SAT0615 THE RS11803505 IL-23R POLYMORPHISM IS A RISK FACTOR FOR THE DEVELOPMENT OF MSUS-DETECTED EROSIONS IN PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSUS

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Background: Articular involvement is a frequent and invalidating manifestation in patients with Systemic Lupus Erythematosus (SLE) ranging from arthralgia to erosive arthritis. Bone erosions are far less common in SLE but require a different management and may display a distinct pathogenesis. Musculoskeletal ultrasonography (MSUS) has proven useful in the assessment of inflammatory arthritis and it is now known that MSUS is more sensitive than radiography in detecting bone erosions.

Objectives: We aimed at analyzing, in a large monocentric cohort of SLE patients who suffered from arthritis, the relationship between 31 polymorphisms in 18 genes previously associated with SLE or other autoimmune/inflammatory arthritis [1] and the presence of bone erosions detected by MSUS.

Methods: Consecutive SLE patients who complained articular involvement (defined by 1997 ACR criteria, i.e. ≥ 2 joints with tenderness, swelling or effusion) were enrolled. Patients RF+ve and/or anti-CCP+ve were excluded. Thirty-one SNPs in 18 genes (STAT4, IL10, IL23R, IRAK1, PSORS1C1, HCP5, MIR146a, PTPN2, ERAP1, ATG16L1, IRGM, TRAF3IP2, VKORC1, CCR5, ATG5, TRAF6, TNFSF4, CD40) were genotyped by allelic discrimination assays. Bilateral high-resolution MSUS of the hands and feet was performed. Erosion number was recorded, the presence of erosive damage was assessed with a dichotomous score.

Results: Clinical and laboratory features of the 95 enrolled patients are described in table.

Clinical and laboratory features	N.	%
Sex	92	96.8
Age (mean $\pm$ SD)	43.38 $\pm$ 11.49	
Age at diagnosis (mean $\pm$ SD)	31.14 $\pm$ 11.83	
Anti-dsDNA	69	72.6
MSUS Erosive score (mean $\pm$ SD)	0.44 $\pm$ 1.34	
MSUS Erosions (yes/no)	15/80	15.8

The presence of MSUS-detected erosions was demonstrated in 15/95 (15.8%) patients, with a mean $\pm$ SD of 0.44 $\pm$ 1.34. Bone erosions were more frequently observed at II metacarpophalangeal joint of the hands (29.4%), followed by the V metatarsophalangeal joint of the feet (16.1%). An association was found between anti-dsDNA and the presence of bone erosions ($P=0.009$, Fisher's exact test); indeed, all the 15 patients (100%) with bone erosions were anti-dsDNA positive vs 54/80 (67.5%) of those without erosions. After Bonferroni correction, IL-23R rs11803505 was found to be associated with the presence of MSUS-detected bone erosions both at genotypic ($P_c=0.045$) and at the allelic level [$P_c=0.022$, OR=8.27 (95% CI 1.02–66.90)]. The association of bone erosions with anti-dsDNA and IL-23R rs11803505 genotype was confirmed in a logistic regression analysis including also age, sex and disease duration ($P=0.005$ and $P=0.02$, respectively).

Conclusions: Our study is the first reporting a contribution of IL-23R gene to MSUS-detected erosive damage in SLE. IL-23R gene is a key player in several inflammatory conditions such as psoriatic arthritis, and variants of this gene have already been associated with peripheral erosive disease [2]. Moreover, we confirmed an association between anti-dsDNA and bone erosions in SLE [3], suggesting that the presence of these antibodies is linked with a more aggressive inflammatory pattern of disease.

References:

- [1] Ciccacci C, Perricone C, et al. PLoS One. 2014.
- [2] Jadon D, Tillett W, et al. Rheumatology (Oxford). 2013.
- [3] Galvão V, Atta AM, et al. Joint Bone Spine. 2009.

