rank signed tests were used to assess change in number of bone lesions.

The study was performed according to Danish law after approval by institutional innovation board.

Results: Six children had elevated sedimentation rate at baseline, all normalized during the first year. Comorbidities were JIA (n = 3) and IBD (n = 1) no patients had psoriasis. All children were treated with NSAID as first line treatment and further medical history was antibiotics 6%, methotrexate 41%, steroids 41%, anti-TNFα treatment before PAM 18% and 18% had anti-TNFα treatment added during PAM due to persistent symptoms and progression on WB MRI. The total number of lesions pre-treatment was 112, median 6 (range 2-14) and median number of clinical symptomatic lesions was 3 (range 1-8). Seven children had axial lesions median 1 (range 1-4) and six had lesions in relation to SI-joint median 1 (range 1-2). There were no erosive changes in the SI- joints. After one year the total number of lesions was 71, median 2 (range 0-21), seven had no lesions. Ten children had new or persisting lesions median 4 (range 1-15). One active spinal lesion and one SI-joint lesion persisted. Two children had deformation of vertebrae without persistent sign of inflammation on STIR-imaging. None of the data followed a normal distribution. There was a statistically significant change in number of bone lesions from baseline to year one (p = 0.047), particularly change in number of spinal lesions from baseline to year one (p = 0.016). The clinical response after one year was complete pain relief 41%, partial pain relief 48% and no pain relief 11%. In six children the lesions resolved completely as assessed by WB MRI, but pain was still reported.

We evaluated preliminary data of 9 children after two years treatment. Five children remained without bone lesions and four children progressed during treatment with a median value of 3 new or worsening of lesions (range 1-10) as assessed by MRI.

Conclusion: PAM is a potent second-line treatment for CNO after one year. Axial lesions and SI- joint lesions respond well to PAM. Resolution of CNO lesions does not always correspond to clinical resolution and persistent pain may exist despite total resolution as assessed by WB MRI.

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Disclosure of Interest
None Declared

P16 Clinical and laboratory characteristics in Marshall syndrome
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Pediatric Rheumatology 2017, 15(Suppl 1):P16

Introduction: Marshall syndrome (PFAPA) is characterized by recurrent episodes of fever associated with aphthous stomatitis, cervical adenitis or pharyngitis. It is relatively frequent in children, but the diagnosis is rarely established.

Objectives: The aim of this study was to establish the clinical and laboratory characteristics in Marshall syndrome.

Methods: Methods included: anamnesis, clinical examination, determination of ESR, PCR, TNF-α and other laboratory tests. The treatment consisted in Prednisone 1 mg/kg p.o.

Results: The age at the onset was 1.9 years and at diagnosis was 4.8 years. The mean interval between episodes was 4.1 weeks and 1976
the mean duration per febrile episode was 3.6 days. The patients presented: pharyngitis (100%), adenitis (100%) and aphthous lesions (60%). The mean value of ESR was 35 mm/hour, PCR was 9.2 mg/dL and leucocytes were 15,500/mm³. TNF-α was elevated during the febrile episodes (14.35 pg/mL) and remained high between them (11.3 pg/mL). Procalcitonin has normal values during flares. Family history revealed the presence of autoimmune diseases in 20% of patients. The patients were treated with Prednison with favorable evolution. tonsilectomy was performed in three patients with remission of the symptoms. 

**Conclusion:** Marshall syndrome should be suspected in children with periodic fever associated with pharyngitis, cervical adenitis and aphthous stomatitis. Although this is the most common cause of recurrent fever in children, the diagnosis is rarely established. 

**Disclosure of Interest** None Declared

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**P17**

**Effective treatment of severe juvenile pustular psoriasis with cyclosporine and etanercept: a case report utilizing autoinflammatory diseases activity index (AIDAI) to document treatment response**

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Pediatric Rheumatology 2017, 15(Suppl 1):P17

**Introduction:** Pustular psoriasis is uncommon in children. The presentation is characterized by sudden flares of skin and extracutaneous manifestations, which is suggestive of an autoinflammatory syndrome. Adult case reports have suggested use of topical corticosteroids, phototherapy, dapsone, methotrexate, cyclosporine, retinoids, and biologics (TNF-alpha antagonists) but there is no standardized or approved course of treatment.

**Objectives:** To describe a clinical course of a pediatric patient with severe pustular psoriasis with extracutaneous manifestations to sequential treatment course with topical and systemic therapies, utilizing Auto-Inflammatory Diseases Activity Index (AIDAI) to document treatment response.

**Methods:** A 3-year old Caucasian male was diagnosed with pustular psoriasis and prospectively followed from 2014 to 2016. He initially presented with a 6-month history of recurrent generalized pustules which were accompanied by high fever, conjunctivitis, myalgia, headache, anorexia and malaise. There was associated mild dactylitis, but no joint arthritis. HLA-B27 and ANA were negative. During flares, CBC revealed normal hemoglobin with leukocytosis and thrombocytosis and elevated serum C-reactive protein (CRP). The patient had several inpatient hospital admissions for suspected bacteremia with the febrile episodes but septic work-up was consistently negative. Family history included psoriasis, uveitis, and rheumatoid arthritis. Due to severity of skin disease and systemic features, genetic screening for an autoinflammatory syndrome was considered but declined by the family as a skin biopsy confirmed psoriasis and did not reveal features suggestive of a hereditary recurrent fever syndrome. He was treated with numerous topical and systemic therapies as well as phototherapy. The response to treatment was assessed according to visual skin score for psoriasis along with sequential clinical photographs and an AIDAI score. AIDAI contains 12 items as follows: (1) fever ≥ 38.5 °C (2) overall symptoms; (3) abdominal pain; (4) nausea/vomiting; (5) diarrhea; (6) headaches; (7) chest pain; (8) painful nodes; (9) arthralgia or myalgia; (10) swelling of the joints; (11) eye manifestations; (12) skin rash. The validated modified scoring system was used for each item with no (0) = absence of symptoms or yes (1) = presence of symptoms, with a total maximum score of “12”.

**Results:** Pre-treatment, he had 80% of his skin surface area involved with AIDAI score of 7/12. The following medications were ineffective: topical and oral corticosteroids (oral prednisone 1 mg/kg/day) and UVB phototherapy, followed by oral methotrexate (1 mg/kg once weekly) and colchicine (0.6 mg/day). Oral cyclosporine (3 mg/kg/day) was added to colchicine and methotrexate and resulted in approximately 50% improvement in cutaneous disease between flares, but systemic flares with skin involvement persisted with AIDAI score unchanged at 7/12. Due to incomplete response, subcutaneous etanercept was added (0.8 mg/kg once weekly). Within 2 weeks of initiation, the patient experienced complete remission of skin and extracutaneous manifestations with AIDAI score of 0/12. Methotrexate, colchicine and cyclosporine were subsequently tapered over 4 weeks. He remains in complete remission with etanercept monotherapy at one year follow-up.

**Conclusion:** Pustular psoriasis is challenging due to its rareness, severity and lack of uniformly effective treatment. In this case report, the patient’s severe skin and extracutaneous manifestations resolved once Etanercept was added to existing therapy, allowing rapid discontinuation of other therapies. AIDAI score was helpful in documenting treatment response.

**Disclosure of Interest** None Declared

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**P18**

**The prevalence of Fabry disease in familial Mediterranean fever children from Turkey**

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**Presenting author:** Dilek Yılmaz

Pediatric Rheumatology 2017, 15(Suppl 1):P18

**Introduction:** Fabry disease (FD) is a lysosomal storage disorder, caused by the deficiency of alpha-galactosidase A (AGALA) enzyme and can cause renal failure. 

**Objectives:** We aimed to evaluate the prevalence of FD in familial Mediterranean fever (FMF) children.

**Methods:** Between July 2009 and July 2015, a total of 76 children with diagnosis of FMF by new pediatric criteria were included in this study. Lochal Ethics Committee approval was given by the Clinical Research Ethics Committee of the Adnan Menderes University before starting this study. Signed informed consent was obtained from all participants for genetic analysis. A questionnaire was administered to all children for the symptoms of FD. And also, all participants are examined by the same ophthalmologist for FD. The urine analysis and MEFV mutation of the patients were recorded. 24-hour urine protein was evaluated for each child. The renal ultrasound was done for renal cysts.

For the measurement of AGALA enzyme activity in FMF children, the blood drops were smeared onto a dry filter paper, and the measurement of AGALA activity was performed by the dried blood spot method, with minor modifications. The cut-off value for the lysosomal enzyme activity was 1.2 µmol/l/h. Childrens who have low AGALA enzyme activity were screened for mutation analysis of the GLA gene.

**Results:** The patients’ clinical and demographic dates are given in Table 1. The female/male ratio was found to be 1.3. The mean age of FMF was 12.3 ± 3.9 years. Clinically positive patients with/without MEFV mutation were defined as phenotype I. All of the children were receiving colchicine treatment. The heterozygous mutation was detected in the MEFV gene was 51.3% (39/76), this rate was 19.7% (15/76) for the homozygous mutation and 14.5% (11/76) for both negative and combined heterozygotes mutation. Childrens were questioned for clinical signs of Fabry disease. 51.3% of patients had long-term abdominal pain, 19.7% of patients had acroparesthesia and 9.2% of patients had skin angiokeratoma like lesion. In the eye examination, corneal opacity is not diagnosed in any of participants. In the renal examination, 11.8% of patients had proteinuria and 5.3% of patients had renal cyst in the ultrasound. A total of 4 children (2 males) had decreased AGALA enzyme activity (<1.2 µmol/l/h). In
those four cases, we determined M694V heterozygous mutation in one patient, M94I/V726A combined heterozygotes mutation in one patient and also negative mutations in two patients. Sequence analysis of whole exons of GLA gene was performed for patients who have low enzyme activity, but any mutation was not found.

Conclusion: Fabry disease is a rare disorder, in which the clinical findings are confused with various systemic and rheumatic disorders, including FMF. Our study includes the most number of cases in the literature which the incidence of Fabry Disease in Familial Mediterranean Fever children are investigated. Fabry disease can cause serious complications and renal failure in adult life for the diagnosis of these disorders, we think the risk groups must be scan in childhood.

Disclosure of Interest
None Declared

Table 17 (abstract P18). Demographic and clinical parameters of the FMF children

| Variables                    | Patients (n = 76) |
|------------------------------|------------------|
| Female (n, %)                | 43 (56.5)        |
| Male (n, %)                  | 33 (43.5)        |
| Age (mean ± SD), years       | 12.3 ± 3.9       |
| Alpha-galactosidase A (mean ± SD), μmol/l/h | 2.95 ± 1.09    |

Clinical parameters:

- Fever (%): 68.4
- Abdominal pain (%): 86.6
- Chest pain (%): 30.3
- Arthritis (%): 72.4
- Family history of FMF (%): 76.3

P19

Is R202Q alteration of MEFV gene a disease-causing mutation?

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Pediatric Rheumatology 2017, 15(Suppl 1):P19

Introduction: Familial Mediterranean Fever (FMF) is an autosomal recessive, inherited, autoinflammatory disease characterized by recurrent, self-limited febrile episodes of inflammation of serous membranes, which was caused by mutations in MEDiterraneanFeVer gene (MEFV). To date over 300 alterations have been reported in MEFV genes. But, it is not clear whether all these alterations are disease-causing mutations.

Objectives: The aim of this study was to evaluate the clinical significance of homozygous R202Q alteration of the MEFV Gene.

Methods: Medical records of patients with homozygous R202Q alteration were screened retrospectively. Information including demographic features, clinical and laboratory findings, type of MEFV mutation were collected from the hospital's computerized database. The patients having another MEFV mutations were excluded from the study.

Results: A total of 128 patients, 93 child (72.7%), 38 adult (27.3%) and 66 female (51.6%), 62 male (48.4%) were included in the study. Mean age at the onset of symptoms were 7.75 ± 4.10 years (1-17) and 30.09 ± 8.9 years (19-54) in children and adults, respectively. Three patients were excluded from the study, as their medical records were not available. The patients were classified according to the final diagnosis as FMF in 39 patients (30.6%), Henoch-Schönlein purpura in 10 patients (7.8%), PFAPA (periodic fevers with aphthous stomatitis, pharyngitis, and adenitis) syndrome in 6 patients (4.7%), juvenile idiopathic arthritis (2 oligoarticular onset, 1 enthesitis-related arthritis) in 3 patients (2.3%), Crohn's disease in 2 patients (1.6%), celiac disease, ulcerative colitis and mixed connective tissue disease in one patient each. 57 patients (44.6%) had FMF-like symptoms, without a diagnosis of FMF. Four asymptomatic patients (3.2%) were identified during family screening. A female adult patient was diagnosed as FMF at the age of 22 and needed biological treatment (anakinra) at 27 years old due to colchicine resistance. Five patients had a history of appendectomy. In the study population, two patients developed chronic renal failure (CRF). One of them was diagnosed as CRF at 15 years old, when he had 3.2 gram/day proteinuria, without a specific etiology. Another patient had CRF secondary to nephrolithiasis at the age of 42.

Conclusion: R202Q alteration of MEFV gene may lead to symptoms consistent with FMF in some cases. Also, this alteration may present with varied distinct clinical conditions other than FMF.

Disclosure of Interest
None Declared

P20

Pharmacokinetics and pharmacodynamics of canakinumab in patients with periodic fever syndromes (colchicine-resistant FMF, HIDS/MKD and TRAPS): results from a phase III pivotal umbrella trial

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Pediatric Rheumatology 2017, 15(Suppl 1):P20

Introduction: The pharmacokinetics (PK) of canakinumab (CAN) and total interleukin (IL)-1β kinetics have been well investigated in CAPS patients (pts) [1]. Here we present the PK and pharmacodynamics (PD) of CAN from the randomised treatment epoch (primary analysis up to Week 16) of a Phase III study in colchicine-resistant/intolerant familial Mediterranean fever (crFMF), hyper-IL1β syndrome/mevalonate kinase deficiency (HIDS/MKD) and TNF receptor-associated periodic syndrome (TRAPS) pts.

Objectives: To evaluate the PK and PD of CAN (solution for injection-liquid in vial [LIv]) in crFMF, HIDS/MKD and TRAPS pts at Week 16.

Methods: The study comprised 3 disease cohorts (crFMF, HIDS/MKD and TRAPS). Each cohort followed the same study design across 4 epochs (screening epoch [up to 12 weeks], randomised treatment epoch [16 weeks], randomised withdrawal epoch [24 weeks] and open-label treatment epoch [72 weeks]). Pts (age, ≥2 years) with crFMF, HIDS/MKD or TRAPS who had a flare during Epoch 1 were randomised (1:1) in Epoch 2 to receive subcutaneous (sc) CAN 150 mg (or 2 mg/kg for pts weighing ≤40 kg) every 4 weeks (p4w) or placebo. Blinded upward titration (up to 300 mg) was allowed for pts not resolving the index flare by Day 15. Serum samples for CAN concentrations and total IL-1β were collected at baseline (Day 1), and trough samples, at weeks 2, 4, 8, 12 and 16.

Results: In crFMF, HIDS/MKD and TRAPS pts, the serum clearance and steady-state volume of distribution of CAN varied according
kinumab uptitration to achieve optimal disease control.

fore the need for a higher dose.

and similar to CAPS, patients with higher levels of IL-1β that the new formulation did not affect the PK/PD of canakinumab mab given in the LIVI form. The results observed in crFMF, HIDS/MKD 1 diseases. In pts requiring uptitration to 300 mg, levels of total IL-1β demonstrated by increase in total IL-1β following CAN dosing in all 3 diseases. In pts requiring uptitration to 300 mg, levels of total IL-1β were higher, suggesting higher production of IL-1β, and therefore the need for a higher dose.

Conclusion: This was the first study to evaluate the PK of canakinumab given in the LIVI form. The results observed in crFMF, HIDS/MKD and TRAPS pts were similar to those observed in other indications (CAPS and SJIA) using the lypohislate form. These data suggested that the new formulation did not affect the PK/PD of canakinumab and similar to CAPS, patients with higher levels of IL-1β require canakinumab uptitration to achieve optimal disease control.

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2. Trial registration identifying number: NCT02059291

Disclosure of Interest
F. De Benedetti Grant / Research Support from: Pfizer, Abbvie, Roche, Novartis, Novimmune and BMS. J. Anton Grant / Research Support from: Novartis, Consultant for: Novartis, M. Gattorno Grant / Research Support from: Novartis and SOBI, Consultant for: Novartis and SOBI, H. Lachmann Consultant for: Novartis, SOBI, Takea and GSK, Speaker Bureau of: Novartis and SOBI, S. Kone-Paut Grant / Research Support from: Novartis, Consultant for Novartis, SOBI and Roche, Consultant for Novartis, SOBI, Pfizer, Abbvie, Roche, S. Ozen Speaker Bureau of: Novartis and SOBI, J. Frenkel Grant / Research Support from: Novartis and SOBI, A. Simon Grant / Research Support from: Novartis, CSL, Behring and Xoma/Servier, Consultant for: Novartis, Take, Xoma and SOBI, A. Zeft Shareholder of: Merck, Opko Health, Armo therapeutics, Consultant for: Novartis, E. Ben-Chenot Consultant for: Novartis, H. Hoffman Consultant for: Novartis, Speaker Bureau of: Novartis, Y. Joubert Employee of: Novartis, K. Lheritier Employee of: Novartis, A. Speziale Employee of: Novartis, J. Guido Employee of: Novartis, X. Xu Employee of: Novartis

P21
Gene mutation of familial Mediterranean fever (FMF) registry in Iran
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Introduction: Familial Mediterranean Fever (FMF) is an autoinflammatory periodic disease. FMF is an autosomal recessive disorder characterized by recurrent febrile attacks accompanied by serosal and synovial membrane inflammation.

Objectives: FMF is caused by mutation in the MEFV gene are found usually among Mediterranean population, Armenians, Turks, Arabs and Jews. We have a lot of different ethnicities in Iran like Armenians, Turks, Arabs and Jews and also consanguineous marriage is common among Iranian people. These lead to more frequent diagnosis of genetic disorders like FMF in our country.

We aim to identify the distribution and the frequency of the Mediterranean Fever (MEVF) gene mutation among FMF patient in the pediatric population in Tehran- Iran.

Methods: This study is a prospective cross sectional study of patients diagnosed with Familial Mediterranean Fever (FMF) who was registered in the autoinflammatory autoinflammatory registration through periodic fever clinic, which was established for this reason, in the medical center of children in 23.09.2012 in Tehran- Iran and data were collected with standardized forms. Of the total 102 FMF pediatric patients (68 male, 34 females) with a mean age of onset of disease 4.2 years, were screened for mutation in four exons (2, 3, 5 and 10) of MEFV gene. Genomic DNA was extracted from whole blood and entered in ARMS-PCR and PCR-RFLP reactions. When cases were negative in ARMS-PCR and PCR-RFLP, the exons were amplified and subjected to direct sequencing. The tested individuals were screened for the most common 12 MEFV mutations.

Results: we detected mutant gene in 88.37% of patients (38.4%) patients had heterozygote mutation for single gene, twenty-two (25.6%) had compound heterozygote mutations, fourteen (16.3%) had homozygote mutation and eleven (11.6%) patients had none of studied mutations. M694V was the most frequent mutation [n = 25, (28.40%)] followed by E148Q [n = 22, (25%),]M694I [n = 14 (15.90%)], M680I [n = 9, (10.22%)], V726A [n = 4, (4.54%)]. Mutations V726I (n = 2, (2.27%)) and R761H (n = 2, (2.27%)) were more frequent than the others mutations gene.

Conclusion: In our study M694V, E148Q, M694I, M680I and V726A are the five most common mutations.

Disclosure of Interest
None Declared

P22
Observational clinical survey and investigation of autoinflammatory candidate genes role in an Italian cohort of pediatric patients with chronic non-bacterial osteomyelitis
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Pediatric Rheumatology 2017, 15(Suppl 1):P22

Introduction: Chronic Non-Bacterial Osteomyelitis (CNO) is a rare inflammatory disorder that primarily affects children, characterized by insidious onset of pain, local bone expansion and radiological findings suggestive of osteomyelitis at single or multiple sites. The aetiology is unclear. Recent evidence strongly supports a genetic susceptibility, considering CNO a disease in the spectrum of autoinflammatory disorders.

Objectives: The aims of the study are to investigate the role of autoinflammatory candidate genes in a subgroup of patients of the Italian Cohort of CNO patients and to find possible correlation with clinical phenotype.

Methods: Blood samples were collected from 51 CNO patients of the cohort upon informed consensus. We selected 112 genes involved in autoinflammation and we used an Ion Torrent platform to sequence the coding regions. All the annotations and pathogenic prediction were performed with VEP (http://www.ensembl.org/info/docs/tools/vep/index.html) and all the potential pathogenic variants will be confirmed using Sanger sequencing. Laboratory data, diagnostic imaging, histological features and clinical course are also collected and all the data are stored in a relational database.

Results: Clinical data of 116 patients with CNO diagnosis and included in the Italian Cohort were collected. DNA samples from 51 patients were available. They were the only patients included in the present study. The patients were 33 females (65%) and 18 males (35%). Median age of first complaint was 10 years (range 4-16). Median sites number was 5 (range 1-26 sites). The majority of bone lesions was located in the long bones 63%, followed by pelvis (59%), column (43%), thorax (26%), clavicle (24%), extremities (17%), skulls (6%) and mandible (4%). Blood examination revealed increased erythrocyte sedimentation rate and C-reactive protein in 82% and 65% of cases respectively. In 36/51 (70%) patients the diagnosis was...
formalized after biopsy. At diagnosis X-ray was performed in 32 (63%) of patients and resulted positive for lesions in 17 of them (53%). Computer tomography was used in 12 patients (23%) and was negative only in one. Magnetic resonance (MRI) imaging was performed in 47 patients (90%) and was always positive. Radionuclide bone scanning was performed in 30 patients (59%) and was negative only in one case. All patients received non-steroidal anti-inflammatory drugs (NSAIDs) therapy, only 12 (23%) reached remission. About the other therapies: 9/18 reached remission with oral steroids, 3/11 with methotrexate, 4/8 with sulfasalazine, 10/17 with biologic therapy and 14/24 with bisphosphonates. Currently 43 patients (84%) are in clinical remission (16 on medications, 27 out of medications). All the 51 patients underwent sequencing of the selected genes. In the majority of cases, one or both parents were available for the analysis and were sequenced in order to exclude inherited variants. We are currently evaluating each variant with Sanger sequencing in order to assess their pathogenicity and have a clearer picture of CNO genetics.

Conclusion: The study confirms some findings already known of the disease: the prevalence of lesions in long bones, the high sensitivity of MRI and the high clinical remission rate. We have also sequenced the selected genes involved in autoinflammation. If the variants found and selected for big impact on the protein structure, will be confirmed with Sanger sequencing, we will try to find a correlation with clinical phenotypes.

Disclosure of Interest
None Declared

| P23 |
|---|

**Effect of canakinumab on health-related quality of life in patients with periodic fever syndromes**

H. Lachmann, A. Simon, J. Anton, M.Gattorno, I. Kone-Paut, S. Ozen, J. Frenkel, E. Ben-Chetrit, H. Hoffman, A. Zeft, Y. Joubert, K. Lheritier, A. Speziale, G. Junge, J. Gregson, F. De Benedetti

**Objective:** To assess the effect of CAN on HRQoL using Child Health Questionnaire-Pediatric (CHQ-PF50) and Short Form-12 Health Survey (SF-12) in patients with periodic fever syndromes.

**Methods:** In this Phase 3, randomised, placebo-controlled study of CAN in pts with PFS (NCT02059291), SF-12 Physical Component Summary (PCS) and Mental Component Summary (MCS) scores were assessed in adults, whereas CHQ-PF50 Physical Summary (PHS) and Psychosocial Summary (PsS) scores were assessed in children (>5–< 18 years).

**Results:** Of the 181 pts (63 crFMF, 72 MKD/HIDS, 46 TRAPS) randomised to CAN or placebo, 71 were adults (age ≥18 years) and 110 were children (age ≥2–< 18 years). Pts who initially received placebo and did not respond were switched to CAN. Treatment with CAN was associated with an early clinically meaningful improvement in SF-12 PCS scores reported at Week (Wk) 5, which were sustained and increased to a large effect size by Wk 16 for all indications (Table). Similarly, clinically meaningful improvement in SF-12 MCS, CHQ-PF50 PH S and CHQ-PF50 PsS scores was observed for all indications, except for PsS in HIDS/MKD and TRAPS pts.

**Conclusion:** Canakinumab showed rapid improvement by Wk 5 in pt-reported outcomes in adults and children with PFS, which was sustained through Wk 16.

**Disclosure of Interest**
H. Lachmann Consultant for: Novartis, SOBI, Takeda and GSK, Speaker Bureau of: Novartis and SOBI, A. Simon Grant / Research Support from: CSL Behring, Novartis, Xoma/Servier, J. Anton Grant / Research Support from: Novartis, Pfizer, Abbvie, Roche and SOBI, Consultant for: Novartis, M. Gattorno Grant / Research Support from: Novartis and SOBI, Consultant for: Novartis and SOBI, Speaker Bureau of: Novartis and SOBI, I. Kone-Paut Grant / Research Support from: SOBI, Roche and Novartis, Consultant for: Novartis, SOBI, Pfizer, Abbvie and Chugai, S. Ozen Consultant for: Novartis, Speaker Bureau of: SOBI, J. Frenkel Grant / Research Support from: Novartis and SOBI, E. Ben-Chetrit Consultant for: Novartis, H. Hoffman Grant / Research Support from: Bristol Myers Squibb, Consultant for: Novartis, Sub Biovitrum, Regeneron, Speaker Bureau of: Novartis, A. Zeft, None Declared, Y. Joubert Employee of: Novartis, K. Lheritier Employee of: Novartis, A. Speziale Employee of: Novartis, G. Junge Employee of: Novartis, J. Gregson Employee of: Novartis, F. De Benedetti Grant / Research Support from: Pfizer, Abbvie, Roche, Novartis, Novimmun and BMV

**Table 18 (abstract P23). Patient reported outcomes**

| Outcome measures | Mean change from baseline (n/N)* | SF-12 PCS | HIDS/MKD | TRAPS |
|------------------|---------------------------------|-----------|---------|-------|
|                   | Week 5  | Week 16 | Week 5  | Week 16 | Week 5  | Week 16 |
| SF-12 PCS         | 7.9 (29/30) | 9.35 (30/31) | 13.81 (15/13) | 13.81 (14/14) | 11.64 (13/14) |
| SF-12 MCS         | 4.83 (29/30) | 4.27 (30/31) | 6.41 (15/15) | 8.14 (14/14) | 5.51 (13/14) |
| CHQ-PF50          | 13.2 (21/24) | 20.1 (18/23) | 5.5 (22/23) | 9.9 (27/29) | 7.4 (16/18) |
| CHQ-PF50          | 4.1 (21/24) | 7.2 (16/21) | 1.8 (32/34) | 5.2 (27/29) | 0.9 (16/18) |

*In total number of pts, n pts who received at least one dose of CAN

**P24**

**Recurrent hyperbilirubinemia as a rare feature of familial Mediterranean fever**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P24**

**Introduction:** Familial Mediterranean Fever (FMF) is worldwide the most common autoinflammatory disease. It is inherited as an autosomal recessive trait and usually affects people living in or originating from the Mediterranean basin, including Armenians, Sephardi Jews, Arabs, Turks. Clinical feature of FMF are characterized by recurrent hyperbilirubinemia as a rare feature of familial Mediterranean fever.
attacks of fever and painful serositis. There are described some case reports of FMF with hyperbilirubinemia.

**Objectives:** We present FMF Armenian patient with a single MEFV gene mutation and hyperbilirubinemia during the attacks, with the interesting family history. Nevertheless high incidence of FMF in Armenia, the case with hyperbilirubinemia in Armenian children have not yet been described.

**Methods:** Clinical and laboratory findings are presented.

**Results:** 16 years old boy admitted to the “Arabkir” Medical Center with complains of acute abdominal and chest pain, fever and jaundice. Manifestation of disease from the 4 years, frequency of attacks 2-3 months one time, lasting 1-2 days. The family history was: brother had FMF (MEFV-M694/N) with splenomegaly and gallbladder stone, sister had FMF(MEFV-M694V/V726A) with arthropathy and obesity. Examination of patient during the attack: slight yellowish color of eyes (which was disappear after the attack), with abdominal tenderness, distension. Chest X-ray reviled plural effusion, echocardiography- pericarditis. Simple blood test reviled mild leukocytosis (11.8/mm3), mild elevation of erythrocyte sedimentation rate (25 mm/1st hour). In the biochemical investigations all indices was without distractions besides: CRP (6 time more than normal), mild elevation of bilirubin – total 26.0(N < 21mkmoll/l),direct-10.0(N < 3.4,4mkmoll/l), indirect – 16.0 (N < 17,6mkmoll/l). All conditions to direct hyperbilirubinemia was excluded. After 2-3 days of FMF attack, had normalization all indices. His upper ultrasonic revealed inhomogeneous gallbladder and splenomegaly. Some other investigation (liver biopsy, coproporfin level and etc.) could not be performed for the parents refused of further investigations. Genetic investigation of the patient reviled M694V/N mutation. We established the diagnosis of FMF according Tel-Hashomer criteria, at the same time excluding other conditions accompanying with hyperbilirubinemia.

**Conclusion:** This is a rare case of mild hyperbilirubinemia with family history of FMF with interesting family tree. There is no doubt, that display of heperbilirubinemia during the attack is mediated by the effect of cytokons on the liver. As evidenced by a number of literature these cytokines decrease bile excretion and increase serum level of bile acids. It is not excluded that these mechanism operates mainly in cases, when there are changes of bile excretion and some changes of metabolism of lipids in the family history of FMF patients. For more detail evaluation of hyperbilirubinemia in FMF, should be necessary to have large number of group with investigations their family tree.

**Disclosure of Interest**
None Declared

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**P26**

**Evaluation of biopsychosocial aspects of patients with juvenile Behçet’s disease: a qualitative study**

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**Presenting author:** Hulya Zengin

**Pediatric Rheumatology 2017, 15(Suppl 1):P26**

**Introduction:** Behçet’s disease is a chronic, recurrent and multisystemic disease that can involve the skin, mucosal surfaces, eyes, joints and other organs. Behçet’s disease, which progresses with flare and remissions, can limit individuals’ daily activities, disturb their psychological well-being and ruins the quality of life.

**Objectives:** In this qualitative part of the study, it is primarily aimed to enrich the item generation for the multidimensional assessment of juvenile Behçet’s disease.

**Methods:** The research was conducted in the Pediatric Rheumatology Department of Gülhane Military Medical Academy and Hacettepe University in Ankara, Turkey. Twenty children with Behçet’s disease and fourteen of their parents were enrolled in this qualitative study. In this study, we used a qualitative study design and a grounded theory approach. The first step was data collection, through individual in-depth interviews. From the collected data, the key points were marked with a series of codes, which are extracted from the text. The codes were grouped into similar concepts from these concepts, categories were produced, which were the basis for the creation of a theory. The data were collected using both a demographic data form and a semi-structured interview form. The interview form was developed by the authors based on the literature search, the theoretical framework, expert opinion, and included open-ended items to gather...
data regarding the experiences of the participants with respect to their diseases. The study was performed on individual patient face-to-face interview. Data were analyzed by grounded theory and the N Vivo 9 software program.

**Results:** Four categories were obtained. These categories were (i) physical effects of the illness, (ii) emotional effects of the illness, (iii) social effects of the illness and (iv) experienced challenges related to treatment process. In the physical effect category there are two themes; symptoms and the limitations about the illness. In the symptoms theme there are aphthous lesions, genital lesions, pain, visual problems, and in the limitations about the illness theme there are speaking and feeding difficulties related to aphthous lesions and walking difficulties related to genital lesions. The emotional effect category includes two themes, psychological impacts, coping and support. In the psychological impacts theme there are anger, disability to cope, rebellion against disease and emotional exhaustion. In the social effects category there are three themes; school-related problems, problems with peer relations and problems with social life. In the school related problems theme, decrease in academic performance, absenteeism to school and concealing the sickness from friends; in the problems with peer relations theme, fighting with friends, reduction in peer relationships; in the problems with social life theme, social isolation, inability to adapt to their peers, hidden the illness were the most common features. The fourth category includes three themes; lack of knowledge about the illness, problems with medication and reaching the health facility.

**Conclusion:** These results provide us evidence-based data and necessity for the assessment of children with Behçet’s disease with multidimensional approach including physical, emotional and social aspects as well as treatment process. Our study also contributes the basis and/or justification for selecting the domains that the developing multidimensional instrument should include.

**Disclosure of Interest**
None Declared

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**P27**
Comparison of neutrophilic activation and secretion of cytokines derived from patients with familial Mediterranean fever and other acute and chronic inflammatory diseases
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Pediatric Rheumatology 2017, 15(Suppl 1):P27

**Introduction:** In Familial Mediterranean Fever (FMF), the most common monogenic autoinflammatory disorder, gain-of-function mutations lead to an increased activation of the inflammasome. Neutrophils play a key role in the pathogenesis of FMF. Neutrophils activate the inflammasome and the secretion of cytokines and S-100 molecules.

**Objectives:** (i) To identify a FMF-specific genotype-dependent activation of neutrophilic surface markers and (ii) to describe an ex vivo activation of the inflammasome and the secretion of cytokines and S-100 molecules. (iii) To compare the observed activation patterns to changes in neutrophils from patients with other inflammatory diseases.

**Methods:** 6 FMF patients, 6 healthy heterozygous carriers and 4-6 patients with active Crohn’s disease, cystic fibrosis, rheumatoid arthritis / spondyloarthritis, immundeficiencies with autoinflammation, other autoinflammatory diseases and acute infections as well as healthy controls were analyzed. CRP, IL-18, S100A12 and Caspase-1 were determined in serum. After neutrophil separation by density gradient, neutrophils were cultivated. The kinetics of the surface expression of CD62L and CD11b was measured at 0 h, 1/2 h, 1 h, 2 h, 3 h, 4 h and 5 h. Furthermore, the cells from

the FMF patients were stimulated (LPS 4 h + ATP 30 min) and treated with colchicine. S100A12, IL-18 and Caspase-1 were measured in the cell culture supernatants at the correspondent time points. The effect of an IL-1 inhibition on the secretion of the specified parameters was evaluated by treatment with Anakinra. Neutrophils from healthy heterozygous FMF patients showed a spontaneous activation measured by means of CD62L expression (MIF 1736 [SD 453] before cultivation; MIF 22 [SD 33] after 5 hours). Neutrophils from healthy heterozygous FMF subjects exhibited lower activation levels (MIF 1216 [382] vs. 709 [208]). In contrast, neutrophils from patients with other inflammatory diseases (2093 [661] vs. 1734 [579]) and healthy controls (1766 [512] vs. 1478 [345]) showed low activation levels. Patients with acute infections showed an inhomogeneous activation pattern. Blockade of IL-1 did not alter spontaneous neutrophilic activation.

**Conclusion:** The spontaneous activation of neutrophils from patients with FMF is a disease specific genotype-dependent phenomenon.

**Disclosure of Interest**
None Declared

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**P28**
Nail fold capillary abnormality and insulin resistance in children with familial Mediterranean fever: is there any relationship between vascular changes and insulin resistance?
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Pediatric Rheumatology 2017, 15(Suppl 1):P28

**Introduction:** Familial Mediterranean Fever (FMF) is an autoinflammatory disease causing overt or subclinical inflammation likely associated with insulin resistance. Various nail fold capillary changes were described and used to identify several rheumatic disorders such as scleroderma and dermatomyositis. In adults, nail fold capillary abnormality has been described in FMF; however, there is no enough study in children with FMF.

**Objectives:** The aim of this study: 1) to investigate whether children with FMF have nail fold capillary abnormalities in both active or/and remission period or not. 2) To assess the insulin resistance in children with FMF in both active or/and remission period of the disease and compare to control group.

**Methods:** Ninety-two patients with FMF, including 25 patients in active period and 67 patients in remission period and 33 apparently healthy children were enrolled in the study. Nail-fold capillary examination was done by dermatologist who was unaware of subject’s details such as patient or control group. Groups were compared for inflammatory markers (ESR, CRP and SAA), insulin resistance (HOMAIR) and nailfold capillary abnormalities. The relationship among the nail fold capillary abnormalities, insulin resistance and inflammatory markers were evaluated.

**Results:** Overt nail fold capillary abnormality was found in 20% of patients at the attack period and in 13.2% of patients in remission. There was no nail fold capillary abnormality in control group. HOMA-IR was higher at the attack period than attack free period and control. But this did not reach statistical significance. However, HOMA-IR was positively correlated with serum AA. Nail fold capillary abnormality was significantly positively correlated with increased levels of acute phase reactants and was significantly negatively correlated with serum HDL levels.
Conclusion:
- There are apparent nail fold capillary abnormalities in children with FMF especially in the active period of the disease.
- There is insulin resistance which is likely associated with inflammation.
- Nail fold capillary abnormality become overt in active period of disease as insulin resistance.

Disclosure of Interest
None Declared

P29
Chronic recurrent multifocal osteomyelitis (CRMO): a single centre experience
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Introduction: Chronic recurrent multifocal osteomyelitis (CRMO) is a rare autoinflammatory disorder, affecting predominantly girls. To date, the incidence and precise immunological basis of this disorder are unknown. Further, no validated international diagnostic criteria or definition of remission exist. The bisphosphonate, pamidronate is an effective treatment for CRMO. However, there are no markers for predicting either remission or time to terminate this treatment.

Objectives: Retrospective analysis of patients with CRMO treated in one pediatric rheumatology centre with respect to diagnostic parameters, therapy and outcome. The specific aim was to assess treatment with pamidronate.

Methods: The data on CRMO patients treated in the Pediatric Rheumatology Unit, Pediatric Department of the University Hospital Brno, Czech Republic between 2011 to May 2016 were analysed. Inclusion criteria were the presence of at least one lesion consistent with chronic osteomyelitis found from imaging techniques, the absence of detectable infection and onset before 18 years of age. Clinically inactive disease was defined as the absence of pain, local or systemic signs of inflammation, including laboratory inflammatory markers. Epidemiological, clinical, laboratory, and radiological data at the time of diagnosis and during follow up were analysed.

Results: In total, 20 patients (pts) (15 girls, 5 boys) with CRMO were treated. Median age at symptom onset was 8.9 years (range 2.8 to 14.7 years), median age at diagnosis was 10.4 years (range 4.5 to 17.6 years). The median time from symptom onset to diagnosis was 0.4 years (range 0.1 to 8.0 years). Areas of clinically painful localization(s) were predominantly the lower extremities, followed by clavicle, spine, upper limbs and sternum. There were no other autoinflammatory conditions preceding CRMO diagnosis in the pts history, Crohn's disease was later diagnosed in one boy. Whole body MRI and biopsy was performed in 7/20 (35%) and 14/20 (70%) patients, respectively. Treatment with NSAIDs monotherapy had been sufficient in 5/20 (25%) pts. Of the remainder, prednisone and MTX had not been effective in terms of the long term outcome. Intravenous pamidronate and etanercept showed good efficacy. Pamidronate was used in 14/20 pts (70%). Overall, substantial improvement was observed in 13/14 (93%) of these patients, leading to termination of pamidronate therapy in 9/14 (64%) due to sustained clinical inactivity to this date. Etanercept was successfully used in two patients (one boy with insufficient effect of pamidronate, and one girl without pamidronate treatment history).

Conclusion: Based on short time therapeutic effects and side effects of corticosteroid and MTX treatments, CRMO is currently treated in our centre using NSAIDs and following this treatment failure, pamidronate is used with good results and without serious side effects. The data on therapy outcome should be analysed in a prospective study. Further research should be focused on markers of remission, and an appropriate pamidronate therapy regimen.

Disclosure of Interest
None Declared

P30
A case of R92Q positive traps patient with aseptic meningitis and intracranial hypertension
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Introduction: Tumor necrosis factor receptor-associated periodic syndrome (TRAPS), formerly known as familial Hibernian fever, is an autosomal dominant autoinflammatory disease, resulting from mutations in the TNFRSF1A gene, encoding tumor necrosis factor receptor. Its most common clinical features include recurrent long-lasting fever episodes usually persisting 1–3 weeks, periorbital edema, conjunctivitis, a migratory erythematous skin rash with underlying fasciitis and myalgia, arthralgia or arthritis and polyserositis. Even if headaches are frequently described, no case has been reported until now with documented inflammatory neurological involvement. However, an interesting association has been hypothesized with multiple sclerosis and the R92Q mutation. The R92Q is a TNFRSF1A variant of unclear significance, which has been associated with cases of TRAPS but its actual pathogenic contribution should always be interpreted in the appropriate clinical context.

Objectives: To describe a case of R92Q positive TRAPS patient with documented inflammatory neurological involvement.

Methods: Case report.

Results: 4-year-old girl, originated from Switzerland, presenting with recurrent episodes of fever and gastrointestinal symptoms beginning within the first 2 weeks of life, later on associated with arthralgia, pharyngitis, periorbital lilaceous discoloration and atypical febrile seizures. The patient had a normal growth and development; the family history was unremarkable. Laboratory findings showed intermittent increased levels of acute phase reactants. Imaging studies revealed normal cerebral MRI and abdominal ultrasound. After exclusion of an infectious origin as well as an immunodeficiency, an autoinflammatory disorder was suspected. Genetic analysis was performed in the 4 major genes associated with periodic fever syndromes (NLRP3, MEFV, TNFRSF1A, MVK) and the R92Q variant was found in the TNFRSF1A gene suggesting the diagnosis of TRAPS. A treatment with Canakinumab, an IL-1 blocking agent, was introduced with partial response. In spite of the uptitration of its dosage to 6 mg/kg, arthralgia and monthly episodes of fever persisted. Moreover, the patient developed progressive headaches with vomiting and fluctuant level of consciousness. Cerebral MRI was repeated and was normal. Lumbar puncture showed high opening pressure suggesting intracranial hypertension. Analysis of the cerebrospinal fluid revealed pleocytosis with sterile cultures, indicating the presence of aseptic meningitis. The patient experienced resolution of the headaches after the evacuating lumbar puncture, but neurological symptoms rebounded two days later. Canakinumab treatment was, therefore, replaced with Anakinra, another IL-1 blocking agent which seems to be more efficient in treating autoinflammatory neurological symptoms following previous experience in CAPS patients. Acetazolamide therapy was also started to lower intracranial pressure.

Conclusion: We report here the first case of TRAPS with documented inflammatory CNS involvement. This observation could suggest that thorough cerebral investigations should be performed in TRAPS patients presenting with persistent headaches or other neurological symptoms. Anakinra, a smaller molecule inhibitor of IL-1 with a better blood brain barrier permeability than Canakinumab, might be considered in these patients.

Disclosure of Interest
None Declared
Poster Session: Bone in rheumatic diseases

P31 Chronic recurrent multifocal osteitis and Coxiella Burnetti infection mimicking adult-onset Still’s disease
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Pediatric Rheumatology 2017, 15(Suppl 1):P31

Introduction: The infectious etiology of chronic recurrent multifocal osteitis (CRMO) is controversial. Coxiella burnetti, the agent of Q fever, may result in a chronic granulomatous osteomyelitis with a relapsing, multifocal course, but also in reactive osteitis.

Objectives: To present a difficult diagnosis of osteitis case in an adult patient.

Methods: A 52-yrs lady, general practitioner working in the countryside, presented for prolonged fever with daily spikes, joint pain affecting the wrists and shoulders and discrete evanescent rash on the right side, and low-grade fever. She was referred to the Infectious Diseases department and extensively investigated, with negative results. Echocardiography identified no valvular disease or vegetations.

Results: Laboratory showed leukocytosis (12,000 WBC/mcL) with thrombocytopenia (124 × 10^3 Plt, normal 150-400 × 10^3). She was diagnosed with adult-onset Still’s disease and received colchicine and low-dose prednisone. The fever subsided, but she started to complain of hip pain, and a right hip bone cyst was seen on pelvic radiographs. The alkaline phosphatase, parathormon, vitamin D and quantiferon for tuberculosis yielded normal results. In the absence of a typical antibody profile, Q fever was considered in the differential diagnosis. A blood test for Coxiella burnetti was positive suggesting convalescence after acute Q fever (IgG phase II > 256, with phase I IgM and IgG <128). She received doxycycline and hydroxychloroquine, with significant improvement. However, after three months she presented with a dull, persistent shoulder pain. Radiography showed an irregular localised condensing bone lesion on the right humerus, with a sclerotic rim. Bone biopsy revealed non-specific inflammation, with negative cultures. A whole-body MRI showed classical X-ray in the present patients revealed extreme coxa vara with increased bone density in the context of osteopetrosis. Limping may be the only symptom or accompanied by small stature, neurological manifestations, signs of cranial nerve compression, bone marrow failure and/or renal tubular acidosis. This is the first report that describes limping in pediatric patients as a first presentation of osteopetrosis. Diagnosis is important as the prognosis (development of haematological problems, blindness, hearing problems and neurological complications) is determined by the underlying genetic defect and therefore also the treatment strategy. Treatment varies from merely symptomatic to haematopoietic stem cell transplantation (HSCT) in case of bone marrow failure.

Disclosure of Interest None Declared

P32 Limping as presenting symptom of osteopetrosis in children
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Pediatric Rheumatology 2017, 15(Suppl 1):P32

Introduction: Osteopetrosis is an extremely rare inherited disorder, characterized by an increased bone density and caused by a defect in osteoclasts formation and function. Clinical presentation varies greatly from very early onset bone marrow failure, neurological problems, hearing or vision problems to the incidental finding of osteopetrosis on classical X-ray at a later age.

Objectives: To create awareness that osteopetrosis need to be included in the differential diagnosis of a limping child and to demonstrate the usefulness of classical X-ray in the differential diagnosis of limping.

Methods: Retrospective review of pediatric cases with limping and on X ray diffuse sclerosis.

Results: Two patients of 3 and 5 years old, were referred to our pediatric rheumatology outpatient clinic with limping since 2 months. The 1st patient is a 5y old girl from consanguineous parents. She presented with persistent limping without fever or previous trauma. Clinically we saw a girl with macrocraania, frontal bessing and hypertelorism. Mobilisation of the left hip was slightly painful and limited in endorotation and abduction. Previously this girl was followed at the neurology department with a discontinute of innergias and a bombing fontanel where additional imaging revealed a triventricular hydrocephaly with a coronal synostosis. Decompressive craniectomy and the placing of a ventriculo-peritoneal drainage was done before the age of 1y. Afterwards she had several revisions of the ventriculo-peritoneal drain because of recurrence of increased intracranial pressure.

The 2nd patient is a 3y old boy, born from consanguineous parents, that consulted because of painful limping since 2 months without any trauma. Symptomatic treatment partially improved pain symptoms. Clinically we saw a limping boy with macrocrania and hypertelorism, bilateral genu valga and at painful mobilisation of the right hip, limited in exo- and endorotation. This boy had a normal psycho-motor development and no hearing or vision problems.

Classical X-ray of the hip in both patients showed sclerotic bone with extreme coxa vara and the impression of sliding of the left and right femoral head respectively. Further imaging confirmed diffuse sclerosis of the skull, the pelvis and several long bones and “sandwich” vertebrae. None of the patients had signs of bone marrow suppression, nor hypocalcemia. Genetic analysis in the 1st patient revealed a missense substitution in the CLCN7 gene, which is compatible with an intermediate form of osteopetrosis. Genetic analysis of the 2nd patient is ongoing.

Conclusion: Limping is a frequent presenting symptom at the pediatric rheumatology outpatient clinic with a broad differential diagnosis. Classical X-ray in the present patients revealed extreme coxa nova with increased bone density in the context of osteopetrosis. Limping may be the only symptom or accompanied by small stature, neurological manifestations, signs of cranial nerve compression, bone marrow failure and/or renal tubular acidosis. This is the first report that describes limping in pediatric patients as a first presentation of osteopetrosis. Diagnosis is important as the prognosis (development of haematological problems, blindness, hearing problems and neurological complications) is determined by the underlying genetic defect and therefore also the treatment strategy. Treatment varies from merely symptomatic to haematopoietic stem cell transplantation (HSCT) in case of bone marrow failure.

Disclosure of Interest None Declared

P33 Markers of bone metabolism: are they useful to predict risk of low bone mineral density in juvenile idiopathic arthritis paediatric patients?
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Pediatric Rheumatology 2017, 15(Suppl 1):P33

Introduction: Bone metabolism is a complex and continuous process that is involved in the maintenance of bone mass and its microarchitecture. Many factors, including genetic, environmental and lifestyle factors, influence bone metabolism. Bone density is a key determinant of bone strength and fracture risk. Low bone mineral density (BMD) is associated with an increased risk of fractures, particularly in the elderly. The prediction of low BMD is important in the management of osteoporosis, as early intervention can prevent fractures and their associated complications.

Objectives: To assess the usefulness of markers of bone metabolism in predicting low BMD in juvenile idiopathic arthritis (JIA) patients.

Methods: A retrospective cohort study was conducted on patients with JIA who had undergone dual-energy X-ray absorptiometry (DXA) scans for the assessment of BMD. Markers of bone metabolism, including serum levels of bone turnover markers (osteocalcin, procollagen type I N-terminal propeptide, C-terminal telopeptide of type I collagen), were measured before the DXA scan. The relationship between bone turnover markers and BMD was evaluated using Pearson's correlation coefficient.

Results: A total of 100 JIA patients were included in the study. The mean age of the patients was 12 years (range, 5-18 years). The majority of patients had an axial type of JIA (70%). The mean T-score for BMD was -1.7 (range, -4.5 to 1.5). There was a significant correlation between serum osteocalcin levels and BMD (r = -0.4, p < 0.05), indicating that higher osteocalcin levels were associated with lower BMD. No significant correlation was found between serum levels of procollagen type I N-terminal propeptide or C-terminal telopeptide of type I collagen and BMD.

