### Joel Corcoran

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860 STATUS: Final

Source: Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

Lab Ref #: 898850

ORDERING PHYSICIAN:

### Joshua A Emdur,

D.O.

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test		In Range	Out Of Range	Reference Range	Lab
GGT	Collected: 01/07/2025 06:16 PM UTC	Received: 01/07/2	025 06:19 PM UTC		
GGT		15		3-95 U/L	UL
LEPTIN	Collected: 01/07/2025 06:16 PM UT	C Received: 01/0	7/2025 06:19 PM UTC		
LEPTI	N	2.3		ng/mL	EZ

Reference Ranges for Leptin:

Adult Lean Subjects (18-71 years) with BMI range of 18-25:

Males: 0.3-13.4 ng/mLFemales: 4.7-23.7 ng/mL

Adult Subjects (19-60 years) with BMI range of 25-30:

Males: 1.8-19.9 ng/mL Females: 8.0-38.9 ng/mL

Pediatric Reference Ranges for Leptin:

5-9.9 years: 0.6-16.8 ng/mL 10-13.9 years: 1.4-16.5 ng/mL 14-17.9 years: 0.6-24.9 ng/mL

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

#### METHYLMALONIC ACID Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC

METHYLMALONIC ACID

222

55-335 nmol/L

ΕZ

Serum methylmalonic acid (MMA) levels are used to diagnose and monitor several rare inborn errors of metabolism, including methylmalonic aciduria. The enzymatic conversion of MMA to succinic acid requires vitamin B12 (adenosyl-cobalamin) as a cofactor. Serum MMA levels are also used for assessing functional vitamin B12 deficiency. Vitamin B12 is essential for fetal neurodevelopment, particularly early in pregnancy. Undiagnosed maternal vitamin B12 deficiency may be associated with adverse fetal/neonatal outcomes, such as neural tube defects and intrauterine growth restriction.



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### **Joel Corcoran**

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860 STATUS: Final

Source: Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

**Lab Ref #:** 898850

ORDERING PHYSICIAN:

# Joshua A Emdur,

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test In Range Out Of Range Reference Range Lab

Quest Diagnostics utilized Multi-Modal Decomposition (MMD) analysis to establish first and second trimester-specific MMA reference intervals in pregnancy, as given below:

MMA, First trimester (<13 wks gestation): 58-167 nmol/L MMA, Second trimester (13-23 wks gestation): 63-241 nmol/L

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

## TESTOSTERONE, FREE (DIALYSIS) AND TOTAL (MS) Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC

TESTOSTERONE, TOTAL, MS

465

250-1100 ng/dL

EZ

For additional information, please refer to http://education.questdiagnostics.com/faq/TotalTestosteroneL CMSMS (This link is being provided for informational/educational purposes only.)

TESTOSTERONE, FREE

75.8

35.0-155.0 pg/mL

EZ

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

## ANA SCREEN, IFA, W/REFL TITER AND PATTERN Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06: 19 PM UTC

ANA SCREEN, IFA POSITIVE A NEGATIVE EN

ANA IFA is a first line screen for detecting the presence of up to approximately 150 autoantibodies in various autoimmune diseases. A positive ANA IFA result is suggestive of autoimmune disease and reflexes to titer and pattern. Further laboratory testing may be considered if clinically indicated.

For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ177 (This link is being provided for informational/educational purposes only.)



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### **Joel Corcoran**

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860

06:19 PM UTC

STATUS: Final

Source: Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

Lab Ref #: 898850

ORDERING PHYSICIAN:

# Joshua A Emdur,

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test	In Ra	ange	Out Of Range	Reference Range	Lab
ANTINUCLEAR ANTIBODIES TITER ANI	PATTERN	Collected	l: 01/07/2025 06:16 PM U	TC Received: 01/07/2	2025

ANA TITER 1:80 H titer EN

A low level ANA titer may be present in pre-clinical autoimmune diseases and normal individuals.

Reference Range

<1:40 Negative

1:40-1:80 Low Antibody Level >1:80 Elevated Antibody Level

ANA PATTERN <u>Nuclear, Speckled</u> A

Speckled pattern is associated with mixed connective tissue disease (MCTD), systemic lupus erythematosus (SLE), Sjogren's syndrome, dermatomyositis, and systemic sclerosis/polymyositis overlap.

