

Marlboro-Chesterfield Pathology, P.C.



· Surgical Pathology · Molecular Pathology · Cytology ·

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Dell A. Dembosky, M.D. Laboratory Director

Pathology Number: S2014-012365

Date Collected: 08 20 2014 Date Received: 08 20 2014

Location: Physician office

Physician (s): Doctor *Test

Patient: Patient, Test

1 Main st

Your town, NC 28374

DOB: 12 12 1970 Sex: F

Invoice/Chart #:

Specimen:

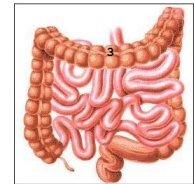
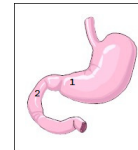
1. Stomach- Antrum
2. Sm Intestine- Duodenum
3. LG Intestine- Transverse Colon

Clinical History:

Screening

Provisional FINAL MICROSCOPIC DIAGNOSIS:

1. **CHRONIC ACTIVE HELICOBACTER GASTRITIS. IP stain POSITIVE for Helicobacter organisms.**
2. **DUODENAL MUCOSA WITH NO PATHOLOGIC DIAGNOSIS. The villous architecture is preserved. No increased intraepithelial lymphocytes seen on H&E.**
3. **TUBULAR ADENOMAS (dt)**



Gross:

1. Received in a specimen container labeled with the patient's name and "antrum" are 2 pieces of tan-pink tissue measuring 2 and 3 mm. The specimen is submitted entirely.
2. Received in a specimen container labeled with the patient's name and "duodenum" are 3 pieces of tan-pink tissue measuring 2-3 mm. The specimen is submitted entirely.
3. Received in a specimen container labeled with the patient's name and "transverse colon" are 4 pieces of tan-pink tissue measuring 4-5 mm. The specimen is submitted entirely.

(DAD/dt)

ICD9 Codes: 211.2 211.3 535.50

CPT Codes: 88305(3) 88342

Note: All IHC tissue is formalin fixed paraffin embedded using Ventana's Benchmark XT's UltraView DAB Detection's System. The degree and percentage of positively stained cells are used in the ASCO/CAP scoring method. Each test is confirmed with known controls. All special stains are confirmed with positive controls. Antibody Clones used: C-ERB-Her-2/neu(4B5), ER(SP1), Ki-67(30-9), P-53(Bp-53-11), PR(1E2), CDX-2 (EPR2764Y), CMV, PCP, HSV I&II, EBV and P504S(P) were developed and the performance characteristics were determined by Marlboro-Chesterfield Pathology, PC. They have not been cleared or approved by the US Food and Drug Administration. FDA does not require these tests to go through premarket FDA review. These tests are used for clinical purposes and are not regarded as investigational or for research. This Laboratory is certified under the Clinical Laboratory Improvement Amendments (CLIA) as qualified to perform high complexity clinical laboratory testing.

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