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Invited lectures

11 Which are the clinical, pathophysiological and therapeutic similarities between migraine & hereditary angioedema with C1-inhibitor deficiency?

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Migraine is the second most disabling chronic neurological disease. Pathophysiologically, migraine is a genetically-driven, cyclic functional disorder affecting several areas of the brain, ultimately leading to an excessive activation of trigeminovascular afferents in the meninges. Through multiple pathways local release of inflammatory peptides as calcitonin gene-related peptide (CGRP), substance P, VIP and bradykinin are presumable. Neurological symptoms of C1-INH-HAE (hereditary angioedema with C1-inhibitor deficiency) as cephalalgia, hemiplegia, and migraine symptoms are similar, which might be due to the role that bradykinin plays. The unpredictable frequency of the attacks is ruled by the extremely variable efficiency of the fine-tuned interplay of the brain with dynamic stimuli originating from the internal or external environment. Unlike several other chronic neurological diseases, migraine can be treated and prevented. However, not all patients respond to treatment.
Both migraine and C1-INH-HAE are genetically-driven paroxysmal disabling diseases, which are characterized by recurrent unpredictable episodic attacks. In rare cases, C1-INH-HAE is manifested with neurological symptoms, including cephalalgia which does not respond to conventional treatment. The complement system has an important function in the regulation of bradykinin release within the brain. Some of the inflammatory molecules in migraine can be measured indirectly (serum, CSF) using ELISAs. This is different from C1-INH-HAE, where conventional laboratory tests are of diagnostic value.

In respect of treatment options: in migraine, treatment is broad, and recently a new class of drugs has been developed, and proven to be effective and safe, targeting CGRP and preventing migraine attacks, however, there may be non-responders.

The pathophysiology shares the presence of inflammatory molecules which lead to pain in migraine and angioedema attacks in C1-INH-HAE. Therapeutic options of both diseases are to be personalized and target-driven.

I2
Analysis of C1-inhibitor deficiency; need for standardization and quality control
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Deficiency of C1-INH results in episodic angioedema without urticarial that is inherited (hereditary angioedema, [C1-INH-HAE]) or acquired (C1-INH-AA). In addition to its role as an inhibitor of C1r and C1s of the classical pathway and MAS1 and MAS2 of the lectin pathway, C1-INH is the major inhibitor of factor XIIa and kallikrein. The lack of inhibition of these enzymes results in excessive bradykinin generation, which in turn mediates increases vascular permeability, leading to angioedema.

In the past decades complement analysis has undergone a tremendous development in part expanding beyond specialized laboratories. However, improving diagnostic strategies requires the correct choice of analytes and the use of well-characterized methods that will yield consistent results between laboratories.

In the diagnosis of the various types of angioedema, as in many fields of immunodiagnostic, there is considerable need for consensus and standardization of analytical methods, which will be a major challenge in the future. In recent years, laboratories specializing in complement analysis have joined with the International Complement Society and the IUIS to coordinate efforts to standardize and improve complement testing, ongoing efforts show first promising results (http://iuisonline.org/index.php?option=com_content&view=article&id=64&Itemid=69).

Since that time eight rounds of external quality assessment, now covering 18 parameters, also including those to better characterize angioedema patients (C4, C1-inhibitor [protein, function], autoantibodies to C1-inhibitor) have been completed. It is recommended to extend this efforts to a more comprehensive analysis of parameters of the clotting and kallikrein-kinin systems for better defining the pathophysiological background and to distinguish angioedema with C1-inhibitor deficiency from primary angioedema.

I3
Prodromes of HAE: scientific evidence or delusional perception?
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Signs and symptoms (Prodromes) which antedate swelling attacks have been noted since early descriptions of Hereditary Angioedema (HAE). Regrettably, no consensual definition of a prodrome exists, contributing to ambiguity and disagreement among medical disciplines and researchers. Heralding signals occurs in other diseases, mostly characterized by a chronic and undulating course. For instance, in Schizophrenia, bipolar disorders, and some neurodegenerative diseases, early signs of abnormal behavior, occurring few years before a psychotic crisis, are regarded as a prodrome. In Migraine, a prodromal stage with subtle symptoms precedes the attacks. Early emotional changes, physical symptoms and visual Aura, are regarded as premonitory and predict oncoming paroxysm of headaches. In Herpes Zoster, a prodrome occurs 48–72 h before the typical vesicular rash, and intense pain in the involved dermatome precedes the rash in more than 90% of cases. Prodromes have also been described in Familial Mediterranean Fever (FMF) and Capillary-leaf Syndrome (Clarkson’s syndrome).

A renewed interest in prodromes preceding HAE took place a decade ago, followed by an extensive literature search and some preliminary explorations. Further studies and expert opinions affirmed the fact that prodromes are more frequent than realized. Our group has recently designed and analyzed a new instrument to evaluate HAE prodromes. In this study 84% of the patients reported ever having a prodrome, and 87% said that they could predict an oncoming attack by experiencing a prodrome. A significant correlation was found between the perception of prodrome and ability to predict an oncoming attack. This data corroborates other studies reporting similarly high rate of association.

Despite consistent patient reports, prodromes’s remains elusive, and their precise nature and mechanisms are unknown. It could be hypothesized that they represent an early surge of angioedema mediators, such as complement fragments or factor XII-dependent contact system kinins. Indeed, bradykinin was detected in the stromal and endothelial cells of prodromal Erythema Marginatum skin rash. Additionally, evidence of early activation of the kallikrein-kinin system, with high kininogenase activity, proenzyme consumption, high-molecular weight kininogen cleavage and C1-INH function alteration, occurring before visible angioedema, was recently demonstrated. In another study, C4 depletion and higher than baseline C4a levels, were detected hours before the onset of an attack.

In summary, albeit frequently reported, prodromes have not been adequately investigated, and systematic tools for their evaluation are missing. Accurate prodrome evaluation is critical for early diagnosis of attacks and timing of medical interventions, particularly in an era when effective drugs are available for self-treatment.

Oral lectures
O1
Parallel comparison of three different assay methodologies for measuring functional C1-inhibitor in HAE plasma
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Objective: We validated three methods for measuring Functional C1-Esterase Inhibitor (FCl-INH) in KEDTA human plasma for Shire clinical studies: conventional chromogenic assay measuring residual C1-esterase activity, Complement C1s component (C1s) binding based Electrochemiluminescent (ECL) assay, and Factor Xlla (FXlla) binding based Enzyme linked immunosorbent assay (ELISA). We performed a side-by-side comparison of all three methods with consented pharmacodynamic (PD) samples from the SAHARA Phase 3, randomized, double-blind, placebo-controlled, two-period, three-sequence, partial crossover study that evaluated the efficacy and safety of subcutaneous administration of 2000 IU of C1 Esterase Inhibitor [Human] Liquid for Injection for the Prevention of Angioedema Attacks in Adolescents and Adults with Hereditary Angioedema (HAE).

Methods and materials: The validation assessments for the assays successfully fulfilled requirements for critical parameters including accuracy, precision, range of quantitation, selectivity, dilution linearity, hook effect, ruggedness and robustness, and analyte stability. The
O2

Assessment of C1-INH function – different methods, different results

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Background: We first reported in C1-INH-HAE type I patients under various treatment regimens, including replacement therapy, a C1-INH concentration > 40% (100% vs. 0.92, and 0.89 for C1s binding vs. Chromogenic, FXIIa binding vs. Chromogenic, and C1s binding vs. FXIIa binding methods, respectively. When the C1-INH PD profiles from three methods of these 15 patients who underwent different treatment regimens in the cross over study were superimposed, the profiles matched for individual patients with an average CV (Coefficient of Variation) of 25% across time-points.

Conclusion: In summary, the results generated in parallel from HAE plasma using three different C1-INH methods are comparable in spite of various principles followed in each of the assays. Since the ELISA results showed a significant correlation to the results from the conventional method, the in-house built ELISAs could serve as good alternates for measuring IC1-INH in HAE plasma with a better dynamic range of detection.

O3

Identification and characterization of large deletions in the SERPING1 gene

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Background: Hereditary angioedema (HAE) is a rare genetic disorder characterized by relapsing, on pruritic swelling of skin and submucosal tissue. HAE is mainly caused by mutation in the SERPING1 gene, resulting in a C1-inhibitor (C1-INH) deficit. Point mutations or small deletions are responsible for the majority of cases. Most of these mutations are private, underlining the high mutation rate for this particular gene. SERPING1 intronic regions have 17 sequences of repeating elements (Alu) which may represent hotspots for non-homologous recombination events, making the gene prone to deletions, insertions and duplications. In approximately 10% of the cases, HAE with C1-INH deficiency is caused by large gene rearrangement not detected by Sanger sequencing. To date, there are few reports with a detailed characterization of the large deletion breakpoints in the SERPING1 gene. The aim of this study was to describe the molecular mechanism leading to these genetic abnormalities.

Materials and methods: C1-INH deficiency was diagnosed in 228 patients from 200 families in the Complement Laboratory, Hôpital Georges Pompidou, Paris, between 1995 and 2018. Genetic screening was performed by amplification by polymerase chain reaction (PCR) of all exons and flanking splices sites of the SERPING1 gene. When no variant was detected by Sanger sequencing, Multiplex Ligation-dependent Probe Amplification (MLPA) was performed.

Results: MLPA has permitted to reveal the molecular mechanisms of the deficiencies in 16 unrelated patients. Partial and total SERPING1 deletions involving one to 6 exons were identified. 49 patients were identified within the 16 families with a mean age at onset of symptoms of 10.7 years, 32% of patients had experienced laryngeal oedema, 82% abdominal crisis and 75% were receiving a long-term prophylactic treatment. The most frequent large deletion observed in the cohort was the exon 4 deletion, found in 6 independent families. Breakpoints were characterized in 5 cases. There were a 2.2 kb and 3.2 kb deletions including the exon 4 which were detected in 2 unrelated alleles. About 8.6 kb and 14 kb of genomic DNA were missing in one case of the exons 4 to 6 deletions and exons 1 to 6 deletions respectively. Altogether, the 16 large deletions presented account for 8% of all SERPING1 mutant alleles investigated in French patients.

Conclusions: In summary, our study highlighted the heterogeneity of the large deletion in the SERPING1 gene associated with C1-inhibitor deficiency.

O4

Diagnosis of bradykinin-mediated angioedema in the emergency department: usefulness of the early biological workup

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crossover SAHARA study had two treatment periods (TP) 1 and 2 of 14 weeks each; the patients were randomized into three of the treatment groups A/B, B/A and A/A for the two periods with A being 200 IU C1-INH and B being Placebo. The seventeen PD time-points were 1a, 8a, 16a, 24a, 27/28a, 27/28b 48 b (in TP1) 1b, 8b, 16b, 24b, 26b, 28b 24b 48 b, 28b 72 b, 28b 96 b (in TP2), 1 week Post and 1 month Post. The PD samples collected from 15 consented patients (out of total 75) were used in this assessment.

Results: Both ELISAs offered a better dynamic range of detection compared to the chromogenic method. A significant correlation was observed between the results from three IC1-INH methods when 219 individual PD data points were correlated; the Pearson ‘R’ was 0.94, 0.92, 0.89 for C1s binding vs. Chromogenic, FXIIa binding vs. Chromogenic, and C1s binding vs. FXIIa binding methods, respectively. When the IC1-INH PD profiles from three methods of these 15 patients who underwent different treatment regimens in the cross over study were superimposed, the profiles matched for individual patients with an average CV (Coefficient of Variation) of 25% across time-points.

Conclusion: In summary, the results generated in parallel from HAE plasma using three different IC1-INH methods are comparable in spite of various principles followed in each of the assays. Since the ELISA results showed a significant correlation to the results from the conventional method, the in-house built ELISAs could serve as good alternates for measuring IC1-INH in HAE plasma with a better dynamic range of detection.
Background: Bradykinin-mediated angioedema (BK-AE) refers to acute and often recurrent swelling of subcutaneous and/or submucosal areas. This condition is typically caused by C1-inhibitor (C1Inh) deficiency or low bradykinin catabolism e.g. under angiotensin-converting enzyme (ACE) inhibitor treatment. Angioedema with normal C1Inh (AE-nC1Inh) is also related to high plasma bradykinin and is assumed to be very rare. We previously showed that activation of the kallikrein-kinin system occurs at the very early phase of BK-AE (i.e. during prodromes). Therefore, we sought to verify if the biological workup performed at admission in the emergency department could help to improve the diagnosis.

Materials and methods: We retrospectively analyzed all patients admitted during 12 months in our adult emergency department for acute angioedema without evidence of anaphylaxis. Plasma samples were analyzed for C1Inh function, spontaneous kallikrein activity and kinin catabolism. All patients have been examined by an allergologist.

Results: Twenty-six patients were admitted among which 4 were admitted during 12 months in our adult emergency department for acute angioedema with normal C1Inh without evidence of anaphylaxis. Plasma samples were analyzed for C1Inh function, spontaneous kallikrein activity and kinin catabolism. All patients have been examined by an allergologist.

Fourteen (41%) patients (9.9–44.7 nmol/min/ml; reference < 10.6 nmol/min/ml males) had low activity of one or more kininases. In 18 (82%) patients, low activity of one or more kininases was seen. The most prevalent kinase deficit was ACE (16 patients), followed by aminopeptidase P (5 patients), carboxypeptidase N (3 patients) and dipeptidylpeptidase IV (2 patients). Multiple kininases deficits were observed in 6 patients (4 with two deficits, 2 with three deficits). ACE inhibitors were taken by 4 patients, one of them having no detectable biological abnormality. No patient took gliclazide treatment. An association between high plasma kallikrein activity and one or more kininases deficits was seen in 7 (32%) patients. C1Inh concentrate or icatibant was administered to 13 patients with rapid clinical improvement. Kallikrein activity was analyzed after and before of C1Inh treatment in 5 patients. This showed high initial kinogenase activity in 4/5 patients with a 58–86% reduction thereafter.

Conclusions: Our experience shows that C1Inh deficiency is not the main cause of BK-AE, high kinogenase activity and/or low kinin catabolism seeming to account for most cases. By performing a biological workup during the acute phase, the diagnosis performance could be enhanced. Knowing that angioedema may lead to life-threatening situations, this issue is essential to improve patient care.

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O6 SGP 120 and contact system in hereditary angioedema. Diagnostic tool in HAE with normal C1-inhibitor?

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Mutations of C1inh are present in some patients with hereditary angioedema with normal C1-inhibitor (HAE-nC1inh), but the underlying disease mechanism remains unclear. There is no accepted biomarker for this disease. In 1989 while developing a new purification procedure for the human complement protein C21 we isolated a then unidentified plasma protein with a molecular weight of 120 Kd. It appeared related to the contact system of coagulation because incubating plasma at 4 °C in glass tubes led to cleavage of the protein. Plasma deficient in high-molecular-weight kininogen (HMWK) and kallikrein when treated similarly did not show cleavage of the protein. If a mixture of HMWK and prekallikrein was added to deficient plasma, the glycoprotein was cleaved after contact activation. In 1994, Xiao Ping Pu and colleagues isolated a similar protein from plasma. Using a partial cDNA sequence, they suggested that their protein had closely related sequence homologies to the heavy chains of the inter-α-trypsin inhibitor (ITI) superfamily. This group of 6 separate protease inhibitors is characterized by the presence of a common light chain, and is susceptible to proteolysis by the enzymes they inhibit. ITIH4 heavy chain 4 (ITIH4) is different from the other members of the group of ITI proteins because it is the only member that has no light chain and circulates as a free isoform. ITIH4 is also the only kallikrein sensitive protein among this group and has a molecular weight of 120 KD. We isolated our protein and identified it as ITIH4-heavy chain 4. We examined fragmentation of the protein in plasma from normal, patients with HAE types 1 and 2 and patients with HAE-nl-C1inh. ITIH4 is fragmented in almost all patients with HAE types 1 and 2 and is intact in almost all normals and patients with HAE-nl-C1inh. On incubation of patient plasma in plastic tubes at 4 °C the protein is intact in normal and cleaved in plasma of patients with HAE-nl-C1inh. We studied C1inh in the same samples. C1inh was at low levels or was cleaved in HAE type 1 or 2 disease. It was normal structurally and functionally in normal and patients with HAE-nl-C1inh. However on incubation in plastic at 4 °C the C1inh was cleaved and lost function in patients with HAE-nl-C1inh but not in normals. Antiserum to this protein is commercially available and we propose that study of its cleavage pattern provides a new means for the laboratory diagnosis of HAE-nl-C1inh.

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O6 Changes of complement parameters during erythema marginatum in patients with hereditary angioedema

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Background: Hereditary angioedema caused by deficiency of the C1-inhibitor protein of the complement system (C1-INH-HAE) is characterized by recurrent episodes of subcutaneous/submucosal edema which may be preceded by erythema marginatum (EM). Our aim was to understand the pathomechanism of EM symptoms by analyzing the parameters of the complement system.
Materials and methods: Eight C1-INH-HAE patients (1 man, 7 women, median age: 45.3 years), followed-up in Angioedema Reference Centers were investigated. These patients experienced EM on several occasions during their lifetime and blood samples were obtained during EM in all cases. The clinical characteristics of EMs were recorded by the Erysthema Marginatum Detailed Questionnaire. In the sera taken from symptom-free patients, during EM and during HAE attack periods, the following complement parameters were measured: C1q, C3, C4, C1-INH, Factor I, Factor B, Factor H, anti-Factor H, anti-C1q, and anti-C1-INH (IgG, IgA, IgM) antibodies. Total activity of the classical-, lectin- and alternative complement pathways and the activity of the C1-INH were investigated as well. Measurements were completed of complement activation products (C3a, C4a, C4d, C5a and sC5b-9) in EDTA plasma. All subjects consented to the study.

Results: We observed the following differences between samples taken during HAE attack vs. during EM: C3 (p = 0.0047), C4 (p = 0.0313), C1-INH concentration (p = 0.003), Factor B (p = 0.0391), and the activation product sC5b-9 (p = 0.0234) levels were significantly lower during HAE attack. Differences in samples during EM vs. symptom-free period were as follows: C3 (p = 0.049), C4 (p = 0.015), Factor B (p = 0.0084) proteins and C4a (p = 0.0114) activation product levels were significantly lower during EM. Levels of sC5b-9 were lower during EM compared to the symptom-free period, however it wasn't significant (p > 0.05) and further decrease were observed during HAE attack in sC5b-9 (p = 0.049) and C4d levels (p = 0.044).

Conclusions: According to our investigations, EM can be considered as the first phase of the HAE attack, as levels of C3, Factor B, C1-INH concentrations and C4 begin to decrease during the prodromal symptom, and this trend continued during HAE attacks. Nevertheless, more patients and further investigations of the kinin-kallikrein, coagulation and fibrinolytic systems are needed for better understanding of the pathomechanism of EM. A new, individualized therapy, administered during EM to prevent the development of HAE attacks seems to be thoroughly grounded. This study was supported by OTKA K124557 and the Pharming Group NV.

O7 Clinical and genetic characteristics of patients with hereditary angioedema at a large tertiary care hospital in Saudi Arabia
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Background: Hereditary Angioedema (HAE) is a rare autosomal dominant disorder characterized by potentially fatal swelling of the larynx. In addition, abdominal angioedema can lead to misdiagnosis and needless surgical intervention, along with narcotic dependence. The prevalence of HAE is estimated at roughly 1:50,000 individuals, however, the overall epidemiological data for this disorder remains inaccurate. The main gene linked to HAE pathogenesis is SERPING1, the gene that encodes for C1-INH. Less commonly observed are mutations in F12, the gene that encodes the Coagulation Factor XII (Hageman Factor). Very recently two other genes have also been described: PLG and ANGPT1

Objectives: We sought to examine the clinical features as well as the molecular genetic defects observed in HAE in the Saudi population.

Methods: Thirty-six (36) patients with a diagnosis of HAE, who were being followed at the Immunology clinics at King Faisal Specialist Hospital (both Riyadh and Jeddah) were included in the study after informed consent. This study is RAC approved. (RAC # 2080 025)

Results: HAE diagnosis was based on the classic clinical presentation of angioedema without urticaria as well as characteristic lab features. 38.8% of the patients were males and the mean age was 31.08 years. Initial molecular screening was performed for the SERPING1 gene. For cases that were negative for this gene, primers were then designed against all coding regions of F12, PLG and ANGPT1 for PCR analysis.

Twenty-six (26) patients were found to have mutations in SERPING1 gene, of which the most common was NM_000062:c.1397G>A:p.R466H (Table 1). In addition, one novel mutation (NM_000062:c.1202T>A:p.I401N) was uncovered in a single Saudi family. All negative cases have been screened for F12, PLG and ANGPT1 gene mutations, but thus far, no such mutations have been found.

Conclusions: Molecular analysis of this HAE cohort has revealed consistent autosomal dominant inheritance and the presence of SERPING1 mutations in > 50% of patients. Although some mutations have been found across multiple families, R466H is the only mutation which appears to dominate the HAE genetic landscape in the region. The lack of observed mutations in F12, PLG and ANGPT1 suggest that there are no prevalent founder mutations in these genes in the local population. Unsolved pedigrees are being prioritized for whole exome sequencing.

O8 D-Dimer and C-Reactive protein in urticaria and angioedema at the Emergency Room
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Background: D-Dimer (D–D) and C-Reactive protein (CRP) have been reported to be increased and correlate with severity of symptoms in urticaria, which is histamine-mediated and may occur with or without angioedema (AE), as well as in histaminergic or bradikynin-induced, primary AE [1–3]. These findings may elicit the suspicion of inflammatory pattern and deep venous thrombosis (DVT) at the Emergency Room (ER), thus prompting to unnecessary and harmful procedures.

Objectives: To assess the prevalence of high D-Dimer and CRP levels in patients with acute urticaria and/or AE and to investigate the possible presence of DVT in subjects with increased plasma D–D concentrations.

Materials and methods: Thirty-five patients (10 male, 25 female; age: mean = 45.51 ± 17.65 years SD, range 15–80 years) admitted to the ER of Padua University Hospital for urticaria/angioedema (urticaria without AE 19 patients, urticaria with AE 7 patients) or primary AE (9 patients) underwent blood sampling for D-dimer, CRP and other factors related to the coagulation pathway. In all the subjects with D–D values higher than normal (referential range of Padua Hospital Laboratory: 0–250 μg/L), a lower limbs venous ultrasonography was
Attacks. Although frequently reported, prodromes have not been adequately investigated, due to lack of systematic tools. HAE prodromes should be better defined and needs a dependable instrument as a PRO metric.

Methods: First phase of the study consisted of personal and health data acquired from HAE patients. Out of 233 patients, 197 (84.5%) responded to a preliminary questionnaire, inquiring if they ever had a prodrome, and if they can predict an oncoming attack by having a prodrome. Mean age was 36.7 ± 20.3 years (Y, range: 2–80, females 42.5%). Mean age of onset was 10.7 ± 10.3Y, age of diagnosis was 15.3 ± 15.1Y. Nearly 62% of this group were diagnosed by age 10 (M > F, p < 0.05).

