

<b>Patient Name</b> TESTINGRNV,RUO	<b>Patient ID</b> SA00062540	<b>Age</b> 45	<b>Gender</b> M	<b>Order #</b> SA00062540
<b>Ordering Phys</b> CLIENT,CLIENT				<b>DOB</b> 10/21/1967
<b>Client Order #</b> SA00062540	<b>Account Information</b>			<b>Report Notes</b>
<b>Collected</b> 09/18/2013 00:00	C7028846-DLMP Rochester SDSC 2 - Client Support Rochester, MN 55901			
<b>Printed</b> 09/26/2013 14:02				

Test	Flag	Results	Unit	Reference Value	Perform Site*
<b>ADAMTS13 Activity and Inhibitor Profile</b>					
<b>RECEIVED:</b> 09/19/2013 15:07 <b>REPORTED:</b> 09/19/2013 15:31					
ADAMTS13 Activity Assay	L	<5	%	>/=70	MCR
ADAMTS13 Interpretation					
Severely reduced ADAMTS13 activity can be observed in congenital/hereditary or acquired thrombotic thrombocytopenic purpura (TTP), or occasionally in other conditions including hemolytic uremic syndrome (HUS), hematopoietic stem cell and solid organ transplantation, liver disease, sepsis, DIC, pregnancy, or effects of certain medications (e.g., ticlopidine, clopidogrel, cyclosporin, mitomycin C, quinine, etc.). The ADAMTS13 inhibitor screening assay is consistent with presence of a neutralizing antibody against ADAMTS13, and the inhibitor titering assay (Bethesda assay) confirms presence of an inhibitor. Altogether these results are consistent with congenital/hereditary TTP or acquired TTP with a neutralizing antibody. Recommend clinical correlation and consider future repeat testing to follow/verify these findings if clinically indicated.					
Non-specific substrate proteolysis by other plasma proteases or recent plasma transfusion or exchange may falsely raise ADAMTS13 activity. Markedly elevated endogenous von Willebrand factor (VWF), hyperlipidemia, hemolysis with plasma free hemoglobin greater than 2mg/dL, hyperbilirubinemia (greater than 6mg/dL) may falsely lower ADAMTS13 activity					
ADAMTS13 Inhibitor Bethesda Titer	H	3.0		<0.4	MCR
<b>RECEIVED:</b> 09/19/2013 15:24 <b>REPORTED:</b> 09/19/2013 15:30					
ADAMTS13 Inhibitor Screen	H	Positive		Negative	MCR
<b>RECEIVED:</b> 09/19/2013 15:24 <b>REPORTED:</b> 09/19/2013 15:30					
PAI-1 Ag, P		PENDING			
<b>RECEIVED:</b> <b>REPORTED:</b>					
Soluble Fibrin Monomer	H	100	mcg/mL	0.0 - 7.9	MCR
<b>RECEIVED:</b> 09/19/2013 15:07 <b>REPORTED:</b> 09/19/2013 15:07					

\* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director:
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<b>Patient Name</b> TESTINGRNV,RUO	<b>Collection Date and Time</b> 09/18/2013 00:00	<b>Report Status</b> Partial
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\* Report times for Mayo performed tests are CST/CDT