

Which is TRUE regarding surveillance of dysplasia in IBD

- A. Standard definition endoscopy is <u>not</u> an acceptable screening method
- B. Virtual chromo-endoscopy is a valid surveillance method
- C. Recommended surveillance intervals for both pancolitis and proctitis are the same
- D. Presence of any dysplasia is indication for urgent total colectomy





Which statement is TRUE:

- A. Most dysplastic lesions in IBD are not endoscopically visible
- B. Colectomy is indicated in all cases of high-grade dysplasia
- C. Disease duration, extent, and activity are associated with risk of dysplasia
- D. Having PSC reduces risk of dysplasia





Clinical Case 7 During surveillance colonoscopy for 74-year-old patient with left sided UC for 12 years in endoscopic remission, a rectal flat 1.5cm lesion with high grade dysplasia is detected on biopsy.



Cancer Risk & Dysplasia Screening in IBD

Bincy P. Abraham, MD, MS, AGAF, FACG, FASGE

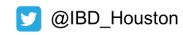
Professor of Clinical Medicine – Houston Methodist Academic Institute

Professor of Clinical Medicine – Weill Cornell Medical College

Distinguished Professor & Director, Fondren IBD Program

Director, Gastroenterology & Hepatology Fellowship







Disclosures

- Governance
 - Member American Board Internal Medicine Board of Directors
 - Member TDDC Clinical Governance Board
- Research
 - AbbVie
 - Gilead
 - o Hoffman-La Roche
 - Janssen
 - Takeda

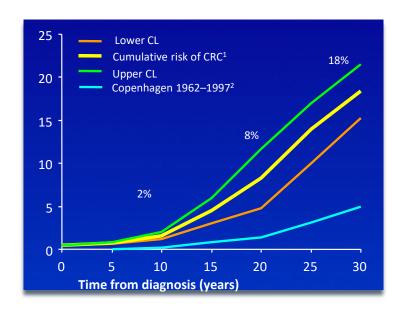


Cumulative Risk of CRC

Meta-analysis: (population-based & referral center studies):

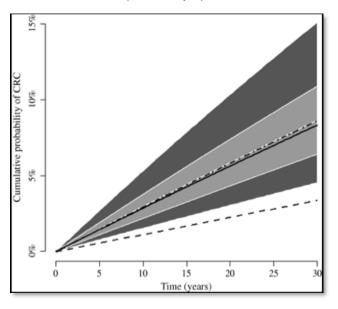
Cumulative Risk	Disease Duration
2.6% (95% CI: 0.8–4.7)	10-20 years
6.6% (95% CI: 1.3–13.8)	>20 years
Up to 21%	>20 years + Extensive Disease

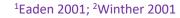
Colorectal Cancer in Ulcerative Colitis: Cumulative Risk



Colorectal & Small Bowel Cancer in CD: Cumulative Incidence:

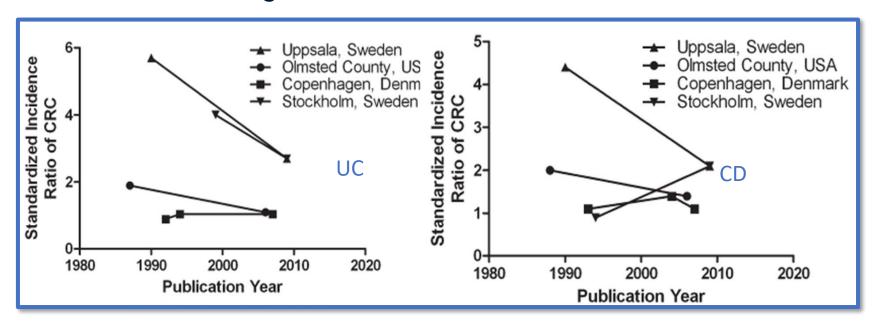
(Meta-Analysis)





Incidence & Rates

- Meta-analysis: 9 population- based studies
 - NS decreasing trend in risk of CRC in IBD over time



However STILL ~ 2X >risk than non – IBD population



Risk Factors for IBD-Associated Colorectal Neoplasia

Patient Factors	Disease Factors	Endoscopic Features
 PSC from time of diagnosis s/p liver transplant s/p proctocolectomy (pouch Ca) 	Disease Duration>7-8 yearsincreases linearly	 CD: >30-50% colon mucosa involved UC: 10-15x: pancolitis 2x: left sided colitis 0 risk: proctitis
Colorectal Neoplasia HxIncreased synchronous/ metachronous lesions	Disease Extent (CD >30-50% colon involvement)	StricturesUC > CDproximal location
Family History (1 st degree relative)	Endoscopic activity	Foreshortened tubular colon
Smoking	Histologic activity	(pseudopolyps)



New Guidelines: When to start surveillance?

