500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex:

Unknown

Specimen Collected: 11/18/2024 07:4	3 MST		
Interstitial Lung Disease H Autoantibody 2	Received: 11/18/2024	07:46 MST	Report/Verified: 11/18/2024 07:57 MST
Procedure	Result	Units	Reference Interval
Rheumatoid Factor	25 ^H	IU/mL	[0-14]
SSA-52 (Ro52) (ENA) Antibody, Ig	G 55 ^{H i1}	AU/mL	[0-40]
SSA-60 (Ro60) (ENA) Antibody, Ig	G 65 ^{H i2}	AU/mL	[0-40]
Jo-1 (Histidyl-tRNA Synthetase)		AU/mL	[0-40]
Ab,IgG			
PL-12 (alanyl-tRNA synthetase)	Positive *		[Negative]
Antibody			
PL-7 (threonyl-tRNA synthetase)	Positive *		[Negative]
Antibody			
EJ (glycyl-tRNA synthetase)	Positive *		[Negative]
Antibody	Devilation *		
OJ (isoleucyl-tRNA synthetase) Antibody	Positive *		[Negative]
SRP (Signal Recognition	Positive *		[Negative]
Particle) Ab	rositive		[Incgacive]
Ku Antibody	Positive *		[Negative]
PM/Scl 100 Antibody, IgG	Positive * ⁱ⁴		[Negative]
MDA5 (CADM-140) Ab	Positive *		[Negative]
NXP2 (Nuclear matrix protein-2)			[Negative]
Ab	j		
Interpretive Information	See Note ¹⁵		
Scleroderma (Scl-70) (ENA)	55 ^{H i6}	AU/mL	[0-40]
Antibody,IgG			
RNA Polymerase III Antibody, IgG	65 H 17	Units	[0-19]
Antinuclear Antibody (ANA), HEp-	Detected *		[<1:80]
2,IgG			
ANA Interpretive Comment	See Note ^{t1 i8}		
Cyclic Citrullinated Peptide Ab	, 55 ^{H 19}	Units	[0-19]
IgG/A			
Ha (tyrosyl-tRNA synthetase) Ab			[Negative]
Ks (asparaginyl-tRNA synthetase) Positive * t3		[Negative]
Ab	Dogitino * t4		[Negative]
Zo (phenylalanyl-tRNA synthetase) Ab	Positive * t4		[Negative]
Antinuclear Ab, Dual Pattern H	Received: 11/18/2024	07:46 MST	Report/Verified: 11/18/2024 07:57 MST
Procedure	Result	Units	Reference Interval
ANA Titer 2	1:640 *		
ANA Pattern	Speckled *		
ANA Titer	1:2560 *		

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500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Office

phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer		Patient	Age/Sex:	Unknown	
Antinuclear Ab, Dual Pattern	Received: 11/18/202	4 07:46 MST	Report/Verifi MST	.ed: 11/18/2024 07:57	
Procedure ANA Pattern 2	Result Nuclear Dot *	Units		eference Interval	
Cytoplasmic Pattern	Received: 11/18/202	4 07:46 MST	Report/Verifi MST	ed: 11/18/2024 07:57	
Procedure Cytoplasmic Titer Cytoplasm Pattern	Result 1:320 * GW Body-like *	Units		eference Interval	
Speckled Pattern Clinical associations: healthy individuals Main autoantibodies: A anti-Topo-1 (anti-Scl- (TIF1g), anti-Ku, ant Nuclear Dots Pattern Clinical associations: Main autoantibodies: A Cytoplasmic discrete d Clinical Associations: sensory neuropathy) Main autoantibodies: N	nti-SSA-52 (Ro52), a 70), Smith, anti-Ul- i-RNA polymerase, an PBC, DM, SjS, SLE, nti-NXP-2, anti-Sp10 ots/GW Body-like pat SjS, SLE, RA and so	nti-SSA-60 (RNP, anti-U2 ti-DFS70/LEI SSc, PM 0 tern	Ro60), anti- 2-RNP, anti-N 9GF-P75	-SS-B/LA, 4i-2, anti-p155/140	
List of Abbreviations Antisynthetase syndrom myopathies (IM) [derma myopathy (NAM)], inter (JIA), mixed connectiv rheumatoid arthritis (syndrome (SjS), system undifferentiated conne t2: 11/18/2024 07:43 MST (Ha (ty Ha positive by line im immunoprecipitation. P t3: 11/18/2024 07:43 MST (Ks (as Ks positive by line im immunoprecipitation. P t4: 11/18/2024 07:43 MST (Zo (pl Zo positive by line im immunoprecipitation. P	tomyositis (DM), pol stitial lung disease e tissue disease (MC RA), systemic autoim ic lupus erythematos ctive tissue disease rosyl-tRNA synthetase) A munoassay. Band cor rofile consistent wi sparaginyl-tRNA synthetas munoassay. Band cor rofile consistent wi henylalanyl-tRNA syntheta munoassay. Bands cor	ymyositis (H e (ILD), juve TD), primary mune rheumat sus (SLE), sy e (UCTD). the th Ha antibo e Ab) rresponding t th Ks antibo se) Ab) rresponding t	2M), necrotiz enile idiopat y biliary cho cic diseases ystemic sclea co 65 KDa obs ody positivit co 65 kDa obs ody positivit	zing autoimmune thic arthritis olangitis (PBC), (SARD), Sjogren cosis (SSc), served by CY. served by CY. KDa observed by	