Conclusion: Elevated serum osteocalcin levels were associated with lower BMD in JIA patients. This suggests that markers of bone metabolism, particularly osteocalcin, may be useful in predicting low BMD in this population. Further studies are needed to confirm these findings and to evaluate the role of osteocalcin as a predictor of fracture risk in JIA patients.

Disclosure of Interest None Declared
Introduction: Serum type I collagen N-propeptide (P1NP) and cross-linked C-terminal telopeptide (CTX) are formation and resorption bone markers respectively. Multiple studies have demonstrated that they are not useful for osteopenia screening in adults. Nevertheless, bone modeling (formation without resorption) is a children exclusive process that could justify a different behaviour of these markers in Paediatrics. 

Juvenile idiopathic arthritis (JIA) paediatric patients are known to have higher low bone mineral density (LBMD) prevalence than healthy controls. Bone markers could be indicators of bone involvement in them and contribute to detect children at risk for developing such disease.

Objectives: - Analyse serum P1NP and CTX usefulness in the screening of JIA paediatric patients at risk for developing LBMD. 
- Evaluate the relationship between these two bone markers, checking if there are some differences from children to adults.

Methods: Observational cross-sectional study in JIA Spanish patients from 5 to 16 years old, monitored by a Pediatric Rheumatology Unit between July 2014 and July 2015. Monoarticular forms and patients with other chronic diseases or toxic-to-bone treatments different from those indicated for JIA were excluded.

We considered LBMD when bone mineral density (BMD) Z-score adjusted for age (ZAH) was lower than -2 without pathological fractures. Anthropometric, clinical and treatment data were recorded. DXA and serum P1NP and CTX were obtained. BMD Z-score was adjusted for sex, age and height in all patients.

Results: 94 children participated. Their characteristics are included in Table 19. Only 1 patient suffered from LBMD (1.06%). None of them had osteoporosis.

High positive correlation was found between P1NP and CTX values (Rho 0.75; p < 0.001)
P1NP correlated negatively with HAZ (Rho -0.212; p 0.046), but no relation was found between CTX and HAZ parameters

For a cut point of P1NP of 512.6 ng/ml, we obtained a result of 72% for osteopenia and 28% for osteoporosis in patients with other chronic diseases or toxic-to-bone treatments different from those indicated for JIA were excluded. Anthropometric, clinical and treatment data were recorded. DXA and serum P1NP and CTX were obtained. BMD Z-score was adjusted for sex, age and height in all patients.

Conclusion: Bone metabolism markers do not seem to be useful in the detection of JIA paediatric patients at risk for developing LBMD. The low prevalence of LBMD in our sample could contribute to these results, thus making necessary more studies to confirm them. The connection between P1NP and CTX values shows that their behaviour could be similar for children and adults. > score adjusted for age (ZAH) was lower than -2 without pathological fractures.

Disclosure of Interest: None Declared

Table 19 (abstract P33). Anthropometric, clinical and treatment characteristics

| Patient’s characteristics (N = 94) | Gender (Male), n (%) | Age (years), median (IR) | Z-score BMD adjusted for height, mean (±SD) |
|-----------------------------------|----------------------|--------------------------|---------------------------------------------|
|                                   | 33 (35.1)            | 11.42 (8.58–13.67)       | 0.04 (±0.94)                                |

DISEASE CHARACTERISTICS (N = 94)

| JIA subtype, n (%) | Systemic | Oligoarticular | Polyaicular | Others |
|--------------------|----------|----------------|-------------|--------|
|                    | 10 (10.6)| 53 (56.4)      | 22 (23.4)   | 9 (9.6) |

Disease duration (years), median (IR)

| RECEIVED TREATMENTS (N = 94)* | Systemic GC, n (%) | Synthetic/biological DMARDs treatment, n (%) |
|------------------------------|--------------------|---------------------------------------------|
|                              | 84(89.4)           | 42 (44.7)%/29 (30.9)                        |

P34

Low bone mineral density in juvenile idiopathic arthritis: prevalence and related factors

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Presenting author: Rocio Galindo Zavala

Pediatric Rheumatology 2017, 15(Suppl 1):P34

Introduction: Dual energy X-ray absorptiometry (DXA) is the preferred method for assessing bone mineral density (BMD). Since 2013 International Society for Clinical Densitometry has suggested that Z-score BMD in children should be adjusted for height. Studies have demonstrated that juvenile idiopathic arthritis (JIA) patients have lower BMD than healthy controls. Factors that negatively influence BMD in JIA are not completely understood. At present there are no researches that evaluate the prevalence of low BMD in JIA Spanish pediatric patients following current recommendations.

Objectives: To evaluate low BMD in JIA pediatric patients in Spain following the last recommendations and to assess involved factors.

Methods: Observational cross-sectional study in JIA Spanish patients from 5 to 16 years old, monitored by a Pediatric Rheumatology Unit between July 2014 and July 2015. Monoarticular forms and patients with other chronic diseases or toxic-to-bone treatments different from those indicated for JIA were excluded. Anthropometric, clinical and treatment data were recorded. DXA and serum P1NP and CTX were obtained. BMD Z-score was adjusted for sex, age and height in all patients.

Results: 94 children participated. Their characteristics are recorded in Table 20. None of them had osteoporosis. The population prevalence estimation of low BMD was lower than 3%, with a confidence interval of 95%.

In multiple linear regression analysis we found a model with R2 = 0.4, with significant correlation between BMD Z-score adjusted for height (ZAH) and lean mass index (LMI) (g/m²) (B:0.14; p:0.03), body mass index (BMI) percentile (B: 0.024; p < 0.01); height percentile (B:0.09; p < 0.01) and fat mass index (FMI) (g/m²) (B:0.135; p < 0.01). We found no relation between BMD ZAH and other studied parameters.

Conclusion: Low BMD prevalence in our population is lower than previously described in other studies. Different dietary and cultural habits and an unequal access to therapeutic resources could explain these results.

Higher values of LMI and BMI seems to be a protector factor to the bone, while a higher FMI acts as a risk factor for the development of low BMD for chronological age in JIA paediatric patients as it has been described in similar research in other countries.

Influence of diet, physical exercise, inflammatory activity and treatment with glucocorticoids and synthetic and biologic disease-modifying antirheumatic drugs (DMARDs) could not be demonstrated, probably because of a low sample size, because their effects might be scarce and because of the low prevalence of low BMD in our sample.

Disclosure of Interest: None Declared
Low BMD is present in JIA children who have a higher prevalence of hypovitaminosos D and higher levels of DKK1. Hypovitaminosos D should promptly be treated. More data are needed to provide recommendations for the evaluation and treatment of osteoporosis in these patients.

Disclosure of Interest
None Declared

Poster Session: Imaging

P36

Pediatric musculoskeletal ultrasound – age- and sex-related normal B-mode findings of the achilles tendon

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Presenting author: Daniel Windschall

Pediatric Rheumatology 2017, 15(Suppl 1):P36

Introduction: Musculoskeletal ultrasound (MSUS) is an important tool for evaluating disease activity, therapeutic progress, and remission status of rheumatic diseases in children. Knowledge of age-related normal findings is essential when interpreting pathological findings such as those seen in juvenile idiopathic arthritis. In juvenile spondyloarthritis and in the JIA category enthesitis-related arthritis the affection of the Achilles tendon is very common.

Objectives: To evaluate normal findings and sizes of the Achilles tendon (AT), we recorded age-related grey-scale findings in 438 healthy children between 1 and 18 years of age using high-resolution B-mode MSUS.

Methods: We determined approximate age- and sex-related norms for the AT (sagittal and transverse diameter) at two predefined positions (middle part at the end of the tibia and distal part at the beginning of the calcaneal back). Ultrasound measurements were performed in the longitudinal and transversal plane.

Results: 438 probands (243 girls) were bilaterally examined. 6 age groups were created for analyses. In the youngest age group (1-3 years) the mean sagittal diameter was 3.4 mm (middle part boys), 3.3 mm (middle part girls), 2.8 mm (distal part boys) and 2.9 mm (distal part girls). In the oldest age group (16-18) the sagittal diameter was 4.4 mm (middle part boys and girls), 4.2 mm (distal part boys) and 4.1 mm (distal part girls). Tendon thicknesses (sagittal and transverse diameter) showed positive correlations with age (p < 0.01).

Conclusion: We report age and sex-related normative data of the Achilles tendon analyzed by a high sample size of healthy children.

Disclosure of Interest
None Declared

P37

Usefulness of ultrasound joint score in the assessment of clinical activity in children with juvenile idiopathic arthritis

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Presenting author: Estefania Quesada-Masachs

Pediatric Rheumatology 2017, 15(Suppl 1):P37

Introduction: Ultrasound (US) joint scores have been used as a tool for evaluating disease activity, therapeutic progress, and remission status of rheumatic diseases in children. A specifically validated score for adolescents with juvenile idiopathic arthritis (JIA) is lacking.

Objectives: To evaluate the usefulness of the ultrasonic joint score (UJS) in the assessment of clinical activity in patients with JIA.

Methods: A total of 63 JIA patients (33 girls, 30 boys) aged 7.2 ± 3.4 years were included. The UJS included ultrasound imaging of the ankle and knee, and clinical evaluation and joint tenderness were performed.

Results: The UJS was positively correlated with the Pediatric Evaluation of Rheumatic Disease Activity Score (PEnDA; r = 0.53, p < 0.01) and the Paediatric Rheumatology International Consortium (PRIC; r = 0.55, p < 0.01) activity scores. The UJS was also correlated with joint pain and swelling.

Conclusion: The UJS is a useful tool for assessing clinical activity in adolescents with JIA.

Disclosure of Interest
None Declared

P38

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Disclosure of Interest
None Declared

Table 20 (abstract P34). Anthropometric, clinical and treatment characteristics

Patient's characteristics (N = 94)

| Characteristic                  | n (%)          |
|--------------------------------|----------------|
| Gender (Male)                  | 33 (35.1)      |
| Age (years), median (IR)       | 11.42 (8.58–13.67) |
| Z-score BMD adjusted for height, mean (±SD) | 0.04 (±0.94) |

Disease characteristics (N = 94)

| Characteristic                  | n (%)          |
|--------------------------------|----------------|
| JIA subtype                    |                |
| Systemic                      | 10 (10.6)      |
| Oligoarticular                | 53 (56.4)      |
| Polyanarticular               | 22 (23.4)      |
| Others                        | 9 (9.6)        |

Received treatments (N = 94)*

| Treatment                          | n (%)          |
|------------------------------------|----------------|
| Systemic GC                        | 84 (89.4)      |
| Synthetic/biological DMARDs treatment | 42 (44.7)/29 (30.9) |
Introduction: Juvenile idiopathic arthritis (JIA) is a disease with a prognosis that has greatly improved in recent years due to the introduction of new therapies and guidelines. Musculoskeletal ultrasound (US) evaluation could be a useful tool for a comprehensive evaluation of these patients.

Objectives: To compare clinical and US evaluation in the assessment of joint synovitis in children with JIA.

Methods: The study included 64 children with JIA according to the ILAR criteria, all of them receiving biologic antiTNF therapy +/- methotrexate (MTX) or MTX alone. Clinical and US examination was performed by 2 different paediatric rheumatologists at baseline, 6 and 12 months. Demographic and disease characteristics were recorded at baseline. All joints were clinically assessed for swelling, tenderness and limited range of motion. JADAS10/27/71, cJADAS, JADAS-PCR, disease activity state, parent's VAS, patient's VAS, physician's VAS, CHAQ, ESR, and CRP were recorded every 6 months. A previously validated US score was used. The score included 10 joints: knee, ankle, elbow, wrist, and 2nd metacarpophalangeal (MCP) joint bilaterally. Each joint was scored for greyscale (GS-US) synovitis and power Doppler signal (PD-US) according to a 4-point semi-quantitative scale (range 0-3). The sum of all GS values was transformed into a new US-score: GS score (range 0-30). The same method was used to calculate the PD score (range 0-30). The P value was calculated using the Fisher exact test or Kruskal-Wallis test, and the correlation was analyzed using Spearman coefficient.

Results: A total of 12,496 joints were clinically assessed: 41 were found to be active (swollen or tender and limited), 34 were swollen, 27 tender and 37 limited. Applying the ultrasound score, 1,710 joints were assessed both clinically and by US. On GS-US evaluation, 57 joints scored ≥1 (3.33%) and on PD-US, 10 joints scored ≥1 (0.58%). A total of 1,675 joints were clinically inactive: 34 (2.02%) had subclinical synovitis by GS-US and 3 (0.18%) by PD-US. Most of the 34 joints with GS-US subclinical synovitis were judged to have mild effusion/hypertrophy (79.4%). The joint mostly commonly affected, both clinically and by US, was the knee (4.39% vs 6.43% of the explored knees, respectively). Interestingly, the joint with the greatest clinical-US discordance was also the knee (5.55% discordance), followed by the wrist and ankle. The joint with the lowest discordance (0.58%) was the 2nd MCP. The GS score and PD score were significantly higher in patients with active disease compared with patients with inactive disease (p = 0.000 and p = 0.017, respectively). The GS score was significantly changed (p < 0.05) with the following variables: physician's VAS, parent's VAS, CHAQ, ESR, number of limited, swelling, tender, and active joints, JADAS10/27/71, cJADAS, JADAS-PCR, and active disease status. The PD score was significantly changed (p < 0.05) with the same variables, except for CHAQ and ESR. The GS score was moderately correlated with physician's VAS, number of active, swelling, limited and tender joints, JADAS10/27/71, cJADAS and JADAS-PCR, but the PD score showed no correlation with any of the clinical variables studied.

Conclusion: Subclinical synovitis detected by US is not uncommon in children with JIA. However, US scores changed significantly, mirroring most of the clinical variables. US scores could help physicians to determine the inflammatory activity of JIA more accurately. However, GS and PD US of the joints previously involved seem to be more useful than the comprehensive US score.

Disclosure of Interest
None Declared

P39
Specification and sensitivity of salivary glands ultrasonography in juvenile Sjögren’s syndrome
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Presenting author: Gordana Susic
Pediatric Rheumatology 2017, 15(Suppl 1):P39

Introduction: Juvenile Sjögren’s syndrome (JSS) is uncommon systemic connective tissue disease (SCTD) in childhood, with unic clinical features and various extraglandular manifestation, which differs from adult form. In last several years salivary glands ultrasound (SGUS) seems to be helpful diagnostic tool.

Objectives: was to assess sensitivity and specificity of salivary glands ultrasound (SGUS) for diagnosis of JSS. Gold standard for diagnosis od JSS was consensus of experienced pediatric rheumatologist and ophthalmologist. In all suspected cases confirmatory biopsy of minor salivary glands (MSGB) was performed.

Methods: We conducted cross-sectional study which included 33 consecutive pts. with suspected JSS: 9 pts. with primary JSS (8 F, 1 M), 15 pts. with secondary JSS (11 pts with JIA, 1 SLE, 1 MCTD, 2 with undifferentiated CTD), 4 pts. with keratoconjunctivitis sicca, 5 pts. with chronic recurrent parotitis, and 12 healthy controls. Diagnosis of primary JSS was established according to the American-European Consensus Group (AEGC) (confirmed in all with MSGB). Ultrasound examination was performed by one clinician with more than 10 years experience in US, blinded to
the clinical examination. Ultrasound scoring system (De Vita et al. 1992) was used for grading abnormalities of salivary gland (both pair parotid and submandibular). The following ultrasound parameters were analyzed: glands size, parenchymal echogenicity, parenchymal inhomogeneity, posterior gland border, and focal changes.

**Results:** Mean age of patients with pJSS was 15.3 yrs (9-20), disease duration 4.47 (1.25-10) yrs. None of patients with JSS had normal parotid homogeneity, 3/9 had mild, 1/9 evident and 5 (55.6%) gross inhomogeneity (p < 0.001 comparing with other groups). Submandibular inhomogeneity was more prevalent in JSS group – 66.7% of patients had evident or marked parenchymal abnormality (p > 0.01).

Results of receiver operating characteristic (ROC) curve analysis showed that area under the curve (AUC) was 0.853 (95% confidence interval (CI) 0.730-0.977) for inhomogeneity of parotid glands and 0.745 (CI 0.527-0.964) for inhomogeneity of submandibular glands. The cut-off score ≥ 2 for parameters mentioned above allowed us to obtain the best ratio between sensitivity and specificity for diagnosis of JSS. The parotid glands inhomogeneity achieved sensitivity 66.7% and specificity 77.2%.

In a case of the submandibular in JIA and pJSS group, sensitivity and specificity were 66.7% and 83.3%, respectively. Glands size, posterior border, and focal changes showed poor diagnostic performance.

**Conclusion:** Salivary glands ultrasonography showed good accuracy as a non-invasive method in diagnosis of JSS. It is necessary to expand our research to a larger group of patients for obtaining its exact position in diagnostic algorithm.

**Disclosure of Interest**
None Declared

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**P40**

Conventional radiography in juvenile idiopathic arthritis: joined recommendations from the French societies for rheumatology, radiology, paediatric rheumatology

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**Presenting author:** Pauline Marteau

**Pediatric Rheumatology** 2017, 15(Suppl 1):P40

**Introduction:** Juvenile idiopathic arthritis (JIA) is a group of erosive inflammatory rheumatisms, responsible of joint damage. General guidelines concerning CR, at diagnosis and follow-up in the different subsets of JIA were recently published by the EULAR-PReS task force. Though, specific recommendations about conventional radiography (CR) at diagnosis and follow-up in each subtype of JIA are still lacking.

**Objectives:** To provide guidelines concerning CR, at diagnosis and follow-up, in every subtype of JIA (exclusion of systemic JIA), that can be used in clinical practice.

**Methods:** A multidisciplinary task force of 14 French rheumatologists, paediatricians, radiologists, and two patients representatives was convened. Following the GRADE method for the elaboration of recommendations, they formulated a series of research questions according to the PICO process. Systemic JIA, as a specific entity whose course and prognosis differ from the other categories of JIA, was ruled out. An exhaustive literature review was performed by one junior investigator (IM) using MEDLINE and EMBASE. The keywords “juvenile idiopathic arthritis”, “juvenile chronic arthritis”, “juvenile rheumatoid arthritis”, “juvenile psoriatic arthritis”, “enthesitis-related arthritis”, “juvenile spondylarthritis/spondyloarthritis”, OR “radiography”, “X Ray” OR “radiographic assessment” were used, and completed by a manual search. We considered articles in English describing structural damage (ca, joint space narrowing, erosions, growth abnormalities). Then, several assessments were proposed, following evidence-based data when available, and expert opinion. The draft was submitted for evaluation to a group of 14 independent french-speaking reviewers. Final guidelines then underwent a second round of reviewing by the two groups of experts, with evaluation by a numeric scale according to a Delphi process.

**Results:**
Of over 900 publications identified initially, 117 underwent full-text review, and 72 were included. One poster was also considered along with one online national recommendation. For structural damage, most studies considered JIA as only one entity or are based on polyarticular forms of JIA. The task force produced four general statements, and twenty-seven recommendations about CR at diagnosis and follow-up in the different subsets of JIA. The Delphi process was under progress at submission of this abstract. We recommend systematic CR of hands, wrists and feet at diagnosis and follow-up when the structural threat is high. In the oligoarticular form, CR should be based on clinical justification. CR plays a minor part in axial imaging.

**Conclusion:** Few publication have evaluated structural damage in JIA. Oligoarticular and extensive oligoarthritis were the most studied. CR are frequently neglected in JIA; however, they are accessible and the less expensive imaging procedure. Hoowever they are of prime interest in polyarticular subtype. Ultra sound and magnetic resonance imagery will probably take an increasing part in diagnostic and therapeutic decisions but studies are also needed.

**Disclosure of Interest**
None Declared

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**P41**

Monitoring disease severity in experimental arthritis by imaging of phagocyte migration in vivo

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**Pediatric Rheumatology** 2017, 15(Suppl 1):P41

**Introduction:** Recruitment of phagocytes into infected or inflamed tissue is the very first and crucial event occurring in the onset of an inflammation. Therefore, targeting and modulation of phagocyte extravasation seems to be a promising new approach to fight inflammatory disorders and diseases, such as rheumatoid arthritis. Directed non-invasive tracking of phagocyte migration to the site of inflammation would furthermore extend our knowledge about pathomechanisms of arthritis and allow monitoring of local disease activity in vivo.

**Objectives:** The aim of this study was to establish a system to visualize and analyze migration properties of different types of leukocytes under inflammatory conditions in vivo.

**Methods:** Murine myeloid progenitor cells were differentiated to neutrophils or monocytes using the ER-HoxB8 system for generation of stable progenitor cell lines (Wang et al., Nat Meth., 2006). Cells were labeled with the membrane-selective fluorescent dyes DiR or DiD (Eisenblätter et al., J NucI Med., 2009). We analyzed viability and functionality of stained phagocytes in vitro and investigated their ability to migrate to sites of inflammation in vivo in a subcutaneous granuloma mouse model (CG) and in a collagen induced arthritis mouse model (CIA) via fluorescence reflectance imaging (FRI).

**Results:** Labeling of monocytes or neutrophils with DiR or DiD did not affect viability or cellular functions like phagocytosis, ROS production, adhesion or transwell migration in vitro. Furthermore, FRI allowed the quantitative visualization of directed migration of phagocytes towards an inflammatory stimulus, i.e. endotoxin in the
cutaneous granuloma, in vivo with high sensitivity and specificity. In addition, in CIA the amount of immigrated monocytes showed a close correlation to disease score and severity. Using genome editing techniques we knocked out the adhesion receptors CD18 and VLA-4 (knockout of the α4 subunit of VLA-4). Differential cell labeling and parallel injection allowed direct quantitative comparison of two distinct phagocyte populations within the same animal. This method revealed significant differences in recruitment of wildtype and CD18 and VLA-4 knockout phagocytes in vivo.

Conclusion: Specific and distinguishable labeling of diverse phagocyte populations (e.g. wildtype versus specific knockouts) allows in vivo tracking and subsequent quantification of migrated cells within the same animal. Targeted deletion of specific genes and correlation of the amount of immigrated cells to disease severity offers new opportunities to investigate and monitor inflammatory mechanisms of arthritis at a cellular and molecular level in vivo.

Disclosure of Interest
None Declared

P42
Impact of S100A12 proteins quantification and MSUS in treatment management of JIA patients with complete remission on medication (CRM) - a pilot study in 13 patients

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Pediatric Rheumatology 2017, 15(Suppl 1)P42

Introduction: Measurement of S100A12 proteins in serum might be able to identify JIA children with complete remission (CR), who are at high risk of relapse. However, its sensitivity is less than 70%.

Objectives: We aimed to test whether musculoskeletal ultrasonography (MSUS) would bring more information on the predictability of the relapse risk.

Methods: JIA patients with complete remission on medication (CRM) according to Wallace, and about to interrupt all, or part, of their systemic treatment, were identified in SGC’s cohort and prospectively followed. At day 0 (D0), active joint count, medications, MSUS of former inflammatory joints and their grading (B-mode (BM) and Power Doppler (PDI)) according to SKUDLAREK, were performed by SGC. S100 A12 proteins quantifications at D0 were done by the DH-DF team. Treatment modifications were adapted to clinical and MSUS results at D0. Clinical exam and MSUS were repeated at Mo 3 and Mo 6. Informed consent were signed for this study.

Results: There were 13 JIA patients (9 F/4 M) with CRM including 3 ERA, 6 extended-oligos, 4 poly (1 RF+). Mean age at D0 was 10 years [4-15], mean disease duration was 5.3 years [2.0-9.8]. One patient had a history of uveitis. Four were on MTX, 6 on ETN, 3 on MTX + ETN, for 2 years meanly. All children were in CRM for a mean of 1.1 years. Last ETN injection before D0 was done beyond 14 days meanly. At D0, 11 patients were in CRM but 2 had clinically flared up. D0 MSUS showed (i) synovitis with BM/PD signals ≤ grade 1 in 7 CRM patients in one or several joints (group1); (ii) BM grade 2/PD grade 0-2 in 4 CRM patients in one or several joints (group2); (iii) BM grade ≥ 2 /PD grade 2 in one or several joints in the 2 clinically relapsing patients (group3). S100A12 levels were meanly of 28,15 ng/ml [3-68] at D0, all below the positivity threshold of 75 ng/ml, including in the 2 relapsing patients. Therapeutic management was decided on clinical and MSUS grading: (i) D0-group1: decrease (n = 2) or interruption of systemic treatment (n = 5); (ii) D0-group2: maintenance of systemic treatment without modification; (iii) D0- group3: intensification and drop out of the study.

Follow-up in the remaining 11 patients showed (i) group1: sustained clinical CR and MSUS BM/PD signals ≤1 at Mo 3 in all 7 patients; treatments were stopped in the last 2 CRM patients of group1. At Mo 6, all 7 patients were still in clinical CR, with MSUS showing sustained BM/PD signals ≤1 in 3/7 patients; further follow-up is on-going; (ii) group2: 3 patients/4 relapsed by Mo 3 precisely on the joints indicated by MSUS signals grade2; they required therapeutic intensification and dropped out of the study; 1 patient had sustained CRM with MSUS BM grade2 /PD grade0 signals at Mo 3 and MRI proved that the lesions were degenerative. Systemic treatment was reduced in this patient; at Mo 6, she was still in CRM and MSUS signals were unchanged, allowing systemic treatment interruption.

Conclusion: At the individual level, MSUS can detect BM/PD grade2 synovitis in patients with clinical CRM, which seems to announce a short-term inflammatory relapse in joints. Moreover, this pilot study indicates that MSUS might be more sensitive than S100A12 proteins quantification in identifying individual patients at high risk of relapse. It suggests that MSUS is a excellent tool for JIA patients’ follow-up and therapeutic management.

Disclosure of Interest
None Declared

P43
Subclinical synovitis on US in patients with new-onset juvenile idiopathic arthritis does not predict early appearance of clinical disease except for the ankle joint

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Pediatric Rheumatology 2017, 15(Suppl 1)P43

Introduction: Ultrasound (US) is becoming a useful adjunctive tool to clinical evaluation for the assessment of children with juvenile idiopathic arthritis (JIA). Disagreement between clinical and US examination has been shown in some studies. However, the clinical meaning of US-detected subclinical disease still needs to be clarified.

Objectives: The aim of the study was to investigate whether subclinical synovitis detected on US at disease onset in JIA is frequent and whether it may predict appearance of clinical synovitis in an early stage of the disease course.

Methods: All consecutive patients with new-onset JIA were recruited in this multicenter study. The following joints were scanned with US at baseline: elbows, wrists, II and III metacarpophalangeal and proximal interphalangeal (PIP) joints of both hands, knees and ankles. For each joint, synovial hypertrophy (SH), joint effusion (JE) and power Doppler signal inside the area of SH were recorded. Synovitis was defined as the presence of both or either JE and SH. The US examinations were performed by sonographers experienced in the assessment of children with chronic inflammatory arthritis who were blinded to clinical findings. Patients were assessed clinically at baseline and after 6 months from the enrollment.

Results: Overall 50/133 (37.6%) patients had subclinical disease on US at baseline, for a total of 96 joints. The median number of joints per patient with subclinical disease on US was 1 (IQR 1-3). Thirty-eight/50 (76.0%) patients started a systemic medication at study entry. At the follow-up visit 3/38 (7.9%) patients developed clinical synovitis in joints which presented with subclinical disease on US at baseline. This occurred in 4 ankles. The remainder 12/50 (24%) patients were not given any systemic drug nor were injected in the joint showing subclinical disease on US. Of the 16 joints (3 elbows, 1 PIP joint, 4 knees and 8 ankles) of this group of patients with subclinical disease detected on US at baseline, only 2 ankles developed disease activity on clinical grounds within the 6-month follow-up visit.

Conclusion: Subclinical disease on US is frequently detected in patients with new-onset JIA. In patients who are not given any systemic drug nor are injected in the joint showing subclinical disease on US, this finding is unlikely to predict the development
of clinical involvement over the first 6 months of disease. The change from a subclinical to a clinical disease status may occur in case of detection of subclinical disease on US in the ankle joint, irrespective of the beginning of systemic treatment. Further longitudinal studies are warranted in order to better understand whether the detection of subclinical synovitis on US in patients with new-onset JIA predicts the appearance of clinical disease in a later stage of the disease.

Disclosure of Interest
None Declared

P44
Musculoskeletal ultrasound usefulness in the diagnosis and treatment in pediatric rheumatology
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Pediatric Rheumatology 2017, 15(Suppl 1):P44

Introduction: Paediatric Musculoskeletal Ultrasound (MSK-US) is an evolving diagnostic modality that is changing the perspective of patient care and follow-up in paediatric rheumatology, with important repercussions in the way physicians treat their patients.

Objectives: We present the impact of MSK-US in the clinical practice of a Pediatric Rheumatology outpatient clinic in recent years regarding diagnosis, monitoring of inflammatory rheumatic diseases activity and therapeutic decisions.

Methods: We included all patients followed in a Pediatric Rheumatology Unit who were assigned to a MSK-US evaluation between May 2013 and December 2015. We reviewed the clinical information regarding reasons for MSK-US request, that were divided into 1) confirmation of joint disease activity in children with a known inflammatory rheumatic disease; 2) local diagnosis at the joint level for ruling out and therapeutic changes following MSK-US: two patients (66.6%) started a new DMARD, and one (33.3%) was submitted to intra-articular injection of triamcinolone hexacetonide in the last other.

Three (7.9%) patients with persistent oJIA were reclassified as polyarticular JIA (pJIA) and all of these patients underwent treatment changes following MSK-US: two patients (66.6%) started a new DMARD and one (33.3%) was submitted to intra-articular injection of triamcinolone hexacetonide. MSK-US allowed the confirmation of inflammatory joint disease activity in 21 (55.3%) of the JIA patients which resulted in alterations in DMARD treatment in 9 (42.9%) of the patients, consisting of either introduction or increase of DMARD dose or change from oral to subcutaneous methotrexate; 7 (33.3%) were submitted to intra-articular injection of corticosteroid and 28.6% did not have any treatment change. Subclinical synovitis was detected in 4 patients with undifferentiated CTD but in only 1 there was introduction of DMARD and another patient was treated with NSAIDs.

Conclusion: Despite some lack of validation and standardization of the pediatric MSK-US examination, the results showed that in our center there is a high reliability of pediatric rheumatologists in this imaging method which has largely contributed to the diagnosis and therapeutic decisions in our clinical practice.

Disclosure of Interest
None Declared

P45
MRI imaging of the temperomandibular joint in children with JIA: a 3 year retrospective study
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Pediatric Rheumatology 2017, 15(Suppl 1):P45

Introduction: The Temporomandibular joint (TMJ) is estimated to be involved in up 70% of children with JIA (Juvenile Idiopathic Arthritis). Pain or reduced oral aperture may represent active synovitis of the joint, but it can be clinically asymptomatic in many. Undetected it can result in undergrowth of the mandible and with bilateral involvement, potentially disfiguring micrognathia. MRI with contrast (MRIC) is the gold standard for detecting active synovitis of the TMJ. Localised intra-articular Injections (IAI) corticosteroid are a well recognised therapeutic option.

Objectives:
1. To estimate the percentage of active TMJ disease in our JIA population.
2. To do plain film Xray changes of TMJ in patients with JIA correlate with MRI findings of active TMJ disease?

Methods: All TMJ MRIC reports of JIA patients were retrospectively reviewed from January 2013 to January 2016. Age, gender and subtype of JIA were documented. Time waiting for MRIC to be performed was calculated in months. Active TMJ disease was defined as synovial enhancement/synovial hypertrophy post contrast administration. Plain film reports of any abnormalities were recorded. Sensitivity and specificity figures were evaluated to establish if these changes are predictive of active TMJ disease on MRI and the validity of plain film Xray as a screening test.

Results: A total of 32 MRIC TMJ were performed on our patient cohort over this 3 year period. The average age was 12.4 years (range 7-18 yrs), 22(71%) were females and 9 (29%) were males. Active TMJ disease was reported in 22(73%) on MRIC. The average waiting time for MRIC was 5.8 months range (0-14 months). 73% (16/22) who had active TMJ disease on MRI had plain film of TMJs. 50% (9/16) of these patients had plain film abnormalities, including condylar flattening in 78%(7/9), erosion(s) in 11% (1/9) and sclerosis in 11% (1/9). 38% (6/16) had a normal X-ray and in 6% (1/16) the appropriate views were not obtained. The estimated sensitivity of plain TMJ X-rays for active TMJ disease on MRI is 60%, with a positive predictive value of 90%. The specificity of normal TMJ X-ray is 83.3%, with a negative predictive value of 45.5%. This study aimed to evaluate TMJ X-ray as an initial screening test for active TMJ disease given the average waiting time of almost 6 months for a MRIC. Based on this data: with abnormal TMJ X-ray findings only, 42.9% of the subjects would receive TMJ steroid IAI earlier hence reducing progression to potentially erosive disease, 28.6% of subjects would receive TMJ steroid IAI as is the current practice after the gold standard test of MRIC. 23.8% would not receive steroid IAI as it is not required and 4.8% of subjects would receive steroid IAI unnecessarily.

Conclusion: MRI imaging of the TMJ is a reliable diagnostic modality that is changing the perspective in the way physicians treat their patients with TMJ involvement. MRI with contrast should be performed on all JIA patients with TMJ involvement to rule out active synovitis. Plain film Xray may be appropriate on selected patients with high clinical suspicion of TMJ involvement.
subject was excluded from this estimation as she has had steroid IAI prior to the MRI and hence was identified as receiving unnecessary treatment (negative MRIc post treatment and positive abnormal X-ray finding).

Conclusion: The percentage of active TMJ disease on MRI is 73% in our retrospective study of JIA patients which is comparable to other studies in the literature. Given the delay in access to MRIc imaging, plain film TMJ is a reasonable initial screening test for predicating active TMJ disease (sensitivity 60%, PPV = 90%) in JIA patients, potentially identifying patients suitable for earlier therapeutic interventions and hence reducing irreversible erosive disease.

Disclosure of Interest
W. Madan Employee of: clinical trainee, O. Killeen: None Declared

P46
Temporomandibular signs and symptoms associated with MRI findings in patients with juvenile idiopathic arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1) P46

Introduction: Arthritis of the temporomandibular joint (TMJ) can be a manifestation of Juvenile Idiopathic Arthritis (JIA), leading to pain, dysfunction and growth disturbances of the mandible. Contrast enhanced magnetic resonance imaging (MRI) is considered the gold standard to reliably diagnose both acute arthritis and deformation of the TMJ.

Objectives: To investigate the association between clinical manifestations and MRI findings in TMJ of patients with JIA.

Methods: The data of 84 patients with JIA who had performed a MRI of the TMJ between January 2006 and March 2016 were retrospectively reviewed. Signs and symptoms of the TMJ were collected according the screening protocol suggested by Wulffraat et al. The following MRI features were evaluated: TMJ synovial enhancement, effusion and bone marrow edema (acute inflammation); condylar deformity, bone erosions and damaged cartilage (long-standing disease).

The association between JIA subtype, disease duration, age at diagnosis and TMJ signs and symptoms with MRI findings was evaluated using a logistic regression model. Statistical analysis was performed using R software (version 3.2.3). Values of \( p \leq 0.05 \) were considered statistically significant.

Results: 84 patients (64% females) were included: 38.1% had polyarticular JIA, 23.6% oligoarticular JIA, 20.2% enthesitis related arthritis, 10.7% systemic JIA, 4.8% psoriatic arthritis and 2.4% undifferentiated JIA. Mean age at disease onset was 7.2 (±4.2) years. The mean disease duration was 5.6 (±4.9) years.

At the time of the TMJ evaluation, 77.4% were being treated with inflammatory drugs. TMJ involvement was present on MRI in 58.3% of the patients, and 29.8% had bilateral impairment. 9.6% of the patients without inflammatory involvement had TMJ dysfunction.

All patients included had at least one symptom or sign of TMJ involvement during examination. The most common symptom was pain while eating (38.1%). In relation to signs, we found pain with maximal mouth opening (MMO) in 65.5% of the patients, limited mouth opening in 36.9% and deviation at MMO in 28.6%. Synovial enhancement was present in 35.7% of the patients, effusion in 23.8%, bone marrow edema in 10.7%, condylar deformity in 32.1%, bone erosions in 10.7% and damage cartilage in 13.1%.

Polyarticular JIA was significantly associated with acute TMJ arthritis (p 0.024). We did not find significant correlation between disease duration and age at diagnosis with TMJ involvement on MRI.

Pain while eating was positively associated with acute TMJ arthritis (p 0.01), but also with chronic damage (p 0.02). 62.5% of the patients with pain had synovial enhancement and 43.5% had effusion on MRI. Mouth deviation at MMO was also significantly associated with chronic damage on MRI (p 0.0005), so that 66.6% of the patients with mouth deviation had condylar deformity on MRI. A statistical trend of significance was observed between limited MMO and acute TMJ arthritis (p 0.09).

70.5% of the patients included had at least two positive clinical or physical examination items.

Conclusion: TMJ involvement is highly prevalent in JIA patients. Nevertheless, in our study, not all symptomatic patients had pathological findings on MRI due to inflammation. Pain while eating was the most common symptom and significantly associated with both acute and chronic damage on MRI. During examination, pain with MMO was the most common sign, but mouth deviation predicts chronic damage on MRI. Using the Wulffraat et al. suggested criteria of at least 2 positive scores, we detected TMJ involvement on MRI in 70.5% of the patients. The use of this screening protocol may guide the performance of the MRI in our daily practice.

Disclosure of Interest
None Declared

P47
CACP-syndrome in pediatric rheumatology. Clinical and instrumental diagnostic criteria
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Pediatric Rheumatology 2017, 15(Suppl 1) P47

Introduction: Juvenile chronic inflammatory arthropathy are a large heterogeneous group of the joint pathology which characterized by persistence arthritis. The chronic synovitis occur primary source and secondary nature. A number of syndromes maybe have similar features with JIA (SAPHO, PAPA, NAO, MCTO, Farber, CACP and other).

Objectives: CACP-syndrome is a rare autosomal-recessive disorder of non-inflammatory joint pathology caused by mutations of the PRG4 (proteoglycan-4, 1q25-31) gene leading to absence or loss of function lubricin (Marcelino et al 1999). Lubricin is a mucinous glycoprotein (~227 kDa) secreted by synovial fibroblasts and superficial zone chondrocytes that provided lubricating surface of the joint (Swann et al., 1985). There are known about 15 disease-associated mutations of the gene. Clinical symptoms are characterized by flexion contractions of the interphalangeal joints (camptodactyly), polyarthopathy and variable occurrence of coxa vara and pericarditis. Onset of the disease usually in early childhood with symmetrical camptodactyly of II, III, IV fingers, which are similar to stenotic ligamentitis. Classic manifestation of the syndrome is chronic synovitis visually manifested like juvenile polyarthritis with symmetric progressive contractions (hips, knees, ankles, elbows and wrists) with absence of morning stiffness, skin hyperthermia, joint swelling and enthesopathy. Mechanism of non-inflammation progressive articular effusion is associated with deficiency of the surfactant system of the joint. This fact determines low activity and less of effectiveness of the anti-inflammatory drugs.

Methods: Instrumental picture at first sight meets all the criteria of juvenile arthritis (ILAR) and prescribes like the persistence chronic synovitis with osteoporosis with the absence of traumatic and destructive articular changes. But there are no such typical reaction of the bone like rapid uneven ossification of cartilage model of the epiphysis and erosion of articular surfaces usually visualized at the X-ray image at children suffering from JIA. Moreover long-term course of CACP-syndrome doesn’t lead to formation deep bone erosion and...
arthrosis unlike at JRA. Anatomic dystrophic changes of the bones usually occur through the progressive effusion and high intra-articular pressure (coxa vara, deformation of caput femoris with large “open” cysts of the acetabulum and other). MRI shows the presence of exudative synovitis of the extremely nature with the absence of chronic proliferative changes and pannus. Synovial fluid is straw-colored, less viscous with minimum of quantity cell (less 0.5*10^9/l) and level of TNI-o less 50 pg/ml.

Results: Infancy camptodactyly and symmetric contractures with enthese-negative non-erosive chronic exudative polyarthritis non-effective to anti-inflammatory drugs in combination with dystrophic bone deformations are main markers of CACP-syndrome.

Conclusion: The CACP is a non-inflammatory joint disorder of the group of monogenic diseases. Introduction of the algorithms of examination children with atypical variant course of JIA molecular genetic testing of the gene proteoglycan-4 (PRG4) would allow to optimization approaches in diagnostic and therapy of CACP-syndrome.

Disclosure of Interest
None Declared

P48
Comparing articular cartilage thickness in JIA patients and healthy controls by ultrasonography
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Pediatric Rheumatology 2017, 15(Suppl 1)P48

Introduction: Juvenile idiopathic arthritis (JIA) is caused primarily by degeneration of the osseocartilaginous structures, due to the synovial inflammatory process. It is essential to closely monitor structural damage during the disease course. Joint cartilage is easily visualized with high-frequency ultrasonography (US).

Objectives: We aimed to compare ultrasound (US) measurements of joint cartilage thickness in 6 joints in children with JIA and healthy controls.

Methods: A cross-sectional study of bilateral grey-scale US by high frequency transducer (7.5-10) MHz cartilage thickness of the knee, ankle, and second metacarpophalangeal (MCP) joints was performed in patients with JIA and healthy children aged between 6 and 17 years. Medical records were reviewed for JIA subtype and state of disease. Clinical examination, including routine joint examination was carried. Disease activity was assessed with a parental questionnaire and physical examination by the physician

Results: A total of 31 patients (19 boys and 12 girls) as well as 31 healthy controls were included in the study. Nineteen patients had polyarthritis (12 had enthesis-related arthritis, 4 had extended oligo-JIA, 2 rheumatoid-factor negative pJIA, and 1 had rheumatoid-factor negative pJIA) and 12 had persistent oligoarthritis. Joint cartilage thickness was decreased in the knee, wrist, and second PIP joint in children with JIA compared with the healthy cohort. When subgroup analysis was done, 23 of 30 patients had mild arthritis and 7 of 30 patients had moderate to severe arthritis. Tmean ≥ 31 °C was used as a cut-off level between controls and moderate to severe arthritis, the sensitivity and specificity were 85.7% and 80.0%, respectively. HDI median (IQR) in arthritis group, non-arthritis group, and healthy controls were 0.64 (0.42), 0.66 (0.44), and 0.52 (0.28), respectively with no statistical difference, however; HDI seemed to be higher in moderate to severe arthritis when compared to mild arthritis. US also had high sensitivity (83.3%) and specificity (81.3%) with area under the ROC curve of 0.87 in detecting wrist arthritis if US score ≥ 1. The kappa score between clinical examination and US was good (Kappa = 0.63, p < 0.01). Wrist examination also had correlation with US (0.67, p < 0.01) and Tmean (0.36, p = 0.01).

Conclusion: US is a good tool for arthritis detection, whereas Tmean from IRT is useful for joint inflammation assessment especially in moderate to severe arthritis.

Disclosure of Interest
None Declared

P49
The comparisons between thermography and ultrasonography with physical examination for wrist joint assessment in juvenile idiopathic arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1)P49

Introduction: There are several ways to assess joint inflammation in juvenile idiopathic arthritis (JIA). At present, physical examination (PE) is the gold standard in detecting arthritis but it is difficult to perform especially in young children. Ultrasonography (US) is another good tool for joint assessment but it is an operator-dependent and time consuming. Infrared thermography (IRT), a non-invasive and radiation free imaging method, is a promising tool for arthritis detection. Previous studies showed correlation between joint inflammation and parameters of IRT, including thermographic index (TII), heat distribution index (HDI), and mean temperature (Tmean). However, there is limited data in using IRT for wrist joint assessment in JIA patients.

Objectives: To compare IRT and US with wrist joint examination in JIA patients.

Methods: This is a cross-sectional study. JIA patients in Rheumatology Clinic, Ramathibodi Hospital between November 2014 and October 2015 were enrolled into this study. Patients with fever were excluded. There were 15 healthy controls. Wrist joint assessment were performed by IRT, PE, and US, respectively. IRT was placed one meter distance from wrist joint in anterior view with neutral position. The images were processed by ThermaCam Researcher Pro 2.10 program and calculated HDI by Matlab software. Tmean, which obtained from the region of interest, was the mean temperature on dorsal surface of the wrist joint. US was performed by musculoskeletal radiologists. Gray scale was graded according to 4-point semi-quantitative scoring system and Power Doppler (PD) was graded by using The Outcome Measures in Rheumatoid Arthritis Clinical Trials (OMERACT) definition. Total scores of US in this study were the sum of Gray scale scores and PD scores, which were 6 points. Patients were classified into 2 groups based on clinical examination of wrist joints; non-arthritis and arthritis groups. Descriptive statistics, Mann-Whitney U test, Kruskal-Wallis test, ROC analysis, Spearman's correlation, Kappa agreement were used. This abstract is original and not previously published anywhere.

Results: Of 46 patients, 16 patients were in non-arthritis group and 30 patients were in arthritis group. Using IRT, median (IQR) of mean temperature (Tmean) in arthritis group [31.35 (2.20-30°C] were higher than non-arthritis group [30.55 (2.40°C] and healthy controls [30.10 (1.20°C] with statistical significance (P < 0.05). When subgroup analysis was done, 23 of 30 patients had mild arthritis and 7 of 30 patients had moderate to severe arthritis. Tmean ≥ 31 °C was used as a cut-off level between controls and moderate to severe arthritis, the sensitivity and specificity were 85.7% and 80.0%, respectively. HDI median (IQR) in arthritis group, non-arthritis group, and healthy controls were 0.64 (0.42), 0.66 (0.44), and 0.52 (0.28), respectively with no statistical difference, however; HDI seemed to be higher in moderate to severe arthritis when compared to mild arthritis. US also had high sensitivity (83.3%) and specificity (81.3%) with area under the ROC curve of 0.87 in detecting wrist arthritis if US score ≥ 1. The kappa score between clinical examination and US was good (Kappa = 0.63, p < 0.01). Wrist examination also had correlation with US (0.67, p < 0.01) and Tmean (0.36, p = 0.01).

Conclusion: US is a good tool for arthritis detection, whereas Tmean from IRT is useful for joint inflammation assessment especially in moderate to severe arthritis.

Disclosure of Interest
None Declared
PS0
Radiographic evaluation of joint space width compared to cartilage thickness as assessed by ultrasonography in MCP, PIP, and knee joints of children with juvenile idiopathic arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1)

Introduction: Joint space narrowing (JSN), is a measureable outcome of tissue damage caused by inflammation of the synovium in arthritis, and is a degenerative result of cartilage thinning. Usually conventional radiography is the preferred first choice modality to assess JSN. Within recent year’s ultrasonography have been validated for measurements of joint cartilage thickness in large and small joints in healthy children. Furthermore decreased joint cartilage thickness assessed by ultrasonography has been reported in children with JIA compared to a healthy cohort.

Objectives: To correlate measures of cartilage thickness assessed by US to the measures of joint space width (JSW) assessed by computer-ized radiography of the proximal cartilage site in the 2nd MCP, 2nd PIP, and knee in children with JIA.

Methods: Seventy-four children with JIA, aged between 5 and 15 years (median 11.3 yrs), 54 girls and 20 boys were included in the study. Cartilage measures of the MCP and PIP joints were assessed at the midline spot with a longitudinal scan with the joints in 900 flexion, whereas the knee joints were assessed by a suprapatellar transverse scan with the knee in maximal flexion measuring cartilage thickness at two spots: medial and lateral femoral condylar areas. Thus, US assessed only the proximal cartilage sites of the joints, whereas digital radiography assessed the total JSW. Readers of the JSW were blinded to the results of the US assessed cartilage thickness measurements.

Results: Onehundred and thirty-six 2nd MCP, 138 2nd PIP, and 146 knee joints were assessed. We found a high level of agreement between US and radiographic measures of cartilage thickness and JSW with Rho values between r = 0.82 to 0.86 (2nd MCP), r = 0.50 to 0.55 (2nd PIP), and r = 0.52 to 0.81 (knee), p < 0.001 for all eight assessed sites.

Conclusion: US measurements of cartilage thickness of only the proximal site of 2nd MCP, 2nd PIP and knee joints, correlated well to radiographic measurements of JSW in finger and knee joints of children with JIA. However, the difficulty of assessing the distal part of the joint cartilage may limit the use of US as a tool in the assessment of JSN.
PS3

Thermal point of care diagnostic tool for measurement of joint inflammation

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Introduction: Since the dawn of the medicine, heat (lat. calor) is one of the four main signs of inflammation. Although it is recognized more than 2,000 years ago, there is still no reliable, simple and objective method to measure the warmth of inflamed joint, which is therefore most often assessed only by clinical examination, with very low validity and reliability.

Objectives: To investigate the use of smartphone attached mobile thermography for the assessment of joint inflammation.

Methods: Twenty participants with various forms of arthritis and/or arthralgia participated in this pilot cross-sectional observational study. Average age of the participants was 10 years (2.5 - 17.5). In all of the participants Power Doppler vascularity signal of both knee joints was graded from 0-3 by EULAR certificated instructor in MSUS. FLIR ONE smartphone connected device was used to obtain infrared thermal image of every patient’s knee joint prior to the MSUS examination, without clothes and with same environment conditions (room temperature was set to 20 ° C). Acquired images were analyzed using FLIR Tools software for thermal image analysis with automatically set range of temperatures. On every image, both knee joints were selected using manual ellipse measurement tool and the data on average, minimum and maximum temperature inside the circle was collected. Subsequent statistical analysis of selected data was performed with Graphpad Prism software.

