Gene expression profiles in primary Sjögren’s syndrome with and without systemic manifestations


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Background: Different phenotypes characterise the clinical spectrum of primary Sjögren’s syndrome (SS). Patients with a clinical expression limited to glandular manifestations (EGMs) are classically distinguished from patients with extra-glandular manifestations (EGMs). The former patients often complain higher level of fatigue and widespread pain (WP). (Segal et al. 2013) This suggests that gene expression pattern may be different in the two subgroups.

Results: The mean age of the patients was 38.9±15.2 years (range, 13–69 years). The study included 85 females (87.6%) and 12 males (12.4%). The clinical presentations were nphrotic syndrome, nephritic syndrome, and asymptomatic urinary abnormalities in 38 (39.2%), 20 (20.6%), 39 (40.2%) patients, respectively. Nine patients were classified Class II (9.3%, including 2 as ClassII +I), 82 as Class IV (84.5%), 10 as Class IV-segmental/IV-S (10.3%) and 72 as Class IV-global (IV-G) (74.2%), including 4 as Class IV-G+V) and 6 as Class V (6.2%). Forty-two(43%) patients presented with acute and 55 (57%) with features of chronic TMA. All patients had received treatment with standard immunosuppressants (55% mycophenolate, 39% cyclophosphamide, 6% other regimen) and steroids.

Abstract OP0201 – Table 1

<table>
<thead>
<tr>
<th>Acute TMA features</th>
<th>Chronic TMA features</th>
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<tr>
<td>Endothelial swelling with partial or complete occlusion of lumina;</td>
<td>Capillary wall thickening with double contours;</td>
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<tr>
<td>Microthrombi, focal or global;</td>
<td>Organizing capillary thrombi;</td>
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<tr>
<td>Fragmented RBC on glomerular subendothelial space and/or mesangial areas;</td>
<td>Glomerular ischaemic collapse with arteriolar occlusion;</td>
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<tr>
<td>Mesangiopathy, focal;</td>
<td>Segmental or global glomerulosclerosis;</td>
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<tr>
<td>Glomerular congestion with effluent arteriolar occlusion;</td>
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At 12 months, CR was observed in 37 patients (38.1%), PR in 22 (22.6%) and no response in 38 (39.1%). Sixty-one patients (62.9%) were antiphospholipid positive (aPL) and 37 (38.1%) received anticoagulation with vitamin-K antagonist (VKA) and/or heparins. Presence of aPLs(OR, 2.4; 95% Cl, 1.2–4.6; p=0.03), anti-DNA positivity (OR, 12.8; 95% CI: 3.0 to 71.3; p=0.002), and chronic features of TMA (OR, 3.0; 95% CI: 1.2 to 7.5; p=0.04) were all found to be associated with no response. When limiting the analysis to aPL positive patients, after adjusting for type of immunosuppressant therapy and LN class on biopsy, variables that were significantly associated with CR +PR were features of acute TMA rather than chronic (OR, 8.62; 95% Cl: 1.4 to 97.1; p=0.03) and the use of VKA/heparins(OR, 2.1; 95% Cl: 1.02–16.2; p=0.046).

Conclusions: In patients with concomitant LN and TMA, the presence of aPL and chronic features of TMA were associated with poorer renal outcomes. In patients with aPL, the use of anticoagulation appeared protective and warrant further investigation as a therapeutic tool, especially in the setting of acute TMA.

Disclosure of Interest: None declared


OBJECTIVES: To investigate the differences of gene expression in SjS patients with GFs and in those with EGMs.

Methods: Nineteen patients with SjS were selected for the study. Gene expression in peripheral blood mononuclear cells (PBMCs) was analysed in 4 patients with EGMs and 4 patients with GFs alone using Clarion D human Affymetrix gene chip (Affymetrix, Santa Clara, CA, USA), and compared to that found in healthy controls. Differences in gene expression were evaluated by analysis of variance (ANOVA) and Step-Up FDR-controlling procedure, being FDR corrected p-values<0.01 and fold change ≥2 considered as statistically significant.

Validation of the gene overexpression was performed by quantitative Real Time (qRT)-PCR in PBMCs from all the selected SjS patients, using the ΔΔCT method for comparing relative fold expression differences.

Results: All the enrolled SjS patients (18 females and 1 male) had a positive lip biopsy, while anti-SSA/Ro antibodies were detected in 10/11 and 6/8 of the patients with EGMs and with GFs alone, respectively. ESSDAI value ranged from 7 to 55 in patients with EGMs (median 17), and from 0 to 2 in patients with GFs alone (median 1).

In both types of patients, the functional analysis of the two transcriptomes showed a large number (>1000) of modulated genes that are involved in the well-known pathological processes of SjS, i.e., apoptosis, inflammatory response, immune response, type I and type II interferons, and Toll-like receptors signalling. Genes modulated only in patient with EGMs showed a significant enrichment of the biological processes associated with immune response (79% of all enriched processes) and, namely, of the molecular pathways related to B cell activation. The analysis of the transcripts expressed only in patients with GFs alone showed instead a preponderant enrichment in different metabolic processes (43%) and in processes associated with the central perception of the stimuli. Indeed, genes involved in sensory perception and in nociceptive signals (i.e., ANPEP, TNFR1, P2RY1, IFNG) were modulated exclusively in patients with GFs alone. The significant differential expression of selected genes in the two SjS subgroups was confirmed by the qR-PCR analysis.

Conclusions: These data indicate that in SjS patients with GFs alone a dysregulation of pain pathways (namely beta-adrenergic receptor and Notch signalling) may play a role in the development of WP that is common in this subset of patients. The biological mechanisms triggering the activation of these genes remain to be completely clarified.

Disclosure of Interest: None declared


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How monogenetic autoinflammatory diseases help to understand and treat rheumatic diseases

OP0203

SAFETY AND EFFICACY OF INTRAVENOUS ADMINISTRATION OF BONE-MARROW DERIVED MESENCHYMAL STEM CELLS IN THERAPY REFRACTORY JUVENILE IDIOPATHIC ARTHRITIS PATIENTS, A PHASE II/III PILOT-STUDY

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Objectives: To compare the total number of adverse events (AE) of allogeneic bone-marrow derived MSC infusion in refractory JIA and to evaluate its effectiveness.

Methods: Single-centre Phase Ib/IIa, open label intervention study in JIA patients previously failing all biologicals registered for their diagnosis. Six patients will receive 2 million/kg intravenous infusions of allogeneic bone-marrow derived MSC. In case of ACR-Ped30-response but subsequent loss of response one and maximal two repeated infusions are allowed. Results: Six JIA patients with 9.2 years median disease duration, still active arthritis and damage were included. All had failed methotrexate, corticosteroids and median 5 different biologicals. MSC were administered twice in 3 patients. No acute infusion reactions were observed and a lower post-treatment than pre-treatment incidence in AE was found. The one sJIA patient had again an evolving macrophage activation syndrome, 9 weeks after tocilizumab discontinuation and 7 weeks post-MSC infusion. Eight weeks after one MSC infusion, 4 patients showed less active joints, 5 patients improved in many clinical parameters and inflammatory parameters decreased in 3/4. After 1 year, we found significantly lower active joint counts.