

New Test

3002479

Autoimmune Liver Disease Reflexive Panel

LIVER PAN



Additional Technical Information

Methodology: Semi-Quantitative Enzyme-Linked Immunosorbent Assay/Semi-Quantitative Indirect Fluorescent Antibody
Performed: Sun-Sat
Reported: 1-8 days

Specimen Required: Collect: Serum Separator Tube (SST).

Specimen Preparation: Separate from cells ASAP or within 2 hours of collection. Transfer 0.5 mL serum to an ARUP Standard Transport Tube. (Min: 0.3 mL)

Storage/Transport Temperature: Refrigerated.

Unacceptable Conditions: Non-serum, heat-inactivated, contaminated, grossly icteric, severely lipemic, grossly hemolyzed specimens or inclusion of fibrin clot.

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	
0050065	Mitochondrial M2 Antibody, IgG (ELISA)	20.0 Units or less	Negative
		20.1-24.9 Units	Equivocal
		25.0 Units or greater	Positive
0051174	F-Actin (Smooth Muscle) Antibody, IgG by ELISA with Reflex to Smooth Muscle Antibody, IgG Titer	Test Number	
		Components	
		Reference Interval	
			F-Actin (Smooth Muscle) Antibody, IgG
		20-30 Units	Weak Positive - Suggest repeat testing in two to three weeks with fresh specimen
		31 Units or greater	Positive – Suggestive of autoimmune hepatitis or chronic active hepatitis
0051244	Smooth Muscle Antibody, IgG Titer	Less than 1:20	
0055235	Soluble Liver Antigen Antibody, IgG	20.0 Units or less	Negative
		20.1-24.9 Units	Equivocal
		25.0 Units or greater	Positive
0055241	Liver-Kidney Microsome - 1 Antibody, IgG	20.0 Units or less	Negative
		20.1-24.9 Units	Equivocal
		25.0 Units or greater	Positive
3000082	Antinuclear Antibody (ANA) with HEp-2 Substrate, IgG by IFA	Less than 1:80	

Interpretive Data:

Refer to report.

Note: If F-Actin, IgG by ELISA is 20 Units or greater, then Smooth Muscle Antibody (SMA), IgG by IFA titer will be added. Additional charges apply.

ANA are determined by indirect fluorescence assay (IFA) using HEp-2 substrate and IgG-specific conjugate at a screening dilution of 1:80. If positive, patterns reported include homogeneous, speckled, centromere, nucleolar, nuclear dots, or cytoplasmic. All positive results are reported with endpoint titers, at no additional charge.

CPT Code(s): 86039, 86376, 83516 x3; if reflexed, add 86256

New York DOH Approved.

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