Warde Medical Laboratory

LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988 34 Y

Referral Testing

Collected: 09/01/2023 11:36 Received: 09/01/2023 11:36

Test Name	Result	Flag	Ref-Ranges	Units	Site
Toxoplasmosis Pregnancy Panel =<16 weeks gestation	See Below				WMPA

Test	Result	Interpretation
Sample Type	Serum	
Sample Date	08/25/2023	
IgG (Dye Test)	1:512	Positive *
IgM ELISA	0.8	Negative
IgG Avidity	High	

*Denotes result outside of reference range.

Reference Ranges and Comments

If you have questions regarding these results, please telephone our consulting physicians at 650-853-4828.

The Sabin-Feldman Dye Test: measures primarily IgG antibodies. Any titer is considered positive. Serum is tested at a 1:16 dilution unless we are notified that patient has eye disease, in which case serum is tested undiluted.

IgM-ELISA: (serum) negative 0.0-1.6, equivocal 1.7-1.9, >=2.0 positive; (CSF) negative 0.0-0.3, positive >=0.4. A negative IgM-ELISA result in an immunologically normal adult almost always excludes recent infection. IgM antibodies may persist for one year or longer following acute infection.

IgG Avidity: A high result in the first 16 weeks of gestation essentially excludes acute infection having been acquired during gestation. A low or equivocal IgG avidity result cannot be interpreted to mean that the patient has had a recently acquired infection since low avidity antibodies may persist for more than five months. IgG avidity results should only be interpreted with results of other tests in the Toxoplasma Serologic Panel. Interpretation can be altered if the patient has previously received anti-toxoplasma therapy, since this will affect IgG maturation kinetics during infection.

IgM-ELISA, IgM-ISAGA, IgA-ELISA, IgE, PCR: These tests

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988 34 Y

Referral Testing

Collected: 09/01/2023 11:36 Received: 09/01/2023 11:36

Test Name	Result	Flag	Ref-Ranges	Units	Site

were developed and their performance characteristic determined by the Toxoplasma Serology Laboratory, Palo Alto Medical Foundation. They have not been cleared or approved by the US Food and Drug Administration (FDA). However, approval by the FDA is not required for use of these tests by our reference laboratory.

Test(s) performed at
PALO ALTO MEDICAL FOUNDATION
795 El Camino Real/Attn: Ames Bldg.
Palo Alto, CA 94301

Reported Date: 2023.09.01 11:36

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED

F301000010
WX0000003826
Printed D&T: 09/01/23 11:36

Ordered By: KAJAL SITWALA, MD, PhD
WX00000000002353

Kajal V. Sitwala, MD, PhD - Medical Director
Form: MM RL1