in diagnosis, although a more aggressive biology cannot be discarded.

Thyroid

THYROID CANCER CASE REPORTS I

Thyroid Extranaonal Diffuse Large B-Cell Lymphoma in Setting of Gastric Large B-Cell Lymphoma
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SUN-475

Background: We report a case of concomitant thyroid and gastric diffuse large B-cell lymphoma (DLBCL). Clinical Case: A 73 year-old man presented with melena of 2 days duration. EGD revealed a 1.5 x 0.7 cm gastric ulcer. Biopsy revealed marked chronic active gastritis with glandular atypia, positive for H. pylori. Repeat biopsy of the same ulcer post H. pylori treatment demonstrated high-grade B-cell lymphoma. Immunostains were positive for CD20, CD10, BCL6, cyclin D and negative for CD 23 and CD 30. FISH was negative for gene rearrangement involving BCL-6/BCL-2 and C-MYC. Staging PET/CT showed hypermetabolic regions involving the stomach wall, large left neck mass and right lung focus. The neck focus had a max SUV 48.5 (8 fold greater than the mach wall, large left neck mass and right lung focus. The arrangement involving BCL-6/BCL-2 and C-MYC. Staging PET/CT showed hypermetabolic regions involving the stomach wall, large left neck mass and right lung focus. The focus had a max SUV 48.5 (8 fold greater than the other sites), that localized to a soft tissue mass measuring 4.9 x 2.7 cm. FNA of the left-sided thyroid mass showed diffuse large B-cell lymphoma with immunostain positive for CD20 and negative for cytokeratin AE1-3, PAX8, and CD 10. The morphology was similar to that of abnormal lymphoid cells in the gastric biopsy. Patient had normal thyroid function tests but positive autoimmune thyroid disease markers (TPO and anti-TG antibody). Treatment with R-CHOP resulted in shrinkage of neck mass. Conclusion: Hypermetabolic regions revealed by PET/CT at distant sites from primary tumor require further evaluation including biopsy as indicated. Clinical correlation and response to chemotherapy can provide supplemental information in the overall assessment of the disease process.

Pediatric Endocrinology

PEDIATRIC ENDOCRINE CASE REPORTS II

A 2-year Old Girl with Turner Syndrome and Neurofibromatosis Type 1
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MON-072

A 2-year old girl with Turner syndrome and Neurofibromatosis type 1
Introduction: Turner syndrome (TS) occurs due to loss of either all or part of the X chromosome, in some or all the cells of the body. The most consistent features of TS are short stature and premature ovarian failure. Neurofibromatosis type 1 (NF1) is an inheritable in an autosomal dominant manner tumor predisposition syndrome and is caused by loss-of-function mutations in the tumor suppressor NF1 gene (neurofibromin 1). Literature review indicated rare cases with NF1 and TS (1). We report the sixth girl with mosaic TS and NF1 who presented with optic nerve glioma.

Case report: A 2-year-old female presented to us due to short stature. Her height was 2,5 SD lower than the mean parental height curve, and her bone age was delayed only by 3 months. She already had a normal (46XX) peripheral blood karyotype (70 mitoses). She had abnormal body proportions and with short limbs with unremarkable café au lait spots. Additionally, to the short stature laboratory investigation we ordered a gene panel to exclude hypochondroplasia, and
a Karyotype in fibroblasts culture from oral cavity sample. The results revealed low IGF-1 and mosaic TS in 14%. We preformed 2 provocative tests which revealed low growth hormone peak < 5 ng/ml. A brain and pituitary MRI to exclude pituitary lesions or structural abnormalities revealed gliomas of the optic chiasma and the right optic nerve with characteristic NF1 “spots” (regions of signal abnormality in T2 sequences) involving the basal ganglia, cerebellum and the right temporal lobe. DNA sequencing targeted to a gene panel related to NF1 and NF2 revealed a novel de novo heterozygous NF1 gene mutation in exon 28 [3764A>G];[=]p.[Gln1255Arg].

**Discussion:** NF1- Gliomas are most commonly seen in young children, (mean 4.5 years). Only 1/3 of affected children will require therapeutic intervention. However early diagnosis, of optic gliomas is important. Our patient was completely asymptomatic by the time of diagnosis and no other symptom or sign of NF1 was apparent. Ophthalmologic examination was normal, but visual electrophysiologic testing was abnormal as far the right optic nerve is concerned. The oncology team decided to preform chemotherapy. In TS impaired growth is related to resistance in GH. Some studies suggested that there could be a relationship between GHD and NF1 even in the absence of an organic pituitary damage. In our patient it has been decided not to treat with GH and closely track the patient’s growth.

