SURGICAL PATHOLOGY REPORT [SYNTHETIC]

ACCESSION #: UC-2025-14859
DATE OF PROCEDURE: 04/23/2025
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REQUESTING PHYSICIAN: Dr. Richard Ellis, Gastroenterology
PATHOLOGIST: Dr. Tammy Stein, Anatomic Pathology

CLINICAL HISTORY:

49 year old female with longstanding history of bloody diarrhea, abdominal pain, and urgency. Colonoscopy showed diffuse erythema, loss of vascular pattern, and contact bleeding from rectum to hepatic flexure. Clinical suspicion for ulcerative colitis.

SPECIMEN RECEIVED:

A. Rectum, biopsy

B. Sigmoid colon, biopsy

C. Descending colon, biopsy

D. Transverse colon, biopsy

E. Ascending colon, biopsy

F. Terminal ileum, biopsy

GROSS DESCRIPTION:

A. Received in formalin labeled "rectum" are 6 tan-pink tissue fragments measuring 2 mm in aggregate.

B. Received in formalin labeled "sigmoid colon" are 4 tan-pink tissue fragments measuring 4 mm in aggregate.

C. Received in formalin labeled "descending colon" are 6 tan-pink tissue fragments measuring 4 mm in aggregate.

D. Received in formalin labeled "transverse colon" are 4 tan-pink tissue fragments measuring 5 mm in aggregate.

E. Received in formalin labeled "ascending colon" are 3 tan-pink tissue fragments measuring 3 mm in aggregate.

F. Received in formalin labeled "terminal ileum" are 3 tan-pink tissue fragments measuring 2 mm in aggregate.

All specimens are entirely submitted in 1 cassette(s).

MICROSCOPIC DESCRIPTION:

A. Rectal mucosa shows fulminant active chronic inflammation with severe cryptitis, crypt architectural distortion, and mucosal ulceration. The inflammatory process is limited to the mucosa without evidence of granulomas. No evidence of dysplasia is identified.

B. Sigmoid colonic mucosa shows moderate active chronic inflammation with marked crypt architectural distortion, numerous crypt abscesses, and complete goblet cell depletion. The inflammatory process is limited to the mucosa without evidence of granulomas. Occasional Paneth cell metaplasia is noted.

C. Descending colonic mucosa shows moderate to severe active chronic inflammation with severe cryptitis, crypt architectural distortion, and Paneth cell metaplasia. The inflammatory process is limited to the mucosa without evidence of granulomas. Lamina propria shows increased plasma cells and lymphocytes.

D. Transverse colonic mucosa shows mild active chronic inflammation with crypt branching, crypt atrophy, and focal crypt abscesses. Mucosal edema and congestion are present.

E. Ascending colonic mucosa shows mild active chronic inflammation with diffuse crypt architectural distortion, crypt abscesses, and goblet cell depletion. Surface epithelium shows reactive changes.

F. Terminal ileal mucosa shows mild reactive changes. No evidence of chronic inflammatory bowel disease identified in this section. Focally, the colonic epithelium shows nuclear enlargement, hyperchromasia, and architectural complexity suspicious for low-grade dysplasia.

DIAGNOSIS:

A. Rectum, biopsy:

- fulminant active chronic colitis with crypt architectural distortion and goblet cell depletion
- fulminant consistent with ulcerative colitis
- No dysplasia identified
- No evidence of cytomegalovirus (CMV) infection

B. Sigmoid colon, biopsy:

- moderate active chronic colitis with crypt architectural distortion and goblet cell depletion
- Focal low-grade dysplasia identified
- moderate consistent with ulcerative colitis
- No dysplasia identified

C-E. Descending, transverse, and ascending colon, biopsies:

- moderate to severe active chronic colitis with crypt architectural distortion
- Features consistent with ulcerative colitis
- No dysplasia identified

F. Terminal ileum, biopsy:

- Mild non-specific inflammation
- No evidence of inflammatory bowel disease

COMMENT:

The histologic findings show a pattern of continuous chronic active colitis with greatest severity in the distal colon and rectum, with relative sparing of the proximal colon. The absence of granulomas, transmural inflammation, and terminal ileal involvement are features favoring ulcerative colitis over Crohn's disease. Correlation with clinical, endoscopic, and radiologic findings is recommended for definitive classification. Clinical correlation and follow-up biopsies are recommended to monitor disease activity and treatment response. The presence of low-grade dysplasia warrants close clinical follow-up and surveillance colonoscopy according to established guidelines for IBD-associated dysplasia.

SPECIAL STUDIES:

Grocott's methenamine silver (GMS) stain is negative for fungal organisms. p53 immunohistochemical stain shows focal overexpression in areas of dysplasia.

This is a synthetic educational pathology report created for AI training purposes. It does not represent a real patient case.