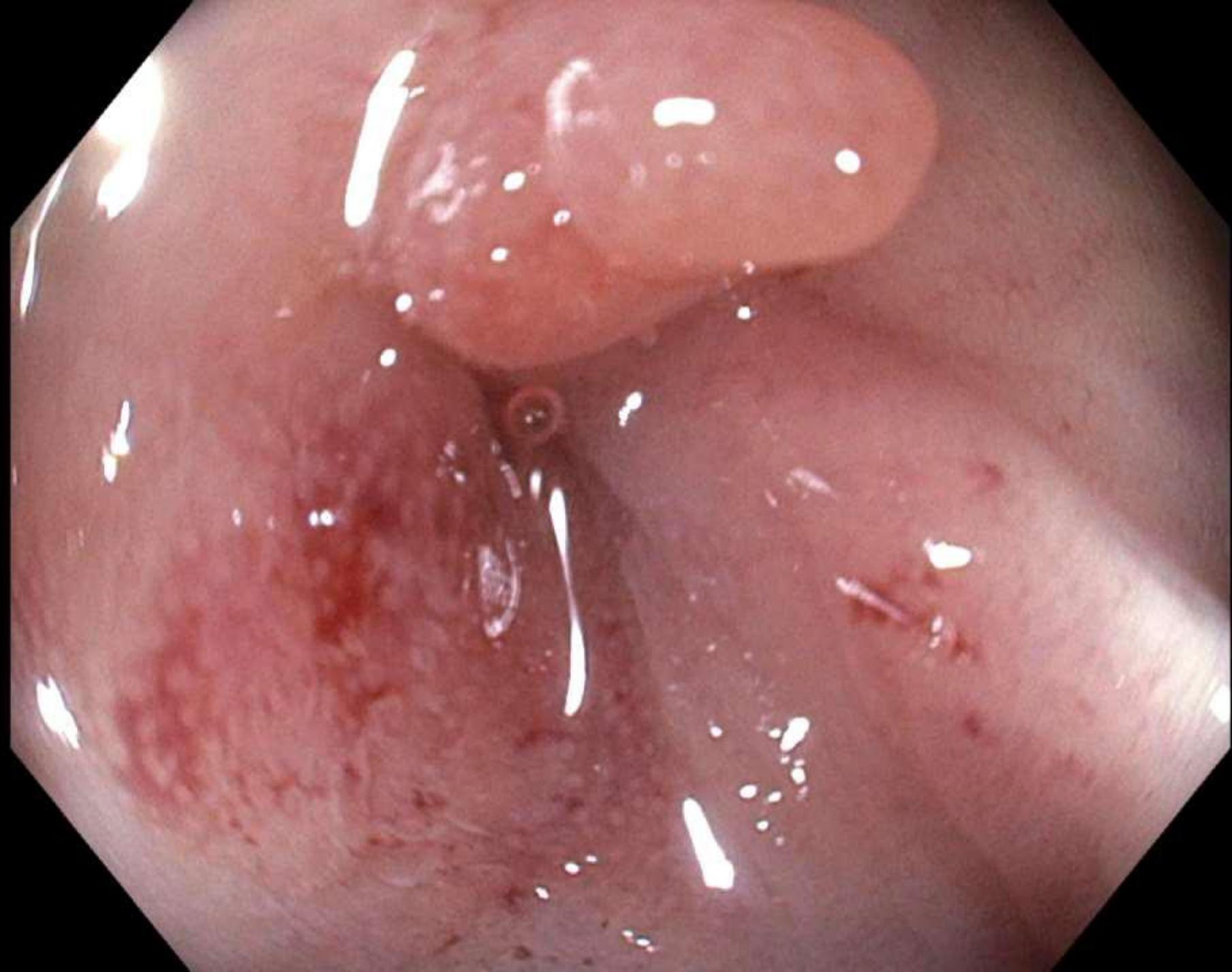


# Case Discussion

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# Case Discussion

- 64 y/o Lady who went for screening colonoscopy. Colonoscopy revealed ICV stenosis, inflammation in the cecum. TI wasn't traversed. Biopsies from the cecum revealed chronic active inflammation.
- She was then traveled to out of town the next day. Two days after her colonoscopy she started to have abdominal pain, bloating, obstipation. She went to ED, CT scan revealed TI stricture with possible fistula, picture of small bowel obstruction.
- She was treated with NGT, Bowel rest, IVF.
- She was referred to me for a second opinion.