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SAT0616 DIAGNOSTIC MODELING OF SYSTEMIC LUPUS ERYTHEMATOSUS BASED ON NEW AND TRADITIONAL AUTOANTIBODIES- DEMONSTRATION OF FEASIBILITY

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Background: Given the heterogeneity of clinical presentations, the diagnosis of Systemic Lupus Erythematosus (SLE) can be challenging, in particular in those patients presenting with early or incomplete disease, or with overlapping or atypical features. Autoantibodies (AABs) are important in aiding the clinical diagnosis of SLE, with some few AABs, anti-double-stranded DNA (dsDNA), anti-Smith (Sm), and anti-ribosomal P (RiboP) being highly associated with SLE. As none of the traditional AABs has sufficient sensitivity to achieve diagnosis of

SLE, current testing is based on measuring multiple AAB assays either in parallel or serial. We have recently identified novel AABs in SLE, which hold promise for improving diagnostic testing of SLE (1). We have developed quantitative ELISA-prototypes for five new AABs, which were tested in combination with traditional AABs.

Objectives: The objectives of this study were to evaluate the diagnostic value of novel AABs and to screen for an optimized combination of novel and traditional AABs using logistic regression to increase the diagnostic accuracy of SLE testing.

Methods: Serum samples were obtained from 156 SLE patients with European ancestry at the rheumatology department of the Heinrich-Heine University (Düsseldorf, Germany), and Hannover Medical School (Hannover, Germany). SLE samples were compared against 126 samples from autoimmune diseases (AID; myositis: n=20; Sjögren's syndrome (SjS): n=31; rheumatoid arthritis (RA) n=36; systemic sclerosis (SSc): n=39), and 77 healthy control samples. Prototype bead based ELISAs were developed for five novel antigens described in (1). Traditional diagnostic AABs were measured using IVD ELISAs and included: SSA/Ro60, SSA/Ro52, La/SSB, Sm, RNP, dsDNA, Scl70, CENPB, Jo-1, CCP, phospholipid and dsDNA. Optimized marker combinations of new and traditional markers were tested using logistic regression and receiver operating curve analysis (ROC).

Results: When comparing 156 SLE patients with 203 control samples, the area under the curve (AUC) of the five novel SLE ELISAs ranged from 0.63 to 0.75. A cut-off was set at a specificity of 95% and yielded a sensitivity ranging from 13.5% to 21.2% for the five novel assays. The sensitivity and specificity of new ELISAs was comparable to traditional ELISAs, which was in this cohort for anti-dsDNA 35% and 97%, anti-Sm 15% and 97%, and anti-RiboP 26% and 97%. A logistic regression model was used to combine the results of multiple tests. Compared to a logistic regression with traditional assays, a logistic regression with novel markers achieved higher sensitivity by pertaining high specificity. The logistic regression model based on a multimarker IVD assay with ten extracted nuclear antigens (ENA) yielded an AUC of 0.87 and a sensitivity of 58% at a specificity of 95%. By contrast, the optimal combination of traditional and novel ELISAs reached an AUC of 0.92 and a sensitivity of 75% at a specificity of 95%.

Conclusions: This study demonstrates the feasibility of combining test results of novel and traditional AABs using logistic regression to increase the diagnostic accuracy for SLE. Further studies are required to assess the impact of different ethnicities on marker selection and algorithm performance.

References:

- [1] Budde P et al. (2016). Lupus. (8):812–22.

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SAT0617 SALIVARY GLAND ULTRASOUND CAN DIFFERENTIATE PRIMARY SJÖGREN'S SYNDROME FROM SYSTEMIC DISEASES WITH SALIVARY GLAND INVOLVEMENT

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Background: Ultrasound of the major salivary glands (SGUS) has shown high diagnostic accuracy for primary Sjögren's syndrome (pSS) [1,2]. None of the existing studies, however, included a control group of patients with sarcoidosis, amyloidosis, human immunodeficiency virus (HIV) or hepatitis C (HCV) infection [2]. These diseases are known to frequently affect the major salivary glands, cause dry mouth or may have histopathology similar to pSS.

Objectives: To assess the diagnostic accuracy of SGUS to differentiate pSS from systemic diseases with salivary gland involvement.

Methods: Twenty consecutive patients fulfilling the AECG criteria for pSS and 20 consecutive patients with well-established systemic diseases, i.e., 5 patients with sarcoidosis, 5 patients with amyloidosis, 5 patients with HIV infection and 5 patients with HCV infection, were included. Echogenicity, parenchymal homogeneity, presence of hypoechogenic areas and clearness of posterior glandular border of both parotid and submandibular salivary glands were scored independently by two blinded observers and rated according to the Hocevar et al scoring system (ultrasound total score (UTS), range: 0–48) [3]. The optimal cut-off point was determined by using ROC curve. The agreement between SGUS positivity and classification as pSS was assessed.

