

Agreement was moderate for the presence of abscess ( $k=0.46$  95% CI 0.14–1.00), acute mucosal inflammatory changes ( $k=0.55$ , 95% CI 0.19–0.90) and fibrotic changes ( $k=0.50$ , 95% CI 0.04–0.95). There was no significant difference in the mean estimated stricture length between MRE and SICUS.

**Conclusion** SICUS compares favourably with MRE in the diagnosis of complications in patients with Crohn's disease. This imaging technique is particularly useful in patients with stricturing (Montreal B2) disease. SICUS is a useful alternative diagnostic technique to MRE, particularly when access to MR may be limited or is poorly tolerated by the patient.

#### REFERENCE

- 1 Chatu S *et al.* Diagnostic accuracy of small intestine ultrasonography using an oral contrast agent in Crohn's disease: comparative study from the UK. *Clin Radiol.* 2012 Jun;67(6):553-9

**Disclosure of Interest** None Declared.

#### PTU-074 ULCERATIVE COLITIS: THE ALPHA-E-BETA-7 INTEGRIN IS ASSOCIATED WITH A HIGH FREQUENCY OF TH17, TH1 AND TH17/TH1 CD4 LYMPHOCYTES

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**Introduction** T lymphocytes expressing the  $\alpha\text{E}\beta 7$  integrin are highly enriched within human intestinal epithelium and lamina propria. Studies exploring pathogenic or protective functions of  $\alpha\text{E}\beta 7$  expressing cells are lacking. Defining this phenotype is critical for our understanding of IBD pathogenesis and of translational importance with the development of etrolizumab, a humanised antibody specific to the  $\beta 7$  integrin that blocks  $\alpha 4\beta 7$ : MAdCAM-1 and  $\alpha\text{E}\beta 7$ :E-cadherin interactions.

**Methods** Lymphocytes within colonic biopsies from a total of 43 UC and 35 non-disease control patients were studied. Multi-colour FACS was optimised to determine surface and intracellular protein expression (CD45, CD3, CD4, CD8,  $\alpha\text{E}$ ,  $\beta 7$ , CD161, IL-17A, TNF $\alpha$ , IFN $\gamma$  and IL-10). qPCR was performed on TCR $\alpha\beta$ + lymphocytes, FACS sorted into CD4<sup>+</sup> $\alpha\text{E}\beta 7$ +, CD4<sup>+</sup> $\alpha\text{E}\beta 7$ -, CD8<sup>+</sup> $\alpha\text{E}\beta 7$ + and CD8<sup>+</sup> $\alpha\text{E}\beta 7$ - prior to gene expression assay. Dual stain IHC for  $\alpha\text{E}$ , plus CD3, CD4, CD8 and FOXP3 was performed using a Ventana Benchmark XT autostainer. Severity of UC was stratified using the Mayo endoscopic score for ulcerative colitis.

**Results** Ulcerative colitis was associated with a significantly increased frequency of T lymphocytes in the intestinal mucosa ( $p < 0.05$ ). IHC revealed the highest expression of  $\alpha\text{E}$  on CD4 and CD8 intraepithelial lymphocytes, although a substantial number of lamina propria lymphocytes also expressed this integrin. In UC, FACS demonstrated CD4<sup>+</sup> $\alpha\text{E}\beta 7$ + lymphocytes had a higher potential to produce the pro-inflammatory cytokines IFN $\gamma$  ( $p < 0.01$ ), TNF $\alpha$  ( $p < 0.001$ ) and IL-17A ( $p < 0.0001$ ) than CD4<sup>+</sup> $\alpha\text{E}\beta 7$ - lymphocytes. In addition, a mean of 31.5% of the CD4<sup>+</sup> $\alpha\text{E}\beta 7$ + lymphocytes produced both IL-17A and IFN $\gamma$  compared to a mean of only 7.7% in the CD4<sup>+</sup> $\alpha\text{E}\beta 7$ - compartment ( $p < 0.001$ ). IL-10 was not differentially expressed between CD4<sup>+</sup> $\alpha\text{E}\beta 7$ + and CD4<sup>+</sup> $\alpha\text{E}\beta 7$ - lymphocytes in controls or UC, and a low frequency of  $\alpha\text{E}\beta 7$ +FOXP3+ cells was observed by IHC. qPCR array confirmed higher mRNA levels of

IFN $\gamma$  ( $p < 0.001$ ), TNF $\alpha$  ( $p < 0.01$ ) and IL-17A ( $p < 0.01$ ), and lower transcription of FOXP3 ( $p < 0.0001$ ) in CD4<sup>+</sup> $\alpha\text{E}\beta 7$ + cells compared to CD4<sup>+</sup> $\alpha\text{E}\beta 7$ - cells.

**Conclusion**  $\alpha\text{E}\beta 7$  expression was associated with an enrichment of pro-inflammatory Th17, Th1 and Th17/Th1 T lymphocytes, and not associated with a regulatory phenotype. These data suggest therapeutic interventions targeting  $\alpha\text{E}$  expressing T cells and the  $\alpha\text{E}\beta 7$  integrin itself may be viable approaches for reducing aberrant inflammatory responses in UC.

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#### PTU-075 SYSTEMATIC REVIEW AND META-ANALYSIS: SENSITIVITY AND SPECIFICITY OF Tc-99m HMPAO LABELLED WHITE CELL SCINTIGRAPHY IN THE DIAGNOSIS OF ACTIVE INFLAMMATORY BOWEL DISEASE

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**Introduction** Technetium-99m hexamethylpropylene amine oxime (Tc-99m HMPAO) labelled white cell scintigraphy (WCS) is frequently used in the assessment of patients suspected of having active inflammatory bowel disease (IBD). However, no previous systematic review and meta-analysis has assessed the sensitivity, specificity, and positive and negative predictive values of this investigation in comparison with colonoscopy and histology. We therefore aim to present these data here.

**Methods** The MEDLINE and EMBASE databases were searched to January 2014. Prospective and retrospective cross-sectional studies recruiting adults suspected of a new diagnosis or flare of IBD, and comparing Tc-99m HMPAO labelled WCS with colonoscopy and histology, were eligible. True positive, false positive, true negative and false negative findings were pooled. A random effects model was used to obtain overall data for sensitivity, specificity, and positive and negative predictive values with a 95% confidence interval (CI).

**Results** We identified 15 eligible studies reporting data from 635 patients (174 Crohn's disease, 164 ulcerative colitis, 136 non-IBD). In total 1300 bowel segments were examined with 698 true positives, 41 false positives, 461 true negatives and 100 false negatives. Sensitivity was 0.90 (95% CI 0.85 to 0.95), specificity was 0.91 (95% CI 0.87 to 0.94), positive predictive value was 0.95 (95% CI 0.92 to 0.97) and negative predictive value was 0.83 (0.76 to 0.89).

**Conclusion** Tc-99m HMPAO labelled WCS is a sensitive and specific test for the diagnosis of active inflammatory bowel disease. Physicians may therefore find this a useful test for those in whom colonoscopy and histology are impractical or contraindicated.

**Disclosure of Interest** None Declared.