# Case Discussion

What are the types of strictures?

# Case Discussion

- What are types of strictures?
  1. Inflammatory
  2. Fibrostenotic
  3. Inflammatory and Fibro-stenotic

# Case Discussion

What is the management plan would you discuss with this patient?

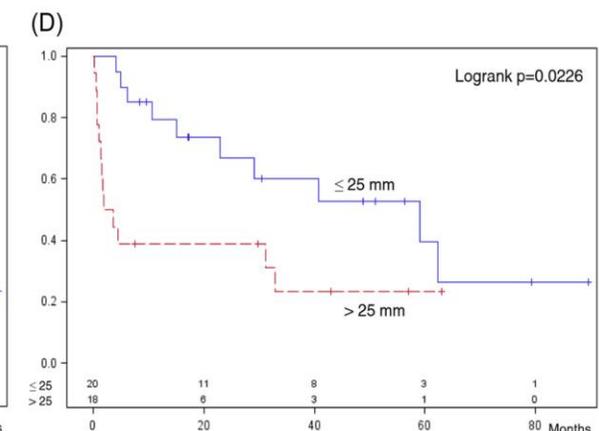
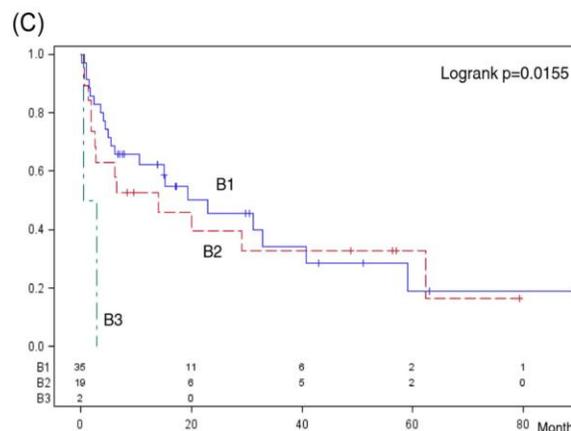
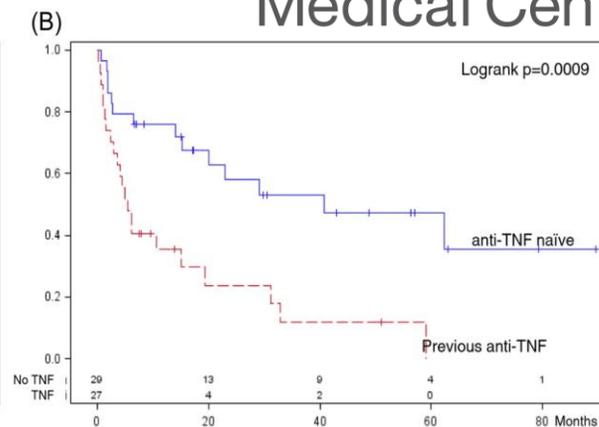
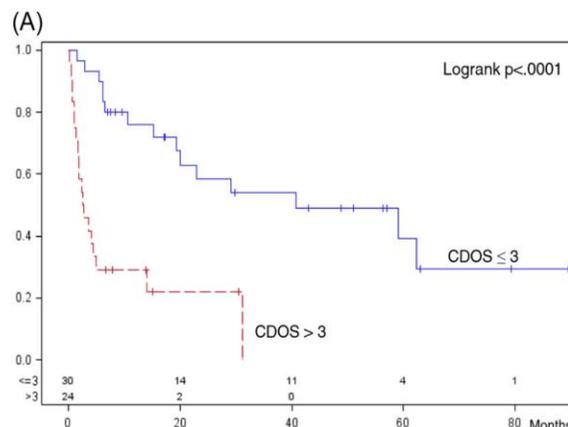
# Case Discussion