Results: The median age of patients with pSS and systemic diseases was 50 and 53 years, respectively. Daily complaints of dry mouth longer than 3 months, need of liquid to swallow food and recurrent or persistent swelling of the major salivary glands were reported by 95%, 85% and 70% of patients with pSS compared to 45%, 30% and 20% of patients with systemic diseases. The accuracy of SGUS to discriminate pSS was excellent, with an area under ROC of 0.91 and optimal cut off point of 15. The agreement between SGUS and diagnosis for pSS was good ($\kappa=0.75$, percentage of absolute agreement 87.5) with a sensitivity of 90%,

specificity of 85%, positive predictive value of 86%, negative predictive value of 89% and diagnostic odds ratio of 51. UTS was positive in 2 patients with HIV infection and one patient with sarcoidosis. Patients with pSS had significantly higher UTS than patients with systemic diseases (median UTS 27 vs. 10, $p < 0.001$) and patients of the various subgroups ($p < 0.05$; Fig).

Figure: UTS in patient (sub)groups. ** $p < 0.001$, * $p < 0.05$. Black horizontal lines indicate median values. The intermittent horizontal line shows the cut-off point. The intermittent black vertical line separates the two major patient groups (pSS vs. systemic diseases) from the subgroups of patients with sarcoidosis, amyloidosis, HIV and HCV infection.

Conclusions: This pilot study indicates that SGUS has a high diagnostic accuracy to discriminate pSS from associated systemic diseases with salivary gland involvement. A minority of HIV and sarcoidosis patients, however, may show SGUS findings mimicking pSS.

References:

- [1] Jousse-Joulin et al. *Rheumatology (Oxford)* 2016;55:789–800.
- [2] Delli et al. *Oral Dis* 2015;21:792–800.
- [3] Hocevar et al. *Rheumatology (Oxford)* 2005;44:768–72.

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SAT0618 DISTURBANCES OF THE ACRAL PERFUSION DETECTED BY FLUORESCENCE OPTICAL IMAGING ARE ASSOCIATED WITH THE DEVELOPMENT OF ISCHEMIC COMPLICATIONS IN PATIENTS WITH SYSTEMIC SCLEROSIS

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Background: Systemic sclerosis (SSc) is a condition causing an impaired microcirculation with the risk of ischemic complications such as digital ulcers and pitting scars (DU/PS). Fluorescence optical imaging (FOI) is an imaging method that detects enhanced microcirculation as a sign of joint inflammation in both hands of patients with rheumatoid arthritis and other arthritides [1, 2]. FOI's impact to record disturbed microcirculation in the hands of patients with systemic sclerosis has not yet been sufficiently investigated.

Objectives: To find associations of disturbed microcirculation initially detected by FOI and the development of new DU/PS throughout a follow-up of 12 months.

Methods: Sixty-three patients with SSc were included and received FOI examination following the Xiralite-System guidelines (ICG 0.1mg/kg BW i.v.; 6 minute duration) as well as capillaroscopy at baseline. After a mean follow-up time of 12 months (min-max: 8–20 months), all participants were followed regarding the development of new ulcers and pitting scars.

Results: A disruption of microcirculation in FOI was defined as a lack of a sufficient fluorescent signal in at least one fingertip over the entire course of the examination and was found in 11 of 63 SSc patients. All patients had a history of DU/PS and frequently presented with a *late pattern* capillaroscopy (9 out of 11) at baseline. Fingers with a disrupted microcirculation also showed a reduced capillary density to a greater extent (96.0%) than fingers with a sufficient signal in FOI (76.0%; $p = 0.0241$).

30 of 60 patients developed digital ulcers or pitting scars during follow up (3 drop outs due to death ($n = 2$) or withdrawal). 81.8% of patients with a disturbed microcirculation in FOI developed these complications during follow-up compared with 42.9% of patients without a disruption in FOI ($p = 0.0419$; OR=6.0 [95% CI 1.2 - 30.7]). A disruption of microcirculation especially increased the risk of developing DU/PS in the same finger: 20.1% of fingers with normal, but 65.4% with a missing FOI signal in the fingertip presented with an ischemic complication during follow-up ($p < 0.0001$; OR=7.5 [95% CI 3.3 - 17.3]).

