

QDetect® Gastrointestinal Panel

CLIA #: 31D2026917



PATIENT INFORMATION:

Patient, Test DOB: 1/2/1990 Gender/Age: M/31 SS#: UN:844008994



SPECIMEN INFORMATION:

Accession #: **TS2133351**Procedure Date: **8/10/2021**Date Received: **8/10/2021**Reported On: **8/11/2021**



PHYSICIAN INFORMATION:

Test Physician, MD
Test Practice
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Gastrointestinal Pathogen Panel (RTPCR) (STOOL) Bacteria Campylobacter Not Detected (C. jejuni/ C. coli) Clostridium difficile **Detected** Toxin A/B gene only * Correlate with C. diffcile Toxin A/B, EIA Escherichia coli O157 Not Detected (E. coli O157) Enteroaggregative E.coli Not Detected (EAEC) Enterotoxigenic E.coli Not Detected (ETEC) It/st Salmonella spp. Not Detected Shigalike toxin producing E.coli Not Detected (STEC) stx1/stx2 Shigella spp. / Enteroinvasive E.coli Not Detected (EIEC) Not Detected Vibrio spp. (V. vulnificus/ V. cholerae) Vibrio parahaemolyticus Not Detected Yersinia enterocolitica Not Detected Viruses Adenovirus F 40/41 Not Detected Norovirus GI/GII Not Detected Rotavirus A Not Detected **Parasites** Cryptosporidium Not Detected Entamoeba histolytica Not Detected Giardia lamblia Not Detected

DISCLAIMER:

BioCode® Gastrointestinal Pathogen Panel (GPP)

The BioCode[®] Gastrointestinal Pathogen Panel (GPP) is a qualitative multiplexed nucleic acid-based in vitro diagnostic test approved by the United States Food & Drug Administration (FDA) for use with the BioCode MDx3000 Instrument. The BioCode GPP is capable of the simultaneous detection and identification of nucleic acids from multiple bacteria, viruses, and parasites extracted from stool preserved in CaryBlair

transport medium obtained from individuals with signs and/or symptoms of gastrointestinal infection.
The BioCode GPP is indicated as an aid in the diagnosis of specific agents of gastrointestinal illness and results are meant to be used in conjunction with other clinical, laboratory, and epidemiological data.

Positive results do not rule out co-infection with organisms not included in the BioCode Gastrointestinal Pathogen Panel. The agent detected may not be the definite cause of the disease. Negative results in the setting of clinical illness compatible with gastroenteritis may be due to infection by pathogens that are not

detected by this test or noninfectious causes such as ulcerative colitis, irritable bowel syndrome, or Crohn's disease. This device is not intended to monitor or guide treatment for C.difficile infection. * Detection of C. difficile Toxin A/B gene does not imply active toxin production. Please correlate with C. difficile Toxin A/B, EIA results.

Fecal Fat, Qualitative:

Castor oil and mineral oil may mimic the appearance of neutral fats. Therefore, the presence of large quantities of neutral fat may indicate that the patient may have ingested either of these oils, thus causing a false positive result.

Calprotectin (Inova QUANTA Flash Chemiluminescent Immunoassay):

This assay is FDA approved and is a quantitative test for detecting fecal concentration of calprotectin. Fecal Calprotectin is an indicator of neutrophilic presence in the stool and is notspecific for inflammatory bowel disease. Values obtained with different manufacturers' assaymethods must not be used interchangeably. Results must be interpreted in conjunction with other clinical and laboratory findings.

OVA + PARASITE, STOOL

Result: No ova, cysts, or parasites seen.

Normal

Comments: The results were obtained using wet preparation and Trichrome stained smear. This test does not include testing for Cryptosporidium, Cyclospora, or Microsporidia. These tests may be ordered separately.

WHITE BLOOD CELLS, STOOL

Result: No white blood cells seen.

Normal

Reference Range: Normal: No white blood cells seen.

FECAL FAT, QUALITATIVE

Neutral Fat: < 60 fat globules / HPF

Normal

Reference Range: Normal: < 60 fat globules/HPF Increased: ≥ 60 fat globules/HPF

CALPROTECTIN, STOOL

Result: 35.5 mg/kg

Normal

Reference Range: Normal: < 50 mg/kg

Borderline: ≥ 50 – < 120 mg/kg Abnormal: ≥ 120 mg/kg

Comments: Followup: Repeat as clinically indicated.

PANCREATICELASTASE ELISA, STOOL

Result: 250 µg/mL

Normal

Reference Range: Normal: > 200 µg/mL

Slight to moderate pancreatic insufficiency: 100200 $\mu g/mL$

Severe pancreatic insufficiency: < 100 µg/mL

C. difficile TOXIN A/B, STOOL (EIA) *

Result: Positive

Abnormal

BILLING CODES:

CPT: 87507, 89055, 82705, 87177, 87209, 87324, 83993, 82656, 85549

ICD10: R19.7

Physician: Test Physician, MD | Medical Director: M. N. Qureshi, MD, PhD | Electronically signed out by: Test Pathologist1