

RIGLIETTI,ROBERT

DOB: 07/30/1995	Age: 29	Specimen: ZD710963E	Collected: 06/03/2025 15:00	Client #: 91436084
Sex: M	Fasting: Y	Requisition: 0000763	Received: 06/03/2025 15:10	SHILIAN,RYAN
Phone: (631) 793-2551		Lab Reference ID: 3965	Reported: 06/10/2025 14:08	ALLERGY IMMUNOLOGY CTR
Patient ID: 1109		Report Status: FINAL / SEE REPORT		STE 409
				16661 VENTURA BLVD
				ENCINO, CA 91436-1914
				Phone: (818) 514-3544
				Fax: (818) 578-0298

FASTING: YES ; COLLECTION KIT GIVEN TO PATIENT. PATIENT ADVISED TO RETURN.

**▲ HISTAMINE, PLASMA**

Analyte	Value
<b>▲ HISTAMINE, PLASMA</b>	<b>9.4 H</b> Reference Range: < OR = 1.8 ng/mL
<p>This test was performed using a kit that has not been cleared or approved by the FDA. The analytical performance characteristics of this test have been determined by Quest Diagnostics Nichols Institute San Juan Capistrano. This test should not be used for diagnosis without confirmation by other medically established means.</p>	

**▲ MANNAN BINDING LECTIN PATHWAY FUNCTION (C4B)**

Analyte	Value
<b>▲ MANNAN BINDING LECTIN PATHWAY FUNCTION (C4B)</b>	<b>&lt;52.0 L</b> Reference Range: >200 ng/mL
<p>The reference range was determined with sera from a population of known deficient patients. A value below 200 ng/mL indicates the patient is deficient in either MBL, MASP-2, or both. A normal healthy population has a lower 95% percentile cutoff of 676 ng/mL.</p> <p>*This test was developed and its performance characteristics determined by Eurofins Viracor. It has not been cleared or approved by the U.S. Food and Drug Administration.</p> <p>FLAG Interpretation: A = Abnormal, H = High, L = Low</p>	

**▲ PROSTAGLANDINS D2 (RANDOM URINE)**

Analyte	Value
<b>SPECIMEN TYPE/CONTAINER:</b>	<b>Urine/Cup</b>
<b>▲ PGD2 (RANDOM URINE)</b>	<b>260 H</b> Reference Range: <175 ng/g Creatinine
<p>This test has not been cleared or approved by the US Food and Drug Administration.</p> <p>This test was developed and its performance characteristics determined by Inter Science Institute. Values obtained with different methods, laboratories, or kits cannot be used interchangeably with the results on this report. The results cannot be interpreted as absolute evidence of the presence or absence of malignant disease.</p>	

**▲ RESPIRATORY ALLERGY PROFILE REGION XIII**

Analyte	Value
<b>DERMATOPHAGOIDES PTERONYSSINUS (D1) IGE</b>	<b>&lt;0.10</b> kU/L
<b>CLASS</b>	<b>0</b>

DERMATOPHAGOIDES FARINAE (D2) IGE	<0.10	kU/L
CLASS	0	
PENICILLIUM NOTATUM (M1) IGE	<0.10	kU/L
CLASS	0	
CLADOSPORIUM HERBARUM (M2) IGE	<0.10	kU/L
CLASS	0	
ASPERGILLUS FUMIGATUS (M3) IGE	<0.10	kU/L
CLASS	0	
ALTERNARIA ALTERNATA (M6) IGE	<0.10	kU/L
CLASS	0	
CAT DANDER (E1) IGE	<0.10	kU/L
CLASS	0	
DOG DANDER (E5) IGE	<0.10	kU/L
CLASS	0	
<b>▲ COCKROACH (I6) IGE</b>	<b>0.15 H</b>	kU/L
CLASS	<b>0/1</b>	
ALDER (T2) IGE	<0.10	kU/L
CLASS	0	
MOUNTAIN CEDAR (T6) IGE	<0.10	kU/L
CLASS	0	
OLIVE TREE (T9) IGE	<0.10	kU/L
CLASS	0	
WALNUT TREE (T10) IGE	<0.10	kU/L
CLASS	0	
COTTONWOOD (T14) IGE	<0.10	kU/L
CLASS	0	
OAK (T7) IGE	<0.10	kU/L
CLASS	0	
ELM (T8) IGE	<0.10	kU/L
CLASS	0	
WHITE MULBERRY (T70) IGE	<0.10	kU/L
CLASS	0	
BERMUDA GRASS (G2) IGE	<0.10	kU/L
CLASS	0	
TIMOTHY GRASS (G6) IGE	<0.10	kU/L
CLASS	0	
JOHNSON GRASS (G10) IGE	<0.10	kU/L
CLASS	0	