Conclusions: Fluorescence optical imaging can reveal an impaired microcirculation in patients with systemic sclerosis, which is associated with microangiopathic changes as seen in capillaroscopy as well as the subsequent development of digital ulcers and pitting scars. Therefore, FOI might help to identify patients at risk for these complications.

References:

- [1] Werner SG et al. Inflammation assessment in patients with arthritis using a novel in vivo fluorescence optical imaging technology. *Ann Rheum Dis.* 2012 Apr;71(4):504–10.
- [2] Werner SG et al. Indocyanine green-enhanced fluorescence optical imaging in patients with early and very early arthritis: a comparative study with magnetic resonance imaging. *Arthritis Rheum.* 2013 Dec;65(12):3036–44.

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SAT0619 ANKLE EVALUATION IN ACTIVE RHEUMATOID ARTHRITIS BY ULTRASOUND: A CROSS SECTIONAL STUDY

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Background: Ankle joint evaluation is underestimated in many clinical and sonographic scores used for evaluation and follow-up of rheumatoid arthritis (RA) patients. Moreover, sonographic scores which included the ankle joint had no agreement on examination parameters. More effort is needed to detect the value of the ankle joint examination in RA and also a description of the earliest and the most frequent ultrasonographic signs that should be considered in ankle assessment (1).

Objectives: detection of ankle affection by ultrasound (US) in active RA and correlate this finding with disease duration, DAS28-ESR 28 score and rheumatoid factor (RF).

Methods: 126 ankle joints and tendons of 63 active RA patients, aged above 18 years old were included in the study. US examination was done to the tibiotalar and talonavicular joints for synovitis and/or effusion on Greyscale (GS) mode and power doppler (PD). The anterior, lateral and posterior ankle tendons were examined for tenosynovitis and tendinosis.

Results: The mean age and $\pm$ standard deviation were 35.1 ± 8.3 with the female-to-male ratio 2:1. The mean disease duration was 22.7 ± 9.6 months. The mean DAS28-ESR 28 score was 3.05 ± 0.66 . The most frequent pathologies detected were tenosynovitis of the flexor, extensor or peroneal tendons (found in 30.2% of the affected ankles); followed by synovitis of the tibiotalar and talonavicular joints (18.3%); next was erosion (8.7%) and lastly tendinosis (4%). The earliest sonographic signs were tenosynovitis, followed by synovitis, erosion, and lastly tendinosis.

Conclusions: It can be stated that ankle evaluation should be considered more in RA assessment. The tibialis anterior and posterior tendons, the tibiotalar and talonavicular joints were the commonest and most frequent sites to be involved in the ankle. Tenosynovitis appears earlier than synovitis. DAS28-ESR score was correlated to synovitis and tenosynovitis but not to erosion. Bilaterality and erosion were correlated with disease duration. RF positivity has a positive correlation with positive US findings in the ankle region.

References:

- [1] Alsuwaidi M, Ehrenstein BP, Fleck M, Hartung W: Ankle Joint Involvement In Rheumatoid Arthritis. An Ultrasound Study Using High Resolution-Colour Doppler Ultrasound. In: *Arthritis and Rheumatism: 2013*: Wiley-Blackwell.

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SAT0620 POTENTIAL ROLE OF METACARPOPHALANGEAL JOINTS ULTRASOUND IN THE DIFFERENTIAL DIAGNOSIS BETWEEN EARLY RHEUMATOID ARTHRITIS AND EARLY SPONDYLOARTHRITIS

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Background: Several studies have demonstrated that musculoskeletal ultrasound (MSUS) is more sensitive in diagnosing arthritis when compared to clinical examination, although, as underlined in a recent review, still remains controversial whether it can improve substantial discriminatory value in an early arthritis (EA) setting. (1) In 2011 Gutierrez M. et al. published preliminary data on high frequency of peritendon extensor tendon inflammation in Psoriatic arthritis (PsA) patients, suggesting a relevant potential role for US in differential diagnosis between Rheumatoid Arthritis (RA) and PsA at metacarpophalangeal (MCP) joints level and recommending additional research in order to confirm these data. (2)

Objectives: To compare MSUS findings between early RA and early Spondyloarthritis (SpA) patients at MCP joints level.

Methods: From a consenting cohort of EA patients presenting to our Rheuma-