Preliminary questionnaire was constructed, based on the literature and investigator’s experience. Six new instruments were used to reach a robust evaluation scale. Data from the preliminary interviews was used to test the instrument’s construct and internal validity. In the 2nd phase, patients were interviewed to obtain data on prodromes and attacks. After final refinement of the instrument, a retrospective and prospective data were obtained and statistical analysis was performed to study prodromes and attacks associations.

Results: In the 1st phase, 165/197 (84%) reported ever having a prodrome, 143/165 (87%) could predict an oncoming attack by experiencing a prodrome. There was a significant correlation between the perception of prodrome and ability to predict an oncoming attack (p < 0.01, r = 0.79). The internal validity of the new questionnaire, constructed based on the preliminary study, was high, as well as its internal reliability (Cronbach’s α = 0.70 to 0.96). Prodromes and attacks, analyzed for each body system cluster, were highly correlated (r = 0.63, p < 0.01). In the 2nd phase, 66 patients (33.5%) completed a questionnaire that covers ‘clusters’ of body systems, affected by both events. Differences between the clinical dimensions of prodromes (i.e. pain, severity, dysfunction, duration), were evaluated by one-way MANOVA with repeated measurements, and shown to be segregated from attacks on all dimensions [F (4, 56) = 45.7, p < 0.001, Eta² = 0.77].

Conclusions: A questionnaire-based PRO instrument for the evaluation of HAE prodromes and attacks was designed and tested. The high internal validity and reliability makes it useful for studying clinical expressions of HAE. A prodrome/attack evaluation instrument can be useful in the diagnosis of oncoming attacks and timing of medical interventions.

O10
Psychosocial burden of hereditary angioedema in a Canadian cohort
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Background: Patients with chronic diseases have increased levels of anxiety, depression and stress – all of which may affect quality of life (QoL) [1, 2]. The unpredictability of swelling in hereditary angioedema (HAE) results in significant anxiety, depression and impaired QoL [3]. The availability of HAE-specific therapies and use of prophylaxis has been associated with an improvement in QoL; however, literature discussing specific aspects of the psychosocial burden of HAE is lacking.

Aim: To assess prevalence of health-related (HR) depression, anxiety and stress

Methods: Seventeen patients, ≥ 18 years, with a confirmed diagnosis of HAE type 1 or 2 (Table 1), completed the following self-reported questionnaires: SF36v2 (generic HRQoL), AE-QoL (disease-specific HRQoL) and both the Depression, Anxiety, Stress Scale (DASS) and the Diagnostic and Statistical Manual of Mental Disorders (DSM5) cross cutting measures (identify/differentiate between HR depression, stress, anxiety and related symptoms) [4–7]. This study was approved by the University of Calgary Ethics Board (REB# 17-0542).

Results: 100% (17/17) of participants reported increased levels of HR fatigue and fear, as well as decreased overall HRQoL. The majority of respondents reported decreased mood (88%) and increased anger (82%) concerning their disease. Disturbed sleep was reported by 76% (13/17) of participants. Most participants reported greater than average amounts of HR stress (76%, 13/17), anxiety (76%, 13/17), and depression (70.6%, 12/17). Somatic symptoms were reported by 64% (11/17) of participants, including feeling that their illness was...
Table 1  Sample characteristics

| Sample (n)     | 17 |
|----------------|----|
| Gender (n, %)  |
| Male           | 4  (24%) |
| Female         | 13 (76%) |
| Age, years (mean, range) | 43 (20–63) |
| Diagnosis (n, %) |
| Type 1         | 11 (64.7%) |
| Type 2         | 6  (35.3%) |
| Treatment protocol (n, %) |
| Prophylaxis    | 11 (64.7%) |
| On-Demand     | 6  (35.3%) |
| Treatment administration (n, %) |
| Self-administered treatment | 16 (94.1%) |
| Provider administered treatment | 1 (5.9%) |

Fig. 1  Trends in Psychosocial and HRQoL Scores

Fig. 2  HAE cohort scores compared to Canadian normative scores (SF36v2)

not taken seriously by others (Fig. 1). Mean scores on all domains of the SF36v2 were significantly lower than Canadian normative data for the entire sample (p < 0.05 for all). Social and physical function, as well as emotional and physical role scores of the SF36v2 were the most impacted (Fig. 2). Female participants tended to have lower scores than male patients in most SF36v2 domains and had significantly higher HAE-related fears (AEQoL; t(5.6) = −2.7, p = 0.035) and significantly more HAE-related stress (DASS; t(15) = −2.2, p = 0.04) than male patients.

Conclusion: This study of Canadian patients demonstrates that HAE negatively influences HRQoL as measured by the AE-QoL. More patients report depressed mood, increased stress, anxiety, anger, and sleep disturbances than not. Patients with HAE have significantly lower emotional and physical functioning than the general population; women with HAE appear to be particularly impacted. This study offers specific, valuable insight into the psychosocial burden of HAE. Future evaluation should include larger samples and further investigation of gender differences.

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O11  Angioedema and urticaria at the Emergency Room: epidemiology and clinical management in a tertiary care center in Italy during the decade 2009–2018

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Background: Primary angioedema (PAE) and urticaria-angioedema syndrome (UA) are common complaints and a frequent cause of hospital admission. However, epidemiological data are still limited, with a few studies published on this topic.

Objectives: To assess the overall impact of PAE and UA on the Emergency Room (ER) of the General Hospital - University of Padua - Italy (GHUPD) and the fluctuations of the trend in a decade.

Methods: We selected 73 ICD9-CM nosologic codes potentially related to allergic symptoms. Thereafter, we read all the 12.897 discharging reports from the ER of the GHUPD associated to these codes in the decade 2009–2018, to confirm the diagnoses and collect detailed information on clinical management and outcome of each admission. All the data were then analyzed by SAS 9.2 program for Windows and statistical analyses performed by χ2 test and Fisher’s exact test. The study was approved by the local Ethical Committee (protocol number 83901/AO/16)

Results: 79.75% of discharging codes were consistent with the clinical features as assessed by reports’ analysis, whilst the remaining 20.25% showed to be wrong. Allergic symptoms accounted for 1.2% of the overall ER admissions (10286/859224, with an increasing trend in their prevalence (2009:1.08% → 2018:1.47%). UA was the main cause of admission (46.43%), followed by asthma (21.58%) and PAE (12.37%) (Fig. 1). Female gender was prevalent for allergy in general (58%), as
well as for UA (60.86%) and PEA (53.87%), whereas women represented 49.35% of total admissions (p < 0.001 for both groups) which showed also an inverse age distribution, with a prevalence of patients older than 60 years. Foods and medicines were the most common causal factor, with use of ACE-inhibitors accounting for 18.84% of all casess (Fig. 2). However, no clear etiology was identified in 72.48% of UA and in 53.26% of PEA. Angioedema showed more severe clinical features than UA, with greater attribution of red/yellow codes (54.12% vs 26.79%, p < 0.001), higher prevalence of mouth involvement (42.9% vs 7.4%, p < 0.001) and more frequent epinephrine administration (3.6% vs 1.01%, p < 0.001). Moreover, AE accounted for all the 32 hospitalizations. Finally, 3 cases of death were reported in the decade, one of whom for UA involving upper airways.

Conclusions: Our data demonstrate the significant impact of AE and UA on the ER, in terms of both patient numbers and resource use. Although a clear etiology is not detectable in most of cases, ACE-inhibitors are frequently associated with AE, which is clearly more severe than UA.

O12 Hereditary angioedema with a specific mutation in the plasminogen gene in 18 families

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Recently a new type of hereditary angioedema, hereditary angioedema with a specific plasminogen gene mutation (HAE-PLG) has been identified by whole exome sequencing and family studies. Aim of the present study was to provide more information about clinical features of HAE-PLG. A total of 18 German families with HAE-PLG were studied. In all families the missense mutation c.988A > G (p.Lys330Glu; K330E) leading to an amino acid exchange in the kringle 3 domain of plasminogen co-segregated with the clinical symptoms of HAE-PLG. None of the investigated family members had HAE symptoms but did not carry this mutation. The size of the families ranged from 2 to 15 affected family members. On average 5 members per family were affected with HAE-PLG or identified as symptom-free mutation carriers. Among 101 relatives in the 18 families there were 96 clinically affected patients and 5 symptom-free individuals with the specific mutation. Sixty-two of the 101 relatives were women (61.4%) and 39 men (38.6%). Two patients from one family had HAE-PLG combined with HAE-C1-INH and were excluded from the symptom evaluation. The remaining group comprised 94 patients who were studied for clinical symptoms of HAE-PLG. The mean age of onset of clinical signs of HAE-PLG was 32.1 ± 17.7 years ranging from 4 to 74 years. The mean onset in 61/94 women was 29.2 ± 16.4 years and in 33/94 men 37.4 ± 19.0 years. Seventy-one of 94 patients (75.57%) had lip or facial swellings and 7/94 (7.4%) swellings of the extremities. Twenty-six of 94 patients (27.7%) had abdominal attacks. One patient reported on genital swellings. Sixty-nine of 94 patients (73.4%) had a total of 1,368 tongue swellings (range 1 to 626). Thirteen of 94 patients (13.8%) had exclusively recurrent tongue swellings (range 1 to 150) and no other clinical signs. Three women died by asphyxiation due to a tongue swelling with upper airway obstruction, one woman at age 25, another one at age 36 and the third woman at age 47. The latter woman had 160 tongue swellings before the fatal upper airway obstruction. One man asphyxiated at age 75 due to a tongue swelling. Lip and facial swelling and potentially life-threatening tongue swellings are the most frequent and important symptoms of HAE-PLG.

O13 Hereditary angioedema with C1-INH deficiency in 96 Brazilian children

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There are scarce data about Hereditary Angioedema (HAE) with C1-INH deficiency in pediatric patients. Although symptoms of HAE begin early in life, diagnosis of the disease is delayed in this period of life. In addition, several new therapies are still restricted to older ages. Considering the restricted access to diagnosis and therapy in developing countries, we evaluated clinical and laboratory characteristics, and therapy of pediatric patients from Brazilian reference centers.

Methods: Medical records of HAE patients aged less than 18 years old, whose diagnosis was confirmed by quantitative and/or functional C1-INH, were included. The following data were collected: age of diagnosis, age at onset of symptoms, prodromes, symptoms, triggering factors, diagnostic tests and treatment. Descriptive statistical analysis was performed. The study was approved by ethical committee.

Results: 96 participants (52 M:44 F) from 17 reference centers on HAE were included: 50% from southeast, 23% from northeast, 16% from midwest, and 11% from south of Brazil. Family history was present in
72/96 patients. Twenty percent of the patients were asymptomatic. Age at onset of symptoms was < 1 year of age in 27%; 1–5 years in 45%; 6–10 years in 21% and 11 to < 18 years in 7%. Median age at diagnosis was 7 years old. Prodromes were reported among 25% of the patients: local burning in 10%; serpiginous erythema in 7%; fatigue and nausea in 2% each and irritability in 5.2%. Angioedema attacks affected: face (5%); lips (13%); tongue (11%); eyelid (6%); ear (2%); neck (1%); hands (19%); arms (15%); legs (5%); genitals (7%); upper airways (3%) epigastric pain and/or abdominal pain (55%). One fourth of the patients had no trigger factor identified. Severity of attacks were mild in 32%; moderate in 40% and severe in 26%. Misdiagnosis included: allergy 16% and helminthiasis 5%. Previous ER visits were reported by 56.3%. Surgical intervention (appendectomy) prior to HAE diagnosis occurred in 4%. Long term prophylaxis was introduced in 54% (53/96): tranexamic acid in 76% (40/53); danazol in 13% (7/53) and oxandrolone in 11% (6/53). Androgens were used in 4/13 under 12 years of age. Short-term prophylaxis was prescribed in 10/96 (tranexamic acid 6/10; danazol 3/10 and plasma derived C1-INH 1/10). 32/96 patients received specific treatment of acute attacks, including icatibant to 4/32; fresh frozen plasma to 15/32; plasma derived C1-INH to 11/32 and tranexamic acid to 13/32 patients.

Conclusions: Although more than 80% of the patients had family history and several members affected, there was a delay in diagnosis. Abdominal pain and surgical interventions were less frequent than reported in adulthood. Attenuated androgens were prescribed for pediatric patients, probably due to restricted access to on demand therapy. Only recently icatibant was licensed for use in children, nevertheless it had been previously used in our population. Educational programs should focus on Pediatricians, aiming at reduced delaying diagnosis and providing appropriate therapy.

Table 1 Results showing significant determinants of breakthrough attacks after short-term prophylaxis in HAE subjects after dental procedures

| Determinant                              | Odds ratio | LCL  | UCL  | p-value   |
|------------------------------------------|------------|------|------|-----------|
| Age                                      | 0.94       | 0.90 | 0.99 | 0.03      |
| H/O procedural-angioedema (yes) vs (no)  | 12.62      | 3.42 | 46.53| 0.0001    |
| Asthma (yes) vs Asthma (no)              | 3.57       | 1.04 | 12.21| 0.042     |

Note: Use of corticosteroids, supplementary oxygen, phenytoin and immunosuppressants (i.e., medications for comorbidities) were significantly associated with breakthrough attacks but were determined to have multicollinearity with history of procedure-related angioedema and therefore, were not included in the multivariate analysis.

Conclusions: Young age (< 26), history of post-procedural HAE attacks and comorbid asthma are significant risk factors for breakthrough HAE attacks following dental procedures despite STP and LTP. The significant association of post-dental procedure HAE breakthrough attacks with asthma implies that HAE subjects with comorbid asthma may represent a distinct HAE phenotype which requires further investigation.
A SERPING1 variant that causes C1-inhibitor deficiency without hereditary angioedema

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More than 400 mutations (around 25% de novo) in SERPING1 have been described to cause functional C1-inhibitor (C1-INH) deficiency and hereditary angioedema (HAE). Angioedema symptoms and C1-INH deficiency co-segregate within the families with autosomal dominant inheritance. Two promoter and two structural SERPING1 variants escape this rule. These four mutations represent a recessive inheritance. Two promoter and two structural variants that cause recessive SERPING1 apply to the other C1-INH alleles.

Here we describe variant g.22006G > A (p.Arg494Gln) in homozygosity. Of the 120 subjects who received STP, a total of 91 subjects reported receiving C1Inh [human] as STP (which includes 79 subjects exclusively on C1Inh [human], 1 subject on C1Inh + C1Inh, 7 on C1Inh + C1Inh + C1Inh, 3 on Other + C1Inh, and 1 on Other + C1Inhibant + C1Inh). 10 received C1Inh only as STP, 1 received Ecallantide only, and 1 received C1Inh only. 17 subjects reported receiving other STP agent (choices were “C1Inh, 130 subjects did not receive STP.

O16

Activation of complement MASP-3 in healthy donors and in patients with C1-inhibitor deficiency

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Activation of the lectin pathway of complement is initiated by mannose-binding lectin (MBL)-associated serine proteases 1 and 2 (MASP-1 and MASP-2). MASP-1 and MASP-2 circulate in the blood as zymogens in complex with pattern recognition molecules (PRMs), such as MBL, other collectins, and ficolins. The third serine protease of the lectin pathway (MASP-3), which is also complexed with PRMs, was shown to be the major physiological activator of pro-factor D (pro-FD) in the blood, linking the alternative and the lectin complement pathways. We have demonstrated earlier that only activated MASP-3 is capable of converting pro-FD to factor D (FD), and indeed the major form of MASP-3 in the blood is the active form. The activation mechanism of MASP-3, however, remains unclear. In vitro MASP-1 can activate MASP-3, and C1-inhibitor is the major physiological regulator of MASP-1. We hypothesized that if MASP-1 is the physiological activator of MASP-3 then individuals with low C1-inhibitor levels would exhibit altered MASP-3 activation.

The activation state of endogenous MASP-3 was detected by Western blot, whereas in other experiments fluorescently labeled recombinant MASP-3 variants were used. We found that a significant portion of full-length, labeled MASP-3 became "activated" (cleaved) in hirudin-plasma in the matter of hours even when the inactive S664A variant was used. The activation was less efficient, but still occurred, when the N-terminally truncated catalytic fragment was used. On the other hand, we found that the ratio of active MASP-3 to type I HAE patients was virtually identical with that in healthy individuals, namely 82 ± 3%, versus 81 ± 4%. This indicates that a protease other than MASP-1 is responsible for the activation of MASP-3. To confirm this assumption we monitored the cleavage of labeled MASP-3 in the presence or absence of a MASP-1-specific inhibitor. Again, no difference was observed.

In conclusion, our results imply that a protease is present in the blood that converts MASP-3 to the active form. The activation is not autoactivation because the inactive variant got cleaved as well. Activation is more pronounced with the full length protein implying that binding of MASP-3 to PRMs might be necessary for efficient activation. Activation of MASP-3 is probably carried out by a protease not inhibited by C1-inhibitor.

O17

Simultaneous determination of human plasma serine proteases complexed with C1-inhibitor in vivo

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No HAE. The variant is located in a region of the molecule that is critical for regulating SERPIN conformational state.

The index cases in each family were identified due to urticaria symptoms that prompted measurements of C1-INH and C4. In none of the four families there was evidence for inherited angioedema without wheals while inherited C1-INH and C4 deficiency were clearly present. Inhibitory function of C1-INH against C1s, kallikrein and FXIIa was below 50% of normal. Repeated measurements of complement parameters in C1-INH deficiency carriers, showed a degree of variability, which is never detected in typical C1-INH-HAE. When these subjects underwent danazol treatment, complement parameters rapidly normalized. All C1-INH deficient subjects in the four families, carried the mutation g.22006G > A (p.Arg494Gln) in SERPING1. When the mutant protein was expressed in a murine hepatoma cell line, analysis of the supernatant failed to detect the protein, which was instead abundant in the insoluble fraction of cell lysate. We hypothesize that intracellular accumulation is due to polymer formation.

Our findings indicate that p.Arg494Gln-C1-INH allows protein synthesis, but impaired cytoplasmic secretion. The same condition seems to apply to the other SERPING1 structural variants that cause recessive forms of HAE. No homozygous p.Arg494Gln has ever been described and we cannot conclude that also this mutant leads to recessive HAE. However, variability of C1-INH plasma levels in this as in other variants that are clinically silent in heterozygous presentation, suggests a lower degree of impairment in the synthesis of functional C1-INH. Whether this is due to the mutant contributing to C1-INH function in plasma or to the mutant that has reduced trans-inhibition effect on the wild type allele remains to be elucidated.

The study was approved by Milano Area 1 Ethics Board, approval number 11846/2017. Informed consent to publish has been obtained from the patients.

Activation of the lectin pathway of complement is initiated by mannose-binding lectin (MBL)-associated serine proteases 1 and 2 (MASP-1 and MASP-2). MASP-1 and MASP-2 circulate in the blood as zymogens in complex with pattern recognition molecules (PRMs), such as MBL, other collectins, and ficolins. The third serine protease of the lectin pathway (MASP-3), which is also complexed with PRMs, was shown to be the major physiological activator of pro-factor D (pro-FD) in the blood, linking the alternative and the lectin complement pathways. We have demonstrated earlier that only activated MASP-3 is capable of converting pro-FD to factor D (FD), and indeed the major form of MASP-3 in the blood is the active form. The activation mechanism of MASP-3, however, remains unclear. In vitro MASP-1 can activate MASP-3, and C1-inhibitor is the major physiological regulator of MASP-1. We hypothesized that if MASP-1 is the physiological activator of MASP-3 then individuals with low C1-inhibitor levels would exhibit altered MASP-3 activation.

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Simultaneous determination of human plasma serine proteases complexed with C1-inhibitor in vivo

Zsófia Jandrasics1,2, Erika Kajdácsi1,2, Nőra Veszeli2,4, Vera Makó4, Anna Koncz1, Kinga Viktória Kőhalmi1, László Cervenak1, Péter Gál1, József Dobó1, Steven de Maat5, Coen Maas5, Henriette Farkas1,2, Lilian Varga1,2,3, Marieke H. Savelkoul1,6, Corrie J. Aalberse1,6, Mikael L. Kruis4, Simon A. Smeets1,6, Steven de Maat5, Coen Maas5, Henriette Farkas1,2, Lilian Varga1,2,3, Marieke H. Savelkoul1,6, Corrie J. Aalberse1,6, Mikael L. Kruis4, Simon A. Smeets1,6

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C1-inhibitor (C1-INH) is an important regulator of complement, coagulation, fibrinolytic and contact systems. The quantity of enzyme/C1-INH complexes in the blood may well be proportional to the level of the in vivo activation of these four cascade-like plasma enzyme systems. Parallel determination of C1-INH-containing activation complexes would be important to understand the role of C1-INH in the regulation of plasma enzyme cascades in diseases like hereditary angioedema due to C1-INH deficiency (C1-INH-HAE). We developed 10 in-house ELISAs for measuring the complexes of C1-INH and the following proteases: Factor Xla (FXla), Factor XIIa (FXIIa), C1r, C1s, thrombin (TR), MASP1 (M1), MASP2 (M2), kallikrein (KK), and for measuring C1-INH concentration and activity. We measured the levels of the complexes in EDTA plasma from 6 healthy controls, from 5 and 5 C1-INH-HAE patients type I and type II in remission, and from 5 C1-INH-HAE patients during attack. We also measured the levels of these complexes in the blood samples taken from one C1-INH-HAE patient, during a subcutaneous attack from the start of the prodromal symptoms during the edematous attack till the spontaneous termination (attack follow up).

There was no significant difference in total level of the measured enzyme/C1-INH complexes with the exception of KK- and M1/C1-INH complexes between the controls and the patients, and of FXla- and M2/C1-INH complexes between C1-INH-HAE type I and II patient. When generated a ratio from the amount of the enzyme/C1-INH complex divided with C1-INH activity we found elevated FXla-, C1r-, KK-, M1-, C1s- and TR/C1-INH complex amounts in patients compared to controls. We found no significant differences among the amount of the complexes compared in attack and remission. During the attack follow up the kinetics of the change of the complexes' amount is very fast.

From this results we can conclude that the change of the C1-INH concentration, activity or the amount of the measured complex levels can not totally explain when and why a patient will have an attack. The pathomechanism of the attack formation may have other important factors which are till unknown. Maybe the local C1-INH production (for example generated by endothelial cells) also take part in the attack formation. The fast changes in the amount of enzyme/C1-INH complexes during the follow up study may reveal that we need a very strict timing if we want to make a good comparison between the amounts of the complexes during attacks.

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O18 Bradykinin, LPS and MASP-1 synergistically regulate endothelial permeability

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Background: Bradykinin is considered as the major mediator of edema in HAE, but we have only limited knowledge about the triggering factors and the initiation of the attacks. Bacterial infections are considered as one of the risk factors for the onset of attacks and during these the complement system becomes activated. We have previously demonstrated that the most abundant enzyme of the complement lectin pathway, mannan-binding lectin-associated serine protease 1 (MASP-1), which is naturally inhibited by C1-inhibitor, directly increases endothelial permeability and activates the expression of permeability related genes. Therefore, we wanted to investigate how bacterial LPS, bradykinin and MASP-1 can interact with one another to influence endothelial permeability.

Methods: We measured the mRNA level of BDKRB1 and 2 with qPCR and found that MASP-1 upregulated the level of BDKRB2, while LPS increased the expression of both bradykinin receptors. In concert with this, the MASP-1 or LPS pretreated cells showed significantly greater Ca²⁺-mobilization to bradykinin than those that were not pretreated (measured with fluorescence microscopy).