**Conclusion:** Coexistence of NF1 with TS is rare. Awareness is needed as early identification and treatment of CNS gliomas can prevent visual loss and severe co-morbidities. 1. Rare Presentation of Neurofibromatosis and Turner Syndrome in a Pediatric Patient. Pediatr Rep. 2017 Jun 26; 9(2): 6810

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**Reproductive Endocrinology**

**MALE REPRODUCTIVE HEALTH - FROM HORMONES TO GAMETES**

**Testosterone Reduces Atherosclerosis and Plaque Specific Inflammatory Markers in the ApoE-/- Mouse Model**

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**SAT-041**

Low serum testosterone in men is an established cardiovascular risk factor and epidemiological evidence demonstrates an association between low testosterone and with coronary events. Clinical evidence suggests that testosterone therapy (Tth) can improve key cardiovascular risk factors in men and surrogate measures of atherosclerosis, the chronic inflammatory process underlying cardiovascular disease. Atherosclerotic plaque-specific testosterone actions are not fully understood. The present study investigates the influence of testosterone on mediators of vascular inflammation and plaque burden in an *in vivo* model of atherosclerosis. ApoE-/- mice were either sham operated, castrated or castrated and received fortnightly intramuscular injections of physiological doses of testosterone (mixed testosterone esters, Sustanon 100) to create 3 experimental groups; normal testosterone, testosterone deficient and testosterone replaced respectively. All groups were fed a high-fat ‘Western’ diet for 16 weeks. Lipid deposition in the aortic root was assessed by Oil Red O as an indication of atherosclerotic burden. Plaque composition was assessed immunohistochemically for indicators of stability including collagen content via Masson’s trichrome, and α-smooth muscle actin (αSMA) as well as markers of inflammation including vascular cell adhesion molecule-1 (VCAM-1), intracellular adhesion molecule-1 (ICAM-1), endothelial-leukocyte adhesion molecule 1 (E-Selectin), and a pan monocyte/macrophage marker (MOMA2). Testosterone deficient castrated mice had significantly increased lipid accumulation in the aortic root compared to testosterone replete sham-operated littersmates (45% intima-media area vs 40%, p<0.05). Tth in castrated mice reversed this effect (39%, p<0.05). Plaque stability was not altered between groups. MOMA2 staining indicated increased infiltration and localisation of monocytes/macrophages in the plaques of castrated mice compared to sham-operated (positive staining (% of plaque) 77% vs 59%, P=0.062) and Tth treatment reduced this (77% vs 63%, P=0.1). Vascular cell adhesion molecule-1 (VCAM-1) and intracellular adhesion molecule-1 (ICAM-1) expression were reduced in castrated mice receiving Tth compared to castrated mice receiving saline (33% vs 46%, p<0.05; and 39% vs 58%, P=0.084 respectively). No significant difference in expression of E-Selectin and αSMA were observed between groups. These findings demonstrate that low testosterone increases aortic root lipid deposition and inflammatory composition in a mouse model of atherosclerosis. Increasing testosterone levels through Tth decreases plaque specific inflammatory markers and atherosclerotic burden. This indicates an anti-inflammatory mechanism by which testosterone can protect against the development and progression of atherosclerosis to reduce cardiovascular risk in men.

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**Diabetes Mellitus and Glucose Metabolism**

**TYPE 1 DIABETES MELLITUS**

**A Case of Autoimmune Polyglandular Syndrome Type 2 and Guillain-Barré Syndrome**

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**SAT-683**

Background: Autoimmune polyglandular syndrome type 2 (APS2) is defined by the occurrence of two or more autoimmune diseases, with Addison’s disease being most prevalent, and autoimmune thyroid disease and type 1 diabetes mellitus also being common. Guillain-Barré syndrome (GBS) is an acute inflammatory demyelinating polyradiculopathy that is also autoimmune in nature, resulting in ascending muscle weakness or paralysis.

Clinical Case: A 49 year old female with past medical history of vitiligo, subclinical hyperthyroidism, and Guillain-Barré Syndrome presented with a 1 month history of paresthesia in her left arm and legs. On exam she was able to raise her arms but had some weakness in the distal lower extremity. Her myalgia was progressive and the patient presented to the hospital with myoglobinuria, fever, and rash. Blood tests revealed an elevated AMO (700 mg/dL) and ANA. An MRI of the brain and spinal cord was normal. The patient was diagnosed with GBS, and was treated with IVIG and physical therapy. She was completely asymptomatic by the time of diagnosis and no other symptom or sign of NF1 was apparent.