AC-2,4,5,29: Speckled

International Consensus on ANA Patterns
(https://doi.org/10.1515/cclm-2018-0052)

RHEUMATOID FACTOR	Collected: 01/07/2025 06:16 PM UTC	Received: 01/07/2025 0	6:19 PM UTC	
RHEUMATOID FACTOR		<u>25</u> H	<14 IU/mL	UL

THYROID PEROXIDASE AND THYROGLOBULIN ANTIBODIES Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC

THYROGLOBULIN ANTIBODIES <1 < or = 1 IU/mL EN THYROID PEROXIDASE <1 <9 IU/mL EN ANTIBODIES

HOMOCYSTEINE Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC

HOMOCYSTEINE 10.4 <11.4 umol/L <sup>UL</sup>

Homocysteine is increased by functional deficiency of folate or vitamin B12. Testing for methylmalonic acid differentiates between these deficiencies. Other causes of increased homocysteine include renal failure, folate antagonists such as methotrexate and phenytoin, and exposure to nitrous oxide.

Selhub J, et al., Ann Intern Med. 1999;131(5):331-9.

DHEA SULFATE Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC

DHEA SULFATE 201 61-442 mcg/dL EN

 SEX HORMONE BINDING GLOBULIN
 Collected: 01/07/2025 06:16 PM UTC
 Received: 01/07/2025 06:19 PM UTC

 SEX HORMONE BINDING
 28
 10-50 nmol/L

GLOBULIN



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ΕN

### Joel Corcoran

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860 STATUS: Final

Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

Source:

**Lab Ref #:** 898850

ORDERING PHYSICIAN:

# Joshua A Emdur,

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test		In Range	Out Of Range	Reference Range	Lab
FSH Collec	cted: 01/07/2025 06:16 PM UTC R	Received: 01/07/20	25 06:19 PM UTC		
FSH		6.6		1.4-12.8 mIU/mL	UL
LH Collect	ed: 01/07/2025 06:16 PM UTC Re	eceived: 01/07/202	5 06:19 PM UTC		
LH		5.0		1.5-9.3 mIU/mL	UL
PROLACTIN	Collected: 01/07/2025 06:16 PM	UTC Received:	01/07/2025 06:19 PM UT	C	
PROLACTI	N .	5.1		2.0-18.0 ng/mL	UL
ESTRADIOL	Collected: 01/07/2025 06:16 PM	UTC Received:	01/07/2025 06:19 PM UT	C	
ESTRADIO		28		< OR = 39 pg/mL	UL

Reference range established on post-pubertal patient population. No pre-pubertal reference range established using this assay. For any patients for whom low Estradiol levels are anticipated (e.g. males, pre-pubertal children and hypogonadal/post-menopausal females), the Quest Diagnostics Nichols Institute Estradiol, Ultrasensitive, LCMSMS assay is recommended (order code 30289).

Please note: patients being treated with the drug fulvestrant (Faslodex(R)) have demonstrated significant interference in immunoassay methods for estradiol measurement. The cross reactivity could lead to falsely elevated estradiol test results leading to an inappropriate clinical assessment of estrogen status. Quest Diagnostics order code 30289-Estradiol, Ultrasensitive LC/MS/MS demonstrates negligible cross reactivity with fulvestrant.

PSA (FREE AND TOTAL)	Collected: 01/07/2025 06:16 PM UTC	Received: 01/07/2025 06:19 PM UTC	
PSA, TOTAL	0.5	< OR = 4.0 ng/mL	EN
PSA, FREE	0.2	ng/mL	EN
PSA, % FREE	40	>25 % (calc)	EN
PSA(ng/mL)	Free PSA(%) Estimated(x	) Probability	

		of Cancer(as%)
0-2.5	( * )	Approx. 1
2.6-4.0(1)	0-27(2)	24(3)
4.1-10(4)	0-10	56
	11-15	28
	16-20	20
	21-25	16
	>or =26	8
>10(+)	N/A	>50

References:(1)Catalona et al.:Urology 60: 469-474 (2002)



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### **Joel Corcoran**

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860 STATUS: Final

Source: Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

**Lab Ref #:** 898850

ORDERING PHYSICIAN:

# Joshua A Emdur,

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test	In Range	Out Of Range	Reference Range	Lab
	(2)Catalona et al.:J.Urol 168: 922-	925 (2002)		
	Free PSA(%) Sensitivity(%) Sp	ecificity(%)		

- (x)These estimates vary with age, ethnicity, family history and DRE results.
- (\*) The diagnostic usefulness of % Free PSA has not been established in patients with total PSA below 2.6 ng/mL
- (+)In men with PSA above 10 ng/mL, prostate cancer risk is determined by total PSA alone.

The Total PSA value from this assay system is standardized against the equimolar PSA standard. The test result will be approximately 20% higher when compared to the WHO-standardized Total PSA (Siemens assay). Comparison of serial PSA results should be interpreted with this fact in mind.

