

change was seen in eosinophil count or in inflammatory markers post treatment. In the sero-positive group 23% had past and current eosinophilia, but this was not statistically different from sero-negative patients. 9/13 reportable charcoal stool cultures were negative. No patients with *Strongyloides* were taking steroids, compared to 23% of sero-negative patients. In both groups, >40% were on two or more immunosuppressants.

Conclusion There is a high sero-prevalence of *Strongyloides* in migrant IBD patients. Patients from Asia demonstrated the highest prevalence. Eosinophilia and raised inflammatory markers were not predictive of positive serology, most likely due to the high rate of immunosuppression. We cannot confirm all sero-positive patients were infected; published data² supports the specificity of *Strongyloides* serology for current infection. We recommend ECCO guidelines and current British Society of Gastroenterology guidelines are adapted to include targeting IBD patients who originate from endemic areas and serological testing be first line. Follow-up of patients is required to assess the impact of treatment on IBD activity.

Competing interests None declared.

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PWE-236 WHITE CELL APHAERESIS (WCA) WITH ADACOLUMN IS EFFECTIVE IN SELECTED CASES OF CHRONIC REFRACTORY COLITIS WITH HIGH HISTOLOGICAL ACTIVITY

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Introduction Treatment options for patients with chronic refractory colitis are limited. White cell aphaeresis (WCA) is effective in inducing clinical remission in chronic refractory colitis in patients with a strong inflammatory burden at baseline and histologically active disease. Previous multinational sham controlled trials have demonstrated significant improvement when patients with high histological activity (modified Rileys score) are selected for treatment.

Methods A prospective study was conducted in 30 patients with severe steroid -dependent or steroid -refractory ulcerative colitis referred for WCA. Inclusion criteria were (i) High disease activity score (partial Mayo score ≥ 6) (ii) Intractable symptoms despite treatment with steroids and/or immunosuppressants (iii) Severe disease at endoscopy and histologically. The aim was to induce clinical and IBD-Q remission at 12 weeks. A Mayo score ≤ 3 defined clinical remission. The 32 item Inflammatory Bowel Disease questionnaire (IBD-Q) was used to assess quality of life prior to treatment and at 12 weeks.

Results Patient Characteristics: Prior to treatment 28 patients (93.3%) were prescribed 5-ASA compounds. 12 patients (40%) were prescribed topical therapies (5-ASA enemas or suppositories/steroids enemas). 27 patients (90%) were steroid dependent (Prednisolone mean dose 21.1 mg, median 20 mg). Three patients (10%) were steroid refractory (no response to high dose oral steroids). 13 patients (43.3%) were prescribed Azathioprine of the remainder all had documented intolerance or a contraindication. One patient (3.3%) was prescribed six Mercaptopurine. Five patients had failed Infliximab (16.6%) and in one patient (3.3%) it was contraindicated. 1 patient (3.3%) had failed intramuscular Methotrexate. **Outcomes:** At week 12 clinical remission (Mayo score ≤ 3) was achieved in 22 patients (73.3%), 18 patients (60%) were no longer prescribed oral steroids. IBD-Q remission at week 12 was achieved in 19 patients (63.3%). Of the remainder, five patients (16.6%) achieved an IBD-Q response. Of eight patients (26.6%) who failed to achieve clinical remission at 12 weeks, one

achieved delayed remission at 20 weeks. Of the remaining seven treatment failures, five underwent colectomy (16.6%).

Conclusion WCA can be effective in inducing clinical remission and improving quality of life (IBD-Q) indices in chronic severe steroid refractory ulcerative colitis with highly active disease histologically. This data series suggests WCA should be considered before colectomy in this challenging patient group.

Competing interests None declared.

PWE-237 CICLOSPORIN IN ACUTE SEVERE ULCERATIVE COLITIS: A META-ANALYSIS

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Introduction The efficacy of intravenous Ciclosporin in the treatment of acute severe steroid refractory ulcerative colitis is variable in several small studies.^{1,2} The benefit of immunosuppression in initial responders for preventing delayed colectomy is unclear and large randomised trials are not forthcoming. The primary aim of the meta-analysis was to evaluate the efficacy of intravenous Ciclosporin in steroid refractory acute colitis. Primary end points were immediate and delayed colectomy rates. Secondary end points were delayed colectomy rates in responders on immunosuppression vs no immunosuppression.

Methods MEDLINE (1966–2011), EMBASE and PubMed databases searched using keywords, “colitis” “acute colitis,” “ciclosporin,” “cyclosporin,” “ulcerative colitis” and “fulminant colitis” from January 1965 to December 2011. Articles were selected/reviewed based on a pre-defined inclusion criteria and independently reviewed by three authors (RK, NG and TT) and data collected. Meta-analysis using random effects model was done.

Results 31 studies involving 1295 patients were included in the final analysis (692 males, average age 37.4 years, range 27–58.8 years). The average duration of colitis prior to admission (25 studies, 1062 patients) was 54 months and 660 patients (26 studies, 1148 patients) had pancolitis. The immediate colectomy rate was 23.6%, (95% CI 20% to >27.5%) and delayed colectomy 37.8% (95% CI 32.8% to 43.1%) at an average follow-up of 28.8 months. On meta-regression analysis the duration of colitis was the only confounding factor for delayed colectomy ($p=0.05$). On sub-group analysis, immunosuppression (Aza/6MP) in initial responders significantly reduced delayed colectomy rates by 20% (OR 0.36, 95% CI 0.61 to 0.81, $p=0.014$). The number need to treat to prevent one delayed colectomy was 5. There was a 13% increase in colectomy rates in the sub-group of patients (5 studies, 113 patients) who were on immunosuppression (failed immunosuppression) prior to iv ciclosporin although this was not statistically significant ($p=0.22$).

Conclusion Intravenous Ciclosporin prevents immediate and delayed colectomy in 76.4% and 62% respectively. There is a non-significant increase in colectomy rates (13%) in patients who failed immunosuppression prior to Ciclosporin. Immunosuppression in initial responders significantly reduces delayed colectomy rates and should be considered unless contraindicated.

Competing interests None declared.

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