## Management

- Disease staging:
  - Lab work, inflammatory markers
  - MRE
  - Colonoscopy with biopsies with attempt to intubate T1
  - Intestinal Ultrasound
- Treatment
  - Medical
  - Surgical
  - Both

# CREOLE ADA in Stricturing Crohn's disease

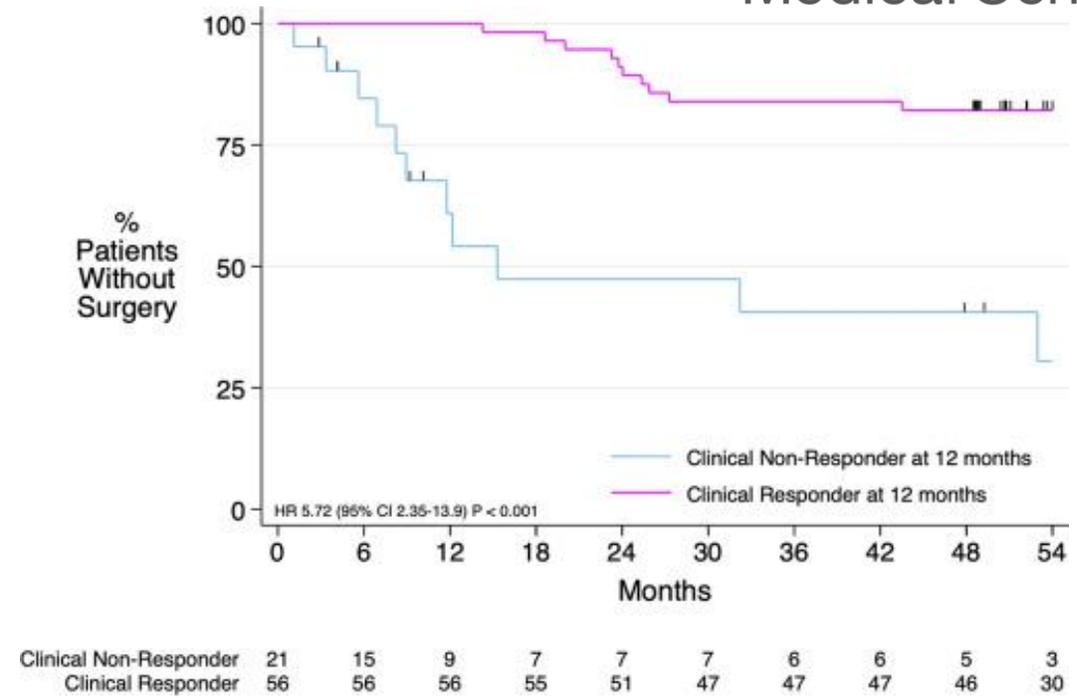
- Multicenter – Observational – Open label study (N=97)
- Patient with Crohn's disease single/multiple small bowel strictures (endoscopy/radiography) and duration of obstructive symptoms <8weeks
- Open-Label ADA (+/- immunosuppressant, taper down prednisone)
- Comparator: nil
- Primary outcome: ADA continuation, with no dilation, surgery, steroids, or severe adverse events or study withdrawals
- Limitations: Uncontrolled, open label, Obstructive symptoms graded with CDOS



- **At wk 24, 64% of patients achieved primary outcome**
- **~ 30% of entire cohort had prolonged success at 4 years**
- **~ 50% of entire cohort didn't require surgery at 4 years**

# STRIDENT Trial ADA in Stricturing Crohn's disease

- RCT, 2:1 of symptomatic Crohn's disease strictures (N=64), open-label, single center.
- First group: High-dose adalimumab induction followed by 40 mg plus thiopurine, with dose increase at 4 and/or 8 months if evidence of persisting inflammation.
- Second group: Standard ADA dosing.
- Primary outcome: Clinical responders had a lower rate of surgical resection compared with nonresponders (20% vs 52%;  $P < .001$ ) followed for 4 years after first 12 months.