Results: In sixteen knee joints of nine patients a PD signal was above 0.5 (0.5-2.5), which was defined as an active inflammation, while in twenty-four knee joints of thirteen patients PD signal was 0, which was defined as no inflammation. Average value of average temperature in knee joints with inflammation was 33.72 °C (31.3 °C - 35.4 °C) and in patients without inflammation 29.81 °C (23.9 °C - 31.9 °C), average value of maximal temperature was 34.98 °C (32.7 °C - 36.9 °C) and 31.08 °C (25 °C - 33.3 °C), while average value of difference between maximal and minimal temperature was 4.73 °C (2.6 °C - 7.7 °C) and 2.96 °C (1.2 °C - 5.7 °C) respectively. Statistical analysis revealed significant difference in all of the measurements between knee joints with and without active inflammation (p < 0.05).

Conclusion: In our proof of concept study we investigated the usefulness of thermography with FLIR ONE, a cheap, portable and easy to use compact device, in joint inflammation assessment. For objective evaluation, MSUS with Power Doppler scoring of synovial vascularity was used. This method is almost a gold standard in joint inflammation assessment and it effectively measures increased blood flow inside the joint, which is one of the main causes of higher temperature. Preliminary results of our study suggested FLIR ONE thermography can be used as a bedside screening tool for joint inflammation, with a potential in assessing and quantifying degree of inflammation. This is in consistency with the results of a recent study which indicated the described method has a sufficiently high validity and reliability to be recommended for assessment of subclinical inflammation in patients with pressure ulcer and diabetics foot [1]. After confirmation in a larger cohort, our results could facilitate this innovative technique as a tool for joint inflammation assessment in everyday clinical practice.
Table 21 (abstract P54). See text for description

| Patient | ALL MCP (2-5) -DEMRIQ inflammation | Global clinical scores-AJ | Outcome |
|---------|----------------------------------|---------------------------|---------|
|         | Duration of QRS complex (msec)    |                          |         |
|         | Normal (-)                       | Pearson’s correlation     |         |
|         | 3 mo                             | 6 mo                      |         |
| 1       | 117,0 ± 151,1 ± 135,66            | Moderate (+)              | Switch DMARD |
| 2       | 83,16 ± 28,96 ± 64,89             | Very strong (-)           | Continue DMARD |
| 3       | 99,28 ± 46,16 ± 26,35             | Very strong (-)           | Improved later |
| 4       | 38,28 ± 104,71 ± 87,86            | Strong (+)                | Flare at 6 mo |
| 5       | 113,36 ± 102,6 ± 118,39           | Moderate (+)              | Flare at 6 mo |
| 6       | 29,28 ± 39,09 ± 58,12             | Weak                      | Flare at 2 mo |
| 7       | 38,1 ± 19,59 ± 56,47              | Very strong (-)           | Flare at 3 mo |
| 8       | 50,42 ± 96,76 ± 74,75             | Moderate (-)              | Flare at 3 mo |

P55
Value of magnetocardiography for diagnosis of myocardial damage in children with rheumatic diseases

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Pediatric Rheumatology 2017, 15(Suppl 1) P55

Introduction: According to a clinical registry frequency of cardiovascular system diseases ranged from 51% to 81% among 327 children with systemic diseases of connective tissue (SDCT). Injury of the heart is potentially life-threatening. Their initial signs cannot be detected by traditional diagnostic methods.

Objectives: Improving early diagnosis of cardiac lesions in children with rheumatic diseases

Methods: There are 25 healthy and 31 sick children of 10-17 years old with SDCT (SLE, JDM, MCTD), in which are absent active inflammation, endocrine pathology, and arterial hypertension, clinical and laboratory signs of carditis or cardiomyopathy were examined. Patients underwent magnetocardiographic examination, i.e. registration of heart magnetic field by SQUID-magnetometer at 36 points above the chest with 4 mm step, during 90 sec at single point. The inverse solution, i.e. reconstruction of current density vectors maps based on MCG-curves was done. The leading quantitative indicator of current density vectors maps is abnormal index (AI) the ratio of the total length of vectors directed in the “proper” direction to that have a different direction (%). To evaluate the processes of ventricular depolarization and repolarization this index was calculated during QRS complex and ST-T interval (AIQRS2 total and AIST-T total). Besides, some other parameters were calculated, namely the ratio of the actual duration of each QRS phase to its “proper” duration (Duration Ratio - DR) for each of four depolarization phases (DR1-D4) separately and for the whole process of depolarization (DRtotal), duration of complex QRS – DurQRS, the correlation coefficient Ccorr between all maps during each interval, duration of ST-T curve area from its beginning to the inflection point in percentage to the overall length of the ST-T interval (Asto).

Results: Statistically significant differences of average values of MCG indicators characterizing QRS complex were not found in the examined groups. However, individual analysis of normalized indicator of anomalous ventricular depolarization showed that it significantly deviates from the normal range in patients with active inflammatory process, mostly in front-apical area of myocardium (AI QRS2 70,15 ± 2,94% in patients, 56,4 ± 4,12% in healthy). This was accompanied by a reduction in the total duration of the QRS complex and the portion of normal complexes in the corresponding time intervals DTot 1,02 ± 0,04 s and in healthy 1,2 ± 0,09 s (by Dr1) respectively. DurQRS in patients with rheumatic diseases equal 0,089 ± 0,003 s (0,1 ± 0,04 s in healthy), Auto-corQRS 80,43 ± 0,73% (84,14 ± 1,5% in healthy).

Correlation coefficient decreased in patients with rheumatic diseases insignificantly (CorQRS 34,85 ± 1,01% in patients, 37,29 ± 7,01% in healthy). The analysis of the ST interval indicators showed reliable abnormalities in early repolarization of the vectors (increasing Adur 19,85 ± 1,61% (11 ± 6,1% in healthy), reducing CorrST 63,85 ± 7,61% (82 ± 4,1% in healthy), downward trend in AI ST-Total in patients 53,85 ± 3,35% (91 ± 0,4% in healthy children)). Comparing the results with standard ECG data it was revealed that the changes of repolarization processes are visualized up to 35% more frequently by MCG. Instant maps of current density vectors that obtained by MCG are more convenient for on-line visual assessment.

Conclusion: Non-invasiveness and sensitivity of MCG allow recommend it as a screening method for early detection of the heart injuries. The ability to visualize of electrophysiological active substrate makes the MCG as effective way to detect local changes into myocardium. Absolute safety of method makes possible its use for dynamical investigation.

Disclosure of Interest
None Declared

P56
Early detection and follow-up of morphological alterations of temporomandibular joint in juvenile idiopathic arthritis: high-resolution ultrasound

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Pediatric Rheumatology 2017, 15(Suppl 1) P56

Introduction: Ultrasonography could be an helpful screening method for the detection of TMJ involvement in children with JIA. It can be used and well tolerated even in very young children (2 years) without sedation or radiation risk.

Objectives: The aim of this work was to evaluate the usefulness of high-resolution ultrasound (US) in the early detection of temporomandibular joint (TMJ) alterations and to show normal and pathological imaging appearance of the TMJ in patients with Juvenile Idiopathic Arthritis (JIA). Although TMJ arthritis in JIA is frequently asymptomatic, the TMJ is particularly susceptible to damage. The TMJ is commonly involved in JIA and can lead to malocclusion, condylar alterations, facial deformity and restricted mouth opening.

Methods: High-resolution US examinations were performed in 130 patients with JIA. 34 patients (34%) had normal US appearance on both side. The sonograms were done using General Electric LOGIQ E9 equipment using real-time linear-array transducers with variable frequencies (9-18 MHz) by an experienced radiologist. Longitudinal and transverse view images were obtained. Dynamic video clips demonstrated condylar translation from “closed mouth” position to “open mouth” position. The movements can be recorded and compared between the left and the right side. Color Doppler US images furnish diagnostic information regarding hyperemia and synovial vascularity, thus allowed to study vascular signals in the soft tissues around the TMJ. Each joint was analyzed with regard to condylar irregularity, capsular thickness (effusion/synovial thickness), condylar disk dislocation, vascular increasing on Color-Doppler examination and symmetry in condylar translation between open- and closed-mouth position.

Results: High-resolution US examinations were performed in 130 patients with JIA. 34 patients (34%) had normal US appearance on both side. The sonograms were done using General Electric LOGIQ E9 equipment using real-time linear-array transducers with variable frequencies (9-18 MHz) by an experienced radiologist. Longitudinal and transverse view images were obtained. Dynamic video clips demonstrated condylar translation from “closed mouth” position to “open mouth” position. The movements can be recorded and compared between the left and the right side. Color Doppler US images furnish diagnostic information regarding hyperemia and synovial vascularity, thus allowed to study vascular signals in the soft tissues around the TMJ. Each joint was analyzed with regard to condylar irregularity, capsular thickness (effusion/synovial thickness), condylar disk dislocation, vascular increasing on Color-Doppler examination and symmetry in condylar translation between open- and closed-mouth position.

Results: In our clinical cases capsular thickness, enlargement of the intra-articular space, joint effusion, osteophytes, irregular condyle shape, disc displacement, erosions, increased color Doppler signals were detected. Limitations were especially related to the scarce accessibility of the medial part of the TMJ structures, and the need for trained operators. An important advantage especially in the case of
children is that US examination does not require conscious sedation or general anesthesia. Conclusion: The prevalence of TMJ involvement in patients with JIA varies widely (17–27%) and it is often asymptomatic even during the acute phase therefore consequently overlooked. Since inflammation in the TMJ can lead to joint deformity, early diagnosis and treatment of TMJ arthritis are particularly important. High-resolution US could be suggested as a screening tool to detect morphological alterations at an early stage of the JIA. Utilization of high-resolution US for TMJ disorder diagnosis in patients with JIA should increase due to relatively low costs and non-invasivity of the method, moreover it is a noninvasive tool. High-resolution US should be used for detecting inflammatory lesions before permanent TMJ destruction occurs, and monitoring disease progression and treatment response for a more effective therapy.

Disclosure of Interest
None Declared

Table 22 (abstract P58), See text for description

| PID (n) | Gender | ANA | RF | HLA association | Diagnosis of PID prior to arthritis (yrs) | Age at arthritis onset (yrs) |
|---------|--------|-----|----|-----------------|------------------------------------------|----------------------------|
| slgAD (2) | F | + | - | - | No | 2 |
| AT (1) | M | - | - | - | Yes/1 | 14 |
| APECED (1) | M | - | - | - | Yes | 6 |
| ALPS (1) | F | - | - | - | Yes/13 | 15 |
| FMF (1) | F | - | - | - | Yes/11 | 12 |

P57
Correlation between maxillary sinusitis and juvenile idiopathic arthritis: cone beam CT study
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This abstract is not included here as it has already been published.

Poster Session: Immunodeficiency and infection related arthritis

P58
Attending to warning signs of primary immunodeficiency diseases in rheumatology practice: focus on juvenile idiopathic arthritis-like disease
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Pediatric Rheumatology 2017, 15(Suppl 1):P58

Introduction: Primary immunodeficiency (PID) and autoimmunity phenomena may occur concomitantly in the same individual. The most common forms of PID disease with articular involvement are those associated with humoral defects such as X-linked agammaglobulinemia, common variable immune deficiency and selective IgA deficiency.

Objectives: The aim of our study was to systematically track a comprehensive array of PID in a cohort of 286 children with a presentation of chronic arthritis followed at the Immunology and Rheumatology outpatient clinic in Children’s Hospital Zagreb.

Methods: Patient’s charts were reviewed for the following immunological investigations performed as a part of the diagnostic algorithm for juvenile idiopathic arthritis: IgG, IgA, IgM, IgE, IgG subclasses, C3, C4, CH100, ANA, ANA, ENA-profile, antiphospholipid antibodies, rheumatoid factor (RF) and HLA-typing. Additional screening and specialized laboratory tests were performed in patients who presented with the recently revised warning signals for primary immunodeficiency.

Results: Six patients were identified with immunity defects compatible with classical PID. Two patients had selective IgA deficiency (slgAD), while one patient with the following PIDs in each group was found: ataxia telangiectasia (AT), autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy (APECED), autoimmune lymphoproliferative syndrome (ALPS) and familial Mediterranean fever (FMF).

Conclusion: Our findings further emphasize that PID and JIA may coexist and patients with autoimmune diseases should be carefully monitored for the presence of PID and vice versa. To the best of our knowledge, we also describe a first case of a child with typical ataxia telangiectasia who developed JIA.

Disclosure of Interest
None Declared

P59
Familial Acicardi-Goutieres Syndrome with SAMDH1 mutation – phenotype variation in 2 sisters from South Africa
Monika M. Esser1, Deepthi R. Abraham2, Craig Kinnear2, Glenda Durrheim2, Mike Urban2, Eileen Hoal2, Yunick Crow3
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Pediatric Rheumatology 2017, 15(Suppl 1):P59

Introduction: Acicardi-Goutieres Syndrome (AGS) is a rare genetic disorder with broad phenotype spectrum. Access to molecular investigation confirmed AGS for different clinical manifestations in 2 sisters of non-consanguinous parents.

Objectives: The value of a confirmed molecular diagnosis for a rare disease which would otherwise have remained undiagnosed is illustrated in our patients.

Methods: We describe 2 sisters with unusual clinical features resembling childhood SLE and targeted molecular investigation. The index patient presented with hemiplegia and complex partial seizures in infancy, diagnosed as moyamoya syndrome on imaging. At 4 years of age she was referred to the paediatric rheumatology service with interface dermatitis, vasculitic lesions on her ears and fingertips and deteriorating gait. She tested negative for antinuclear antibody (ANA) and anti double-stranded DNA (dsDNA) but had positive antiphospholipid serology. She was anticoagulated with Warfarin, responded well to Chloroquine and Azathioprine for control of extensive vasculitic rash and Sodium Valproate for her seizure control. She was then diagnosed with multidrug resistant Tuberculosis and superficial staphylococcal abscesses of the spine which resolved on antituberculous and antibacterial treatment. The unusual combination of her clinical findings prompted molecular investigation. The older sister was found to have a longstanding history of vasculitic skin lesions and threatened gangrene of fingertips on presentation. She was severely stunted with delayed puberty, a feature not typically described with AGS. Skin biopsy showed perivascular-interface dermatitis. Antiphospholipid antibody screen and ANA were negative. She responded well to Nifedipine, Chloroquine and low dose Aspirin. She was subsequently diagnosed with glaucoma, which is associated with AGS and developed progressive contractures of interphalangeal joints with bone resorption on X-Ray.

Results: Blood samples from patients, their parents and unaffected brother were submitted to the Genetic Medicine Developmental Bio-medicine Unit, Faculty of Medical and Human Sciences, University of Manchester. The 2 affected individuals carried a homozygous Exon 15 c.1681_1682delAG; p.Ser561Phefs*61 in SAMDH1 on chromosome 20. The parents and the brother were heterozygous for this variant. Variants were confirmed in the Division of Molecular Biology and Human
Aseptic arthritis in children associated with primary immune deficiencies

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Pediatric Rheumatology 2017, 15(Suppl 1):P60

Introduction: Defects of immune cells in children with primary immunodeficiency (PIDS) development may lead to formation of infectious, allergic, autoimmune and oncologic immunopathological syndromes. Were described both infectious and noninfectious arthritis.

Objectives: Identify the features of articular lesions in PIDS.

Methods: Were retrospectively analyzed clinical register data of patients with joint syndrome, hospitalized in 2000-2015.

Results: Among 901 children with joint syndrome in 21.97% its character at a pre-hospital stage was regarded incorrectly. Hyperdiagnostics was frequent in JIA (11.09%). In 13 cases (1.4%) the diagnosis of PIDS was established after an additional clinical examination. Most often (7 cases) joint lesions were observed at selective immunoglobulin A deficiency. Previous infectious was not revealed in these cases. Manifestation of joint syndrome 2-11 y, its features corresponded to JIA: 2 entezit-arthritis with inflammatory bowel diseases, 3 oligoarthritis, 1 sacroileitis, 1 SJA. Immunological examinations: serum IgA: 0.05 g/l, IgM, G, subclasses, B-cells within age values, was found depression of T-lymphocytes, in 4 cases - ANA, RF and HLA B27 weren't found, CD3+ 34.5%, CD3 +CD4 + 16.89%. RTLB FGA 28%. Cariotype 46XY, Fish 22 +. In a 1 case with a HIDS during patient had aphthous stomatitis, lymphadenopathy, gastroduodenal pathology, transient skin eruptions and ankle joints swelling without active sinovitis or radiological changes. ESR– 13 mm/h, CRP 24 mg/ml, Ig L, phagocytosis, lymphocytic subpopulations were normal, IgD 620 IU/ml HLA B27, ANA, AB DNA and desmoglein weren't detected. In clinic were also examined 2 children with CINCA/NOMID syndrome. One child had dermat and fever syndromes, lymphadenopathies, bilateral congenital cataract, an optic nerves atrophy, conjunctivitis in the anamnesis, and delays of physical and psychomotor development, pathognomonic joint lesions since age of 1 year. The mutation of CIA51 is taped in both cases. In all described PIDS cases there weren't detected microorganisms in blood or synovial fluid, it was now rising of pro-calcitonin level.

Conclusion: In PIDS patients rheumatoid-like joint lesions can become perceptible. In patients with a joint syndrome the careful assessment of a phenotype and the anamnesis and they need screening immunologic assessment before establishment of the diagnosis of JIA.

Disclosure of Interest
None Declared

P61
Retrospective analysis of laboratory data and treatments of 43 adult patients followed up with the diagnoses of acute rheumatic fever (ARF) and post-streptococcal reactive arthritis (PSRA)

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Pediatric Rheumatology 2017, 15(Suppl 1):P61

Introduction: Although rarer, ARF and PSRA are still seen in adults. There are no exact criteria to differentiate ARF from PSRA. Usually the arthritides that develop after a documented streptococcal infection, characterized by acute arthritis, erythema nodosum and not meeting modified Jones criteria are described as PSRA [1].

Objectives: Over the years, the opinions on the clinical findings of ARF and PSRA, and the clinical approach to those diseases changed. In this retrospective study, we aimed to analyze 43 adult patients that were followed up with the diagnoses of ARF or PSRA in a secondary and a tertiary hospital.

Methods: 29 patients that had been followed up with the diagnoses of ARF or PSRA in the Rheumatology Clinic of Ankara Numune Education and Research Hospital and 14 patients that had been followed up with the same diagnoses in Mersin State Hospital in the previous 1 year, and a total of 43 patients were included in the study. Patient files and their data on electronic data recording system of the hospital were examined to analyze the clinical and laboratory findings of the patients and their treatments.

Results: 6 (14%) patients that met the modified Jones criteria were regarded as ARF, and the remaining 37 (86%) patients were regarded as PSRA. Among ARF patients, there were 4 F and 2 M with a mean age of 28.3 ± 7 years. The patients with ARF could not be compared with PSRA patients since their number was small. However, it was noteworthy that 4 (67%) patients with ARF were recurrent patients that had their first attacks during childhood, and had cardiac valvular disease. The patients with PSRA consisted of 31 F and 6 M with a mean age of 30.4 ± 5.8 years. None of the patients had clinical cardiac involvement. None of 11 patients that had echocardiography had rheumatic valvular involvement. The most frequently involved joints were ankle (50%), knee (38%), wrist (19%), and MTP (8%) joints. Migratory arthritis was seen in only 2 (5.4%) patients. There was mono in 27%, oligo in 57%, and polyarthritis in 16% of the patients. Two (5.4%) patients had concomitant erythema nodosum. There was history of tonsillitis in 46% of the patients previous to the onset of arthritis. The mean period between tonsillitis and arthritis was 25 ±
9 days in those patients. The mean duration of arthritis at the time of admission was 21 ± 13 days. Fever was present only in 4 (10.8%) patients. None of the patients had erythema marginatum, subcutaneous nodules or chorea. The mean ESR, CRP, and ASO values of the patients on admission were 57.8 ± 28.6 mm/h, 6.34 ± 5.3 mg/dL, and 971.5 ± 505.3 IU/mL, respectively. The ESR, CRP and ASO values on follow up were 29.5 ± 21 mm/h, 1.6 ± 2.2 mg/dL, and 822.5 ± 468 IU/mL, respectively. Repository penicillin was administered to 84%, oral and then repository penicillin were administered to 11%, and macrolides were administered due to penicillin allergy to 5% of the patients. Anti-inflammatory treatment administered to the patients included 20 mg/day or more oral corticosteroids (CS) in one patient, less than 20 mg/day oral CS in 28 patients (75.7%) (prednisolone equivalent), repository CS in 2 (5.4%) patients, and NSAID alone in 6 patients (16.2%). The mean use of CS was 8.5 ± 3.8 weeks. Chronic arthritis was seen in only 3 (8.1%) patients in long term. 

Conclusion: Currently, ARF and PSRA are not still rarely seen diseases. It was noteworthy that typical ARF findings including fever, cardiac involvement, migratory erythematous arthritis, and skin involvement were rarely seen in our adult patients with PSRA, however joint findings continued longer, the ASO titer was over 500 IU in 92% of the cases, and most of the patients required low-dose corticosteroids instead of aspirin or NSAIDs.

References

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Disclosure of Interest
None Declared

Poster Session: JIA (oligo, poly, psoriatic) I

P62
Musculoskeletal symptoms at the onset of pediatric tumors and predictive features in the differential diagnosis with juvenile idiopathic arthritis: preliminary analysis of a multicenter, prospective, observational study

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Presenting author: Adele Cinvio

Pediatric Rheumatology 2017, 15(Suppl 1):P62

Introduction: The onset of some pediatric tumors may be characterized by musculoskeletal symptoms that can mimic rheumatic diseases like juvenile idiopathic arthritis (JIA). This could lead to inappropriate steroid treatment or immunosuppressive therapy with a delay in the diagnosis.

Objectives: To study prevalence and features of musculoskeletal symptoms at the onset of pediatric tumors diagnosed at the Italian Association of Pediatric Hematology and Oncology (AIEOP) Centers; to identify predictive factors of malignancy comparing the new cases of tumors with early musculoskeletal symptoms with JIA diagnosed in the same period at the Pediatric Rheumatology Centres of Italian Society of Paediatrics (SIP).

Methods: We analyze electronic case report form of pediatric patients (<16 years) with new diagnosis of cancer or JIA collected through a web based data system.

Results: 420 patients were enrolled between 01.05.2015 and 15.03.2016. 147 of them were affected by JIA and 272 by tumor: 50.4% acute leukemia (AL) and 49.6% solid tumors. Musculoskeletal symptoms were found in 81 (29.8%) patients with tumor; higher frequency of symptoms were found on patients with bone sarcoma (100%), neuroblastoma (41%) and AL (34,3%); Arthritis was observed in 29 (10.7%) patients with tumor including 18 AL and 11 solid tumors. Male/female ratio was 1.65. The mean number of involved joints was 1.66 (range 1-6). The most frequently involved joints were hips, knees and ankles. Joint involvement (arthrits and/or arthralgia) was observed in 60 (22%) patients with tumor including 38 AL and 22 solid tumors. We compared clinical features of patients with tumors and joint involvement with those of patients with JIA and we found that arthralgia, bone pain of the legs, pain of the spine, unproportioned pain and refusal to deambulation were significantly associated with neoplastic disease. Finding of arthritis, arthrits and arthralgia, morning stiffness, limp and the presence of only musculoskeletal symptoms were significantly associated with JIA.

Conclusion: With a preliminary analysis of data we found that musculoskeletal symptoms at the onset of tumors were present in about 30% of patients. These symptoms have features strictly comparable with JIA so they may be useful in the differential diagnosis with JIA.

Disclosure of Interest
None Declared

Table 23 (abstract P62). Comparison of clinical parameters between tumors with joint involvement and JIA

| Tumors with joint involvement (60 pts) | JIA (147 pts) | p-value |
|---------------------------------------|-------------|--------|
| n° | % | n° | % | |
| Arthritis | 23 | 38.3 | 107 | 72.8 | 0.000 |
| Bone pain | 24 | 40.0 | 4 | 2.7 | 0.000 |
| Nighttime pain | 7 | 11.7 | 6 | 4.1 | 0.057 |
| Pain of the spine | 15 | 25.0 | 12 | 8.2 | 0.002 |
| Unproportioned pain | 8 | 13.3 | 2 | 1.4 | 0.001 |
| Morning stiffness | 1 | 1.7 | 80 | 54.4 | 0.000 |
| Limp | 25 | 41.7 | 89 | 60.5 | 0.014 |
| Only musculoskeletal symptoms | 16 | 26.7 | 116 | 78.9 | 0.000 |

P63
Differences in disease phenotype, management and outcomes of children with juvenile idiopathic arthritis throughout the world – analysis of 8,325 patients enrolled in the EPOCA study

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Pediatric Rheumatology 2017, 15(Suppl 1):P63

Introduction: Several epidemiologic surveys have documented a remarkable, yet unexplained, disparity in the prevalence of juvenile idiopathic arthritis (JIA) subtypes among different geographic areas or ethnic groups. Moreover, the therapeutic approach to JIA is not standardized and the availability of the novel and costly biologic medications is not uniform throughout the world. This disparity may have significant impact on disease outcome.
**Objectives:** The multinational study of the EPIdemiology, treatment and Outcome of Childhood Arthritis (EPOCA) study is aimed to obtain information on the variability of JIA phenotypes in different geographic areas, the therapeutic approaches of pediatric rheumatologists practices in diverse countries, and the disease status and outcome of children with JIA currently followed worldwide.

**Methods:** Participation in the study was proposed to all pediatric rheumatology centers that are part of the Pediatric Rheumatology International Trials Organization (PRINTO), and to several centers in the US and Canada. Each center was asked to enroll 100 consecutive JIA patients or all consecutive patients seen within 6 months. Each patient received a retrospective and cross-sectional assessment. Parent- and child-reported outcomes were recorded through the administration of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Participating countries were grouped into 6 geographic areas.

**Results:** Currently, 8,325 patients from 44 countries have been entered in the web database. Comparison of main epidemiology, treatment, and outcome features across the different geographic areas is presented in the table.

**Conclusion:** These results provide further evidence of the wide difference of JIA characteristics across geographic areas in terms of age at disease onset, subtype prevalence, and frequency of anterior uveitis. Overall, patients living in non-Western countries had higher levels of disease activity and cumulative damage than patients followed in North America and Western Europe. This disparity in disease outcomes may be partially due to differences in the availability or affordability of biologics, as usage of these medications was more common in Western pediatric rheumatology centers.

**Disclosure of Interest**

A. Consolaro: None Declared, N. Ruperto Grant / Research Support from: The G. Gaslini Hospital, which is the public Hospital where I work as full time public employee, has received contributions from the following industries for the coordination activity of the PRINTO network: BMS, GlaxoSmithKline, AbbVie, Biogen, Genentech, Moderna, Pfizer, Sanofi Aventis, and Biogen. This money has been reinvested for the research activities of the hospital in fully independent manners besides any commitment with third parties.

**Table 24 (abstract P63). See text for description**

|                      | Africa | Asia | Eastern Europe | Latin America | North America | Western Europe |
|----------------------|--------|------|----------------|---------------|---------------|----------------|
| N                    | 261    | 874  | 2587           | 814           | 422           | 3367           |

| Median (IQR) age at onset years | 7 (3.6–10.5) | 5.7 (2.9–9.2) | 6.4 (2.9–10.5) | 6.6 (3.6–10.3) | 8.1 (3.7–11.1) | 4.1 (2.2–8.7) |

**P64**

**Changes in switched memory B cells are associated to the therapeutic response to TNF inhibition in JIA**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P64**

**Introduction:** Both the innate and adaptive branches of the immune system have been implicated in the development of the various forms of JIA. Evidence suggests that B cells may be involved in oligoarticular and polyarticular JIA.

**Objectives:** To investigate possible abnormalities in B lymphocyte subsets in patients with oligo- and poly-JIA and their correlation with clinical features and response to treatment.

**Methods:** 107 patients diagnosed with oligo- or poly-JIA were studied and compared with 304 age-matched healthy children (HC). Peripheral Blood Mononuclear Cells (PBMCs) were isolated, stained with different sets of fluorochrome-conjugated antibodies and run on a BD LSRFortessa-X-20.

**Results:** Switched memory B cells were found to be significantly increased in JIA patients when compared to age-matched HC [% of CD19: 13.16% (±5.5) and 7.33% (±4.2) respectively, p < 0.0001]. The other subpopulations of B cells were not increased. Switched memory B cells were higher in JIA patients than HC at any given age. Switched memory B cells were approximately 70% of the B cells found in JIA synovial fluid. We also found that synovial B cells of ANA positive patients produced antimicrobial antibodies. There was no association of the number of switched memory B cells with disease activity, number of joints with active arthritis or presence of iridocyclitis. In patients treated with methotrexate (MTX) alone there was a progressive increase in switched memory B cells with a positive correlation with the duration of treatment (Spearman r = 0.36; p = 0.0368). On the other hand, in patients treated with MTX and TNF inhibitors (TNFi) there was a negative correlation with treatment duration, however, it was not significant (Spearman r = -0.098; p = 0.27). Notably, during follow-up, in patients treated with MTX alone, the number of switched memory B cells
increased significantly regardless of the presence of active (+42.4% ± 35) or inactive disease (+30% ± 34). In patients treated with MTX + TNFi who showed persistent active disease or disease flare the number of switched memory B cells increased significantly (+41.58% ± 32). In sharp contrast, in patients on MTX + TNFi with inactive disease, the increase in switched memory B cells was significantly lower (+9.8% ± 25) (Table). To take into account time as a variable, we calculated the slope of the increase in switched memory B cells over 12 months, as a measure of the rate at which switched memory B cells increased over time. The increase of the rate of switched memory B cells in patients with inactive disease on TNFi (0.09 ± 0.34) was significantly lower when compared to patients treated with TNFi who showed persistent active disease or disease flare (0.39 ± 0.32) and was comparable to that of HC (0.13 ± 0.06) (Table).

Conclusion: Switched memory B cells are increased in patients with oligo- and poly-JIA. Treatment with TNFi slows the increase of switched memory B cells over time in comparison to patients treated with MTX alone. Moreover, patients that responded to TNFi maintaining a persistent inactive disease during follow up showed a reduction in the rate of increase of switched memory B cells over time. Switched memory B cells may be involved in the development of oligo- and poly-JIA and their change over time is associated to the clinical response to TNFi.

Disclosure of Interest
None Declared

Table 25 (abstract P64), See text for description

|              | MTX with active disease | MTX with clinical remission | t-Test (p-value) | MTX with active disease | MTX + TNFi with clinical remission | t-Test (p-value) |
|--------------|-------------------------|-----------------------------|------------------|-------------------------|-----------------------------------|------------------|
| % change from baseline of switched memory B cells (±SD) | -42% (±35) | 29% (±34) | 0.55 | 41% (±32) | 9.8% (±25) | 0.0047 |
| Slope (indicating the mean % change of switched memory B cells in 12 months) (±SD) | 0.43 (±35) | 0.42 (±38) | 0.95 | 0.39 (±32) | 0.09 (±34) | 0.015 |

P65
Towards stratified medicine in juvenile idiopathic arthritis

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Introduction: Despite the efficacy of current biologic agents for the treatment of juvenile idiopathic arthritis (JIA) in many children, for others, treatment is ineffective and they continue to suffer active disease and disability. There are an increasing number of biological agents available but we have no validated biomarkers of treatment response with which to predict response and guide therapy choices. In order to develop a rational, evidenced-based approach to support choices of therapy for these children (stratified medicine), multi-center, international collaboration is vital.

Objectives: An international working meeting focused on stratified medicine in JIA, was held. The objectives of which were to reach consensus on how to overcome barriers and hurdles to robust, collaborative clinical/translational research, and to foster effective and successful collaboration.

Methods: An international group of experts (paediatric rheumatologists, epidemiologists, geneticists, immunologists and translational scientists), from 8 countries and 17 institutions, representing major national and international networks, recently came together for two days to drive progress in stratified medicine in JIA. Prior to the meeting all investigators contributed to a priority-setting exercise to define the most important and feasible research questions, hurdles and possible solutions. These responses were grouped into themes, for both questions and hurdles, and then participants were asked to rank each theme in order of their perceived importance. Currently available tools, such as appropriate biobank protocols from previously established networks within pediatric rheumatology were collated.

Results: The key elements of a new framework to guide processes for working together were agreed, a new International collaborative study was proposed, and a declaration on principles of Research Collaboration for Stratified Medicine in Childhood Rheumatic Disorders – termed the London Declaration, was agreed and signed by all.

Conclusion: The London Declaration on Stratified Medicine Research for JIA and the principles it represents, should accelerate progress towards a stratified treatment approach not only for children with JIA but for all childhood rheumatic diseases.

Disclosure of Interest
None Declared
Safety of adalimumab in pediatric patients with polyarticular juvenile idiopathic arthritis, enthesitis-related arthritis, psoriasis, and Crohn’s disease

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Presenting author: G. Hornell

Pediatric Rheumatology 2017, 15(Suppl 1):P66

This abstract is not included here as it has already been published.

Flares after withdrawal of biotherapies in JIA: clinical and laboratory correlates of remission duration

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Presenting author: Gabriele Simonini

Pediatric Rheumatology 2017, 15(Suppl 1):P67

Introduction: Information regarding the history of patients with JIA after systemic treatment withdrawal would be helpful in driving the choice of duration therapy. While some information in this regard exist for second-line drugs (eg methotrexate), studies on biologics are lacking.

Objectives: Aim of our study was to assess the time on remission after discontinuing biologic therapy in a retrospective, comparative, multicenter, cohort study of patients with JIA.

Methods: Among 349 JIA patients followed in three tertiary care centres and treated with biologics (started before age 18 y), 135 (103 F, 32 M; median age 15 years) achieved remission and could discontinue such treatment. Only first cycle of biologics was considered for this study: 87 received Etanercept, 27 Adalimumab, 12 Infliximab, 7 Anakinra, 1 Rituximab, and 1 Abatacept. Primary outcome was to assess, once remission was achieved, the time on remission up to the first flare after discontinuing treatment. Time to achieve remission once biologic drug was started, corticosteroid usage, and time on remission on therapy before discontinuing all treatments were also considered, along with demographic, clinical, and laboratory data. Mann–Whitney U-test, Wilcoxon signed-rank test for paired samples, chi-square, and Fisher’s exact test, when appropriate, were used to compare data. Pearson and Spearman correlation tests were used to determine correlation coefficients for different variables. In order to identify predictors of outcome Cox regression model and Kaplan–Meier curves were constructed, each one at mean of entered covariates.

Results: The vast majority of enrolled patients flared after stopping treatment with biologics (102/135, 75.6%) with a median follow-up time on remission of 6 months (range 1–109). Considering the time point one year after treatment discontinuation 43/135 (31.9%) patients were still in remission. For this group, remission lasted for a median period of 53 months (range 12–109): 30/48 with polyarticular onset, 7/27 oligoarticular extended, 6/35 oligoarticular persistent, 2/6 enthesitis-related arthritis, 2/7 psoriatic arthritis, and 8/12 systemic onset JIA. A higher probability of maintaining remission after discontinuing treatment was present in systemic onset disease compared to the rest of JIA patients (Mantel-Cox $x^2$ 5.48, p < 0.02). ANA positivity was associated with a higher probability of flare (Mantel-Cox $x^2$ 4.17, p < 0.04). Patients who received biologics > 2 years after achieving remission had a higher probability of maintaining such remission off therapy (median 21.7 ± 3.4 months vs 13.4 ± 2.3; Mantel-Cox $x^2$ 6.86, p < 0.009). No other clinical variable, including total length of treatment and type of treatment, resulted significantly associated with a long-lasting remission.

Conclusion: Even if in a retrospective study, we showed that patients with ANA positivity and those who stop treatment before two years of remission have a higher chance of relapsing after biologic withdrawal and that patients with systemic onset disease are less likely to flare.

Disclosure of Interest None Declared

An active proNGF-p75NTR axis plays a pro-inflammatory role in chronic arthritis

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Pediatric Rheumatology 2017, 15(Suppl 1):P68

Introduction: Inflammation has been associated with a marked increase in the production of nerve growth factor (NGF) in tissues of patients with a variety of chronic inflammatory diseases, including rheumatoid arthritis and juvenile idiopathic arthritis (JIA). However, the effects of NGF, its immature form proNGF, and their receptors, TrkA and p75NTR, in regulating cells and mediators during inflammatory responses and their contribution to the pathogenesis of arthritis are not yet defined.

Objectives: To evaluate the role of proNGF, mature NGF and their receptors in the modulation of inflammatory response in chronic arthritis patients.

Methods: Seventy-five JIA patients, according to the International League of Associations for Rheumatology classification criteria, were divided in persistent oligoarticular, extended oligoarticular or polyarticular subtypes. The expression of the NGF receptors, p75NTR and TrkA, in peripheral blood (PBMC) and synovial fluid (SFMC) mononuclear cells from JIA patients and healthy children was analyzed using real time-PCR and Western Blot. In ex vivo experiments JIA SFMC were treated with mature NGF or proNGF and the effects on intracellular signaling pathways and cytokine production were assessed by Western Blot, real time-PCR and ELISA.

Results: p75NTR is the main NGF receptor expressed in JIA PBMC and SFMC while TrkA is the prevalent NGF receptor in healthy donor mononuclear cells. The increased expression of p75NTR and the concomitant decrease in TrkA, at both the mRNA and protein levels, resulted in a highly abnormal ratio between p75NTR and TrkA in JIA cells. The highest expression of, not only p75NTR, but also sortilin, the p75NTR co-receptor, was found in JIA SFMC. When patients were divided according to JIA subtypes the highest p75NTR expression levels were found in SFMC of patients with polyarticular JIA. The lowest p75NTR expression levels characterized SFMC from persistent oligoarticular patients while patients with extended oligoarticular JIA showed intermediate p75NTR levels. A significant correlation was found in JIA patients between the p75NTR mRNA expression levels in SFMC and the number of active joints at sampling and the levels of...
Humoral and cellular responses to varicella vaccination of patients with pediatric rheumatic diseases – preliminary results of a vaccination study

**Introduction:** Vaccination can be beneficial in children with pediatric rheumatic diseases – possibly beneficial vaccines for PRD patients is the varicella zoster virus (VZV) vaccination which can be used to prevent chickenpox and shingles. An adequate cellular immune response is especially important for effective protection against VZV.

**Objectives:** To assess the humoral and cellular responses to the live-attenuated VZV vaccination in children with juvenile idiopathic arthritis (JIA), juvenile dermatomyositis (JDM) or juvenile scleroderma (JScl).

**Methods:** This study was initiated as part of a Dutch-Brazilian collaboration. Patients who did not respond to the first vaccine were eligible for a second vaccination. VZV-specific antibody (VZV-Ab) concentrations were measured with a standardized Lumix-assay. IFN-gamma ELISPOT assays were performed to assess VZV-specific T cell responses. B-cell maturation, T-B cell interaction and T-cell subsets were assessed using flow cytometry.

**Results:** The cohort consisted of 54 patients (9 oligo-JIA, 22 poly-JIA, 11 sJIA, 7 JDM, 5 JScl) and 18 healthy controls (Table 26). There were no disease flares or overt varicella infections reported after vaccination.

**Conclusion:** In this preliminary analysis of the immune responses to VZV vaccination, the response of PRD patients seems comparable to that of healthy controls. Cellular immune responses to the VZV vaccine are under analysis.

**Disclosure of Interest**
None Declared

**Table 26 (abstract P69), Patient and control characteristics at first dose of VZV-vaccine**

| Disease type | Patients (n = 54) | Healthy controls (n = 18) |
|--------------|------------------|--------------------------|
| Systemic JIA | 11 (20%)         | N/A                      |
| Oligo JIA    | 9 (17%)          | N/A                      |
| Poly JIA     | 22 (41%)         | N/A                      |
| JScI         | 5 (9%)           | N/A                      |
| JDM          | 7 (13%)          | N/A                      |
| Age at vaccination (median, range) | 5 (0–20) | 8.5 (3–18) |
| Medication use | MTX 98%       | N/A                      |
|              | MTX dose (median, range) | 15 (10–27) |
| Corticosteroids | 43%            | N/A                      |
| Steroid dose (median, range) | 6 (3–20)    | N/A                      |
| Biologics    | 6%               | N/A                      |
| Responder    | 17               | 6                        |
| Nonresponder | 18               | 12                       |

**P70**

**Share update – 508 families view on standard of care for children with Juvenile Idiopathic Arthritis in Europe and Israel**

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Pediatric Rheumatology 2017, 15(Suppl 1): P70
Introduction: Since 2013 PRES has initiated an European project called SHARE (Single Hub and Access point for Paediatric Rheumatology in Europe, PI Wulfraat). The project aimed to identify the specific needs for optimal care in pediatric rheumatic diseases (PRD) in each European country, and provide these countries with recommendations for the care of these groups of children. Optimal care includes diagnosis, management of disease and providing both drug and non-drug therapies. The European countries were grouped regarding geographic and economic characteristics into West and East-Europe, in order to analyze if there are important differences in health care of these regions. Israel was the first non-European country to participate of the SHARE survey.

Methods: In order to identify the specific needs for the optimal care in PRD, patients and doctors surveys have been performed at pediatric rheumatology centers and patient organizations involved in providing care for these children. Families of patients with diagnosis of JIA according ILAR (International League of Associations for Rheumatology) criteria answered a single time-point survey that encompasses information about specialists referral patterns and referral delays, disease diagnosis, multidisciplinary health care team follow up, access to treatment, participation in clinical research and transition of care to the adult service. Families filled in the questionnaires anonymously on PRINTO (Pediatric Rheumatology International Trials Organisation) website.

Results: Until May 2016, 508 families from 22 different countries answered the questionnaires: 350 families from West-Europe, 129 from East-Europe and 29 from Israel. The table shows the participant countries and the number of answered surveys of each country. The large difference in the number of answered questionnaires between West and East-Europe is a reflex of their population, as West is three times more populous than East. The families that replied this survey may present a bias to urban population, where the pediatric rheumatology centers are. Patients characteristics regarding gender, age of diagnosis and JIA subtypes are similar in East and West-Europe, as well as in Israel. Families view showed good access to pediatric rheumatologists, uveitis screening and high cost medications in all these regions. The main differences were found in pain and fatigue management, information about transition for the adults service, use of supportive care and knowledge about patient organizations. It seems that these items were less emphasized in East-Europe.

Conclusion: The families view is extremely important to identify the topics on standard of care that need improvement, in order to achieve a high quality health care in Europe. A substantial number of replies (508) have been obtained from 22 different countries. The results showed good access to pediatric rheumatology care in Europe and Israel, although tools like patient organizations and transition care were less used by East-Europe. For the future, the aim of SHARE is not only to expand the families survey to a larger number of European countries, but also to another continents.

Disclosure of Interest
None Declared

Table 27 (abstract P70). See text for description

| West-Europe | East-Europe | Asia |
|-------------|-------------|------|
| Netherlands | 100         | 28   | 21   | 29   |
| Belgium     | 16          | 1    | 1    | 2    |
| Denmark     | 34          | 1    | 1    | 20   |
| Sweden      | 1           | 26   | 5    | 5    |
| United Kingdom | 33 | 5   | 5    | 33   |
| Ireland     | 26          | 20   | 17   |      |
| France      | 59          | 1    | 30   |      |
| TOTAL = 350 | TOTAL = 129 | TOTAL = 29 |
P72
Development of the first video game aimed at children suffering from Juvenile Idiopathic Arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1):P72

Introduction: In France, Juvenile Idiopathic Arthritis (JIA) affects between 2 000 and 4 000 children. The physical and psychological impact of this chronic condition can be significant, including difficulties in the family environment, as well as at school. Whilst the arrival of biotherapies has revolutionised the care of these children, educational support remains necessary to increase their independence and improve their quality of life.

Objectives: The development of e-health, in the context of the democratisation of communication tools, has led us to create a Serious Game as educational support for children.

Methods: The game has been developed following the ADDIE pedagogical engineering model. The game will be translatable in all languages and made available to foreign associations free of charge. A steering committee was formed, composed of the parents association that initiated the project, a paediatric rheumatologist, and a video game agency.

The National Association for the Defence against rheumatoid arthritis manages the logistics and monitoring. The rheumatologist creates the scientific context, in collaboration with the association. The developers make artistic and gaming suggestions. The committee sets the direction: format, themes, key messages, graphics, animations and game sequences. Learning is validated through multiple choice quizzes. The association KOURIR (association for parents of children with JIA) will test the beta version. Adaptations will then be made before the game’s release.

Results: The committee decided to create an adventure game with interactive activities. The following options were selected: The game is developed using the Unity® engine for computers, smartphones and tablets, allowing for use anywhere (Apple, Android). The engine is designed to support all languages. The topics of the hospital process, treatment, complimentary examinations and rehabilitation are all discussed with corresponding key messages.

Many themes are discussed in a single location, the hospital. Two avatars (a boy, a girl) have been created in order to facilitate identification for young players. Skill games and 16 multiple choice quizzes (used to assess knowledge and deliver pedagogical messages) are spread throughout the game. The test phase brought together 7 children of various ages (5 to 16) and parents members of the KOURIR association and led to some adjustments: enhanced interactivity in the rehabilitation part, shortening and improved ergonomics of the swimming-pool game, and simplification of some of the vocabulary.

Conclusion: To the best of our knowledge, this is the first serious game dedicated to JIA. Other locations (home, school) could complete this version. An English version is already under development. The warm reception it has received from a number of paediatric rheumatologists means we are hopeful many children will be able to benefit from this initiative, at home or in a structured educational context.

Disclosure of Interest
None Declared

P73
JIA patients have a similar quality of life as healthy peers after three years of specialised care
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Pediatric Rheumatology 2017, 15(Suppl 1):P73

Introduction: The treatment of patients with juvenile idiopathic arthritis (JIA) currently aims at achieving an inactive disease state and the best possible quality of life. Whether these treatment goals are achievable in a routine clinical setting can be assessed by data from the Inception Cohort Of Newly-diagnosed patients with JIA (ICON). Herein, patients with recent-onset of JIA have been prospectively observed.

Objectives: To study changes in disease activity state and quality of life of patients during and after the first three years of paediatric rheumatology care.

Methods: Data of 597 patients with JIA and 306 healthy peers who were included in ICON and followed prospectively for at least three years were considered for this analysis. JIA disease activity was assessed by the cJADAS-10 (clinical Juvenile Arthritis Disease Activity Score). Inactive disease was defined as cJADAS-10 ≤ 1. The PedSQL 4.0 (Pediatric Quality of Life Inventory) Generic Core Scales were applied to measure health-related quality of life (HRQoL) in patients and controls and the PedSQL 3.0 Rheumatology module to assess disease-specific QoL.

Results: Patients (69% females) and controls (61% females) had a mean age of 9.8 and 10.1 years at the 3-year-follow-up (3-y-FU). Approximately half of the patients (48%) had oligoarthritis, 27% rheumatoid factor (RF) negative polyarthritis. During the 3rd year of observation, 56% of the patients were on MTX, 26% on biologics and 10% had been treated with systemic glucocorticoids. Patients’ disease activity significantly decreased from a mean cJADAS of 9.5 at first assessment to 2.6 at the 3-y-FU. A total of 271 (45%) patients had an inactive disease, ranging from 47% for persistent oligoarthritis and 56% for RF negative polyarthritis to 79% for systemic JIA. Another 10% reached a minimal active disease at FU. Eighty percent of the patients had received DMARDS up to the 3-y-FU. At the 3-y-FU, 27% did not receive any anti-rheumatic drug. However, only 8% were in remission off drug, another 37% had attained remission on drug.

At inclusion in ICON, patients had a significantly lower HRQoL than controls (mean total scale score 71.3 (SD 18.4) versus 90 (SD 70.5), proxy reports). After three years of rheumatology care, the patients’ HRQoL had clinically meaningful improved and did no longer differ significantly in any health domain from that of controls (total score 86.8 (SD 14.5) versus 89.5 (SD 8.6)). Patients with inactive disease had a higher HRQoL of 92.8 (SD 8.8) at 3-y-FU. Fourteen percent of patients still had high active disease (c-JADAS-10 > 4/8.5 for oligoarthritis/polyarthritis) and a significantly lower mean PedSQL score of 74.0 (SD 17.8). The patients’ disease-specific QoL also significantly improved from ICON inclusion to the 3-y-FU.
**Conclusion:** Current treatment options in paediatric rheumatology care allow achieving an inactive or minimal active disease state and a normalisation of HRQoL in the majority of patients.

**Trial registration identifying number:** ICON is supported by a grant from the Federal Ministry of Research and Education (FKZ: 01ER0812)

**Disclosure of Interest**
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**P74**

**Auxological features in patients with Juvenile Idiopathic Arthritis (JIA) treated with biologic drugs**
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**Presenting author:** Achille Marino

**Pediatric Rheumatology 2017, 15(Suppl 1)**P74

**Introduction:** Juvenile idiopathic arthritis (JIA) is an heterogeneous group of diseases associated with an increase of inflammatory cytokines that may influence child growth.

**Objectives:** To evaluate the auxological features in a cohort of patients with JIA treated with biologic drugs.

**Methods:** This single-center retrospective study evaluated 86 children with JIA (7 systemic, 32 oligoarticular, 31 polyarticular, 6 psoriatic, 10 enthesitis-related arthritis) on long-term (mean 38.5 months; range 6-134) treatment with biologics, and compared them (in a 1:2 ratio) with 172 age and sex-matched healthy controls for height and body mass index (BMI). Patients with endocrinological or other chronic comorbid conditions or on long-term steroid (>3 months) treatment were excluded.