LPS also induced the mRNA level of PAR2, which is a receptor of MASP-1 on endothelial cells, and we demonstrated that MASP-1 elicited greater Ca²⁺-mobilization after LPS pretreatment. To measure the endothelial permeability, we used the modified X-permeability assay. The LPS pretreatment could significantly increase endothelial permeability in response to MASP-1.

Conclusion: Our findings highlight that significant interaction can occur amongst endothelial cell activators (between MASP-1 and LPS, LPS and bradykinin and MASP-1 and bradykinin) in the regulation of endothelial cell permeability. These synergistic interactions may give us a more detailed picture on the pathogenesis of HAE and highlight the importance of MASP-1 as a potential additional factor triggering edematous attacks.

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is a significant antagonist of the B1 receptor expressed in the vein, whereas PHA-022121 is not.  

**Conclusions:** The human umbilical vein bioassay confirms and extends the characterization of PHA-022121 in a tissue system: the drug is a competitive, reversible B2 receptor antagonist that is considerably more potent and selective than icatibant.

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**O20**

Changes of coagulation parameters during erythema marginatum in patients with hereditary angioedema  

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**Background:** Hereditary angioedema caused by deficiency of the C1-inhibitor protein of the complement system (C1-INH-HAE) is characterized by recurrent episodes of subcutaneous/submucosal edema which may be preceded by erythema marginatum (EM) as a prodromal symptom. Our aim was to analyze the changes in parameters of the coagulation and fibrinolytic systems during the development of EM and HAE attack.

**Materials and methods:** Eight C1-INH-HAE patients (1 man, 7 women, median age: 45.3 years), followed-up in Angioedema Reference Center, had a personal history of EM and were investigated. These patients experienced EM on several occasions during their lifetime and blood samples were obtained during EM or HAE attack.

**Results:**  
C1-inhibitor (C1Inh) function, spontaneous kallikrein activity, kininogen, and high molecular kininogen (HK) were determined in citrate plasma samples from 4 individuals during symptom-free period, during EM and during HAE attack. The basic coagulation tests (PT, PTT, INR) were normal. D-dimer activity was low in all subjects.

**Conclusions:**  
Our study confirmed that the development of HAE attacks seems to be thoroughly grounded. This study was supported by OTKA K124557, EFP-3.6-VEKOP-16-2017-00009 and the Pharming Group NV.

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**O21**

Plasminogen missense mutation p.Lys330Glu: altered plasminogen glycoforms type I & II and activation susceptibility  

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**Background:** Hereditary angioedema with normal C1-inhibitor (nC1inh-HAE) may be associated to specific mutations, e.g. F12 and ANGPT1 gene variants. Recently the new variant c.988A>G altering the plasminogen gene (PLG, NM_000301.3) in exon 9 was associated to nC1inh-HAE. This variant led to the missense mutation p.Lys330Glu (K330E) in the plasminogen (PLG) kringle 3 domain. PLG has two glycoforms, type I PLG (~33% of circulating PLG), which is O-glycosylated at Thr346 and N-glycosylated at Asn289 and type II PLG, with an O-glycosylation on Thr346 and that comprises ~67% of circulating forms. Type I PLG has 10-fold less affinity for cells compared to the type II (Gonzalez-Gronow et al., 2002). Conversely, type I PLG appears to function more efficiently than type II in the degradation of fibrin clots. We aimed to investigate biological phenotype of the variant pertaining to angioedema.

**Materials and methods:** Citrate plasma samples from 4 individuals from a Greek family were harvested. We investigated the index case, a 57-year old man, homozygous carrier and presenting with severe angioedema, her two healthy daughters and her healthy granddaughter, carriers of the heterozygous mutation. C1-inhibitor (C1Inh) function, spontaneous kallikrein activity, kinin catabolism and kininogen cleavage were investigated. PLG activation and its circulating species, with clot lysis time and basic coagulation tests were carried out.

**Results:** All individuals displayed normal C1Inh function and kallikrein activity. High molecular kininogen (HK) was found cleaved (CHK), at nearly 17% in homozygous patient in samples out of angioedema attack (reference 1–15.6%). During the active period, CHK abundance was found nearly 25% and 21%, 5 h and 24 h, after the attack, respectively. All subjects had normal activity for carboxypeptidase N, dipetidylpeptidase IV and angiotensin converting enzyme. But aminopeptidase P activity was low in all subjects. The basic coagulation tests (PT, PTT, INR) were normal.

Interestingly, the anti-PLG immunoblot showed that the glycoform profiles are reversed in the homozygous patient, with ~60% of type I PLG. The other subjects had also altered glycoform patterns, with two bands of approximately equal intensity. PLG activation was found enhanced by PLG activators urokinase (UK) and streptokinase (SK), more than by tissue plasminogen activator (tPA).

**Conclusion:** Different glycosylation patterns of circulating PLG have important biological impact. The enhanced PLG activation susceptibility may be related to altered patterns, in agreement with the observation of Takada et al. (Takada and Takada, 1983) showing that type I PLG is more susceptible to activation by UK or SK than type II PLG. These observations are congruent with an increased cleaved HK, a situation likely to be associated with angioedema development.

**Consent to publish:** Written, informed consent for publication was obtained from the patient (or parent/guardian for patients under 16)
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O22

Who and when: the analysis of the molecular mechanisms of C1-inhibitor deficiency induced angioedema for the best therapeutic choice

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Angioedema (AE) due to inherited or acquired deficiencies of C1-inhibitor (C1-INH) is characterized by self-limiting localized swelling of deeper layers of the skin or submucosal tissues, becoming particularly life threatening if it occurs in the upper respiratory tract. C1-INH regulates the release of bradykinin which can enhance permeability of post-capillary venules interacting with two different type of receptors, B1 and B2. Since there are different therapeutic options we investigate the molecular mechanisms that lead to the attacks, in particular the ability of C1-INH and bradykinin receptor antagonists to block the ongoing permeabilizing effect of the acute attack plasma collected from patients, in order to identify the most effective therapeutic strategy.

For this purpose, we used a transwell in vitro model with a filter covered by primary human endothelial cells (EC), in the upper chamber we add the fluorescent-BSA and the stimuli and the BSA leaked into the lower chamber was evaluated using a Fluorescence reader. We found that the presence of C1-INH (BehrinertP) was able to block the endothelial permeability induced by the plasma collected from patients during attack (APL) in the majority of the patients. To mimic the in vivo situation we stimulated the EC with the APL for 30 min and then the SN was collected and used to stimulate the ECs in the transwell model. In that case the inhibition of the leakage by C1INH was not seen in all the patients. This observation was further confirmed by using the plasma collected from patients before and 1 h after the clinical treatment with C1-INH. The addition of C1-inhibitor from 10 min before the addition of APL and till 10 min after the addition of APL resulted efficient in the inhibition of endothelial leakage, while the use of C1-inhibitor 20 min after the contact between the APL and the EC resulted completely inefficient. Then we added the antagonist of B1 BK receptor (R054) or the antagonist of B2 BK receptor (Icatibant) and they both resulted able to block the permeabilizing effect of the SN. This inhibition resulted even stronger using a combination of both antagonists.

On the basis of these results, we conclude that the inhibition of endothelial leakage induced by APL stimulation by C1-INH indicates the involvement of that molecule in controlling the onset of AE attacks, although the inability of C1-INH to completely block the permeabilizing effect of the SN indicates that after the activation of the cells there are other molecules involved. The most plausible is BK but also other related metabolites can interact with specific receptors. The effectiveness of the treatment seems to be correlated with the time of treatment, as soon is treated the patient C1-INH is perfectly working but after the activation of the kinin system antagonists of kinins receptors seem to be more efficient in reducing the vascular permeability.

O23

PHA-022121, the first-in-class orally active bradykinin receptor B2 antagonist for on-demand and prophylactic treatment of HAE

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Patients are eagerly awaiting next generation treatments for hereditary angioedema (HAE), asking for oral treatment to replace the burden of current injectables. Pharvaris is developing PHA-022121 as a first-in-class novel proprietary small-molecule antagonist of the B2 receptor, for oral on-demand treatment of acute HAE attacks and for prophylactic prevention of attacks. PHA-022121 is entering clinical phase I studies in Q2 2019, was optimized and developed by Pharvaris, a company founded by the team which successfully developed Firazyr the only approved and widely used B2 antagonist for on demand treatment. Based on preclinical studies, PHA-022121 demonstrates excellent drug-like physicochemical properties, primary activity, oral bioavailability and metabolic stability. PHA-022121 shows sub-nanomolar potency in a calcium mobilization assay using recombinant human B2 receptors expressed in a mammalian cell line (0.15 nM) and at endogenous human B2 receptors in the human umbilical vein model (pA2 value corresponding to 0.35 nM). The compound is several thousand-fold selective for the B2 receptor versus the B1 receptor as well as against 130 other molecular targets (including GPCRs, ion channels, enzymes and transporters). Oral bioavailability is high in rat and monkey. In a proof of concept study, PHA-022121 potently inhibits bradykinin-induced haemodynamic changes in freely moving monkeys. The onset of activity was 1 h or less (first time point measured), which was faster than icatibant. PHA-022121 also partially prevents carrageenan-induced paw edema in rat with a longer duration of action as compared to icatibant. Based on experimental data and modeling, Pharvaris expects that a single daily pill of less than 30 mg will provide therapeutic efficacy for at least 24 h. Pharvaris plans to develop PHA-022121 as an oral on-demand and prophylactic treatment of HAE attacks.

O24

Clinical evaluation of pharmacokinetics, pharmacodynamics, safety, and efficacy dose-response of BCX7353 as an acute treatment for angioedema in patients with hereditary angioedema (HAE)

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Background: Approved acute treatments for angioedema attacks in hereditary angioedema (HAE) are administered parenterally, including plasma kallikrein inhibitors. Guidelines recommend at-home on-demand treatment. BCX7353, an orally administered kallikrein inhibitor with fast onset and sustained duration of action, may improve access to self-administered treatment and allow earlier dosing after symptom onset.

Methods: BCX7353 doses for evaluation of efficacy in HAE subjects were selected using pharmacokinetic (PK) and pharmacodynamic (PD) evidence from the first-in-human trial in healthy subjects (dose range: 30 to 1000 mg). Separately, plasma concentration–time profiles, plasma kallikrein inhibition (KKI)–time profiles, and PK-PD relationships of orally administered BCX7353 (750 mg) were evaluated in 6 HAE subjects. We separately conducted a proof-of-concept clinical trial (ZENITH-1) using a 3-part, dose-de-escalation, randomised-sequence, double-blind, placebo-controlled, 3-period crossover design. Subjects were randomised in each part to 2 treatments with BCX7353 (at the same dose level) and 1 with placebo; for each treatment, subjects confirmed the attack with the investigator before dosing. This trial tested on-demand at-home treatment of angioedema attacks with 750, 500, and 250 mg doses of BCX7353 in 58 subjects with HAE. Subjects recorded symptoms and interventions using standardised questions and visual analogue scales (VAS) for skin swelling, skin pain, and abdominal pain in a diary.

Results: Drug levels (Table 1) exceeded 8 × half-maximal effective concentration (EC50) for KKI (approximately lower limit of normal plasma sample 4-fold, BCX7353 concentration > 80% inhibition in vivo corresponds to approximately > 94% inhibition in vivo).

Table 1 PK and PD parameters in healthy subjects and HAE subjects after single oral single doses of BCX7353

| Population          | N  | Dose of BCX7353 mg | CEEmin ng/mL | AUGcET ng·h/mL | [7353] × EC50 at 0.5, 1, 2, 4 h (%) | KNI at 30 min (%) | KKI at 24 h (%) |
|---------------------|----|-------------------|--------------|----------------|-------------------------------------|------------------|-----------------|
| Healthy subjects    | 6  | 250               | 104 (24)     | 995 (25)       | 0.3,1,0                             | 12 (44)          | 24 (18)         |
| Healthy subjects    | 6  | 500               | 245 (59)     | 2700 (43)      | 0.4,6,0                             | 50 (26)          | 72 (18)         |
| Healthy subjects    | 6  | 750               | 584 (26)     | 5670 (17)      | 6,6,6,6                             | 80 (7)           | 81 (11)         |

* PK parameters are geometric mean (% coefficient of variation)

b PD parameters are mean (standard deviation) in an in ex vivo assay (note that n = 5 in HAE subjects), and results reported are percent reduction in specific amidolytic activity from pre-dose sample. As the reagent volume dilutes the plasma sample 4-fold, BCX7353 concentration ex vivo is 25% of that present in vivo postdosing. The reported ex vivo % inhibition therefore underestimates the in vivo inhibitory activity achieved. For example, a value of 80% inhibition ex vivo corresponds to approximately 94% inhibition in vivo.

c PK study in subjects with HAE was conducted at an HAE reference center at University Hospital, Frankfurt, Germany

Table 2 Efficacy endpoints (difference, active vs placebo) for each dose level in ZENITH-1. Phase 2 trial of oral BCX7353

| Endpoint                     | 250 mg | 500 mg | 750 mg |
|------------------------------|--------|--------|--------|
| Number of attacks, active/placebo | 21/11  | 25/11  | 64/31  |
| Change from baseline in VAS score at 4 h, mm³ | 0.57   | -2.1   | -6.98*** |
| Improved or stable VAS at 4 h, % | 6.4    | 18.5   | 21.0*   |
| Improved or stable VAS at 24 h, % | 14.5   | 23.6   | 27.0*   |
| Improved or stable symptoms at 4 h, % | 12.6   | 12.4   | 22.3*   |
| Improved or stable symptoms at 24 h, % | 11.6   | 27.6   | 28.6**  |
| Standard of care rescue treatment at 24 h, % | -7.4   | -13.5  | -31.6*** |
| No or mild symptoms at 24 h | 16.4   | 23.6   | 31.8*** |

* LS mean difference to placebo reported. P-value generated from a mixed effect linear model including treatment, period, and sequence as fixed effects, subject within sequence as a random effect, and predose 3-symptom composite VAS score as a covariate.

** Difference to placebo in proportion of attacks achieving the endpoint reported. P-value generated from a generalised logistic model including treatment, period, and sequence as fixed effects, and subject within sequence as a random effect. Use of standard of care HAE medication = failure for endpoint.

b No significant difference from placebo in any efficacy endpoint.

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O25
KVD900, a new oral on-demand treatment of hereditary angioedema attacks achieves complete plasma kallikrein suppression: safety, tolerability, pharmacokinetic and pharmacodynamic results from a phase 1 first-in-human study
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Background: Attacks of swelling and pain in hereditary angioedema are attributed to increased vascular permeability due to excessive and uncontrolled formation of the proinflammatory peptide hormone bradykinin (BK). BK is generated through cleavage of high molecular weight kininogen (HK) by the serine protease plasma kallikrein (PKa). KVD900 is a novel, potent (Ki 3 nM), selective inhibitor of PKa. An orally available and rapidly absorbed PKa inhibitor could provide a new therapeutic opportunity to halt and resolve HAE attacks early.

Methods: We conducted a first in human study to evaluate the safety, tolerability, pharmacokinetics and pharmacodynamics of KVD900 in healthy adult males aged 18 to 55 years. In Part A we investigated single ascending doses (5 to 600 mg) of KVD900 capsule (a powder in capsule formulation); 8 cohorts of 8 subjects with 6 active treatment and 2 placebo subjects per cohort. In Part B we compared single 100 mg doses of KVD900 tablet and KVD900 capsule in a crossover design involving 8 subjects. In Part C we compared single 600 mg doses of KVD900 100 mg tablets under fed and fasting conditions in a crossover design involving 12 subjects. Plasma kallikrein enzyme activity was measured in dextran sulfate stimulated whole plasma samples using a fluorogenic enzyme assay and a capillary based high molecular weight kininogen (HK) cleavage immunoassay.

Results: Overall, 68 healthy males received KVD900. KVD900 was generally safe and well tolerated without any severe or serious adverse events and without related gastrointestinal adverse events. All adverse events were mild, except for one AE (headache) in the 10 mg cohort of Part A, which was considered moderate. Mean maximum plasma concentration with the 600 mg capsule reached 4.830 ng/mL (± 1.080). Similar exposures were reached with 600 mg tablets under fed and fasted conditions (area under the curve – 0 to infinity [AUC 0–t] 21,200 h*ng/mL vs. 19,800 h*ng/mL). Complete PKa inhibition (99.3%) was achieved on 600 mg, fasted and fed (CI: 99.0% - 99.5%); >85% PKa inhibition was maintained for 8 to 10 h. KVD900 rapidly provided HK cleavage protection for at least 10 h consistent with the PKa enzyme inhibition data.

Conclusion: This first-in-human study of KVD900 showed that a single oral administration of up to 600 mg KVD900 is generally safe and well tolerated without any severe adverse events. KVD900 achieves rapid suppression of PKa activity.

O26
Pharmacokinetics, safety, and potency of ATN-249, a novel oral plasma kallikrein inhibitor for hereditary angioedema
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Background: Hereditary angioedema (HAE) is a rare, potentially life-threatening disease characterized by acute skin and mucosal oedema. Currently available treatments are administered s.c. or i.v., thus there is a need for well tolerated, orally-administered therapies. ATN-249 is a novel oral plasma kallikrein inhibitor designed to treat HAE by blocking kalikrein-mediated production of bradykinin. ATN-249 showed dose linear pharmacokinetics and minimal food effects in a phase 1 single ascending dose study in healthy volunteers. Here we report ATN-249’s pharmacokinetics (PK) and safety from a 14-day multiple ascending dose study and its in vitro potency relative to an approved plasma kallikrein inhibitor.

Materials and methods: Healthy participants received multiple doses of ATN-249 100 mg QD, 200 mg QD, 400 mg QD, 300 mg BID, or placebo for 14 days (6 ATN-249:2 placebo in each dose cohort). Serial blood draws and urinalysis were conducted to calculate PK parameters. Adverse events (AEs) were assessed. Separately, the potency of ATN-249 and lanadelumab, a subcutaneously administered plasma kallikrein inhibitor, were tested in biochemical inhibition and contact activation assays in human plasma as well as in a semi-quantitative Western blot assay evaluating attenuation of cleaved kininogen.

Results: PK parameters following 14-day QD regimens were dose proportional (Table 1). With repeated dosing over 14-days, trough concentrations were on average approximately 200 ng/mL (460 nM) and generally exceeded 527 ng/mL (1.2 µM) for 400 mg QD and 300 mg BID regimens, respectively. Average concentrations at steady-state (Cavg) were approximately 683 ng/mL (1.57 µM) and 1198 ng/mL (2.7 µM) for the 400 mg QD and 300 mg BID regimens, respectively. Across treatment and placebo cohorts, the most common AEs were headache, contact dermatitis secondary to ECG lead placement, and nausea. All AEs were self-limited and not related to study drug. Mean ATN-249 trough concentrations for the 400 mg QD and 300 mg BID dosages exceeded expected therapeutically relevant concentrations and were greater than those demonstrating attenuation of cHMWK suppression of PKa activity.

Table 1 Mean (% CV) PK parameters by dose on day 14 in MAD study (n = 6 per cohort)

| Parameter       | Dose (mg) | 100 QD | 200 QD | 400 QD | 300 BID |
|-----------------|-----------|--------|--------|--------|---------|
| AUCtau (ng*hr/mL) | 4592 (26.9) | 9749 (27.2) | 16390 (47.8) | 28740 (26.4) |
| Cavg (ng/mL)    | 191 (26.9) | 406 (27.2) | 683 (47.8) | 1198 (26.4) |
| Cmax (ng/mL)    | 496 (22.7) | 1202 (26.1) | 2010 (60.5) | 2040 (28.6) |
| Cmin (ng/mL)    | 47 (36.0) | 92 (44.5) | 153 (45.1) | 527* (28.2) |
| Tmax (Hours)    | 2.4 (35.6) | 2.2 (49.9) | 1.9 (43.2) | 11.1 (58.8) |
| Half-Life (Hours) | 10.9 (8.1) | 10.5 (30.3) | 11.7 (20.4) | 7.7 (7.6) |

*η = 5

Fig. 1 Inhibition of plasma kallikrein via biochemical inhibition (percent inhibition)
Population pharmacokinetic analysis of C1-esterase inhibitor functional activity in the COMPACT open-label extension study

Conclusions: ATN-249's PK were dose-linear with low to moderate between-subject variability. Repeat dose trough ATN-249 concentrations were above predicted therapeutic concentrations in ex vivo assays of contact activation. ATN-249 was well-tolerated and no AEs were drug related. ATN-249 demonstrated potent kallikrein inhibition comparable to lanadelumab in biochemical and ex vivo contact activation assays, including Western blot detection of cleaved kininogen. These results, along with predictable PK, suggest ATN-249 may be a potent, safe, oral plasma kallikrein inhibitor for prophylactic treatment of HAE.

O27
Population pharmacokinetic analysis of C1-esterase inhibitor functional activity in the COMPACT open-label extension study is consistent with previous COMPACT studies

Background: In patients with hereditary angioedema (HAE), the relationship between increased risk of attacks and deficiency in complement component 1 (C1) esterase inhibitor (C1-INH[f]) is well established. The objective of this study was to characterise the pharmacokinetics (PK) of subcutaneous (SC) C1-INH in patients with HAE enrolled in the COMPACT open-label extension (OLE) study using a population PK model previously developed using data from prior COMPACT studies. Variation in PK of C1-INH(SC) in different age groups was also evaluated.

Materials and methods: C1-INH(f) was measured in four trials (COMPACT Phase I, II, III and OLE study [NCT1760343, NCT01912456, NCT01576523 and NCT02316353, respectively]). Previously, a one-compartment model with first-order elimination and bodyweight effect on clearance was used to describe the PK of C1-INH(f) after C1-INH(SC) administration in patients with HAE. One thousand profiles for subjects with HAE were simulated based on the distribution of individual weights from COMPACT OLE to assess whether the previously described model could characterise the PK of C1-INH(SC) in patients who completed COMPACT OLE. Post-hoc clearance estimates were visually evaluated. Age groups evaluated for variation were <12, 12–17, >17–<65 and ≥65 years old.

Results: The previously developed population PK model was able to characterise observed C1-INH(f) following C1-INH(SC) administration in patients completing COMPACT OLE. Simulations of steady-state C1-INH(f) following C1-INH(SC) administration at 40 IU/kg and 60 IU/kg doses captured most data from the COMPACT OLE study within the prediction interval, suggesting that the previously developed model was consistent with the observed C1-INH(f) in patients in COMPACT OLE. Individual posthoc clearance estimates were also similar across investigated age groups for paediatric, adolescent, adult, and geriatric subjects. The relationship between clearance and body weight in the previously developed model was found to be applicable to the relationship observed in COMPACT OLE.

Conclusions: Overall, the final analysis of COMPACT OLE was consistent with the analysis performed for prior COMPACT studies. Population PK analysis with data from the COMPACT OLE demonstrated that the previously developed population PK model could predict observed C1-INH(f) in the COMPACT OLE without bias. Population PK parameters remained unchanged and C1-INH(f) was similar across all COMPACT studies following C1-INH(SC) administration. In addition, the PK of C1-INH(SC) was similar in paediatric, adolescent, adult, and geriatric subjects.

