

P1005: Dose escalated Infliximab in Inflammatory Bowel Disease - Crohn's Colitis Cure (CCC) Data Insights Program

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BACKGROUND

- Infliximab (IFX) is a monoclonal antibody targeting TNF- α
- Some people with IBD on standard dosing of IFX therapy have persistent disease activity, necessitating dose escalation (DE).
- Globally, pharmaceutical funding schemes variably fund escalated doses of drug, making access to adequate dosing insecure.
- Aim: to evaluate the extent of IFX DE and patient outcomes in a real-world cohort.

METHODS

- Crohn's Colitis Care is a cloud-based IBD-specific electronic medical record (EMR) used at IBD centres across Australasia since 2018.
- Data from CCCare flow across to a de-identified clinical quality registry, which was interrogated in November 2023.
- DE for IFX was defined as dosing interval <8 weeks and/or dosing >5mg/kg.

RESULTS

- Across 14 private and public IBD care centres, 1725 people were receiving IFX, of whom 44.2% were on DE therapy.
- Of the entire IFX cohort, 65% had Crohn's Disease, 33% had ulcerative colitis, and 2% were IBD-unclassified.
- Even gender distribution (52.5% male)

In the DE cohort (n=762):

- Median age: 37 years (IQR 27-49)
- Median age at diagnosis: 23 years (IQR 16-32)
- Median disease duration: 11.6 years (IQR 5.8-17.3).
- DE and standard dose cohorts did not differ by IBD diagnosis, gender, disease duration, age or age at diagnosis.

RESULTS (CONT.)

- Median drug level was higher 12-months post DE (8.16 μ g/ml vs 3.65 μ g/ml).
- Faecal calprotectin (FCP) remission rate (FCP <250 μ g/g) was higher 12 months post DE, as was endoscopic remission rate and patient reported outcome remission (Table 1).
- The rate of systemic corticosteroid use was lower 12 months post-DE than pre-DE.
- In the DE-IFX cohort, measures of healthcare utilisation were lower 12-months post-DE; hospitalisation (56 vs 146; p<0.001), surgical interventions (41 vs 61; p=0.040), endoscopic procedures (210 vs 397, p<0.001) and radiological investigations (243 vs 343; p<0.001).

| Outcome Measure (%) | Pre-DE | 12 months post-DE | p-value |
|--|--------|-------------------|---------|
| Faecal Calprotectin Remission (<250ug/g) | 50.8 | 69.6 | <0.001 |
| Endoscopic Remission | 23 | 39.2 | 0.047 |
| Radiologic Remission | 73.2 | 69.5 | 0.512 |
| PRO2 Remission | 70.5 | 78.1 | 0.015 |
| Systemic Steroid Therapy | 5 | 2.1 | 0.002 |

CONCLUSIONS

These prospectively collected data from large Australian and New Zealand IBD treatment centres show:

- A considerable proportion of people receiving IFX required DE, which appeared to be effective.
- DE IFX was associated with improved measures of remission and a reduction in several measures of healthcare utilisation.
- Further analysis can be performed to determine price points at which DE is cost-effective; additional data on quality of life and indirect healthcare costs would allow for a robust, holistic assessment of value in care.