Long-term surgery-free survival according to 12-month clinical response

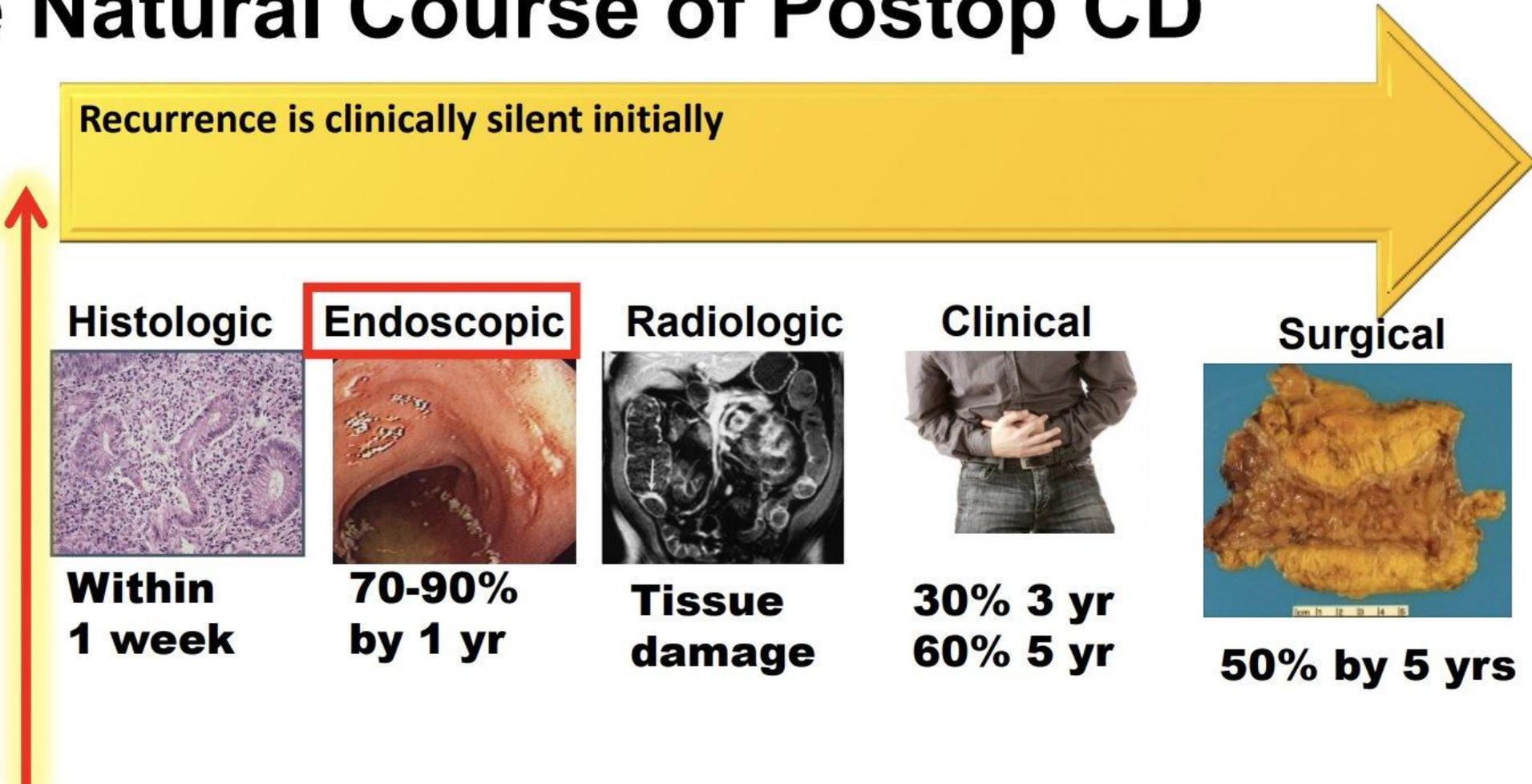
# Case Discussion

What surgery would you offer this patient?

