ARUP LABORATORIES | A nonprofit enterprise of the University of Utah and its Department of Pathology

Effective Date: October 20, 2025

TEST CHANGE

Factor XIII Activity

2006182, F13 A

Specimen Requirements:	
Patient Preparation:	
Collect:	Lt. blue (sodium citrate). Special~~Refer to~~ Specimen Collection and Handling Hemostasis/Thrombosis Specimens guide located at https://www.aruplab.com/Specimen-Handling/SpecialSpecimenCollection/Hemostasis-Thrombosis.pdf ~~for hemostasis/thrombosis specimen handling guidelines.~~
Specimen Preparation:	Transfer 2 mL platelet-poor plasma to an ARUP standard transport tube.~~Standard Transport Tube.~~ (Min: 1 mL)
Transport Temperature:	CRITICAL FROZEN. Separate specimens must be submitted when multiple tests are ordered.
Unacceptable Conditions:	Serum. EDTA plasma, clotted or hemolyzed specimens.
Remarks:	
Stability:	Ambient: 4 hours; Refrigerated: Unacceptable; Frozen: ~~at -20 Degrees C or below: 1 month; Frozen at -70 Degrees C or below:~~ 3 months
Methodology:	Chromogenic Assay
Performed:	Tue
Reported:	1-8 days
Note:	
CPT Codes:	85290
New York DOH Approval Status:	This test is New York DOH approved.
Interpretive Data:	

~~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.~~

Reference Interval:

Factor XIII Activity 69-143%

Deleted Cells