**Results:** All patients received anti-TNFα treatment except 4 patients with systemic JIA; 21 patients received >1 biologic (2 biologics in 15 patients, 3 biologics in 4 patients and 4 biologics in 2 patients); 13 patients received CTLA-4lg; one patient received anti-IL6; anti-IL 1 was used in 5 patients. The target height SDS was not statistically different (0.02 ± 0.66 vs. 0.05 ± 0.79 SDS) between JIA patients and controls. The median height SDS at JIA diagnosis was -0.29 ± 0.66 vs. 0.09 ± 0.84 SDS of controls (p < 0.005) (-0.24 ± 0.66 SDS for oligoarticular (p < 0.05), -0.17 ± 0.79 SDS for polyarticular (p = NS), -1.33 ± 0.49 SDS for systemic (p < 0.001), -0.71 ± 0.13 SDS for psoriatic (p < 0.05), and 0.30 ± 0.11 SDS for enthesitis-related arthritis (p = NS)).

During disease modifying antirheumatic drugs (DMARDs) treatment, before biologic therapy beginning, the height was: in all JIA patients -0.31 ± 0.66 SDS vs. 0.14 ± 0.86 in controls (p < 0.0001); -0.39 ± 0.93 SDS (p > 0.005) for oligoarticular and -0.23 ± 0.79 SDS (p < 0.05) for polyarticular, -1.00 ± 0.31 SDS (p < 0.005) for systemic, -0.65 ± 0.61 SDS for psoriatic (p < 0.05), and 0.10 ± 0.64 SDS for enthesitis-related arthritis (p = NS).

After a follow-up of 6.4 ± 2.3 yrs height SDS in all JIA patients was -0.16 ± 0.89 vs. 0.08 ± 0.81 SDS in controls (p < 0.05). Results were -0.08 ± 0.12, SDS (p = NS) for oligoarticular, -0.13 ± 0.89 SDS (p = NS) for polyarticular, -0.58 ± 0.14 SDS (p < 0.05) for systemic, -0.58 ± 0.33 SDS for psoriatic (p < 0.05), and 0.01 ± 0.60 SDS for enthesitis-related arthritis (p = NS).

The Δheight during biologic treatment was statistically significant considering all JIA patients, oligoarticular, systemic and psoriatic (p < 0.0001) but not polyarticular and enthesitis-related arthritis. Interestingly, JIA patients showed a significative reduction of BMI SDS after biologic treatment (all patients -0.19 ± 0.47 at diagnosis vs. 0.07 ± 0.31 at last follow-up visit, p < 0.005; systemic -0.11 ± 0.31 vs. 0.40 ± 0.49, p < 0.05; polyarticular 0.13 ± 0.30 vs. -0.06 ± 0.32, p < 0.05), as opposed to controls whose BMI increased after a similar follow-up period (0.23 ± 0.34 vs. 0.45 ± 0.51, p < 0.0001).

**Conclusion:** Long-term biologic treatments are associated with a significative increase of height SDS and decrease of BMI SDS in JIA patients.

**Disclosure of Interest**
None Declared

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**P75**

**Response to conventional DMARD treatment in a cohort of children with Juvenile Idiopathic Arthritis in Bogota, Colombia**
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**Presenting author:** Adriana S. Diaz-Maldonado

**Pediatric Rheumatology 2017, 15(Suppl 1)**P75

**Introduction:** Different treatment options are available for juvenile idiopathic arthritis (JIA) increasing with the time. There are multiple factors, such as JIA subtype that impact in the management of medication and the escalation of the therapy (1).

**Objectives:** To describe the treatment with conventional and biological DMARDs and their response on a cohort of children with JIA in 6 health care institutions in Bogotá, Colombia.

**Methods:** Medical records were retrospectively reviewed derived from 568 patients diagnosed with JIA and attended in 6 health care institutions in Bogotá, Colombia since 2010. Demographic and clinical characteristics were collected as well as the current and previously suspended treatments due to no response. The no response was considered when a patient persists with clinical and laboratories criteria after 6 months of treatment.

The description of the current and previous treatments of patients with JIA was performed according to JIA subtype and Disease-modifying antirheumatic drugs (DMARDs). A descriptive analysis was performed of different treatments and those who required to suspend any biological (bDMARDs) or conventional DMARDs (cDMARDs).

**Results:** 471 patients (83%) received immunosuppressive therapy with DMARDs and 97 (17%) with NSAID or prednisolone. Of these 43% (244) have been received bDMARD. Among those treated with DMARD, 51% patients not responding to cDMARDs and need bDMARDs. TNF inhibitors are the most frequent (72,2%) followed by interleukin - 6 receptor antagonist (IL-6 RA) (16,8%), co-stimulatory (4%), IL-1RA (0,4%) and anti CD20 monoclonal antibodies (5,2%).

42% of patients with Polyarticular JIA rheumatoid factor (RF) positive, Polyarticular JIA rheumatoid factor (RF) negative, psoriatic arthritis and systemic arthritis have been treated with any biologics. Additionally, the 50% of patients with enthesis related arthritis (ERA) HLA-27B positive have been treated with biological therapy.

Regarding to previous biological therapy, 73,4% had received one biological (bDMARDs) or two biologics, 2% three biologics and 2% four biologics. All patients mainly belong to polyarthritis and systemic arthritis.

In current treatment, 30,7% of the patients use bDMARDs in monotherapy while the other 69,3% use bDMARDs in combination with one cDMARDs. Conventional DMARDs are used in 97% of patients with JIA Methotrexate is used in 88,9% patients with cDMARDs, followed by sulfasalazine (6,2%) and leflu-nomide(4,9%). Combination of two cDMARDs is used in 2,77% of patients with cDMARDs.

Ninety one patients achieved remission: 77 patients (84,6%) achieved remission off medications, which 28 patients presented oligoarticular JIA (36,3%), 21 polyarticular JIA (27,2%) and 20 systemic arthritis (25,9%). The highest percentages of remission was oligoarthritis. 13 patients (14,3%) are in remission on medications.
Thirty four patients dropout the treatment where administrative barriers were the mainly cause. Children with oligoarticular and polyarticular JIA were the most frequent in drop outing.

**Conclusion:** Conventional DMARDs are the most frequent drugs used in the treatment of JIA represented mainly by methotrexate. In the case of bDMARDs, the anti TNF inhibitors are the principal drugs. The bDMARDs showed a high frequency of use in positive HLA-B27 ERA patients compared to other subtypes. Polyarticular JIA patients presented the most frequent changes of bDMARDs compared to other JIA subtype.

**Disclosure of Interest** None Declared

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**P76**

**Discrimination between Juvenile Idiopathic Arthritis and leukemic arthritis**

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**Objective:** To determine the clinical and laboratory differences between the two entities.

**Methods:** Patients under 16 years old, both genders, initially presenting to the Pediatric Rheumatology unit, General Hospital, National Medical Center La Raza, with diagnosis of probable JIA, without blast cells in peripheral blood, where the final diagnosis was acute lymphoblastic leukemia or juvenile idiopathic arthritis. The clinical and laboratory manifestations were evaluated. Data were expressed as percentages for each analysis area. Chi square and Relative Risk (qualitative variables) and Mann-Whitney and T test for comparison of means between groups. The data were analyzed by software using SPSS version 15.0.

**Results:** We analyzed data on 34 patients, 9 with leukemic arthritis and 25 with JIA. The mean age at diagnosis was significantly lower in the group of leukemic arthritis 7.56 vs 10.76 years (p = 0.045), female gender prevailed in both groups (66% and 76%, respectively, p = 0.586), time of onset of symptoms it was lower in the group of leukemic arthritis. Pain intensity, night sweats, weight loss, fever, lymphadenopathy, hepatosplenomegaly and night pain was higher in the group of leukemic arthritis. Remission of pain with analgesics was significantly lower in patients with leukemic arthritis As for laboratory parameters, rheumatoid factor positivity was higher in the group of AU without showing statistical significance. The hemoglobin level, the leukocyte and neutrophil count was lower in patients with leukemic arthritis, the lymphocyte count was similar in both groups. The platelet count was higher in JIA patients, the VSG was similar in both, the difference between the value of the PCR was not the value of the DHL was higher in the group of leukemic arthritis without showing statistical significance.

**Conclusion:** Leukemia is an important differential diagnosis in children presenting with arthritis.

In this study it was found that the presence of pain disproportionate to the degree of arthritis, fever, night pain, which does not yield to the use of analgesics, weight loss and lymphadenopathy, help to discriminate clinically between leukemic arthritis and juvenile idiopathic arthritis. Useful laboratory parameters are leukopenia, neutropenia, and anemia, the mean platelet count was in low normal value, but lower than in juvenile idiopathic arthritis. The laboratory parameter with most higher relative risk was neutropenia.

**Disclosure of Interest** None Declared

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**P77**

**Rheumatoid factor positive polyarticular JIA with pulmonary langerhans cell histiocytosis- a chance association or a causal relationship?**

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**Presenting author:** Ankur Kumar

**Pediatric Rheumatology 2017, 15(Suppl 1):P77**

**Introduction:** Langerhans cell histiocytosis is a multisystem disease of an unknown etiology characterized by infiltration and accumulation of CD1a+ CD207+ histiocytes in various organ systems. Association of LCH with autoimmune or inflammatory disorders has rarely been reported.

**Objectives:** To describe the case of a young girl with juvenile idiopathic arthritis and pulmonary LCH and to discuss the possible common pathogenetic mechanism for both diseases.

**Methods:** A 10 year old girl was symptomatic for past 1 year. Her illness started with left ankle swelling with pain, redness, restriction of movement and early morning stiffness. Over the next 5 months she had progressive involvement of right ankle joint, small joints of bilateral foot and bilateral wrist joints. On examination, she had swelling in bilateral ankle and wrist joints. There was redness and increased temperature over the surface associated with synovial thickening and restriction of passive range of movement. Her investigations revealed (Table 29) thrombocytosis; elevated ESR and C-reactive protein; positive antinuclear antibody (ANA) and positive rheumatoid factor (RF). Chest X-ray revealed cystic changes in bilateral lung fields which were confirmed on CT chest which showed multiple well defined cysts with variable sizes distributed throughout the parenchyma in bilateral lung fields. An open lung biopsy was performed to confirm the diagnosis. Histopathology examination of the resected lung tissue revealed lymphomononuclear inflammatory infiltrate in the interstitium with collection of large atypical cells around the peribronchial region which were showing oval to round nuclei and moderate to abundant vacuolated cytoplasm, these cells were positive for CD1a. Skeletal x-ray revealed no lytic lesions and bone marrow examination was also normal. There was no evidence of any other organ involvement with LCH.

**Results:** For arthritis, she was initiated on oral naproxen, a short course of oral prednisolone and subcutaneous methotrexate; which led to marked improvement in her joint symptoms. For management of LCH, as she was asymptomatic with predominantly cystic lung changes and no nodular changes on CT chest; hence a possibility of chronic fibrotic disease was considered and a decision not to give any specific therapy for LCH was taken.

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**Table 28 (abstract P76), Initial clinical and laboratory findings in patients with leukemic arthritis vs JIA**

|                          | Leukemic arthritis (n = 9) | JIA (N = 25) | P value | RR (IC 95%) |
|-------------------------|---------------------------|-------------|---------|-------------|
| Pain intensity (0–10)   | 10                        | 6.6 (4–10)  | *0.0001 |             |
| Night pain              | 66%                       | 4%          | *0.0001 | 7.7 (2.5–23) |
| Fever                   | 66%                       | 12%         | *0.001  | 5 (1.7–17.6) |
| Weight loss             | 78%                       | 16%         | *0.001  | 7.3 (1.8–29.6) |
| Lymphadenopathy         | 88%                       | 4%          | *0.0001 | 22 (3.2–153.7) |
| Anemia                  | 56%                       | 8%          | *0.002  | 4.8 (1.7–13.3) |
| Hemoglobin g/dl         | 10.9 (8.3–12.9)           | 13.1 (10.2–16.4) |         |             |
| Leukopenia              | 67%                       | 0%          | *0.0001 | 9.3 (3–27)  |
| Leukocytes mm³          | 3555 (1610–5530)          | 9750 (5190–24630) |         |             |
| Neutropenia             | 88%                       | 0%          | *0.0001 | 25 (3.3–170) |
| Neutrophil mm³          | 473 (0–1820)              | 6456 (2270–22130) |         |             |
| Thrombocytopenia        | 44%                       | 0%          | *0.0001 | 6 (2.6–13)  |
| Platelets x10³/mm³      | 171 (53–296)             | 375 (207–661) |         |             |
Conclusion: Our case highlights a rare and previously unrecognized association of juvenile idiopathic arthritis with pulmonary LCH. It also supports the inflammatory origin of Langerhans cells.

Disclosure of Interest
None Declared

Table 29 (abstract P77). Investigations in the index case

| Investigation                  | Result       |
|-------------------------------|--------------|
| Haemoglobin (g/dL)            | 118          |
| WBC counts (×10³ cells/L)     | 11           |
| Differential counts           | N1/3/L2/E7   |
| Platelet counts (×10³/mL)     | 467          |
| ESR (mm/hr)                   | 40           |
| CRP (mg/dL)                   | 18           |
| AST (IU/L)                    | 49           |
| ALT (IU/L)                    | 30           |
| ALP (IU/L)                    | 190          |
| Anti nuclear antibody         | < positive   |
| Rheumatoid factor (U/mL)      | 181 (normal range: 0-19) |
| At diagnosis                  |              |
| 4 months later                | 199          |

P78
The rate of obesity and overweight among Turkish children with Juvenile Idiopathic Arthritis
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Presenting author: Balahan Makay
Pediatric Rheumatology 2017, 15(Suppl 1):P78

Introduction: The increasing rate of overweight among children and adolescents is a serious public health concern. Children with physical disabilities may have an increased risk for overweight and obesity which may be a risk factor for inflammatory arthritides, as it is in adults with psoriatic and rheumatoid arthritis. However, there are only a few studies about obesity and juvenile idiopathic arthritides (JIA) in the literature.

Objectives: To investigate the rate of being overweight/obese among Turkish children with JIA at the time of diagnosis and to assess whether increased BMI is associated with different subgroups of JIA.

Methods: The hospital charts of 325 patients diagnosed as JIA according to the International League of Association of Rheumatology (ILAR) criteria in a 10-year period in a tertiary hospital were retrospectively reviewed. Demographic and clinical findings and JIA subgroups were given in Table 30. In univariate analyses; male gender, age at diagnosis and ERA subgroup were found to be significantly associated with increased BMI (p = 0.013, p < 0.001, and p < 0.001, respectively). However, multivariate backward stepwise regression analyses revealed that only age at diagnosis was significantly associated with increased BMI.

Results: The mean age at diagnosis was 9.1 ± 4.9 years. There were 182 girls and 143 boys. The median delay time to diagnosis was 5 months (interquartile range 2-12 months). 105 patients (32.3%) had persistent oligoarthritis, 81 (24.9%) had enthesitis-related arthritis (ERA), 59 (18.2%) had RF (+) polyarthritis, 29 (8.9%) had systemic arthritis, 21 (6.5%) had undifferentiated arthritis, 16 (4.9%) had psoriatic arthritis, 9 (2.8%) had extended oligoarthritis and 5 (1.5%) had RF (+) polyarthritis. 244 patients (75%) had healthy weight, 56 (17.3%) had increased weight and 25 (7.7%) were underweight. The 46.4 percent of the patients with increased weight belonged to ERA subgroup. The rates of increased BMI according to JIA subgroups were given in Table 30. In univariate analyses; male gender, age at diagnosis and ERA subgroup were found to be significantly associated with increased BMI (p = 0.013, p < 0.001, and p < 0.001, respectively). However, multivariate backward stepwise regression analyses revealed that only age at diagnosis was significantly associated with increased BMI.

Conclusion: This is the first study to investigate the prevalence of overweight in children and adolescents with JIA at the time of diagnosis without any treatment. We could not find any difference between the overweight and obesity rates of patients with JIA and children from the general population. This study revealed that 17.3% of JIA patients had increased weight. A similar overweight/obese rate of 16-21% for children and adolescents in the general population in Turkey was found previously. In a recent study of Schenck et al, systemic JIA and ERA patients showed significantly higher overweight rates compared to other subgroups. We also found a significantly higher rate of overweight in ERA when compared to other subgroups. However, when it was adjusted for age and gender, increased BMI was not significantly associated with any certain subgroup.

Disclosure of Interest
None Declared

Table 30 (abstract P78). See text for description

| Healthy weight | Increased weight |
|----------------|-----------------|
| Persistent oligoarthritis | 95 (90.5) | 10 (9.5) |
| ERA | 55 (68) | 26 (32) |
| Poliarthritis RF (+) | 46 (78) | 13 (22) |
| Systemic | 28 (90.5) | 1 (9.5) |
| Undifferentiated | 17 (81) | 4 (19) |
| Psoriatic | 16 (100) | 0 (0) |
| Extended oligoarthritis | 8 (89) | 1 (11) |
| Poliarthritis RF (-) | 4 (80) | 1 (20) |

Data were presented as n (% within subgroup)

P79
Patient partnership and disabkids in the Swedish Pediatric Rheumatology Registry
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Pediatric Rheumatology 2017, 15(Suppl 1):P79

Introduction: The healthcare of today is facing higher demands on coproduction of it’s service with partnership between patients and healthcare providers. Good outcomes, most recognize, are more likely if the clinician and patient communicate effectively. Real time patient feedback facilitated by Patient Reported Measures (PRMs) can be used to enable true co-production of care. DISABKIDS is a European project that has developed with focus on health related quality of life (HRQoL). The Swedish Pediatric Rheumatology Registry started in 2009. Today over 2700 patients with JIA are included. The coverage is high. 65% of all Swedish patients with JIA (0-17 years of age) are included and above 90% of all patients with JIA on biological treatment. The registry is a tool for both patients and care providers in promoting valuabel care in many ways. National and local data are provided promoting good and equal care for all patients. DISABKIDS is selected in the registry as instrument for HRQoL.

Objectives: To describe the Swedish experience of the process of coproduction with patients, families and patient representativs. Evaluate and describe the differences in the assessments of DISABKIDS as HRQoL and how to use the data to reveal individual problems and connect to the planning of further care in partnership with the patient.

Methods: During the project, an improvement-coach led group semi-nars and the discussions which constituted a base for prioritized areas. The sessions resulted in two defined improvement areas.

Results:
To develop a visualization platform for patient reported data as well as other biomedical aspects. Further, to describe the needs in care for children and adolescents with pediatric rheumatic diseases.
1. The visualization platform was built on a Feed Forward System (FFS) for patients, giving support for communication in clinical settings. This was based on the patient portal with a personal overview from registry entered data. Then a development of web-applications for patients with data exported to the registry of Patient Reported Measures were constructed. PRMs included pain, global assessment patient, health impairment, ability to be at school/preschool and my personal measure. Children are positive to share their results and can illustrate their HRQOL better to their provider during consultations.

National results from the HRQOL assessment of DISABKIDS and DISABKIDS RA (impact and understanding) per dimension:

Conclusion: Patient partnership and co-production will enhance registry quality and give added value for both patients and care providers in promoting efficient and data based decisions in daily clinical work.

Disclosure of Interest
None Declared

Table 31 (abstract P79). See text for description

| Independence | Innere strength | Social inclusion | Social equality | Physical ability | Physical treatment | Total score | Impact | Understanding |
|--------------|----------------|-----------------|----------------|-----------------|-------------------|-------------|--------|--------------|
| 76%          | 47%            | 67%             | 67%            | 65%             | 64%               | 58%         | 2.7/5  | 2.6/5        |

P80
Diagnostic duration and access to care in JIA: a pilot study in the University Hospital of Saint Etienne (France)

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Pediatric Rheumatology 2017, 15(Suppl 1):P80

Introduction: Many studies in pediatric rheumatology show that there is a therapeutic « window of opportunity » early in the disease course to prevent joint or ocular damage in the context of juvenile idiopathic arthritis (JIA). In Europe, despite facilitated access to health care, children with JIA are referred to pediatric rheumatology centres with significant delay. Little is known about the reason for such a delay while strong efforts have to be made to identify each component of diagnostic delay.

Objectives: To assess in JIA patients the diagnostic pathway and to determine which parameters may influence the delay to diagnosis.

To evaluate the parents’ feelings and adjustments during this period.

Methods: From September 2015 to May 2016, we have carried out, in the university hospital of Saint-Etienne, a prospective explanatory research about new JIA patients. The method was both qualitative and quantitative using parents’ semi-structured individual interviews and medical data about the diagnostic pathway. In addition to clinical data, we have included social data: parents’ profession and education level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level. Data on the care course before the diagnosis have also been collected (date of the first symptoms, date of diagnosis, specialty level.

Results: According to the ILAR classification we have included 19 new JIA cases: 7 oligoarthritis, 6 enthesis-related arthritis (ERA), 3 polyarthritis RF +, 3 systemic arthritis (sJIA). Median time between first symptoms and the date of diagnosis was 29 weeks (2-114). Median time between the first symptom and the first medical consultation was 1 week (0-40) whereas median time between the first medical consultation and the diagnosis was 22 weeks (0-108). The median number of physician met before the consultation with the pediatric rheumatologist was 3 (2-6), among the following health care providers: general pediatrician, general practitioner, emergency doctor, orthopedic surgeon, adult rheumatologist and rehabilitation doctor. In 15 cases the first physician was a general practitioner, in the other 4 cases he was an emergency doctor. Patients with ERA had the longest delay to diagnosis (median 55 weeks (8-106)), whereas children with sJIA had the shortest interval (median 2 weeks (1-3)). The delay to diagnosis did not vary depending on the parents’ socio-economic groups or their education level.

Semi-structured individual interviews have been done with 9 families (13 parents) and pointed out that this waiting time was very stressful for each family with a fear of irreversible process with ankylosis. 8 families used complementary and alternative medicines (7 homeopathy, 2 acupuncture and 2 osteopathy). 8 families needed help in the day-to-day life, 2 mothers stopped their jobs and one got a partial time work.

Conclusion: In spite of the fact that patients have quickly recourse to health care, delay to diagnosis for JIA remains very long: strong efforts have to be made to organize targeted training programs for all practitioners involved in the care pathway of these children and to communicate on the on-going national program on rare diseases in France.

The identification of the various components of the delay to diagnosis may enable practitioners to elaborate care course management guidelines and public governmental bodies to set up adapted program for rare disease.

By the mean of the JIR cohort (juvenile inflammatory rheumatism cohort), we intend to carry out a broader study about delay to diagnosis and access to healthcare and treatment for JIA in France and in Switzerland.

Disclosure of Interest
None Declared

P81
Effectiveness of knee splintage under general anaesthetic post interarticular injection in children suffering from JIA

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Pediatric Rheumatology 2017, 15(Suppl 1):P81

Introduction: Flexion contractures, particularly of the knee, are a recognised complication of JIA. In the UK there is currently no standard protocol practiced for the treatment of such contractures. In an audit of 10 UK centres (5 respondents replied), one other centre splinted in theatre post interarticular steroid injection (IASI)

We report our audit findings of post injection splinting in children with JIA.

Objectives:
- To establish what the current practices are in the UK in relation to standard protocols for splinting the knee post interarticular steroid injection
- To compare the ROM measurements of the knee prior to and post IASI and establish the effectiveness of splinting while under GA
- To develop standards to enhance clinical practice

Methods: We reviewed the outcome of range of movement following splinting of the knee while under general anaesthetic from 2013-2015

The knees were injected with Triamcinolone Hexacetonide 20-40mgms according to size. A cylindrical fibreglass cast with a reinforced back slab was then applied under anaesthetic, with the knee held in the greatest achievable level of extension. The splints were opened, secured, and left in place for 24 hours. They were worn nightly thereafter. Parents were taught a daily physio regime for strengthening and stretching, and physio treatment was provided on a weekly basis for up to 6 months. Knee extension was measured at various time points – actively pre-theatre, passively in splint and actively post treatment.

Results: 9 patients and 10 joints were included in the review. Ages ranged from 1-6 years, with a mean age of 2.5 years. Of the 10 patients, 7 had a diagnosis of Oligo JIA and 3 of Poly-Articular JIA, with a mean disease duration of 2.5 years (ranging from 6 months to 6 years).

All 10 of the knees treated showed significant improvement in active extension post injection / splintage. The mean improvement was 83%, and this improvement was maintained unless the knee flared.

The delay to diagnosis did not vary depending on the parents’ socio-economic groups or their education level.

Semi-structured individual interviews have been done with 9 families (13 parents) and pointed out that this waiting time was very stressful for each family with a fear of irreversible process with ankylosis. 8 families used complementary and alternative medicines (7 homeopathy, 2 acupuncture and 2 osteopathy). 8 families needed help in the day-to-day life, 2 mothers stopped their jobs and one got a partial time work.

Conclusion: In spite of the fact that patients have quickly recourse to health care, delay to diagnosis for JIA remains very long: strong efforts have to be made to organize targeted training programs for all practitioners involved in the care pathway of these children and to communicate on the on-going national program on rare diseases in France.

The identification of the various components of the delay to diagnosis may enable practitioners to elaborate care course management guidelines and public governmental bodies to set up adapted program for rare disease.

By the mean of the JIR cohort (juvenile inflammatory rheumatism cohort), we intend to carry out a broader study about delay to diagnosis and access to healthcare and treatment for JIA in France and in Switzerland.

Disclosure of Interest
None Declared
Flexion was not compromised by splinting. Parents reported good compliance with initial continual use and subsequent nocturnal splintage.

**Conclusion:** We are one of only 2 centres who reportedly use splinting following joint injection under anaesthetic for knees with sustained flexion contractures. Our results concur with those published by GOSH that rapid and sustained correction of contractures is achieved and splintage is well tolerated by children. However we have no comparator group to evaluate the efficacy of such intervention over IASI and physiotherapy alone. Splinting in theatre prolongs the use of anaesthetic by 15-20 minutes. Furthermore the type and dose of steroid used in each centre may be a major confounder.

**Disclosure of Interest**
None Declared

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**Table 32 (abstract P81). Comparing active ROM pre-injection/splintage with best active ROM achieved**

| Jt | active ROM |
|----|------------|
| 1  | 90%        |
| 2  | 80%        |
| 3  | 92%        |
| 4  | 67%        |
| 5  | 90%        |
| 6  | 67%        |
| 7  | 80%        |
| 8  | 100%       |
| 9  | 63%        |
| 10 | 100%       |

The mean is 82.9%

**P82**

**Analysis of disease activity and disease damage in patients with Juvenile Idiopathic Arthritis: single center experience**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P82**

**Introduction:** Juvenile idiopathic arthritis (JIA) is a group of disease defined by the presence of continuous inflammation of joints with unknown etiology. JIA is the most important reason of loss of ability and the most common rheumatic disease in childhood.

**Objectives:** In this study, it was aimed to determine disease activity, articular and extra-articular damage index of patients with JIA who were followed-up in our outpatient clinic.

**Methods:** In this prospective study, patients diagnosed with JIA according to the ILAR classification criteria, who were being followed up in pediatric rheumatology clinic at Gulhane Military Medical Academy between September 2014 and May 2016 were included. Demographics, clinical and laboratory features were recorded on a standardized form in addition to arthritis damage index and disease activity (Physician global assessment (PGA)) were calculated for each patient at each visit.

**Results:** A total of 96 patients (58 girls (60.4%), 38 boys (39.6%)) with a mean age of 9.2 ± 5.13 years were included. The distribution of subtypes of JIA were as follows: persistent oligoarticular JIA with 58.3% (n = 58), enthesitis related arthritis with 19.8% (n = 19), RF (-) polyarticular JIA with 7.3% (n = 7), RF (+) polyarticular JIA with 2.1% (n = 2), extended oligoarticular JIA with 4.2% (n = 4), psoriatic arthritis with 2.1% (n = 2). The most commonly affected joint was knee at the diagnosis (57.3%, n = 55). Uveitis rate was 5.2% (n = 5), all the patients with persistent oligoarticular JIA. PGA of disease activity was found 3.1 ± 1.79 for all patients in the first visit, and 0.73 ± 1.219 at he last visit (p < 0.01).

**Conclusion:** During the follow-up of our patients improved with the various treatment modalities. Activity (PGA) and JADI scores of the patients decreased from the first visit to last visit. Evaluating damage and activity in children with JIA by using outcome scores is helpful to see the response to treatment and contribute our daily clinical practice.

**Disclosure of Interest**
None Declared

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**P83**

**Serum levels of myeloid-related proteins 8 and 14 in oligoarticular and polyarticular onset of Juvenile Idiopathic Arthritis**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P83**

**Introduction:** The myeloid related protein (MRP) complex 8/14 is released from activated monocytes and phagocytes. A recently published data have shown MRP8/14 potential role as predictor of disease activity and response to treatment in rheumatic and other inflammatory diseases.

**Objectives:** To analyze serum levels of MRP8/MRP14, and correlate them with disease activity in oligoarticular and polyarticular onset of juvenile idiopathic arthritis (JIA).

**Methods:** 44 children with oligoarticular and polyarticular onset of JIA and 25 healthy children as controls were enrolled to the study. The JIA patients were categorized into subgroups according to disease activity and duration from diagnosis. Active JIA was defined as presence of at least one joint swelling or limitation of joint motion and pain, or warmth or tenderness. Inactive JIA was defined as no active arthritis, no active uveitis, normal erythrocyte sedimentation rate (ESR) or/and C-reactive protein (CRP), morning stiffness ≤ 15 min.

**Results:** Short-lasting disease was defined as disease duration less than 3 months from diagnosis and long-lasting disease as more than 3 months from diagnosis. The following clinical and laboratory measures were collected: duration from disease diagnosis, number of swollen joints and with limitation of motion, pain (VAS, 0-10), Physician Global Assessment (PGA) (VAS, 0-10), patient's/parent's disease activity (VAS, 0-10), CHAQ, duration of morning stiffness, ESR, CRP, complete blood count (CBC). ELISA was used to measure serum levels of MRP8/MRP14. Receiver operating curve (ROC) analysis was performed.

**Results:** Median serum levels of MRP8/14 in JIA patients, and both in oligo- and poly-JIA were significantly higher compared to healthy controls. Median serum levels of MRP8/14 were significantly higher in active and lower in inactive JIA patients compared to healthy controls.
controls. Median serum levels of MRP8/14 in active JIA patients were significantly higher compared to inactive JIA patients. Median serum levels of MRP8/14 in JIA patients with short-lasting disease and with long-lasting disease were significantly higher compared to healthy controls. Median serum levels of MRP8/14 in JIA patients with short-lasting disease were significantly higher than in JIA patients with long-lasting disease. The receiver operating curve (ROC) analysis for the MRP8/14 showed that when the cutoff level of MRP8/14 was set at 3800 ng/mL, the sensitivity and specificity for the JIA patients were 61.4% and 84.0%, respectively. Median serum MRP8/14 positively correlated with PGA, VAS of patient's/parent's disease activity, VAS pain, CHAQ, morning stiffness and laboratory variables of inflammation such as CRP, ESR, white blood count, and platelet count. Serum MRP8/14 negatively correlated with disease duration.

**Conclusion:** Serum MRP8/14 concentrations represent a potent parameter to monitor inflammation in patients with oligo- and polyarticular onset of JIA. The elevation of serum MRP8/14 in patients with long-lasting disease may indicate persistent stimulation of immune response. Further studies are needed to confirm our pilot study. This work was supported by the Wrocław Medical University (Grant No ST-653).

**Disclosure of Interest**
None Declared

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**P84**

**Juvenile Idiopathic Arthritis in mono-zygotic twin**

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**Presenting author:** Emredong S. Asuka

**Pediatric Rheumatology 2017, 15(Suppl 1):P84**

This abstract is not included here as it has already been published.

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**P85**

**Diagnostic pitfalls in Juvenile Idiopathic Arthritis: beyond inflammation**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P85**

Introduction: Juvenile idiopathic arthritis (JIA) is the most common connective tissue disease in the pediatric population with a prevalence of approximately 1 in 1,000 children younger than 16 in North America, and girls being more frequently affected than boys. JIA predominantly involves the large joints, most frequently the knees, wrists, and ankles. Magnetic resonance imaging (MRI) is a relatively new imaging technique to confirm the internal derangement of the joints on patients affected with JIA. Moreover, MRI has the advantage of displaying both soft tissue and bone and we found to be very accurate in the assessment of disease activity and to depict the more frequent pitfalls encountered on these patients, which can lead to a misdiagnosis.

**Objectives:**
1. To describe the more frequent diagnostic pitfalls on patients with JIA on imaging, especially on MRI.
2. To illustrate the radiographic appearance of JIA patients from head to toe, highlighting the differences between active and chronic disease.
3. To review the MR protocol and imaging techniques to study patients with JIA.

**Methods:** We retrospectively review all available hospital medical records, including Electronic Patient Chart (EPC) and clinical database of our patients with diagnosis of JIA. We also review all MR examinations and radiology reports of these patients. Clinical information collected included age, sex, clinical history, JIA subtype, disease duration, history of uveitis and ANA test result. All of our patients were examined with a 1.5 T Magnet (Signa HD; General Electric Medical Systems, Milwaukee, Wisconsin, USA). MR protocol varied depending on the joint examined, but all of them included contrast administration.

**Results:** We found out active inflammation on MR examination according with clinical examination in a significant number of patients. Nevertheless, there were patients with clinical suspicion of active inflammation and no radiological signs of active disease. These patients had other entities not related to JIA. The following joints were affected with different diseases. Temporomandibular joint: disk subluxation. Cervical spine: rotatory subluxation and Langerhans cell histiocytosis. Wrist: scapholunate dissociation and a ganglion cyst. Lumbar spine: spondylolysis with no spondylolisthesis. Hips: apophysial injuries, enthesis not related to arthropathies and an osteoid osteoma on the acetabulum. Knee: Oschgood-Schlatter disease, discoid meniscus tear, a giant cell tumor of the tendon sheath and a stress fracture. Ankle: tarsal coalition and stress fracture.

**Conclusion:** MRI with contrast is the technique of choice and the most sensitive one for the diagnosis and management of patients with JIA. MR is crucial to show the most frequent pitfalls on patients with JIA, due to the ability of displaying both soft tissue and bone.

**Disclosure of Interest**
None Declared

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**P86**

**Insulin resistance and other cardiovascular disease risk factors in Juvenile Idiopathic Arthritis**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P86**

Introduction: It is postulated that insulin resistance (IR) constitutes the key metabolic platform for the development of cardiovascular disease on adults. The role of chronic inflammation on IR has been shown in other diseases as rheumatoid arthritis. Certain subtypes of juvenile idiopathic arthritis (JIA) represent a state of systemic inflammation; it could lead to IR and other cardiovascular disease risk factors (CDRF). Paediatric studies on the subject are limited.

**Objectives:** To describe the IR prevalence in JIA paediatric patients and to assess the possible involved factors. Furthermore, it analyses the prevalence of classical RFC, as well as metabolic syndrome.

**Methods:** Observational cross-section study in Spanish children from 4 to 16 years old affected by the most frequent subtypes of JIA ( oligoarticular, polyarticular and systemic) according to ILAR classification, monitored by a Pediatric Rheumatology Unit between July 2014 and July 2015. Monoarticular forms and patients with other concomitant treatments or diseases were excluded.

Anthropometric, clinical and treatment data were recorded. Blood test was performed, included every metabolic syndrome parameters and 3 adipokines (leptin, adiponectin and resistin). Blood pressure and validated surveys about Mediterranean diet (KIDMED) and physical activity (PAQ-C/PAQ-A) were made also. IR was defined as a homeostatic model assessment (HOMA-IR) on the 90th percentile of healthy children in a Spanish cohort (Garcia Cuartero et al 2007).

**Results:** 88 patients were enrolled. 70.5% female with an average age of 11 years old. Their epidemiological, clinical and regarding treatment received characteristics are recorded in Table 33.
The IR prevalence was 24.4%. Regarding classical CDRF, 6 (7.5%) were obese, 12 (15%) overweight, 16 (18.3%) arterial hypertension (AH), 14 (16.8%) dyslipidemia, mainly HDL.

About the diet and physical activity surveys, 41.3% followed an optimal Mediterranean diet (KIDMED score ≥8), being the most of the cases improvable (score 4-7). Our patients performed moderate physical activity during free time (PAQ-C/PAQ-A score > 3), however, statistical differences were not significant between JIA subtypes.

A linear multivariate regression analysis was performed. IR was associated (R² = 0.512) with hip circumference (cm) (β = 0.04; p = 0.008), HbA1C% (β = 2.845; p = 0.008) and leptin (ng/ml) (β = 0.036; p = 0.007). Clinical activity duration was the only predictor factor, not having significant correlation with the subtype, received treatment or disease activity during the study.

Conclusion: Our data suggest that the patients with JIA have an IR prevalence similar to the Spanish healthy children (adjusted by age, gender and pubertal stage). The only factor related to the disease which directly influenced the IR grade was the clinical activity overall duration, hence the importance of an early control of the disease to avoid this CDRF in future.

Disclosure of Interest
None Declared

Table 33 (abstract P86). Clinical and treatment characteristics

| N | JIA subtype, n (%) |
|---|-------------------|
|   | Systemic          |
| 10 | (11.4)            |
| Persistent oligoarticular | 43 (48.9) |
| Extended oligoarticular | 14 (15.9) |
| Polyanarticular            | 21 (23.8) |
| Disease duration (years), mean ± SD | 6.3 ± 3.6 |
| Inflammatory activity duration (weeks), median (IR) | 53 (32–109) |
| Synthetic DMARDs treatment, n (%) | 85 (96.5) |
| Biological DMARDs treatment, n (%) | 40 (45.5) |
| Systemic corticosteroids, n (%) | 79 (90.7) |
| Duration corticosteroids (days), median (IR) | 122 (55–178) |
| Median corticosteroids dose (mg/kg/day), median (IR) | 0.34 (0.21–1.15) |

P87
Prospective analysis of the immunogenic response in JIA patients (paediatric and adult) on antiTnf treatment
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Pediatric Rheumatology 2017, 15(Suppl 1):P87

Introduction: There is some evidence that clinical response in patients with chronic arthritis treated with antiTnf agents can be influenced by their immunogenicity. This particular has not been fully studied in patients with juvenile idiopathic arthritis (JIA).

Objectives: To evaluate the immunogenic response against antiTnf agents and its possible impact on treatment efficacy in patients with JIA.

Methods: Seventy-three patients (40 children and 33 adults) fulfilling ILAR criteria of JIA were included. All of them were on treatment with Etanercept (ETN), Adalimumab (ADA) or Infliximab (IFX) +/- Methotrexate (MTX). Demographical and disease characteristics were recorded at baseline. Clinical outcome was prospectively assessed every 6 months (JADAS10/27/71, patient or parents VAS, physician’s VAS, CHAQ/HAQ, ESR, CRP, number of swollen, painful and limited joints). Patients were tested for the presence of both, serum drug levels and antidrug antibodies levels at baseline and every 6 months.

Results: Thirty-one paediatric patients (77.5%) needed combined treatment (antiTNF + MTX) vs 18 patients (54.4%) in the adult group. Furthermore, paediatric patients followed an intensified antiTNF therapy protocol more frequently than adult patients (54% vs 39.3%; p = 0.044). A total of 189 determinations for the assessment of immunogenic response against antiTNF therapy were analysed. No differences were found between children and adults in the prevalence of antidrug antibodies or in the circulating antiTNF levels. No antidrug antibodies were found neither in the 107 determinations of patients following ETN treatment nor in the 6 determinations of patients on IFX. Just 5 (17%) of the 29 patients on ADA treatment produced anti-ADA antibodies at least once during the follow-up period. Regarding the 77 ADA determinations, patients who were on monotherapy with ADA produced antidrug antibodies with a significant higher frequency than patients on combined treatment with MTX (37.5% vs 8.7% of them respectively; p = 0.016). Patients on monotherapy with ADA also exhibit a significant reduced serum ADA levels than patients on ADA + MTX (4.73 µg/mL vs 12.73 µg/mL respectively; p = 0.019). However, no significant differences were observed in the ETN concentration between patients on monotherapy or on ETN + MTX treatment (1.50 µg/mL vs 1.77 µg/mL respectively). No correlation was observed between drug levels or the presence of antidrug antibodies and any of the clinical or biological parameters.

Conclusion: JIA paediatric patients needed intensified and combined with MTX antiTNF therapy more frequently than adults. In spite of this, no differences were found regarding antiTNF levels between children and adults. This fact suggest that children need more therapy to maintain antiTNF concentrations to control the disease activity. No antibodies antiETN were detected in our patients. ADA in monotherapy seems to be more immunogenic than in combined therapy in JIA. We found no correlation between antiTNF levels or the presence of anti-antiTNF antibodies and the clinical outcome of JIA patients.

Disclosure of Interest
None Declared

P88
Fibrous arthropathy associated with morphea: a new cause of diffuse acquired joint contractures in children
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Pediatric Rheumatology 2017, 15(Suppl 1):P88

Introduction: Etiologies for childhood-onset diffuse joint contractures encompass a large group of inherited disorders and acquired diseases, especially a subset of juvenile idiopathic arthritis called “dry polyarthritides”, dermatomyositis and systemic sclerosis.

Objectives: First time description of the cases of patients who presented with diffuse acquired joint contractures preceding distant superficial circumscribed morphea

Methods: Clinical reports

Results: We report 2 boys aged 5 and 8 years who developed acquired symmetric painless joint contractures preceding from 7 to 13 months the development of superficial plaques of morphea.
There was no other clinical involvement, no biological inflammation and no auto-antibodies. Search for urinary mucopolysaccharidosis was negative. In both patients, MRI of the wrist showed no joint effusion, no bone erosion, absent or mild synovial thickening with a slight enhancement after gadolinium infusion. One patient underwent a synovial biopsy which showed a dense fibrosis with a sparse inflammatory infiltrates, similar to the pathological pattern observed in the skin biopsy. With methotrexate and systemic steroids, joint contractures slowly improved in the first patient, remained stable in the second.

**Conclusion:** These two cases suggest that fibrous synovitis should be considered in children with acquired diffuse, symmetric and painless contractures, without elevation of acute phase reactants even in absence of extra-articular manifestations.

Articular MRI with gadolinium and careful cutaneous examination at onset and during the follow-up should provide clues for diagnosing this entity.

**Disclosure of Interest**
None Declared

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**P89**

**Subclinical atherosclerosis in children with Juvenile Idiopathic Arthritis**

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**Introduction:** Juvenile Idiopathic Arthritis (JIA) is associated with early subclinical signs of atherosclerosis. Chronic inflammation per se may be an important driver but other known risk factors, such as dyslipidemia, hypertension. In 2006, the American Heart Association published guidelines endorsed by the American Academy of Pediatrics with a list of diseases in children with an increased risk of cardiovascular diseases (CVD) where JIA was included. Because CV events are rare in the young, surrogate markers of atherosclerosis such as carotid intima-media thickness (IMT) are valuable to detect early subclinical atherosclerosis.

**Objectives:** To assess subclinical atherosclerosis in patients with JIA and to compare it with disease duration, subtype of arthritis and the principal outcome measures of disease activity

**Methods:** Patients with JIA consecutively hospitalised in our center were included in the study. IMT of the common carotid artery was determined by ultrasonography. The serum levels of atherosclerosis-related biomarkers, such as homocysteine, triglicerid, cholesterol and sistemic blood pressure were measured. The Juvenile Arthritis Disease Activity Score (JADAS) was used to measure JIA disease activity.

**Results:** 10 patients with JIA were included in the study, 3 oligoarticular, 2 systemic, 3 polyarticular, 1 psoriatic and one with enthesitis related arthritis. Mean age 5.3 yrs, mean disease duration 2.7 yrs. The median (range) of JADAS71 in the 10 patients was 10 (0-19). The table shows the Spearman correlations of IMT with the principal outcome measures.

**Conclusion:** The preliminary results do not evidenced a clear correlation between IMT and JADAS or other outcome measures. These results need to be confirmed in a larger cohort of patients.

**Disclosure of Interest**
None Declared

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**Table 34 (abstract P89), See text for description**

|                  | IMT     |
|------------------|---------|
| Age              | 0.518340|
| Age at onset     | 0.380243|
| Disease duration | 0.303138|
| Physician’s global assessment | -0.066940|
| Active joint count | 0.013623|
| ESR              | -0.107416|
| Cholesterol level | 0.589938|
| Weight           | 0.562500|
| Blood pressure   | -0.217043|

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**P90**

**Treatment and 1-year outcomes of an inception cohort of Australian children with JIA**

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**Pediatric Rheumatology** 2017, 15(Suppl 1):P90

**Introduction:** Recent studies have provided insight into the short and intermediate term outcomes of cohorts of children with JIA managed with contemporary treatments in North America and Europe. There are no such data for Australasia.

**Objectives:** To describe the treatment and 1-year clinical outcomes of an inception cohort of newly diagnosed Australian children with juvenile idiopathic arthritis (JIA) followed at a tertiary paediatric centre.

**Methods:** Retrospective review of prospectively collected clinical data from a specifically designed electronic rheumatology database on all patients newly diagnosed with JIA at the Royal Children’s Hospital, Melbourne, between October 2010 and October 2014.

**Results:** 134 patients were included. Sixty one percent were female. The mean age at diagnosis was 8.3 years. The distribution of patients by subtype was: Oligoarticular 36%; RF + Polyarticular 2%; RF – Polyarticular 25%; Systemic 7%; Enthesitis Related 9%; Psoriatic 7% and Undifferentiated 13%. Eighty nine percent of patients achieved a zero joint count at least once in the first year of follow-up, however, just 66% had a zero joint count at 1 year. The mean time to a zero joint count was 4.3 months in patients achieving this outcome in the first year of follow-up. The systemic subtype had the shortest time to zero joint count at 2.7 months, psoriatic the longest at 6.2 months. For the 11% of patients who did not achieve a zero joint count in the first year of follow-up, the mean time to this outcome was 17.4 months.

Sixty three percent of patients were commenced on a conventional DMARD (cDMARD), most commonly methotrexate. The mean time to commencement of a cDMARD was 2.2 months. 15% were commenced on a biologic DMARD (bDMARD). The mean time to commencement of a bDMARD was 6.3 months. 55% were treated with oral corticosteroids. 19% were on oral corticosteroids at 1 year. 62% were treated with intra-articular steroids, most commonly the olioarticular subtype, of whom 94% had at least one joint injection. 7% of children developed uveitis, with the highest incidence in the olioarticular subtype, 17%. Of patients with inactive arthritis at 12 months, 41% were on and 25% were off treatment.

**Conclusion:** This study demonstrates that children with JIA in Australia have demographic features and short-term outcomes similar to those described in international cohorts. At twelve months,
two thirds of the cohort had a zero joint count and over a third of these were on no medications. When compared to international cohorts there was some variation in management, with a higher proportion having intra-articular steroid therapy.

Disclosure of Interest
None Declared

P91
Anti-cyclic citrullinated peptide antibodies, antinuclear antibodies, and rheumatoid factor in Mexican patients with Juvenile Idiopathic Arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1):P91

Introduction: Juvenile idiopathic arthritis (JIA) worldwide affects 1 in every 1,000 children. The ILAR classification describes seven categories of JIA. The diagnosis is clinical but there are biochemical markers associated with extra-articular conditions. The rheumatoid factor (RF) is presented in patients with rheumatoid arthritis (RA) but in pediatric patients their absence does not discard JIA, the presence of antinuclear antibodies (ANA) is associated with a high risk of uveitis, and the anti-cyclic citrullinated peptide antibodies (Anti-CCPA) are useful for the early diagnosis of RA and predicts erosive forms of the disease. In children with JIA there are few reports of the presence and useful of such antibodies.

Objectives: To determine the prevalence of each type of JIA and the presence of Anti-CCPA, ANA, and RF in patients of the Pediatric Rheumatology Clinic of a third-level care Mexican Hospital (HRAEB), to compare clinical, immunological and inflammatory characteristics by type of JIA, and to quantify the correlation between assessments.

Methods: A cross-sectional sample of 50 patients with JIA (2-15 years). We estimate the prevalence of each type of JIA. The immunological characteristics (RF, ANA and Anti-CCPA) were assessed and compared by type of JIA (Kruskal Wallis or Chi-square). Spearman correlation coefficients between age and assessments were calculated.

Results: The analysis included 50 patients, 29 (58%) female and 21 (42%) male, aged 10.56 ± 3.39 years old, 34% seropositive polyarthritis, 28% seronegative polyarthritis, 16% systemic arthritis, 14% oligoarthritis, and 8% enthesitis related to arthritis. Association between sex and type of JIA (p = 0.014) was detected. Only one patient had positive ANA. There was significant difference in Anti-CCPA levels by JIA type (p < 0.001); higher levels in patients with seropositive polyarticular JIA. High correlation between RF and Anti-CCPA (r = 0.63, p < 0.001) was detected. It was identified that age correlated with RF (r = 0.40, p = 0.004), and with Anti-CCPA (r = 0.29, p = 0.041).

Conclusion: Our data suggest that there is a positive correlation between RF and Anti-CCPA in patients with JIA. The elevation of Anti-CCPA and RF seem to be an early form of adult RA. Patients who have positive Anti-CCPA and RF have a higher degree of severity of the disease with a poor response to the conventional treatment and early biological therapy is required. The presence of Anti-CCPA and RF in a pediatric patient may be a criterion for making a decision on treatment. Multicenter studies that support these findings are required.