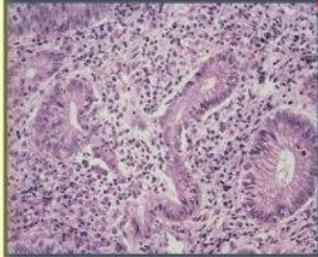
# Case Discussion

What is post-operative management?

# The Natural Course of Postop CD



Histologic



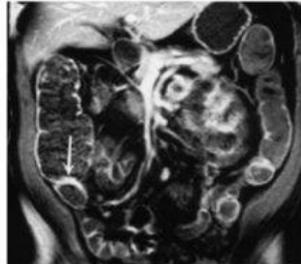
**Within  
1 week**

Endoscopic



**70-90%  
by 1 yr**

Radiologic



**Tissue  
damage**

Clinical



**30% 3 yr  
60% 5 yr**

Surgical



**50% by 5 yrs**

D'Haens GR, et al. *Gastroenterology*. 1998;114(2):262-267. Olaison G, et al. *Gut*. 1992;33(3):331-335. Rutgeerts P, et al. *Gastroenterology*. 1990;99(4):956-963. Sachar DB. *Med Clin North Am*. 1990;74(1):183-188.

# MODIFIED RUTGEERTS SCORE

i2a anastomosis

i2b before anastomosis

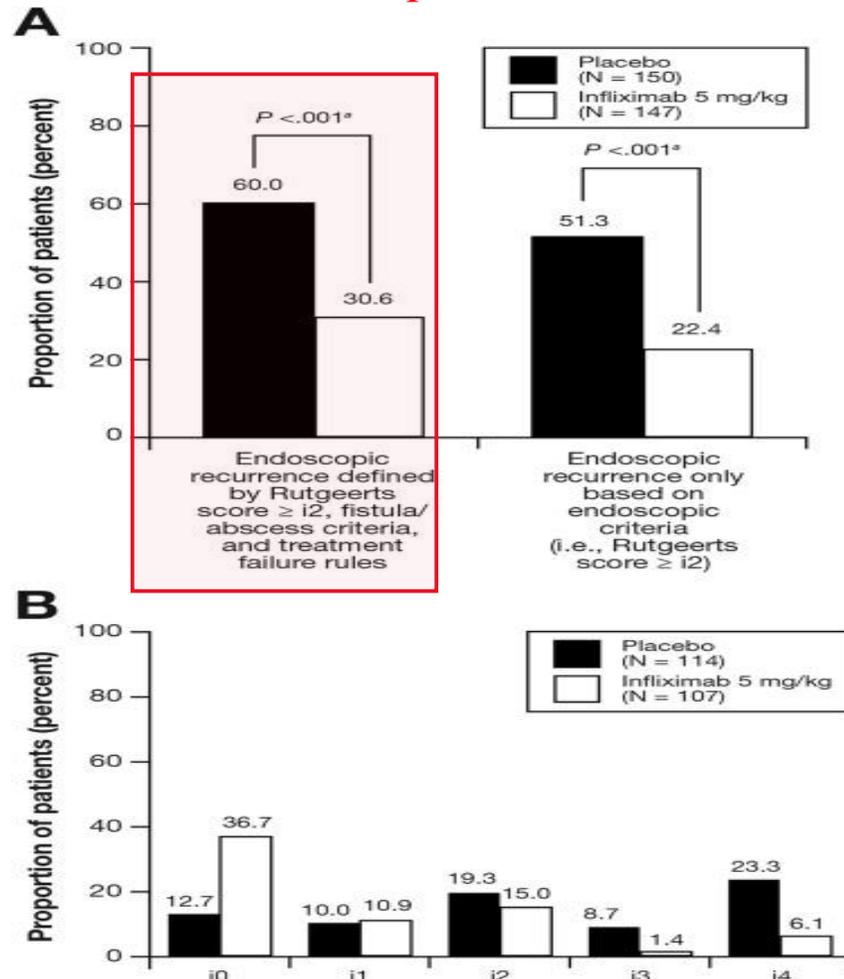
Rutgeerts score		
i0	No lesions	
i1	≤ 5 aphthous ulcers	
i2	> 5 aphthous lesions with normal mucosa between the lesions, or skip area of large lesions, or lesions confined to ileocolonic anastomosis	
i2a	Lesions confined to anastomosis (including anastomotic stenosis)	
i2b	> 5 aphthous ulcers or larger lesions, with normal mucosa in-between, in the neoterminal ileum (with or without anastomotic lesions)	
i3	Diffuse aphthous ileitis with diffusely inflamed mucosa	
i4	Diffuse inflammation with large ulcers, nodules, and/or narrowing	

