Multicenter randomised clinical trial.

Patients with CD after surgery were randomised to:
oral probiotic LA1 (Lactobacillus johnsonii 10¹⁰ CFU) or placebo.

Primary endpoints: Endoscopic recurrence at 12 months.

Results: N=70

- At 3 months endoscopic recurrence, LA1 vs placebo, p=0.48
- Severe endoscopic recurrence (i3/i4) 21% in LA1 vs 15% in placebo, p=0.33
- No differences in clinical recurrence

Conclusion:

Oral administration of the probiotic LA1 in patients with CD failed to prevent early endoscopic recurrence at 12 weeks after ileocaecal resection.

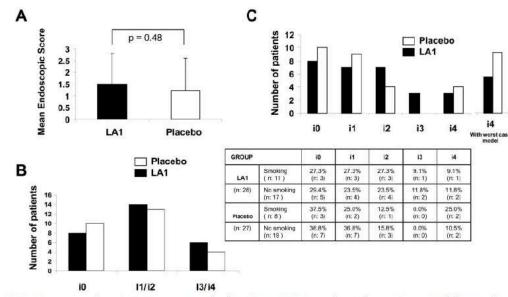


FIGURE 2. A: Mean endoscopic (Rutgeerts) scores in both treatment groups (ITT analysis without worst-case model). B: Stratification of patients in each treatment group according to endoscopic recurrence severity (ITT analysis without worst-case model). C: Stratification of patients in each treatment group per endoscopic score (ITT analysis with or without worst-case model) and according to smoking status (ITT analysis without worst-case model).

