

TEST CHANGE

Paraneoplastic Reflexive Panel, CSF

3004517, PNSPAN CSF

Specimen Requirements:

Patient Preparation:

Collect: CSF.

Specimen Preparation: Transfer 2 mL CSF to an ARUP standard transport tube. (Min: 1

mL).

Transport Temperature: Refrigerated

Unacceptable Conditions: Contaminated, heat-inactivated, hemolyzed, or lipemic

specimens

Remarks:

Stability: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month

Methodology: Semi-Quantitative Indirect Fluorescent Antibody

(IFA)/Qualitative Immunoblot/Semi-Quantitative Cell-Based

Effective Date: April 21, 2025

Indirect Fluorescent Antibody

Performed: Wed

Reported: 1-9 days

Note: Purkinje Cell (PCCA) antibody and neuronal nuclear Neuronal

Nuclear (ANNA) antibody IgG are screened by IFA. If the IFA screen is indeterminate, then a Neuronal Nuclear Antibodies (Hu, Ri, Yo, Tr/DNER) IgG by Immunoblot will be added. If the IFA screen is positive at 1:1 or greater, then a PCCA/ANNA antibodies titer and Neuronal Nuclear Antibodies (Hu, Ri, Yo, Tr/DNER) IgG by Immunoblot will be added. Additional charges apply. If CV2 Antibody IgG Screen by IFA is positive, then CV2 Antibody IgG Titer by IFA will be added. Additional charges

apply.

CPT Codes: 86255 x2; 84182 x3x2; if reflexed, add 86256 and/or-84182 x4;

if reflexed add 86256 per titer

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

Refer to report



Reference Interval:

Test Number	Components	Reference Interval
	Paraneoplastic Abs (PCCA/ANNA) IgG, CSF	None Detected
	Ma2/Ta Antibody, IgG by Immunoblot, CSF	<u>Negative</u>
	CV2 Ab IgG CBA-IFA Screen, CSF	Less than 1:1
	SOX1 Antibody, IgG by Immunoblot, CSF	Negative
	Amphiphysin Antibody, CSF	Negative

Effective Date: April 21, 2025

HOTLINE NOTE: There is a component change associated with this test. One or more components have been added or removed. Refer to the Hotline Test Mix for interface build information.