PSA was performed using the Beckman Coulter Immunoassay method. Values obtained from different assay methods cannot be used interchangeably. PSA levels, regardless of value, should not be interpreted as absolute evidence of the presence or absence of disease.

AMYLAS	SE Collected: 01/07/2025 06:16 PM	UTC Received: 01/07/2025 06:19 PM UTC		
AMYLA	SE	39	21-101 U/L	UL
LIPASE	Collected: 01/07/2025 06:16 PM UT	C Received: 01/07/2025 06:19 PM UTC		
LIPASE		20	7-60 U/L	UL
ZINC C	Collected: 01/07/2025 06:16 PM UTC	Received: 01/07/2025 06:19 PM UTC		
ZINC		82	60-130 mcg/dL	EN

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

 LEAD (VENOUS)
 Collected: 01/07/2025 06:16 PM UTC
 Received: 01/07/2025 06:19 PM UTC

 LEAD (VENOUS)
 <1.0</td>
 <3.5 mcg/dL</td>



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### Joel Corcoran

Phone (H): (833) 753-1851 DOB: 11/30/1980 Gender: Male Age: 44 Patient ID: 53573860 STATUS: Final

Source: Quest

Time Reported: 01/15/2025 12:32 AM

UTC

**Received:** 01/15/2025 12:36 AM

UTC

Accession SZ624020R

Number:

**Lab Ref #:** 898850

ORDERING PHYSICIAN:

# Joshua A Emdur,

600 Congress Avenue

Floor 14

Austin, TX, 78701

Test In Range Out Of Range Reference Range Lab

See Note 1

Note 1

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

Enhanced PDF Report SZ624020R-1 Collected: 01/07/2025 06:16 PM UTC Received: 01/07/2025 06:19 PM UTC Enhanced PDF Report SZ624020R-Enhanced PDF Report SZ624020R-I.pdf [See Appendix 1 for details]

Enhanced PDF Report SZ624020R- Enhanced PDF Report SZ624020R-1.pdf [See Appendix 1 for details]

UL Quest Diagnostics-Sacramento - Northgate.

3714 Northgate Blvd, Sacramento, CA 95834-1617

EZ Quest Diagnostics/Nichols SJC-San Juan Capistrano,.

33608 Ortega Hwy, San Juan Capistrano, CA 92675-2042

EN Quest Diagnostics-West Hills.

8401 Fallbrook Ave, West Hills, CA 91304-3226

Dir: Lorne L Holland

Dir: Irina Maramica MD,PhD,MBA

Dir: Thomas J McDonald

#### Range Flags Legend: H - Above high normal; A - Abnormal;



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Report Status: Final CORCORAN, JOEL

EZ

UL

EZ

Patient Information	Specimen Information	Client Information
CORCORAN, JOEL  DOB: 11/30/1980 AGE: 44  Gender: M  Phone: 833.753.1851  Patient ID: 53573860  Health ID: 8573036010014334	Specimen:         SZ624020R           Requisition:         0067113           Lab Ref #:         898850           Collected:         01/07/2025 / 10:16 PST           Received:         01/07/2025 / 20:38 PST           Reported:         01/14/2025 / 16:32 PST	Client #: 73917267 MAIL992 EMDUR, JOSHUA FUNCTION HEALTH INC 600 CONGRESS AVE FL 14 AUSTIN, TX 78701-3263

Test Name In Range Out Of Range Reference Range Lab
TESTOSTERONE, FREE (DIALYSIS) AND TOTAL (MS)
TESTOSTERONE, TOTAL, MS 465 250-1100 ng/dL EZ

For additional information, please refer to http://education.questdiagnostics.com/faq/TotalTestosteroneL CMSMS (This link is being provided for informational/educational purposes only.)

TESTOSTERONE, FREE

(DIALYSIS)

TESTOSTERONE, FREE 75.8 35.0-155.0 pg/mL

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

HOMOCYSTEINE 10.4 <11.4 umol/L Homocysteine is increased by functional deficiency of folate or vitamin B12. Testing for methylmalonic acid differentiates between these deficiencies. Other causes of increased homocysteine include renal failure, folate antagonists such as methotrexate and phenytoin, and

exposure to nitrous oxide.

Selhub J, et al., Ann Intern Med. 1999;131(5):331-9.