O28
Functional C1-Esterase inhibitor and complement protein 4 levels were not altered by lanadelumab treatment in HAE patients in the phase 3 HELP study

Background: The HELP Study® is a multicenter, randomized, double-blind, placebo-controlled Phase 3 study to evaluate the efficacy and safety of lanadelumab for long-term prophylaxis against acute attacks of hereditary angioedema (HAE) in subjects with Type-I and Type-II HAE. All three lanadelumab treatment arms demonstrated statistically significant and clinically meaningful reductions in the number of HAE attacks over the 26-week treatment period compared to placebo. HAE results from variations in the SERPING1 gene that encodes the C1-inhibitor (C1INH), a serine protease inhibitor. Reduced plasma levels of C1INH lead to enhanced activation of the contact system, triggering high levels of bradykinin and increased vascular permeability; this could lead to a feedback loop resulting in further consumption of C1INH by proteases. To investigate the hypothesis whereby plasma kallikrein inhibition minimizes C1-INH consumption and thereby affects complement protein 4 (C4) levels, we assessed the effect of
lanadelumab treatment on functional C1-INH (fC1-INH) and C4 levels in HAE patients in the HELP study.

Methods and materials: The pharmacodynamic samples collected at Predose, Day(D)56, D98, D140 and D182 from all treatment arms: Placebo, lanadelumab 150 mg Q4 W, 300 mg Q4 W and 300 mg Q2 W arms had 41, 28, 29 and 27 subjects, respectively. The average pre-dose fC1-INH measured in HAE subjects (n=115) was 207 mU/mL (SEM of 11 mU/mL) and 306 mU/mL (SEM of 20 mU/mL) in the ELISA and the chromogenic methods, respectively. Average C4 measured in pre-dose samples was 108 µg/mL (SEM of 11 µg/mL).

No statistically significant difference was observed in fC1-INH and C4 levels in the treatment arms when multiple comparisons were made for each of the arms to placebo at all timepoints tested, and/or when pre-dose levels compared to each of the other time-points. The comparisons were also made on baseline normalized percentage changes.

Conclusion: Long term lanadelumab treatment does not seem to affect fC1-INH and C4 levels in HAE subjects in the HELP study suggesting that specific inhibition of plasma kallikrein may not reduce C1-INH consumption resulting in a notable increase in C4 levels. Ongoing research for other HAE pathway biomarkers in the HELP study might provide additional understanding of disease mechanisms.

O29 Hereditary angioedema with normal C1-inhibitor: first report of an Argentinian family with factor XII mutation

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Hereditary angioedema (HAE) is a rare genetic disease associated with either a quantitative or qualitative deficiency in C1-inhibitor (C1-INH) or normal C1-INH.

HAE with normal C1-INH levels can be caused by mutations in different genes. Until now, there are 3 known proteins whose genes mutations lead to HAE: F12 (HAE-FXII), plasminogen (HAEPLG) and angiotensin 1 (HAEANGPT1).

Approximately 30% of these cases are due to F12 gene (HAE-FXII) mutations. Point mutation (Thr328Lys or Thr328Arg), a large deletion (deletion of 72 base pairs: c.971_1018 + 24del172*) or an 18-bp duplication in the F12 (FXII) gene are detected in HAE-FXII.

A 42-year-old woman consulted for facial angioedema. She had 2 episodes of facial angioedema, that lasted 3 days and does not respond to treatment with corticosteroids and antihistamines, in the last 2 years without any recognized trigger.

She referred frequent abdominal pain associated with diarrhea and one of them with hypotension. A diagnosis of gastritis and irritable bowel diagnosis was provided. She was taking levothyroxine 50mcg/day and contraceptives (drospirenone 3 mg, ethinyl estradiol 0.03 mg).

She reported that her sister suffered from facial angioedema after a dental procedure. After the suspicion of HAE, contraceptives were discontinued.

Materials and methods: Serum, citrated plasma and EDTA-blood was collected from the patients. Antigenic values of C4 and C1-INH was assayed by turbidimetric immunoassays in serum samples from patients. Functional C1-INH activity was assayed by a chromogenic assay in plasma samples from patients.

Genetic test for F12 was assayed by sanger sequencing of exon 9 and intron–exon boundaries.

Results: The results of quantitative and qualitative C4 (40.15 mg/dl), C1-INH (24.75 m/dl) and functional C1-INH (154%) were normal.

Genetic test for F12 gene revealed the patient is heterozygous for the common missense mutation c.983C > A (p.Thr328Lys). All symptoms related to angioedema disappeared after contraceptives discontinuation.

After the confirmation of diagnosis of HAE of this patient, pedigree analysis in her family led to diagnosis of 2 other patients. Time to diagnosis since first symptom was 3 years.

Conclusions: We described the first Argentinian family with mutation in F12 gene. Interestingly we found that the clinical episodes in the case study are mild and are related to hormonal levels. All family members are less symptomatic than other types of HAE and the delay in diagnosis was 3 years.

Consent to publish: Written, informed consent for publication was obtained from the patient (or parent/guardian for patients under 16)

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of acquired angioedema type I was confirmed (with haematopoietical background) and following 2 facial attacks were successfully treated with icatibant with rapid relief of symptoms within 2 h after application. Besides typical symptoms of AAE, recurrent symptoms of generalised urticaria associated with intense pruritus were observed. Treatment with desloratadine was initiated (from single dose up to 4-times increased dose), but without any clinical effect. UAS7 score was permanently above 28 points and therefore a treatment with omalizumab was initiated. After second application of omalizumab, complete remission of urticaria was succeeded.

Hereby, we present a unique case of two separated diseases – acquired angioedema of I type and chronic spontaneous urticaria. Despite their similarities, two therapeutic strategies were used for achievement of clinical control over the symptoms. We would like to point out the possible clinical co-existence of two rare diseases, what should be taken into account during the differential-diagnostic algorithm of angioedema in clinical practice.

Consent to publish: Written, informed consent for publication was obtained from the patient [or parent/guardian for patients under 16]

O31 Oligoarticular juvenile idiopathic arthritis in a child with type I hereditary angioedema: a case report
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Introduction: Hereditary angioedema due to C1-inhibitor deficiency is a rare autosomal dominant disorder affecting the C1-inhibitor gene affecting about 1 in 50,000 individuals. The hallmark of HAE is recurrent angioedema. Autoimmune disorders may complicate genetic C1-inhibitor deficiency.

Case report: A 7-year-old boy with a family history of angioedema in his father (diagnosis not established, deceased—globolectasia multiforme) and C1-inhibitor deficiency in paternal grandmother, developed painless swellings of his face following minor trauma from age 6 years. Type I HAE was diagnosed, C4 0.06 g/l (0.16–0.54) C1-inhibitor 0.069 g/l (0.22–0.38), 0% function.

Two months later he developed spontaneous painful swelling of his right knee without trauma or preceding infection. Examination showed warm swollen right knee with minimal restriction of movement and no deformity. No associated systemic or ocular symptoms. FBC and CRP were normal. Autoantibodies and HLA B27 negative. MRI showed large joint effusion with no structural cause. He was referred to Paediatric Rheumatology where a diagnosis of oligoarticular juvenile idiopathic arthritis was made. He made a good response to aspiration and intra-articular triamcinolone. To date there has been no recurrence of JIA. His HAE is under control.

Discussion: HAE I and II may be complicated by autoimmune disorders. Case series report 2–12% of the cohort developing a spectrum of autoimmune disorders. Brickman (JAC 1986,77,749) reported a 6-year-old boy with seronegative non-deforming polyarthritis, IgA deficiency and micrognathia. Our patient is clinically similar except for the single joint involvement. Our patient fulfils the criteria for JIA. We believe JIA in this boy is a complication of HAE and are surprised that we have not encountered a similar situation earlier in his disease.

Consent to publish: Written, informed consent for publication was obtained from the patient [or parent/guardian for patients under 16]

O32 Cardiac tamponade following cardiac surgery in type 2 HAE patient
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Introduction: Invasive and surgical procedures are well known to precipitate angioedema in hereditary angioedema. Cardiac surgery has been successfully performed with a number of prophylactic treatment. We report a patient with HAE who had cardiopulmonary bypass mitral valve replacement surgery with C1-inhibitor prophylaxis complicated by pericardial effusion and cardiac tamponade which was successfully treated.

Case report: MK (female, date of birth 03/09/1972) who has been resident in UK since 1991 was diagnosed with Type 2 HAE at age 22 years and on danazol prophylaxis. Sjogren’s syndrome was diagnosed in 2005. She developed carcinoma breast in 2013 which was treated by lumpectomy followed by chemotherapy and radiotherapy and long term tamoxifen. Mitral valve disease was diagnosed in 2014. Mitral valve replacement surgery was performed in February 2017. She received 1000 units C1-inhibitor before and 1500 units after cardiopulmonary bypass surgery. She was discharged a week later to represent with palpitations and peripheral oedema. Atrial fibrillation, large pericardial effusion and early tamponade was diagnosed. She underwent needle pericardiocentesis under 1000 units C1-inhibitor which was complicated by puncture of the pulmonary artery. Pericardial effusion recurred. Emergency surgical drainage was performed under 1000 units C1-inhibitor before and 1500 units after surgery. Pericardial drain was in situ for 3 days following which she had successful cardioversion under 1000 units C1-inhibitor prophylaxis. The patient who is followed up in another hospital has had exacerbations of angioedema since discharge.

Discussion: The patient had a major surgical procedures done in 2 hospitals with no resident immunologist. There is no specific evidence based guidance on the prophylactic treatment of HAE in major complex surgery. The treatment was successful but highlights the need for a better prophylaxis and treatment protocol and multidisciplinary approach in the management of complex invasive procedures in HAE.

Consent to publish: Written, informed consent for publication was obtained from the patient [or parent/guardian for patients under 16]

O33 Clinical characteristics and therapeutic modalities in Polish C1-INH–HAE patients. A pilot cohort study in adults population
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Introduction: Hereditary angioedema (HAE) is a rare, autosomal dominant, genetic disorder caused by defects in the C1 inhibitor (C1-INH). It is characterized by recurrent attacks of subcutaneous, mucosal, and visceral angioedema, and, in some patients, severe internal bleeding. The prevalence of HAE is estimated at 1:10,000 in the general population. The majority of patients have type I HAE, which is caused by mutations in the C1-INH gene leading to C1-INH deficiency. Type II HAE is caused by mutations in the genes encoding the plasma kallikrein (PK)- and relaxin-2-related peptide (RELAP)-1 receptor (KLK12). Treatment options for HAE include self-administered plasma-derived C1-INH concentrate (C1-INH-HAE) or recombinant C1-INH (rC1-INH) for acute attacks, and prophylactic treatment with C1-INH-HAE or C1-INH–induced conversion factor (C1-INH-ICF) for prevention of acute attacks and surgical procedures.

Background: Our study aimed to determine current management approaches and clinical characteristics of Polish population of adults patients with hereditary angioedema due to C1-inhibitor deficiency (C1-INH-HAE).

Materials and methods: We undertook a survey of consenting C1-INH-HAE patients with a structured medical interview addressing i.a. patients’ sex, date of first symptoms and diagnosis, frequency and localization of angioedema, medication use for on-demand and prophylactic treatment.

Results: Ninety patients, at the mean age of 41.7 years (range 18–77), were available for analysis at the present stage (females: 57, males: 33; HAE type I: 90%, HAE type II: 10%). The mean age at first symptoms was 13 years (range 1–32) and the mean delay in diagnosis was 14.8 years (range 0–56). Family history was present in 89.8% of subjects, while de-novo mutation was presumed in 10.2%. Family history of HAE related death, unnecessary surgeries...
and intubations due to laryngeal attacks were reported in 33%, 11.1% and 4.9% of the patients, respectively. The most common symptoms were (i) peripheral swellings: no attacks, < 12 and ≥ 12 attacks in 16.1%, 59.8%, 24.1% of patients respectively, median: 5, mean: 7.4, range: 0–30 attacks/6 months per patient and (ii) abdominal swellings: no attacks, < 12 and ≥ 12 attacks in 20.1%, 58.6%, 21.3% of patients respectively, median: 4, mean: 6.8, range: 0–30 attacks/6 months per patient. Laryngeal attacks, potentially life threatening, were reported in 28.9% of patients within 6 months before a survey. Additionally, 75.4% of patients reported attacks in multiple localizations at the same time. 61.7% of the patients report self-administration of on demand treatment and 78.3% of them carry emergency treatment when travelling. 8.1% of the patients do not have on demand drugs at home. The others have pdC1-INH (83.5%), icatibant (46.8%), rhC1-INH (8.9%) – percent do not add up to 100% due to patients having simultaneously more than one drug, 19.8% of the patients didn’t use on demand treatment within the last 6 months, the others used pdC1-INH (76.8%), icatibant (49.3%), rhC1-INH (8.7%). Long-term prophylaxis with danazol and/or tranexamic acid were used by 9.3% of the patients within the last 6 months and by 51.5% of the patients, ever.

Conclusions: Clinical profile of the investigated patients is similar to other reported populations. Collected data gives further insight into unmet medical needs, e.g. self-administration and may help to identify certain patient groups which require additional focus in the future.

O34
Treatment of patients with hereditary angioedema with normal C1-inhibitor: evaluation of 295 patients
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Background: Hereditary angioedema with normal C1-inhibitor (HAE-nlC1-INH) is a rare condition and clinical features are like those of HAE with C1-INH deficiency. Hormones have special role as triggering factors. There is no biomarker for diagnosis, requiring a consistent clinical and family history and/or identification of associated mutation (Factor 12, Angiopoietin 1 and Plasminogen).

Methods: 304 patients of 101 unrelated families were evaluated as suspected HAE-nlC1-INH. Questionnaires were filled out by 16 Brazilian Centers. HAE-nlC1-INH criteria was fulfilled by 225 patients of 78 families. One of the families (4 members) showed Angiopoietin 1 mutation. Detailed clinical data including therapy were recorded. Genetic mutations were performed for the 3 known HAE-nlC1-INH mutations. This study was approved by the Ethics Committee of Centro Universitario Saúde ABC (CAAE:51896015.0.1001.0082).

Results: 44/225 (19.5%) were asymptomatic and 181/225 (80.5%) were symptomatic. Out of these, 141/181 (77.9%) had F12 mutation, 2/181 (1.1%) had Angiopoietin 1 mutation, 21/181 (11.6%) the mutation was unknown and 17/181 (9.4%) weren’t tested yet (Table 1). Eight symptomatic patients with F12 mutation (8/141, 5.6%) underwent surgical procedures due to angioedema: laparotomy, 4 (2.8%); laparoscopy, 3 (2.1%); tracheostomy, 1 (0.7%). Among symptomatic patients with unknown mutation, 2 were submitted to laparotomy (2/38; 5.3%). Thirteen (13/181, 7.2%) were admitted in ICU during attacks at least once and 3/181 (1.7%) died due to HAE attacks. Of the symptomatic patients, 38/181 (21%) did not receive any specific treatment. The patients treated (143/181, 79%) were 116/143 (81.1%) with F12 mutation, 25/143 (17.5%) with unknown mutation and 2/143 (1.4%) with Angiopoietin 1 mutation. Among the patients who under went specific treatment: 45/143 (31.5%) only stopped or modified contraceptives; 32/143 (22.4%) were medicated during the attacks only and 66/143 (46.1%) received long term prophylactic treatment. Short term prophylaxis was performed for 28/143 (19.6%). Tables 3 and 4 refer to drugs used for on-demand treatment and prophylaxis.

Conclusions: Our registry represents one of the greatest HAE-nlC1-INH casuistry reported. Clinical manifestations were similar both in F12 or ANGPT1 mutated and unknown mutation HAE-nlC1-INH patients. Most of the patients were symptomatic and almost half of them needed prophylactic treatment.

Keywords: Hereditary angioedema; C1-inhibitor; therapy; f12 mutation.

Table 1 Clinical data of patients with HAE-nlC1-INH according to the identification of mutation

| HAE-F12 | HAE-unknown |
|---------|-------------|
| Patients | 183 | 38 |
| Families | 62 | 15 |
| Family history | 170/183 (92.9%) | 38/38 (100%) |
| Sex | 153F:30M (83.6%;16.4%) | 33F:5M (86.8%;13.2%) |
| Onset of symptoms (age) | 2–68 (136/141) (median 18.5) | 2–52 (31/38) (median 16) |
| Site of edema | | |
| Subcutaneous | 84/141 (59.6%) | 26/38 (68.4%) |
| Facial | 119/141 (84.4%) | 31/38 (81.6%) |
| Genital | 21/141 (14.9%) | 10/38 (26.3%) |
| Abdominal | 109/141 (77.3%) | 28/38 (73.7%) |
| Tongue | 32/141 (22.7%) | 5/38 (13.2%) |
| Upper airways | 65/141 (46.1%) | 17/38 (44.7%) |
| Trigger factors | | |
| Hormones (symptomatic women) (n = 128) | | |
| Oral contraceptives | 87/128 (68.0%) | 17/33 (51.5%) |
| Pregnancy | 25/128 (19.5%) | 6/33 (18.2%) |
| Menses | 1/128 (0.8%) | 3/33 (9.1%) |
| HRT | 2/128 (1.6%) | 1/33 (3.0%) |
| All symptomatic patients | | |
| Stress | 87/141 (61.7%) | 18/38 (47.4%) |
| Trauma | 70/141 (49.6%) | 16/38 (42.1%) |
| Surgery | 11/141 (7.8%) | 4/38 (10.5%) |
| Dental procedures | 21/141 (14.9%) | 5/38 (13.2%) |
| Infection | 6/141 (4.3%) | 4/38 (10.5%) |
| Weather | 11/141 (7.8%) | 4/38 (10.5%) |
| Exercise | 4/141 (2.8%) | 5/38 (13.2%) |

HRT: hormonal reposision therapy
Table 3 Treatment on demand in patients with HAE-nlC1-INH

| Medication       | Total (32/143 = 22.3%) | HAE-F12 (24/116 = 19.8%) | HAE-Unknown (8/25 = 28%) |
|------------------|-------------------------|--------------------------|--------------------------|
|                  | n  | %            | n  | %            | n  | %            |
| Icatibant        | 7  | 21.9         | 3  | 12.5         | 4  | 50.0         |
| C1-inhibitor     | 3  | 9.4          | 1  | 4.2          | 2  | 25.0         |
| Tranexamic acid  | 25 | 78.1         | 21 | 87.5         | 4  | 50.0         |
| Danazol          | 1  | 3.1          | 0  | -            | 1  | 12.5         |

Table 4 Medications used for continuous prophylactic treatment by patients with HAE-nlC1-INH

| Medication       | TOTAL (66/143 = 46.1%) | HAE – F12 (53/116 = 45.7%) | HAE – UNKNOWN (12/25 = 48%) | Angiopoetin 1 (1/2 = 50%) |
|------------------|-------------------------|-----------------------------|-----------------------------|---------------------------|
|                  | n =66 | %        | n =53 | %        | n =12 | %        | n =1  | %        |
| Tranexamic acid  | 52   | 78.8     | 43   | 81.1     | 8     | 66.7     | 1     | 100      |
| Danazol          | 20   | 30.3     | 12   | 22.6     | 7     | 58.3     | 1     | 100      |
| Oxandrolone      | 14   | 21.2     | 8    | 15.1     | 5     | 41.7     | 1     | 100      |
| Epsilon aminocaproic acid | 4  | 6.1     | 0    | 3         | 25    | 1        | 100     |

O35 Long-term prophylaxis with C1-inhibitor concentrate in patients with hereditary angioedema

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15(Suppl 4):2019, O36

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Objectives:introduction: Hereditary angioedema due to C1-inhibitor deficiency (C1-INH-HAE) is a rare autosomal dominant inherited disease. The recurrent swelling attacks such as subcutaneous edema and colic-like abdominal pain negatively affect quality of life (QoL). Laryngeal edema is rare, but life-threatening if untreated. C1-INH is currently approved for prophylaxis to routinely prevent attacks in patients aged ≥ 6 (EU) and ≥ 12 years (US). Real life data in nine pediatric patients with HAE who received C1-INH concentrate for the routine prevention of angioedema attacks was documented and followed up.

Methods: After giving informed consent the following data was collected and analyzed from patient’s diaries and records 1 year before onset of and after introduction of prophylactic treatment: age at first manifestation and diagnosis, age at first treatment, frequency and location of attacks, prophylactic respectively on-demand therapy regimen. Initial standard prophylactic treatment (SP) consisted of 1000 U C1-INH, (bw > 40 kg) or 500 U C1-INH (bw < 40 kg) every 3–4 days i. v. and was intensified (individualized prophylaxis—IP) in case of > 2 breakthrough attacks per month. In 2 patients the prophylactic regimen had to be intensified to every 2 days regimen.

Results: Six patients (3 male/3 female) aged 4.5–17.2 years with HAE-C1-INH who received pdhC1INH for the routine prevention of angioedema attacks were documented and followed up. Conclusion: LTP with pdhC1INH was more frequently needed by women, proved to be an effective, safe and well tolerated alternative in patients with contraindications for administration of conventional LTP, including pregnancy and lactation.

O36 The needs of individually tailored prophylaxis with C1-INH concentrate in pediatric patients with hereditary angioedema (HAE) – real life data from 6 pediatric patients

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Background: Hereditary angioedema due to C1-inhibitor deficiency (HAE-C1-INH) is a rare autosomal dominant inherited disease. The recurrent swelling attacks such as subcutaneous edema and colic-like abdominal pain negatively affect quality of life (QoL). Laryngeal edema is rare, but life-threatening if untreated. C1-INH is currently approved for prophylaxis to routinely prevent attacks in patients aged ≥ 6 (EU) and ≥ 12 years (US). Real life data in nine pediatric patients with HAE who received C1-INH concentrate for the routine prevention of angioedema attacks was documented and followed up.

Methods: After giving informed consent the following data was collected and analyzed from patient’s diaries and records 1 year before onset of and after introduction of prophylactic treatment: age at first manifestation and diagnosis, age at first treatment, frequency and location of attacks, prophylactic respectively on-demand therapy regimen. Initial standard prophylactic treatment (SP) consisted of 1000 U C1-INH, (bw > 40 kg) or 500 U C1-INH (bw < 40 kg) every 3–4 days i. v. and was intensified (individualized prophylaxis—IP) in case of > 2 breakthrough attacks per month. In 2 patients the prophylactic regimen had to be intensified to every 2 days regimen.

Results: Six patients (3 male/3 female) aged 4.5–17.2 years with HAE-C1-INH who received pdhC1INH for the routine prevention of angioedema attacks were documented and followed up. Conclusion: LTP with pdhC1INH was more frequently needed by women, proved to be an effective, safe and well tolerated alternative in patients with contraindications for administration of conventional LTP, including pregnancy and lactation.

O36 The needs of individually tailored prophylaxis with C1-INH concentrate in pediatric patients with hereditary angioedema (HAE) – real life data from 6 pediatric patients

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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):O35

Background: Hereditary angioedema due to C1-inhibitor deficiency (HAE-C1-INH) is a rare autosomal dominant inherited disease. The recurrent swelling attacks such as subcutaneous edema and colic-like abdominal pain negatively affect quality of life (QoL). Laryngeal edema is rare, but life-threatening if untreated. C1-INH is currently approved for prophylaxis to routinely prevent attacks in patients aged ≥ 6 (EU) and ≥ 12 years (US). Real life data in nine pediatric patients with HAE who received C1-INH concentrate for the routine prevention of angioedema attacks was documented and followed up.

