

Laboratory Service Report

1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #	
SAMPLEREPORT,FX13M	SA00049072	46	M	SA00049072	
Ordering Phys			•	DOB 05/25/1966	
Client Order # SA00049072	Account Information	1		Report Notes	
Collected 09/17/2012 13:00	C7028846-DLMP RO 3050 SUPERIOR DR	IVE			
Printed 09/28/2012 13:54	ROCHESTER,MN 55	901			

Test	Flag	Results	Unit	Reference Value	Perform Site*
Factor XIII, Qual, w/Reflex 1:1 Mix Factor XIII, Qualitative	AB	Lysis	REPORTED 09/1	8/2012 13:19 No Lysis	Y00
Factor XIII, 1:1 Mix	AB	No Lysis			6 Y00 6

Clot lysis was observed in the qualitative factor XIII screening test. Clot lysis was not observed when the test was repeated using a 1:1 mix of patient plasma and pooled normal plasma. This pattern suggests severe factor XIII deficiency (less than approximately 1% of normal activity).

False-positive results (lysis) can be caused by heparin (therapy with unfractionated or low molecular weight heparin or contamination from a line), decreased or abnormal fibrinogen, increased fibrinolysis (inherited or acquired fibrinolytic disorders), fibrinolytic drugs, or other factors that affect clot structure or stability. Interpretation of these results requires correlation with clinical information and the results of additional laboratory testing. Quantitative factor XIII testing (factor XIII activity) is recommended for confirmation (ARUP sendout test).

* Performing Site:

Γ	Vooc	ARUP Laboratories	Lab Directory
1	Y006	500 Chipeta Way Salt Lake City, UT 84108	Lab Director:

	Collection Date and Time 09/17/2012 13:00	Report Status Final
	03/11/2012 13:00	
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^{*} Report times for Mayo performed tests are CST/CDT