

Randomized open-label.

Aim: Safety and efficacy of IFX in paediatric CD.

Induction (w0,2,6) 5mg/kg IFX then patients responding at w10 were randomized to IFX q8 or q12 through w46.

#### Outcomes:

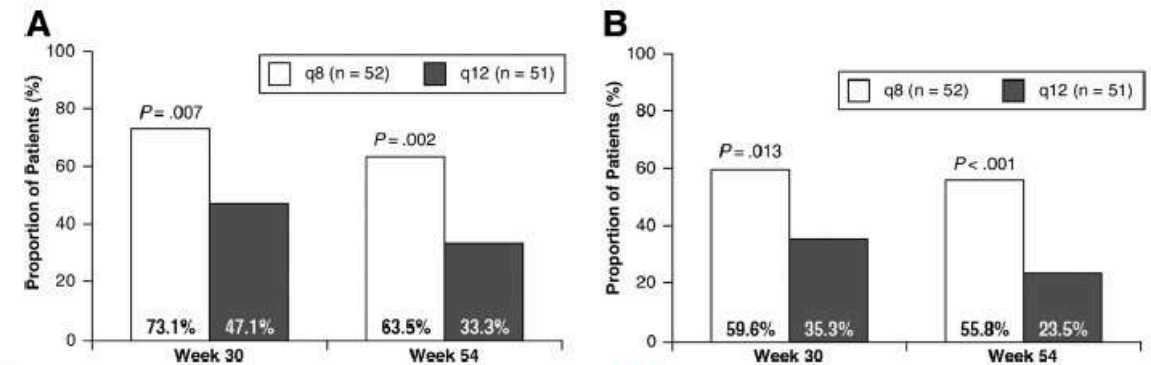
- Efficacy of induction regimen with IFX
- Clinical response and remission at w54

#### Results:

- Overall response to IFX w10: 88.4% clinical response; 58.9% clinical remission
- W54 clinical response: 63.5% IFXq8 vs 33.3% IFXq12 :  $p=0.002$
- W54 clinical remission: 55.8% IFXq8 vs 23.5% IFX q12;  $p<0.001$

#### Conclusion:

Paediatric patients responding to induction with IFX are more likely to be in clinical response and remission at w54 when therapy was given every 8 weeks rather than 12.



**Figure 2.** Clinical response (decrease from baseline to week 10 in the total PCDAI score of at least 15 points and a total PCDAI score of no more than 30 points (A) and clinical remission (PCDAI score of 10 points or lower; (B) for week 10 responders at weeks 30 and 54. All randomized patients.

