

Specimen Collected: 10-Dec-21 06:48

Procedure	Result	Units	Reference Interval
DPPX IgG Ab, CSF, with Rflx Received: 10-Dec-21 06:48 Report/Verified: 10-Dec-21 06:55			
DPPX Ab IgG CBA IFA Screen, CSF	Detected * t1 i1		< 1:1
DPPX IgG Ab Titer, CSF Received: 10-Dec-21 06:48 Report/Verified: 10-Dec-21 06:55			
DPPX Ab IgG CBA IFA Titer, CSF	1:40 * i2		< 1:1

Interpretive Text

t1: 10-Dec-21 06:48 (DPPX Ab IgG CBA IFA Screen, CSF)
DPPX Antibody, IgG is detected. Titer results to follow.

Test Information

i1: DPPX Ab IgG CBA IFA Screen, CSF
INTERPRETIVE INFORMATION: DPPX IgG Ab, CSF, with Rflx

Anti-DPPX IgG antibody is found in a subset of patients with autoimmune encephalitis and may occur with or without associated tumor. Decreasing antibody levels may be associated with therapeutic response; therefore, clinical correlation must be strongly considered. A negative test result does not rule out a diagnosis of autoimmune limbic encephalitis.

This indirect fluorescent antibody cell-based assay (CBA) utilizes dipeptidyl aminopeptidase-like protein 6 (DPPX) transfected cells for the detection 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 US 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, CSF
INTERPRETIVE INFORMATION: DPPX IgG Ab Titer, CSF

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: Tracy I. George, MD

ARUP Accession: 21-344-900017

Report Request ID: 15067588

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