	Previous:	Current:
Extensive Colitis	≥8-10 yrs from dx	≥8 yrs from dx
Left-sided Colitis	≥10-15 yrs from dx	≥8 yrs from dx
Repeat	1-2 years	1-3 years
Proctitis	Ş	?
PSC	Annually from diagnosis	



Dysplasia Surveillance

Ideal:

- Perform by experienced gastroenterologist / at IBD center
- Disease in remission
- *Do not delay if active disease
- Adequate bowel preparation
- Minimal pseudopolyps

Standard of Care:

- Standard definition scopes no longer acceptable!
- High-definition white light colonoscopy:
 - 2 to 4 random biopsies q10cm from cecum to rectum (minimum 32 biopsies)
 - If lesions found: add targeted biopsies + biopsies from surrounding "normal" mucosa (to evaluate for invisible dysplasia or inflammation)
- Chromoendoscopy (indigo carmine/ methylene blue)
 - Targeted biopsies of lesions + random biopsies (to evaluate for invisible dysplasia or inflammation)



To Chromo or Not to Chromo? Dye Chromoendoscopy (CE) v. High Definition – White Light Endoscopy (HD-WLE)

Meta-analysis (3 RCTs & 3 observational studies)

- 1358 IBD patients undergoing surveillance: (670 CE, 688 HD-WLE)
- More dysplasia found on CE vs. HD-WLE (18.8% vs. 9 %, P=0.08)

Systematic analysis of 3 RCTs

- (242 CE, 151 HD-WLE)
- More dysplasia found on CE (12.4%) vs. HD-WLE (10.4%)



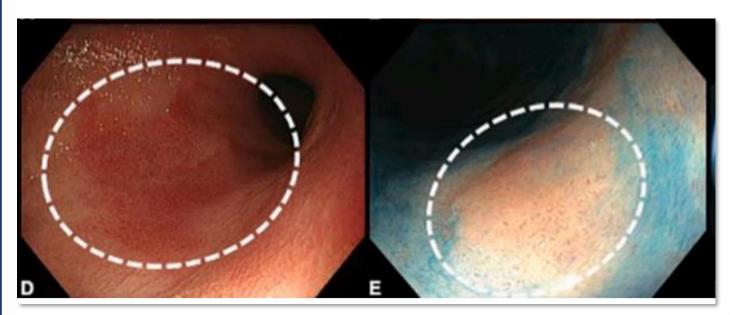
Virtual Chromoendoscopy?

- Fuji Intelligent Chromo Endoscopy (FICE) and iSCAN (Pentax)
- Narrow band imaging:
 - Currently not recommended:
 - Does not increase yield of dysplasia detection



Dye Chromoendoscopy

Methylene blue or Indigo Carmine







Endoscopically "Invisible" Dysplasia

Invisible Dysplasia on Random Biopsies

Confirm by expert GI / IBD Pathologist

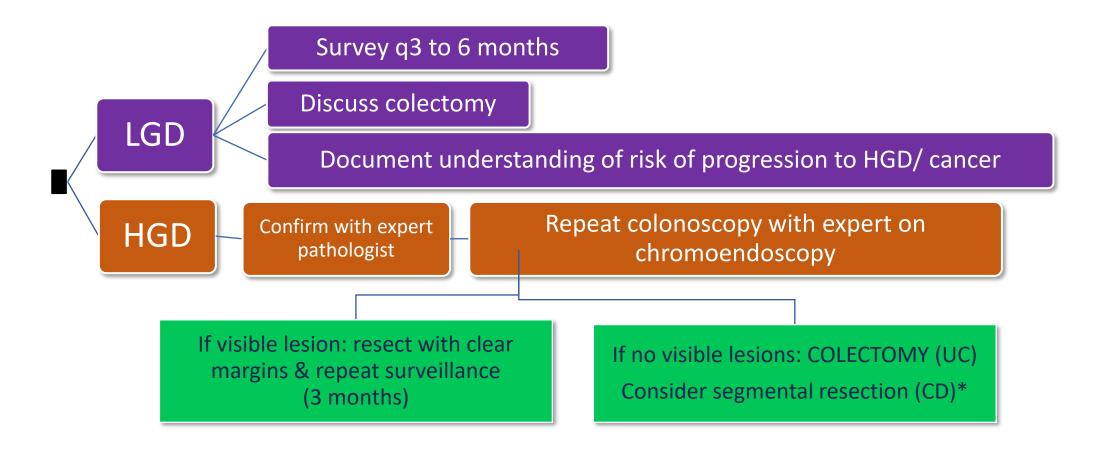
Repeat Enhanced Colonoscopy (chromoendoscopy) by Expert GI/ IBD specialist

Up to 1/3 of "invisible" dysplasia is actually visible!