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500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer PATIENT REPORT

Unknown

Patient Age/Sex:

<u>Test Information</u> i1: SSA-52 (Ro52) (ENA) Antibody, IgG INTERPRETIVE INFORMATION: SSA-52 (Ro52) (ENA) Antibody, IgG 29 AU/mL or Less Negative 30 - 40 AU/mL Equivocal 41 AU/mL or Greater Positive SSA-52 (Ro52) and/or SSA-60 (Ro60) antibodies are associated with a diagnosis of Sjogren syndrome, systemic lupus erythematosus (SLE), and systemic sclerosis. SSA-52 antibody overlaps significantly with the major SSc-related antibodies. SSA-52 (Ro52) antibody occurs frequently in patients with inflammatory myopathies, often in the presence of interstitial lung disease. i2: SSA-60 (Ro60) (ENA) Antibody, IgG REFERENCE INTERVAL: SSA-60 (Ro60) (ENA) Antibody, IgG 29 AU/mL or Less Negative 30 - 40 AU/mL Equivocal 41 AU/mL or Greater Positive i3: Jo-1 (Histidyl-tRNA Synthetase) Ab, IgG INTERPRETIVE INFORMATION: Jo-1 Antibody, IgG 29 AU/mL or less.....Negative 30-40 AU/mL.....Equivocal 41 AU/mL or greater.....Positive Presence of Jo-1 (antihistidyl transfer RNA [t-RNA] synthetase) antibody is associated with polymyositis and may also be seen in patients with dermatomyositis. Jo-1 antibody is associated with pulmonary involvement (interstitial lung disease), Raynaud phenomenon, arthritis, and mechanic's hands (implicated in antisynthetase syndrome). i4: PM/Scl 100 Antibody, IgG INTERPRETIVE INFORMATION: PM/Scl-100 Antibody, IqG by Immunoblot The presence of PM/Scl-100 IgG antibody along with a positive ANA IFA nucleolar pattern is associated with connective tissue diseases such as polymyositis (PM), dermatomyositis (DM), systemic sclerosis (SSc), and polymyositis/systemic sclerosis overlap syndrome. The clinical relevance of PM/Scl-100 IgG antibody with a negative ANA IFA nucleolar pattern is unknown. PM/Scl-100 is the main target epitope of the PM/Scl complex, although antibodies to other targets not detected by this assay may occur. This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug

Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

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500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex:

Unknown

Test Information

i5: Interpretive Information INTERPRETIVE INFORMATION: Interstitial Lung Disease Autoantibodies

If present, myositis-specific antibodies (MSA) are specific for myositis, and may be useful in establishing diagnosis as well as prognosis. MSAs are generally regarded as mutually exclusive with rare exceptions; the occurrence of two or more MSAs should be carefully evaluated in the context of patient's clinical presentation. Myositis-associated antibodies (MAA) may be found in patients with CTD including overlap syndromes, and are generally not specific for myositis. The following table will help in identifying the association of any antibodies found as either MSAs or MAAs.

Antibody SpecificityMSASSA 52 (Ro) (ENA) Antibody IgGX	
SSA 52 (RO) (ENA) ANLIDODY IGG	
SSA 60 (Ro) (ENA) Antibody IgG X	
Smith/RNP (ENA) Ab, IgG X	
Jo-1 (histidyl-tRNA synthetase) Ab, IgG X	
PL-12 (alanyl-tRNA synthetase) Antibody X	
PL-7 (threonyl-tRNA synthetase) Antibody X	
EJ (glycyl-tRNA synthetase) Antibody X	
OJ (isoleucyl-tRNA synthetase) Antibody X	
SRP (Signal Recognition Particle) Ab X	
Ku Antibody X	
PM/SCL 100 Antibody, IgG X	
U2 sn (small nuclear) RNP Antibody X	
Fibrillarin (U3 RNP) Ab, IgG X	
Mi-2 (nuclear helicase protein) Antibody X	
P155/140 Antibody X	
TIF-1 gamma (155 kDa) Ab X	
SAE1 (SUMO activating enzyme) Ab X	
MDA5 (CADM-140) Ab X	
NXP2 (Nuclear matrix proten-2) Ab X	

