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PATIENT REPORT

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

Specimen Collected: 13-Jun-23 06:10

DPPX IgG Ab CBA, Serum, with Rflx | Received: 13-Jun-23 06:18 Report/Verified: 13-Jun-23 06:49
Procedure Result Units Reference Interval

DPPX Ab IgG CBA-IFA Screen, Serum Detected * t1 i1 [<1:10]

DPPX Ab IgG Titer by CBA-IFA, Ser | Received: 13-Jun-23 06:18 Report/Verified: 13-Jun-23 06:49 Procedure Result Units Reference Interval

DPPX Ab IgG CBA-IFA Titer, Serum 1:640 * i2 [<1:10]

Interpretive Text

t1: 13-Jun-23 06:10 (DPPX Ab IgG CBA-IFA Screen, Serum)
DPPX Antibody, IgG is detected. Titer results to follow.

<u>Test Information</u>

i1: DPPX Ab IgG CBA-IFA Screen, Serum INTERPRETIVE INFORMATION: DPPX Ab IgG CBA IFA Screen, Serum

DPPX antibody is found in a subset of patients with autoimmune encephalitis, and is often associated with prodromal gastrointestinal symptoms and unintentional weight loss. It may occur with or without associated tumor. Decreasing antibody levels may be associated with therapeutic response. A negative test result does not rule out a diagnosis of autoimmune neurologic disease. Results should be interpreted in correlation with the patient's clinical history and other laboratory findings.

This indirect fluorescent antibody assay utilizes DPPX transfected cells for the detection and semiquantification of the DPPX IgG antibody.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

i2: DPPX Ab IgG CBA-IFA Titer, Serum
 INTERPRETIVE INFORMATION: DPPX Ab IgG CBA IFA Titer, Serum

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.

*=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

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