

Patient ID SA00128831	Patient Name TESTRNV, IMPLEMENTATION	Birth Date 1986-11-13	Gender M	Age 33
Order Number SA00128831	Client Order Number SA00128831	Ordering Physician CLIENT,CLIENT	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 01 May 2020 00:00		

FVIII Inhib Profile Prof Interp

Reviewed by

KATHLEEN MEYERS

MCR

FVIII Inhib Profile Prof Interp

MCR

IMPRESSION: Positive. Assays confirm the presence of a specific inhibitor of factor VIII (FVIII) of 0.6 Bethesda Units.

COMMENTS: Decreased FVIII coagulant activity. Result of qualitative FVIII inhibitor screening assay is consistent with presence of an inhibitor. Reflexive titrating assay (Bethesda), performed after pre-analytical heat inactivation of patient plasma, confirms the presence of a specific inhibitor of coagulation FVIII of 0.6 Bethesda Units.

Received: 01 May 2020 11:33

Reported: 01 May 2020 11:38

FVIII Bethesda Units, P

Result Name	Value	Unit	Reference Value	Performing Site
 FVIII Bethesda Units, P	0.6	BU	≤0.5	1 MCR

Received: 01 May 2020 11:33


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Factor VIII Inhib Profile, P

FVIII Inhib Profile Tech Interp

See FVIII Inhib Profile Prof Interp.

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Result Name	Value	Unit	Reference Value	Performing Site
 Coag Factor VIII Activity Assay, P	54	%	55–200	1 MCR

Received: 01 May 2020 11:32

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
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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Coag Factor VIII Assay Inhib Scrn,P

Result Name	Value	Unit	Reference Value	Performing Site
  Coag Factor VIII Assay Inhib Scrn,P	Positive		Negative	1 MCR

Received: 01 May 2020 11:33

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Test Environment
EVP Template

Laboratory Notes

- 1** This test has been modified from the manufacturer's instructions. Its performance characteristics were 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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