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Conclusions: These real life data strongly indicate that prophylaxis regimes should be tailored according to the individual patient's needs in order to achieve favorable results.

Posters

P01
A case of hereditary angioedema associated with rheumatoid arthritis: treatment challenges
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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P01

Background: Patients with hereditary angioedema (HAE) have an increased incidence of autoimmune diseases, such as systemic lupus erythematosus, rheumatoid arthritis (RA), autoimmune thyroiditis, glos-merulonephritis, etc.

We here present a case of severe RA in a female patient with HAE and the impact of specific RA treatments on the severity of HAE.

Case report: 34-years old female patient in whom diagnosis of HAE was established after diagnosing of her uncle and mother. She is a member of the largest HAE family in Serbia, in which all offspring inherited the disease. Disease-causing missense mutation in SERP-ING1 gene (Pro377Ser-c.1195C>T on exon 7) was identified in all family members. The first presentation of HAE was laryngeal angi- oedema at the age of 24. Since then moderate and severe attacks occurred with the frequency of 1–2 per year. She was diagnosed with seropositive erosive RA at the age of 20 and initially treated with prednisolone and metotrexate. Since she did not go into remis-sion the treatment with tocilizumab (humanized recombinant anti-interleukin-6 receptor antibody) was initiated two years later. She was treated with tocilizumab 480 mg i.v. monthly, daily oral methylprednisolone 4 mg and alfalcacidol between 22 to 33 years of age with good clinical response. For arterial hypertension since the age of 26 treatment consisted of amlopidine and bisoprolol. As she planned to become pregnant therapy was changed to methyldopa and etanercept (tumor necrosis factor inhibitor) at a dose of 50 mg s.c. weekly. Shortly after the introduction of new therapy she experienced flare-ups of RA and severe HAE attacks 1–3 per month including laryngeal edema. When pregnancy was confirmed etanercept was discontinued. Therapy with methyldopa was continued. She miscarried when she was 6 weeks pregnant at October 2018. Since etanercept was stopped, she has not experienced any severe HAE attack. Tocilizumab treatment was reintroduced in January 2019 and proved to be effective.

There is no literature data of tocilizumab usage in HAE. Etanercept was used in two patients with psoriatic arthritis and angioedema (one with HAE, the other with AAE) with favorable effects. Two cases of severe angioedema, were described in patients treated with etanercept. (one with RA and one with refractory adult-onset Still's disease).

Conclusion: To the best of our knowledge, this is the first case of successful treatment of RA with tocilizumab in HAE patient. Moreover, it's needed to further evaluate the effect of etanercept on possible wors-ening of HAE.

Consent to publish: Written, informed consent for publication was obtained from the patient [or parent/guardian for patients under 16].

P03
Tranexamic acid plus sodium bemiparin as long term prophylaxis in a patient with FXII-HAE during pregnancy: a case report
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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P03

Background: Hereditary angioedema (HAE) is characterized by recurrent attacks of severe swelling with involvement of multiple organs, which are induced by genetic mutations that result in increased bradykinin levels. Patients with a mutation in F12 gene (FXII-HAE) especially show worsening of symptoms under hyperestrogenic conditions, such as pregnancy. Therapy for HAE is limited during pregnancy, delivery, and postpartum (1). Plasma-derived C1-INH (pdcC1-INH) concentrate is the election treatment recommended during these periods (2).

Materials and methods: We present the evolution and management of repeated angioedema attacks during pregnancy in a woman with HAE, with normal levels and function of C1-INH and a mis-sense muta-tion in F12 gene. She had been diagnosed in 2012, during her first pregnancy trimester the patient was treated several times with intravenous plasma derived human C1-INH concentrate (pdcC1INH) (Berinert™, CSL-Behring, Marburg, Germany) in the Emer-gency Room, sometimes needing two or three 1,500U doses to resolve

Conclusions: Treatment with rhC1-INH in pregnant women with HAE was generally safe and well tolerated. All 14 deliveries were defined in full-term babies at full-term without complications. Supported by Pharming Healthcare Inc.
the attack. pdhC1INH was discarded as LTP due to lack of efficacy in the treatment of acute angioedema attacks. The patient was evaluated by a haematologist before starting oral tranexamic acid (TXA), who recommended co-treatment with sodium bempiparin 7,500U daily and hematoletic controls every few weeks. She was asymptomatic for 4 months immediately after starting oral TXA 500 mg every 8 h, having just 3 mild attacks in the last 3 months of pregnancy and treating 2 of them with 1,500U of pdhC1INH with acceptable response. A caesarean delivery was scheduled because of a prior caesarean and short term prophylaxis with IV pdhC1INH (1,000U) was administrated. She discontinued TXA the night before the delivery, which went through without incidences, and a healthy female baby was born. A tubal ligation was also performed. She was breastfeeding and had to continue the treatment with heparin for 6 weeks after the delivery. No new angioedema attacks were observed postdelivery.

**Conclusions:** In our patient, a successful management of the angioedema attacks during pregnancy was carried out with TXA and anticoagulant treatment. No side effects were observed.

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**PO4**

**Metabolic complications of late diagnosis in hereditary angioedema**

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**Allergy, Asthma & Clinical Immunology** 2019, 15(Suppl 4):P04

**Background:** Type I hereditary angioedema (type I HAE) is a rare genetic disease characterized by episodes of subcutaneous or mucosal edema, without urticaria or pruritus, that can be fatal due to laryngeal swelling. In Mexico the diagnosis is usually delayed due to lack of clinical suspicion in first-contact physicians. The most frequent previous diagnosis is chronic urticaria, whereby patients receive recurrent and prolonged treatment with antihistamines and corticosteroids. Metabolic syndrome (MS) is traditionally an adult disease, however its prevalence in pediatrics is increasing; is characterized by a combination of dyslipidemia, abnormal glucose regulation, central adiposity and hypertension. It has defined diagnostic criteria for children and adolescents by the International Diabetes Federation. Metabolic syndrome in children and adolescents is associated with a higher risk of developing cardiovascular disease and type 2 diabetes in this group of patients.

**Case report:** We present the case of 3 patients with type I HAE who before diagnosis received multiple cycles of systemic corticosteroids as treatment for angioedema, emesis and abdominal pain. Two patients met obesity criteria by body mass index (BMI); within their diagnostic approach multiple metabolic disorders were evidenced: hypertriglyceridemia, low high density protein (HDL) cholesterol, increased abdominal circumference and hypertension for age, fulfilling criteria for metabolic syndrome. The third patient was obese, with full moon face, hirsutism, dorsal fat pad and total cholesterol level in 232 mg/dl, diagnosis of Cushing syndrome was made.

**Conclusion:** Metabolic syndrome, obesity and cushing syndrome are serious complications of the excessive use of steroids in patients with type 1 HAE. As allergologist we must intervene quickly and aggressively seeking to avoid the increased in cardiovascular risk and type 2 diabetes in this group of patients.

**Consent to publish:** Informed consent to publish has been obtained from this patients parents.

**Table 1 Definition of metabolic syndrome in children and adolescents by the International Diabetes Federation**

| Parameters                          | 10–16 years |
|-------------------------------------|-------------|
| Waist circumference                 | ≥ 90th percentile* |
| Number of abnormalities             | ≥ 2 of the following |
| Triglycerides                       | ≥ 150 mg/dl |
| High density protein (HDL) cholesterol | < 40 mg/dl |
| Blood pressure                      | Either |
| Systolic                            | ≥ 130 mmHg |
| Diastolic                           | ≥ 85 mmHg |
| Fasting glucose                     | ≥ 100 mg/dl |

*Fernandez JR, et al. Waist circumference percentiles in nationally representative samples of African-American, European-American, and Mexican-American children and adolescents. J Pediatr 2004; 145:439

**Table 2 Metabolic abnormalities**

| Parameters                          | Patient 1 | Patient 2 | Patient 3 |
|-------------------------------------|-----------|-----------|-----------|
| Age                                 | 12 years  | 14 years  | 7 years   |
| Weight                              | 60 kg     | 74 kg     | 40        |
| BMI                                 | 26.3 (P97) | 27.9 (P99) | 24 (P99)  |
| Waist circumference                 | 89 cm     | 102 cm    | 72 cm     |
| Triglycerides                       | 162 mg/dl | 216 mg/dl | 132 mg/dl |
| High density protein (HDL)           | 33.3 mg/dl | 36.1 mg/dl | 41       |
| Blood pressure                      |           |           |           |
| Systolic                            | 125 (P > 95) | 120 (P50) | 100 (P50) |
| Diastolic                           | 75 (P90-95) | 84 (P99)  | 60 (P50)  |
| Fasting glucose                     | 93 mg/dl  | 106 mg/dl | 83 mg/dl  |

**Case report:** We present the case of 3 patients with type I HAE who before diagnosis received multiple cycles of systemic corticosteroids as treatment for angioedema, emesis and abdominal pain. Two patients met obesity criteria by body mass index (BMI); within their diagnostic approach multiple metabolic disorders were evidenced: hypertriglyceridemia, low high density protein (HDL) cholesterol, increased abdominal circumference and hypertension for age, fulfilling criteria for metabolic syndrome. The third patient was obese, with full moon face, hirsutism, dorsal fat pad and total cholesterol level in 232 mg/dl, diagnosis of Cushing syndrome was made.

**Conclusion:** Metabolic syndrome, obesity and coughing syndrome are serious complications of the excessive use of steroids in patients with type 1 HAE. As allergologist we must intervene quickly and aggressively seeking to avoid the increased in cardiovascular risk and type 2 diabetes in this group of patients.

**Consent to publish:** Informed consent to publish has been obtained from these patients parents.

**PO5**

**HAE Patients in Ukraine: frequency and localization of attacks in 2018**

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**Allergy, Asthma & Clinical Immunology** 2019, 15(Suppl 4):P05
**Table 1 Localization and frequency of HAE attacks in 2018**

| Edema localization | Man | Women |
|--------------------|-----|-------|
| Peripheral         | 30  | 40    |
| Face               | 15  | 7     |
| Abdominal          | 25  | 27    |
| Upper airways      | 5   |       |

**Background:** Ukraine has HAE 37 patients: 5 children (mean age 9 y.o., age range 2 to 16 y.o.), 2 men and 20 women (mean age 41 y.o., age range 21 to 68 y.o.). The patients have no access to the contemporary treatment, but the situation is expected to change in 2019.

**Objective:** This study focuses on the anamnesis, localization, severity and frequency of HAE attacks.

**Method:** Analysis of the patients’ answers to the questionnaire about time of the first attack, date of HAE-established diagnosis, frequency, severity, most frequent localization of the attacks, triggers and family anamnesis.

**Results:** 30 patients indicated that they had attacks between 0 and 5 years old, 5 patients developed attacks in the puberty, 2 patients started suffering after 30 years old. Out of these 30 patients 21 adult patient and 1 boy has severe throat and abdominal attacks that require hospitalization. The boy was diagnosed at 5 years old with no family history (parents have had no blood tests yet).

20 patients consider psychological and physical stress, respiratory viral infections, physical traumas, hypothermia, overheating as triggers to develop an HAE attack. Taking ACE inhibitors is a trigger as well: an HAE patient took ACE inhibitor (captopril) to decrease the blood pressure, and in 4 h he developed extremely severe abdominal attack, hypotony and loss of consciousness. In 6 h of intensive care (FFP and other treatment) he was back to normal.

5 adults (3 men and 2 women) had upper airway attacks in 2018. They were hospitalized and transfused FFP [fresh frozen plasma]. FFP was effective in all cases, the edema and pain decreased within one hour and in 4–6 h the patients were transferred from intensive to a usual unit. Yet, after the FFP 2 patients developed rash and scleral icterus (probably due to not sufficient plasma purification) that went back to normal in 3–5 days of antiallergic and detoxication therapy. In Ukraine, FFP is also transfused upon a severe abdominal attack when a patient is in intensive care for an average of 3 days. The attacks of the peripherals are not treated because they are not life-threatening.

As shown in Table 1, during 2018 men and women had HAE attacks mainly of peripherals and abdominal parts. Men had twice more facial and upper airway attacks. When compared with others, the patients that suffer HAE from their childhood have more severe and frequent attacks.

**P07**

**Higher annual rate of angioedema attacks in HAE-C1-INH patients above the age of 65 compared to patients aged 18 to 64 years**

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**Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4): P07**

**Background:** Hereditary angioedema with C1-inhibitor deficiency (HAE-C1-INH) is a rare genetic disease that usually starts to manifest in childhood. Almost all adult patients with HAE-C1-INH who are under clinical care are symptomatic. Data on the course of the attack frequency with increasing age in adulthood are scarce. In this cross-sectional study we report data on the annual attack rate in patients over 65 years of age compared to the attack rate in patients at the age of 18 to 64.

**Methods:** Attack characteristics in 147 adult patients with HAE-C1-INH followed at the HAE Comprehensive Care Center at the University Hospital Frankfurt, Germany, were investigated. Patients on exclusive on demand therapy were included. The main exclusion criteria were long-term prophylaxis or pregnancy during the investigational period.

After approval by the Ethics Committee of the University Hospital Frankfurt, Germany, data on the clinical course of the disease including frequency and sites of HAE attacks, triggering factors and therapy documented in the patients’ records and symptom diaries were analyzed. The frequency of HAE attacks in the entirely documented previous year was assessed in two age groups of patients, age ≥ 65 years (47) and age 18 to 64 (100). To analyze the effect of age in more detail, a subset was analyzed further: male and female patients aged 18–30 (28), 31–50 (49) and 51–64 (23).

**Results:** The mean annual attack rate of angioedema attacks was higher in patients aged 65 or more versus patients aged 18 to 64 years (37.7 vs 31.5 attacks per year, p = 0.000). The mean annual attack frequency in the subsets of patients aged 18–30 years and aged 31–50 years was 28.1 per year and 29.9 per year (p = 0.000). The highest annual number of attacks was found in patients aged 51 to 64 (39.5 per year) (p = 0.000).

**Conclusions:** In this cross-sectional study, patients with HAE-C1-INH aged 65 years or more had more attacks than younger adults. Apart from more diligence in the documentation of attacks or possible selection of more severe cases in the older patient population, this may be an indication of an individual increase of the attack rate over a life-time.
P08
A questionnaire study to determine association of dental hygiene practices in hereditary angioedema subjects with the incidence of post-procedural angioedema attacks

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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P08

Rationale: Dental hygiene behaviors and practices in HAE subjects (i.e., the frequency of routine dental care visits and use of dental hygiene products) may be influenced by severe life-threatening episodes triggered by routine professional dental procedures. We utilized a questionnaire to analyze perceptions by HAE subjects about personal dental care practices compared to non-HAE (control) populations.

Methods: A self-reported questionnaire linked to REDCap® server was distributed to gather information on differences in dental care perceptions and behaviors in HAE (n = 250) and non-HAE populations (n = 256) matched by age, gender, race, and ethnicity. Chi square analyses and a generalized linear model (SAS) were used to determine the significance of association between several parameters related to dental care beliefs and utilization of dental care products, using the questionnaire responses from both groups.

Results: He frequency of routine dentist visits did not differ significantly between groups, but among the HAE group it was significantly less in subjects with previous post-procedure angioedema (AE) attacks. Interestingly, antibacterial toothpaste usage was higher among HAE subjects (9.6% vs. 4.3%, p = 0.02). The odds of using anti-bacterial toothpaste was significantly higher among HAE subjects concerned about AE attacks and actually experiencing attacks than those not concerned or experiencing such attacks (OR: 3.3 [1.1, 9.7], p = 0.03).

Conclusion: Experiencing angioedema attacks after prior dental procedures was the most significant determinant in HAE subjects resulting in less frequent routine dentist visits and preference for using anti-bacterial toothpaste. (P08)

P09
The relationship between disease activity and quality of life – a first–time survey in hereditary angioedema

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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P09

Background: Prodomes of Hereditary Angioedema (HAE) are frequently reported in close association with the swelling attacks. Since attacks usually follow prodomes in a close proximity, patients may not be able to tell the difference between the two events. For practical purposes, especially self-administration and timing of treatments, it is important to distinguish prodomes from attacks. We sought to investigate the differences and correlations between prodomes and attacks, by using a new validated PRO instrument.

Methods: We designed and tested a questionnaire-based PRO instrument for the evaluation of HAE prodomes and attacks. The high internal content validity and reliability (Cronbach’s α = .70 to .96) render it suitable for studying clinical expressions of HAE. A cohort of 66 HAE patients completed a questionnaire, developed specifically to address their most recent experience with prodomes and attacks. It incorporates five ‘clusters’ of body systems: limbs, abdomen, face, laryngeal, genitalia. This instrument was powered to evaluate if they could distinguish between five dimensions of prodomes and subsequent attacks. Items included: location, pain, intensity, impairment and dysfunction. Patients were asked to grade their experience in both events on a Visual Analog Scale (Likert) of 0–10 cm.

Results: One-way MANOVA with repeated measurements, on all dimensions, showed significant differences between the clinical dimensions of prodomes and attacks, in all body clusters [F (4, 56) = 45.7, P < .001, Eta² = .77]. Patients could distinguish between the two and consider them separately, but associated. Additionally, one-way ANOVA for each dimension demonstrated very high differences. For example, differences in abdominal pain [F (1, 59) = 104.1, P < .001, Eta² = .64], in the limbs [F (1, 59) = 57.2, P < .001, Eta² = .47], and less in the face [F (4, 60) = 30.7, P < .001, Eta² = .33]. Association between the events, examined by Pearson’s coefficient, showed high statistical correlation (i.e. abdominal pain r = .45, p < .001, laryngeal pain r = .46, p > .001, facial pain r = .36, p < .01). Therefore, we could demonstrate that higher intensity of a prodrome was followed by higher intensity of an attack. Collectively, the intensity, severity, impairment and loss

enrolled, who completed the Angioedema Quality of Life (AE-QoL) questionnaire on occasion of their annual follow-up visit (95 in 2016, 97 in 2017, and 94 in 2018). The findings from the survey were subjected to statistical analysis in order to establish whether the AE-QoL total score is correlated with the annual number of HAE episodes, or with the levels of complement parameters (classical total complement, C3, C4, C1-INH concentration, and C1-INH functional activity) measured in blood samples obtained upon completion of the questionnaire.

Results: We found a significant positive correlation between AE-QoL total score and the annual number of HAE attacks in each year of the survey (2016: p < 0.0001, r = 0.41; 2017: p < 0.0001, r = 0.51, 2018: p < 0.0001, r = 0.51). However, a significant relationship could not be established in any of these years between AE-QoL total score and the level of classical total complement, C3, C4, and C1-INH functional activity.

Conclusion: AE-QoL is a valuable tool for the assessment of quality of life in C1-INH-HAE patients, because our findings established AE-QoL as a good indicator of disease severity. This makes us to believe that the patients’ quality of life can be reliably assessed based on the annual number of HAE episodes. However, when AE-QoL reflects a greatly reduced quality of life, but the number of HAE attacks is low, individualized assessment is necessary to develop an appropriate therapeutic strategy.

This study was supported by OTKA K124557

P10
Are HAE patients able to distinguish prodomes from attacks, and are they correlated?

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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P10

Background: Prodomes of Hereditary Angioedema (HAE) are frequently reported in close association with the swelling attacks. Since attacks usually follow prodomes in a close proximity, patients may not be able to tell the difference between the two events. For practical purposes, especially self-administration and timing of treatments, it is important to distinguish prodomes from attacks. We sought to investigate the differences and correlations between prodomes and attacks, by using a new validated PRO instrument.

Methods: We designed and tested a questionnaire-based PRO instrument for the evaluation of HAE prodomes and attacks. The high internal content validity and reliability (Cronbach’s α = .70 to .96) render it suitable for studying clinical expressions of HAE. A cohort of 66 HAE patients completed a questionnaire, developed specifically to address their most recent experience with prodomes and attacks. It incorporates five ‘clusters’ of body systems: limbs, abdomen, face, laryngeal, genitalia. This instrument was powered to evaluate if they could distinguish between five dimensions of prodomes and subsequent attacks. Items included: location, pain, intensity, impairment and dysfunction. Patients were asked to grade their experience in both events on a Visual Analog Scale (Likert) of 0–10 cm.

Results: One-way MANOVA with repeated measurements, on all dimensions, showed significant differences between the clinical dimensions of prodomes and attacks, in all body clusters [F (4, 56) = 45.7, P < .001, Eta² = .77]. Patients could distinguish between the two and consider them separately, but associated. Additionally, one-way ANOVA for each dimension demonstrated very high differences. For example, differences in abdominal pain [F (1, 59) = 104.1, P < .001, Eta² = .64], in the limbs [F (1, 59) = 57.2, P < .001, Eta² = .47], and less in the face [F (4, 60) = 30.7, P < .001, Eta² = .33]. Association between the events, examined by Pearson’s coefficient, showed high statistical correlation (i.e. abdominal pain r = .45, p < .001, laryngeal pain r = .46, p > .001, facial pain r = .36, p < .01). Therefore, we could demonstrate that higher intensity of a prodrome was followed by higher intensity of an attack. Collectively, the intensity, severity, impairment and loss
of functionality were much lower in the prodromes than in the attacks, particularly in the skin and abdomen clusters. 

Conclusions: By using HAE-specific instrument, we found that patients could identify the prodromes, and describe them accurately by using the questionnaire items. We demonstrate that prodromes and attacks in various locations are distinguishable but associated. 

P11 Psychological processes in the adaptation to disease in adults with hereditary angioedema due to C1-inhibitor deficiency: a pilot study from Italian referral centers

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Background: In a previous research with pediatric C1-inhibitor Angioedema (C1-INH-HAE) patients we found higher levels of perceived stress and alexithymia, evaluated by self-report scales, compared to the normative validation samples. 

Objectives: The aim of this study is to evaluate the relation between psychological processes and: a) severity of the disease; b) patient’s adaptation to it, in a sample of adults with C1-INH-HAE. In particular, we evaluated the processes of emotion regulation, alexithymia, perception of stress and patient’s health engagement. 

Method: Within a broader mixed-method research design in clinical health psychology, n. 60 people affected by HAE from the main Italian referral centers and n. 30 controls – affected from different chronic conditions- will be administered the following psychometric scales: Toronto Alexithymia Scale (TAS-20-Taylor, 2004), Emotion Regulation Questionnaire (ERQ, Gross & John, 2003), Perceived Stress Scale (PSS, Cohen, and Williamson, 1988), Patient Health Engagement Scale (PHE-S, Graffigna et al., 2015). 

Results: Preliminary results on 19 subjects are reported. 79% of the sample are females. The mean age is 43.95 (±10.47) years and the mean of the years elapsed from the diagnosis is 14.64±(±14.42). The levels of perceived stress are sensibly higher (20.26(±7.5) than the normative sample (12.1(±5.9)). Alexithymia and Emotion Regulation scores do not differ significantly from the normative validation samples. The scores to Patient Health Engagement Scale (PHE-S) on the adaptation to the disease show that the majority of the subjects lays on a level of “alert” or “adhesion” toward the disease. 