If no lesions: Random biopsies q10cm (minimum 32)



Endoscopically "Invisible" Dysplasia





Management of Visible Dysplasia:

Multimodal Management

Resect (endoscopic/surgical)

Obtain/ Maintain disease quiescence (medical management)

Modify risk factors (smoking)

Continue close colonoscopic surveillance (unless total proctocolectomy).

Goal: en bloc resection, negative margins

If unable (EMR, ?ESD) → then surgery

Challenges/ Risks

- Increased risk submucosal fibrosis (inflammation)
- Incomplete resection
- Perforation
- Bleeding
- Recurrence



Approach to Visible Dysplasia

DALM: Dysplasia Associated

Lesion/ Mass

ALM:

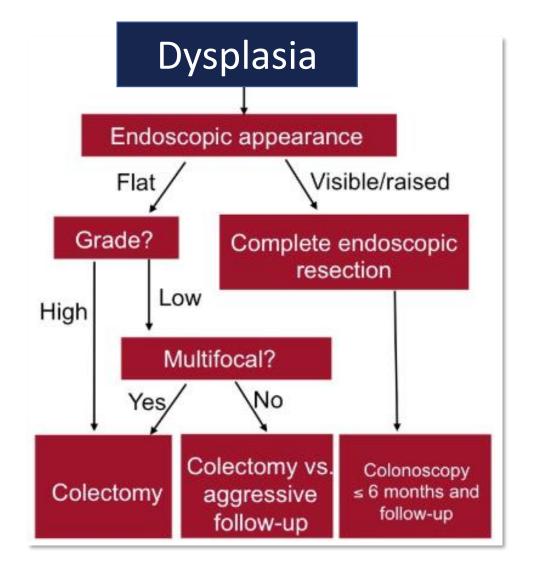
Adenoma

Like Mass

The terms "DALM" and "ALM" are being replaced by:

- "polypoid"
- · "non-polypoid"
- "flat"
- · "invisible" dysplasia

Paris Classification





Risks of Unresected Disease

- Always recommend total proctocolectomy when dysplasia is found (visible/invisible)
- High risk of synchronous dysplasia/cancer
- High risk of later development of metachronous neoplasia

Retrospective study:

- 75 CD patients w/ localized colon cancer undergoing resection:
- 39%: at least one metachronous cancer
- Mean time to new dysplasia 5 years
- Mean time to cancer 6.8 years, respectively.



Dysplasia in IBD

- Dysplasia risk is decreasing but still 2x higher than the general population
- Disease duration, extent, and activity conveys increased risk
- Most dysplasia is now "visible" therefore use high definition and consider chromoendoscopy in high-risk patients
- Endoscopically visible dysplasia (LGD/ HGD) if resected with clear margins may avoid colectomy





Clinical Case 7 During surveillance colonoscopy for 74-year-old patient with left sided UC for 12 years in endoscopic remission, a rectal flat 1.5cm lesion with high grade dysplasia is detected on biopsy.



Which is TRUE regarding surveillance of dysplasia in IBD

- A. Standard definition endoscopy is not an acceptable screening method
- B. Virtual chromo-endoscopy is a valid surveillance method
- C. Recommended surveillance intervals for both pancolitis and proctitis are the same
- D. Presence of any dysplasia is indication for urgent total colectomy





Which is TRUE regarding surveillance of dysplasia in IBD

- A. Standard definition endoscopy is not an acceptable screening method
- B. Virtual chromo-endoscopy is a valid surveillance method
- C. Recommended surveillance intervals for both pancolitis and proctitis are the same
- D. Presence of any dysplasia is indication for urgent total colectomy





Which statement is TRUE:

- A. Most dysplastic lesions in IBD are not endoscopically visible
- B. Colectomy is indicated in all cases of high-grade dysplasia
- C. Disease duration, extent, and activity are associated with risk of dysplasia
- D. Having PSC reduces risk of dysplasia





Which statement is TRUE:

- A. Most dysplastic lesions in IBD are not endoscopically visible
- B. Colectomy is indicated in all cases of high-grade dysplasia
- C. Disease duration, extent, and activity are associated with risk of dysplasia
- D. Having PSC reduces risk of dysplasia