COMMON RAGWEED (SHORT) (W1) IGE	<0.10	kU/L
CLASS	0	
MUGWORT (W6) IGE	<0.10	kU/L
CLASS	0	
RUSSIAN THISTLE (W11) IGE	<0.10	kU/L
CLASS	0	
ROUGH PIGWEED (W14) IGE	<0.10	kU/L
CLASS	0	
MOUSE URINE PROTEINS (E72) IGE	<0.10	kU/L
CLASS	0	
IMMUNOGLOBULIN E	101	Reference Range: <OR=114 kU/L

## INTERPRETATION

Analyte	Value
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### INTERPRETATION

Specific IGE Class	kU/L	Level of Allergen Specific IGE Antibody
0	<0.10	Absent/Undetectable
0/1	0.10-0.34	Very Low Level
1	0.35-0.69	Low Level
2	0.70-3.49	Moderate Level
3	3.50-17.4	High Level
4	17.5-49.9	Very High Level
5	50-100	Very High Level
6	>100	Very High Level

The clinical relevance of allergen results of 0.10-0.34 kU/L are undetermined and intended for specialist use.

Allergens denoted with a "\*\*\*" include results using one or more analyte specific reagents. In those cases, the test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the U.S. Food and Drug Administration. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

## N METHYLHISTAMINE, RANDOM, U

Analyte	Value
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**N METHYLHISTAMINE, RANDOM, U** 95 Reference Range: 30-200 mcg/g Cr

-----ADDITIONAL INFORMATION-----  
 This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

**CREATININE, RANDOM, U** 100 Reference Range: 16 - 326 mg/dL

## TRYPTASE

Analyte	Value
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**TRYPTASE****6.1** Reference Range: <11.0 mcg/L

The Tryptase test, fluorescent enzyme immunoassay (FEIA), measures both the Alpha and Beta forms of Tryptase. Measuring both forms of Tryptase increases sensitivity for the diagnosis of mastocytosis, and mast cell degranulation as a cause of anaphylaxis.

**IMMUNOGLOBULINS**

Analyte	Value	
IMMUNOGLOBULIN A	<b>297</b>	Reference Range: 47-310 mg/dL
IMMUNOGLOBULIN G	<b>1055</b>	Reference Range: 600-1640 mg/dL
IMMUNOGLOBULIN M	<b>112</b>	Reference Range: 50-300 mg/dL