Conclusion: First of all, our results seem to confirm the higher levels of stress found in pediatric C1-INH- HAE patients. Moreover, the adaptation to the disease measured by the PHE-S, that refers to the level of eudaimonic wellbeing, is low regardless to the years elapsed from the diagnosis. These results seem to suggest that the clinical features of C1-INH-HAE, such as the high unpredictability of the attacks may impair patients’ capacities to adapt to the disease and to cope with stress. An enlargement of the sample is needed to confirm and enrich such results. In a second phase, a broader aim of our project will be focused on developing an ad hoc counseling intervention focused on the disease management. 

P12 Depression in hereditary angioedema can cause sexual morbidity

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Introduction: Hereditary angioedema (HAE) is a rare disease characterized by recurrent attacks involving hand-arm-leg, face, bowels, genitals and upper airways. Although symptoms usually begin in the first 10 years of life, the frequency and severity of attacks increase in adolescence. In this study we analyzed that how the sexual life quality is affected by HAE attacks’ severity and frequency, attacks after sexual intercourse and anxiety and depression in HAE. 

Material and method: Forty-eight patients who have been followed at Egé University Medical Faculty Internal Medicine HAE special clinic were enrolled in this study. Socio-demographic characteristics of the patients, disease onset age, attack localization and frequency, gender-specific sexual life questions were assessed. Hospital Anxiety and Depression Scale (HAD) and new sexual satisfaction scale (NSSS) were also applied. 

The adaptation and reliability studies of HAD for the Turkish population showed that 7 were cut-off scores for depression and 10 were cut-off scores for anxiety. Higher values than these cut-off points were defined as depression risk and anxiety disorder risk. 

Results: Of the 48 patients with the average age 39.89±13.68; 26 (54.2%) were female. In females, the mean score of NSSS was 65±14.24 and the mean anxiety score 8.3±3.16 and the mean depression score was 6.3±3.85; whereas mean scores of males were 72.95±17.71, 7.4±3.69 and 5.36±3.27, respectively. The mean NSSS score of the patients which had attacks depending on sexual intercourse (n = 10; 63.8±25.02) was lower than the patients which did not have attacks (n = 32; 71±12.96) (t = 0.87, p = 0.402). The NSSS mean score of the patients with depression risk (n = 14) was significantly lower than those who had no risk of depression (n = 29); [61.57±15.04 vs 72.69±16.06; (t = 2.17; p = 0.036)]. When both sexes were examined separately for NSSS score; in only males, the risk of having depression affected the mean NSSS total scores [with depression risk n = 7; 60.42±17.94 and without depression risk (n = 15; 78.8±14.75) (t = 2.543; p = 0.019)]. 

Discussion: The sexual life of HAE patients with depression risk are affected negatively, therefore an intervention for this group should be planned. We believe that this study shed light on the related problem and will lead to further studies. We did not find any significant finding in women. Most probably female patients had not been able to answer the questions honestly due to the privacy of sexual satisfaction scale questions. 

P13 Immigrants’ perspective on living with hereditary angioedema in Denmark – a qualitative study

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Background: In the last decades several migrants with C1-inhibitor hereditary angioedema (C1-INH-HAE), a rare and potentially life-threatening disease, has moved to Denmark. HAE is known to impact quality of life (QoL) and patients often face a high disease burden, but there is a lack of data pertaining immigrants living with this rare disease. The objective of this study is to investigate and improve our understanding of how patients of other ethnic backgrounds experience living with HAE in Denmark, and to gain deeper insight into these patients’ understanding of disease, challenges and management of symptoms in their everyday life. We also explore their past and familial experiences from their countries of origin. 

Methods and methods: This study is designed as a qualitative study, based on semi-structured interviews, to ensure a deep and broad understanding of the patients’ experiences. The participants are recruited from the group of adult immigrants that are treated at the national HAE center. In addition to the interviews, patients are asked to complete two quality of life questionnaires, EQ-SD and HAE-QoL. The anonymized data will be analyzed descriptively.
Results: 9 out of 13 adult immigrants with HAE in Denmark have chosen to participate in the study (age 21–60 years). Collection of data is ongoing until April 2019 and results of selected thematic areas will be presented. Preliminary results show that one main concern for several migrant patients is not receiving accurate treatment at emergency departments, when they present with severe attacks and emergency medication.

Conclusion: This study is expected to provide unique information about migrants experiences living with a rare disease in Denmark. Preliminary data highlights the challenges, but also importance of teaching migrant patients home therapy, as is also the case for ethnic Danish patients.

P14
Description of angioedema episodes prompting a call on the bradykinin mediated angioedema reference centre on-call hotline
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Introduction: Angioedema (AE) are acute, localized subcutaneous swellings that regress over hours to days. Their severity and treatment differ depending on the underlying cause: mast cell (MC) or bradykinin (BK) mediated angioedema. To this date, no reliable diagnostic tests are available to rapidly differentiate these forms of AE.

Objective: This study aimed to describe the possible underlying cause of an AE episode after a call on the French national reference center for AE hotline.

Methods: Physicians calling on the CREAK hotline between March and August 2018 were asked to fill a clinical description form for the AE episode. Patients were classified in the groups AE in the presence of a renin-angiotensin system inhibitor (ACE inhibitor or Angiotensin receptor blocker) (Drug-AE), mast cell mediated AE (MC-AE) and possibly bradykinin mediated AE in the absence of a renin-angiotensin inhibitor (BK-AE).

Results: 88 patients were included. 41 (48.8%) in the Drug-AE group, 39 (46.4%) in the MC-AE group, and 4 (4.8%) in the BK-AE group. The incidence of a renin-angiotensin system inhibitor (ACEi) from an authentic ACEi induced BK-AE render the use of costly trand 북ic acid; others treat symptoms with icatibant or C1 esterase inhibitor concentrate (C1INH). C1INH (plasma-derived or recombinant) is available for short-term prophylaxis (pre-surgery, dental) or longer term (school/college exams) according to the UK HAE consensus document [1].

Results: The questionnaire comprises 25 questions reflecting the previous 6 months. The answers are graded out of 5 or 6 with higher score reflecting ‘not a problem’ and lower score ‘extremely’. Answers are captured into 7 dimensions: physical functioning and health, disease related stigma, emotional role and social functioning, concern about offspring, perceived control over illness, mental health, and treatment difficulties. The results are presented here by gender and age ranges for each dimension. Note: no female patients in this cohort fell into the age range 35 – 50 years.

Conclusion: High scores across all dimensions (mean and median scores) suggest this group of patients has reasonably good quality of life apart from one outlier, a female aged 50+. However lower scores for perceived control over illness highlight the unpredictable nature of HAE attacks even if few patients experience treatment difficulties. The author intends to repeat this study when new ‘pipeline’ prophylactic medications become available for UK patients.

Acknowledgements: Alex Symons, MSc Student Archaeological Science, University of Oxford for help with data presentation.

P15
Quality of life among HAE patients in South West England (Devon and Cornwall) using HAE-QoL questionnaire designed by Foundation for Biomedical Research of La Paz University Hospital Madrid (FIBHULP)
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Background: Hereditary Angioedema (HAE) is a rare but potentially life-threatening inherited condition. HAE is characterised by episodes of angioedema affecting various body parts including the hands, feet, face and airway. Incidence is approximately 1 in 50,000. The Department of Clinical Immunology and Allergy at University Hospitals Plymouth has 43 patients with HAE from across Devon and Cornwall: an incidence of approximately 1 in 42,000. Dedicated HAE clinics are held 3 times a year where patients are seen, clinically assessed, have treatment reviews and are encouraged to enroll in clinical trials and research studies or have ongoing review by the research team.

Method: In June 2017 permission was granted to the lead author to use the HAE-QoL v.2 questionnaire, and the project was registered with UHPNT Audit Department.

Between October 2017 and February 2019 patients attending HAE clinics were approached to complete the questionnaire and 24 agreed. All HAE patients in our clinics have access to a range of therapies. Some prefer to take traditional oral prophylactic therapies (attenuated androgens, tranexamic acid); others treat symptoms with icatibant or C1 esterase inhibitor concentrate (C1INH). C1INH (plasma-derived or recombinant) is available for short-term prophylaxis (pre-surgery, dental) or longer term (school/college exams) according to the UK HAE consensus document [1].

Results: The questionnaire comprises 25 questions reflecting the previous 6 months. The answers are graded out of 5 or 6 with higher score reflecting ‘not a problem’ and lower score ‘extremely’. Answers are captured into 7 dimensions: physical functioning and health, disease related stigma, emotional role and social functioning, concern about offspring, perceived control over illness, mental health, and treatment difficulties. The results are presented here by gender and age ranges for each dimension. Note: no female patients in this cohort fell into the age range 35 – 50 years.

Conclusion: High scores across all dimensions (mean and median scores) suggest this group of patients has reasonably good quality of life apart from one outlier, a female aged 50+. However lower scores for perceived control over illness highlight the unpredictable nature of HAE attacks even if few patients experience treatment difficulties. The author intends to repeat this study when new ‘pipeline’ prophylactic medications become available for UK patients.

Acknowledgements: Alex Symons, MSc Student Archaeological Science, University of Oxford for help with data presentation.
Hereditary angioedema with C1-inhibitor deficiency (C1-INH-HAE) is characterized by recurring and spontaneously resolving edematous attacks with not fully understood pathomechanism. Previously many studies published on the activation of plasma enzyme systems during edematous attacks, nevertheless kinetic follow-up has never been performed. For the first time, we aimed to study the kinetics of parameters in the coagulation and fibrinolytic systems in a spontaneously resolved edematous attack of a C1-INH-HAE patient.

In a 56-year-old female with C1-INH-HAE we monitored the severity of the symptoms during the observation period and altogether twelve blood samples were obtained. Blood samples collected from a healthy control volunteer at 5 different times during a 24–hour period. We measured Factors XI and XII activities (FXa, FⅪa), prothrombin time (PT) and activated partial thromboplastin time (aPTT) as well as concentration of Factor V, VII and X (FV, FVII and FX respectively), prothrombin fragment 1 + 2 (F1+2), thrombin-antithrombin (TAT)-complex, D-dimer, fibrinogen.

After a 24-hour symptom-free period and another 19-hour prodromal period (characterized by erythema marginatum), the patient had a 29-hour-long edematous attack in multiple skin locations, and was followed up for another day. During prodromal stage—the levels of D-dimer, F1 + 2 and TAT-complex were as constantly low as those levels measured in the healthy control, whereas fibrinogen level increased. Levels of F1 + 2 and TAT-complex were significantly elevated at the onset of edematous symptoms whereas level of D-dimer was elevated after 6 h. Levels of all three parameters reached maximum 12 h after reaching the maximum severity score of symptoms. FXⅪa and FXⅪa as well as the levels of FV, FⅦI and FX did not show unidirectional changes during the observation period. PT and aPTT did not shorten before or during the edema.

Real-time monitoring of F1 + 2 and TAT complex suggest that thrombin may contribute to edema formation. Whereas the generation of thrombin probably not related to the activation of neither the intrinsic nor the extrinsic pathway of coagulation but related to other factors such as MASP-1 or MASP-2. We confirmed that D-dimer is a prominent biomarker of an ongoing edematous attack while fibrinogen is could be a biomarker of prodromal period. This study was a part of a project aimed to better understand the mechanisms leading to the onset and to the resolution of edematous attack. This study was supported by OTKA 112110 and the UNKP-16-3 New National Excellence Program of the Ministry of Human Capacities.

Consent to publish: Written, informed consent for publication was obtained from the patient and control person.
and/or function, lack of family history, no C1NH gene mutation, age at onset of angioedema symptoms and low or undetectable C1q levels.

**Objectives:** To test the diagnostic value of a novel ELISA assay for the detection of circulating complexes between C1NH and C1INH-autoantibodies (C1NH-anti-C1NH) in serum.

**Methods:** An in-house sandwich ELISA was designed to measure free anti-C1INH Abs and complexed C1NH-anti-C1NH Abs. Twenty European patients with AAE were included and characterized on the basis of their complement levels and function.

**Results:** Free anti-C1INH Abs were detected in 9/20 patients (6 of IgG class, 2 of IgM class and one simultaneously presenting IgG and IgM classes), whereas C1NH-anti-C1NH complexes were found in 90% (18/20) of the AAE cases, regardless of the presence or absence of detectable free anti-C1INH Abs. It is of note that 9/20 (45%) patients showed negative serum free anti-C1INH Abs, but positive C1INH-anti-C1NH complexes in their first measurement. In the cohort presented, IgM-class C1INH-Abs are specifically and strongly associated to undetectable C1q serum levels.

**Conclusion:** Detection of C1NH-anti-C1NH Abs provides a diagnostic added value for AAE diagnosis, especially in those cases in whom no free anti-C1NH antibodies are detected. The link between IgM-class C1NH-anti-C1NH complexes and C1q consumption could have further implications for the development of autoimmune manifestations in AAE.

**P19**

**The Romanian Hereditary Angioedema Center started the Global Hereditary Angioedema Registry activities**

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**Introduction:** Taking into consideration that hereditary angioedema is a rare disorder, studies on a large number of patients are still wanted. The Global Hereditary Angioedema Registry is a common, worldwide platform that collects demographic and clinical data from consenting people with hereditary angioedema, in accordance with agreed inclusion criteria and definitions.

**Methods:** The Global Angioedema Registry was developed in 2017, by the Italian hereditary angioedema working group. Romania joined this registry in December 2018, the fifth country after France, Greece, and Hungary. The minimum requirements for affiliation included recruiting at least ten patients with a follow-up visit or phone contact with the center during the last two years and obtaining the signed informed consent from the patients. For each patient the following data were recorded: demographic characteristics (date of birth, gender, date of diagnosis, the age at diagnosis), plasma levels of C4 and C1-INH (antigenic and function), presence of family history of angioedema, comorbid diseases, frequency, location and severity of attacks, use of on-demand and prophylactic treatments. The data source was the medical record of individual patients. The acquisition of data was performed by the treating physician.

**Results:** Till January 2019, a total of 14 patients were included by the Romanian Hereditary Angioedema Center: nine female and five men with C1-INH-HAE (10 type1 and four type 2), median age 45, median age at diagnosis 26. Eight patients had positive family history for angioedema. Based on patient diary, between April-November 2018, 144 attacks were recorded: 45 peripheral, 86 abdominal and 13 laryngeal. Sixty-six of the attacks were severe, 71 moderate and seven mild. One hundred and twelve attacks were treated with subcutaneous injection of Icatibant and four with additional plasma-derived C1-INH. Ten patients were on prophylactic therapy.

**Conclusions:** A prospective long-term registry is a critical tool in building a broad and comprehensive knowledge base. By continuing to introduce valid data from as many patients as possible, useful information can be obtained for the better knowledge of this rare disease.

**P20**

**Hereditary angioedema: quality of life in 19 patients**

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**Background:** Hereditary angioedema (HAE), is characterized by recurrent episodes of angioedema that expose patients to the risk of asphyxiation and have considerable impact on their quality of life (QoL). New therapies have been developed for treating or preventing attacks. On demand treatment is the first option, however many of them need long-term prophylaxis (LTP). The aim of this study is to evaluate the QoL of patients with HAE according to their treatment schedules.

**Materials and methods:** 24 patients with HAE attended at the Gregorio Marañón University hospital were asked to complete the AE-QoL questionnaire. Nineteen returned the fulfilled document. AE-QoL questions are grouped into four domains (Functioning, Fatigue/Mood, Fears/Shame and nutrition). Individual domain scores were transformed into a linear 0–100% scale where higher scores are indicative of a higher QoL impairment. Patients were classified in Group A: patients on "on demand" treatment; Group B: received LTP (plasma-derived C1INH 63%, attenuated androgens 28% or tranexamic acid 9%); Group B1: i.v. treatment; Group B2: LTP with oral drugs.

**Results:** Group A, n = 8, (50%), average age of 53y (range: 26–86). Group B, n = 11, (64%), average age of 53y (range: 28–73). 62.5% patients from group A have type 1 HAE-C1INH, 25% type 2 and 12.5% nC1-INH (HAE-FXII). 100% group B have type 1 HAE-C1INH. The average age at the time of the first symptoms was 15.75y in group A and 4.9y in group B, (p = 0.17). The average time between onset of symptoms and diagnosis was 12.12y in group A and 9.82y in group B (p = 0.9). AE-QoL scores from group A and group B were: Functioning domain: 16% vs 19.31%; Fatigue/Mood dimension: 23.33% vs 36.36%; Fears/Shame domain: 18.97% vs 24.98%; Nutrition domain: 22.22% vs 24.5%. No score exceeded >40%. No significant differences in QoL were found between both groups. A comparison between B1 and B2 and B1 and A groups was performed. Significant increased differences were only found in the functioning domain between group B1 and B2 17.85% vs 21.87% (p < 0.05).

**Conclusion:** To achieve an optimal control of symptoms, a precise treatment adapted to the severity of the disease is needed. According to this questionnaire, patients had an impaired QoL most prominent in the fatigue/Mood dimension. Surprisingly, no significant differences in QoL were found between patients requiring LTP, more severe cases, and patients in "on demand" treatment. The long interval between the onset of symptoms and the confirmation of the diagnosis was notori-ous in both groups.

**P21**

**The French side of the Global Angioedema Registry**

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**Rationale:** Angioedema is a recurrent localized swelling of cutaneous and mucosal tissues. Potentially life-threatening, it causes temporarily impaired function which deteriorates quality of life. Seven inherited or acquired forms of angioedema without wheals are yet classified, including hereditary or acquired C1-inhibitor deficiency (C1-INH-HAE and C1-INH-AAE). This year, the French angioedema network (CRAEK) joined the registry of angioedema without wheals (Cloud-R HAE). Here we present the contribution of the Grenoble Alpes University Hospital (CHUGA) to this disease registry.

**Methods:** Study population is composed of C1-INH-HAE/AAE patients with a proved diagnosis. The following items are collected: patients' personal-demographic data, medical/hospitalization/genetic characteristics, major comorbidities, treatments (prophylaxis/acute attacks).

As from Cloud-R HAE structure, patients can directly provide information on angioedema attacks and their treatment through a dedicated electronic app, web connection or paper support, which is then transferred into the registry at CHUGA.

The Study Protocol has received approval from the Ethical Committee.

**Results:** Since February 2018, 23 C1-INH-HAE patients have been included (informed consent signed). Within C1-INH-HAE, median age is 44 years (range 12–72), sex ratio: 6/17 (M/F). Seventeen percent of them provide prospective data on angioedema attacks. 71% of attacks have been treated. Due to the frequency of symptoms, 52% of them are on long-term prophylaxis (LTP) with tranexamic acid (8%), Danazol (8%), C1-INH concentrate or recombinant (16%) and 34% with progestin. Since February 2019, 8 additional centers have joined the registry: they included 13 more patients without event yet.

**Conclusions:** Angioedema registry gives the possibility to gather information to define natural history of angioedema and to evaluate treatment efficacy in real life. The possibility that data from single countries merge into a global structure facilitates improvement and dissemination of the knowledge on this rare disease and its treatment.

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**P23**

**Designing and delivering Educational Therapeutic Program training kit for HAE patients across France**

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**Rationale:** In 2016, the French Educational Program "EDUCREAK" was assessed by the French Health Care Agencies (ARS) and further approved for an additional 4 years. This pilot program was initially developed in cooperation with staff from four sites in France. After the program's implementation and success in four years, our new objectives are to scale up the program on a national level and streamline best practices among HAE healthcare professionals (HCP).

**Methods:** To reach these objectives, meetings and workshops were held to interactively share the different experiences and methods used by HCP. From these sessions we reflected and drew conclusions on the optimal way to cater to both adult and children HAE patients. We weighted costs and grants required to deliver adequate outputs. In a participatory approach, patients and HCP were asked to work together to attain common objectives, guided by a step-by-step explanatory sheet. Training kit components were then tested by HAE advocate patients and HCP in the last five months to ensure satisfaction. An evaluation system has been established to keep measuring the kit's efficacy. Indicators are not only based on autonomous and safe patient skills but also on quality of life (HAE-QoL and AE-QoL scores).

**Results:** According to the above criteria, an educational training kit was designed and delivered to assist HCP in providing skills for quality care for HAE patients. It has now been streamlined in 12 sites across the country (an additional 8 compared to the previous 4). We thank the French Ministry of Health for their financial support to the National Reference Centre for Angioedema "CRAEK".

**Conclusions:** We are pleased to present a valuable training kit for HAE patients and their HCP. In May, the training kit will be launched in Paris. Next steps include ensuring the training of HCP on using the training kit and systematically evaluating the 10 different sites where they will be used.
P24
Genetic segregation study in angioedema with normal C1-inhibitor (n-C1-INH-HAE) in Southern Spanish population
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Background: In the last two decades have been more frequently described families with HAE with normal C1-INH levels and function. The pathophysiologic mechanism and the genetic basis of this type of HAE were unknown, but subject of recent studies. The field of medical genomics is rapidly growing and interpret genetic and genomic data is driving a new era of healthcare. It constitutes, therefore, a key component of personalized medicine.

Our objective was to study the possible genetic mutations in 8 unrelated families with nC1-INH-HAE.

Materials and methods: We have included 24 patients from 8 unrelated families with n-C1-INH-HAE. In all of them, the index case had been previously analyzed and carried all the prevalent variants of a panel of 55 selected genes encoding proteins related with Kallikrein-Kinin system. Some of the studied relatives presented angioedema and some of them were asymptomatic. For the familial study, we studied 45 genes out of these 55 previously selected. The genetic analysis was carried out through a customized NGS platform (Ampliseq, Thermo Scientific), examining the coding regions and the exon-intron splice junctions of these genes. Subsequently, the results were compared with the Ion Reporter v5.2 software (Thermo Scientific).

Results: Of the 45 genes studied, we have found mutations in 22 out of 45 genes: SERPINA1, XPNPEP2, A2 M, KLK3, ESRR, BDKRB2, PLG, PLAT, PLAU, MME, TNF, F12, ACE, KLKB1, SERPING1, MPO, TLR4, ELANE, HSP90AA1, CPN1, SERPINE1, F13B. A big number of the detected mutations are not described previously. Most mutations occurred in heterogeneous and we can not know at present their clinical significance. We have found a mutation in homozygosis for PLAUR, present in one patient. The clinical situation as the age of the studied members varies between the different families, as well as between members of the same family, although most episodes of angioedema involve facial, peripheral and abdominal areas.

Conclusion: In the genetic segregation analysis we describe a series of heterogeneous mutations in our patients of n-C1-INH-HAE and in some of their asymptomatic relatives. Further studies are needed to find correlation among the detected mutations and clinical expression.

P25
A deep intronic SERPING1 variant associated with hereditary angioedema due to C1-inhibitor deficiency (C1-INH-HAE)
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In about 5% of C1-INH-HAE cases no mutation in the coding region of SERPING1 can be detected. Bearing in mind that non-coding variants can potentially have deleterious effects on a transcript through the regulation of splicing or transcription, defects located in an intronic or an untranslated region of the gene have been considered responsible for modifying C1-INH expression in these cases. However, until today no such alteration has been described. Here, we present a deep intronic SERPING1 variant associated with type I C1-INH-HAE.

