

Quarterly HOTLINE: Effective November 12, 2018

3000601 Antinuclear Antibody (ANA) with HEp-2 Substrate, IgG by IFA

with Reflex by Pattern



New Test

Additional Technical Information

Methodology: Semi-Quantitative Indirect Fluorescent Antibody/Qualitative Enzyme-Linked Immunosorbent Assay/Semi-Quantitative Enzyme-

Linked Immunosorbent Assay/Semi-Quantitative Multiplex Bead Assay/Qualitative Immunoblot

Performed: Sun-Sat Reported: 1-8 days

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

Specimen Preparation: Separate serum from cells ASAP or within 2 hours of collection. Transfer 1 mL serum to an ARUP Standard

ANA AB PAT

Transport Tube. (Min: 0.6 mL)

Storage/Transport Temperature: Refrigerated

Unacceptable Conditions: Non-serum specimens. Contaminated, grossly hemolyzed, heat-inactivated, 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			
3000082	Antinuclear Antibody (ANA), HEp-2, IgG	Less than 1:80			
2003040	PM/Scl-100 Antibody, IgG by Immunoblot	Negative			
2012173	Fibrillarin (U3 RNP) Antibody, IgG	Negative			
0050215	Double-Stranded DNA (dsDNA) Antibody, IgG by	Test Number	Components Double-Stranded DNA (dsDNA) Antibody, IgG by ELISA		Reference Interval
	ELISA with Reflex to dsDNA Antibody, IgG by IFA				None Detected.
		2002693	Double-Stranded DNA (dsDNA) Antibody, IgG by IFA (using Crithidia luciliae)		Less than 1:10
2005287	Chromatin Antibody, IgG	19 Units or less	Units or less Negative		
		20-60 Units		Moderate Positive	
		61 Units or greater		Strong Positive	
2001601	RNA Polymerase III Antibody, IgG	19 Units or less		Negative	
		20-39 Units		Weak Positive	
		40-80 Units		Moderate Positive	
		81 Units or greater		Strong Positive	
0050599	Scleroderma (Scl-70) (ENA) Antibody, IgG	29 AU/mL or less		Negative	
		30-40 AU/mL		Equivocal	
		41 AU/mL or greater		Positive	
0050470	RNP (U1) (Ribonucleic Protein) (ENA) Antibody, IgG	29 AU/mL or less		Negative	
		30-40 AU/mL		Equivocal	
		41 AU/mL or greater		Positive	
0050085	Smith (ENA) Antibody, IgG	29 AU/mL or less		Negative	
		30-40 AU/mL		Equivocal	
		41 AU/mL or greater		Positive	
2012074	SSA 52 and 60 (Ro) (ENA) Antibodies, IgG	Test Number	Components		Reference Interval
					29 AU/mL or Less: Negative 30-40 AU/mL: Equivocal 41 AU/mL or greater: Positive
			30-40 AU/mL: Equiv 41 AU/mL or greater		29 AU/mL or Less: Negative 30-40 AU/mL: Equivocal 41 AU/mL or greater: Positive
0050692	SSB (La) (ENA) Antibody, IgG	29 AU/mL or less		Negative	
		30-40 AU/mL		Equivocal	
		41 AU/mL or greater		Positive	

Interpretive Data: Refer to report.



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Note: The Antinuclear Antibody (ANA), IgG by IFA with Reflex by IFA Pattern begins with Nuclear Antibody (ANA) by IFA, IgG. Depending on findings, one or more reflexive tests may be required. Tests added may include Double-Stranded DNA (dsDNA) Antibody, IgG by ELISA; Double-Stranded DNA (dsDNA) Antibody, IgG by IFA (using Crithidiae luciliae); Chromatin Antibody, IgG; RNP (U1) (Ribonucleic Protein) (ENA) Antibody, IgG; Fibrillarin (U3 RNP) Antibody, IgG; Smith (ENA) Antibody, IgG; SSA 52 (Ro) (ENA) Antibody, IgG; SSA 60 (Ro) (ENA) Antibody, IgG; SSB (La) (ENA) Antibody, IgG; Scleroderma (Scl-70) (ENA) Antibody, IgG; PM/Scl-100 Antibody, IgG, by Immunoblot; and/or RNA Polymerase III Antibody, IgG. Additional charges apply.

ANA determined by indirect fluorescence assay (IFA) use 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.

CPT Code(s): 86039; if homogenous or speckled pattern add 86235 x6, 86225, and 83516; if reflexed add 86256; If nucleolar pattern add 86235 x3

and 83516

New York DOH Approved.

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