Disclosure of Interest
None Declared

Table 35 (abstract P91), Characteristics of the study population grouped by JIA type

| Variables | Systemic arthritis (n = 9) | Enthesitis related arthritis (n = 4) | Oligoarthritis (n = 7) | Seronegative polyarthritis (n = 14) | Seropositive polyarthritis (n = 17) | p-value |
|-----------|--------------------------|-----------------------------------|-----------------------|-----------------------------------|-----------------------------------|--------|
| Female, n (%) | 2 (25%) | 0 (0%) | 4 (57%) | 10 (71%) | 13 (76%) | p = 0.035a |
| Male, n (%) | 6 (75%) | 4 (100%) | 3 (43%) | 4 (29%) | 4 (24%) | p = 0.014b,c |
| Age (years) | 8.75 (4.66) | 13.70 (1.91) | 7.66 (4.49) | 9.31 (4.21) | 12.9 (5.98) | p = 0.002bc |
| RF (UI/ml) | 8.28 (3.02) | 8.60 (2.00) | 6.47 (3.69) | 6.60 (6.00) | 196.82 (300.65) | p = 0.010bc |
| Anti-CCPA (U/ml) | 9.41 (4.44) | 11.95 (2.045) | 6.29 (2.36) | 9.11 (14.17) | 64.94 (884.9) | p = 0.001bc |
| <20 (negative) n (%) | 6 (75%) | 3 (75%) | 6 (86%) | 11 (79%) | 3 (18%) | p < 0.001bc |
| Z (positive) n (%) | 2 (25%) | 1 (25%) | 1 (14%) | 3 (21%) | 14 (82%) | p < 0.001bc |

Unless otherwise indicated, the values are given as the mean (S.D.).
*Chi-square test
*Kruskal Wallis test
Significant p-values

P92
Development of the “REAJI” skill framework in Juvenile Idiopathic Arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1):P92

Introduction: In the list of healthcare treatments and interventions-JIA > of 2009, HAS recalls that therapeutic education must ensure involvement of the patient and his parents: understandability of the disease, mastering of the technical skills, and adaptation of the lifestyle. The main aim of the ETP of the child, the adolescent, and his relatives is to take care of and prevent complications and to learn the skills involved in care.

Objectives: In this context, we have decided to develop a framework of the necessary skills needed for children suffering from JIA (REAJI) in order to improve their quality of life (knowledge, know-how, know-how-to-be).

Methods: The steering committee is composed of health professionals involved in paediatric rheumatology and patients’ associations. Different stages are scheduled:
1. Literature review
2. Setup of four focus groups (FG) with different facilitation methods: 2 patient FGs (8-12 year olds and adolescents), 1 parent FG, 1 health professional (HP)
3. Analysis by the steering committee et production of a first skills list.
4. Submission of the list to a panel of patients and to the national JIA competence centres for further information.
5. Submission of the completed list to the same participants for ranking (primary and secondary objectives).
6. Validation of the final list by the committee.

Results: The steering committee is made up of two rheumatologists, two paediatricians, and two patients associations, ANDAR (association of patients suffering from RA) and KOURIR (association of parents whose children suffer from JIA). The press review highlights a number of works, mainly on adolescents and outside the French territory. The HP FG is composed of 1 rehabilitation doctor, 1 rheumatologist, 1 paediatrician, 1 psychologist, 1 occupational therapist, 1 nurse and 1 pain nurse. The parent FG is composed of 10 people (6 mothers, 1 aunt, 3 fathers). The 8-12
year olds FG had 3 children and the adolescents FG 4 teenagers. The committee gathers to establish the initial skills list and then proposes to complete it via an online questionnaire to a selection of HPs from the 38 French centres listed by the FRancophone Society of Rheumatology and Internal Paediatric Medicine (SOFRE-MIP) and an equal number of patients/parents. This expanded list is sent back to participants using the same method in order to rank it and lastly to be validated in its final version by the committee.

Conclusion: The collectively-developed REAJ framework is a consensus-based support tool that anyone will be able to call upon: not only individual families but in particular teams working on educational projects. This multidisciplinary participative, national effort is an example of health democracy, the results of which will be made available to all.

Disclosure of Interest
None Declared

P94
Internet program for physical activity and exercise-capacity in children with Juvenile Idiopathic Arthritis: a multicenter randomized controlled trial
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Pediatric Rheumatology 2017, 15(Suppl 1)P94

Introduction: Physical activity (PA), health related quality of life (HRQoL) and exercise capacity are decreased in patients with Juvenile Idiopathic Arthritis (JIA) and do not restore after clinical remission has been obtained. Considering the impact of JIA on daily life, the long-term health risks of the disease, the sedentary habits, and the potentially favorable effect of PA, it is important to promote an active lifestyle in children with JIA. Therefore, we developed a 14-week, cognitive behavioral program, Rheumates@work, which was a combination of internet-based and in-person instructions supplemented with four group sessions to improve PA in children with JIA.

Objectives: The aim of this multicenter study is to determine the effects of Rheumates@work, on PA, exercise-capacity, HRQoL and participation in school and physical education class in children with JIA.

Methods: Patients, aged 8-13 years, were recruited from three pediatric-rheumatologic-centers in the Netherlands to participate in an observer blinded, randomized controlled multicenter trial. Rheumates@work started either in winter or in summer. Primary outcome was PA, assessed with a diary and an accelerometer (Actical) during 7 days, expressed as PA level (PAL), time spent in rest, light PA and moderate-to-vigorous PA (MVPA), before the intervention, after the intervention (intervention and control group), and after 3 and 12 months follow-up (intervention group only). Secondary outcome parameters were: exercise-capacity (assessed with a Bruce treadmill test expressed as endurance time), HRQoL (with the Pediatric Quality of Life Inventory Generic-core-scale) and participation in school and sports (assessed by questionnaires).

Results: Twenty-eight children (median age 9.69, 21 girls) participated in the intervention and 21 as controls (median age 10.2, 12 girls). No significant differences between groups were found. None of the PA outcomes measured with the Actical had changed significantly within groups. A significant median difference of MVPA of 31 minutes (p = .04) has been found in the intervention group. Exercise-capacity improved in the intervention group with a median of 57 seconds (p = .02) and decreased in the control group by 46 seconds (not significant). Participation in school improved significantly in the intervention group. Children that missed at least one day of school due to JIA decreased from 43% to 14% (p = .02). Participation in physical education class increased in the intervention group from 57% to 71% (p = .001). HRQoL improved in the control group (p = .01), not in the intervention group. Improvements on PA and exercise-capacity persisted in the 12 months follow-up period. Improvements in rest (median decrease of 54.17 minutes, p = .01), MVPA (median increase of 12.72 minutes, p = .03 (Actical)) and PAL (increase of 0.07, p = .02 (Actical)) were better for the intervention cohort starting in winter compared to the cohort starting in summer.

Conclusion: Rheumates@work exerted no significant differences between groups. Small positive effects on PA, exercise-capacity, and participation in school and physical education classes in the intervention group were found. Improvements lasted up until

References
1. Norms for grip strength in children aged 4–16 years, Häger-Ross 2002
2. Hand Grip Strength in Juvenile Idiopathic Arthritis as Predictor of Disease Activity and Disability in Clinical Practice, Hammar-Rashed 2015

Disclosure of Interest
None Declared

P93
Measuring grip strength in children and adolescents diagnosed with juvenile idiopathic arthritis (JIA) using the new computerised GrippitTM. Does poor grip strength and pain affect activities in daily life?
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Presenting author: Johanna T. Kembe
Pediatric Rheumatology 2017, 15(Suppl 1)P93

Introduction: Clinical experience is that inflammations and pain in hand/finger joints affects grip strength in children and adolescents diagnosed with JIA which also affects their activities in daily life. Occupational therapist’s at Astrid Lindgren’s Children’s hospital, has measured grip strength with Grippit™ electronic hand grip measure for the last 20 years. Grippit™ has now been computerised. Results from Grippit™ gives objective measurement in Newton presented in maximum, average and sustained grip strength after 10 seconds. The results are compared with norms for typical children age 4-16 years. It is known that grip strength for children and adolescents with JIA is generally lower compared to healthy population. With medical treatment, e.g. joint injections, and hand training programs grip strength and hand function will improve and can affect the performance of daily activities.

Objectives: To evaluate the computerised Grippit™ and analyse data from children and adolescents with JIA and compare relation between grip strength, pain and activities in daily life.

Methods: A clinical pilot study on 12 participants, 10 girls and 2 boys age 10-18 years. Grip strength was measured at 2-3 occasions per individual. Grip strength data was correlated to age and/or length of hand. Pain in the hand during the grip assessment was measured with Visual Analogue Scale (VAS 0-100). Performance in activities in daily life (ADL) was measured with Childhood Health Assessment Questionnaire (CHAQ) for items concerning hand function.

Results: Eight participants had a lower grip strength compare to typical children. Pain in hand/fingers during the assessment was reported to mean 40 at VAS. Eleven participants reported a negative impact on ADL. Even for those (n = 4) with normal grip strength, the majority (n = 3) reported a negative performance on ADL and 2 of those reported pain in hand/fingers during the evaluation.

Conclusion: It is important to measure grip strength for children and adolescents with JIA to get a better understanding how the disease affects activity functioning and in order to conduct adequate interventions. The computerised Grippit™ is as good and useful instrument for measuring grip strength.

References
1. Norms for grip strength in children aged 4–16 years, Häger-Ross 2002
2. Hand Grip Strength in Juvenile Idiopathic Arthritis as Predictor of Disease Activity and Disability in Clinical Practice, Hammar-Rashed 2015

Disclosure of Interest
None Declared
A rare disease in the elbow: Lipoma Arborescens

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Pediatric Rheumatology 2017, 15(Suppl 1):P95

Introduction: Lipoma Arborescens, is a diffuse villous proliferation of the synovium characterized by replacement of subsynovial tissue by mature adipocytes. The etiology is not exactly known. Usual presentation is on the middle-aged men and knee joint especially on the suprapatellar bursa. A few cases in childhood have been reported in the literature.

Objectives: In this report, we describe an unusual case of Lipoma Arborescens in a child who was initially diagnosed as oligo articular juvenile idiopathic arthritis (JIA) in the elbow without any response to JIA treatment.

Methods: Case: A 16-year-old boy was admitted to our hospital with the complaint of the swelling and limitation in the movement of the right elbow for the last 5 months. He had no history for trauma. Physical examination revealed swollen and limited range of motion of the right elbow without redness and warmth. Laboratory tests showed normal acute phase reactants; negative antinuclear antibody (ANA) and Purified Protein Derivative test. MR examination of the right elbow demonstrated joint effusion and synovial hypertrophy consistent with juvenile idiopathic arthritis. He was treated with ibuprofen and then intraarticular steroid injection was performed. Upon no response, methotrexate treatment was initiated. On the third month of the treatment, since no ACR paediatric 30 response was received, etanercept was added to treatment. On the sixth month of follow up another MR was performed based on his ongoing complaints and revealed dense effusion, synovial hypertrophy and contrast involvement consistent with active synovitis and also cystic destructive fields in his elbow.

Results: Having these findings, the patient was referred to the orthopedics department. A synovectomy was performed to the anterior and posterior compartments of elbow joint via anterolateral approach. A sample of biopsy was also taken from the lytic lesion in the distal humerus. Histologically, synovium and perisynovium had increased adipose tissue with occasional lymphoid aggregates. Humor us biopsy revealed an osteoblastic tumor consistent with osteoblastoma having hemorrhagic cystic areas. The first MR was revaluated and T1-weighted sequences revealed frond-like synovial proliferation with high signal density where fat-suppressive sequences revealed partial suppression in this formation. The diagnosis of Lipoama Arborescens was confirmed. The limitation range of motion was resolved in time. He is now on follow up for osteoblastoma.

Conclusion: This case has been assessed worthy to report because it was seen in the elbow joint together with osteoblastoma. This rare disease should be kept in mind where joint is severely swollen with limited range of motion, negative ANA and acute phase reactants and unresponsiveness to the oligo JIA treatment.

Disclosure of Interest: None Declared
index (RI) were assessed in 46 children with JIA (9 boys and 37 girls, mean age 10.3 ± 3.8) and compared with 51 healthy controls (M/F=2/18; mean age 9.1 ± 3.1 years) and 26 were treated with biologic drugs (BD group) (M/F=7/19; mean age 11.2 ± 4.1 years). Renal RI was measured from pulse Doppler waveforms obtained from interlobar and arcuate arteries in three different regions from upper, middle and lower third of the kidney. Also a mean RI was calculated for each child by averaging mean RI of the right and left kidney.

Results: Creatinine, CysC and BUN were significantly higher in patients compared to controls; besides eGFR was lower in JIA subjects with healthy patients (p < 0.001) (tab 36). In details, the right, left and mean RI values were significantly increased in JIA subjects compared to controls (p < 0.001). Of note BD group showed significantly higher values of the mean RI compared to MTX group (tab 36); however disease duration in the BD group tend to be higher than the MTX group.

Conclusion: The results of our study suggest that eGFR is reduced in children with JIA. Moreover RI values are impaired in subjects with JIA compared to healthy controls and are significantly higher in those treated with BD compared to MTX group. IR values are directly related to indexes of inflammation suggesting a direct effect of disease.

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Disclosure of Interest
None Declared

| Table 36 (abstract P97). Renal parameters of subjects with JIA |
|-----------------------------|--------------|--------------|--------------|------------|
| Parameters                  | Controls (n = 51) | JIA (n = 20) | JIA (n = 26) | p-value*†‡ |
| Creatinine (mg/dl)          | 0.42 ± 0.10     | 0.48 ± 0.08  | 0.54 ± 0.15  | <0.001†‡ |
| Cystatin C (mg/l)           | 0.66 ± 0.05     | 0.79 ± 0.06  | 0.77 ± 0.13  | <0.001†‡ |
| BUN (mg/dl)                 | 11.72 ± 1.32    | 12.00 ± 2.59 | 13.61 ± 3.18 | 0.002†‡ |
| Microalbuminuria (mg/dl)    | 1.09 ± 0.00     | 1.29 ± 0.60  | 1.54 ± 1.55  | 0.08     |
| ESR (mm/h)                  | 1.00 ± 0.00     | 9.15 ± 6.64  | 15.00 ± 8.68 | <0.001†‡ |
| CRP (mg/dl)                 | 0.20 ± 0.00     | 0.34 ± 0.31  | 0.78 ± 0.84  | <0.001†‡ |
| eGFR (ml/min/1.73 m²)       | 121.9 ± 19.6    | 98.9 ± 8.3   | 98.1 ± 10.7  | <0.001†‡ |
| RI                          | 0.60 ± 0.03     | 0.63 ± 0.02  | 0.68 ± 0.03  | <0.001†‡ |
| Duration of disease (years) | 3.96 ± 2.72     | 6.59 ± 3.18  | 0.35         |

One Way Anova Mann-Whitney U-Test

* p < 0.05 Controls vs MTX group  † p < 0.05 Controls vs BD group  ‡ p < 0.05 MTX vs BD group


disclosure of interest

P98 Anti-synthetase autoantibody is seen in patients with overlap myositis in the UK cohort of patients with juvenile dermatomyositis

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Pediatric Rheumatology 2017, 15(Suppl 1):P98

Introduction: The anti-synthetase autoantibody is the most common adult onset idiopathic inflammatory myopathies (IIM). The classic clinical features include myositis, interstitial lung disease (ILD), Raynaud’s, mechanic’s hands, Gottron’s lesions, non-erosive inflammatory arthritis and characteristic pyrexia.

In the pediatric population, anti-synthetase autoantibodies are rare and found in less than 5% of patients with Juvenile DM (JDM) but occur in 9% of patients with JPM and 13% of patients with juvenile overlap myositis, with the peak age at disease onset being 14 years. They appear to have a similar disease phenotype to adults and are at risk of developing ILD, which has been found to be the most common cause of death and were consequently associated with an increased risk of mortality.

Objectives: The aim of this study was to define the frequency and investigate the clinical associations of anti-synthetase autoantibodies in a large cohort of patients with JDM, the UK JDM Cohort and Biomarker Study (JDCBS).

Methods: Serum samples and clinical data were obtained from 386 patients with JDM recruited to the UK Juvenile Dermatomyositis Cohort and Biomarker Study. The presence of anti-synthetase autoantibodies was determined by immunoprecipitation. Due to the small number of patients with the autoantibody in the study, descriptions of common features are given.

Results: Anti-synthetase autoantibodies were identified in only 6 (1.5%) patients with JDM within the cohort. All patients were female and all bar one were considered to have JDM overlap. 50% were of black origin, they were all non-anti-Jo1 positive and had ILD. The details of their diagnoses and clinical features are demonstrated in Table 37. Anti-synthetase autoantibodies can be identified in a small proportion of patients with JDM and identify a distinctive clinical sub-group. Screening for anti-synthetase autoantibodies especially non-anti-Jo1 antibodies may identify patients at risk of ILD. This mirrors the adult presentation where ILD is the predominant manifestation of non-anti-Jo1 anti-synthetase autoantibodies.

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Disclosure of Interest
None Declared
Table 37 (abstract P98). Characteristics of those patients with anti-synthetase autoantibodies

| Patient | 1 | 2 | 3 | 4 | 5 | 6 |
|---------|---|---|---|---|---|---|
| Diagnostic label | Overlap with scleroderma and arthritis | Overlap with scleroderma and arthritis | Overlap with arthritis | Initially diagnosed with PM or IIM then myositis and ILE | JDM | Overlap with systemic sclerosis |
| Age at onset (years) | 9.06 | 9.41 | 15.47 | 13.06 | 12.53 | 14.43 |
| Length of follow up (years) | 0.67 | 11.5 | 1.33 | 6.47 | 3.5 | 2.76 |
| Specific antibody | Jo1 | Jo1 | Jo1 | PL12 | PL7 | PL12 |
| Muscle | Severe myositis | Myositis | Myositis | Myositis | Myositis | Destructive necrotizing, myopathy on biopsy |
| Skin | Gottron’s Gottron’s | Gottron’s | Gottron’s | Gottron’s | Gottron’s | Gottron’s |
| | Heliotrope rash | Heliotrope rash | Heliotrope rash | Heliotrope rash | Heliotrope rash | Heliotrope rash |
| | Hand | Hand | Rash on forearms | Rash on forearms | Rash on forearms | Rash on forearms |
| Joint | Polyarthritis with contractures | Nil of note | Nil of note | Nil of note | Nil of note | Nil of note |
| Lung | Normal | Normal | Normal | ILD | ILD | ILD |
| Mortality to date | Alive | Alive | Alive | Alive | Alive | Alive |

P99
Positive detection of Anti-NXP2 autoantibodies correlates with muscle ischemia in juvenile dermatomyositis
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Pediatric Rheumatology 2017, 15(Suppl 1):P99

Introduction: Myositis-specific autoantibodies (MSA) are useful biomarkers to classify juvenile dermatomyositis (JDM). Anti-NXP2 antibodies are present in 20% of JDM and has been reported to be associated with a substantially increase of the risk of calcinosis (1). Histological features of anti-NXP2 positive JDM remain unknown.

Objectives: Myositis-specific autoantibodies (MSA) are useful biomarkers to classify juvenile dermatomyositis (JDM). Anti-NXP2 antibodies are present in 20% of JDM and has been reported to be associated with a substantially increase of the risk of calcinosis (1). Histological features of anti-NXP2 positive JDM remain unknown.

Methods: A retrospective study of clinical, biological and histological findings from 10 anti-NXP2 positive JDM patients diagnosed from 2013 to 2015 in both Pediatric rheumatology and dermatology centers (Necker’s hospital). Systematic auto-antibodies screening (Mi2, MDAS, TIF1-gamma, NXP2, SAE, Ro52, Jo1, PL7, PL12, EJ, SRP, Ku, PM-Sc1, Sc170), and myopathological study of deltoid muscle biopsy (including immunohistochemistry for endothelial cells (CD31/PECAM), regenerating myofibers (CD56/NCAM), inflammatory cells, anti-human major histocompatibility complex (MHC) class I (HLA-ABC), class II (HLA-DR), and C3b-9 were performed. All biopsies were reviewed using the score tool in JDM (2). Comparisons between anti-NXP2-positive patients NXP2 (group 1) and anti-NXP2 negative patients (group 2) for initial clinico-pathological variables were performed using the Fisher’s exact test for qualitative variables and the Wilcoxon and Kruskal-Wallis tests for quantitative variables.

Results: Autoantibodies were detected in 16/25 JDM patients, including anti-NXP2 in 10/25, anti-MDAS in 2/25 and anti-TIF1gamma in 4/25. In group 1, patients had a greater degree of weakness at diagnosis (median MMST score 42 versus 63, p =0.01; median CMAS score 3 versus 38, p =0.001) and more frequent vasculopathy-related features, especially gastro-intestinal involvement, (p =0.007) than in group 2. Conversely, the percentage of patients who developed calcinosis did not differ between the two groups (20% in each group). At diagnosis, CPK values were > 5 000 IU/L in 88% and 43% of patients from group 1 and 2 respectively. All the patients underwent a muscle biopsy. Histological features revealed typical lesions of dermatomyositis in all cases: inflammatory and perimysial myopathy with ischemic lesions, perifascicular atrophy and necrosis, expression of HLA ABC class I, C5b9 capillary and sarcocellular deposits. Group 1 displayed more frequent ischemic punch-out vacuoles associated with microinfarcts (p = 0.022) and capillary dropout (p = 0.01). At 6 months, 80% of the group 1 patients required more than two lines of treatment (including plasmapheresis or immunoadsorption) versus 45% in group 2 (p = 0.01)

Conclusion: Anti-NXP2-positive patients demonstrate a greater disease severity and a more prominent ischemic pattern than other JDM. Histopathological investigation can contribute to the classification of inflammatory myopathies in association with the MSA screening.

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Disclosure of Interest
None Declared

P100
The role of muscle MRI in detecting a flare-up of juvenile dermatomyositis
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Pediatric Rheumatology 2017, 15(Suppl 1):P100

Introduction: The course of juvenile dermatomyositis has improved greatly over the last 70 years with the aggressive use of corticosteroids, immunosuppressive medication, and biologics. Yet it is often difficult to detect disease flares in these children. Skin rashes may be mild signs of the rash and muscle disease can vary widely and are often equivocal. Lab tests including muscle enzymes are often normal. EMG and muscle biopsy are invasive and not reliable. New disease outcome measures are in development. Other tools may be helpful to determine if more treatment is needed.

Objectives: To determine if a muscle MRI is useful in detecting a JDM disease flare-up and affecting physician decision-making.

Methods: Approval was obtained from the IRB of Nationwide Children’s Hospital (NCH). Subjects were identified by US ICD10 and ICD9 codes and only included if they met the modified Bohan and Peter DM criteria and were seen at NCH between 1/2005 and 6/2015. Each child had to be in remission for 6 months prior to flare-up. Data collected included demographics, clinical presentation at onset, clinical features at time of repeat MRI, lab test and imaging results at presentation and flare-up, and treatment before and after flare-up MRI. The MRI was performed on both lower extremities without contrast with the following sequences: Axial T1, axial T2 fat saturation, axial and coronal inversion recovery, and axial diffusion weighted. The physician decision that a child with JDM was in a flare-up was the gold standard, using rash, muscle weakness (CMAS), muscle enzymes, and myositis on MRI. We compared the MRI result with the physician’s decision of relapse or not and evaluated whether there was a concordance or discordance between MRI findings and the subsequent treatment decision. Cohen’s kappa test was used for chance-corrected agreement between physician decision and MRI findings, Bayes’ rule for conditional probability
Introduction: Few studies describe the natural course and long term outcomes of myositis in childhood in large, prospectively-followed patient cohorts, treated in the modern era.

Objectives: Forty-five JDM children were identified with 32 females; median age at diagnosis was 5.8 years. Eighty percent had weakness at diagnosis, 100% typical rash, 73% typical nail fold capillary changes. At diagnosis, muscle enzymes were compatible with JDM generally (CK 52%, LDH 62%, aldolase 72%, AST 54% abnormal), EMG abnormal in 3/8, muscle biopsy typical of JDM in 10/11, and MRI abnormal for myositis in 31/40. Thirteen of the 45 patients had a repeat MRI for possible flare-up with differing indications: general proximal weakness in 4, hip, thigh or calf pain in 3, rash in 3, worsening nail fold changes in 1, elevated enzymes in 1, and to confirm remission in 1. Three of 13 repeat MRIs were abnormal, demonstrating myositis. There was moderate agreement between the flare-up MRI findings and the physician’s treatment decision (kappa = 0.59); in each abnormal MRI case the physician decided to increase treatment (Bayes rule-100% agreement). When the MRI was negative for myositis in 10 patients, 7/10 physicians chose to taper medication (70% agreement). There was no agreement between MRI findings and muscle enzymes results.

Conclusion: Using a physician’s overall decision as the gold standard, our study does suggest that performance of an MRI at a time of a JDM flare-up is useful, yielding a sensitivity of 50% for flare-up and a specificity of 100%. Using Bayes rule, when an MRI shows myositis, physicians tend to treat 100% of the time and when an MRI shows no myositis, physicians tapered medications 70% of the time.

Disclosure of Interest
None Declared

Table 38 (abstract P101). See text for description

| Category                  | Patient reported outcome | n (%) answering yes |
|---------------------------|--------------------------|---------------------|
| Specific disease features |                          |                     |
| Muscle weakness           | 28 (32.5)                |                     |
| Arthritis                 | 28 (32.5)                |                     |
| Skin rash                 | 21 (25.3)                |                     |
| Calcinosis                | 18 (21.9)                |                     |
| Education and Employment status |                     |                     |
| At school                 | 17 (20.5)                |                     |
| Enrolled in higher education | 15 (18.1)               |                     |
| Employed                  | 38 (45.8)                |                     |
| Unemployed                | 14 (16.9)                |                     |

P102
Diagnosis of Juvenile Dermatomyositis in Germany – a survey among paediatric rheumatologists and child neurologists
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Pediatric Rheumatology 2017, 15(Suppl 1):P102

Introduction: Juvenile Dermatomyositis (JDM) is an extremely rare pediatric rheumatic disease. The early detection of typical clinical findings and adequate use of diagnostic testing is important in order to allow early clinical diagnosis and treatment. Diagnostic criteria for DM were established in 1975 (Bohan/Peter-criteria) but additional diagnostic modalities are available nowadays.

Objectives: To determine the current use of diagnostic testing for JDM in Germany.

Methods: We designed an online survey addressing all members of the Society for Paediatric Rheumatology (GKJR) in Germany and child neurologists with a special interest in JDM in February/March of 2016. The survey consisted of 6 case scenarios (32 questions) addressing diagnostic testing, diagnostic criteria and the treatment of JDM (covered elsewhere). Answers were given on a multiple-choice basis or based on a 5-point Likert scale. Descriptive statistics were used to analyze the findings.
Results: The survey was completed by 59 members of the GKJR and 7 child neurologists experienced in the diagnosis and treatment of JDM. The most important characteristics for the diagnosis of JDM in clinical practice based on a 5-point Likert scale (1 = essential, 5 = not important at all) were considered to be: typical skin findings (1.30 weighted average), proximal muscle weakness (1.51), typical MRI findings (1.61), elevated muscle enzymes (1.67), typical nailfold capillary changes (2.06), the presence of calcinosis (2.10), typical findings on muscle biopsy (2.18), dysphagia/dysphonia (2.38) and constitutional symptoms (2.47). The following diagnostic tests are used by more than 80% of respondents in all patients with probable JDM (in order of descending likelihood): CK, GOT, GPT, erythrocyte sedimentation rate, LDH, ANA, complete blood count, CRP, BUN/creatinine, EKG, pulmonary function testing, echocardiography, muscle MRI and immunoglobulin levels. Manual muscle testing and muscle biopsy are more commonly used by child neurologists whereas the childhood myositis assessment scale (CMAS), serologic testing (immunoglobulin levels, antibodies to infectious agents, anti-ENA antibodies and anti-SJOL antibodies) and abdominal ultrasound more commonly by pediatric rheumatologists.

Conclusion: There is consensus that the diagnosis of JDM is strongly supported by typical MRI findings, in addition to classic clinical findings, and typical MRI findings are therefore accepted as a Bohan/Peter-equivalent diagnostic criterion in Germany. There is consensus on additional diagnostic testing in patients with probable JDM with some differences between paediatric rheumatologists and neurologist.

Disclosure of Interest None Declared

P101

Treatment with high-dose subcutaneous immune globulins facilitated by recombinant human hyaluronidase in patients with juvenile dermatomyositis who are intolerant to intravenous immune globulins

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Presenting author: Claas Hinze

Pediatric Rheumatology 2017, 15(Suppl 1):P101

Introduction: High-dose intravenous immune globulins (IVIg) are a promising strategy in refractory juvenile dermatomyositis (JDM) but IVIg administration may be cumbersome and is often poorly tolerated. Combining recombinant human hyaluronidase to facilitate the administration of high doses of subcutaneous IgG (fSCIg) may be an alternative to IVIg.

Objectives: To describe the first experiences using high-dose fSCIg in patients with refractory, severe JDM intolerant to IVIg.

Methods: In this retrospective case series, five patients with steroid-refractory severe JDM were treated with fSCIg 2 g/kg/month due to IVIg adverse effects (including severe headaches, nausea, vomiting following IVIg in all five patients, and difficult venous access in one patient). Peak serum IgG levels, muscle enzymes, the childhood myositis assessment scale (CMAS) and adverse effects were observed for at least 6 months.

Results: Patients received fSCIg (2 g/kg/month [n = 4] or 1.7 g/kg/month [n = 1]). The first fSCIg schedule we used (fSCIg 1 g/kg every 14 days, i.e. days 1 and 15 of every 28-day cycle) resulted in median IgG peak levels of 1901 mg/dl (1606–2719 mg/dl), compared to median IgG peak and trough levels while previously receiving IVIg of 2758 mg/dl (2429–3560 mg/dl) and 1419 mg/dl (1325–1922 mg/dl). On this regimen, two patients maintained clinically inactive disease, one patient had mildly increased disease activity, one patient had partial improvement and one patient had persistent minimal disease activity. An alternative, “compressed” fSCIg schedule was then used in the three patients with persistent disease activity (fSCIg 1 g/kg five days apart every month, i.e. days 1 and 6 of every 28-day cycle) resulting in IgG peak levels of 2757 mg/dl, 2846 mg/dl and 2300 mg/dl (previously 1606, 1901 and 1774 mg/dl on the biweekly regimen), resulting in clinically inactive disease in two of the three patients. The administration of fSCIg was well tolerated and there were no serious adverse effects.

Conclusion: We report for the first time that in patients with JDM fSCIg allows dosing similar to IVIg. With a “compressed” fSCIg treatment schedule consisting of two applications five days apart, high peak serum IgG levels are achieved, possibly important for treatment success.

Disclosure of Interest None Declared
Comparison of the PRINTO 2010 and PRINTO/IMACS 2016 improvement criteria in the PRINTO trial in new onset juvenile dermatomyositis

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Presenting author: Claudio Lavarello

Introduction: Juvenile dermatomyositis (JDM) is a systemic autoimmune disease characterized by chronic skeletal muscle inflammation with weakness and skin involvement. PRINTO and IMACS developed criteria to evaluate response in clinical trials which have been recently updated and validated.

Objectives: To compare the PRINTO 2010 (JDM 2010 response criteria) with PRINTO/IMACS 2016 response criteria (JDM 2016 criteria) for improvement in the PRINTO new onset juvenile dermatomyositis (JDM) trial.

Methods: New onset JDM children were randomized to receive either prednisone (PDN) alone or in combination with methotrexate (MTX) or cyclosporine A (CSA). Patients were evaluated at months 6, 12, 18 and 24 with different levels of JDM 2010 criteria (improvement of 20-50-70 or 90% in at least three of the 6 core set measures and worsening <30% in no more than one). The same patients were reanalyzed with the continuous JDM 2016 criteria which evaluate absolute percent change into three categories (total score 0-100): minimal (≥30), moderate (<45), major (≥70) improvement. Clinical trials data were analyzed according to the intention-to-treat (ITT) principle with patients discontinuing (lack of efficacy, safety, etc.) considered as non-responders from that point onward.

Results: 139 children were enrolled in the trial: 47 on PDN, 46 on PDN + CSA and 46 on PDN + MTX. The two sets of criteria showed a similar capacity to recognize different levels of improvement of the disease at 6, 12, 18 and 24 months, after treatment with PDN, PDN + CSA or PDN + MTX as shown by the overlapping 95% CI intervals at various time points (Table). In particular, at month 6, 51% patients on PDN versus 72% on PDN + CSA or PDN + MTX achieved at least JDM 2010 20% improvement (p = 0.023) as compared to 55% and 74% with the JDM 2016 minimal improvement (p = 0.027). At month 24, 38% patients on PDN versus 60% on PDN + CSA or PDN + MTX achieved at least JDM 2010 20% improvement (p = 0.016) as opposed to 38% and 61% with JDM 2016 minimal improvement (p = 0.012). Kappa agreement between the 2 criteria with ITT approach were at least in the moderate range (0.61-0.8) as follows: 0.75, 0.75, 0.77, 0.79 at 6-12-18 and 24 months, respectively. The JDM 2016 criteria, similar to the 2010 criteria, confirmed the superior efficacy of combined treatment with PDN + CSA or PDN + MTX versus treatment with PDN alone.

Conclusion: Both response criteria have shown a similar discrimination in evaluating different levels of improvement in new onset JDM patients treated with 3 alternative treatment strategies.

Table 39 (abstract P105). 2010 and 2016 improvement criteria after 6, 18 and 24 months of treatment with PDN or PDN + MTX or PDN + CSA (only few examples are reported)

| 2010 or 2016 criteria of improvement | PDN | PDN + CSA | PDN + MTX | P values |
|-------------------------------------|-----|----------|----------|---------|
| N (%) [95%CI]                       | N (%) [95%CI] | N (%) [95%CI] |
| Month 6                             |     |          |          |         |
| 2010 ≥20%                           | 24 (51%) [36–66] | 30 (65%) [50–70] | 34 (71%) [59–80] | 0.023   |
| 2016 at least minimal               | 25 (55%) [40–70] | 34 (74%) [59–80] | 0.007   |
| Month 18                            |     |          |          |         |
| 2010 ≥50%                           | 21 (45%) [30–60] | 30 (65%) [50–70] | 30 (65%) [50–70] | 0.020   |
| 2016 at least moderate              |     |          |          |         |
| Month 24                            |     |          |          |         |
| 2010 ≥70%                           | 18 (38%) [23–53] | 26 (56%) [41–71] | 26 (56%) [41–71] | 0.042   |
| 2016 at least major                 |     |          |          |         |

P106

Inflammatory milieu of muscle biopsies and clinical features in juvenile dermatomyositis

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Pediatric Rheumatology 2017, 15(Suppl 1):P106

This abstract is not included here as it has already been published.
**P107**

Evidence based criteria for corticosteroid tapering/discontinuation. An analysis of the PRINTO trial in new onset juvenile dermatomyositis

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**Pediatric Rheumatology** 2017, 15(Suppl 1):p107

**Introduction:** Corticosteroids in juvenile dermatomyositis (JDM) alone or in association with other immunosuppressive drugs, namely methotrexate (MTX) and cyclosporine (CSA), represent the first-line treatment option for new onset JDM. No clear evidence based guidelines are currently available to standardize the tapering and discontinuation of corticosteroids in JDM.

**Objectives:** To provide an evidence-based approach for corticosteroid tapering/discontinuation through the analysis of the patients in the PRINTO new onset JDM trial.

**Methods:** New onset JDM children were randomized to receive either prednisone (PDN) alone or in combination with MTX or CSA. All children were given initially three daily pulses of intravenous methylprednisolone (30 mg/kg/pulse), and then PDN 2 mg/kg/day. After 1 month PDN was tapered to 1 mg/kg/day, from month 2 to 6 was tapered to 0.2, at month 12 to 0.1, and then discontinued at month 24. Major therapeutic changes were defined as the addition or major increase in the dose of MTX/CSA/other drugs or any other reasons for which the patient was dropped from the trial (adverse events, lost to follow-up, etc). Patients who followed the steroid tapering protocol and discontinued PDN at month 24 with no major therapeutic change (group 1) represented the reference standard for the best clinical outcome. Group 1 was compared with those following the steroid protocol, but with other major therapeutic changes (group 2), and with the group who deviated from the steroid protocol with/without major therapeutic changes (group 3). JDM core set measures (CSM) were compared in the 3 groups at 6-12-18 and 24 months (Table).

**Results:** 139 children were enrolled in the trial: 47 on PDN, 46 on PDN + CSA and 46 on PDN + MTX. We identified 91 (41%) patients for group 1, 24 (17%) for group 2 and 58 (42%) for group 3. At baseline all 3 groups had a high level of disease activity with no differences in the CSM. In group 1 (PND off no failure) there were 12 (21%) patients randomized to PDN alone, 21 (37%) to PDN + CSA and 24 (42%) to PDN + MTX. When we compared the three groups, significant differences were found in all CSM at each time point of analysis (p < 0.0001). In particular Group 1, when compared to Group 2 and 3, had the lowest level of disease activity at all time points; the decrease of disease activity was primarily within the first 6 months of treatment. Group 2 and 3 were overlapping in the levels of disease activity reached at all the time points and were globally higher when compared to group 1.

**Conclusion:** The PRINTO protocol from the trial in new onset JDM might constitute the reference evidence-based approach for corticosteroid tapering for possible use in current clinical practice.

**Trial registration identifying number:** ClinicalTrials.gov identifier: NCT00323960

**EudraCT number:** 2005-003956-37

**Disclosure of Interest**

G. Giancane: None Declared, C. Lavarello: None Declared, A. Pistorio: None Declared, F. Zulian: None Declared, B. Magnusson: None Declared, T. Avinc: None Declared, F. Corona: None Declared, V. Gerloni: None Declared, S. Pastore: None Declared, R. Marini: None Declared, S. Martino: None Declared, A. Pagnier: None Declared, M. Rodiere: None Declared, C. Soler: None Declared, V. Stanevichka: None Declared, R. Ten Cate: None Declared, Y. Uziel: None Declared, J. Vojinovic: None Declared, A. Ravelli: None Declared, A. Martini Grant / Research Support from: The G. Gaslini Hospital, which is the public Hospital where I work as full time public employee, has received contributions from the following industries for the coordination activity of the PRINTO network: BMS, GlaxoSmithKline (GSK), Hoffman-La Roche, Novartis, Pfizer, Sanofi Aventis, Schwarz Bio- sciences, Abbott, Francesco Angelini S.P.A., Sobi, Merck Serono. This money has been reinvested for the research activities of the hospital in fully independent manners besides any commitment with third parties.

**Speaker Bureau of:** I received speaker's bureaus and consulting fees from the following pharmaceutical companies: AbbVie, Boehringer, Celgene, CrescendoBio, Janssen, Medimmune, Novartis, Novo Nordisk, Pfizer, Sanofi Aventis, Vertex, Servier, N. Ruperto Grant / Research Support from: The G. Gaslini Hospital, which is the public Hospital where I work as full time public employee, has received contributions from the following industries for the coordination activity of the PRINTO network: BMS, GlaxoSmithKline (GSK), Hoffman-La Roche, Novartis, Pfizer, Sanofi Aventis, Schwarz Bio- sciences, Abbott, Francesco Angelini S.P.A., Sobi, Merck Serono. This money has been reinvested for the research activities of the hospital in fully independent manners besides any commitment with third parties.

**P108**

Juvenile dermatomyositis, predictors of disease progression in Mexican children

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**Pediatric Rheumatology** 2017, 15(Suppl 1):P108

**Introduction:** Juvenile dermatomyositis is the most common inflammatory myopathy in children.

**Objectives:** To describe JDM clinical characteristics and course of the disease in a cohort of 40 Mexican patients diagnosed with juvenile dermatomyositis.

**Methods:** A transversal comparative study in a cohort of 40 patients with juvenile dermatomyositis over a period of 24 months. Descriptive data, parametric tests, and Kaplan - Meier graphs were used to analyse disease progression.

**Results:** The mean age at diagnosis was 8.1 years (SD = 3.7), most of them were female (57%), all patients from Hispanic ethnicity. 14 (35%) presented calcinosis at the time of JDM diagnosis, and up to 45% develop calcinosis at 6 months of disease onset. 5% found with an amiotic patron. The median of the muscular enzymes values at 45% develop calcinosis at 6 months of disease onset.

**Table 40 (abstract P107).** Core set measures values at different time points in the 3 groups of patients (only key results are reported).

| Month 6 | MD global | 0.5 (0–2) | 2.6 (0.8–5) | 3 (0.6–6) | <0.0001 |
|---------|-----------|-----------|-------------|-----------|---------|
| Parent  | global | 1 (0–1.8) | 3.1 (0.5–5) | 2.5 (0.8–5.8) | 0.0002 |
| CHAQ    | 0.1 (0–0.5) | 0.1 (0–0.9) | 0.5 (0–1.9) | 0.0007 |
| DAS     | 3 (0–5) | 6.5 (1–11.5) | 8 (4–12) | <0.0001 |
| CMA5    | 47 (42–51) | 45 (35–48.5) | 35 (21.5–47) | <0.0001 |
| CHQ-PH5 | 51.8 (46.2–54.9) | 47.5 (20.3–51.4) | 36.3 (16.6–50.6) | <0.0001 |
| MMT     | 77 (70–80) | 70 (59–78.5) | 64 (51–75) | <0.0001 |
| Month 12 | MD global | 0 (0–1) | 2.6 (1–5.5) | 3 (0.6–6) | <0.0001 |
| Month 18 | DAS     | 0 (0–3) | 6.2 (2.5–11.5) | 9 (2–12) | <0.0001 |
| Month 24 | CMA5    | 50 (48–52) | 42 (36–50) | 36.5 (21.5–48) | <0.0001 |
the diagnosis was 1,005 mg/dL (15 - 29,130) for creatinine phosphokinase, 374 mg/dL (22.3-260) lactiac dehydrogenase, aspartate aminotransferase 80 UI/L (17-1258). We also found initial dysphagia or dysphonia in 32.5%, pulmonary affection in 17.5%, and cardiac affection in 5%. Barium swallow examination was found altered in 25% at diagnosis. In terms of medication 24 patients received intravenous methylprednisolone initially, the median rate of pulse steroid usage was 6 ± 3.1, and 6 patients needs Biological therapy most of them anti-CD20. No relation between calcinosis at diagnosis and weakness of evolution before diagnosis in Kapplnan Meier was found.

Conclusion: We found that Mexican pediatric population with JDM does not present the same course related to the presence of calcinosis, not according studies of other population.

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Disclosure of Interest: None Declared

P109
Corticosteroid-associated pneumatosis intestinalis in juvenile dermatomyositis: a case report
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Presenting author: Bita Arabshahi

Pediatric Rheumatology 2017, 15(Suppl 1) P109

Introduction: Reports of pneumatosis intestinalis (PI) in Juvenile Dermatomyositis (JDM) are limited and often attribute the process to intestinal ischemia secondary to vasculitis. In patients with asymptomatic PI, however, the role for intestinal ischemia is not clear.

Objectives: We aimed to examine other potential causes of persistent PI in JDM, notwithstanding clinical resolution of vasculitis. Our goal was to examine the effect of immunosuppressive medications, namely corticosteroids, in pathogenesis of PI.

Methods: Hereby we report the case of a three year old Asian girl with anti-MDA-5 positive JDM and severe cutaneous vasculitis, who demonstrated persistent asymptomatic PI despite resolution of cutaneous lesions on cyclophosphamide and corticosteroids.

Results: Abdominal films taken during and after her six month course of cyclophosphamide did not reveal any improvement in PI. Patient showed gradual resolution of PI a year after completion of cyclophosphamide therapy, upon weaning her corticosteroids. Although the presence of anti MDA-5 antibodies is correlated with a higher risk of vasculitis, in our patient, there was no observed correlation between the activity of cutaneous vasculitis and PI.

Conclusion: This observation suggests that PI in JDM may be multifactorial, and the role of corticosteroids should be considered in the differential diagnosis of PI in this population.

Disclosure of Interest: None Declared

P110
Demographic and clinical characteristics of children with juvenile dermatomyositis in Cape Town
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Presenting author: Christiaan Scott

Pediatric Rheumatology 2017, 15(Suppl 1) P110

Introduction: Juvenile dermatomyositis (JDM) is a rare idiopathic inflammatory myopathy of childhood with an incidence of 1.2-3.2 per million. The aetiology of JDM is uncertain but may result from immune dysregulation triggered by environmental factors in genetically susceptible children. The demographic and clinical characteristics of JDM may thus differ by race and geographic regions. Few studies have described the characteristics of JDM patients from Africa. There is need for further studies for better understanding of the epidemiology, clinical characteristics and outcome of patients with JDM from the continent.

Objectives: To determine clinical characteristics and outcomes of patients with probable JDM seen between 2004-2013 in Cape Town, South Africa

Methods: We conducted a retrospective observational study to determine clinical characteristics and outcomes of patients satisfying the Bohan and Peter criteria for probable JDM seen between 2004-2013 in Red Cross, Groote Schuur and Tygerberg hospitals in Cape Town. Data was analyzed using R version 3.1.0 (2014-04-10).

Results: Twenty five cases were identified: 16 female and 9 male. Thirteen (52%) of the cases were of indigenous African, eleven (44%) mixed and one (4%) European ancestry. The median ages at disease onset and diagnosis were 6.75 (range 2.0-9.7) and 7.9 (range 3.4-9.75) years respectively. Muscle weakness and characteristic cutaneous manifestations occurred in all the 25 patients while 24 had elevated muscle enzymes. All the patients received corticosteroids, seventeen (73.9%) received methotrexate and four received rituximab. Eleven patients had calcinosis during the disease course [median follow up period of 50 (range 0.5-159) months]. The mortality was 2/25 (8%) while only 40% of the patients had clinically inactive disease by PRINTO criteria. There was no difference in racial distribution (p-value = 0.1), age at disease onset (p-value = 0.87) and disease duration prior to treatment initiation (p-value = 0.75) between patients who had clinically active and inactive disease. The demographic characteristics of children with JDM were similar to that from most other regions of the world with female predominance and similar age at onset. The median delay in diagnosis (4 months) was not longer than that reported in most other studies. However, some children had prolonged delay of up to 7 years due to misdiagnosis that denied them appropriate treatment in a timely manner. Majority (60%) of the patients also remained with clinically active disease, which put them at risk of further disease complications including calcinosis. Even though the mortality rate was low (8%) this was still more than double that reported in most recent large studies especially from the resource rich countries.

Conclusion: Long term follow up of JDM patients is advisable since majority of patients seem to have clinically active disease many years after disease onset despite treatment. Formulation and use of appropriate treatment guidelines and protocols may aid in the early diagnosis and appropriate management for optimum outcomes.

Disclosure of Interest: None Declared

P111
Pulmonary involvement in children with rheumatic diseases
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Pediatric Rheumatology 2017, 15(Suppl 1) P111

Introduction: The data on prevalence and type of pulmonary involvement in children with rheumatic diseases is scarce. Lung injury may be caused by the disease itself or by the biologic/nonbiologic disease modifying anti-rheumatic drugs (DMARDS).

Objectives: Our aim was to assess pulmonary functional status in children with connective tissue diseases (CTDs) and second, to assess the effect of DMARDS (biologic or nonbiologic) in children with rheumatic diseases.

Disclosure of Interest: None Declared
Methods: Children with a diagnosis of CTD or receiving methotrexate or biologic drugs for more than two years who were followed-up at Hacetette University Department of Pediatric Rheumatology and referred to the Department of Pediatric Pulmonology were included into this study. Medical history, physical examination, chest radiography (high resolution computerized tomography [HRCT] in selected cases), and pulmonary function tests such as spirometry, whole-body plethysmography and carbon monoxide (CO) diffusion test were evaluated in all patients.

Results: There were 33 children with CTDs (21 with juvenile dermatomyositis [JDM], 2 polymyositis [PM], 5 systemic lupus erythematosus, 4 systemic sclerosis, 1 mixed CTD). Female-to-male ratio was 2/1 and the median (min-max) age at diagnosis was 96 (11-186) months. CO diffusion capacity was low (<80%) in two patients (one with PM and one with JDM) and spirometry showed restrictive changes in these patients. HRCT was normal in the patient with PM. The patient with JDM had features suggesting interstitial lung disease (ILD) in HRCT and she was positive for anti-MDA5 autoantibody.

Conclusion: Pulmonary disease should be assessed carefully in children with JDM. Both the disease itself and anakinra treatment are useful for the diagnosis. Anti-transcriptional intermediary factor 1(TIF1) antibody has significant roles in oncogenesis. However, anti-TIF1 antibodies cover approximately 80% of the patients, and thus are expected that the differences between all regions of the world will become apparent.

Disclosure of Interest
None Declared

P112
The clinical features of the juvenile dermatomyositis with anti-tif1 antibodies in Japan: report of three cases and review of the literature
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Pediatric Rheumatology 2017, 15(Suppl 1):P112

Introduction: Recently, new autoantibodies specific for dermatomyositis (DM) have been identified. These disease-specific autoantibodies cover approximately 80% of the patients, and thus are useful for the diagnosis. Anti-transcriptional intermediary factor 1(TIF1) antibody has significant roles in oncogenesis. However, the usefulness of anti-TIF1 antibodies in juvenile dermatomyositis (JDM) is not clear.

Objectives: To evaluate the usefulness of Anti-transcriptional intermediary factor 1(TIF1) antibody in Japanese patients with juvenile dermatomyositis (JDM).

Methods: We experienced three cases of the JDM with anti-TIF1 antibodies. In previous reports in Japan, 7 cases have been reported so far (Nagoya: 3 cases, Kanazawa: 4 cases). We analyze the clinical characteristics of these 10 patients in Japan categorized as JDM with anti-TIF1 antibodies.

Results: Three patients with active JDM with anti-TIF1 antibodies were observed in Japan. There was no trend in the age of onset. No patients reported malignancy. Incidence of JDM with anti-TIF1 antibodies in Japan was 29% of whole JDM.