Six members of a Greek family with type I C1-INH-HAE were examined. Conventional genotyping of the family members by sequencing all SERPING1 translated regions and intron-exon boundaries, long-range PCR and multiplex ligation dependent probe amplification (MLPA) did not reveal any defect. Analysis by next-generation sequencing (NGS) platform targeting the entire SERPING1 gene [1] uncovered a novel mutation (c.-22-155G>T) in intron 1 that was confirmed by Sanger sequencing. All four affected family members (3 men, 1 woman, mean age at disease onset > 6 years) were carriers of the mutation while none of the two healthy members did. The same mutation was also detected in an additional unrelated Greek male patient and was not detected in 6 additional C1-INH-HAE families without defects in translated regions.

Bioinformatics analysis by the use of Neural Network (score 0.96, range 0–1), NetGene 2 Server (confidence 0.79, range 0.5–0.95), Alternative Splice Site Predictor (ASSP) (confidence 0.458, range 0–1) and FSPLICE (threshold 10.16) bioinformatics tools corroborated that c.-22-155G>T mutation creates an alternative donor site and as a result alters the splicing process. Analysis of the effect of the mutation on the SpliceAid 2 bioinformatics tool showed that it eliminates the number of possible protease acting in the specific region. TraP score evaluating a single nucleotide variant’s ability to cause disease by damaging the final transcript, classified the mutation at the intermediate pathogenic range, akin to possibly damaging classifications (score 0.686, range 0–1). TraP evaluates the mutation based on DANN (score 0.9507, range 0–1) and GERP conservation score 4.8, range –12.3 to 6.17) prediction tools.

Up to now, 49 different intronic mutations in SERPING1 gene have been associated with hereditary angioedema but all are located in the donor and acceptor site or a few nucleotides from these regions. Provided that it will be confirmed by appropriate functional tests, the c.-22-155G>T is the first deep intronic mutation associated with C1-INH-HAE.

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P26
The silo effect in the annotation of SERPING1 variation
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Defining the clinical validity and the pathogenicity of SERPING1 variants is a particularly challenging task. Reporting of variants by reputable variant databases is considered as a criterion for their classification (criteria PP5 and BP6 of ACMG-AMP guidelines) [1]. We investigated the usefulness of public databases in regard to the interpretation of SERPING1 variants. The ClinVar, an NCBI-funded primary centralized database for archiving clinically relevant variants for many diseases, and the HAEdb, a C1-inhibitor gene mutation database were examined.

Using an appropriate broad PubMed search query we retrieved 213 publications referred to 574 distinct SERPING1 variants the great majority of which (95.3%) are associated with hereditary angioedema due to C1-inhibitor deficiency (C1-INH-HAE). Among these, 42.9% were missense/morphens, 9.1% splice site and 0.6% regulatory variants, 35.5% were indels, and 11.9% large defects. At the same time, 544 out of the 567 SERPING1 variants listed in the obtainable by subscription professional version of Human Gene Mutation Database (HGMD) [2] are reported to be disease-causing and five as likely disease-causing.
By December 2018, 82 and 302 SERPING1 variants had been reported in ClinVar and HAEdb, respectively. All variants reported in HAEdb but only 51% of those in ClinVar were reported in the literature (Fig. 1). In order to examine whether classifiable SERPING1 variants reported in other databases were included in ClinVar and HAEdb, we filtered the 351 SERPING1 variants reported in the Exome Aggregation Consortium (ExAC) database v.0.3.1 in regard to their global frequency, according to the BS1 criterion of ACMG-Amp guidelines. Variants with an allele frequency lower than the prevalence of angioedema (<0.002%), and non-canonical transcript variants were excluded. ACMG-Amp criteria were applied to the remaining 99 variants using the available open-access evidence. 15/99 variants were found reported in ClinVar (2 benign, 8 likely benign, 2 variants of unknown significance -VUS- and 2 with multiple assertions) while only 4/99 in HAEdb (1 pathogenic, 1 unknown and 2 polymorphisms).

Public databases include less than 50% of classifiable SERPING1 variants with ClinVar containing approximately 4-times less than HAEdb references. However, the ClinVar database allows for the deposition of variants with clinical observations and assertions, with review status tracked to enable a more transparent view of the levels of quality of the curation [3]. Therefore, clinical laboratories and researchers involved in genotyping angioedema patients are encouraged for real-time submission in ClinVar of variant data, including clinical assertions and evidence used for the variant classification.

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P27

Variant pathogenicity curation in primary angioedema with normal C1-inhibitor (nl-C1-INH-HAE)

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Background: Last year, genomic analysis allowed the discovery of two variants pathogenic for nl-C1-INH-HAE in two different genes (PLG and ANGPT1). However, the implementation of high throughput DNA sequencing results in a dramatic increase of variants identification the classification of which represents a challenging task. Here, we present the variant pathogenicity curation following the next-generation sequencing (NGS) genotyping of nl-C1-INH-HAE patients.

Materials and methods: 130 unrelated patients with nl-C1-INH-HAE (53 Hungarian, 32 Italian, 27 Spanish, 9 Polish, 9 Greek) were submitted to targeted genotyping focused on 55 genes possibly involved in angioedema pathogenesis. The gene list was compiled from literature data on angioedema and genetic predisposition, protein–protein interaction networks, and pathway analysis.

169 patients with HAE due to C1-inhibitor deficiency (C1-INH-HAE) were genotyped as controls. A NGS custom platform (Ampliseq, Thermo Scientific) was developed by which all coding regions and exon–intron splice junctions of these genes (coverage > 90%) were analyzed. Analysis of primary data was conducted with Ion Reporter software v.S.2 (Thermo Scientific). Variants with worldwide frequency >1% (1000 Genomes Global Minor Allele Frequency, ExAC) and polymorphisms (UCSC Common SNPs) for which no disease associations are reported in the ClinVar database were excluded. SIFT and PolyPhen2 tools were used for bioinformatics prediction of the pathogenicity of the remaining uncommon variants. Family segregation studies were performed where feasible.

Results: Among nl-C1-INH-HAE patients, no carriers of the ANGPT1 p.Ala119Ser variant were found while the PLG p.Lys330Glu was detected in 4 (2.9%) unrelated probands (one homozygote) along with other uncommon PLG variants (p.Arg89Lys, p.Arg253His, p.Arg490Gln, p.Arg523Trp). 259 uncommon variants were filtered in amongst which 35 mutations were not previously reported in population databases. Notably, the novel p.Tyr94Ser in KRT1 was detected twice. 7/130 and 3/130 nl-C1-INH-HAE patients, but none of C1-INH-HAE patients, were found to be heterozygous of KNG1 and XPNPEP1 uncommon variants, respectively. Amongst them, the novel KNG1 p.Pro574Ala variant was found segregated with the disease in an Italian family, while segregated compound heterozygosities were also detected in two other families.

Conclusions: We provide evidence indicating the existence of a heterogeneous genetic background linked with nl-C1-INH-HAE cases. Family segregation and functional studies are in progress for the pathogenicity or the disease modifying effect of these genetic alterations to be confirmed.

P28

Hereditary angioedema: a report from the Republic of Belarus

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Background: Hereditary angioedema due to C1-inhibitor deficiency (Type I) or dysfunction (Type II) is a rare genetic condition characterized by recurrent episodes of edema with an estimated frequency of 1:50,000 in the global population without racial or gender differences. HAE Type III is even less common, and unlike Types I and II, does not appear to be connected with the levels of C1-inhibitor. Recently three genes associated with HAE type III have been found. As long as U-HAE is present in the classification, it will encourage scientists to further searches for new genes.

Materials and Methods: For patients with angioedema of unclear etiology, with or without a family history, the following tests are performed according to need: measuring levels of C3c, C4, C1-INH, C1q; C1 function test and expression of C1-INH. NGS-seq amplicons of SERPING1 gene applies to confirm HAE Types I and II. To confirm HAE Type III, the analysis of up to 201 amplicons of 18 genes using Nextera XT (Illumina) is required. All clinically significant observations are confirmed by Sanger sequencing, MLPA.

Results: Overall 149 patients (56.37% female; 43.63% male) were included in this analysis. For 45 patients (64.44% female; 35.56% male) from 19 unrelated families C1-INH-HAE were confirmed. 17 splicing (37.7%), 15 missense (33.3%), 8 frameshift (17.8%), 3 large deletion (6.7%), 2 nonsense (4.5%) mutations have been found. It was decided to compare whether there is a pattern between the type of mutation and the expression of C1-INH. A statistically significant dependence has not been found.

Conclusion: Belarus is a country in Eastern Europe with a population of 9,508 million. Comparing the frequency of occurrence of HAE to the total number of population, there should be about 190 patients in the register, that is, about 76% of HAE patients are still undiagnosed. Improving the HAE diagnosis by enhancing the education of physicians and patients as well as raising awareness in the society is our main goal to the upcoming years. The NGS-sequencing could be a useful in determining the exact genetic alteration. Using NGS technology in genetic diagnostic is much cheaper and easier than SSCP and Sanger sequencing of all exons.

P29
A national audit of hereditary and acquired angioedema in New Zealand
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Hereditary Angioedema (HAE) is a rare potentially life threatening genetic condition, but access to effective therapies can reduce mortality and improve quality of life. Patients with this condition in New Zealand remain uncharacterized by number, geographical distribution, severity or treatment experience. New Zealand Immunologists were invited to recruit patients with HAE and AAE to the audit or those identified as having Berinert® for hereditary or acquired angioedema. Participants were consented, their angioedema related health information was collected and they were invited to take part in telephone or face to face interviews about their experience of healthcare. Twenty one patients with hereditary angioedema were recruited, three of whom had acquired angioedema C1-inhibitor deficiency. Three patients were diagnosed prior to the onset of symptoms due to the diagnosis of a family member with the disease by screening family members. The average diagnostic delay was 12.9 years. Variation in delay between different types of HAE, with the greatest mean delay being in Type 1 HAE at 18.5 years, 13 years in type 2 HAE, and 3.6 years in AAE. Within the cohort of 21 patients there were reports of 4 deaths of family members due to HAE. The majority of patients 19/21 (90%) had a written plan to present to the emergency department. Few (24%) had a Medic Alert® bracelet. In 2015 there were a total of 217 HAE attacks in 16 patients. Five patients (24%) were asymptomatic. Only one patient had angioedema of the upper airway in 2015 but did not require intubation. Six patients had 136 abdominal attacks; some with high frequency (range 1–52). 4 patients said HAE had no impact on their life, 10 had minor impact, and 4 moderate and 3 described it as severe. This study characterizes a cohort of AAE and HAE patients in New Zealand.

P30
Situational analysis of diagnosis and treatment of hereditary angioedema in Latin America
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Background: Hereditary angioedema (HAE) is an autosomal dominant disease resulting in unpredictable attacks of swelling that can be debilitating or life-threatening for affected patients, so early diagnosis and appropriate therapy are essential. Since tests and drugs needed for diagnosis and treatment are not always available, management of this condition remains a challenge both for the treating physicians their patients. HAE affects an estimated of 1 in 50,000 individuals, although this may vary in different regions.

Objective: To develop a situational analysis regarding the availability to perform diagnostic tests and the access to treatment in patients with HAE in Latin America.

Methods: Cross-sectional survey in which 45 expert physicians in HAE from fourteen countries in Latin America were asked about the availability and access to diagnostic tests and treatment for hereditary angioedema in their country of origin. Three patient organizations were also included (Brazil, Chile and Mexico).

Results: The percentage of patients with diagnosis related to the estimated prevalence can be observed in Table 1. The diagnostic laboratory tests and the treatments available by country, at the time of the survey are shown in Tables 2 and 3, respectively.

Table 1 Estimated patients by population vs patients with diagnosis

| Country | Population 2018* | Estimated prevalence 1:50,000 | Patients diagnosed | % With diagnosis |
|---------|------------------|------------------------------|-------------------|-----------------|
| Argentina | 44,556,277 | 891 | 430 | 48.3 |
| Brazil | 212,664,367 | 4,253 | 578 | 13.6 |
| Chile | 18,433,065 | 368 | 11 | 2.98 |
| Colombia | 49,436,892 | 989 | 227 | 22.9 |
| Costa Rica | 4,945,674 | 99 | 19 | 19.1 |
| Dominican Republic | 10,859,463 | 217 | 22 | 10.1 |
| Ecuador | 16,783,322 | 335 | 5 | 1.49 |
| El Salvador | 6,170,519 | 123 | 8 | 6.5 |
| Guatemala | 17,206,382 | 344 | 3 | 0.87 |
| Mexico | 131,452,016 | 2629 | 296 | 11.2 |
| Panama | 4,092,816 | 82 | 13 | 15.8 |
| Paraguay | 6,663,728 | 137 | 12 | 8.75 |
| Peru | 32,424,843 | 648 | 15 | 2.31 |
| Puerto Rico | 3,669,093 | 73 | 90 | 123 |
| TOTAL | 559,558,457 | 11,188 | 1729 | 15.45 |

*World population prospects 2017 United Nations

@Bolivia and Venezuela were invited but, they did not present data
Table 2 Diagnostic laboratory tests available by country

|                      | fC1-INH | AgCl-INH | C4 | C1q | Anti-C1-INH Abs |
|----------------------|---------|----------|----|-----|-----------------|
| Argentina            | Y       | Y        | Y  | Y   | N               |
| Brazil               | Y       | Y        | Y  | Y   | Y               |
| Chile                | Y       | Y        | Y  | N   | Y               |
| Colombia             | Y       | Y        | Y  | Y   | Y               |
| Costa Rica           | Y       | Y        | N  | Y   | N               |
| Dominican Republic   | Y       | Y        | N  | Y   | N               |
| Ecuador              | Y       | Y        | Y  | Y   | Y               |
| El Salvador          | N       | N        | Y  | N   | N               |
| Guatemala            | N       | N        | Y  | N   | Y               |
| Mexico               | Y       | Y        | Y  | Y   | Y               |
| Panama               | Y       | N        | Y  | N   | N               |
| Paraguay             | N       | N        | Y  | N   | N               |
| Peru                 | Y       | N        | Y  | Y   | N               |
| Puerto Rico          | Y       | Y        | Y  | Y   | Y               |

fC1-INH C1-inhibitor function, AgCl-INH C1-inhibitor concentration (antigen), Anti-C1-INH Ab autoantibodies against C1-inhibitor, Y yes, N no

Table 3 Treatments available by country

|                      | Berin-  | Cyn-  | Fira- | Rucon- | Kalbi- | LMWH* | Tranexamic | FFP** | Danazol |
|----------------------|---------|-------|-------|--------|--------|--------|------------|-------|---------|
|                      | ert®    | ryze® | yzyn®| est®   | tor®   |        |            |       |         |
| Argentina            | Y       | N     | Y     | N      | N      | Y      | Y          | Y     | N       |
| Brazil               | Y       | N     | Y     | N      | N      | Y      | Y          | Y     | N       |
| Chile                | Y       | N     | N     | N      | Y      | Y      | Y          | Y     | N       |
| Colombia             | Y       | N     | N     | N      | N      | N      | Y          | Y     | Y       |
| Costa Rica           | N       | N     | N     | N      | N      | N      | Y          | Y     | Y       |
| Dominican Republic   | N       | N     | N     | N      | N      | Y      | Y          | Y     | Y       |
| Ecuador              | N       | N     | N     | N      | N      | Y      | Y          | Y     | N       |
| El Salvador          | N       | N     | N     | N      | Y      | Y      | Y          | Y     | Y       |
| Guatemala            | N       | N     | N     | N      | Y      | Y      | Y          | Y     | Y       |
| Mexico               | Y       | N     | Y     | N      | Y      | Y      | Y          | Y     | Y       |
| Panama               | N       | N     | Y     | Y      | Y      | Y      | Y          | Y     | N       |
| Paraguay             | N       | N     | N     | N      | N      | Y      | Y          | Y     | Y       |
| Peru                 | N       | N     | N     | Y      | N      | Y      | Y          | Y     | Y       |
| Puerto Rico          | Y       | N     | Y     | Y      | Y      | Y      | Y          | Y     | Y       |

*Low molecular weight heparins, **Fresh frozen plasma, Y yes, N no

Conclusion: In Latin America, the availability diagnostic tests of HAE seem to be related to the degree of development of the country and treatment depends of the health laws of each. In some countries first-line treatments are not offered to all patients and even non recommended treatments continue to be used, despite the known adverse effects. The need for better access to diagnosis and treatment of HAE in Latin American patients cannot be underestimated.

P31 Hereditary angioedema in Belarus: epidemiology, clinical characteristics and access to diagnosis and treatment

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Background: Hereditary angioedema due to C1-inhibitor deficiency (C1INH-HAE) is a rare disease caused by deficiency of complement C1-INH. Only few states in developing countries have an adequate management of HAE, but none of them belongs to the former USSR area. This study analyses data from C1INH-HAE patients from Belarus, as example of the region.

Materials and methods: In Belarus were identified 3 referral centers and 2 associations for the evaluation of patients with HAE (Belarusian Research Center for Pediatric Oncology, hematology and immunology; Jeffrey Modell Belarusian Center for Primary Immunodeficiency (Minsk); Health Care Institution with 4 City Children’s clinical Hospital and 10 City clinical Hospital (Minsk); Republican Scientific Center for Radiation Medicine and Human Ecology (Homel); Republican Association of parents of patients with primary immunodeficiency “save the immunity”; republican association “help HAE patients.” Clinical characteristics and access to diagnosis and treatment were collected from 2014 by the Belarusian Research Centre for Paediatric Oncology, Haematology and Immunology in Minsk. A questionnaire about disease’s severity, prophylactic and on-demand therapy was administered to patients. Data about attacks refers to year 2016.

Results: We identified 37 C1-INH-HAE patients, 22 female (59%); 35 type 1 (95%), 2 type 2 (5%) HAE; mean age 31 years (min 4- max 67, median 32); mean age at onset of symptoms 1y (min 1-max 63 y) and a median age at the diagnosis of 29 y (min 1-max 63 y), with a diagnostic delay of 18 y. Mean levels of C1-INH and C4 were 0.04 g/L and 0.03 respectively. The estimated prevalence was 1:255000. Twenty patients accepted to compile the questionnaire and to collect data about attacks. 271 attacks were reported with an attacks mean/patients of 16 (min 3, max75, median 22). 181 were peripheral, 111 abdominal, laryngeal attacks were not reported separately. 8 patients defined their disease severity as severe, 8 moderate, 4 mild. 7 patients used on-demand therapy during attacks (2 patients used fresh-frozen plasma, 2 C1-INH concentrate, 2 icatibant, 1 tranexamic) although C1-INH concentrate nor icatibant are registered; 5 patients used prophylactic therapy with attenuated androgens.

Conclusions: In Belarus the possibility to diagnose and manage patients with C1-INH-HAE is scarce, effective on-demand treatments for acute attacks are still lacking. Nonetheless the growing knowledge of the disease, the registration of patients associations among the Ministry of Health and the identification and development of center of reference are the first steps to guarantee patients prompt diagnosis and adequate treatment.

P32 Comparison of the C1-INH productions of different endothelial cells

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Endothelial cells (ECs) play a key role in edema formation. In the case of hereditary angioedema (C1-INH-HAE), the edema formation is the
consequence of the permeability increasing effect of the elevated bradykinin (BK) level. Because of this key role of ECs in the edema formation, the question emerges: can ECs somehow downregulate the permeability increasing effect of BK?

Although C1-INH production of endothelial cells was reported, the data are not consistent and sometimes are controversial. Since different tissues behave quite differently during edematous attacks, we aimed to map the C1-INH producing capabilities of various ECs. We studied the regulation of C1-INH production by several potential factors – known to trigger edematous attacks- on endothelial cells. We measured the C1-INH mRNA production of primary ECs, such as human umbilical vein and arterial ECs (HUVECs and HUAEcs), human dermal microvascular ECs (HDMECs) and human glomerular ECs (GECs), as well as of EC line, human brain microvascular ECs (HCMEC-D3), and of HepG2 cell line as a positive control. After this we used different stimuli: thrombin (TR), BK, TGF-beta, interferon gamma (IFNg), and TR and BK together, and measured the change of mRNA and protein levels of C1-INH after 24 and 48 h. We used qPCR to measure mRNA levels, and an in-house ELISA to detect C1-INH protein production.

We found that all investigated ECs can produce C1-INH at mRNA level. Moreover, HUVEC and HDMEC produced and secreted C1-INH into the cell culture supernatant (2.41 ± 0.34 ng/10^5 cells for HUVEC, 0.496 ± 0.018 ng/10^5 cells for HDMEC). IFNg treatment caused a significant increase in the expression of C1-INH at both mRNA and protein level in HUVECs and in HDMECs. Although TGF-beta, and BK together with TR or TNF, and in some HUVEC lines BK alone could also induced the expression of C1-INH, these changes were minor compared to the effect of IFNg.

All endothelial cell types produced C1-INH, which suggests that ECs can actively regulate the plasma serine protease cascades, thus the pathophysiological process of angioedema.

The difference between C1–INH expression upon stimulation with several potential trigger factors highlights that initiation routes of HAE attacks may implicate distinct predisposition for edema formation as well as to distinct efficiency to resolve the attacks. Taken together, we propose that the pathophysiology of HAE attacks may depend on the integrative function of bradykinin metabolism, C1-INH metabolism and the actual phenotype of endothelial cells. Supported by OTKA 112110.

P34 Increased fibrinolysis-induced bradykinin formation in hereditary angioedema confirmed using stored plasma and biotechnological inhibitors

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Background: We recently investigated the pathways of immunoreactive bradykinin (iBK) formation in fresh blood of normal volunteers and of patients with hereditary angioedema due to C1-esterase inhibitor deficiency (HAE-1/2) [1]. Since we did not detect iBK formation following platelet or neutrophil activation in whole blood, we adapted the techniques to small volumes (200 μl) of previously frozen plasma and further analyzed the mechanisms of iBK formation with additional biotechnological inhibitors.

Materials and methods: Each experimental point was obtained using 200 μl of thawed citrated plasma transferred to a 1.5 ml conical test tube; activators or inhibitors were added to test various pathways of kinin generation. All tubes contained enalaprilat to protect BK from rapid inactivation. The tubes were incubated under rotary agitation in a pre-equilibrated (37 °C) Thermomixer apparatus. The ethanol extraction, sample evaporation and enzyme immunoassay were then performed as described [1].

Results: iBK formation was observed under stimulation with tissue kallikrein (KLK1, 10 nM), the particulate material Kontakt-APTT (concentration reduced to 2% v/v) or recombinant tissue plasminogen activator (tPA, 169 nM), with little background in unstimulated plasma incubated for up to 2 h. Plasma samples from HAE-1/2 patients responded earlier to tPA than those from controls, as previously reported with whole blood. Lanadelumab inhibited iBK formation induced by Kontakt-APTT and tPA. A highly specific plasmogen inhibitor, DX-1000, abolished tPA-induced iBK formation in plasma but had no
Small effect against Kontact-APTT, confirming the role of fibrinolysis in tPA-induced kinin formation. The anti-lanadelumab neutralizing antibody M293-D02 reversed the inhibitory effects of lanadelumab.

