scaffold protein involved in the proteasomal processing of p105 into the p50 subunit. Inhibition of this process impairs the nuclear translocation of NF- κ B and the consequent p65/p50 regulated gene transcription. Hence, TRUSS may be a novel target for modulating the inflammatory functions of TNF α -TNF-R1 signalling.

S31 CHARACTERISATION OF CELL ADHESION MOLECULE-1 IN LUNG MAST CELLS

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Introduction and Objectives Cell adhesion molecule 1 (CADM1) is implicated in several diseases and a prognostic factor for lung cancer. In human lung mast cells (HLMC) it contributes to cell-cell adhesion and proliferation. Functional CADM1 isoforms arise from alternative splicing between exons 7 and 11. Exon 9 may be functionally important because it encodes a specific cleavage site for TNF- α -cutting enzyme, TACE, in the longest SP1 and SP6 isoforms. Our aim was to characterise the isoforms of CADM1 expressed in HLMC.

Methods CADM1 expression in isolated HLMC and human mast cell lines (HMC-1 and LAD2), was investigated using RT-PCR, cloning, and transfection.

Results Multiple highly glycosylated CADM1 isoforms were found in HLMCs and cell lines. The SP4 isoform containing exon 8 represented ~80% of clones in both HLMCs and 'differentiated' LAD2 cells, and ~96% in non-differentiated HMC-1 cells. The SP1 isoform with exons 8+9 represented $\leq$ 20% of clones in HLMCs and LAD2 cells. A novel SP6 isoform with exons 8+9+10 was found only in HLMCs (<5% of clones). More sensitive PCR analysis detected all these isoforms and an additional isoform SP3, lacking exons 8–10, in HLMCs and in cell lines. In contrast to these functional isoforms, non-functional isoforms were also found with two cryptic exons between exons 1 and 2, which cause a translational frame shift and premature termination of the protein. The cryptic exon B is located within the hot spot for SNPs. When the isoforms SP4, SP1 and SP6 were expressed in the HMC-1 cell line and the epithelial cell line HEK293 as fusion CADM1-GFP proteins, they were found at the cell periphery as well as in the cytoplasm. They were concentrated in cell boundaries in clumps of HMC1 cells.

Conclusions Multiple and novel CADM1 isoforms are found in HLMCs. The longest SP1 and SP6 isoforms might be involved in the negative regulation of HLMC adhesion following secretion of TACE after activation. Some cryptic exons are likely to be a result of specific SNPs, creating new branching points for splicing. Non-functional cryptic CADM1 isoforms may reduce mature protein expression and affect the function of CADM1 expressing cells.

S32 CYCLICAL MECHANICAL STRETCH ENHANCES THE PRO-FIBROTIC RESPONSES OF PRIMARY EMBRYONIC FOETAL FIBROBLASTS, BUT NOT ADAM33 EXPRESSION

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Rationale A Disintegrin And Metalloprotease (ADAM)33 is a susceptibility gene associated with asthma, bronchial hyperresponsiveness (BHR) and reduced lung function in young children. It is selectively

expressed in mesenchymal cells, including bronchial smooth muscle and fibroblasts. We have previously shown that ADAM33 expression increases during lung development when spontaneous peristaltic contractions of the airways commence. Therefore we hypothesised that mechanical strain induces ADAM33 expression and affects smooth muscle/myofibroblast differentiation.

Methods Primary human embryonic fibroblasts from the pseudo-glandular stage of lung development were cultured on flexible collagen-coated membranes and exposed to cyclical mechanical stretch (30% amplitude, 12 cycles per minute) for 48, 96 and 168 h. Control cells were cultured on the same membranes without mechanical stretch. Quantitative RT-PCR was performed for ADAM33 and α -smooth muscle actin (α -SMA), a marker of smooth muscle and myofibroblast differentiation. We also measured collagen III and IL-8 mRNA. Soluble collagen protein levels in culture supernatants were measured using soluble collagen assay.

Results ADAM33 and α -SMA mRNA expression were not significantly affected by mechanical strain. In contrast, collagen III mRNA expression was increased fourfold by cyclical mechanical strain and there was a threefold increase in soluble collagen proteins in culture supernatants of stretched cells. Unexpectedly, IL-8 expression was also increased by cyclical mechanical strain.

Conclusion Mechanical strain did not appear to influence markers of smooth muscle differentiation. However, the increase in ECM production may indicate a requirement for stiffening of the airways as the tubular structures develop. We postulate that IL-8 is not proinflammatory in the context of airway development, and may be a paracrine growth factor for developing epithelial cells.

S33 SUPPRESSION OF CONSTITUTIVE AND STIMULATED SECRETION OF HISTAMINE FROM HUMAN LUNG MAST CELLS BY A SECRETED FACTOR FROM LUNG EPITHELIAL CELLS

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Introduction Constitutive and IgE-dependent secretion of histamine from human lung mast cells (HLMC) is suppressed by direct contact with the BEAS-2B bronchial epithelial cell line, but not conditioned epithelial cell media. This suggests that direct contact or a low concentration/labile secreted factor is involved. We explored this relationship further using a Transwell co-culture system of HLMC with BEAS-2B monolayers or air liquid interface primary bronchial epithelial cultures (ALI) derived from asthmatic and healthy subjects.

Methods ALI and BEAS-2B were grown on Transwell membranes. IgE-sensitised HLMC were then cultured i) on the luminal surface of a BEAS-2B monolayer, ii) separated from the BEAS-2B basal surface by a Transwell membrane, or iii) on the well bottom with BEAS-2B and ALI sitting on the Transwell insert. Cells were co-cultured for 16 h, media removed, and HLMC stimulated with anti-IgE for 30 min. Parallel controls without epithelial cells were performed. Histamine concentrations were determined by radio-enzymic assay.

Results BEAS-2B ($n=3$): Compared to no epithelium control, constitutive histamine secretion from HLMC in direct contact with BEAS-2B was suppressed by a mean ($\pm$ SEM) $57 \pm 15\%$ ($p=0.04$), and by $55 \pm 9\%$ when separated by Transwells ($p=0.02$). IgE-dependent secretion was suppressed by $79 \pm 8\%$ ($p=0.04$) from HLMC in direct contact with BEAS-2B and by $88 \pm 7\%$ ($p=0.03$) when separated by Transwells, compared to control. ALI-culture