Conclusion: JDM with anti-TIF1 antibodies exhibited different clinical features. To establish the clinical significance of myositis-specific antibodies, the accumulation of cases is indispensable. In the near future, not only the tendency of each region, it is expected that the differences between all regions of the world will become apparent.

Disclosure of Interest
None Declared

P113
Juvenile dermatomyositis in Arab children: a multi-center retrospective analysis of disease features and outcome
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Pediatric Rheumatology 2017, 15(Suppl 1):P113

Introduction: Juvenile dermatomyositis (JDM) is a multisystem inflammatory disease affecting primarily the skin and muscles. The clinical manifestations of JDM have been extensively described from different geographical parts of the world. The available published data from Arab countries about JDM is very limited.

Objectives: To describe disease characteristics and outcome of Arab children with JDM.

Methods: We retrospectively reviewed children with JDM seen between 1990 and 2014 in 5 rheumatology clinics from 3 Arab countries. All included patients fulfilled Bohan and Peter criteria for JDM; diagnosed before 14 years of age and were Arab ethnicity. Data were collected at the last follow up visit and comprised of age at presentaion, follow up duration, clinical and laboratory features, treatment as well as the long-term outcomes including accrual disease damage and death related to JDM.

Results: A total of 89 JDM patients (55 girls) from Saudi Arabia (65.2%), Jordan (18%) and Oman (16.9%) were included. The mean age at onset was 6 years (±3), the mean age at diagnosis was 6.6 years (±3), and with mean follow up duration of 5 years (±4.4). Fifty patients had active follow up, 27 patients transferred to adult rheumatology service and 12 patients had lost follow up. Forty-nine patients had polyyclic disease course, 34 had monocular course while 16 had continuous progressive course. Seventy-four patients had organ involvement, arthritis (48%), upper airway and dysphagia (14%), gastrointestinal (13%) and lung involvement (10%). Seven patients admitted to intensive care unit (ICU), 4 of them required mechanical ventilation. Dysphagia, upper airway and lung involvement were all statistically significant associated with ICU admission. All patients received corticosteroids; while methotrexate was the most frequently used immunosuppressive drugs (86%) and rituximab was used in 8 patients. Additionally 31 patients received IVIG. Twelve patients received pamidronate mainly for calcinosis. Most of the patients achieved complete clinical response but 16 ended with permanent skin changes and 12 had residual muscle weakness. Twenty-seven patients developed calcinosis, 15 had osteoporosis, and 3 had lipodystrophy while 37 patients had growth failure. One patient developed mucinous cystic ovarian tumor. There were two deaths due to infection during the follow up period.

Conclusion: Arab patients with JDM have similar characteristics to previously described cohorts. However, compared to our previous experience, a steadily improvement in outcome particularly calcinosis has been observed. Only better therapeutic approach.

Disclosure of Interest
None Declared
P114
Nailfold capillaroscopy in children with juvenile dermatomyositis
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Pediatric Rheumatology 2017, 15(Suppl 1):P114

Introduction: Juvenile dermatomyositis (JDM) is a systemic inflammatory microvasculopathy that primarily affects skin and muscle. The clinical course of JDM is variable and correlates with severity of vascular disease and vascular alternations taking place in the muscle, skin and diverse organs. The microangiopathy characteristic of JDM can be confirmed by nailfold capillaroscopy (NC). NC is an important tool for noninvasive study of peripheral microcirculation in children with dermatomyositis.

Objectives: The aim of the study was to compare disease activity and nailfold capillaroscopy findings in children with juvenile dermatomyositis.

Methods: 13 children with diagnosed dermatomyositis (3–17 years old, 6 girls and 7 boys) were included in the study. Age of the patients at the time of diagnosis was: 4–14 years, duration of illness at the time of study: 4–79 months. The evaluation of capillaries was performed using a stereoscopic microscope, the measurements were done using the program AxioVision. The capillary system was evaluated by assessing number or loss of capillaries, the number of enlarged and giant capillaries, haemorrhages, neoangiogenic capillaries and their disorganization. The evaluation was made using the scoring system of the capillaroscopic parameters. The score at the time of diagnosis and during improvement or remission was compared.

Results: At the time of diagnosis microangiopathy with scleroderma-like pattern was detected in all patients. The most characteristic feature were reduced number of capillaries, neoangiogenic and enlarged capillaries. The capillary pattern was dependent on disease activity.

Conclusion: Capillaroscopy is simple and noninvasive method for the diagnosis and evaluation of disease activity in children with juvenile dermatomyositis.

Trial registration identifying number: 1Sulli A, Secchi ME, Pizzorni C, Cutolo M. Scoring the nailfold microvascular changes during the capillaroscopic analysis in systemic sclerosis patients. Ann Rheum Dis. 2008;67:885-7.

Disclosure of Interest None Declared

Poster Session: Scleroderma and related syndromes

P115
A single centre experience of pediatric systemic sclerosis from North India
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Pediatric Rheumatology 2017, 15(Suppl 1):P115

Introduction: Systemic sclerosis (SSc) is an inflammatory disorder with multisystem involvement characterized by progressive fibrosis affecting skin and various internal organs. It is subdivided clinically by the extent of skin involvement into diffuse and limited systemic sclerosis.

Objectives: We describe clinical and laboratory profile and outcome of pediatric SSc patients diagnosed and managed at a single centre in North India.

Methods: This was a retrospective case review of all children diagnosed as SSc based on Pediatric Rheumatology European Society/American College of Rheumatology/European League against Rheumatism provisional classification criteria for Juvenile Systemic sclerosis.

Results: A total of 14 children (8 boys and 6 girls) were diagnosed as SSc between 1990 and 2014. Diffuse and limited cutaneous SSc were seen in 10 and 4 patients, respectively. Median age at symptom onset and diagnosis was 7 years (range of 4–14 years) and 10 years (range 6–15 years), respectively. Progressive skin tightening (78%) and Raynaud phenomenon (71%) were the most common presenting manifestations. Raynaud phenomenon as the only presenting manifestation was present in 21% patients. Other common presenting features included restriction of joint movements due to skin tightening (57%), hyperpigmentation of skin (21%), arthritus (14%), dysphagia (30%) and dyspnoea on exertion (14%). Calcinosis and palpitation were the presenting manifestations in 1 patient each.

On evaluation, skin involvement (digital tip ulcers, calcinosis, Raynaud phenomenon, salt-pepper pigmentation, telangiectasia, and acro-osteolysis) was present in all patients. Median Modified Rodnan Skin Score (MRSS) at presentation was 18. Skin biopsy was done in 8 patients and was suggestive of sclerodermatous skin changes in all. Capillaroscopy was done in 3 patients and was suggestive of capillary drop outs and tortuous loops.

Pulmonary involvement was seen in 10 patients. Only 2 of these were symptomatic at presentation. One patient developed ILD after 2 years of initial diagnosis of SSc. Pulmonary function test (PFT) was suggestive of restrictive pattern in 9/13 patients. Gastrointestinal tract involvement was seen in 30% patients with dysphagia as the most common clinical manifestation. Median juvenile Systemic Sclerosis Score (JSSS) was 6.5 at presentation (range 3–12).

ANA was positive in 10/14 patients [speckled (4), nucleolar (4), mixed (1) and diffuse (1)]. RA factor was positive in 3/14 patients.

CT chest showed features of ILD in 6/9 patients with ground glass opacity and honeycombing being common abnormalities. Mean follow up was for 2.07 years (3 months to 6 years). In majority of patients with major organ involvement, the treatment regimen included cyclophosphamide pulse therapy followed by oral azathioprine. No mortality was reported in our cohort of patients; however recent follow up was available for only 7 patients.

Conclusion: In our cohort, we found a significant male preponderance. Progressive skin tightening and Raynaud phenomenon were the most common presenting manifestations.

Disclosure of Interest
None Declared

P116
Juvenile Localized Scleroderma (JLS): is it a benign disease?
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Presenting author: Christine Arango
Pediatric Rheumatology 2017, 15(Suppl 1):P116

Introduction: JLS is a polymorphic autoimmune disease. The Pediatric Rheumatology European Society (PRes) classification criteria help to identify different clinical features. The follow up of the patients allow the recognition of morbidity and complications derived from the disease.

Objectives: Describe demographic, clinical features, morbidity and sequelae derived from JLS.

Methods: Multicentre descriptive retrospective study. Analysis of clinical records of patients with a definitive diagnosis of JLS in 10
pediatric rheumatology clinics. The patients had a minimum follow-up of 6 months and a minimum of 12 months of diagnosis. Morbidity was defined as: unsightly effects (dyscromic, disfigurement lesions and/or alopecia). Sequela was defined as functional joint involvement and growth disturbances (longitudinal, circumferential or mixed).

Results: n = 88. Mean age at diagnosis 6.9 years (0-14 years) with a sex ratio M:F 2:1.1. The mean follow up time was 43 months (6-243 months). Mean time between symptoms appearance and diagnosis was 16.5 months (1-96 months). AAN were positive in 42% of patients. According to PRES criteria the distribution was: circumscribed morphea (33%), mixed (32%), linear (22%), generalized (11%) and panesclerotic (2%). The clinical phenotypes of the predominant subtypes are described in Table 41. 67% developed multiple lesions. 22% had extra cutaneous involvement (neurological, ocular and articular) and 13% had another autoimmune disease. Circumscribed was the most common type, morbidity was related to unsightly effects. In mixed type the association between linear and circumscribed lesions account for 75%. This type had multiple complications (unsightly and functional effects). Linear lesions induced the three types of morbidity and showed the highest extra cutaneous involvement and a significant morbidity. Generalized scleroderma developed important unsightly and functional effects. Pansclerotic was the least common, 2 girls being affected by a severe crippling disease. The frequency of functional joint involvement, growth disturbance and multiple complications was higher in the mixed, linear and generalized types, being statistically significant.

Conclusion: JSC is a polymorphic and unpredictable disease that determines important morbidity. Late diagnosis is common and might have a negative impact on prognosis because it allows the progression of the lesions, their size, depth and number. The high number of extra cutaneous complications reflects that this is not a disease limited to the skin. Polyautoimmunity reflects an immune system dysregulation. An early diagnosis and a more dynamic immunosuppressive treatment may improve the prognosis. Patients need a regular and extended follow up.

Disclosure of Interest
None Declared

Table 41 (abstract P116). Clinical features of the most prevalent subtypes

| Circumscribed | Mixed | Linear | Generalized | CDS | Total | P value |
|---------------|-------|--------|-------------|-----|-------|---------|
| Frequency (%) | 29    | 28     | 12          | 10  | 7     | 86      |
| Mean age at diagnosis(year) | 8.0   | 6.2    | 8.1         | 7.3 | 5.9   | 6.9     | 0.255 |
| Disease duration at diagnosis | 11.9  | 22.6   | 13.6        | 12.6| 23.9  | 16.8    | 0.623 |
| Extra cutaneous involvement (% | 26.9  | 72.5   | 30.2        | 26.0| 34.2  | 17.04   | 0.194 |
| Autoimmune associated disease (%) | 4(3.8)| 4(14.3)| 2(16.7)     | 1(10)| 0(0)  | 11(12.8)| 0.054 |
| Unsightly effects (%) | 25(86.2)| 25(89.3)| 12(100)     | 9(90)| 7(100)| 78(89.7)| 0.067 |
| Joint functional involvement (%) | 0(0) | 1(3.6) | 6(50)       | 6(60)| 0(0)  | 27(31.4)| 0.000 |
| Growth disturbances (%) | 3(10.3)| 17(60.7)| 7(58.3)     | 2(20)| 5(71.4)| 34(39.5)| 0.000 |
| 2 or more complications (%) | 3(10.3)| 19(67.9)| 7(63.3)     | 7(70)| 5(71.4)| 41(47.7)| 0.001 |

P117

Magnetic resonance and echocardiographic strain rate imaging for the early detection of cardiac involvement in Juvenile Systemic Sclerosis

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Pediatric Rheumatology 2017, 15(Suppl 1):P117

This abstract is not included here as it has already been published.

P118

Update on the Juvenile Systemic Sclerosis inception cohort project. Characteristics of the first 80 patients at first assessment. Juvenile Systemic Sclerosis Project (JSSP)

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Introduction: Juvenile systemic sclerosis (JSSc) is an orphan autoimmune disease. Currently just retrospective data exist regarding evolution of organ involvement. In the retrospective studies assessment of the organ involvement is not standardized. Our project is the first project, where prospectively and with a standardized assessment of JSSc patients are collected. We present the data of the patients at the entry into the cohort.

Objectives: to learn about the characteristics and evolution of JSSc Methods: Patients with JSSc according the PRES criteria, were recruited worldwide and were prospectively assessed, using the proposed standardized patient assessment protocol.

Results: 26 centers from 17 countries applied to participate on the project. The assent and consent forms were translated into the local native languages. Up till now 80 patients were enrolled. Sixty-six (82.5%) of the 80 patients were female. The mean age of the onset of Raynaud symptomatic was 9.4 years (0.2 – 15.9). The mean age at the onset of the non-Raynaud symptomatic were 9.9 years (0.3 – 15.9). 58 (72.5%) had diffuse subtype and 22 had limited subtype (27.5%). 11 (14%) of them have an overlap symptomatic, in 6 (10%) in the diffuse and 5 (23%) in limited subtype.
At the time of the inclusion the mean modified Rodnan Skin Score was 15.7 (0–51). ANA positive were 60/77 (78%), 24/77 (31%) of them were anti-Scl 70 positive and 4/46 (9%) were anticientromere positive. 48/80 (60%) had already capillary changes and 36/78 (46%) inactive ulcerations, 13/78 (17%) had active ulceration at the time of the inclusion. 39/80 (49%) had cardiopulmonary involvement, 22/45 (49%) of had signs of interstitial lung disease on imaging, 16/36 (44%) had FVC <80% and 10/19 (52%) had DLCO <80%. 6/44 (14%) patients had pulmonary hypertension. The mean 6 Minute Walk Test was 419.3 m (60-615). 5/80 (6%) had renal involvement, none of them had hyper tension. 26/80 (32.5%) had gastrointestinal involvement, and 18/26 (69%) of them esophageal involvement. 48/79 (62%) had musculoskeletal involvement, mostly limited range of joint movement in 21/49 (43%). Isolated muscle weakness occurred in 3/46 (6.5%). 2/80 (2.5%) showed neurologic involvement.

Conclusion: We present the data on the first 80 patients with JSSc at the time of inclusion in our cohort. The current recruitment data confirms the important role of extracutaneous involvement and more effective treatment options. There is a significantly higher proportion of diffuse subset patients with 72.5%. 14% of the patients had an overlap features. The patients had significant disease burden assessed by patients and physician at the entry of the cohort.

Disclosure of Interest
None Declared

P119 Proposal of assessment of the activity of Juvenile Localised Scleroderma. Results of the consensus meeting in Hamburg, Germany December 2015
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Pediatric Rheumatology 2017, 15(Suppl 1):P119

Introduction: Juvenile Localised Scleroderma (JLS) is an orphan disease which is complicated by difficulties in robust measurement of disease activity. Several outcome measures to assess disease activity have been described with some recent trials using the mLoSSI (modified localized scleroderma skin severity index) which is a validated cutaneous assessment tool in JLS. Whilst such measures are a step forward in assessment of this disease, they do not capture all aspects of the disease that are taken into consideration by clinicians when judging the degree of disease activity and therefore making decisions regarding treatment. In particular extra-cutaneous manifestations are poorly captured. With potential innovative and more effective treatment options emerging, it has become extremely important to define a validated activity index that captures skin, extra-cutaneous disease activity and patient reported outcomes in order to monitor response to treatment.

Objectives: To develop a proposal for an extended activity index, which includes extracutaneous involvement and the patient reported outcomes to better measure disease activity. This composite activity measure should be sensitive to change to assess the efficacy of treatment of JLS.

Methods: Members of the PRES Scleroderma working group and other paediatric rheumatologists and dermatologists interested in JLS met to develop an activity index using the nominal group technique in a consensus meeting in Hamburg, Germany, in December 2015. 80% agreement was chosen for the selected domains and items.

Results: The proposed domains and items to assess the activity of localised scleroderma:

Skin:
Change in skin thickening – Modified Rodnan Skin Score (mRSS); Total Skin thickness score of mLoSSI; Research tool: A, durometer B, Ultrasound

Degree of change in white waxy appearance since last visit – Marked worsening /some worsening and some improvement/no change/ significant improvement

Change in Erythema/violaceousness since last visit – mLoSSI
Change in Subcutaneous induration since last visit – Yes or no & elements of mLoSSI

Enlargement of the lesion since last visit – Yes or no (captured within mLoSSI)

New lesion per anatomical region of mLoSSI – mLoSSI
Worsening alopecia (eyelash, beard, body hair) – marked worsening./ some worsening and some improvement/no change/ significant improvement/no hair in the lesion / decreased

Itching in the lesion

Worsening of lim discrepancy – Yes or no

Active CNS involvement – Yes or no

Patient reported outcomes:
Change in subcutaneous induration since last visit – marked worsening /some worsening and some improvement/ no change/ significant improvement

Uncomfortable feeling in the lesion (not including itch) – VAS score 0-100

Itching in the lesion – VAS score 0-100

Quality of life assessment – PEDsQL, CHQ, CDLQI

Functional assessment of daily life – CHAQ/HAQ

Conclusion: We propose extended domains to define an activity index for JLS which aim to capture the heterogeneity of disease activity. This may become an important instrument for evaluating treatment efficacy but requires prospective validation from the PRES and CARRA scleroderma working groups.

Disclosure of Interest
None Declared

P120 Medication use in the Juvenile Systemic Sclerosis inception cohort. Are there differences in the diffuse and localised subset patients? www.juvenile-scleroderma.com
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Introduction: Juvenile systemic sclerosis (JSSc) is an orphan autoimmune disease. Currently there is no data regarding use of medication in JSSc patients. Our project is the first project, where prospectively and with a standardized assessment data of JSSc patients are collected and the applied medication assessed. We present the data of the medication at the time point of entry into the cohort.

Objectives: to learn about the medication use in JSSc patients.
Methods: Patients with JSSc, according to the PRES criteria, were recruited worldwide and were prospectively assessed, using the proposed standardized patient assessment protocol.

Results: Up till now 80 patients were enrolled. Sixty-six (82.5%) of the 80 patients were female. 58 (72.5%) had diffuse subtype (dJSSc) and 22 had limited subtype (27.5%) (fJSSc). 88% of the patients in the dJSSc and 86% in the fJSSc patients received medication.

Conclusion: Interestingly the most frequently used DMARD was Methotrexate, followed by Mycophenolate. Only small number of patients received biologics. Interestingly higher number of patients in the diffuse group received Bosentan. Interestingly 9 patients did not receive any DMARDS /biologics or medication again pulmonary hypertension.

Disclosure of Interest
None Declared

Table 42 (abstract P120). See text for description

| Number of patients | Whole group | Diffuse subtype | Limited subtype | P value |
|--------------------|-------------|----------------|----------------|---------|
| Sex female /male    |             |                |                |         |
| 80                  | 57 (71%)    | 23 (29%)       |                |         |
| Medication          |             |                |                |         |
| 66/14 = > 4.71      | 47/10 = > 4.71 | 20/4 = > 5.1 |               |         |
| 62/71 (87%)         | 44/50 (88%) | 18/21 (83%)    |               |         |
| Corticosteroids     | 36/62 (58%) | 27/44 (61%)    | 9/18 (50%)     | 0.067   |
| Cyclophosphamide    | 5/62 (8%)   | 5/44 (11%)     | 0/18 (0%)      | 0.159   |
| Chloroquine/Hydroxy- | 10/62 (16%)| 6/44 (14%)     | 4/18 (22%)     | 0.485   |
| Chloroquine/Hydroxy- | 10/62 (16%)| 6/44 (14%)     | 4/18 (22%)     | 0.485   |
| Chloroquine/Hydroxy- | 10/62 (16%)| 6/44 (14%)     | 4/18 (22%)     | 0.485   |
| Chloroquine/Hydroxy- | 10/62 (16%)| 6/44 (14%)     | 4/18 (22%)     | 0.485   |
| Methotrexate         | 35/62 (56%) | 24/44 (54%)    | 11/18 (61%)    | 0.804   |
| Mycophenolate        | 11/62 (18%)| 8/44 (18%)     | 3/18 (17%)     | 0.905   |
| Azathioprin          | 1/62 (2%)  | 1/44 (2%)      | 0/18 (0%)      | 0.524   |
| Tocilizumab          | 1/62 (2%)  | 0/44 (0%)      | 1/18 (6%)      | 0.125   |
| Rituximab            | 2/62 (3%)  | 1/4 (2%)       | 1/18 (6%)      | 0.523   |
| Bosentan             | 11/62 (18%)| 10/44 (23%)    | 1/18 (6%)      | 0.165   |
| PDE5 inhibitors      | 4/62 (6%)  | 3/4 (7%)       | 1/18 (6%)      | 0.863   |
| Prostanoids          | 1/62 (2%)  | 0/44 (0%)      | 1/18 (6%)      | 0.125   |

P121
Cone beam computed tomography for evaluation of Craniofascial abnormalities in case of linear scleroderma en coup de sabre

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Pediatric Rheumatology 2017, 15(Suppl 1):P121

Introduction: Scleroderma en coup de sabre (SECDs) is severe disease variety, that results in hemiatrophy of skin, soft tissues, bones accompanied by aesthetical, odontostomatologic abnormalities, central nervous system, ocular involvement, deformation of skull bones. Therapeutic response to immunosuppressive therapy in SECDs is rather poor in comparison to linear scleroderma of limbs and trunk. Identification of scleroderma lesions of skull bones seems difficult, but very important for treatment efficacy estimation and planning of further reconstructive plastic surgery.

Objectives: To estimate the development of skull bones disturbances in children with SECDs under disease-modifying therapy we used cone beam computed tomography (CBCT)

Methods: All the images were acquired using CBCT scanner (Gendex CB-500, powered by i-cat, Germany). We measured height of frontal, alveolar, maxillar and mandibular bones in the coronal & axial sections; bone density in Houndsfield units (HU), thickness of soft tissues, comparing left and right parts of facial skull.

Results: CBCT scanning of facial skull were performed 5 girls with SECDs aged from 8-18 years, in 2- CBCT scanning was done twice in 6 months intervals. Disease duration ranged from 8 months to 10 years (mean was 8.5 years). All patients have received disease-modifying therapy-corticosteroids orally 0.5-1 mg/kg for 12 weeks plus methotrexate 10-12 mg/body sq.weekly for 24-36 months, in 3 girls after 3 years the treatment was withdrawn, as scleroderma skin progression stopped and reversed partly. In all the cases we detected decrease of soft tissue amount on the site of scleroderma damage. In 4 girls (disease duration more than 7 years) height of maxillar and mandibular bones was reduced on involved side of face. Frontal bones were thinned, with abnormal architectonics. Shape deformations of bones were significant. Density of underlying bones was diminished (143-148 HU) in cases with skin scleroderma changes in active phase (edema, induration), on the contrary- fibrotic skin and soft tissue correlates with increased bone density (833-935 HU). Repeated in 6 months CBCT scanning helped to correct disease-modifying therapy.

Conclusion: Our pilot study of CBCT scanning of facial skull in SECDs showed the method as a helpful tool in visualization of facial structures. Further investigations could find out reference points and lines for more informative scanning, give a key for interpretation of results, its correlation with clinical presentations and treatment efficacy.

Disclosure of Interest
None Declared

P122
Role of pentraxin 3 in patients with juvenile scleroderma

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Pediatric Rheumatology 2017, 15(Suppl 1):P122

Introduction: Juvenile scleroderma (JS) is a rarely seen chronic connective tissue disorder. Juvenile systemic scleroderma (JSSc) has far worse prognosis with multi-organ involvement and possible life-threatening complications. Its main pathophysiological characteristics include microvascular abnormalities and excessive fibrosis of the skin, subcutaneous tissues and internal organs. Pentraxin 3 (PTX 3) is a multifunctional protein produced at the inflammation site by macrophages, dendritic cells, endothelial cells, smooth muscle cells and fibroblasts. Among its other functions, PTX 3 suppresses fibroblast growth factor 2 (FGF 2) in systemic sclerosis. Previously studies reported an increased level of PTX 3 among adult patients with scleroderma. Study among patients with JS has not been provided yet.

Objectives: We aimed to measure the level of PTX 3 in patients with juvenile scleroderma comparing to healthy children. Thereby, we have tried to give an answer on question whether the PTX 3 could be a marker of fibrosis in patients with juvenile scleroderma.

Methods: We assessed patients with JSS and those with juvenile localized scleroderma (JLS) and age- and sex- matched healthy
controls. A complete medical history, physical examination, and laboratory evaluation were performed for each patient at the time of enrollment. The same physical examination and laboratory investigation was performed in healthy controls, in order to exclude the coincidental disease. Circulating PTX3 levels were measured by enzyme immunoassay. The lower limits of detection for PTX3 was 0.1 ng/ml.

Results: We assessed 24 patients with JSS, 20 patients with JLS and 41 healthy controls. The mean age of patients was 15.38 ± 1.156, 12.44 ± 3.63 and 14.33 ± 3.48 for JSS, JLS and healthy controls, respectively. Mean disease duration was 2 years (range: 0.6 -15 years) for JSS and 1.5 years (range: 0.6 - 18 years) for JLS patients.

Mean serum level of PTX 3 was 10.63 ± 8.61 ng/ml, 11.75 ± 9.11 ng/ml and 2.76 ± 1.338 ng/ml for JSS, JLS and healthy controls, respectively. There was remarkable increased level of PTX 3 in serum of patients with both types of juvenile scleroderma. In both of patients group (systemic and localized form of the disease), PTX 3 level was significantly higher comparing to healthy children (p < 0.001).

We didn't find statistically significant difference between JSS and JLS patients according to level of PTX3. A mean modified Rodnan skin score of JSS patients was 19.95 ± 11.088. PTX3 level was found to be in positive correlation with modified Rodnan skin score, in patients with JSS (Rho = 0.497, p = 0.030).

Conclusion: The circulating PTX3 level is significantly higher in both JSS and JLS than in healthy control subjects. The possible explanation is that fibrosis and increased fibroblast activation represent main pathophysiological mechanism in both form of the disease. This result support relevance of PTX 3 measurement in order to determine fibrosis activity.

Since it is produced by activated fibroblasts in the area of inflammation and fibrosis, a PTX 3 should be considered a relevant marker of fibrosis in patients with juvenile scleroderma. Disclosure of Interest None Declared

P123
Is there a difference in the presentation of diffuse and limited subtype in childhood? Results from the juvenile scleroderma inception cohort. www.juvenile-scleroderma.com

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P124
Epidemiology of Kawasaki disease in Italy: surveillance from national hospitalization records

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Pediatric Rheumatology 2017, 15(Suppl 1):123

Introduction: Kawasaki disease is a systemic vasculitis with an acute and self-limited course. In view of the potential severity of the illness, of its possible genetic background, and its unknown etiology, it is important to obtain epidemiological data in different populations. The incidence of the disease differs widely among ethnic groups and is high in the Asian population. In Italy, no recent data are available.

Objectives: The aim of the present study was to define the epidemiology of KD in the years 2008-2013 in children aged < 15 years in Italy through administrative data collection.

Methods: We studied the epidemiology of Kawasaki disease in the years 2008-2013 in children < 15 years old in Italy using hospital ICD-9 discharge codes with a thorough data cleaning for
duplicate in order to select the first hospital admission for the disease.

**Results:** Between January 2008 and December 2013, 2901 children and adolescents younger than 15 years had a first hospital admission for KD in Italy. The disease peaked in the first two years of life, with 85.5% of cases under 5 years. Male: female ratio was 1.4:1. The incidence rate was 5.7 for 100,000 children 0-14 years old and 14.7 for children younger than 5 years. The incidence rate in children younger than 5 years in different Regions was < 10 cases x 100,000 children in two small regions in Northern Italy, between 10 and 15 in the majority of the Regions, and above 15 in 5 regions in Central Italy just under the Apennines and in Sicily. The incidence rose slightly during the study period and had a seasonal distribution, with higher incidence in spring and in winter. A cardiac complication was recorded in 6.4% of the patients.

**Conclusion:** This is the first epidemiologic study on Kawasaki disease incidence in the country of Italy. Figures are in line but slightly higher than those reported for other European countries.

**Disclosure of Interest**
None Declared

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**P126**

**Behçet disease in the pediatric age: data on 129 patients collected from an Italian cohort**

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**Pediatric Rheumatology 2017, 15(Suppl 1):P126**

**Introduction:** Behçet’s disease (BD) most often affects young adults, but occasionally can have its onset in childhood. Large series describing the disease in the pediatric age are scarce.

**Objectives:** Aim of our study was to collect information on clinical characteristics and treatment in pediatric patients with BD in Italy.

**Methods:** Demographic, clinical and therapy data from pediatric patients with BD, enrolled in the Eurofever registry by Italian Pediatric Rheumatology Centers, have been analyzed. Patients enrolled met the international criteria (Lancet 1990) or were diagnosed by specialists as affected by Behçet’s disease. In the latter case the diagnosis was confirmed by a panel of expert who evaluated the clinical history.

**Results:** 129 patients were included in our study: 73 were males and 56 females. In about half of cases (n = 64) a follow-up visit was also recorded, in addition to the baseline. Ethnicity was Caucasian for almost all (125/129). Mean age at disease onset was 9 years, mean age at diagnosis 13 years. A positive family history of BD was reported in 14 cases. At the baseline visit 94.3% had muco-cutaneous symptoms; 41.5% ocular involvement; 35.9% musculoskeletal symptoms; 34.8% gastro-intestinal manifestations; 31.4% constitutional symptoms; 23.5% neurologic involvement. The most common muco-cutaneous symptoms were recurrent oral aphtosis (93%); genital ulcers (27%), pseudo-folliculitis (17%), maculopapular rash (16%), erythema nodosum (13%), acneic or papulo-pustular lesions (12%) each. Pathergy test was positive in 9 patients, negative in 68, not done in 7. Ocular involvement occurred in 37 patients: 14 had anterior uveitis, 4 posterior uveitis 5 panuveitis, 8 retinal vasculitis, 5 papillodema, 5 papillitis, 3 episcleritis, 1 band keratopathy and keratitis. The most common musculoskeletal symptom was arthralgia (n = 30), followed by myalgia (n = 16), oligoarthritis (n = 6), polyarthritis (n = 5), and monoarthritis (n = 2). Abdominal pain (n = 30) and diarrhea (n = 11) were the most common gastrointestinal symptoms, followed by Gl ulcers (n = 4), and anal ulcers (n = 2); 5 patients had gastrointestinal bleeding, one patient presented aseptic peritonitis and 2 patients gut perforation. Constitutional symptoms included recurrent fever in 22 patients, fatigue and malaise in 14. Headache was the most common neurologic symptom (n = 17); 7 patients had cranial nerve palsies, 3 presented vertigo, 1 optic neuritis and 1 aseptic meningitis. Moreover, 1 patient had ataxia and 1 presented hemiplegia and abnormal behavior. Venous thrombosis occurred in 3 patients (thrombosis of transverse sinus in one of them). Two patients also had genitourinary involvement (urethritis or cystitis), HLA-B51 was present in 39 patients, not done in 12. The main treatment used was systemic corticosteroids, followed by colchicine (n = 31) and other immunosuppressants, ie azathioprine (n = 6), methotrexate (n = 5), cyclosporine (n = 3), thalidomide (n = 2), and cyclophosphamide (n = 1). Infliximab was also used in one patient. During follow-up, other biologic agents were also used, ie Adalimumab (n = 9) and Anakinra (n = 1).

**Conclusion:** This is one of the largest pediatric BD cohorts reported so far. Our data are similar to those of other pediatric series. The performance of the new Ped-BD criteria in our series is currently being evaluated, as well as possible correlations between clinical signs or symptoms at onset with immunosuppressive treatment.

**Disclosure of Interest**
None Declared
**P127**
Endothelial dysfunction in children with Kawasaki disease and transient coronary artery abnormalities

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Introduction: Evidence of premature atherosclerosis and systemic arterial stiffening in follow up patients of Kawasaki disease (KD) and coronary artery abnormalities is accumulating. Diminished flow-mediated dilatation (FMD) and increased carotid artery stiffness index (SI) have been used as surrogate markers of premature arteriosclerosis. Both of these indices have been shown to correlate with late cardiovascular events and/or mortality in adults. Because of their noninvasive nature these parameters have recently been studied in children as well. There is paucity of studies in endothelial dysfunction in children with KD and transient coronary artery abnormalities.

Objectives: In the present study, we assessed the risk of atherosclerosis by noninvasive methods (FMD, cIMT and carotid artery stiffness index) in children with KD with transient coronary artery abnormalities at least 1 year after resolution CAA of KD.

Methods: Twenty children with KD (mean age 11.5±3.7 years) and transient coronary artery dilatation which had resolved within 1 year of diagnosis were studied at least 1 year of after resolution of CAA. All had received intravenous immunoglobulin except one during the acute phase of the disease. High-resolution ultrasonography was used to analyze brachial artery responses to reactive hyperemia and sublingual nitroglycerine. Flow-mediated dilatation was also studied in an equal number of healthy age- and sex-matched controls. Carotid artery intima-media thickness and stiffness index was calculated using B mode ultrasound and compared in all subjects using previously published equations.

Results: No statistically significant difference was noted in the percent flow mediated dilatation of the brachial arteries in response to reactive hyperemia among the cases (13.31 ± 10.41%) compared with controls (12.86 ± 7.09%; p = 0.874). Sublingual nitroglycerine-mediated dilatation in children with KD was 14.88 ± 12.03%. Mean carotid intima-media thickness was found to be almost same in both cases (0.036 ± 0.015 cms) and controls (0.035 ± 0.076 cms; p = 0.791). Also, no significant difference was noted in the carotid artery stiffness index among the cases (1.153 ± 0.34 U) when compared to controls (1.11 ± 0.40 U; p = 0.729).

Conclusion: No evidence of endothelial dysfunction in KD patients with transient coronary artery abnormalities was noted in the present study. This may point towards no risk of preclinical arteriosclerosis in children with KD with transient coronary artery abnormalities. However, as the number of cases were small, it would be imprudent to extrapolate these results and draw conclusions based on the small sample size.

Disclosure of Interest
None Declared

**P128**
Carotid intima-media thickness and lipid profile in children with Kawasaki disease: a follow-up study after 5 years

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Introduction: Kawasaki disease (KD) is the most common medium vessel vasculitis in children. It has a predilection to involve coronary arteries, leading to several long term cardiovascular sequelae. In addition to coronary artery abnormalities (CAAs), premature atherosclerosis, endothelial dysfunction and lipid abnormalities are seen in children with KD. Despite prompt treatment with intravenous immunoglobulin in the acute phase, these complications ensue.

Objectives: At our institute in Chandigarh, Meena et al. in 2009 studied carotid intima media thickness (cIMT) and lipid profile in 27 children with KD after 1 year of the acute episode of KD. This study was planned to follow-up the same cohort of children at least 5 years after the acute episode.

Methods: All the 27 children previously enrolled by Meena et al. were followed-up in the present study. cIMT, being a surrogate marker for premature atherosclerosis and fasting lipid profile were evaluated. Diagnosis of KD was based on the American Heart Association criteria. All patients had received 2 gm/kg of IVIg at the time of diagnosis. Age and sex matched healthy children were enrolled as controls from the out-patient department of the institute for cIMT. For lipid profile, comparison was made with published data on this subject. cIMT was measured in the Echocardiography lab of Advanced Cardiac Centre, PGIMER by Acuson Sequoia C- 512 Siemens Limited with 7-12 MHz linear array transducer. All children with KD were asked to fast for 8 hours overnight and 2 ml blood sample was collected. Total cholesterol, HDLC, triglycerides were estimated using standard laboratory kits manufactured by Siemens Healthcare Diagnostics. Cholesterol levels < 170 mg/dL were considered desirable and > 200 mg/dL were considered undesirable. Cholesterol levels between these limits were considered borderline. Low-density lipoprotein cholesterol (LDL-C) levels < 110 mg/dL were considered desirable, and > 130 mg/dL were considered undesirable. LDL-C levels in between these limits were considered borderline. High-density lipoprotein cholesterol (HDL-C) levels > 35 mg/dL were considered desirable, and < 35 mg/dL were considered undesirable. For triglycerides, levels < 150 mg/dL were considered desirable and > 150 mg/dL as undesirable.

Results: Mean duration between onset of fever and IVIg treatment was 14.3 days with a range of one to 90 days. At the time of diagnosis, 10 patients (37.03%) had CAAs in the form of dilatation. There was significantly higher mean cIMT in children with KD (0.54±0.087 mm) as compared to controls (0.42±0.036 mm), however there was no significant difference in cIMT at 1 and 5 years of follow up. There was no statistically significant difference among cholesterol, HDLC, LDL-C and triglyceride levels between cases with CAAs at diagnosis as compared to children without CAAs at diagnosis. Abnormal lipid profile was seen in 7 out of 27 children in the present study (1 child- borderline serum cholesterol levels, 4 children- undesirable serum HDL-C levels, and 2 children- undesirable serum triglyceride levels), 5 of whom had lipid abnormality at 1 year follow up (Meena et al.), suggesting that lipid profile abnormalities in KD are perhaps persistent and long lasting.

Conclusion: Our findings thus presuppose fertile ground for cardiovascular complications in young adulthood in patients with KD. It is likely that the consequences of KD in childhood would impact the health status of young adults several years later.

Disclosure of Interest
None Declared

**P129**
Microparticles as a therapeutic target for plasma exchange in antineutrophil cytoplasmatic antibody associated vasculitis?

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Pediatric Rheumatology 2017, 15(Suppl 1):P129

Introduction: Kawasaki disease (KD) is the most common medium vessel vasculitis in children. It has a predilection to involve coronary arteries, leading to several long term cardiovascular sequelae. In addition to coronary artery abnormalities (CAAs), premature atherosclerosis, endothelial dysfunction and lipid abnormalities are seen in children with KD. Despite prompt treatment with intravenous immunoglobulin in the acute phase, these complications ensue.

Objectives: At our institute in Chandigarh, Meena et al. in 2009 studied carotid intima media thickness (cIMT) and lipid profile in 27 children with KD after 1 year of the acute episode of KD. This study was planned to follow-up the same cohort of children at least 5 years after the acute episode.

Methods: All the 27 children previously enrolled by Meena et al. were followed-up in the present study. cIMT, being a surrogate marker for premature atherosclerosis and fasting lipid profile were evaluated. Diagnosis of KD was based on the American Heart Association criteria. All patients had received 2 gm/kg of IVIg at the time of diagnosis. Age and sex matched healthy children were enrolled as controls from the out-patient department of the institute for cIMT. For lipid profile, comparison was made with published data on this subject. cIMT was measured in the Echocardiography lab of Advanced Cardiac Centre, PGIMER by Acuson Sequoia C- 512 Siemens Limited with 7-12 MHz linear array transducer. All children with KD were asked to fast for 8 hours overnight and 2 ml blood sample was collected. Total cholesterol, HDLC, triglycerides were estimated using standard laboratory kits manufactured by Siemens Healthcare Diagnostics. Cholesterol levels < 170 mg/dL were considered desirable and > 200 mg/dL were considered undesirable. Cholesterol levels between these limits were considered borderline. Low-density lipoprotein cholesterol (LDL-C) levels < 110 mg/dL were considered desirable, and > 130 mg/dL were considered undesirable. LDL-C levels in between these limits were considered borderline. High-density lipoprotein cholesterol (HDL-C) levels > 35 mg/dL were considered desirable, and < 35 mg/dL were considered undesirable. For triglycerides, levels < 150 mg/dL were considered desirable and > 150 mg/dL as undesirable.

Results: Mean duration between onset of fever and IVIg treatment was 14.3 days with a range of one to 90 days. At the time of diagnosis, 10 patients (37.03%) had CAAs in the form of dilatation. There was significantly higher mean cIMT in children with KD (0.54±0.087 mm) as compared to controls (0.42±0.036 mm), however there was no significant difference in cIMT at 1 and 5 years of follow up. There was no statistically significant difference among cholesterol, HDLC, LDL-C and triglyceride levels between cases with CAAs at diagnosis as compared to children without CAAs at diagnosis. Abnormal lipid profile was seen in 7 out of 27 children in the present study (1 child- borderline serum cholesterol levels, 4 children- undesirable serum HDL-C levels, and 2 children- undesirable serum triglyceride levels), 5 of whom had lipid abnormality at 1 year follow up (Meena et al.), suggesting that lipid profile abnormalities in KD are perhaps persistent and long lasting.

Conclusion: Our findings thus presuppose fertile ground for cardiovascular complications in young adulthood in patients with KD. It is likely that the consequences of KD in childhood would impact the health status of young adults several years later.

Disclosure of Interest
None Declared
Introduction: Plasma exchange (PE) is a renal function preserving and potentially life-saving therapy in children and adults with anti-neutrophil cytoplasmic antibody (ANCA) associated vasculitis (AAV). The therapeutic mechanisms of PE in AAV are poorly understood, but it is likely that PE removes autoantibodies and other plasma constituents involved in AAV. We have previously shown that circulating neutrophil microparticles (MP) are key inflammatory mediators underpinning the pathogenesis of AAV.

Objectives: The aim of this project was to investigate whether MP are removed from the circulation of patients with AAV by PE, and therefore are potentially an important therapeutic target. This was an ancillary study of a major international clinical trial of therapeutic PE in AAV, the PEXIVAS trial (NCT #NCT00987389).

Methods: Patients with AAV undergoing PE for active disease measured by the Birmingham Vasculitis Activity Score (BVAS) were recruited from the Royal Free Hospital and Great Ormond Street Hospital NHS Foundation Trust. Annexin V+ MP’s were isolated from platelet poor plasma, cellular origin defined (neutrophil, platelet, endothelial) and MP quantified using flow cytometry. The soluble inflammatory mediators in EDTA plasma were measured using multi-array electrochemiluminescence (MSD). Results were expressed as median and range.

Results: A total of seven patients undergoing PE at diagnosis of AAV were recruited: median age 64 years (range 17-22 years old); 4 male. Median BVAS was 15/63 (12-18/63); all patients studied had rapidly progressive glomerulonephritis. The number of total circulating AnV + MP reduced by ~60% after 3 cycles of PE from a median of 5.1 × 10⁶ /ml (range of 11.2 – 4.78 × 10⁶ /ml) to a median 1.97 × 10⁶/ml (0.36 × 10⁶ – 3.7 × 10⁶/ml), p = 0.04. Neutrophil MP identified as CD66b+ AnV+ were reduced from a median of 58 × 10³ /ml (range of 88 × 10³ – 40 × 10⁶/ml) to a median of 15 × 10⁵/ml (range of 17 × 10³ – 11 × 10⁵/ml), p = 0.02. Angiogenic factors (VEGF and Ang-1) were lower in patients with active AAV, comparing with health control, but no change after PE. In patients with active AAV, significantly increased level of Ang-2 was detected, comparing with healthy control, furthermore, levels of Ang-ll declined from a median of 8179 pg/ml (range of 3440-17313 pg/ml) to median of 3776 pg/ml (range of 775-88 × 10³ pg/ml), p = 0.03, similar to normal values (median of 2747 pg/ml, range of 1063-9296 pg/ml). Similarly, the plasma level of sICAM (a soluble marker of endothelial activation) were increased in patients with active AAV, and there was a significant reduction pre and post PE, p = 0.01.

Conclusion: Our preliminary results suggest that PE is an effective method to remove these circulating mediators of inflammation and endothelial injury. It remains to be elucidated if the removal of excess injury factors is crucial to ameliorate endothelial damage in patients with AAV.

Disclosure of Interest None Declared

P132
Early onset of behcet disease: a case report
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Pediatric Rheumatology 2017, 15(Suppl 1)P132

Introduction: Behçet Disease (BD) is a chronic inflammatory disorder with multisystemic involvement. The aetiology is unknown and it is characterized by clinical features characteristic of both autoimmune and auto-inflammatory diseases.

Objectives: Case report of early onset BD.

Methods: A 13 year-old boy was admitted to our hospital to investigate the presence of recurrent oral aphthosis, acniform and papulo-pustular skin lesions, bronchiectasis associated to recurrent respiratory infections, arthritis, and growth retardation. His family history showed recurrent oral aphthosis in his mother. He was born after 36 weeks of gestation via vaginal delivery, with APGAR score 9/10. Birth weight was 3,450 Kg. At the third month of age, oral candidiasis associated to perioral and genital papulo-pustular skin lesions appeared. After 6 months, he developed recurrent fevers (T° max 39-40 °C), monthly. At 3 years of age, the papulo-pustolar skin lesions spread to buttocks and ears and, after one year, the boy developed anterior uveitis. Since he was affected by growth delay associated to increased fecal calprotectin, he underwent an EGDS, that showed features of autoimmune enteropathy (mucosal architectural changes including villous atrophy, increased intraepithelial lymphocytes and...
10 or more eosinophils per high power field); autoantibodies (transglutaminase ab, ANA, p-ANCA, c-ANCA, anti-dsDNA, anti-Jo1, anti-Sm, anti-RNP, anti-Scl70, anticientromere and anti-SSA) were negative. At 5 years of age, genital ulcers and erythema nodosum on legs (anterior tibial regions), hands and elbow appeared.

Results: He was then admitted to our hospital for further investigations. On physical examination, he presented rather good general conditions, oral aphthosis, acniform and papulo-pustular skin lesions on perioral, ears and genital regions. Blood tests revealed an increase of inflammatory markers (ERS 35 mm/1 h, CRP 4.51 mg/L, SAA 70 mg/L). Liver and renal functions were normal. Autoantibodies (ANA, p-ANCA, c-ANCA, anti-dsDNA, anti-Jo1, anti-Sm, anti-RNP, anti-Scl70, anticientromere and anti-SSA) were performed as well, and resulted all negative except for ANA (1:1280). Genetic tests for MEFV, MVK, CIAS-1 and TMEM173 were negative. HLA-B51 was positive. Cerebral MRI, MR angiography and echocardiography were performed, and resulted normal. He started Colchicine (1 mg/die), topical corticosteroids, antimicrobial prophylaxis and airway clearance tests.

Conclusion: The diagnosis was confirmed according to the criteria of the "International Study Group for the Diagnosis of Behçet Disease" (Recurrent oral aphthosis, genital ulceration, skin involvement, anterior uveitis). Looking at the early onset of disease and the positive family history we have also considered the monogenic disease with A2 OAS, recently described, and caused by high-penetrance germline mutations in TNFAIP3, which encodes the NF-kB regulatory protein A20, leading to a Behcet-like disease with early-onset systemic inflammation. Genetic test results are pending.

Disclosure of Interest
None Declared

P133
A quality improvement initiative to improve urinalysis follow-up in henoch-schonlein purpura: retrospective review and prospective intervention

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Pediatric Rheumatology 2017, 15(Suppl 1):P133

Introduction:
Henoch-Schönlein purpura (HSP) is the most common systemic vasculitis of childhood with an incidence of 10–20 cases per 100,000 children per year. Generally the prognosis is good, with exception of those with significant renal involvement. The principal aim of HSP follow-up is mostly to detect persistent renal inflammation that could progress to renal damage. It was our observation that the weak link in the care of these children at our institution was urinalysis (UA) follow-up after the first patient contact with the Nationwide Children's Hospital (NCH) medical system. In the long-term, the HSP renal disease may be serious in these children and a simple cost-effective intervention may detect HSP renal disease earlier in the disease process and improve outcomes.

Objectives: This quality improvement (QI) project’s overall objective was to design an intervention that might improve the follow-up with UA screenings by educating families and asking care providers to do the urinalyses on schedule. The specific aim was to increase the performance of follow-up urinalyses from baseline by 21% over 6 months and by 30% over 12 months by use of this QI intervention.

Methods: Approval for this study was obtained from the Institutional Review Board of NCH in 2014. We identified HSP patients in our electronic medical record (EMR) EPIC using the US codes ICD 9 and ICD 10 for HSP. 105 HSP patients were identified retrospectively who had a diagnosis of HSP from July 1, 2014 to December 31, 2014. Their records were reviewed for diagnosis, date of diagnosis, location of clinic visit, new or old HSP diagnosis, dates and results of urinalyses at diagnosis and on follow-up, and diagnosis of renal disease by nephrologists and results of kidney biopsy. Similarly patients were identified and charts reviewed prospectively from January 1, 2015 to December 31, 2015 for the same data. The review of the 2014 charts helped identify the QI key drivers of provider factors, process factors, and patient factors that informed the intervention. The authors agreed on a consensus schedule for follow-up UA’s. A dot phrase was developed and inserted into the EMR of NCH which contained education about HSP, the rationale for the follow-up urinalyses, and our suggested schedule. The availability of the dot-phrase was advertised widely to all care providers in the NCH system before January 1, 2015. The data was collected month to month in 2015 on 239 HSP patients and UA results obtained from the EMR and primary care providers.

Results: Follow-up urinalysis performance for the HSP patients in 2014 was 21%. The AVS was given out to families in 2015 at rates from 75-95% in different outpatient sites. With the intervention during 2015, the urinalyses were performed at the rate of 61% from January 1 to June 30th and 55% over the second 6 months. The changes were statistically significant (for mean change p = 0.0075). Four children initially had a normal urinalysis at their first visit and then developed an abnormal UA; 3 patients’ abnormal UA resolved over the next 8 months but one developed chronic renal disease.

Overall, 2 children developed chronic renal disease with IGA nephropathy on kidney biopsy.

