

LABORATORY REPORT

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108 **EXAMPLE, REPORT W**

WX0000003827 M 07/08/1978 45 Y

Referral Testing

Collected: 09/22/2023 15:25 Received: 09/22/2023 15:25

Test Name Result Flag Ref-Ranges Units Site

Cutaneous Immunofl Ab IgG, Serum

Basement Membrane IgGNegativeNegativeMMRLCell Surface Ab IgGNegativeNegativeMMRLOtherSee CommentMMRL

The absence of serum cell surface and basement membrane zone IgG antibodies by cutaneous immunofluorescence testing (test code CIFS) does not support a diagnosis of an autoimmune mucocutaneous blistering disorder. However, this test may occasionally be negative in patients with these disorders and so the final definitive diagnosis should not be based solely on the results of serum cutaneous immunofluorescence testing (test code CIFS). If the clinical suspicion for an autoimmune mucocutaneous blistering disorder is high, other tests including (a) lesional skin or mucous membrane biopsy for histopathological evaluation on sections from formalin-fixed, paraffin-embedded tissue; (b) perilesional skin or mucous membrane biopsy for cutaneous immunofluorescence testing (test code CIB) on frozen section; and (c) serum testing for desmogleins 1 and 3 (test code DSGAB) and/or bullous pemphigoid BP180 and BP230 (test code BPAB) antibodies by ELISA technique are recommended.

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.

Test Performed by:
Mayo Clinic Laboratories - Rochester Main Campus
200 First Street SW, Rochester, MN 55905
Lab Director: William G. Morice M.D. Ph.D.; CLIA# 24D0404292

Human Split Skin IgGNegativeNegativeMmRLMonkey Esophagus IgGNegativeNegativeMMRL

Performing Site:

MMRL: MAYO MEDICAL REFERENCE LAB 3050 Superior Drive NW Rochester MN 55901

Reported Date: 2023.09.22 15:25 CIABG

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

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