

## **TEST CHANGE**

Paraneoplastic Reflexive Panel, CSF 3004517, PNSPAN CSF		
Specimen Requirements:		
Patient Preparation:		
Collect:	CSF.	
Specimen Preparation:	Transfer 2 mL CSF to an ARUP <u>standard transport</u> <u>tube.</u> 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 <u>(IFA)/</u> 4Qualitative Immunoblot <u>/Semi-Quantitative Cell-Based</u> Indirect Fluorescent Antibody	
Performed:	Wed	
Reported:	1-9 days	
Note:	Purkinje Cell (PCCA) antibody and 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 CV2.1 Antibody IgG Screen by IFA is positive, then CV2.1 Antibody IgG Titer by IFA will be added. Additional charges apply.	
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CPT Codes:	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.	
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CPT Codes: New York DOH Approval Status:	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 CV2.1 Antibody IgG Screen by IFA is positive, then CV2.1 Antibody IgG Titer by IFA will be added. Additional charges apply.	



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.

Reference Interval:

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

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