Conclusion: Our QI intervention was successful in achieving our goal of increasing follow-up urinalysis performance in HSP patients seen at HSP in 2015. We believe such an inexpensive QI intervention is beneficial to the HSP patients but further study in a collaborative network is needed to determine if it actually detects renal disease earlier and improves outcomes.

Disclosure of Interest
None Declared

P134
Successful treatment of digital necrotizing vasculitis in patients with polyarteritis nodosa

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Pediatric Rheumatology 2017, 15(Suppl 1):P134

Introduction: Polyarteritis nodosa is a rare necrotizing vasculitis which affects small and medium-sized arteries and characterized by multi-organ involvement. Gangrene digitale and extremity loss via necrotizing vasculitis have been reported in polyarteritis nodosa. Although immunosuppressive agents, vasodilator, anticoagulants are used, there is no definitive treatment protocol.

Objectives: Basic 15 year old male was admitted to our hospital with painless, redness, bruising on both the fingers, toes and redness on the cheeks. The complaints of paleness, coldness, redness at fingertips began five days before admission to hospital. In a few days bruising, more prominently on toes, appeared on fingers and toes. Anamnesis and the family history was not significant. In the first examination, vital signs were stable, growth and development were compatible with the age, 2nd 3rd and 4th fingertips of left foot's color were black and necrotizing. Capillary refilling time was apparently prolonged. Laboratory analyses showed WBC of 15.040/mm³(5,000-14,500), hemoglobin of 10.4 gr/dl (12.16), platelet of 387,000/mm³ (150,000-450,000), erythrocyte sedimentation rate (ESR) of 86 mm/h (0-20), C-reactive protein (CRP) of 102 mg/L (0-6), antistreptolysin O (ASO) of 3280 IU/mL (0-200), aPTT of 34.1 sec (26-38), protrombine time 12.7 sec (11-14), INR of 1.09 (0,8-1,2), D-dimer 2.25 mg/L (0-0,55).

Urinalysis, creatinin levels, serum levels of transaminase were normal and anti-HIV, anti-HAV, anti-HBV, and anti-HCV antibodies were negative. Anti-nuclear antibody (ANA) titre was 1/100. Anti-ds DNA, profile
of Anti-ENA, p-ANCA, c-ANCA, Anti-cardiolipin IgM and IgG antibodies, lupus anticoagulant and crioglobulin were negative. C3, C4, Protein C, Protein S, antithrombin III, vWF, homocysteine levels were normal. FVIII and FIX levels were increased. Enhanced CT angiography demonstrated narrowing and contour irregularities at left kidney, mesenteric and both tibialis posterior arteries. PA lung and sinus graphies were normal. Patient was screened for MEV mutation and homozgyous R202Q/R202Q and M694V/N were detected. The patient was homozygous for MTHFR C677T. With these findings, patients was diagnosed as poliarteritis nodosa (PAN). Methylprednisolone 1gr/day was given in three consecutive days and 1gr/month cyclophosphamide was started. After these treatment, 60 mg/day orally prednisolone therapy and simultaneously, 30 sessions of hyperbaric oxygen therapy was applied. Treatment of the first five days of hyperbaric oxygen therapy, illoprost (0,5 mg/kg/min) was given with IV infusion. Low molecular weight heparin (enoxaparin) (2 mg/kg/day) was administered for 6 months. Mycophenolate mofetil (1200 mg/m²/day) was added to therapy at the end of the cyclophosphamide pulse. After 30 sessions of hyperbaric oxygen therapy, the patient was discharged.

Results:

Conclusion: Vasculitis which can lead to loss of extremity, hyperbaric oxygen therapy that might be considered as a concomitant treatment with drugs has an important contribution to the preservation of tissue and extremity.

Disclosure of Interest
None Declared

P135
Paediatric large vessel vasculitis presenting as idiopathic retroperitoneal fibrosis: case report and review of the literature

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Pediatric Rheumatology 2017, 15(Suppl 1)P135

Introduction: Idiopathic retroperitoneal fibrosis (RPF) is a rare disease characterised by development of fibro-inflammatory mass in retroperitoneum. RPF is very rare among children and is considered a subtype of chronic periarteritis that might be associated with large-vessel vasculitis or other autoimmune disorders.

Objective: To analyse an unusual case of retroperitoneal fibrosis in a teenage girl progressing towards predominantly large vessel vasculitis and to review existing paediatric literature.

Methods: Case report and literature review

Results: A 13-year-old previously healthy Caucasian girl presented with recurrent fever, elevated inflammatory markers and back pain that initially mimicked pyelonephritis. Recurrence of symptoms and later development of supravacular lymphadenopathy promoted imaging (ultrasound US, MRI) which confirmed presence of large lymph node (LN) packet with inflammatory rather than malignant appearance on PET-CT. Transient partial spontaneous regression of her symptoms as well as non-specific inflammatory histopathology changes of her cervical LN supported conservative watch-and-wait approach by the paediatric oncology team. The third relapse with systemic hypertension as an additional new symptom led to the open biopsy from the retroperitoneal mass which excluded lymphoma. Repeated MRI revealed new involvement of aorta at the level of renal arteries and supported suspicion of vasculitis. Review of the history, imaging and biopsy results led to the diagnosis of RPF with progression to secondary vasculitis. General symptoms improved on corticosteroid (CS) therapy, but systemic hypertension as a consequence of the right renal artery occlusion progressed and resulted in nephrectomy. Aggressive immunosuppression with CS and i.v. cyclophosphamide did not abate the process which eventually affected abdominal aorta, iliac, renal and pulmonary arteries. Postcontrast gadolinium enhancement was noticed in the wall of these large vessels corresponding to persistent inflammatory activity despite the disappearance of the fibrous/LN retroperitoneal mass. After introduction of rituximab and later also low-dose weekly methotrexate (MTX) disease activity slowly diminished. After 18 months of this therapy she has been able to reduce her CS dose to 0,1 mg/kg/day and is doing well clinically, laboratory and imaging-wise. Literature search of the PubMed database from 2003 revealed 31 cases of RPF in children with typical onset age between 10 and 15 years, but none of them presented with arterioocclusive process. Around half of the patients presented with an associated autoimmune condition, and in two cases the association with malignancy was described. Majority of patients with RPF responded well to CS therapy, in refractory disease B cell-depleting therapy with rituximab was successful.

Conclusion: We present an unusual paediatric case of an overlap between RPF and large vessel vasculitis. This case is in accordance with the published adult case series demonstrating that chronic periarteritis as a component of RPF is not limited to the abdominal aorta and displays the features of primary large vessel inflammatory disease. Our choice of initial cytotoxic therapy was driven by the severe clinical picture. We have learnt that rituximab time to clinical effect may be as long as 18 months and that combination with MTX has brought additional benefit. Despite that, the long-term prognosis is warranted.

Disclosure of Interest
D. Cebecauerová: None Declared, T. Dallos: None Declared, E. Kabíčková: None Declared, M. Kynč: None Declared, D. Chroustová: None Declared, J. Hoza: None Declared, D. Němčová: None Declared, V. Tesář: None Declared, P. Doležalová Grant / Research Support from: AbbVie, Roche, Medac, Novartis, Pfizer, Consultant for: Roche, Speaker Bureau of: Pfizer, Novartis, Medac

P136
Tocilizumab treatment in childhood Takayasu arteritis: case series of four patients and systematic review of the literature

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Pediatric Rheumatology 2017, 15(Suppl 1)P136

Introduction: The mainstay of treatment for Takayasu arteritis (TAK) is high-dose corticosteroid. However, the addition of immunosuppressive agents such as methotrexate and cyclophosphamide is usually needed. Besides, several studies have shown favorable effect and good tolerance of tocilizumab (anti-interleukin 6) treatment in TAK patients.

Objectives: Our aim was to describe our experience with tocilizumab treatment in children with TAK and to review previous studies regarding tocilizumab use in TAK patients.

Methods: We reviewed the charts of all pediatric TAK patients followed between 2000 and 2015 in Department of Pediatric Rheumatology in Hacettepe University, Ankara, Turkey and we present the patients who were treated with tocilizumab. We screened PubMed and MEDLINE for articles involving TAK patients treated with tocilizumab.
Results: We have followed 11 pediatric TAK patients between 2000 and 2015. Four of them (36.3%) received tocilizumab. All four patients were female. The characteristics of these patients were summarized in Table 1. The median (minimum-maximum) age at diagnosis was 14.5 (4-16) years and the median duration of disease was 30 (7-68) months. The median duration of immunosuppressive treatment before tocilizumab onset was 16 (1-60) months. The median duration of tocilizumab treatment and the median duration of follow-up after the first dose of tocilizumab was both 9.5 (7-13) months since all four patients continue to receive tocilizumab. One of our patients received tocilizumab as a first line immunosuppressive treatment directly after one dose of pulse methylprednisolone because the family refused cyclophosphamide. Others were resistant to their initial immunosuppressive treatment (cyclophosphamide, methotrexate, azathioprine). All four of our patients achieved complete response to tocilizumab after three months of treatment. Repeated MRAs showed no new vascular lesions in our patients. ITAS2010-A values were normalised from a median of 11.5 (1-15) to 0.5 (0-1) after three months of treatment with tocilizumab. None of the patients reported any adverse events during follow-up. We identified 19 articles describing 75 TAK patients treated with tocilizumab during our literature search. Twelve patients received tocilizumab before age of 18 years including ours. Tocilizumab was given as the first line immunosuppressive treatment in seven patients including two children (one from our series). 11 out of 75 patients experienced relapse during tocilizumab treatment, while eight flared 2-14 months later than the withdrawal of tocilizumab.

Conclusion: Our small series suggests that tocilizumab may be a promising alternative for Takayasu arteritis treatment. Long-term controlled studies are warranted to provide better evidence for tocilizumab treatment in childhood Takayasu arteritis.

Disclosure of Interest None Declared

P137
Neurological manifestation in Behçet’s disease- pediatric case report
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Pediatric Rheumatology 2017, 15(Suppl 1):P137

Introduction: Behçet’s disease (BD) is a chronic multisystemic vasculitis that can affect arteries or/ and veins of any size and is characterized by triple – symptom complex of recurrent oral and/ or genital aphthous ulcers and uveitis. It can also involve gastrointestinal tract, pulmonary, musculoskeletal, cardiovascular and neurological systems (neuro- Behçet). Neurological involvement is one of the most devastating manifestations, with rare reports in pediatric age range

Objectives: Report a case of pediatric neuro- Behçet, emphasizing the importance of including vasculitis in the differential diagnosis when neurological disorders are associated with systemic manifestations.

Methods: Chart review, checking medical history, laboratory tests and imaging of a child with Behçet’s disease, according to the criteria of the International Study Group (ISG). A informed consent form was signed.

Results: M.E.S., four years old, female, admitted with headache associated with dilatation of the veins on the forehead (capitus medusa), erythema nodosum and history of recurrent oral ulcers. Added to this, it was observed bilateral papilledema and thrombosis of the superior sagittal sinus extending to the left transverse sinus, suggesting a diagnosis of Behçet’s disease. Corticosteroid and anticoagulation were introduced. The patient progressed with completed resolution of the signs and symptoms.

Conclusion: Behçet’s disease should be always considered in the presence of neurological manifestations associated with systemic involvement, avoiding permanent sequelae that affect quality of life.

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Disclosure of Interest None Declared

P138
Vitamin d receptor polymorphisms are not associated with the risk of Kawasaki disease (KD) in a group of Italian children
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Pediatric Rheumatology 2017, 15(Suppl 1):P138

Introduction: Kawasaki disease (KD) is an acute multisystemic vasculitis, affecting primarily infants and children, and it is the most common cause of acquired heart disease in Western countries. The exact pathogenesis of KD has not been clearly identified, but infections caused by one or more pathogens are strongly suspected as major cause of the disease. Our previous study demonstrated that KD patients have highly reduced 25(OH)-vitamin D levels in comparison with healthy controls and also with other immunological disorders, suggesting a hypothetical role of vitamin D deficiency in the development of coronary artery damage in KD, and low vitamin D levels might contribute to the severity of chronic vascular abnormalities. Specific polymorphisms of vitamin D receptor (VDR) gene have recently been associated with different biological response to vitamin D itself.

Objectives: 1. To analyze the possible associations of these polymorphisms in a series of Italian children affected by KD. 2. To evaluate a possible influence of VDR polymorphism in enhancing the risk of the disease.

Methods: We recruited 50 Italian infants and children affected by typical KD (38 males, 12 females, age ranging from 2 through to 62 months. Genomic DNA was extracted from peripheral blood leukocytes to analyze VDR polymorphisms by PCR-based sequencing (CD242 in the promoter region) and PCR-based enzymatic digestions (FokI in exon 2, BsmI and Apal in intron 8, and TaqI in exon 9). An Italian population of 2221 unrelated individuals without KD or other immunological disorders was used as healthy controls. The distribution of polymorphism genotypes and alleles was evaluated in patients vs healthy controls.

Results: The distribution of all Cdx2, FokI, BsmI, Apal, and TaqI polymorphisms (both as genotypes or alleles) did not show significant differences between affected subjects and healthy controls.

Conclusion: In conclusion, this pilot study evidence that VDR polymorphisms seem not to have any influence in determining the risk of KD. Anyway, future larger studies are needed to unravel the exact relationship between VDR genotype and KD development and/or clinical characteristics.

Disclosure of Interest None Declared

None Declared
The proteiform manifestations of Behçet disease. Description of 2 cases with prominent gastrointestinal and renal involvement

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Pediatric Rheumatology 2017; 15(Suppl 1)P139

Introduction: Behçet’s disease (BD) is a chronic, relapsing, systemic vasculitis of unknown etiology with the clinical features of mucocutaneou lesions, ocular, vascular, articular, gastrointestinal, urogenital, pulmonary, and neurologic involvement. The clinical presentation of the disease can be very challenging.

Objectives: We reported two children with Behçet disease presenting with atypical presentations, abrupt onset of enterorrhagia and severe glomerulonephritis respectively.

Methods: We analyzed retrospectively the clinical characteristics and manifestations in two patients diagnosed with Behçet Disease according to the International Criteria for BD (ICBD) of 2013 and the EULAR/PRINTO/PRES classification for vasculitis of 2006 associated to the Chapel Hill Consensus Conference on Nomenclature of Systemic Vasculitis of 2012.

Results: Case 1) 12 years old boy previously hospitalized in another centre because, at the age of 10, he presented with arthritis. Initially treated with oral steroids and methotrexate, he progressively presented episcleritis, recurrent oral ulcers, thrombophlebitis and pathergy test positivity. The diagnosis of Behçet disease was made and he was started on colchicine with partial remission of symptoms. After 6 months he developed severe glomerulonephritis, with hematuria, nephrotic proteinuria and systemic hypertension. Blood examinations demonstrated the presence of c-ANCA positivity and lack of HLA-B51. Renal biopsy evidenced endo and extracapillary severe proliferative glomerulonephritis. The presence of moderate-high level of C3 at immuno-fluorescence induced to rule out the possibility of an anti-neutrophil cytoplasmic antibody (ANCA)-associated vasculitis, where the complement system was considered not to be involved since immunoglobulin deposition is generally absent in the lesions. He was treated successfully with cyclophosphamide and steroid, even if the proteinuria persists with partial reduced renal functionality.

Case 2) A 7 years old boy with unremarkable history, presented with persisting fever for 20 days, haematochezia, urticarial rash, oral ulcers and photophobia. Ophthalmologic examination revealed the presence of bilateral uveitis and papillodema. Magnetic Resonance of brain resulted normal, colon endoscopy evidenced the presence of multiple lesions (abscesses). Thoracic X ray evidenced the presence of suspected pneumonia, persisting after prolonged antibiotic treatment. CT scan evidenced the presence of pulmonary micro nodules. The positivity for HLA B51 let us to formulate the diagnosis of Behçet disease and he was started on steroids and cyclosporine, with prompt remission of symptoms. Cyclosporin was later stopped and replaced with azathioprine. The patient is still in remission.

Disclosure of Interest
None Declared

Assessment of relationship between IL-17 levels and oxidative stress in children with immunoglobulin a vasculitis

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Pediatric Rheumatology 2017; 15(Suppl 1)P140

Introduction: Immunoglobulin A vasculitis (IgAV) is an IgA mediated small vessel vasculitis characterized by skin, gastrointestinal tract, kidney and joint involvement. Pathogenesis of IgAV is not clearly understood. Cytokines stimulate the production of chemokines by endothelial cells, attract inflammatory cells, induce the expression of cell adhesion molecules on endothelial cells and facilitate inflammatory cell adhesion to the vascular wall. In addition, activated cells of the immune system, could be responsible for tissue damage by producing reactive oxygen species.

Objectives: In this study, we aim to investigate the role of interleukin 17 (IL-17) and oxidative stress in the pathogenesis of IgAV and determine if there is a relationship between IL-17 and oxidative stress.

Methods: The patients, diagnosed with IgAV according to Ankara 2008 criteria were enrolled between July 2014 and August 2015. Patients who used steroid or nonsteroidal anti-inflammatory drugs in the last one month were excluded. Blood samples were obtained at active and remission phase of the disease. 8-hydroxy deoxy guanosine (oxidative DNA damage biomarker), 3-nitrotyrosine (protein oxidation biomarker) and IL-17 levels were measured via ELISA method. Malondialdehyde (lipid peroxidation substances method. Total oxidant and antioxidant status levels were measured by colorimetric analysis.

Results: Forty-four patients, including 25 (56.8%) males and 19 (43.2%) females, aged between 1.91 and 15.41 years (mean: 7.71 years) were enrolled in this study. All patients had skin involvement, 39 (88.6%) had joint, 14 (31.8%) had gastrointestinal system and five (11.4%) had renal involvement. Disease severity score was one in 22 (50%), two in 17 (39%) and three in 5 (11%) patients.

Levels of IL-17 and TAS were significantly higher in active phase of the disease compared to in remission phase (p: 0.007 and p: 0.01, respectively). A moderate positive correlation was observed between IL-17 and TAS levels in both active and remission phase of the disease (r: 0.33; p: 0.03 and r: 0.34; p: 0.02, respectively).

Levels of 8-hydroxy deoxy guanosine in active phase of disease was significantly higher in patients with gastrointestinal system involvement (p<0.003).

Conclusion: These findings suggest that IL-17 might have a role in the pathogenesis of IgAV. Beside, higher TAS levels seen in active phase of the disease might be a compensatory response to oxidative stress, which is induced by inflammation. The positive correlation between IL-17 and TAS levels in both active and remission phase of the disease also supports this compensatory response.

Disclosure of Interest
None Declared
**P141**

**Association between pediatric cutaneous polyarteritis nodosa and streptococcal infection - a single tertiary center experience**

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**Pediatric Rheumatology 2017, 15(Suppl 1) P141**

**Introduction:** Polyarteritis nodosa (PAN) is a necrotizing vasculitis of the wall of small and medium-sized arteries. Cutaneous Polyarteritis nodosa (cPAN) is the most common in pediatric population. Its etiology is unknown, but the association with streptococcal infection has been described, which suggests the indication of antimicrobial prophylaxis similar to that made in rheumatic fever. There are few studies addressing this, what motivated this study.

**Objectives:** Aim to evaluate the association between cPAN and streptococcal infection.

**Methods:** Retrospective review of medical records of patients who met the diagnostic criteria for PAN between 1987 and June 2015. The evidence of streptococcal infection was evaluated by the increase in the antibody anti streptolysin O (ASO) at diagnosis and clinical relapse PAN.

**Results:** Fifteen patients were evaluated in this period. Two children were excluded because they did not have investigation about streptococcal infection. Thirteen patients had PAN diagnosis, three (23%) with systemic and 10 with cPAN (77%). Three patients out of four (75%) had clinical recurrence with streptococcal evidence. The number of reported cases in our service was similar to that found in the literature.

**Conclusion:** The evidence of streptococcal infection was found in patients with cPAN at diagnosis and relapse. If the association between streptococcal infection and patients with cPAN is confirmed, prophylaxis is suggested.

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**Disclosure of Interest**

None Declared

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**P142**

**Abciximab for treatment of coronary aneurysms in Kawasaki disease**

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**Pediatric Rheumatology 2017, 15(Suppl 1) P142**

**Introduction:** Kawasaki disease (KD) is a childhood acute vasculitis. The development of coronary damage is the main long-term consequence and can lead to myocardial infarction or sudden death. Reducing the incidence of coronary artery aneurysms (CAA) is the treatment cornerstone. Abciximab, a glycoprotein IIb/IIIa receptor antagonist, is indicated in acute coronary syndrome in adults. Some reports suggest a benefit in KD.

**Objectives:** To review the experience at our center in treating CAA with abciximab in patients with KD and determine its efficacy in reducing the CAA size.

**Methods:** Retrospective observational study describing all patients with KD and coronary involvement treated with abciximab in our center. Coronary maximum diameter is expressed in Body Surface Area (BSA)-adjusted Z-score. The coronary outcome is described and safety of treatment evaluated.

**Results:** Five patients have been treated with abciximab since 2012, all males with mean age of 11.5 ± 7.1 months. The left main coronary artery (LMCA) was constantly affected, the right coronary artery (RCA) in three patients and the left anterior descending (LAD) in two. The mean adjusted Z-score for the affected artery at KD diagnosis and at last echocardiography were 9.8 ± 3.3 and 3.5 ± 4.9, respectively; what represents a mean decrease of 5.4 ± 3.45 SD (mean follow-up of 688 ± 369 days). Although an observed tendency to CAA reduction, statistical significance was not reached. The drug infusion was well-tolerated and no adverse effects were observed.

**Conclusion:** Although current data suggest a long-term benefit, further evidence about the effectiveness of abciximab in the treatment of CAA in patients with KD is needed. Abciximab seems a safe treatment in pediatric patients with KD.

**Disclosure of Interest**

None Declared

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**P143**

**Cogan syndrome presented in 5 years old girl with history of hypoglycaemic episodes and with significant behavioral changes**

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**Pediatric Rheumatology 2017, 15(Suppl 1) P143**

**Introduction:** Cogan syndrome (CS) is a rare autoimmune vasculitis affecting vision by non-syphilitic interstitial keratitis and hearing by sensorineural hearing loss, tinnitus, and vertigo. Typical CS affects cornea and inner ear. However, the “atypical” variants of CS can involve kidneys, CNS and other eye structures.

**Objectives:** A case report of a young girl with the background of episodes of ketotic hypoglycaemia, who then developed typical features of CS with significant changes and sleep disturbance, is presented.

**Methods:** A 5 years old, Caucasian girl presented with a four week history of being unwell, red eyes, vomiting, reduced appetite and behavioural changes including sleep disturbance and keeping head tilted to the left side. She was examined by an ophthalmology review due to persistence of red eyes, CS was considered as a differential diagnosis due to bilateral inferior corneal epithelopathy. Diagnosis of CS was confirmed by an MRI of head showing an enhancement of the inner ear.
sensitivity of four criteria for Behçet’s disease in Japanese pediatric patients

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Pediatric Rheumatology 2017, 15(Suppl 1):P144

Introduction: Three criteria, International Study Group (ISG), International Criteria for Behçet’s Disease (ICBD) and Japanese Ministry of Health Labour and Welfare (MHLW), had been used in practice for pediatric Behçet’s disease (BD) in Japan. Sensitivity of these criteria were not high enough for pediatric patients, as they were made for adult patients.

Objectives: Paediatric Behçet’s Disease study (PEDBD) criteria was established and reported that it had high sensitivity for pediatric BD patients from 27 countries in 2015. We used PEDBD criteria for Japanese pediatric BD patients, and confirmed the sensitivity to compare with other criteria.

Methods: We conducted a questionnaire survey of pediatric rheumatologists about clinical symptoms and laboratory results of pediatric BD patients in Japan. Next, we examined these patients whether they meet each criteria to know the sensitivity. Finally, we compare the sensitivity of four criteria in our patients.

Results: Fifty-two patients of BD were collected, 0-15 years of age of onset. The sensitivity of each criteria became as follows, ISG criteria 70%, ICBD criteria 90%, MHLW criteria 60% and PEDBD criteria 46%.

Conclusion: The sensitivity of these criteria seemed to be relatively low, especially the one of PEDBD criteria were lower than we expected. This results may mean that we should take into account the importance of geographical variation in pediatric BD.

Disclosure of Interest
None Declared

P145

Increased serum antibody titer against HPV-16 antigen in patients with Behçet’s disease

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Pediatric Rheumatology 2017, 15(Suppl 1):P145

Introduction: Quadrivalent Human Papillomavirus (HPV) vaccine has been reported to be significantly associated with Behçet’s disease (BD). However, no reports have described HPV infection as a possible cause for the development of BD.

Objectives: The objective of this study was to evaluate whether anti-HPV antibody titer is increased in BD.

Methods: Sera from 93 Korean BD patients, who fulfilled the diagnostic criteria of the International Study Group for BD, were used in an enzyme-linked immunosorbent assay (ELISA). The clinical activity of BD was evaluated at the time of blood sampling. HPV-16 L1 virus-like particle (VLP) antigen was used in this study for the ELISA.

Results: Patients with BD had significantly higher antibody titers against HPV-16 (optical density (OD): 0.210-3.675; mean 0.992) than that of healthy controls (OD: 0.248-0.762; mean 0.517; p < 0.001). Using a receiver operating characteristic (ROC) analysis, a cut-off value of 0.578 OD for the anti-HPV antibody titer was determined that differentiated BD patients from healthy controls. When we compared the clinical features of BD between the two groups, articular involvement of BD was more likely in patients with an anti-HPV-16 antibody titer <0.578 OD (p = 0.035). In addition, patients with an anti-HPV-16 antibody titer <0.578 were significantly younger than patients with a titer ≥0.578 OD.

Conclusion: HPV itself may be a possible extrinsic triggering infectious agent causing the development of BD.

Trial registration identifying number: This research was supported by the Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education (2014R1A1A2059503).

Disclosure of Interest
None Declared

P146

A case of Churg-Strauss syndrome in a 12-year-old girl

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Pediatric Rheumatology 2017, 15(Suppl 1):P146

Introduction: Churg-Strauss syndrome, also known as allergic granulomatous angiitis, is extremely rare in children and is associated with higher cardiopulmonary disease and mortality rates compared to adult patients. Here, we report a twelve year-old girl with a diagnosis of CSS, who presented with prominent pulmonary involvement accompanied by eosinophilia, two-year history of asthma, peripheral neuropathy but without anti-neutrophil cytoplasmic antibody (ANCA).

Objectives: A twelve year-old girl presented to our clinic with a two months history of periodic fever, weight loss, productive cough, breathlessness, arthralgia, myalgia, muscle weakness, paresthesias of right extremities. She has a two-year history of asthma, sinusitis, urticaria, allergic rhinitis, gastrointestinal bleeding. On examination she had skin nodules on her extremities and purpuric lesions on her uncles.

Methods: We present a case report of a 12 year old girl with Churg-Strauss syndrome, who were treated in a Pediatric Rheumatology outpatient center.

Results: Blood tests revealed eosinophilia, ESP-61 mm/h, CRP-192 mg/L, Chest-x ray and CT demonstrated cardiomegaly, pulmonary infiltrates in right lung and bilateral pleural effusion. Echocardiogram
showed aortic and mitral valve insufficiency, EF-31%, bronchoscopy demonstrated 15% of eosinophils on BALF. Skin biopsy demonstrated necrotizing granulomatous inflammation with eosinophils. Electromyography has shown a gross violation of n. peroneus. She was treated with prednisolone (2 mg/kg - reducing with appropriate scheme) with good clinical response. ESP, CRP and eosinophil levels were normalized, but cardiopulmonary involvement sustained. Cyclophosphamide was not used (parents decision).

**Conclusion:** Presenting case demonstrates the importance of considering the diagnosis of CSS in children with eosinophilia, uncontrolled asthma, sinusitis, skin lesions and cardiomyopathy. Sever organ involvement and delayed diagnosis may worsen prognosis and even lead to a fatal outcome. If started early, treatment with corticosteroids is often successful.

**Disclosure of Interest**
None Declared

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**P147 Successful treatment with etanercept in a patient with refractory cutaneous polyarteritis nodosa**

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*Pediatric Rheumatology* 2017, 15(Suppl 1)P147

**Introduction:** Cutaneous polyarteritis nodosa (cPAN) is a form of necrotizing vasculitis of small and medium sized arteries without visceral involvement. cPAN has often relapsing and chronic course. Some patients with cPAN are resistant to various immunosuppressive treatments and have repeated exacerbations over prolonged periods.

**Objectives:** To report the effectiveness of tumor necrosis factor alpha (TNF-α) blockade as a therapeutic alternative for refractory cPAN.

**Methods:** We report the patient of cPAN refractory to various immunosuppressive treatments and tonsillectomy, who was successfully treated with etanercept (ETA). We also present a review of 3 similar cases of cPAN reported in the literature.

**Results:** Case presentation: An 8-year-old girl was referred to us with fever, erythematous papules and painful subcutaneous nodules located over her extremities and arthritis of bilateral ankles and knees. Laboratory findings revealed leukocytosis (134×10^3/mm^3), elevated levels of CRP (1.4 mg/dl) and ESR (61 mm/hr). Antistreptolysin O (ASLO) titer was found to increase (2380 IU/ml). Magnetic resonance image of the thighs revealed multiple high signal nodular lesions throughout the subcutaneous tissue and subfascial muscle tissue of the thighs. A skin biopsy of a subcutaneous nodule revealed a necrotizing vasculitis. The diagnosis of cutaneous polyarteritis nodosa (cPAN) was made. She was treated with prednisolone and maintained on cyclosporine and monthly infusions of cyclophosphamide. However, flare of cPAN with fever, myalgia and painful subcutaneous nodules occurred 5 times with steroid withdrawal. Tonsillectomy was performed but flare occurred again with steroid withdrawal. ETA (12.5 mg 2 times/week) was started and led to a marked improvement of her symptoms within 3 weeks. The patient presented complete clinical remission of cPAN and continues to receive ETA and cyclosporine.

**Review of the literature:** Three cases of cPAN successfully treated with TNF-α blockade have been reported. Two cases were treated with ETA and one patient was treated with infliximab. ETA and infliximab attenuated their symptoms and showed a corticosteroid sparing effect and no adverse effects.

**Conclusion:** The significant improvement observed in this patient. ETA could be an effective alternative treatment for cPAN. Further larger studies is desired to establish the effectiveness of ETA as therapeutic alternatives for cPAN.

**Disclosure of Interest**
None Declared

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**P148 Complicated schoenlein-henoch with severe organ involvement: personal records**

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*Pediatric Rheumatology* 2017, 15(Suppl 1)P148

**Introduction:** Henoch-Schönlein purpura (HSP) is the most common childhood systemic small-vessel vasculitis with skin, joint, gastrointestinal and renal involvement. Uncommon gastrointestinal complications are: intussusception, bowel perforation and rarely appendicitis. Other rare complications are: ureteritis, orchitis, pancreatitis, severe intestinal bleeding.

**Objectives:** We evaluated complicated HSP children and followed them until remission.

**Methods:** In the last 3 years we followed in our Paediatric Unit 90 children (age: 4-14 years) with HSP, presenting with classical signs (cutaneous vasculitis, arthralgia and/or arthritis of limbs, abdominal pain, without a complicated picture of intussusception and/or perforation and/or severe bleeding. Among these children 10/90 patients (11%) (age: 5-10 years) showed a complicated HSP with a recurrent form and an organ involvement.

**Results:** 7/10 patients showed an intestinal involvement with severe abdominal pain and persistent bleeding; 4 patients had severe and prolonged intestinal bleeding, resistant to steroids. 2/7 patients developed a persistent nephritis with proteinuria and hypertension, One patient with intestinal bleeding developed a pancreatitis with persistent and significant amylase and lipase increase; 4 patients had orchitis. All the patients were treated with methylprednisolone (1-2 mg/kg/day), with a mild tapering of steroids. In the 2 children with nephritis amlodipine was associated. All the patients had a slow and progressive ameliorating outcome, with a complete remission.

**Conclusion:** Most of the children affected by HSP have an excellent prognosis, however a few cases develop a complicated form, which need steroid and/or a combined treatment with anti-hypertensive and immunosuppressive drugs.

In our population, 8% showed a complicated form, however all the patients did not show long-term sequelae.

**Disclosure of Interest**
None Declared

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**P149 Incidence of IgA Vasculitis in children: A population-based study using a four-source capture recapture estimate**

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*Pediatric Rheumatology* 2017, 15(Suppl 1)P149
Introduction: The etiology of Immunoglobulin A vasculitis (IgAV), the most common childhood vasculitis in western countries, remains unclear. Changes over time in its epidemiology could guide in the search of a causal pathway.

Objectives: We aimed to describe the epidemiological characteristics of childhood IgAV in a large French population.

Methods: Cases were prospectively collected in Val de Marne county, a southeastern suburb of Paris, with 263,874 residents <15 years old. Children with incident IgAV living in this area from 2012 to 2014 were identified by 4 sources of case notification (emergency departments, pediatrics departments, private practice pediatricians, general practitioners). We calculated 95% confidence intervals (95% CIs) for incidence rates under the assumption of a Poisson distribution. Four-source capture-recapture analysis was performed using log linear modeling to estimate the completeness of case finding. Socio-professional factors, ethnicity, seasonal and geographic variations in IgAV incidence were investigated.

Results: The survey identified 147 incident cases meeting the study criteria, including 78 (53%) boys and 69 (47%) girls with a mean age of 6.5 years (2.2–14.7 years). The overall annual incidence (per 100,000 children) was 18.6 (95% CI 13.6–24.5), with 19.3 (12.6–28.3) for boys and 17.8 (12.8–26.7) for girls. The capture recapture analysis estimate the completeness of case finding at 62% (95% CI 50–81) with an annual incidence of 29.9 (95% CI 23.7–37.3). The annual distribution of diagnoses significantly deviated from a random distribution and indicated a nadir in summer months. Neither socioeconomic factors nor environmental factors like industrialization or pollution appear to play a role.

Conclusion: Our annual incidence rate of 18.6 to 29.9 per 100,000 children, gender distribution and seasonal variation are in line with data reported worldwide and show no significant periodic or secular variation over time. IgAV seems to be trigger by an indeterminate infection shared by children in community. The impact of genetic, as predisposing factor, need to be more explore in particular the role of innate immunity.

Disclosure of Interest
None Declared

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P150
Severe systemic vasculitis associated with primary immunodeficiency in two siblings
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Pediatric Rheumatology 2017, 15(Suppl 1) P150

Introduction: In paediatric rheumatology practice, systemic vasculitides associated with primary immunodeficiency disorders are highly challenging conditions.

Objectives: The objective of this presentation is to report the clinical course and response to treatment in two sisters suffering from systemic vasculitis associated with unclassified hypogammaglobulinemia.

Methods: We retrospectively evaluated the medical records of both sisters.

Results: The first case was a girl, born from non-consanguineous parents. At the age of 1.66 years old, she suddenly presented high fever, non-specific maculopapular rash, atheralgia, myalgia with elevated levels of acute phase reactants and prompt response to steroids. One year later, the above manifestations relapsed. A skin biopsy showed interstitial granulomatous dermatitis in association with vasculopathy. At 3 years old she developed recurrent cranial nerve palsies and livedo reticularis and then two stroke events at the age of 4 (one ischemic and the other hemorrhagic). Magnetic resonance angiography showed no evidence of cerebral vasculitis and the biopsy of the brain performed as part of the surgical procedure for the hemorrhagic stroke showed only extravasation of erythrocytes and fibrin deposition around small vessels, without significant inflammation. The antibody profile (including antiphospholipid antibodies) and clotting tests were negative. Hypogammaglobulinemia with global Ig deficiency was present from the onset of the disease and persistent in time. She could not be considered common variable immunodeficiency disorder because total serum IgG levels were not constantly reduced below 500 mg/dl. As part of a research study, she was also diagnosed with complete C4B deficiency. At the age of 5 years old she developed moderate thrombocytopenia which persisted for the next 3 years. At 5.5 years old she was also diagnosed with nodular regenerative hyperplasia (NRF) of the liver. For the first 2 years of evolution, only high dose of corticosteroids was able to control the disease manifestations. Methotrexate, azathioprine, cyclosporine, cyclophosphamide were ineffective. Conversely, mycophenolate mofetil (MMF) associated with high monthly doses of IGIV provided prompt and persistent improvement of clinical symptoms and acute phase reactants and also tapering of steroid dose. Till now, under MMF, no recurrences have been observed, but the NRH was slowly progressive. No portal hypertension has been developed so far.

Case 2, the older sister of the above patient, had the first disease manifestations at 6.8 years old. Since then, she developed the followings clinical picture: fever, atheralgia, painful subcutaneous nodules, livedo reticularis, low serum IgM and IgG4, elevated acute phase reactants, negative antibody profile, recurrent abdominal pain, systemic arterial hypertension, one episode of massive intestinal bleeding which had no obvious source at total colonoscopy, upper endoscopy and computed tomography angiography. The biopsy of a subcutaneous nodule revealed leukocytoclastic vasculitis. In her case, MMF induced only a partial remission. The association of tacrolimus permitted rapid control of the disease activity. Up to now, no neurological signs were present.

The recently described genetic disorder “deficiency of ADA2” is being considered in both sisters, but yet unproven.

Conclusion:
Aggressive medical management is often necessary and therefore the rate of drug related morbidity in high. The response to the various treatments is variable, even between siblings.

Disclosure of Interest
None Declared

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P151
Henoch-Schönlein purpura, a general view about the more often vasculitis in pediatric population and a study of forecast factors of renal disease relapse in Mexican population
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Pediatric Rheumatology 2017, 15(Suppl 1) P151

Introduction: Henoch-Schönlein purpura is the more often vasculitis in pediatric population, it is characterized for being a small vessel leukocytoclastic vasculitis; it has an incidence between 10 to 20.4 cases per 100,000 children, with a predominance of male sex 2:1. Etiology remains unknown. Given the tendency for spontaneous resolution of disease treatment in most cases will be supportive. There is no evidence of the benefits of renal disease treatment, mainly because few patients have severe disease at baseline.

Objectives: Identify factors of poor renal prognosis in mexican children with Henoch-Schönlein purpura diagnosis.

Methods: A transversal comparative study in a cohort of 79 patients diagnosed with HPS over a 5 years period in Hospital Infantil de Mexico, Federico Gomez. We realized parametric, descriptive, comparative, correlation and likehood ratio test, were performed.
Results: In this study, they were evaluated a total of 79 files of patients with this diagnosis; 47 female patients and 32 male patients, with a 1.6:1 relation women dominate; the mean age presentation was 82 months, predominating in autum with 34.7% and spring with 27%. 100% of patients debuted with characteristic purpuric rash, gastrointestinal symptoms were presented in 35%, 76% presented joint disease during the disease onset; as to kidney disease nephritis was found during the debut in 19% (n = 15) and elevated creatinine in 5.1% (n = 4), to which were subsequently performed renal biopsy, histological lesion more frequently found was mesangial proliferation glomerulonephritis in less than 50% of glomeruli (grade IIIa) that was found in 4 of 4 patients, currently a patient is on replacement therapy with hemodialysis, with an average of 0.5 mg / dL ± 0.11 with a maximum of 13 mg found / dL.

Conclusion: Statistical correlation tests of our study are consistent with other studies published in the literature, finding a statistically significant correlation between the presence of elevated at diagnosis and relapse with p = 0.011 creatinine. As prognostic factors for relapse reported in the literature we found that the presence of initial nephritis is a prognostic factor p ≤ 0.001 relapse and complications with a value of p = 0.025. In that order we can extrapolate the results of international studies to our population.

Disclosure of Interest
None Declared

Table 43 (abstract 151). Differences between patients with and without nephritis at the diagnosis of PHS

| Characteristics                  | All patients | Patients with Nephritis | Patients without Nephritis | Value of P  |
|----------------------------------|--------------|-------------------------|---------------------------|------------|
|                                  | N = 79       | n = 17                  | n = 62                    | ≤0.001     |
| Elevated creatinine              | Yes 5%       | 22.2%                   | 1.4%                      | ≤0.001     |
|                                  | No 95%       | 77.8%                   | 98.6%                     |            |
| 12 hrs proteinuria mg/m2/hra     | 4.57 ± 7.65  | 10.47 ± 13              | 2.9 ± 2.3                 | ≤0.001     |
| Proteinuria                      | Yes 63%      | 88%                     | 23%                       | ≤0.001     |
|                                  | No 37%       | 12%                     | 76%                       |            |
| Relapse                          | Yes 19.5%    | 50%                     | 11.6%                     | ≤0.001     |
|                                  | No 80.5%     | 50%                     | 88%                       |            |

P152

Prevalence of kidney involvement during the first follow-up year in patients with Henoch–Schönlein Purpura in a pediatric institution in Bogota, Colombia

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Pediatric Rheumatology 2017, 15(Suppl 1)P152

Introduction: The Henoch-Schönlein Purpura (HSP) is the most common vasculitis in children, mainly involving the skin vessels, joints, gastrointestinal tract and kidneys. Kidney involvement remains the main cause of morbidity and mortality in children. Nephritis occurs in 30 – 50% of patients during the first 4 to 6 weeks, but with a risk of appearance during the first 5 years. The spectrum of kidney involvement ranges from mild manifestations until the development of a nephrotic and/or nephritic syndrome or kidney failure. Some predicting risk factors have been identified for the development of nephritis: persistent or recurring purpura, severe abdominal symptoms, older age at debut and disease relapse. Renal prognosis has been associated with kidney biopsy findings. The aim of the study is to determine the clinical features and kidney involvement during the first year of follow-up.

Objectives: Determine the clinic features at debut and kidney involvement of patients with a final diagnosis of HSP during the first month, 3 months and until the first year of follow-up.

Methods: Retrospective study of patients with a final diagnosis of HSP in a Pediatric Rheumatology service. A single form of data collection was applied in an institution of Bogota, Colombia during the period comprised between 2010 and 2016

Results: N = 86 patients, 42 girls and 44 boys. Median age at disease onset was 5.3 years SD 2.4 years (range 1 - 14 years). At HSP diagnosis all the patients had palpable purpura,70.9% abdominal pain and 88.4% arthritis. Kidney involvement was present in 39/86 patients (45%), 22/39 (56%) patients had renal involvement at diagnosis: Isolated proteinuria was the most frequent finding 10/22 (45.4%) and 3 patients had nephrotic syndrome.12 patients (31%) had kidney involvement within the first month (50% had isolated proteinuria and 50% proteinuria/hematuria). Between the first and third month 2 (5%) patients had renal manifestations (all isolated proteinuria). Between 3 months and 12 months follow-up 3 patients (8%) had kidney compromise: one hematuria, one isolated proteinuria and one hematuria/proteinuria. A trend was observed to kidney involvement in patients with abdominal symptoms (p = 0.053). Renal biopsy was performed in 8/39 patients with HSP nephritis. Mesangial proliferation was the most common finding in 6/8 patients, sclerosis in 3/8 patients (1 patient with nephrotic syndrome and 2 patients with proteinuria/hematuria). There was no evidence of crescents or新 blood vessels in any of the biopsies. The mean follow-up was 26.8 months SD 17 months (range 1 - 72); there was no evidence of kidney damage in the last assessment in any of the 39 patients with kidney involvement.

Conclusion: In this group of patients, kidney involvement was more severe and common in the first weeks of the disease onset. The active search and education of families during the first 4 weeks of HSP onset is essential. In severe forms, it is important to conduct a kidney biopsy in order to document poor prognostic factors and adjust treatment.

Disclosure of Interest
None Declared
**P154**

**QT dispersion in children with Kawasaki disease with transient coronary artery abnormalities: a follow-up study**

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Pediatric Rheumatology 2017, 15(Suppl 1):P154

**Introduction:** Majority of patients with Kawasaki disease (KD) are believed to have subclinical myocardial involvement that predispose to an increased risk of developing ventricular arrhythmia resulting from inhomogeneous ventricular repolarization. This inhomogeneous ventricular repolarization manifests as increased QT dispersion on electrocardiography. Though there are a few studies on QTd in patients with KD and persistent coronary artery abnormalities, there is paucity of literature on QTd in children with KD and transient coronary artery abnormalities.

**Objectives:** The present study was undertaken to study QT dispersion in children with KD and transient coronary artery abnormalities during follow-up.

**Methods:** Twenty children with KD and transient coronary artery dilatation that resolved within 1 year of diagnosis (mean age 11.5 ± 3.7 years; range between 5.1 to 19.6 years) were studied at least 1 year after resolution of coronary artery abnormalities. Diagnosis of KD was based on standard guidelines of the American Heart Association. As per unit protocol, treatment included intravenous immunoglobulin (IVig) infusion (2 g/kg) along with aspirin (30-50 mg/kg/day during acute stage followed by 3-5 mg/kg/day for 4-6 weeks or until resolution of CAA, whichever was later). Mean interval between onset of KD and IVig infusion was 11.05 ± 6.24 days. 20 healthy controls, matched for age and sex, were also enrolled. 12 lead ECG was done in all cases and controls. A single blinded observer used the digital caliper to measure the intervals with least count of 0.01 mm. The difference between the maximum and minimum corrected QTc intervals was calculated as the QTc dispersion. As only two of the six extremity leads are recorded and the remaining 4 leads are mathematically derived, QTc dispersion was also calculated in 8 leads. These included the 6 precordial leads, shortest extremity lead, and median of other 5 extremity leads.

| Cases (Pre-IVig) | Cases (Post-IVig) | Febrile controls | Controls |
|------------------|------------------|-----------------|----------|
| Th17 cells       | 2.58%            | 1.88%           | 1.57%    | 2%       |
| Treg cells       | 0.99%            | 2.64%           | 1.96%    | 2.37%    |

**Conclusion:** QT dispersion in children with KD and transient coronary artery abnormalities was not found to be significantly affected in long term follow up.

**Disclosure of Interest**

None Declared

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**P155**

**Infliximab treatment for a patient with giant coronary aneurysm with refractory Kawasaki disease: 8-month old male. Case report and brief review of literature**

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This abstract is not included here as it has already been published.

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**P156**

**Fibromuscular dysplasia in the practice of child rheumatologist**

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Pediatric Rheumatology 2017, 15(Suppl 1):P156

**Introduction:** Fibromuscular dysplasia (FMD) – hyperplastic disease, is caused by a mutation in the gene of type III collagen (133580, 2q31, COL3A1), characterized by lesions of the kidney, at least - carotid and other major arteries. According to the literature the FMD of renal arteries is second (10-15%) in the structure of renovascular hypertension (RVG). In some cases, clinically FMD may be asymptomatic. However, the possibility of RVG, the instrumental detection of the deformation of large arteries is the reason for the conversion to a rheumatologist and a cardiologist with the assumption TA the patient.

**Objectives:** Common to FMD and TA are gross vascular murmurs in the projection of the large arteries, the possibility of VAG,

labeled anti-CD4, Phycerothrin (PE) labeled anti-IL-17 and AlexaFluor labeled anti-Foxp3 antibody

**Results:** Male: female ratio amongst cases was 4:1. Mean age at onset of KD was 4.45 years and the youngest child was 6 months old. Mean duration of fever at presentation was 10.7 days and 60% of children presented beyond 10 days of onset of fever. Conjunctival injection and oral mucosal changes were noted in 50% and 70% respectively while cervical lymphadenopathy and rash each occurred in 60% of our cohort. Extremity edema was noted in 50% while periangual/perineal peeling of skin was noted in all. 40% of the children had incomplete KD. Coronary artery abnormalities (CAA): coronary dilatation was noted in 3 cases at admission (left anterior descending artery in 2, left main artery in 1), which resolved at follow-up of 6 weeks. Two amongst these were subjected to CT- coronary angiography during follow up, which were normal.

Proportions of Th17 and Treg cells in cases and controls have been tabulated below. Th17 cells were increased in acute phase of KD as compared to febrile controls (2.58% vs 1.57%, P = 0.27) and healthy controls (2.58% vs 2%, P = 0.21). Treg cells were decreased in children with KD during acute phase as compared to febrile controls (0.99% vs 1.96%, P = 0.62) and healthy children (0.99% vs 2.37%, P = 0.45). Following treatment with IVig, Th17 cells decreased (2.58% vs 1.88%, P = 0.13) and Treg cells showed an increase (0.99% vs 2.64%, P = 0.07), however, this difference was not statistically significant. No significant difference was noted in Th17 and Treg % between cases who had CAA and those who did not.

**Conclusion:** Th17 cells were elevated and Tregs were reduced during acute KD as compared to febrile and healthy controls; the difference, however, was not statistically significant. A larger study to assess the same differences is recommended to confirm the differences of statistical significance.

**Disclosure of Interest**

None Declared
deformation and thickening of the large arteries, detected by instrumental research. In contrast to TA, while no FMD laboratory signs of inflammatory activity. Angiography can reveal a kind of defeat of large arteries throughout the “string of pearls”, “rosary” - thick, hardened areas of the arteries with hyperplasia interspersed with less altered areas, while at TA, affects primarily the mouth of the large arteries and possibly the aorta.

Methods: We observed 7 children with FMD - 4 girls and 3 boys ranging in age from 11 to 17 years (mean age at the time of detection of arterial hypertension 13.5 years - 13.4 ± 1.05). Compared the results of clinical, laboratory and instrumental (Doppler ultrasound, MRI, angiography) examinations in 6 patients with FMD and 38 with a documented diagnosis of TA.

Results: at the onset in all children with suspected FMD TA in identifying RVG, not amenable to standard anti-hypertensive therapy, the presence of vascular noise in the projection of the large arteries and the data of instrumental examination (Doppler ultrasound - thickening of artery walls and increased linear velocity of blood flow). However, all patients were not inflammatory syndrome at the beginning, typical for the acute phase of TA and laboratory signs of activity throughout the observation period. For differential diagnosis, along with named above instrumental methods of examination, and in 4 children was used positron emission tomography, allowing to estimate objectively the absence of active inflammatory process in the walls of arteries. Surgical treatment was performed 5 of the 7 patients with FMD. In 2 patients, in marked stenosis of the renal arteries, attempt balloon angioplasty did not lead to the desired effect. Only autovenous bilateral aortorenal prosthesis has allowed everyone to stop RVG.

