

Client: Example Client ABC123 123 Test Drive Salt Lake City, UT 84108 UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB	8/1/1976
Gender:	Male
Patient Identifiers:	01234567890ABCD, 012345
Visit Number (FIN):	01234567890ABCD
Collection Date:	00/00/0000 00:00

West Nile Virus Antibodies, IgG and IgM by ELISA, CSF

ARUP test code 0050228

West Nile Virus Antibody IgG CSF	2.83 IV H	(Ref Interval: <=1.29)			
	The CSF specimen shows evidence of blood contamination and is, therefore, likely contaminated with serum antibodies. Antibody testing from this specimen is not recommended as the blood may interfere, causing a false-positive result that does not represent intrathecally produced antibodies. False-negative results are also possible. It is highly recommended that serum testing for the same analyte also be performed in order to aid in interpreting the CSF test result.				
	INTERPRETIVE INFORMA	TION: West Nile Virus Ab IgG by ELISA, CSF			
	1.30 - 1.49 IV	 Negative: No significant level of West Nile virus IgG antibody detected. Equivocal: Questionable presence of West Nile virus IgG antibody detected. Repeat testing in 10-14 days may be helpful. Positive: Presence of IgG antibody to West Nile virus detected, suggestive of current or past infection. 			
	detecting West Nile there is a clinical test should not be u should the results b history or other dat family, such as St. cross-reactivity wit	d to be used as a semi-quantitative means of virus-specific IgG in CSF samples in which suspicion of West Nile Virus infection. This sed solely for quantitative purposes, nor e used without correlation to clinical a. Because other members of the Flaviviridae Louis encephalitis virus, show extensive h West Nile virus, serologic testing pecies should be considered.			
	fluid may indicate c consideration must b	ibodies to West Nile virus in cerebrospinal entral nervous system infection. However, e given to possible contamination by blood antibodies across the blood-brain barrier.			
	determined by ARUP L approved by the US F	ped and its performance characteristics aboratories. It has not been cleared or ood and Drug Administration. This test was certified laboratory and is intended for			
West Nile Virus Antibody IgM CSF	1.71 IV H	(Ref Interval: <=0.89)			

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:



Specimen is repeatedly POSITIVE for anti-West Nile virus, IgM using the Focus Diagnostics ELISA assay. A false positive rate of 2-3% has been demonstrated with the Focus Diagnostics ELISA assay.

Repeated and verified.

The CSF specimen shows evidence of blood contamination and is, therefore, likely contaminated with serum antibodies. Antibody testing from this specimen is not recommended as the blood may interfere, causing a false-positive result that does not represent intrathecally produced antibodies. False-negative results are also possible. It is highly recommended that serum testing for the same analyte also be performed in order to aid in interpreting the CSF test result.

INTERPRETIVE INFORMATION: West Nile Virus Ab IgM by ELISA, CSF

0.89 IV or less	Negative - No significant level of West Nile virus IgM antibody
	of West Nile virus IgM antibody detected.
	Equivocal - Questionable presence
	of West Nile virus IgM antibody
	detected. Repeat testing in
	10-14 days may be helpful.
1.11 IV or greater	Positive - Presence of IqM
5	antibody to West Nile virus
	detected, suggestive of current
	or recent infection.
	or recent infection.

This test is intended to be used as a semi-quantitative means of detecting West Nile virus-specific IgM in CSF samples in which there is a clinical suspicion of West Nile virus infection. This test should not be used solely for quantitative purposes, nor should the results be used without correlation to clinical history or other data. Because other members of the Flaviviridae family, such as St. Louis encephalitis virus, show extensive cross-reactivity with West Nile virus, serologic testing specific for these species should be considered.

The detection of antibodies to West Nile virus in cerebrospinal fluid may indicate central nervous system infection. However, consideration must be given to possible contamination by blood or transfer of serum antibodies across the blood-brain barrier.

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.

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

ARUP LABORATORIES | 800-522-2787 | aruplab.com 500 Chipeta Way, Salt Lake City, UT 84108-1221 Jonathan R. Genzen, MD, PhD, Laboratory Director Patient: Patient, Example ARUP Accession: 23-361-137136 Patient Identifiers: 01234567890ABCD, 012345 Visit Number (FIN): 01234567890ABCD Page 2 of 3 | Printed: 2/2/2024 11:34:36 AM 4848



VERIFIED/REPORTED DATES						
Procedure	Accession	Collected	Received	Verified/Reported		
West Nile Virus Antibody IgG CSF	23-361-137136	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00		
West Nile Virus Antibody IgM CSF	23-361-137136	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00		

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

ARUP LABORATORIES | 800-522-2787 | aruplab.com 500 Chipeta Way, Sati Lake City, UT 84108-1221 Jonathan R. Genzen, MD, PhD, Laboratory Director Patient: Patient, Example ARUP Accession: 23-361-137136 Patient Identifiers: 01234567890ABCD, 012345 Visit Number (FIN): 01234567890ABCD Page 3 of 3 | Printed: 2/2/2024 11:34:36 AM 4848