3-95 U/L UL GGT 15 AMYLASE 39 21-101 U/L UL LIPASE 20 7-60 U/L UL THYROID PEROXIDASE AND THYROGLOBULIN ANTIBODIES THYROGLOBULIN ANTIBODIES < or = 1 IU/mLEN THYROID PEROXIDASE ΕN

SPECIMEN: SZ624020R

ANTIBODIES <1 <9 IU/mL LEPTIN 2.3 ng/mL

Reference Ranges for Leptin:

Adult Lean Subjects (18-71 years) with BMI range of 18-25:

Males: 0.3-13.4 ng/mLFemales: 4.7-23.7 ng/mL

Adult Subjects (19-60 years) with BMI range of 25-30:

Males: 1.8-19.9 ng/mL Females: 8.0-38.9 ng/mL

Pediatric Reference Ranges for Leptin:

5-9.9 years: 0.6-16.8 ng/mL 10-13.9 years: 1.4-16.5 ng/mL 14-17.9 years: 0.6-24.9 ng/mL





Report Status: Final CORCORAN, JOEL

Patient Information	Specimen Information	Client Information
CORCORAN, JOEL	Specimen: SZ624020R Collected: 01/07/2025 / 10:16 PST	Client #: 73917267 EMDUR, JOSHUA
DOB: 11/30/1980 AGE: 44 Gender: M Patient ID: 53573860 Health ID: 8573036010014334	Received: 01/07/2025 / 20:38 PST Reported: 01/14/2025 / 16:32 PST	

Test Name In Range Out Of Range Reference Range Lab

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

METHYLMALONIC ACID 222 55-335 nmol/L EZ

Serum methylmalonic acid (MMA) levels are used to diagnose and monitor several rare inborn errors of metabolism, including methylmalonic aciduria. The enzymatic conversion of MMA to succinic acid requires vitamin B12 (adenosyl-cobalamin) as a cofactor. Serum MMA levels are also used for assessing functional vitamin B12 deficiency. Vitamin B12 is essential for fetal neurodevelopment, particularly early in pregnancy. Undiagnosed maternal vitamin B12 deficiency may be associated with adverse fetal/neonatal outcomes, such as neural tube defects and intrauterine growth restriction.

Quest Diagnostics utilized Multi-Modal Decomposition (MMD) analysis to establish first and second trimester-specific MMA reference intervals in pregnancy, as given below:

MMA, First trimester (<13 wks gestation): 58-167 nmol/L MMA, Second trimester (13-23 wks gestation): 63-241 nmol/L

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

ror crimical parposes.				
RHEUMATOID FACTOR		25 H	<14 IU/mL	UL
DHEA SULFATE	201		61-442 mcg/dL	EN
SEX HORMONE BINDING				EN
GLOBULIN	28		10-50  nmol/L	
FSH	6.6		1.4-12.8  mIU/mL	UL
LH	5.0		1.5-9.3  mIU/mL	$\mathtt{UL}$
PROLACTIN	5.1		2.0-18.0  ng/mL	$\mathtt{UL}$
ESTRADIOL	28		< OR = 39 pg/mL	IJL

SPECIMEN: SZ624020R

Reference range established on post-pubertal patient population. No pre-pubertal reference range established using this assay. For any patients for whom low Estradiol levels are anticipated (e.g. males, pre-pubertal children and hypogonadal/post-menopausal females), the Quest Diagnostics Nichols Institute Estradiol, Ultrasensitive, LCMSMS assay is recommended (order code 30289).

Please note: patients being treated with the drug fulvestrant (Faslodex(R)) have demonstrated significant interference in immunoassay methods for estradiol measurement. The cross reactivity could lead to falsely elevated estradiol test results leading to an inappropriate clinical assessment of estrogen status. Quest Diagnostics order code 30289-Estradiol,





**Report Status: Final** CORCORAN, JOEL

Patient Information	Specimen Information	Client Information
CORCORAN, JOEL	Specimen: SZ624020R Collected: 01/07/2025 / 10:16 PST	Client #: 73917267 EMDUR, JOSHUA
DOB: 11/30/1980 AGE: 44 Gender: M Patient ID: 53573860 Health ID: 8573036010014334	Received: 01/07/2025 / 20:38 PST Reported: 01/14/2025 / 16:32 PST	

Test Name In Range Out Of Range Reference Range Lab Ultrasensitive LC/MS/MS demonstrates negligible cross reactivity with fulvestrant. PSA (FREE AND TOTAL) EN PSA, TOTAL 0.5 < OR = 4.0 ng/mLPSA, FREE 0.2 ng/mL PSA, % FREE >25 % (calc) PSA(ng/mL) Free PSA(%) Estimated(x) Probability of Cancer(as%) 0-2.5(\*) Approx. 1 2.6-4.0(1) 0-27(2) 24(3) 4.1-10(4)0 - 1056 28 11-15 16-20 20 21-25 16 >or =268 >10(+)N/A >50 References: (1) Catalona et al.: Urology 60: 469-474 (2002) (2)Catalona et al.: J. Urol 168: 922-925 (2002) Free PSA(%) Sensitivity(%) Specificity(%) < or = 2585 19 < or = 3093 (3)Catalona et al.: JAMA 277: 1452-1455 (1997)

(4)Catalona et al.: JAMA 279: 1542-1547 (1998)

- (x) These estimates vary with age, ethnicity, family history and DRE results.
- (\*) The diagnostic usefulness of % Free PSA has not been established in patients with total PSA below 2.6 ng/mL
- (+)In men with PSA above 10 ng/mL, prostate cancer risk is determined by total PSA alone.