**STREPTOCOCCUS PNEUMONIAE AB (IGG) (23 SEROTYPES)**

Analyte	Value	
SEROTYPE 1 (1)	<b>0.6</b>	
SEROTYPE 2 (2)	<b>5.5</b>	
SEROTYPE 3 (3)	<b>0.6</b>	
SEROTYPE 4 (4)	<b>&lt;0.3</b>	
SEROTYPE 5 (5)	<b>2.6</b>	
SEROTYPE 8 (8)	<b>6.1</b>	
SEROTYPE 9 (9N)	<b>2.8</b>	
SEROTYPE 12 (12F)	<b>&lt;0.3</b>	
SEROTYPE 14 (14)	<b>&lt;0.3</b>	
SEROTYPE 17 (17F)	<b>0.9</b>	
SEROTYPE 19 (19F)	<b>1.8</b>	
SEROTYPE 20 (20)	<b>3.0</b>	
SEROTYPE 22 (22F)	<b>2.4</b>	
SEROTYPE 23 (23F)	<b>&lt;0.3</b>	
SEROTYPE 26 (6B)	<b>&lt;0.3</b>	
SEROTYPE 34 (10A)	<b>&lt;0.3</b>	
SEROTYPE 43 (11A)	<b>1.8</b>	
SEROTYPE 51 (7F)	<b>1.0</b>	
SEROTYPE 54 (15B)	<b>0.9</b>	
SEROTYPE 56 (18C)	<b>0.3</b>	
SEROTYPE 57 (19A)	<b>5.9</b>	
SEROTYPE 68 (9V)	<b>2.9</b>	

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**SEROTYPE 70 (33F)**

0.5

Serologic correlates of protection against pneumococcal disease have not been rigorously established for all patient populations. Published data and expert consensus (including WHO) suggest protection from invasive disease usually occurs at levels  $\geq 0.3$ - $0.50$  mcg/mL for healthy children receiving pneumococcal conjugate vaccines. Higher titers may be necessary to protect from non-invasive infection (e.g., pneumonia, otitis, sinusitis). Expert opinion suggests that a cut-off of  $\geq 1.3$  mcg/mL may be a more relevant value to assess antibody responses after pneumococcal polysaccharide vaccines or for immunocompromised patients. In addition to antibody quantity, protection also depends on antibody avidity and opsonophagocytic activity. Some experts consider that post-vaccination (4-6 weeks) IgG seroconversion and/or 2- to 4-fold rise in IgG titers for  $>50\%$  to  $70\%$  of vaccine serotypes demonstrates a normal post-vaccine serologic response. Persons with high initial serotype-specific titers may have less robust responses.

Quest Diagnostics uses a multi-analyte immunodetection (MAID) method. The method employs the Luminex flow cytometric system which measures multiple analytes simultaneously. The FDA standard reference serum 89-S is used as the calibration standard. Results are reported in mcg/mL.

This assay detects all of the 23 of the serotypes in the 23-valent polysaccharide vaccine and 12 of the 13 serotypes in the 13-valent conjugate vaccine.

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 used for clinical purposes.


For additional information, please refer to <http://education.questdiagnostics.com/faq/FAQ181> (This link is being provided for informational/educational purposes only.)

**Performing Sites**

ACER Eurofins Viracor, 18000 W 99Th Street, Lenexa, KS 66219-1233 Laboratory Director: Brock Neil PhD BCLD (ABB)  
EN Quest Diagnostics-West Hills, 8401 Fallbrook Ave, West Hills, CA 91304-3226 Laboratory Director: Thomas J McDonald  
EO Mayo Clinic Laboratories, 200 First Street SW, Rochester, MN 55905-1770 Laboratory Director: Nikola A Baumann M.D. Ph.D.  
EZ Quest Diagnostics/Nichols SJC-San Juan Capistrano,, 33608 Ortega Hwy, San Juan Capistrano, CA 92675-2042 Laboratory Director: Irina Maramica MD,PhD,MBA  
INS INTERSCIENCE INSTITUTE, 944 W Hyde Park Blvd, Inglewood, CA 90302-3308 Laboratory Director: James Lee MD,PhD  
MYM Mayo Clinic Laboratories, 3050 Superior Dr Nw, Rochester, MN 55905-1770 Laboratory Director: Nikola A Baumann M.D. Ph.D.

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**Key**

 Priority Out of Range  Out of Range

These results have been sent to the person who ordered the tests. Your receipt of these results should not be viewed as medical advice and is not meant to replace discussion with your doctor or other healthcare professional.

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