HOTLINE: Effective August 16, 2021

New Test **3003748** **Inflammatory Bowel Disease Differentiation Panel** **IBD-PAN**

Methodology: Semi-Quantitative Enzyme-Linked Immunosorbent Assay/Semi-Quantitative Indirect Fluorescent Antibody
Performed: Sat-Sun
Reported: 1-4 days

Specimen Required: Patient Prep: N/A
Collect: Serum Separator Tube (SST).
Specimen Preparation: Separate from cells ASAP or within 2 hours of collection. Transfer 1.5 mL serum to an ARUP Standard Transport Tube. (Min: 0.6 mL)
Storage/Transport Temperature: Refrigerated.
Remarks: N/A
Unacceptable Conditions: Contaminated, heat-inactivated, hemolyzed, or severely lipemic specimens.
Stability (collection to initiation of testing): After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 year (avoid repeated freeze/thaw cycles)

Reference Interval:

Test Number	Components	Reference Interval	
	Saccharomyces cerevisiae Antibody, IgG		
		20.0 Units or less	Negative
		20.1-24.9 Units	Equivocal
		25.0 Units or greater	Positive
	Saccharomyces cerevisiae Antibody, IgA		
		20.0 Units or less	Negative
		20.1-24.9 Units	Equivocal
		25.0 Units or greater	Positive
	ANCA IFA Titer	Less than 1:20	
	ANCA IFA Pattern	None Detected	

Interpretive Data:
Refer to report.

Note: This test may be a useful tool for distinguishing ulcerative colitis (UC) from Crohn disease (CD) in patients with suspected inflammatory bowel disease. ANCA IFA is simultaneously tested on ethanol- and formalin-fixed slides to allow differentiation of C- and P-ANCA patterns.

CPT Code(s): 86671 x2; 86255

New York DOH Approved.

HOTLINE NOTE: Refer to the Test Mix Addendum for interface build information.