

Randomized. Open-label.
 CD patients controlled with IFX were randomized to switch to ADA.
 Dose optimisation allowed, loss of response/intolerance could cross over to the alternative group.

Outcomes: proportion of patients in the ADA group preferring ADA over IFX and the proportion of patients who needed rescue therapy with steroids, intensified antiTNF or need to stop treatment.

Results:

- Dose optimization or interruption 47%ADA vs 16% IFX; $p=0.006$
- Interruption due to loss of response: 2%IFX vs 28% ADA; $p=0.003$
- Injection site reactions more frequent than infusion reactions; $p=0.01$
- AntiTNF levels were stable during the 1 year period

Conclusion:

Elective switching from IFX to ADA associated with loss of tolerance and loss of efficacy within 1 year. Adherence to the first antiTNF is recommended.

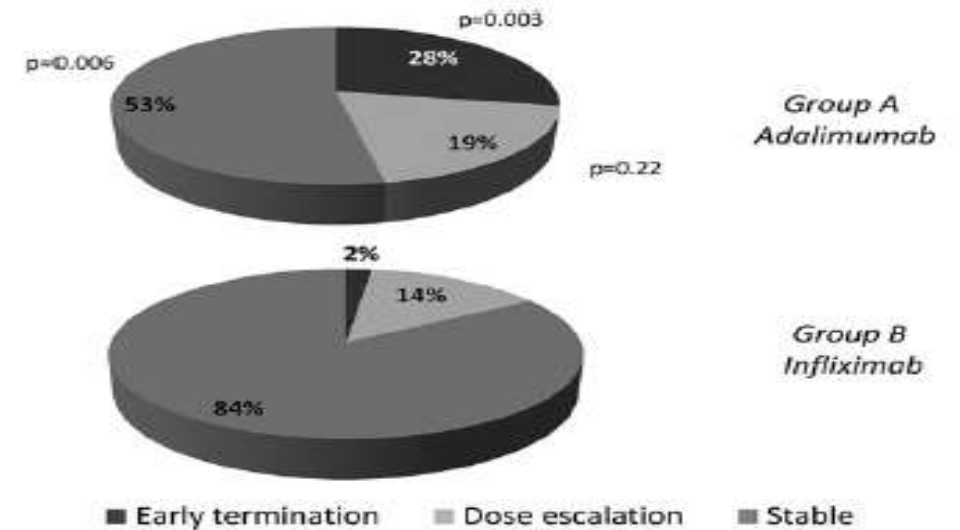


Figure 3 Graphical overview of the need for dose adjustments and the proportion of patients discontinuing the assigned treatment throughout the trial.