



Dementia, Autoimmune/Paraneoplastic Evaluation, Serum

Patient ID	Patient Name		Birth Date	Sex	Age
321	IMPLEMENTATION, TEST T		1973-05-01	F	50
Order Number X100457662	Client Order Number X100457662	Ordering Physician TESTING	Report Notes		
Account Information C7028846 DLMP Roches	ter	Collected 08 Mar 2024 05:00			

Dementia, Autoimm/Paraneo, S

Dementia, Interpretation, S

MCR

No informative autoantibodies were detected in this evaluation. However, a negative result does not exclude autoimmune dementia, idiopathic or paraneoplastic. Sensitivity and specificity of antibody testing are enhanced by testing both serum and CSF.

IFA Notes

None.

Result Name	Result	Unit	Reference Value	Performing Site
AMPA-R Ab CBA, S	Negative		Negative	1 MCR
Amphiphysin Ab, S	Negative		Negative	1 MCR
AGNA-1, S	Negative		Negative	1 MCR
ANNA-1, S	Negative		Negative	1 MCR
ANNA-2, S	Negative		Negative	1 MCR
ANNA-3, S	Negative		Negative	1 MCR
CASPR2-IgG CBA, S	Negative		Negative	1 MCR
CRMP-5-IgG, S	Negative	10	Negative	1 MCR
DPPX Ab CBA, S	Negative		Negative	1 MCR
GABA-B-R Ab CBA, S	Negative		Negative	1 MCR
GAD65 Ab Assay, S	0.00	nmol/L	≤ 0.02	1 MCR
GFAP IFA, S	Negative		Negative	1 MCR
IgLON5 CBA, S	Negative		Negative	1 MCR
LGI1-IgG CBA, S	Negative		Negative	1 MCR
mGluR1 Ab IFA, S	Negative		Negative	1 MCR
Neurochondrin IFA, S	Negative		Negative	1 MCR
NIF IFA, S	Negative		Negative	1 MCR
NMDA-R Ab CBA, S	Negative		Negative	1 MCR
PCA-2, S	Negative		Negative	1 MCR
PCA-Tr, S	Negative		Negative	1 MCR
PDE10A Ab IFA, S	Negative		Negative	1 MCR
TRIM46 Ab IFA, S	Negative		Negative	1 MCR

Received: 12 Mar 2024 15:03 Reported: 12 Mar 2024 15:18

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
MCR	Mayo Clinic Laboratories - Rochester Main Campus	200 First Street SW, Rochester, MN 55905	William G. Morice M.D. Ph.D	24D0404292



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X100457662	X100457662	TESTING			
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C7028846 DLMP Rochester		08 Mar 2024 05:00			



Laboratory Notes

1 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.

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