# INFLIXIMAB IN POST OPERATIVE CROHN'S DISEASE

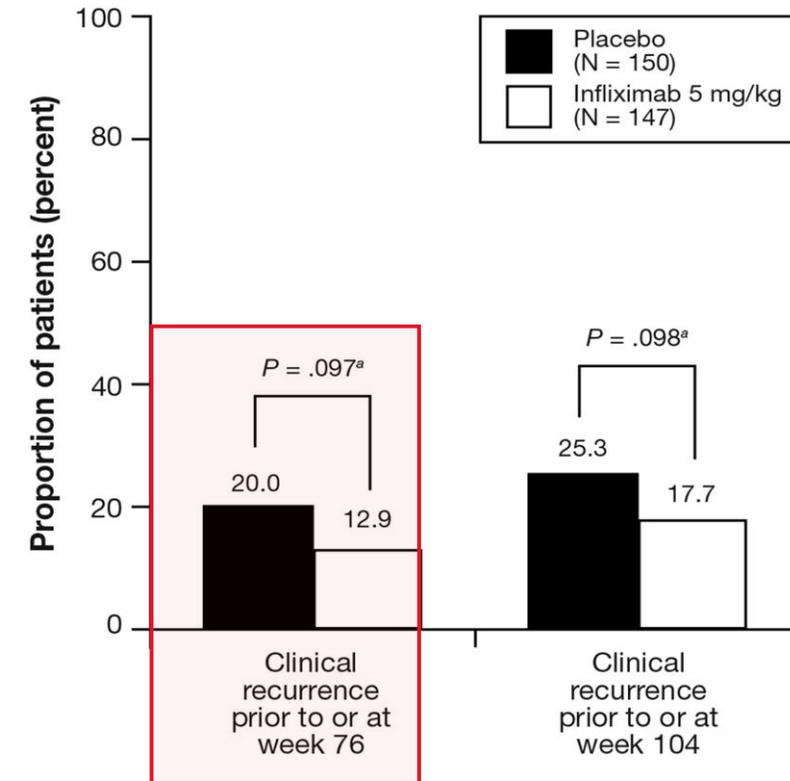
## PREVENT Study

- RCT comparing IFX vs placebo
- 297 patients: Patients were randomly assigned (1:1) to groups given infliximab (5 mg/kg) or placebo every 8 weeks for 200 weeks
- Infliximab reduces endoscopic, but not clinical recurrence, post-operative
- No induction or Therapeutic drug monitoring