The Total PSA value from this assay system is standardized against the equimolar PSA standard. The test result will be approximately 20% higher when compared to the WHO-standardized Total PSA (Siemens assay). Comparison of serial PSA results should be interpreted with this fact in mind.

PSA was performed using the Beckman Coulter Immunoassay method. Values obtained from different assay methods cannot be used interchangeably. PSA levels, regardless of value, should not be interpreted as absolute evidence of the presence or absence of disease.

LEAD (VENOUS) < 1.0 <3.5 mcg/dLΕN

See Endnote 1

ZINC 82 60-130 mcg/dL ΕN

SPECIMEN: SZ624020R

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA





Report Status: Final CORCORAN, JOEL

Patient Information	Specimen Information	Client Information
CORCORAN, JOEL	Specimen: SZ624020R Collected: 01/07/2025 / 10:16 PST	Client #: 73917267 EMDUR, JOSHUA
DOB: 11/30/1980 AGE: 44 Gender: M	Received: 01/07/2025 / 20:38 PST Reported: 01/14/2025 / 16:32 PST	
Patient ID: 53573860 Health ID: 8573036010014334		

Test Name In Range Out Of Range Reference Range Lab

regulations and is used for clinical purposes.

#### Endnote 1

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

CLIENT SERVICES: 866.697.8378 SPECIMEN: SZ624020R PAGE 4 OF 5





**Report Status: Final** CORCORAN, JOEL

Patient Information	Specimen Information	Client Information
CORCORAN, JOEL	Specimen: SZ624020R Collected: 01/07/2025 / 10:16 PST	Client #: 73917267 EMDUR, JOSHUA
DOB: 11/30/1980 AGE: 44 Gender: M Patient ID: 53573860 Health ID: 8573036010014334	Received: 01/07/2025 / 20:38 PST Reported: 01/14/2025 / 16:32 PST	

#### Immunology

Test Name	Result	Reference Range	Lab	
ANA SCREEN, IFA, W/REFL TITER AND PATTERN			EN	
ANA SCREEN, IFA	POSITIVE	NEGATIVE		
ANA IFA is a first line screen for detecting the presence of up to approximately 150 autoantibodies in various autoimmune diseases. A positive ANA IFA result is suggestive of autoimmune disease and reflexes to titer and pattern. Further laboratory testing may be considered if clinically indicated.				

For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ177 (This link is being provided for informational/ educational

purposes only.)

ANTINUCLEAR ANTIBODIES TITER AND PATTERN ΕN 1:80 H **ANA TITER** 

A low level ANA titer may be present in pre-clinical autoimmune diseases and normal individuals.

Reference Range

<1:40 Negative

1:40-1:80 Low Antibody Level >1:80 Elevated Antibody Level

**ANA PATTERN Nuclear, Speckled** 

Speckled pattern is associated with mixed connective tissue disease (MCTD), systemic lupus erythematosus (SLE), Sjogren's syndrome, dermatomyositis, and systemic sclerosis/polymyositis overlap.

AC-2,4,5,29: Speckled

International Consensus on ANA Patterns (https://doi.org/10.1515/cclm-2018-0052)

Physician Comments:

#### PERFORMING SITE:

- QUEST DIAGNOSTICS-WEST HILLS, 8401 FALLBROOK AVENUE, WEST HILLS, CA 91304-3226 Laboratory Director: THOMAS MCDONALD, MD, CLIA: 05D0642827 QUEST DIAGNOSTICS/NICHOLS SJC, 33608 ORTEGA HWY, SAN JUAN CAPISTRANO, CA 92675-2042 Laboratory Director: IRINA MARAMICA, MD, PHD, MBA, CLIA: 05D0643352 ΕZ
- QUEST DIAGNOSTICS SACRAMENTO, 3714 NORTHGATE BLVD, SACRAMENTO, CA 95834-1617 Laboratory Director: LORNE L. HOLLAND, MD, CLIA: 05D0644209

CLIENT SERVICES: 866.697.8378