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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

Specimen Collected: 11/18/2024 07:45 MST

Patient Age/Sex:

Unknown

specimen corrected: 11/18/2024 07:4	5 MSI		
Extended Myositis Panel 2 F	Received: 11/18/20	24 07:46 MST	Report/Verified: 11/18/2024 07:57 MST
Procedure	Result	Units	Reference Interval
SSA-52 (Ro52) (ENA) Antibody, Ig0	G 65 ^{H i1}	AU/mL	[0-40]
SSA-60 (Ro60) (ENA) Antibody, Ig0	G 75 ^{H i2}	AU/mL	[0-40]
Smith/RNP (ENA) Ab, IgG	60 H i3	Units	[0-19]
Jo-1 (Histidyl-tRNA Synthetase)	55 ^{H i4}	AU/mL	[0-40]
Ab,IgG			
PL-12 (alanyl-tRNA synthetase)	Positive *		[Negative]
Antibody			
PL-7 (threonyl-tRNA synthetase)	Weak Positive *		[Negative]
Antibody			
EJ (glycyl-tRNA synthetase)	Positive *		[Negative]
Antibody			
OJ (isoleucyl-tRNA synthetase)	Positive *		[Negative]
Antibody			
SRP (Signal Recognition	Positive *		[Negative]
Particle) Ab	Desibility *		[Negotive]
Ku Antibody	Positive * Positive * ⁱ⁵		[Negative]
PM/Scl 100 Antibody,IgG Fibrillarin (U3 RNP) Ab,IgG		16	[Negative]
Mi-2 (nuclear helicase protein)	High Positive *	10	[Negative]
· · · ·	Positive		[Negative]
Antibody P155/140 Antibody	Positive *		[Negative]
TIF-1 gamma (155 kDa) Ab	Positive *		[Negative]
SAE1 (SUMO activating enzyme) Ak			[Negative]
MDA5 (CADM-140) Ab	High Positive *		[Negative]
NXP2 (Nuclear matrix protein-2)	_	L	[Negative]
Ab	LOW POSICIVE		[Negative]
Myositis Panel Interpretive Data	aSee Note ⁱ⁷		
Antinuclear Antibody (ANA), HEp-	Detected *		[<1:80]
2,IgG			
ANA Interpretive Comment	See Note ti i8		
Ha (tyrosyl-tRNA synthetase) Ab	Positive * t2		[Negative]
Ks (asparaginyl-tRNA synthetase)) Positive * t3		[Negative]
Ab			
Zo (phenylalanyl-tRNA synthetase) Ab	Positive * t4		[Negative]
Antinuclear Ab, Dual Pattern F	Received: 11/18/20	24 07:46 MST	Report/Verified: 11/18/2024 07:58 MST
Procedure	Result	Units	Reference Interval
ANA Titer 2	1:320 *		
ANA Pattern	Speckled *		

*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

Unless otherwise indicated, testing performed at: ARUP Laboratories 500 Chipeta Way, Salt Lake City, UT 84108 Laboratory Director: Jonathan R. Genzen, MD, PhD
 ARUP Accession:
 24-323-900011

 Report Request ID:
 20183802

 Printed:
 11/19/2024 12:56 MST

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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

Unknown

Patient Age/Sex:

Jonathan R. Genzen, MD, PhD, Chief Medical C	Ifficer			
Antinuclear Ab, Dual Pattern	Received: 11/18/20	24 07:46 MST	Report/Verified: MST	11/18/2024 07:58
Procedure ANA Titer ANA Pattern 2	Result 1:1280 * Centromere *	Units	Refer	ence Interval
Cytoplasmic Pattern	Received: 11/18/20	24 07:46 MST	Report/Verified: MST	11/18/2024 07:58
Procedure Cytoplasmic Titer Cytoplasm Pattern	Result 1:320 * Rods and Rings	Units		ence Interval
Speckled Pattern Clinical associations: healthy individuals Main autoantibodies: An anti-Topo-1 (anti-Scl- (TIF1g), anti-Ku, ant Centromere Pattern Clinical associations: Main autoantibodies: An	nti-SSA-52 (Ro52), 70), Smith, anti-U1 i-RNA polymerase, a SSc, PBC	anti-SSA-60 -RNP, anti-U nti-DFS70/LE	(Ro60), anti-SS- 2-RNP, anti-Mi-2	-B/LA,
Rods and Rings pattern Clinical Associations: pegylated interferon-a Main autoantibodies: II	commonly found in lpha/ribavirin comb	ination ther		created with
List of Abbreviations Antisynthetase syndrom myopathies (IM) [derma myopathy (NAM)], inter (JIA), mixed connective rheumatoid arthritis (I syndrome (SjS), system undifferentiated connec	tomyositis (DM), po stitial lung diseas e tissue disease (M RA), systemic autoi ic lupus erythemato ctive tissue diseas	lymyositis (e (ILD), juv CTD), primar mmune rheuma sus (SLE), s e (UCTD).	PM), necrotizing enile idiopathic y biliary cholar tic diseases (SA	g autoimmune c arthritis ngitis (PBC), ARD), Sjogren
 t2: 11/18/2024 07:45 MST (Ha (ty Ha positive by line immi immunoprecipitation. P: 11/18/2024 07:45 MST (Ks (as 	munoassay. Band co rofile consistent w	rresponding ith Ha antik		ved by
 11/18/2024 07.45 Mai (Rs (as Ks positive by line imminoprecipitation. Particular 11/18/2024 07:45 MST (Zo (ph) Zo positive by line imminoprecipitation. Particular 	munoassay. Band co rofile consistent w menylalanyl-tRNA syntheta munoassay. Bands co	rresponding ith Ks antik Ase) Ab) rresponding	ody positivity. to 68 and 58 KDa	

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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

<u>Result Footnote</u>

f1: NXP2 (Nuclear matrix protein-2) Ab

> Low positive reactivity to nuclear matrix protein (NXP2) detected. Strong clinical correlation is recommended.

Test Information

i1:

i3:

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.

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i2:
      SSA-60 (Ro60) (ENA) Antibody, IgG
      REFERENCE INTERVAL: SSA-60 (Ro60) (ENA) Antibody, IgG
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29 AU/mL or Less	Negative
30 - 40 AU/mL	Equivocal
41 AU/mL or Greater	Positive
Smith/RNP (ENA) Ab, IgG	
INTERPRETIVE INFORMATION: Smith/	RNP (ENA) Antibody, IgG

19 Units	or Less	Negative
20 to 39	Units	Weak Positive
40 to 80	Units	Moderate Positive
81 Units	or greater	Strong Positive

Smith/RNP antibodies are frequently seen in patients with mixed connective tissue disease (MCTD) and are also associated with other systemic autoimmune rheumatic diseases (SARDs) such as systemic lupus erythematosus (SLE), systemic sclerosis, and myositis. Antibodies targeting the Smith/RNP antigenic complex also recognize Smith antigens, therefore, the Smith antibody response must be considered when interpreting these results.

i4: 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

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Laboratory Director: Jonathan R. Genzen, MD, PhD

ARUP Accession: 24-323-900011 Report Request ID: 20183802 Printed: 11/19/2024 12:56 MST Page 3 of 6

phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex:

Unknown

Test Information

i4: Jo-1 (Histidyl-tRNA Synthetase) Ab, IgG 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).

i5: PM/Scl 100 Antibody, IgG INTERPRETIVE INFORMATION: PM/Scl-100 Antibody, IgG 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.

i6: Fibrillarin (U3 RNP) Ab, IgG Interpretive Information: Fibrillarin (U3 RNP) Antibody, IgG

The presence of fibrillarin (U3-RNP) IgG antibodies in association with an ANA IFA nucleolar pattern is suggestive of systemic sclerosis (SSc). In SSc, these antibodies are associated with distinct clinical features, such as younger age at disease onset, frequent internal organ involvement (pulmonary hypertension, myositis and renal disease). Fibrillarin antibodies are detected more frequently in African American patients with SSc compared to other ethnic groups. Strong correlation with ANA IFA results is recommended.

In a multi-ethnic cohort of SSc patients (n=98), U3-RNP antibodies detected by immunoblot had an agreement of 98.9 percent with the gold standard immunoprecipitation (IP) assay. Approximately 71 percent (5/7) of the borderline U3-RNP results with ANA nucleolar pattern in this cohort were IP negative.

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.

i7:

INTERPRETIVE INFORMATION: Extended Myositis Panel

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Laboratory Director: Jonathan R. Genzen, MD, PhD

Myositis Panel Interpretive Data

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phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex:

Unknown

Test Information

i7: Myositis Panel Interpretive Data

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

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.

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 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

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Patient Age/Sex:

Unknown

Test Information

i8: ANA Interpretive Comment

(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.

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