Secondary end point:  
Endoscopic recurrence



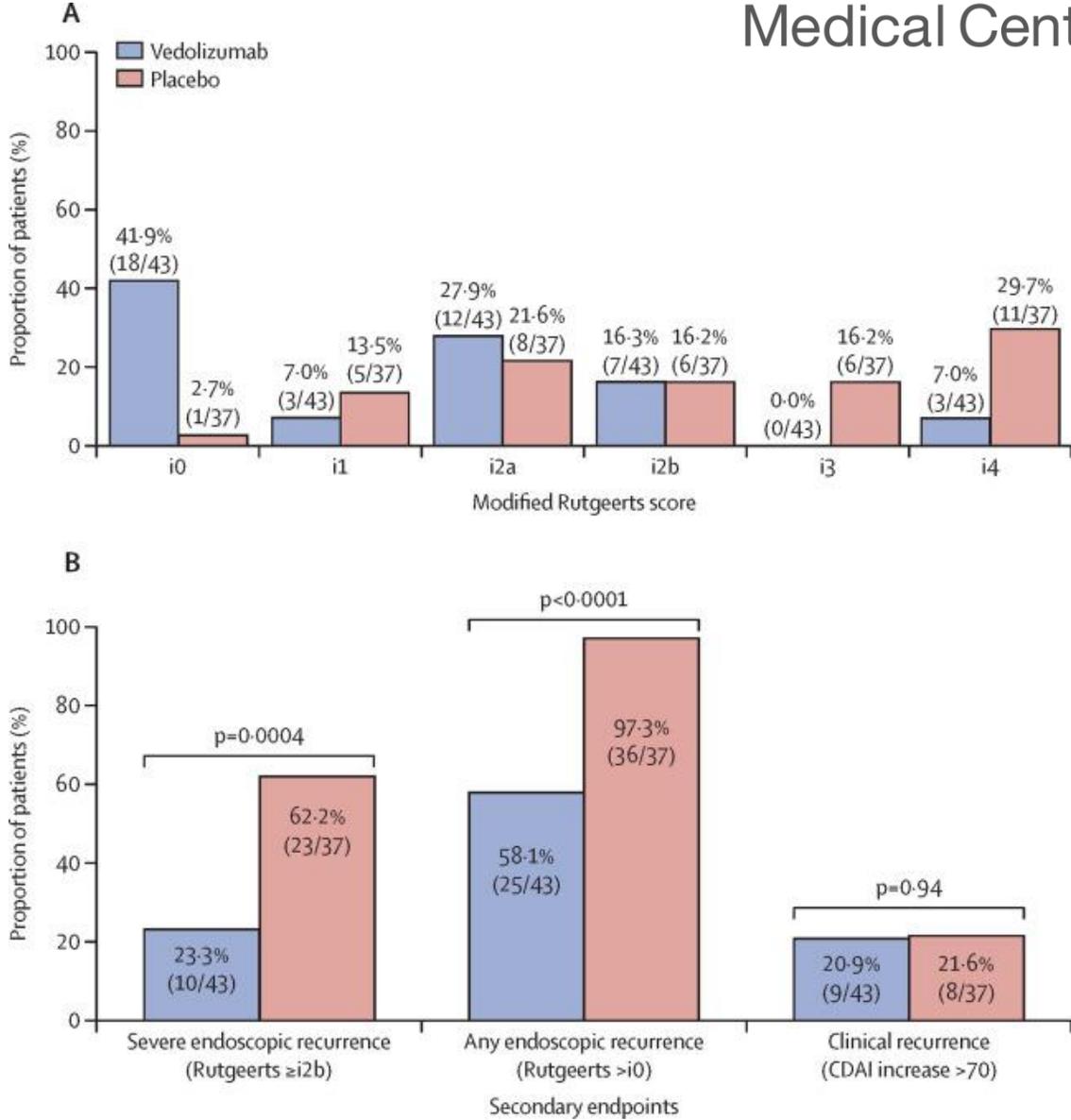
Primary end point:  
Clinical recurrence



Regueiro M Gastroenterology. 2016 Jun

# REPERIVO Study VDZ in Postoperative CD

- RCT – Double blinded
- Patient were randomly assigned, 1:1 ratio, within 4-weeks post-operative to either receive VDZ 300mg IV or Placebo at week 0,8,16,24
- No induction
- VDZ reduces endoscopic, but not clinical recurrence, post-operative



**THANK YOU**