

von Willebrand Disease Profile, Plasma

Patient ID 321	Patient Name TESTRNV, IMPLEMENTATION		Birth Date 1970-11-13	Gender M	Age 48	
Order Number X100288762	Client Order Number X100288762	Ordering Physician TESTING	Report Notes	Report Notes		
Account Information C7028846 DLMP Rochester		Collected 23 May 2019 08:00				

von Willebrand Disease Prof

von Willebrand Disease Tech Interp

MCF

IMPRESSION: No laboratory evidence of von Willebrand disease (VWD). COMMENTS: Normal or elevated factor VIII coagulant activity and/or von Willebrand factor (WWF) antigen and/or VWF activity [latex immunoassay] provide no evidence for von Willebrand disease (VWD). NOTE: VWF antigen and/or VWF latex immunoassay activity and/or factor VIII may be increased above baseline levels by acute or chronic inflammation, stress or adrenergic stimuli, pregnancy or estrogen and oral contraceptive (OCP) therapy, liver disease or recent infustion of plasma, cryoprecipitate, desmopressin (DDAVP) or VWF concentrates and mask the diagnosis of mild von Willebrand disease (VWD).

Desuit Name	Value	1144	Deference Value	Performing Site	
Result Name	Value	Unit	Reference Value		
Coag Factor VIII Activity Assay, P	55	%	55–200	1 MCR	
von Willebrand Factor Ag, P	55	%	55–200	1 MCR	
von Willebrand Factor Activity, P	55	%	55–200	1 MCR	

Received: 18 Jul 2019 11:19

Reported: 18 Jul 2019 11:20

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.

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