distance of ≈ 1500 nucleotides, indicating that overlap with the promoter site is most likely. A common sequence motif around these CpG sites was not immediately detectable but requires more detailed analysis.

Conclusions Microarray based comparative analysis of transcriptional and epigenetic differences suggests a detailed picture of methylation associated gene regulation and enables to generate an epigenetic map of relevant CpG site for genes expressed and regulated in immune cell types. As many of the microarray based suspected CpG sites of a defined gene did not match with differential gene expression, epigenetic profiling with microarrays has to be interpreted carefully.

A7.6 COMPARATIVE TRANSCRIPTOME ANALYSIS OF HUMAN AND MOUSE SYNOVIAL FIBROBLAST RESPONSES TO TNF

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¹Thomas Häupl, ¹Till Sörensen, ¹Bijana Smiljanovic, ¹Marc Bonin, ²Andreas Grützkau, ³Christoforos Nikolaou, ³Ioannis Pandis, ³George Kollias, ⁴Anthony Rowe. ¹Dept. of Rheumatology and Clinical Immunology, Charité University Medicine, Berlin; ²German Arthritis Research Center, Berlin; ³Biomedical Sciences Research Centre 'Alexander Fleming', Institute of Immunology, Vri; ⁴Janssen Research and Development

Background and Objectives Animal models are essential for development of new therapeutics to test their effectiveness and to control for side effects. Targeted therapies with biologics reveal differences in responsiveness and suggest substantial differences in the molecular pathomechanisms of human as well as murine arthritis. In order to characterise such differences, we started a pilot investigation based on published transcriptomes to develop bioinformatic strategies of interspecies gene expression comparison.

Materials and Methods A GEO repository search for compatible data sets on arthritis in human and mouse revealed the Affymetrix based transcriptomes of synovial fibroblasts from rheumatoid arthritis (RA) and osteoarthritis (OA) patients (GEO-accession: GSE13837) and mouse BALBc wt (GSE17160) with similar stimulation (TNF-alpha) and time point after stimulation (12 hours). MAS5 algorithms of signal extraction and pairwise comparison were performed. The BioRetis database was used for group comparisons, candidate gene selection and data sharing. Differentially expressed genes were exported for interspecies comparison in a file-maker database. Affymetrix annotation files were used for mapping of orthologous genes.

Results The number of differentially expressed genes induced by TNF in synovial fibroblast from RA, OA and BALBc were 1584, 1476 and 503 respectively. Comparison of genes overexpressed in RA and OA revealed up to 70% (1034 genes) overlap. Murine and human response was similar for 197 genes in all three groups, 45 only between BALBc and RA and 24 only between BALBc and OA. Ranking the genes by dominance in change call and/or fold change, 7 genes were overlapping between the top 10 of both human cells, OA and RA, and 13 out of the top 20. All overlapping genes in human revealed a high correlation ($R = 0.795$). In contrast, overlap between RA and BALBc SFbl responses revealed only 4 genes out of the top 10 and 5 out of the top 20 in each and a weaker correlation (0.392). Pathway tools including IPA may identify TNF as one potential upstream regulator. However, the reference list of expected gene candidates for scoring in IPA seemed to be about 2.5 fold longer in human and to include less than 60% of the murine reference candidates.

Conclusions Interspecies comparison in this defined experiment revealed first insight in common responses but also demonstrates that differences can be observed even in most dominant candidate genes. Therefore, systematic investigations are needed to provide a reliable basis for comparative mapping of the molecular pathomechanisms in mouse models and human disease.

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A7.7 DIFFERENT GENETIC BACKGROUND OF DERMATOMYOSITIS AND POLYMYOSITIS IN A SINGLE CENTRE COHORT

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¹M Remakova, ¹M Skoda, ¹T Svitalkova, ¹M Faustova, ¹L Plestilova, ²Z Betteridge, ¹H Mann, ¹J Vencovsky, ¹O Krystukova, ¹P Novota. ¹Institute of Rheumatology, Prague, Czech Republic; ²Royal National Hospital for Rheumatic Diseases, Bath, UK

Background and Objectives The idiopathic inflammatory myopathies (IIM) are systemic connective tissue diseases in which autoimmune pathology is responsible for promotion of chronic muscle inflammation and weakness. As in many other autoimmune diseases, the development of IIM is also associated with genes of HLA complex. The aim of this study was to determine the basic relation between alleles of HLA genes and IIM.

Materials and Methods We have performed low to high resolution genotyping to characterise the allelic profiles of HLA-DRB1, -DQB1 and -DQA1 loci in a large group of single centre cohort of patients suffering from IIM ($n = 269$). The genomic DNA was prepared by standard DNA extraction methods and the HLA typing was done using the commercial LABType® SSO kit (One Lambda, USA). Statistical evaluation of results was done with chi-2 test and Fisher exact test. Autoantibody profiles were analysed with radioactive immunoprecipitation.

Results The frequencies of HLA-DRB1*03:01 and -DRB1*16:01 alleles were increased in IIM patients and the difference reached statistical significance when compared to healthy controls ($P < 0.01$ for DRB1*03:01; $P < 0.05$ for DRB1*16:01). Different alleles were associated with dermatomyositis (DM) or polymyositis (PM). The frequency of DRB1*16:01 was significantly higher in DM patients ($P < 0.01$), whereas the frequency of DRB1*03:01 was higher in patients suffering from PM ($P < 0.01$), when compared to controls.

Presence of HLA-DRB1*03:01 allele was associated with anti-Jo-1, anti-Ro52, or anti-Pm-Scl positivity in all IIM patients ($P < 0.05$). Interestingly, the DRB1*03:01 allele was also associated with the negativity for anti-p155/140 in our patients ($P < 0.01$).

The DRB1*16:01 allele was associated with negativity of all studied autoantibodies, particularly in subgroup of DM patients ($P < 0.05$).

Conclusions This study identifies different genetic background between patients with dermatomyositis and polymyositis in a homogenous population of patients from a single centre.

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A7.8 DNA DEMETHYLATION IN SALIVARY GLAND EPITHELIAL CELLS FROM PATIENTS WITH PRIMARY SJÖGREN'S SYNDROME MAY BE AScribed TO INFILTRATING B CELLS

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^{1,2}Yves Renaudineau, ^{1,3}Yosra Thabet, ¹Christelle Le Dantec, ³Ibtissem Ghedira, ^{1,4}Valérie Devauchelle, ^{1,4}Divi Cornec, ¹Jacques-Olivier Pers. ¹Research Unit EA2216 "Immunology, Pathology and Immunotherapy", European University of Brittany, Brest University Medical School Hospital, Brest, France; ²Laboratory of Immunology and Immunotherapy, CHRU Morvan, Brest, France; ³Research Unit O3UR/07-02 "Autoimmunity and Allergy", Faculty of Pharmacy, Monastir, Tunisia; ⁴Unit of Rheumatology, CHRU Cavale Blanche, Brest, France

Background and Objectives Sjögren's syndrome (SS) is an autoimmune exocrinopathy characterised by an epithelium injury surrounded by dense lymphocytic infiltrates composed of activated T and B cells. Present at the interface of genetic and environmental risk factors, epigenetic modifications are suspected to play a key role in SS. Accordingly, we decided to further characterise DNA methylation in SS.