Advanta Analytical Laboratories
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Patient: Race Bannon
DOB: 02-05-1960
Gender: Male
Accession: 770519
Specimen: Fecal

Advantage GastroIntestinal PCR Panel
FINAL

Referring Physician: Doe, John
Client: Demo Account
Collection Date: 04-05-2022
Received Date: 04-05-2022
Report Date: 04-05-2022

Organism(s) Tested	*Qualitative	Quantitative DNA copies / mL	Qualitative Result
Ancylostoma duodenale	N/A	N/A	NEGATIVE
Campylobacter spp.	N/A	N/A	NEGATIVE
Clostridium difficile **	N/A	N/A	NEGATIVE
Cryptosporidium spp.	N/A	N/A	NEGATIVE
Entamoeba histolytica	N/A	N/A	NEGATIVE
Enterobius vermicularis	N/A	N/A	NEGATIVE
Enterohemorragic/verotoxin producing Escherichia coli	N/A	N/A	NEGATIVE
Enteroinvasive Escherichia coli	**MEDIUM**	**2.31E+05**	**POSITIVE**
Giardia lamblia	N/A	N/A	NEGATIVE
Helicobacter pylori	N/A	N/A	NEGATIVE
Human adenovirus	N/A	N/A	NEGATIVE
Human astrovirus	N/A	N/A	NEGATIVE
Necator americanus	**MEDIUM**	**1.54E+05**	**POSITIVE**
Norovirus G1	N/A	N/A	NEGATIVE
Norovirus G2	N/A	N/A	NEGATIVE
Rotavirus	N/A	N/A	NEGATIVE
Salmonella spp.	N/A	N/A	NEGATIVE
Sapovirus	N/A	N/A	NEGATIVE
Shigella spp.	N/A	N/A	NEGATIVE
Yersinia enterocolitica	N/A	N/A	NEGATIVE

Testing Disclaimer **

1. Neonates + infants < 12 months with diarrhea: Due to the high prevalence of asymptomatic carriage of toxigenic C. difficile in infants, testing for C. difficile is not recommended.
2. Children 1-2 years of age with diarrhea: C. difficile testing should not be routinely performed unless other infectious or noninfectious causes have been excluded.
3. Children > 2 years of age with diarrhea: Testing is recommended for patients with prolonged or worsening diarrhea and risk factors (eg, underlying inflammatory bowel disease or immunocompromising conditions) or relevant exposures (eg, contact with the healthcare system or recent antibiotics).

Laboratory Notice

This nucleic acid amplification test was developed, and its performance characteristics determined with polymerase chain reaction (PCR) by Advanta Analytical Laboratories, LLC. This test is a lab developed test and has not been cleared or approved by the FDA. This laboratory is regulated under CLIA as qualified to perform high complexity testing and accredited by the College of American Pathologists (CAP). These tests are used for clinical purposes. They should not be regarded as investigational or for research purposes.