TEST CHANGE

Inflammatory Bowel Disease Differentiation Panel

3003748, IBD-PAN

Specimen Requirements:	
Patient Preparation:	N/A
Collect:	Serum separator tube~~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)
Transport Temperature:	Refrigerated.
Unacceptable Conditions:	Contaminated, heat-inactivated, hemolyzed, or severely lipemic specimens.
Remarks:	N/A
Stability:	After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 30 days (avoid repeated freeze/thaw cycles)
Methodology:	Semi-Quantitative Indirect Fluorescent Antibody (IFA)/Semi-Quantitative Enzyme Immunoassay (EIA)
Performed:	Mon, Wed, Fri ~~Sun-Sat~~
Reported:	1-4 days
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 Codes:	86036; 86671 x2
New York DOH Approval Status:	This test is New York DOH approved.
Interpretive Data:	
Refer to report.	
Reference Interval:	

Test Number	Components	Reference Interval		
	S. cerevisiae Antibody, IgG			
		20.0 Units or less	Negative	
		20.1-24.9 Units	Equivocal	
		25.0 Units or greater	Positive	
	S. 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		