<u>The Year in Review Series</u>: Case 2. Anemia Case-based NBME review



Howard J. Sachs, MD <u>www.12DaysinMarch.com</u> E-mail: Howard@12daysinmarch.com <u>The Year in Review Series</u>: Case 2. Anemia Case-based NBME review







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Tutorial Services



- 1. HCT 30%, MCV 72, low iron, low iron binding capacity
- 2. HCT 30%, MCV 105, low B-12, (+) intrinsic factor antibodies
- 3. HCT 30%, MCV 72, low iron, high iron binding capacity
- 4. HCT 30%, MCV 82, elevated indirect bilirubin









- 1. Crypt abscess with neutrophilic infiltrate
- 2. Granulomatous infiltrate with transmural inflammation
- 3. Thin walled vessel lined by epithelium and little smooth muscle
- 4. Poorly differentiated cells with few glands
- 5. Tubular adenoma with evidence of DNA mismatch repair

The biopsy reveals the presence of granulomas. Which of the following epiphenomenon is the patient most likely to experience?





Video Capsule Endoscopy, Indication: GI blood loss of unclear origin

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Case 2: Another instance where you don't need the graphic to answer the question

So which of the choices are consistent with GI blood loss?

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<u>IDA</u>: Iron loss (GI/GYN) Nutrition/Absorption (e.g. Celiac disease)

<u>ACD</u>: IL-6 release (↑ Hepcidin; ↓ EPO)

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Macrocytic Anemia 2° to Autoimmune Gastritis

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Type II Hypersensitivity Reaction <u>Complication</u>: Gastric Carcinoma $2^{\circ} \uparrow$ Gastrin

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Normocytic anemia with elevated unconjugated bilirubin = hemolysis



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1. Crypt abscess with neutrophilic infiltrate

<u>Ulcerative (Neutrophilic) Colitis:</u>

Continuous involvement from anorectal region "Colitis" – doesn't involve small bowel Not associated with aphthae (macroscopic: pseudopolyps)

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Angiodysplasia (AVM)



The vascular channels may be separated from the intestinal lumen only by the vascular wall and a layer of attenuated epithelial cells.

Minor injury \rightarrow significant bleeding.



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Adenocarcinoma

Lynch Syndrome (hereditary non-polyposis): DNA mismatch repair (MMR) a/w microsatellite instability. Right sided CRC, young age (a/w endometrial cancer)

> <u>Familial adenomatous polyposis</u>: a/w APC mutation (Adenomatous Polyposis Coli) 100's of adenomas and colon cancer at young age



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<u>Crohn's (Granulomatous) Colitis</u> Terminal ileum > ileocolonic > colonic Aphthae: earliest lesion → progress, coalesce Skip Lesions



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Cutaneous



Entero-Visceral

Anal

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This question is asking about both complication and extraintestinal manifestations.

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Very dilated colon <u>Implication</u>: Toxic Megacolon (clinical diagnosis) Associated with UC

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IBD question with cholangiogram = PSC

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MSK Extraintestinal Manifestation: Fusion of SI joint c/w sacroiliitis

Inflammatory LBP

- 1. Indolent
- 2. Persistent
- $3. \ Worse \ in \ AM$
- 4. Better with activity
- 5. Young male



Steatorrhea Fat soluble vitamin deficiencies Renal stones (↑ oxalate absorption)

> <u>Skin</u> Erythema nodusum Enterocutaneous fistulae

 $\frac{\text{Loss of Terminal Ileum}}{\text{B-12 deficiency}}$ Gall stones (\$\forall \end{terminal enterohepatic circulation})



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