Conclusions: Frozen plasma is a suitable material for measuring iBK formation kinetics, with possible applications such as investigating the effect of rare disease states on the kallikrein-kinin system and monitoring the effect of HAE prophylactic treatments. The instability of the kallikrein-kinin system in HAE-1/-2 may reside in the upstream fibrinolytic system. Common attack triggering factors are compatible with this hypothesis.

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Consent to publish: The CHU de Québec-Université Laval ethical review board approved the study (file no. 2018-3857). All study subjects gave written informed consent.

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P35
High plasma exposures of KVD900 achieved in First in Human study markedly inhibit plasma prekallikrein activation; early blockade of plasma kallikrein (PKa) may halt attacks in hereditary angioedema (HAE) by reducing contact system activation

Introduction: Early treatment of acute attacks in HAE is associated with improved clinical outcomes. These clinical benefits are attributed to the reduction of plasma kallikrein-mediated high molecular weight kininogen (HK) cleavage and bradykinin action. The effects of PKa inhibition on contact system activation has received relatively little attention. This study examines the effects of KVD900, a rapidly acting oral PKa inhibitor with high clinical exposure, on plasma prekallikrein cleavage and PKa activity during contact system activation.

Methods: Single ascending doses of KVD900, a potent and selective PKa antagonist icatibant in hereditary angioedema – a real-life study

Objective: The synthetic peptide icatibant – a bradykinin B2-receptor antagonist administered subcutaneously – is intended for the acute treatment of angioedema attacks in patients with hereditary angioedema with C1 inhibitor deficiency (C1-INH-HAE). Our study analyzed the efficacy and the adverse effects of icatibant, as well as patient satisfaction with treatment.

Materials and methods: We analyzed 546 angioedema attacks experienced by 40 C1-INH-HAE patients. By completing a questionnaire, the patients recorded treatment-related information. The severity of the individual angioedema attacks was graded. The patients also recorded any adverse effects, and rated their satisfaction with the treatment.

Results: The distribution of the analyzed attacks: 278 subcutaneous, 178 gastrointestinal, 8 upper airway, and 82 multiple locations. Icatibant was injected at a median interval of 65.0 min after the onset of the angioedema attack, the severity of which was 66 (median), according to the visual analog scale (VAS). The symptoms started to improve 35.0 min (median) and resolved 426.0 min (median) after treatment. The time between the administration of the injection and the resolution of the attack was correlated with time until injection (r = 0.2322, p < 0.0001). On 39 occasions, the symptoms failed to improve or to resolve after the administration of a single dose. A second dose was administered in 23/39 instances, and this eliminated the symptoms in 20/23. The angioedema attack reoccurred in 9.3% of the instances (n = 51). We did not find any significant difference in time to injection between icatibant-responsive and non-responsive vs. rebound attacks. However, the latter occurred in 16.32% of instances when the time to injection was 30 min or less, and in 7.71% when it was longer than 30 min. Thirty-three patients experienced a skin reaction at the injection site. On average, satisfaction with the treatment was 88.65 on a 100-mmVAS, and a mean score of 9.05 was assigned on a scale of 1 to 10.

Conclusion: Icatibant is an effective and safe medicine with a rapid onset of action in C1-INH-HAE. We can conclude that early treatment resulted in quicker resolution of the symptoms, however 30 min or less administration time showed higher percent of rebound attacks. Local skin reactions were common, but possibly drug-related systemic adverse effects did not occur.

This study was supported by OTKA K124557.

P36
The treatment of angioedema attacks by bradykinin B2-receptor antagonist icatibant in hereditary angioedema – a real-life study

Objective: The synthetic peptide icatibant – a bradykinin B2-receptor antagonist administered subcutaneously – is intended for the acute treatment of angioedema attacks in patients with hereditary angioedema with C1 inhibitor deficiency (C1-INH-HAE). Our study analyzed the efficacy and the adverse effects of icatibant, as well as patient satisfaction with treatment.

Materials and methods: We analyzed 546 angioedema attacks experienced by 40 C1-INH-HAE patients. By completing a questionnaire, the patients recorded treatment-related information. The severity of the individual angioedema attacks was graded. The patients also recorded any adverse effects, and rated their satisfaction with the treatment.

Results: The distribution of the analyzed attacks: 278 subcutaneous, 178 gastrointestinal, 8 upper airway, and 82 multiple locations. Icatibant was injected at a median interval of 65.0 min after the onset of the angioedema attack, the severity of which was 66 (median), according to the visual analog scale (VAS). The symptoms started to improve 35.0 min (median) and resolved 426.0 min (median) after treatment. The time between the administration of the injection and the resolution of the attack was correlated with time until injection (r = 0.2322, p < 0.0001). On 39 occasions, the symptoms failed to improve or to resolve after the administration of a single dose. A second dose was administered in 23/39 instances, and this eliminated the symptoms in 20/23. The angioedema attack reoccurred in 9.3% of the instances (n = 51). We did not find any significant difference in time to injection between icatibant-responsive and non-responsive vs. rebound attacks. However, the latter occurred in 16.32% of instances when the time to injection was 30 min or less, and in 7.71% when it was longer than 30 min. Thirty-three patients experienced a skin reaction at the injection site. On average, satisfaction with the treatment was 88.65 on a 100-mmVAS, and a mean score of 9.05 was assigned on a scale of 1 to 10.

Conclusion: Icatibant is an effective and safe medicine with a rapid onset of action in C1-INH-HAE. We can conclude that early treatment resulted in quicker resolution of the symptoms, however 30 min or less administration time showed higher percent of rebound attacks. Local skin reactions were common, but possibly drug-related systemic adverse effects did not occur.

This study was supported by OTKA K124557.
P38 Characterizing the minority of hereditary angioedema attacks that require more than a single injection of icatibant
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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P38

Background: The Icatibant Outcome Survey (IOS; NCT01034969), an ongoing international observational registry, monitors the safety and effectiveness of icatibant, a bradykinin B2 receptor agonist approved for the treatment of hereditary angioedema with C1-inhibitor deficiency (C1-INH-HAE). We evaluated reinjection data from more than 6000 attacks that required icatibant reinjection.
Methods: Descriptive, retrospective analyses were conducted on data obtained from patients reporting ≥1 icatibant-treated attack between July 2009 and January 2019 to determine patient and attack characteristics associated with icatibant reinjection. For these analyses, patients were stratified by number of icatibant injections administered per attack.
Results: Included in this analysis were 501 patients with C1-INH-HAE (HAE type I and II; 58.9% females) who had 6047 icatibant-treated attacks. Most attacks were treated with a single injection of icatibant. Of 405 attacks requiring icatibant reinjection, 224 (55.3%) were abdominal and 197 (48.6%) were severe or very severe (in any location). Few laryngeal attacks (21/225 [9.3%]) required reinjection. Reinjection was associated with longer median attack duration. The second injection was administered 12 h after the first dose in most attacks (86.3%). BMI (<25 kg/m² vs ≥25 kg/m²) was not a significant predictive factor for icatibant reinjection (P = 0.169). There was no significant difference for C1-INH rescue medication use after one icatibant injection (5.8%) versus more than one injection (5.4%; P = 0.751).
Conclusions: Reinjection was required in less than 10% or 1 in 10 attacks treated with icatibant. In these very severe or severe attacks, patients were more likely to require reinjection. Most patients requiring more than one dose waited at least 12 h after the first dose to reinject.
Trial registration: The Icatibant Outcome Survey, NCT01034969

P39 Long-term prophylaxis in hereditary angioedema patients followed in ITALian Centers for Angioedema (ITACA) and enrolled in HAE Global Registry (HGR)
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Long-term prophylaxis (LTP) is targeted to prevent the recurrence of acute attacks in patients affected by hereditary angioedema due to C1-inhibitor deficiency (C1-INH-HAE). Approved for LTP in Italy are plasma derived C1-INH (pdC1-INH), attenuated androgens (dana-zol/stanozolol), tranexamic acid. According to guidelines, patients start LTP when disease control is not satisfactory with on-demand treatment. This study aims at evaluating prevalence and efficacy of LTP at ITACA centers.
Four hundred and four patients with C1-INH-HAE (mean age 51.2 yo, range 8 to 88 yo) at 6 ITACA centers are listed in the HGR, a multi-center disease registry (ClinicalTrials.gov NCT03828279). The study considered patients on stable LTP at July 1, 2018. ITACA policy for stable LTP is presence of more than one attack per month. All patients in the study recorded angioedema attacks prospectively; the number of attacks between July-December 2018 was analyzed.
Primary efficacy endpoint was number of attack-free days. Side effects were recorded during the observation period. Ninety nine patients of 404 (24.5%) were on LTP (61 F, 38 M); 60 with danazol, 9 with stanazolol, 22 with pdC1-INH and 8 with tranexamic acid. Androgen posology was highly variable, from 200 to 2800 mg/week (mean 860) for danazol and from 4 to 14 mg/week for stanozolol (mean 9.2). The mean dose for pdC1-INH was 1985 IU/week (from 1000 IU to 6000 IU/week). Tranexamic acid dose fluctuated between 0.7 and 21 g/week (mean 6.54 g/week). 38/60 patients (63.3%) on danazol LTP were attack-free throughout the 6 months observation period, the remaining 22 (36.6%) had 139 HAE attacks, with a mean frequency of 6.3 attacks/semester ranging from 1 to 30 attacks/semester. Finally, 3/8 patients (37.5%) treated with tranexamic acid were attack-free and 5 had total of 20 attacks in the semester (range 1–9; mean 4).

Conclusions: In our population, attenuated androgens represent the main approach to LTP. All LTP approaches are effective (attacks reduction over 50%) in the large majority of patients. Pooling patients’ data in HGR allows rapid evaluation of drug efficacy, improves knowledge on the disease and facilitates treatment personalization.

P40
Shortage in France of plasma derived C1-INH concentrates: state of play and consequences for patients
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Introduction: In France, several treatments are available for Hereditary angioedema: two plasma derived C1Inh (Berinert, Cinryze), a recombinant C1Inh (Ruconest), Firazyr, danazol and tranexamic acid. Since 2017, France has been experiencing a shortage of blood products that affect C1Inh concentrates. This shortage has triggered a healthcare crisis that has altered the care of patients. The CREAK (National Reference Center for Angioedema) and the ANSM (National Agency for Health and Drugs) have had to take exceptional measures, including those to offer Ruconest, including pregnant women and children, as well as long term prophylaxis.

Methods: CREAK conducted a survey within its network and referring physicians to assess the impact of this health crisis.

Results: The shortage induced a therapeutic change in 61 patients, half of whom had a very severe illness and one-third a severe illness. For all these patients, it was a modification of their long term prophylaxis with C1Inh concentrate (Cinryze or Berinert). In 70% of cases, patients were switched to Ruconest with an imbalance of pathology in 40% of cases. 40% of doctors could not apply international recommendations on short-term prophylaxis. 5 pregnant women had difficulties of care because of this shortage. The doctors appreciated the help of CREAK in 80% of the cases and followed its recommendations. The image of laboratories has been altered with 70% of doctors who no longer trust them. 70% of physicians consider that the lives of their patients have been jeopardized.

Conclusion: The shortage of plasma derived C1Inh has impaired patients’ quality of life and sometimes put their lives at risk. CREAK and ANSM had to advocate off-label uses of certain products. Since October 2018, France has obtained the availability of lanadelumab for these severe patients in order to depend less on plasma derived C1Inh. The first encouraging results give new hope to patients and doctors.

P41
Limelight on erythema marginatum: a review of clinical features and the introduction of a new management strategy in hereditary angioedema
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Allergy, Asthma & Clinical Immunology 2019, 15(Suppl 4):P41

Objective: To assess the incidence and clinical characteristics of erythema marginatum (EM) along with the efficacy of the therapies administered during episodes of EM to patients receiving follow-up care for C1-inhibitor-deficient hereditary angioedema at the Hungarian Angioedema Reference Center.

Materials and methods: During the first stage of our study, we surveyed the incidence of EM using the Erythema Marginatum Basic Questionnaire (EMBQ) developed by our team. In the second stage, we evaluated therapeutic efficacy with the Erythema Marginatum Detailed Questionnaire (EMDQ).

Results: According to the EMBQ, 72 out of 134 C1-INH-HAE patients from 60 families (mean age 40 years, min.: 6 years, max.: 82 years) experienced EM during their lifetime. Forty-three of these 72 patients were females (59.7%). In 41.7% of families with C1-INH-HAE, EM occurred in all family members, whereas 20% of the families have never experienced this skin lesion. In these 72 patients, EM appeared in the following locations: on the upper extremities in 79.2%, on the chest in 63.9%, on the lower extremities in 33.3%, on the back in 25%, in the abdominal region in 19.4%, and on the face in 12.5%. EM first occurred at the age of 17 years on average, and as an isolated cutaneous lesion in 47.2% of the patients. EM was followed by an acute episode of hereditary angioedema in 8 out of 10 instances. As shown by the EMDQ, 16 C1-INH-HAE patients (15 females and 1 male) administered the following treatments on 160 occasions: plasma-derived C1-INH concentrate (pdC1-INH – n=73), recombinant C1-INH concentrate (rhC1-INH – n=79), and icatibant (n=8). The onset of the HAE attack was prevented by pdC1-INH in 94.5%, by rhC1-INH in 93.6% and by icatibant in 50% of cases.

Conclusion: EM is characterized by a variegated clinical picture, as regards both the time of the onset and the location of symptoms. Being an objective prodromal sign, its occurrence creates an opportunity to administer acute treatment for HAE attacks earlier and thereby to prevent their onset. This new therapeutic strategy is cost-effective, and it might significantly improve the patients’ quality of life. This study was supported by OTKA K124557.
were women, with HAE-C1INH type I and mean age of 47 years. Five patients (58%) showed signs of periodontal disease. Forty-three dental procedures were performed, the most common being tooth extractions, dental restorations and supra-gingival removal of calculus. Six patients (67%) were submitted to dental procedures without modification of long-term prophylaxis, whereas an increased dose of AA and/or C1-INH concentrate was administered before the dental procedures in three patients (33%), especially in those undergoing dental extractions. Angioedema attacks were not observed.

Conclusion: This is the first large series on dental management of Brazilian patients with HAE. Interdisciplinary evaluation is crucial for proper management of patients with HAE.

P43
Development and validation of the self-efficacy assessment questionnaire in the management of hereditary angioedema for patients and family caregivers (HAE-SES)
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Purpose: Research suggests that most patients and family caregivers lack competence in the management of Hereditary Angioedema (HAE). Self-efficacy is considered to be paramount in the development of people's competence. The aim of this study is to design and psychometrically assess the pilot version of a scale to measure self-efficacy in the management of HAE by patients and their family caregivers.

Materials and methods: The study followed an observational, cross sectional design. Fifty-one participants (30 patients and 21 family caregivers) and 16 experts participated in the study. Firstly, a 32-item version of the scale was developed by the researchers. Secondly, a panel of 16 experts in the topic were asked to score the relevance of each item using a Likert-type scale (1 = not relevant; 2 = somewhat relevant; 3 = quite relevant; 4 = very relevant) in order to determine its content validity. The individual item’s content validity index (i-CVI) and the scale’s content validity index (S-CVI) were calculated. Thirdly, the tool’s reliability was tested by exploring its internal consistency calculating the Cronbach alpha coefficient (a) once 51 participants had completed the scale.

Results: After the analysis, the pilot version of the HAE-SES was comprised of 29 items grouped into 3 dimensions: “Identification of symptoms and initial decision making”, “Treatment management” and “Post-treatment crisis management and waste disposal”; The HAESES’s content validity proved to be excellent (i-CVIs ranged between 0.8 and 1). All the dimensions of the questionnaire obtained CVI scores higher than 0.9, with an overall score of S-CVI = 0.93. In addition, the excellent reliability of the HAE-SES was evidenced by its internal consistency (a = 0.957).

Conclusion: The pilot version of the HAE-SES showed excellent psychometric properties to measure self-efficacy in the management of HAE for patients and family caregivers, allowing the validation process to continue.

P44
Headache as a symptom of hereditary angioedema
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Background: Headache is among the most common symptoms, and a possible accompanying sign of a number of diseases. In the family of primary headache disorders, bradykinin — a vasodilatory substance, which also enhances vascular permeability — might be involved in the pathomechanism of migraine. Bradykinin is the key vasoactive mediator responsible for the symptoms of hereditary angioedema resulting from C1-inhibitor deficiency (C1-INH HAE). In addition to the frequent recurrence of subcutaneous and submucosal edema formation, angioedema has been reported to occur in rare locations including the central nervous system, and cause headache along with neurological symptoms.

Conclusion: This study assessed the incidence and properties of headache, as well as the possible efficacy of HAE therapies in its relief.

Materials and methods: All C1-INH HAE patients attending the annual follow-up visits between 2014 and 2018 were enrolled because since 2014, our practice for taking the medical history also involves administering a questionnaire on headache.

Results: In the past five years, 156 patients (63 males and 93 females, mean age 39.2 (1.2 to 86.7 years) returned to the National Angioedema Reference Center for follow-up visits yearly. During this period, 84 of these 156 patients – mostly (71.4%) women – reported headache. Migrain-like features were identified in 17 patients including 13 women.

During follow-up, C1-inhibitor treatment was administered for 39 migraine-like episodes. Five of the six patients (1 male and 5 female, with a mean age of 38.1 years) received plasma-derived C1-INH concentrate (500 to 1500 IU), whereas a single patient was treated with 500 IU plasma-derived and 4200 IU recombinant C1-inhibitor. Except for a single episode during which the patient did not experience any therapeutic effect, the symptoms exhibited substantial regression, and resolved completely in 2 h. No adverse effects occurred. According to the patients, without treatment with C1-INH, migraine-like headache persisted for about half a day or for a full day, and this had a rather negative effect on their well-being.

Conclusion: As shown by our experience accumulated during the past 5 years in patients with C1-INH HAE, treatment with C1-INH might prove beneficial in the management of episodic migraine-like headache unresponsive to treatment and accompanied by unilateral neurological signs. Further studies relying on imaging modalities or other biomarkers are necessary to decide whether these episodes are the manifestations of HAE, or the components of a separate, co-occurring disorder.

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P45
Short-term prophylaxis in patients with angioedema due to C1-inhibitor deficiency undergoing dental procedures
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Background: Patients affected by angioedema due to hereditary and acquired C1-inhibitor deficiency (C1-INH-HAE and C1-INH-AAE, respectively) report troubles in accessing dental care. In patients with C1-INH deficiency, dental procedures can trigger life-threatening laryngeal attacks. Dentists often are not familiar with the management of angioedema attacks and do not dare to treat dental disease in these patients. As a consequence, patients with C1-INH deficiency suffer a lack of proper dental care. Short-term prophylaxis (STP) is recommended by international guidelines [1] before dental procedures, but prospective clinical trials are lacking. It is well known that infections, also localized in the oral cavity, may trigger angioedema attacks [2]. Therefore, improving dental care may be a useful strategy to reduce the frequency of angioedema attacks [3]. The primary endpoints of this study were the assessment of the presence of hurdles in receiving dental care and the effectiveness of short-term prophylaxis...
in preventing angioedema attacks. The secondary endpoint was the impact of dental care in angioedema course.

**Materials and methods:** All patients affected by angioedema due to C1-INH deficiency treated in the dentistry outpatient department of ASST Fatebenefratelli Sacco hospital in the period 2009–2017 were considered for the analysis. Data on oral status, dental procedures, STP, and disease course were collected prospectively.

**Results:** Twenty-nine patients were analysed (27 with C1-INH-HAE and 2 with C1-INH-AAE). Of them, 58.6% reported hurdles in accessing dental care. At the first visit, 55.17% patients had moderate-to-severe oral disease. Sixty-three dental procedures were performed in 20 patients. Fifty procedures were preceded by STP with plasma derived C1-INH (pdC1-INH) in patients with/without long-term prophylaxis (LTP). One procedure in one C1-INH-HAE patient was preceded by short course of androgens (danazol). Post-procedural attack occurred in 2 patients. One C1-INH-HAE patient undergoing a tooth extraction without STP/LTP experienced a laryngeal attack. The other post-procedural attack occurred in a C1-INH-AAE patient with anti-C1-INH antibodies with STP. Angioedema course did not worsen in any patient after dental care, but improved in 4 of them.

**Conclusions:** Patients had encountered hurdles in accessing dental care, responsible also for the high rate of moderate-to-severe oral disease at the first visit. STP is protective towards attacks after dental procedures in C1-INH-HAE patients. In C1-INH-AAE patients with anti-C1-INH antibodies, STP with pdC1-INH may be nonprotective. Treating oral diseases tends to reduce the frequency of attacks.

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**Consent to publish:** Informed consent to publish has been obtained by CSL Behring, Italy.

**Ethics approval:** The study was approved by Comitato Etico Interaziendale Milano Area A in February 26, 2016 with protocol number 3431/2016.

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**P46**

**Mass spectrometry based screening for hereditary angioedema disease**

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**Background:** Hereditary angioedema (HAE) is an autosomal dominant disease resulting from mutations in the SERPING1 gene, leading to the deficient (type 1) or nonfunctional (type 2) C1-inhibitor protein. Clinical manifestation in all HAE types include acute attacks of edemases affecting the upper airway, face, extremities, genitals, and gastrointestinal system. In 93% of patients with HAE-caused abdominal pain this is the only manifestation of the disease. Centogene’s epidemiological data show a HAE frequency of 1:17,525 in Turkey and 1:36,600 in Germany. Centogene conducts several HAE related studies to update the prevalence of the disease and develop specific biomarker.

**Analytical methods:** The standard diagnosis procedure of HAE in our laboratory is based on Dry Blood Spot (DBS) filtercard technology that simplifies both the sample collection and the logistics. The diagnostic workflow is a two-tier approach: firstly, the complement proteins are quantified by tandem mass spectrometry (MS/MS) and, secondly, the results are confirmed by genetic analysis. The quantification of the complement proteins is performed in situ by quantifying unique complement peptides for each protein (Complement C4 and Complement 1-Inhibitor peptides) using LC/MS/MS—liquid chromatography multiple reaction monitoring mass spectrometry.

**Epidemiological studies:** Centogene currently runs the “Epidemiological Study on Hereditary Angioedema” (EHA Study) that aims to test subjects with unclear abdominal pain for HAE and to search for new biomarkers in the blood of positive patients. In a total of 4 years, 5000 participants from 7 countries worldwide will be included. To further strengthen the biomarker development, the “Biometabolic HAE” Study (BioHAE) will include diagnosed HAE patients and check for potential new biomarkers via LC/MS/MS. The “Hereditary Angioedema Kininogen Assay” Study (HAEKA) plans to characterize cleaved high molecular weight kininogen (cHMWK) as a promising biomarker and to further investigate the effect of a novel medication (lanadelumab) in this setting. Participants will donate blood in timely intervals during an edema attack to understand the cHMWK kinetics in the HAE course.

**Conclusion:** The 2 tier approach (LC/MS/MS followed by confirmation using NGS) was proven to be a rapid and precise way of identifying type 1 and type 2 HAE patients. Several studies by Centogene will use these techniques to search and characterize novel HAE specific biomarkers.

The studies were approved by University Rostock’s Ethics Board, approval numbers A2017-0007, A2018-0057, and A2019-0046.

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