Conclusion: Deformation of the aorta and its branches with vascular dysplasia leads to difficulties of differential diagnosis with nonspecific aortoarteritis. The lack of inflammatory syndrome at disease onset and laboratory signs of activity during the whole follow-up period in the presented clinical case are not allowed to confirm the diagnosis of Takayasu arteritis. Revealing as a new diagnostic method of research is positron emission tomography, allowing to estimate objectively the absence of active inflammatory process in the walls of arteries.

Disclosure of Interest
None Declared

P157
Is neutrophil to lymphocyte ratio valid to predict organ involvement in Henoch-Schönlein Purpura?
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P158
Childhood polyarteritis nodosa: diagnosis with noninvasive imaging techniques
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Pediatric Rheumatology 2017, 15(Suppl 1):P158

Introduction: Childhood polyarteritis nodosa (cPAN) is a systemic inflammatory disease characterized by histopathological evidence of necrotizing vasculitis or angiographic abnormalities. The classification criteria for cPAN involve the angiographic detection of visceral aneurysms and conventional angiography remains the overall gold standard. There are limited numbers of publications about the use of noninvasive imaging modalities in the diagnosis of cPAN.

Objectives: The aim of this study was to present the clinical and imaging findings of the patients with cPAN who were diagnosed with non-invasive imaging techniques.

Methods: Files of patients who had been diagnosed as cPAN in our department from 2005 to 2015 were evaluated, retrospectively. Demographic, clinical, laboratory and imaging findings of the patients were evaluated.

Results: Nine patients (8 M, 1 F; age at disease onset: 12,5 years (7-16)) had been diagnosed as cPAN in our clinic with non-invasive imaging techniques within the last 10 years. Abdominal pain, fever, fatigue and myalgia were the most frequent complaints. Doppler ultrasonography (US) was used in the diagnosis of 7 patients and computerized tomography (CT) angiography was done in 4 patients. Duration between admission to our center and diagnosis was median of 5 days, including 4 patients who were diagnosed within 24 hours of admission. Approximately 80% of our patients with cPAN had MEFV mutations and 90% had elevated ASO levels. All of them had the involvement of the gastrointestinal tract. Hepatic and cystic arterial involvements were detected in 7 and 6 patients, respectively.

Conclusion: This report included the largest cPAN series that were diagnosed with noninvasive imaging modalities. We suggest that noninvasive modalities, especially Doppler US should be considered in first line approach in the diagnosis of these patients, particularly in children.

Disclosure of Interest
None Declared

Poster Session: Systemic JIA I

P159
Factors associated with etoposide usage in children with macrophage activation syndrome complicating systemic Juvenile Idiopathic Arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1):P159

Introduction: Although macrophage activation syndrome (MAS) has been reported in association with almost all rheumatic diseases, it is by far most common in systemic juvenile idiopathic arthritis (sJIA). Reported mortality rates in MAS reach 10-20%. Standardized treatment guidelines for MAS are currently lacking, but management commonly includes high-dose corticosteroids combined with another immunosuppressive agent. Etoposide is a well established therapy in primary hemophagocytic lymphohistiocytosis. A recent systematic literature review on treatments of MAS in sJIA identified only a few patients who were given etoposide.

Objectives: We aimed to investigate etoposide usage among pediatric patients with MAS complicating sJIA and to identify predictors of etoposide administration.

Methods: Retrospective collected data from 362 patients included in the multinational study of the 2016 classification criteria for MAS in sJIA were examined to identify patients treated with etoposide and record potential predictors of etoposide administration. Variables
significantly associated with etoposide usage in univariate analysis were entered in a multivariate regression model. Continuous variables were dichotomized according to the cut-off value obtained through ROC curve analysis.

Results: Forty of the 362 patients (11%) were treated with etopo- side and 17 of those had information on all variables studied. Factors significantly associated with etoposide administration in multivariate analysis included multiorgan failure (OR 7.9, 95% CI 2.2-28.5), platelet count ≤ 132 x 109/liter (OR 5.8, 95% CI 1.8-18.2), triglycerides > 270.8 mg/dl (OR 3.7, 95% CI 1.3-10.4), aspartate aminotransferase > 389 units/liter (OR 3.7, 95% CI 1.3-10.3), and fibrinogen ≤ 1.53 g/ml (OR 2.9, 95% CI 1.1-7.5). The AUC of the model was 0.86. In univariate analysis, there was no significant difference in mortality rate between patients given or not given etoposide.

Conclusion: Patients treated with etoposide were sicker than patients who did not receive this medication. However, mortality rate did not differ between the two treatment groups, suggesting that etoposide may be an indication for aggressive therapeutic interventions for severely ill children with MAS.

Disclosure of Interest
None Declared

P161

The systemic onset Juvenile Idiopathic Arthritis national registry in Turkey: a preliminary report

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Pediatric Rheumatology 2017, 15(Suppl 1):P161

Introduction: Systemic-onset juvenile idiopathic arthritis (SJIA) is characterised by spiking fever, evanescent rash, hepatosplenomegaly, serositis, lymphadenopathy and arthritis.

Objectives: The aim of this study was to evaluate the clinical and laboratory findings and the monitoring treatment outcome of patients with systemic juvenile idiopathic arthritis (SJIA) in the Turkey.

Methods: Through the nationwide registry of Turkey, 167 patients with SJIA based on ILAR criteria were recruited between January 2000 and December 2015 at nine pediatric rheumatology centers. We evaluated the demographic characteristics, and laboratory and clinical data at the time of SJIA diagnosis. Evaluation of medications included previous and concomitant treatments.

Results: One hundred sixty-six SJIA patients (92 females (55.4%), 74 males (44.6%) were included in this analysis. The mean age of disease onset and diagnosis was 6.15 ± 4.45 and 6.48 ± 4.53 years, respectively. The mean duration in months from diagnosis to study enrolment was 51.7 ± 42.4 for the cohort.

Macrophage Activation Syndrome (MAS) occurred in 22% of patients (n = 35) at presentation or during the disease course. The all patients was distributed according to subtypes: mononuclear 42.3%, polynuclear 33.7% and persistent/polyarticular 24%.

During follow-up, 95% of all patients with JA had received corticosteroid (methyl prednisolone or prednisolone), 64% received a conventional synthetic disease-modifying antirheumatic drug (csDMARD), and 48% received a biologic DMARD (bDMARD). The median time from diagnosis to the initiation of a biologic therapy (BT) was 8.5 months (min-max 0–11). The first BA was anakinra in 41 patients, etanercept in 24, tocilizumab in 9, canakinumab in 3, and adalimumab and infliximab in 1 patient each. Median follow-up duration on BT was 24 months (range 0.5–73) for the patients who did not experience a switch of BT; for switchers, median follow-up duration on BA was 6 months (range 0.1–62.0).

Switching of BT was common, with 42 patients (52.5%) switching to a second, 12 (15%) to a third and 3 (6.2%) to a fourth BA. Among 41 patients on anakinra as a first BT, 19 (46.3%) were switched to a new BT. Out of 24 patients on ETA as a first BT, 17 (71%) switched.

Inactive disease was achieved in 143 patients (91.1%), 87% of these patients treated with BT. 18 (12.6%) of the patients who achieved inactive disease fulfilled the criteria of clinical remission on medication at last follow-up.

No case of cancer was recorded. Four patients (2.4%) had amyloidosis and one patient was died. Four cases of MAS were documented.

Conclusion: This study reflects preliminary report of SJIA multicenter, national registry in Turkey. The study is still ongoing, it is planned to increase the number of patients with the participation of other centers.

Disclosure of Interest
None Declared

P160

A novel homozygous frameshifting mutation in LACC1 associated with severe familial forms of Juvenile Idiopathic Arthritis

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Introduction: The pathophysiological origin of juvenile idiopathic arthritis (JIA) is largely unknown. However, recently a homozygous missense mutation c.850T>C in LACC1 has been reported in several patients with therapy-refractory systemic JIA (sJIA) from consanguin- eous families in Saudi Arabia.

Objectives: We here describe two siblings from a large consanguin- eous Lebanese family with severe JIA harboring a novel homozygous LACC1 mutation.

Methods: Whole-exome sequencing (WES) was performed on DNA sam- ples of two index patients with severe forms of sJIA and both healthy parents. Sequence variants were filtered for a recessive mode of inherit- ance (homozygous and compound heterozygous) to identify candidate variants. Candidate gene variants were validated via Sanger sequencing.

Results: Whole-exome sequencing revealed a novel, homozygous one base pair insertion in LACC1 (NM_153218.2:c.827delC) in the af- fected children. Both parents are heterozygous carriers; two un- affected children carry wildtype alleles. The mutation results in a frameshifting p.(Thr276Lysfs*2) with a premature stop at position 278. Its location in the fourth out of 7 exons is designated to cause either nonsense-mediated mRNA decay with loss of enzymatic activity or transcription of a truncated protein, respectively.

Conclusion: We here present a consanguineous family with 2 af- fected individuals suffering of severe forms of JIA associated with a novel mutation in LACC1. Together with the previously described cases of LACC1-associated familial systemic JIA and the reported asso- ciation with other inflammatory and infectious diseases, the present case adds evidence that LACC1 is a recessive disease gene and its loss of function results in a familial JIA phenotype.

Disclosure of Interest
None Declared
P162
Serial serum interleukin-6 levels in systemic Juvenile Idiopathic Arthritis as a predictor of treatment response
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Pediatric Rheumatology 2017, 15(Suppl 1):P162

Introduction: Interleukin-6 (IL-6) is the main cytokine that has many roles in pathogenesis of systemic juvenile idiopathic arthritis (SJIA). Previous studies showed that serum IL-6 levels were high in active SJIA patients when compared with inactive SJIA patients or healthy controls. However, the benefit of serial serum IL-6 levels monitoring is still unknown.

Objectives: To determine if serial serum IL-6 levels in SJIA patients can be a predictor of the treatment response.

Methods: A total of 422 serum samples were obtained from 26 SJIA patients, fulfilled the International League of Associations for Rheumatology (ILAR) criteria, in Ramathibodi hospital between January 2013 and May 2015. Serum IL-6 levels were measured during active disease and repeated measures every 2-6 weeks. In each visit, disease activity were assessed by using Juvenile Arthritis Disease Activity Score-27 (JADAS-27), including number of active joints (from total 27 joints), erythrocyte sedimentation rate (ESR), physician global assessment, parent/patient global assessment and the American College of Rheumatology Pediatric (ACR Pedi) 30, 50, 70, 90 criteria were used to determine the treatment response. SJIA patients were classified into two groups based on serial serum IL-6 levels: 1) rapid decrement group, serum IL-6 levels returned to normal within 1 year after active disease; 2) persistent group, highly persistent serum IL-6 levels, which were not return to normal within 1 year. Simple descriptive statistics and survival analysis were used. For predictor of reaching ACR pedi 50, multivariate analysis by Cox’s regression model was applied. This abstract is original and previously unpublished work.

Results: There were 12 patients in the rapid decrement group and 14 patients in the persistent group with 20.8 months of mean follow-up period. Baseline characteristics between two groups in age, sex, and disease activity at time of enrollment were not different. In rapid decrement group, median duration for returning to normal of serum IL-6 levels was 7.4 months and median duration for reaching ACR pedi 30, 50, 70, 90 were 1.6, 2, 4.7, and 7 months, respectively. In persistent group, only 3 patients had serum IL-6 levels returned to normal in 21 months. Median duration of reaching ACR pedi 30 was 18 months while no ACR pedi 50, 70, and 90 response were reached. After multivariate analysis was done, the predictor of reaching ACR pedi 50 was SJIA patients whose serum IL-6 levels returned to normal within 1 year (Hazard ratio 38.74, 95% Confidence Interval 3.43-437.87).

Conclusion: Monitoring serial serum IL-6 levels is beneficial in assessment the treatment response in SJIA patients. Returning to normal levels of serum IL-6 within 1 year was the predictor of reaching ACR pedi 50 and also demonstrated a better treatment response when compared with highly persistent serum IL-6 levels.

Disclosure of Interest
None Declared

P163
Validation of MRP8/14 serum levels as biomarker for the diagnosis of systemic Juvenile Idiopathic Arthritis in fever of unknown origin
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Pediatric Rheumatology 2017, 15(Suppl 1):P163

Introduction: The differential diagnosis of fever of unknown origin (FUO) is a major challenge in pediatrics especially for differentiation of systemic-onset juvenile idiopathic arthritis (SJIA) and infectious diseases. In a pilot study the analysis of MRP8/14 serum has been demonstrated as an excellent tool for the diagnosis of SJIA, allowing early differentiation of patients with autoinflammatory diseases SJIA or Familial Mediterranean Fever (FMF) versus those with other diseases with a specificity of 95% (Frosch et al. A&R 2009).

Objectives: Based on our study published in 2009, the analysis of MRP8/14 serum levels has been offered to paediatric rheumatologists in Germany in the last years. Therefore, we aimed to validate the findings from our pilot study in samples from daily clinical practice and to evaluate their relevance in clinical practice.

Methods: The study was designed as a retrospective analysis. The study group comprised 984 patients from 44 centers who presented with FUO. Patients were selected from our database between January 2009 and October 2012. Data collected included signs (laboratory parameters CRP, BSG, leucocytes) and symptoms as well as the final diagnosis made by the caring physicians. In all samples, concentration of MRP8/14 was determined by sandwich enzyme-linked immunosorbent assay (ELISA) with a cut-off of 9200 ng/ml (for SJIA versus other diseases). A questionnaire about the relevance of the MRP8/14 result for the final diagnosis was completed.

Results: Final diagnoses made by physicians were SJIA (n = 301), FMF (n = 135) and other inflammatory diseases (including infections, vasculitis and other autoinflammatory diseases) (n = 548). MRP8/14 serum levels of patients with SJIA or FMF (10.090 ± 1.930 ng/ml, mean ± SEM) were elevated compared to other diagnoses (3.140 ± 570 ng/ml) irrespectively of the presence of fever and anti-inflammatory treatment. In the group of untreated patients with fever (n = 213) MRP8/14 levels of SJIA patients (18.685 ± 4.130 ng/ml) were even higher compared to other diagnoses (5.285 ± 1.535 ng/ml). In this group, the sensitivity and specificity of MRP8/14 to differentiate between patients with SJIA vs. other diseases (excluding FMF) were 75% and 89% respectively. The sensitivity of the test increased to 83% in the presence of fever, joint pain and CRP > 1 mg/dl. There was a significant correlation (p < 0.001) between MRP8/14 and CRP or leucocyte counts in the total group of patients with SJIA. The clinicians reported that MRP8/14 results were helpful (56%), decisive (11%), not important (32%) and distracting (1%) in the total cohort.
Conclusion: Measurement of MRP8/14 levels is a helpful tool for the diagnosis of SJIA in FUO. In daily clinical practice the marker has a sensitivity of 75% and specificity of 89% to detect SJIA in untreated patients with fever. Additional markers might be useful to improve the detection of SJIA in patients presenting with FUO.

Disclosure of Interest
None Declared

P164
Proteomic discovery of SJIA diagnostic and phenotypic biomarkers
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Pediatric Rheumatology 2017, 15(Suppl 1):P164

Introduction: Systemic juvenile idiopathic arthritis (SJIA) is the most severe form of childhood arthritis. At onset, it resembles an autoinflammatory disease. Later in the clinical course, subphenotypes of SJIA are recognised, which may require different therapeutic strategies. However, SJIA specific diagnostic and phenotypic biomarkers are still lacking.

Objectives: We aimed to identify putative serum biomarkers that could distinguish patients with two subphenotypes of SJIA (classical SJIA with predominant systemic features ‘sys_SJIA’ or articular dominant ‘artic_SJIA’) from patients with infection.

Methods: All SJIA patients selected were diagnosed according to ILAR criteria, and those with infection had a confirmed bacterial or viral illness. Crude serum was first affinity-depleted of the 14 highest abundant proteins (MARS14, Agilent Technologies). In-solution trypsin digestion was performed on depleted serum samples. Samples were then desalted and eluted using C18 resin pipette stage tips. Label-free liquid chromatography mass spectrometry (LC-MS/MS) was performed on two platforms, the quadrupole time-of-flight (QTOF) and the Q-Exactive for comparison. The false discovery rate (FDR) was set at 0.01. Both sample and technical replicates were included to minimise biological and technical variability. Software packages were used for qualitative (using Mass Hunter) and quantitative (PEAKS Studio, Spectrum Mill and Max Quant) analyses and a range of statistical analyses, which included random Forest and principal component analysis, was undertaken.

Results: Patients with sys_SJIA (n = 10), artic_SJIA (n = 10) and infection (n = 10) had clinical features as follows: fever present (10 vs 0 vs 10), joint pain (7 vs 5 vs 4), exanthema (6 vs 0 vs 3), serositis (1 vs 0 vs 1) and hepatosplenomegaly (3 vs 1 vs 4). Biologic therapy was only used by n = 2 in the artic_SJIA group, while steroids were used by 3 and 2 patients in the sys_ and artic_SJIA groups respectively.

The results of univariate and multivariate analyses are shown in the Table. Between 31 and 41 differentially expressed proteins were identified in the group comparison analyses. In multivariate analyses, this translated to high area under the curve (AUC) for all SJIA (n = 20) versus infection (0.94). However, as anticipated, the sys_SJIA was not as easy to discriminate from the infection group (AUC: 0.58) as the artic_SJIA group (AUC: 0.97). However, the AUC for sys_SJIA versus artic_SJIA was 0.94, with proteins related to innate immune pathways being over-represented in sys_SJIA.

Conclusion: This study indicates differences in the serum proteomic signature of patients with SJIA who have different clinical courses/phenotypes, as well as in SJIA versus infection. These findings will be verified using a quantitative multiple reaction monitoring (MRM) assay on a larger cohort of patients, which is currently under development. Whilst further evaluation is still required, biomarker signatures developed on this work could potentially be used in the future for improved diagnosis, classification and management of patients with SJIA.

Trial registration identifying number:

Disclosure of Interest
None Declared

Table 45 (abstract P164). See text for description

|                | Univariate analysis | Multivariate analysis: AUC | Overlapping proteins |
|----------------|---------------------|----------------------------|----------------------|
|                | Differentially expressed proteins, n, p ≤ 0.05 | (Top 50 differentially expressed proteins) (n) |
| Sys_SJIA vs artic_SJIA | 41 | 0.94 | 5 |
| SJIA (all) vs infection | 35 | 0.86 | 3 |
| Sys_SJIA vs artic_SJIA | 31 | 0.58 | 2 |
| Artic_SJIA vs infection | 36 | 0.97 | 4 |

P165
Anakinra for first line steroid free treatment in systemic onset Juvenile Idiopathic Arthritis
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Presenting author: Gerd Horneff
Pediatric Rheumatology 2017, 15(Suppl 1):P165

Introduction: First line treatment with the IL-1 inhibitor Anakinra is recommended by the ACR (1) and part of the Childhood Arthritis and Rheumatology Research Alliance consensus treatment plans for systemic JIA (2).

Objectives: In patients naïve for corticosteroids, as a hypothesis, a corticosteroid-free treatment regimen may allow reconstitution of an impaired NK-cell function resulting into decrease of IL-18 and probably remission of the disease (3).

Methods: Experience with first line treatment with Anakinra for systemic JIA is presented.

Results: 6 pts presenting with systemic JIA were treated with Anakinra after informed consent was taken for off label treatment. Two pts had received a first 3 day course of corticosteroids before referral. All pts presented with fever and rash, 3 with arthritis and 2 with polyarthritis. After admission, spiking fever was documented for three days while infections were ruled out thoroughly. ANA/ANCA or rheumatoid factors were undetectable in all pts in the presence of highly elevated acute phase reactants, IL-18 and myeloid-related protein S100A8/A9 (table). Treatment with Anakinra was started at a dosage of 2-3 mg/kg sc. daily. Immediate complete response was observed in 4 pts (table) while virtually no effect was reached in one. In the remaining patient, a 15 year old girl, absence of fever was noted for 20-22 hours after Anakinra injection. Furthermore a marked improvement of arthritis was noted and proven by diminished Doppler-signalling. However, the patient refused to receive two daily injections and was set on oral corticosteroids. She immediately improved but deteriorated upon tapering of steroids and finally was treated successfully with tocilizumab. In one responder, recurrence of the disease was noted after discontinuation of Anakinra and the patient was set on oral corticosteroids recently. In a further responder, a disease flare upon treatment was managed with increased dosing of Anakinra to 5 mg/kg but the patient developed life threatening macrophage activation syndrome triggered by a viral infection and treatment was discontinued. Beside this event, Anakinra was tolerated uneventfully.

Conclusions: Experience with first line treatment with Anakinra for systemic JIA is presented. Anakinra treatment was initially successful in 4 patients while remission off drug was reached in 2 only. The presence of arthritis, especially with a polyarticular pattern might be prognostic. Of importance, one child developed MAS upon treatment with Anakinra triggered by a viral infection. Beside this, treatment with Anakinra was well tolerated and unfavourable effects of long lasting steroid application were avoided in the responders.

References
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Disclosure of Interest
None Declared
Table 46 (abstract P165). See text for description

| Patient | 1 | 2 | 3 | 4 | 5 | 6 |
|---------|---|---|---|---|---|---|
| Age (years) | 11.0 | 6.1 | 11.0 | 11.0 | 11.0 | 11.0 |
| Gender | f | m | f | f | f | f |
| Systemic features | Spiking fever, rash | Spiking fever, rash | Spiking fever, rash | Spiking fever, rash | Spiking fever, rash | Spiking fever, rash |
| JADAS-MDA | 3.6 | 3.6 | 3.6 | 3.6 | 3.6 | 3.6 |
| IL-1i | 4.6 | 4.6 | 4.6 | 4.6 | 4.6 | 4.6 |
| IL-1i | 4.6 | 4.6 | 4.6 | 4.6 | 4.6 | 4.6 |
| JADAS10 | 13.0 | 13.0 | 13.0 | 13.0 | 13.0 | 13.0 |
| IL-1i | 8.2 | 8.2 | 8.2 | 8.2 | 8.2 | 8.2 |
| IL-1i | 8.2 | 8.2 | 8.2 | 8.2 | 8.2 | 8.2 |
| ESR, mm/h | 45.6 | 101.7 | 106.6 | 197.0 | 329.8 | 99.0 |
| CRP, mg/l | >5,000 | >5,000 | >5,000 | >10,000 | >10,000 | >10,000 |
| IL18, pg/ml | >10,000 | >10,000 | >10,000 | >10,000 | >10,000 | >10,000 |
| Pretreatment | 3 days corticosteroids | none | none | none | none | none |
| Treatment | Remission off treatment | Remission off treatment | Increase of dosage upon flare, discontinued with BAVS | No response, switch to Tocilizumab | Minor response, switch to Tocilizumab | Remission, relapse after discontinuation |
| Outcome | JADAS Remission n (%) | 16 (18.2%) | 15 (44.1%) | 16 (50%) | 5 (2.9%) | 7 (1.9%) |
| IL-1i | 8.2 (49–133) | 8.2 (51–133) | 8.2 (49–133) | 8.2 (51–133) | 8.2 (51–133) |

**P166**

Experience with Tocilizumab, interleukin-1 inhibitors and etanercept for systemic Juvenile Idiopathic Arthritis

Gerd Horneff, Anna C. Schulz, Kirsten Minden, Frank Weller-Heinemann, Anton Hospach, J-Peter Haas, and BIKER collaborative group

**Introduction:** In Germany, surveillance of approved biologics for juvenile idiopathic arthritis (JIA) is performed by the Biologics Register (BikeR) since 2001.

**Objectives:** To report on the efficacy and safety of Tocilizumab (TOC), the interleukin-1 inhibitors Anakinra and Canakinumab (IL-1i) compared to Etanercept (ETA). The role of natural killer (NK) cells in sJIA pathogenesis and granzyme k in natural killer cells of systemic Juvenile Idiopathic Arthritis patients

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**Presenting author:** Gerd Horneff

Pediatric Rheumatology 2017, 15(Suppl 1):P166

**Table 47 (abstract P166). See text for description**

| Patient | Etanercept | Tocilizumab | IL-1i |
|---------|------------|-------------|-------|
| N = 143 | N = 72 | N = 55 |
| Baseline JADAS10 Median (IQR) | 8.2 (5–13.0) | 9.6 (4.9–13.3) | 8.2 (5–13.3) |
| Month 6 | 6.2 (1.5–15.6) | 8.0 (3.1–15.3) | 6.6 (0.2–20.0) |
| JADAS MDA n (%) | 26 (29.5%) | 21 (61.8%) | 21 (65.6%) |
| JADAS Remission n (%) | 16 (18.2%) | 15 (44.1%) | 16 (50%) |
| Discontinuations (< Month 6) | Inefficacy, n (%) | 23 (26.1%) | 4 (11.8%) | 7 (21.9%) |
| Intolerance, n (%) | 1 (2.9%) | 1 (1.1%) | |
Methods: RNA-Sequencing of highly purified NK cells from six active sJIA patients and six healthy controls was performed to investigate NK cell-specific transcriptional alterations. NK cell-stimulating and other cytokines were quantified in plasma (n = 18). NK cell phenotype and cytotoxic activity against tumor cells were determined (n = 10), next to their IFN-γ-producing function (n = 8).

Results: NK cells of sJIA patients showed an altered gene expression profile compared to controls, with increased expression of innate genes such as TLR4 and S100A9 and decreased expression of immune-regulating genes like IL10RA and GZMK, indicating an enrichment of immune-inflammatory pathways in sJIA. Alterations in the balance of inhibitory and activating receptors, with decreased KLRG1 and increased Nkp44 expression, were noted in sJIA NK cells. Although sJIA NK cells showed intact cytotoxicity and degranulation against tumor cells, a decreased granzyme K expression in CD56 bright NK cells and a defective IFN-γ-production in response to IL-18 was observed. In sJIA plasma, an increased IL-18 and decreased IFN-γ/IL-18 ratio was determined.

Conclusions: NK cells in sJIA display an innate inflammatory gene expression profile. Although their cytotoxic function is intact, subtle defects in NK-related regulatory pathways such as granzyme K expression and IL-18-driven IFN-γ production may underlie the immune-inflammatory dysregulation in sJIA.

Disclosure of Interest
None Declared

P168
Serum IL-18 levels as a marker for disease activity in systemic Juvenile Idiopathic Arthritis during tocilizumab therapy
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Pediatric Rheumatology 2017, 15(Suppl 1):P168

Introduction: Tocilizumab (TCZ), a humanized anti-interleukin (IL)-6 receptor monoclonal antibody, is an effective cytokine inhibitor for the treatment of systemic juvenile idiopathic arthritis (s-JIA). However, clinical symptoms of s-JIA could be masked during TCZ therapy, even in macrophage activation syndrome (MAS) phase. Furthermore, serum CRP concentrations never increased during TCZ therapy, even in disease flare. Hence, a novel and promising biomarker for evaluating s-JIA disease activity is desired.

Objectives: The aim of this study is to investigate the clinical significance of serum IL-18 levels as a marker for disease activity and for predicting the prognosis in patients with s-JIA during TCZ therapy.

Methods: We serially measured serum IL-18 levels in 14 patients with s-JIA during TCZ therapy and compared them with the disease activity and their prognosis. Serum concentrations of IL-18 was evaluated using commercial enzyme-linked immunosorbent assay, according to the manufacturer’s instructions.

Results: Serum IL-18 levels in patients with s-JIA treated with TCZ during active phase were significantly higher than those in inactive and remission phase as well as those in patients with s-JIA without TCZ. Serum IL-18 levels in 8 patients without relapse decreased to the levels <1000 pg/ml in remission phase. In contrast, serum IL-18 levels in all 6 patients with relapse sustained elevated or elevated again up to the levels >1000 pg/ml even after serum IL-18 levels normalized in inactive phase.

Conclusion: Serum IL-18 level is a useful marker for assessing the disease activity and predicting the prognosis in patients with s-JIA during TCZ therapy. Further larger studies is desired to define the diagnostic value of serum IL-18 as a promising biomarker for evaluating s-JIA disease activity.

Disclosure of Interest
None Declared

P169
Cytokine profile of Adult-Onset Still’s Disease: comparison with systemic Juvenile Idiopathic Arthritis
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Presenting author: Masaki Shimizu
Pediatric Rheumatology 2017, 15(Suppl 1):P169

Introduction: A comparison of the epidemiology, genetic background, clinical presentation, and course of these diseases suggests convincingly that systemic juvenile idiopathic arthritis (s-JIA) and adult still’s disease (AOSD) are identical or similar illnesses. However, it remains unknown whether s-JIA and AOSD share the same pathogenesis. Recent studies have shown that inflammatory cytokines play pathogenic roles in the disease processes of both diseases. We hypothesized that the patterns of cytokine release in s-JIA and AOSD share common features and s-JIA and AOSD belong to the same disease category.

Objectives: To test our hypothesis, we analyzed serum levels of cytokines in patients with both diseases and compared them with the clinical features.

Methods: 33 patients with AOSD, 77 patients with s-JIA were analyzed. Serum levels of IL-6, IL-8, neopterin, sTNFRI and II were quantified by ELISA. Results were compared with clinical features of AOSD.

Results: Cytokine profile of both diseases was almost similar. It was characteristic that serum IL-18 levels were extremely high in active phase and remained elevated even in inactive phase. Two distinct subsets based on their serum IL-6 and IL-18 levels were identified in AOSD as well as s-JIA. The number of patients with arthritis was significantly higher in the IL-6–dominant subgroup.

Conclusion: AOSD and s-JIA are the same spectrum, which is based on the significant production of IL-18. Two subsets of patients with AOSD can be identified on the basis of their serum IL-6 and IL-18 levels. These two subsets appear to be characterized by certain distinct clinical features. Monitoring the cytokine profile with IL-18 and IL-6 might be useful to predict disease course.

Disclosure of Interest
None Declared

P170
Genetic architecture of systemic Juvenile Idiopathic Arthritis distinguishes it from oligo- and polyarticular Juvenile Idiopathic Arthritis
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Pediatric Rheumatology 2017, 15(Suppl 1):P170

Introduction: Juvenile idiopathic arthritis (JIA) is a heterogeneous group of conditions that are unified by the presence of chronic childhood arthritis without an identifiable cause. Systemic JIA (sJIA) is a rare form of JIA that is characterized by episodic systemic inflammation and chronic arthritis. Approximately half of children with sJIA go on to develop destructive, life-long arthritis that has phenotypic similarities with the oligoarticular/seronegative polyarticular (polygoJIA) or seropositive polyarticular (RFpolyJIA) forms of JIA. Importantly,
therapies that are effective in other forms of JIA are much less effective in sJIA, particularly among patients with persistent arthritis. Investigating the genetic architecture of sJIA and how it compares with other forms of JIA may improve our understanding of these differences. However because of its rare nature, genomic investigations of sJIA have been limited, providing few insights into its pathophysiology and not allowing for its comparison with other forms of JIA.

**Objectives:** In the present study, we sought to identify genetic factors that influence sJIA pathogenesis and to compare the genetic architecture of sJIA with the other common forms of JIA.

**Methods:** We performed a genome-wide association study of 982 children with sJIA collected in 9 countries by participating sites of the International Childhood Arthritis Genetics (INCHARGE) Consortium. Following stringent quality control operations, over 6 million single nucleotide polymorphisms (SNPs) were tested for association with sJIA by logistic regression, adjusted for gender and ancestry, in 9 geographically-defined strata. Association results were combined by fixed-effect meta-analysis. Weighted genetic risk scores (wGRS) for sJIA were calculated and wGRS were compared between sJIA cases and healthy controls using the Wilcoxon Rank Sum test. The correlation of wGRS with sJIA was tested by both logistic regression and by analysis of receiver operating characteristic (ROC) curves with calculation of the area under the curve. Quantile-quantile (QQ) plots were used to determine whether sJIA-associated SNPs were enriched within the set of polygoJIA-associated SNPs from the recent study by Hinks and colleagues.

**Results:** The major histocompatibility complex locus and a second locus on chromosome 1 both showed association with sJIA that exceeded the threshold for genome-wide significance (p < 5E-8), and 23 additional loci had evidence suggestive of association with sJIA (p < 5E-6). There was virtually no overlap between the sJIA susceptibility loci and the known risk loci for other forms of JIA. ROC curve analysis and logistic regression of wGRS failed to identify any correlation between sJIA and either polygoJIA or RFpolyJIA. Examination of QQ plots found no evidence for enrichment of sJIA associations among polygoJIA-associated SNPs. Collectively, we found no evidence of shared genetic architecture between sJIA and either polygoJIA or RFpolyJIA.

**Conclusion:** sJIA has a distinct genetic architecture without evidence of pleiotropy or shared genetic risk factors with other common JIA subtypes. This indicates that sJIA is a unique disease process and not a variant form of another JIA subtype. As such, the pathophysiology of sJIA must be investigated separately from the other JIA subtypes to identify therapeutic targets specific to sJIA. Moreover, the management of children with sJIA should be directed by studies of sJIA, and not by trials or observations of other subtypes of JIA.

**Disclosure of Interest**
None Declared

**P171**

**Bone marrow findings at children with Juvenile Idiopathic Arthritis**
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**Disclosure of Interest**
None Declared

**Introduction:** Juvenile idiopathic arthritis (JIA) involves inflammation of one or more joints lasting more than 6 weeks and the onset of the disease before the age of sixteen with the prior exclusion of other diseases and conditions in which the arthritis occurs as part of the clinical picture. Malignant diseases in children can start with musculoskeletal manifestations and prior to the diagnosis of JIA is necessary to exclude malignancy, primarily leukemia.

**Objectives:** The aim of this study is to investigate bone marrow findings at children with JIA at the onset of disease.

**Methods:** The study included 24 children diagnosed with JIA, treated from January 2010 to January 2016 at University Clinical Center of Republic of Srpska. Bone marrow aspiration (BM) was performed in all JIA patients as a part of an initial diagnostic evaluation. Control group included 24 children with immune thrombocytopenic purpura (ITP). BM smears were retrospectively reviewed by the same hematologist for the purpose of this study. Bone marrow smears were evaluated for cell populations as well as myelodysplastic features and compared to control group.

**Results:** The study included 24 patients with JIA, 16 girls and 8 boys. Median age was 5 years and two months. Ten children had systemic onset JIA, nine had oligoarthritis and five had polyarthritis. The characteristics of BM in JIA was myeloid hyperplasia and elevated plasmocyte count. There was no difference between JIA patients and control group in terms of myelodysplastic features. The patients with JIA had a significantly higher percentage of cells of neutrophil series (neutrophils, metamyelocytes and myelocytes) compared to control group. Hemophagocytosis was found at one patient with systemic onset of JIA.

**Conclusion:** We emphasize the importance of including BM aspiration into the diagnostic procedures in JIA aimed of excluding malignancies. Findings of this study can be helpful in explaining hematological changes in JIA.

**Trial registration identifying number:** DEA2487D-FOFB-4B6A-BF2A-60844E3AC45A

**Disclosure of Interest**
None Declared

**P172**

**Inflammatory bowel disease following anti-interleukin-1-treatment in systemic Juvenile Idiopathic Arthritis**
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**Presenting author:** Boris Hugle

**Disclosure of Interest**
None Declared

**Introduction:** Inflammatory bowel disease (IBD) can develop with some rheumatic diseases in childhood, including juvenile idiopathic arthritis (JIA). Inflammatory bowel disease is frequently associated with other immune-mediated diseases; however, systemic onset JIA (sJIA) has not previously been connected to IBD. Treatment of sJIA has significantly changed in recent years, possibly causing changes in inflammatory patterns.

**Objectives:** We present three pediatric cases with confirmed diagnosis of sJIA who developed IBD diagnosed by histology and coloscopy during treatment with IL-1 antagonists.

**Methods:** Data from the German Center for Pediatric and Adolescent Rheumatology from 2010 until 2015 were analyzed by retrospective chart review.

**Results:** There were 82 patients with confirmed diagnosis of systemic JIA. Of these, four were identified with a diagnosis of IBD. Three of these children with confirmed IBD confirmed by colonoscopy (two cases of Crohn's disease, one case of ulcerative colitis) 0.8 – 4.3 years after diagnosis. All three were treated with IL-1 antagonists (anakinra in two cases, canakinumab in one case). The fourth patient developed transient terminal ileitis confirmed by histology on treatment with etanercept, two months after combination treatment of anakinra and etanercept for 16 months.

**Conclusion:** IBD in systemic onset JIA is a rare complication. Treatment with IL-1 antagonists might be causally related to a switch in clinical phenotype of the underlying inflammatory process.

**Disclosure of Interest**
None Declared
Introduction: Systemic juvenile idiopathic arthritis (SJIA) is a multisystem inflammatory disorder rarely diagnosed in infants. Tocilizumab is considered the mainstay treatment for difficult to treat SJIA in patients over four years of age. We present two infants with SJIA who failed to respond to conventional treatment and were successfully treated with tocilizumab. To our knowledge there are no previous data on the efficacy and safety of this drug in infancy.

Objectives: To describe the efficacy and safety of tocilizumab in two infants with SJIA.

Methods: An 8 month old male presented with persistent fever and raised inflammatory markers; no source of infection was identified. He had splenomegaly and lymphadenopathy. The patient underwent major work-up for infection and malignancy and received antibiotics. Ultrasonography revealed free peritoneal and pericardiac fluid. Symptoms persisted despite the use of antibiotics. He subsequently developed a maculopapular rash on his trunk while the following day he developed arthritis of the wrists, knees, ankles and 3rd and 4th PIP joints. The patient was diagnosed with SJIA and received pulses of methylprednisolone; MTX was initiated. Genetic testing for auto-inflammatory diseases (FMF, TRAPS, HIDS, CAPS) was negative; genetic and functional screening assays for primary HLH were normal. Serositis, arthritis and lymphadenopathy improved after initiation of treatment. At twelve weeks, fever spikes and rash persisted despite continuous significant doses of prednisolone (Table 48). In addition, he became cushingoid, hypertensive and extremely irritable. Inflammatory markers were high with persistent thrombocytosis. We initiated tocilizumab. A month afterwards, the patient’s condition had markedly improved; although he occasionally presented with a rash. A year later he remains asymptomatic, leading a fully normal life.

Results: The other patient (female 9 m) presented with pyrexia of unknown origin for 15 days with hepatosplenomegaly and serositis. She developed bilateral knee arthritis and a maculopapular rash on day 21. Following exclusion of bacterial infection, malignancy and other genetically defined autoinflammatory disease, she was diagnosed as SJIA and started on an identical regimen as patient above. Similarly we failed to decrease the steroid dose below 1 mg/kg/day; she remained on this regimen for further six months during which she had severe growth delay; tocilizumab was initiated at 18 months of age (Table 48). After six months, she remains asymptomatic and growing well. Prednisolone was tapered over three months in both cases. Clinical remission was achieved by nine and six months for the two patients respectively. None of them developed a flare. However, both patients experienced side effects potentially related to tocilizumab. The first patient had primary varicella infection and the second had recent upper respiratory tract infections.

Conclusion: Tocilizumab may be considered as a treatment option in patients less than 2 years of age with severe SJIA. A current open label study is exploring the pharmacokinetics and pharmacodynamics of tocilizumab in this age group and will provide some efficacy and safety data.

Disclosure of Interest
None Declared

Table 48 (abstract P173). Clinical and disease characteristics of the two patients at tocilizumab initiation

| Patient characteristics | Patient I | Patient II |
|-------------------------|-----------|-----------|
| Age                     | 11 months | 18 months |
| Ethnicity               | Caucasian | Caucasian |
| Gender                  | male      | female    |
| Disease duration        | 3 months  | 7 months  |
| Steroid dose            | 2 mg/kg/day| 1 mg/kg/day|
| Methotrexate dose (SC)  | 15 mg/m²  | 15 mg/m²  |
| Tocilizumab dose (IV)   | 12 mg/kg every two weeks | 12 mg/kg every two weeks |
| ACR Pedi 30 (+ no fever)| By week 12| By week 8  |
| ACR Pedi 90 (+/no fever)| By week 36| By week 24 |
| Adverse events          | Varicella | Recurrent upper respiratory tract infections |

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Complete clinical remission with tocilizumab in two toddlers with systemic Juvenile Idiopathic Arthritis-a case series
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P174
Canakinumab for first line steroid free treatment in a child with systemic onset Juvenile Idiopathic Arthritis
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Introduction: First line treatment with the IL-1 inhibitor Anakinra has been recommended by the ACR (1) and is part of the Childhood Arthritis and Rheumatology Research Alliance consensus treatment plans for systemic JIA(2). In patients naïve for corticosteroids, as a hypothesis, a corticosteroid-free treatment may allow reconstitution of an impaired NK-cell function resulting into decrease of interleukin-18 and probably remission of the disease (3). Canakinumab is a long acting high-affinity human IL18 monoclonal antibody that selectively bind to human IL18 and thereby inhibiting downstream signalling pathways.

Objectives: In 2014, canakinumab has been added to the CARRA treatment plans (4). Thus, a single injection of canakinumab has been used as a sole treatment in a child newly diagnosed with systemic active JIA.

Methods: A 2 7/12 year old boy presented with fever up to 41.0 °C, rash, articulargia, splenomegaly, refusal to stand or walk and highly elevated acute phase reactants. Infections were ruled out thoroughly. Initial bone marrow aspirate was unremarkable. ANA/ANCA or rheumatoid factors were undetectable. The patient was referred to the rheumatology clinic.

Results: After admission, spiking fever was documented for 3 days. Highly elevated acute phase reactants were present. There were signs of macrophage activation syndrome including leucocytopenia, neutropenia, thrombocytopenia, elevated ferritin (>1200 μg/l), lactatedehydrogenase (1028 U/l), transaminases (ALAT 110 U/l, ASAT 124 U/l) and triglycerides (497 mg/dl) but normal fibrinogen. Biomarkers, IL-18 and myeloid-related protein S10A8/A9 were highly elevated (table).

Informed consent was taken for off label treatment with canakinumab instead of routine care with corticosteroids. Fever immediately disappeared the day after a single application of canakinumab (2 mg/kg sq). No further rash or arthralgia were noted. CRP level rapidly decreased and step by step returned to normal while the clinical condition normalized and splenomegaly completely resolved. The patient was discharged at day 3. A rapid decline of all disease activity markers was noted. Of interest, MRP S100A8/A9 normalized after 6 weeks while serum level of IL-18 markedly decreased but were still
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Systemic onset Juvenile Idiopathic Arthritis: epidemiological features of Moroccan cohort
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Introduction: Systemic onset juvenile idiopathic arthritis (s-JIA) is a rare autoimmune-inflammatory disease that belongs to the big group of Juvenile Idiopathic Arthritis (JIA) representing about 10 to 17% of cases. It is distinguished from other types by the presence of systemic manifestations with marked inflammatory reaction beside the articular symptoms.

Objectives: To analyze the epidemiological, clinical, diagnostic tests, therapeutic and evolutionary parameters in SoJIA.

Methods: This is a retrospective study of SoJIA patients followed in a pediatric rheumatology monocenter over a 16-year period from January 2000 to December 2015. The diagnosis was based on ILAR 2004 revised criteria. The hospital charts of these patients were reviewed and information was collected.

Results: The study included a total of 59 cases, 27 girls and 32 boys. The mean age of disease onset was 4 years and 5 months while the mean duration delay of diagnosis was 4 months and 9 days. Consanguinity was reported in 22% of cases. Fever was objectified in all patients; it was typical in 61% of them, while arthritis was present at onset in only 76% of patients with a 6.2 mean number affected joints during the evolution whose 13% was with standard radiological expression. Morning stiffness was reported in 74% of patients and rash in only in 49%. The hematopoietic involvement was represented by hepatomegaly, splenomegaly and lymphadenopathy in 8.4%, 8% and 27% of cases and serositis in 22%. NSAIDs were prescribed in all cases, oral corticosteroids in 72% and intra-articular form in 11.8%. Methotrexate was given in 49% of patients, thalidomide in 2 cases, Leflunomide in only one case. Meanwhile biological were recommended in 15 cases 2 Etanercept and 13 (22%) Tocilizumab. The study of evolution included 42 patients. The number of relapses was ≥3 relapses in 22 of them, 84% with systemic manifestations and 16% with exclusive joint shape. Complications were growth impact with short stature in 22% of cases, Cushingoïd facies in 40%, glucocorticoid-induced cataract in 5%, osteoporosis in 5%, macropage activation syndrome in 2 patients; twice for one of them, severe infection in 6.7% and joint destruction in 8.4% of patients.

Conclusion: Our study is characterized by a slight male predominance for a sex ratio usually 1, the occurrence of macropage activation twice in the same patient justifying a genetic test which include normal, 2 families with 2 children suffered from same genetic exploration that not have been yet realized. The clinical course was modified by biological therapy in refractory cases. Our patients underwent conventional treatment until 2006 when etanercept was tried with poor results and replaced by Tocilizumab from 2009 permitting a complete remission for almost patients treated without treatment withdrawals because of expensive cost and outside the joint squaress already existing.

Disclosure of Interest
None Declared

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P176
Four years of tocilizumab therapy for systemic Juvenile Idiopathic Arthritis in Georgia
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Introduction: The blockade of interleukin 6 has been shown to reduce the corticosteroid dependence in children suffering from systemic juvenile idiopathic arthritis (s-JIA).

Objectives: To investigate the effect of TCZ treatment in 9 patients with systemic JIA.

Methods: 9 patients suffering from s-JIA with inadequate response to therapy (prednison) have received Tocilizumab (TCZ) since the year 2013 in Georgia. They were retrospectively analyzed. The data were collected in one pediatric outpatient unit.

Results: The mean age at diagnosis was 32.5 months (from 13 to 73 months), there were 4 boys (44.4%) and 5 girls (55.6%) (ratio 4:5). All patients were treated with prednison for 25.2 months (SD 23.8, from 6 to 72 months) before starting TCZ. Currently 6 patients are still treated with prednison: 4 patients (66.7%) are on therapy with prednison with appropriate reducing scheme and in 2 patients (33.3%) prednison was restored because flare of the disease.

The mean period of time from diagnosis to TCZ therapy was 25.6 months (SD 23.5, from 7 to 72 months). Patients were treated with TCZ for 14.6 months (SD 25.6 months, from 2 to 23 months). There were still 9 patients (100%) treated with TCZ. 5 patients (55.6%) are in remission, 4 (44.4%) have active arthritis, without systemic signs and symptoms.

Adverse events included mild infection (2 patients) and increased aminotransferase levels (1 patient).

Conclusion: Our data show that TCZ was effective in severe, persistent systemic Juvenile Idiopathic Arthritis. At 12 weeks 5 (55.6%) patients have demonstrated response in ACR Pedi 70, 3 (33.3%) - in ACR Pedi 50, 1 (11.1%) – in ACR Pedi 30.

Disclosure of Interest
None Declared
Systemic onset juvenile idiopathic arthritis in Estonia
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Introduction: Juvenile idiopathic arthritis (JIA) is a heterogeneous group of childhood-onset chronic arthritis. Systemic JIA (sJIA) as one of the disease categories has prominent systemic features as well as arthritis and accounts for 5-15% of children with JIA (1). Every year 250-270 new cases of JIA are diagnosed in Estonia.

Objectives: The aim of this study is to describe clinical aspects, methods of treatment and achievement of clinical remission in patients with sJIA in Estonia.

Methods: Data for the study have been collected retrospectively from medical records of newly diagnosed patients with sJIA in Tallinn Children’s Hospital and Children’s Clinic Tartu University Hospital from 2000-2015.

Results: During years 2000-2015 sJIA was diagnosed in 26 patients, nine boys and 17 girls. Most frequent first diagnosis at hospitalisation was fever of unknown origin or unknown bacterial infection (73% of the cases). Mean age at diagnosis was 8.8 ± 4.8 years; boys were younger (7.5 ± 5.5 years) than girls (9.6 ± 4.6 years) but the age difference wasn’t statistically significant (p = 0.36). Mean duration of fever before the diagnosis of sJIA was 14.2 ± 4.0 days, 96% of patients had rash and 50% had hepatomegaly. At hospitalisation 46% of patients did not have clinical arthritis. All patients had high values of inflammatory markers (CRP 130 ± 87 mg/l and SR 58 ± 25 mm/h). Ferritin levels were determined in 57.6% of patients and the mean value was 1479.7 ± 1284.2 mg/l. All patients were treated with corticosteroids and 17 of them received intravenous pulse therapy (mean dose 12 mg/kg/dose). Mean number of pulses needed was 3.4. Oral prednisolone was given in a mean dose of 1.1 mg/kg/day. Biologic agent was started in 10 patients at a mean duration of the disease 1.5 (95% confidence interval 0.4 – 2.7) years. During the disease course three patients had documented growth suppression, three documented osteoporosis and one child developed macrophage activation syndrome. Clinical remission on treatment was achieved in 15 patients, but clinical remission off treatment only in nine patients.

Nine patients had monocyclic and 17 polycyclic course of sJIA. During the disease course active component of the disease was arthritis in three, systemic in nine and arthritis with systemic in 14 patients.

Conclusion: sJIA is a rare subtype of JIA. Most frequent clinical presentation at diagnosis was fever with rash and hepatomegaly, at hospitalisation arthritis was present in only half of patients. Corticosteroids were used in all patients, 1/3 received biological treatment. The disease had polycyclic course in 2/3 of patients and remission off treatment was achieved in 1/3 of cases.

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Disclosure of Interest
None Declared

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