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes. Scleroderma (Scl-70) (ENA) Antibody, IgG

i6:

INTERPRETIVE INFORMATION: Scleroderma (Scl-70) (ENA) Ab, IgG

29 AU/mL or Less Negative 30 - 40 AU/mL Equivocal 41 AU/mL or Greater Positive

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Patient Age/Sex:

Unknown

Test Information

i6: Scleroderma (Scl-70) (ENA) Antibody, IgG

The presence of Scl-70 antibodies (also referred to as topoisomerase I, topo-I or ATA) is considered diagnostic for systemic sclerosis (SSc). Scl-70 antibodies alone are detected in about 20 percent of SSc patients and are associated with the diffuse form of the disease, which may include specific organ involvement and poor prognosis. Scl-70 antibodies have also been reported in a varying percentage of patients with systemic lupus erythematosus (SLE). Scl-70 (topo-1) is a DNA binding protein and anti-DNA/DNA complexes in the sera of SLE patients may bind to topo-I, leading to a false-positive result. The presence of Scl-70 antibody in sera may also be due to contamination of recombinant Scl-70 with DNA derived from cellular material used in immunoassays. Strong clinical correlation is recommended if both Scl-70 and dsDNA antibodies are detected.

Negative results do not necessarily rule out the presence of SSc. If clinical suspicion remains, consider further testing for centromere, RNA polymerase III and U3-RNP, PM/Scl, or Th/To antibodies.

i7: RNA Polymerase III Antibody, IgG INTERPRETIVE INFORMATION: RNA Polymerase III Antibody, IgG

19 Units or lessNegative
20 - 39 UnitsWeak Positive
40 - 80 UnitsModerate Positive
81 Units or greater ...Strong Positive

The presence of RNA polymerase III IgG antibody, when considered in conjunction with other laboratory and clinical findings, is an aid in the diagnosis of systemic sclerosis (SSc) with increased incidence of skin involvement and renal crisis with the diffuse cutaneous form of SSc. RNA polymerase III IgG antibody occur in about 11-23 percent of SSc patients, and typically in the absence of anti-centromere and anti-Scl-70 antibodies.

A negative result indicates no detectable IgG antibodies to the dominant antigen of RNA polymerase III and does not rule out the possibility of SSc. False-positive results may also occur due to non-specific binding of immune complexes. Strong clinical correlation is recommended.

If clinical suspicion remains, consider additional testing for other antibodies associated with SSc, including centromere, Scl-70, U3-RNP, PM/Scl, or Th/To. i8: ANA Interpretive Comment INTERPRETIVE INFORMATION: ANA Interpretive Comment

Presence of antinuclear antibodies (ANA) is a hallmark feature of systemic autoimmune rheumatic diseases (SARD). However, ANA lacks diagnostic specificity and is associated with a variety of diseases (cancers, autoimmune, infectious, and

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Patient Age/Sex:

Unknown

Test Information

i8: ANA Interpretive Comment

inflammatory conditions) and may also occur in healthy individuals in varying prevalence. The lack of diagnostic specificity requires confirmation of positive ANA by more specific serologic tests. ANA (nuclear reactivity) positive patterns reported include centromere, homogeneous, nuclear dots, nucleolar, or speckled. ANA (cytoplasmic reactivity) positive patterns reported include reticular/AMA, discrete/GW body-like, polar/golgi-like, cytoplasmic speckled or rods and rings. All positive patterns are reported to endpoint titers (1:2560). Reported patterns may help guide differential diagnosis, although they may not be specific for individual antibodies or diseases. Mitotic staining patterns not reported. Negative results do not necessarily rule out SARD.

i9: Cyclic Citrullinated Peptide Ab, IgG/A INTERPRETIVE INFORMATION: Cyclic Citrullinated Peptide Ab, IgG/A

19 Units or	less	Negative
20-39 Units		Weak Positive
40-59 Units		Moderate Positive
60 Units or	greater	Strong Positive

A positive result for cyclic citrullinated peptide (CCP) antibodies in conjunction with consistent clinical features may be suggestive of rheumatoid arthritis (RA). Anti-CCP, IgG/IgA antibodies are present in about 66-74 percent of RA patients and have specificities of 96-99 percent. Detection of IgA antibodies in addition to the usual IgG antibodies enhances the sensitivity due to some RA patients having IgA antibodies to CCP in the absence of IgG. These autoantibodies may be present in the preclinical phase of disease, are associated with future RA development, and may predict radiographic joint destruction. Patients with weak positive results should be monitored and testing repeated.

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