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

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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

^{1,2}CA Lamb*, ^{1,2}JC Mansfield, ³GW Tew, ⁴D Gibbons, ²AK Long, ⁴PM Irving, ³L Deihl, ³J Eastham Anderson, ⁵G O'Boyle, ¹DE Jones, ⁴A Hayday, ³M Keir, ³JG Egen, ¹JA Kirby. ¹Newcastle University, UK; ²Newcastle Upon Tyne Hospitals NHS Foundation Trust, Newcastle Upon Tyne, UK; ³Genentech, South San Francisco, USA; ⁴King's College London, London; ⁵University of Sunderland, Sunderland, UK

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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, αE , $\beta 7$, CD161, IL-17A, TNF α , IFN γ and IL-10). qPCR was performed on TCR $\alpha\beta$ + lymphocytes, FACS sorted into CD4⁺ $\alpha\text{E}\beta 7$ +, CD4⁺ $\alpha\text{E}\beta 7$ -, CD8⁺ $\alpha\text{E}\beta 7$ + and CD8⁺ $\alpha\text{E}\beta 7$ - prior to gene expression assay. Dual stain IHC for αE , 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 αE on CD4 and CD8 intraepithelial lymphocytes, although a substantial number of lamina propria lymphocytes also expressed this integrin. In UC, FACS demonstrated CD4⁺ $\alpha\text{E}\beta 7$ + lymphocytes had a higher potential to produce the pro-inflammatory cytokines IFN γ ($p < 0.01$), TNF α ($p < 0.001$) and IL-17A ($p < 0.0001$) than CD4⁺ $\alpha\text{E}\beta 7$ - lymphocytes. In addition, a mean of 31.5% of the CD4⁺ $\alpha\text{E}\beta 7$ + lymphocytes produced both IL-17A and IFN γ compared to a mean of only 7.7% in the CD4⁺ $\alpha\text{E}\beta 7$ - compartment ($p < 0.001$). IL-10 was not differentially expressed between CD4⁺ $\alpha\text{E}\beta 7$ + and CD4⁺ $\alpha\text{E}\beta 7$ - 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 γ ($p < 0.001$), TNF α ($p < 0.01$) and IL-17A ($p < 0.01$), and lower transcription of FOXP3 ($p < 0.0001$) in CD4⁺ $\alpha\text{E}\beta 7$ + cells compared to CD4⁺ $\alpha\text{E}\beta 7$ - 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 αE 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

¹CJM Williams*, ²AC Ford, ¹A Poullis. ¹Department of Gastroenterology and Hepatology, St George's Hospital and Medical School, London, UK; ²Leeds Gastroenterology Institute, Leeds General Infirmary, Leeds, UK

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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.