

NEW TEST - Available Now

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Vitamin C, Plasma (High-Dose Therapy)

3017651, VIT C IV

Specimen Requirements:			
Patient Preparation:			
Collect:	Green (sodium or lithium heparin). Place specimen in ice bath immediately. Also acceptable: Plasma separator tube.		
Specimen Preparation:	Protect from light, centrifuge, transfer plasma, and freeze within 1 hour of collection. Transfer 0.5 mL plasma to an ARUP amber transport tube. (Min: 0.3 mL)		
Transport Temperature:	CRITICAL FROZEN AND LIGHT PROTECTED. Separate specimens must be submitted when multiple tests are ordered.		
Unacceptable Conditions:	EDTA plasma, whole blood, or body fluids. Grossly hemolyzed specimens.		
Remarks:	Thawing and refreezing of the specimen and exposure to light will result in decreased vitamin C concentration.		
Stability:	After separation from cells: Ambient: Unacceptable; Refrigerated: Unacceptable; Frozen: 1 month		
Methodology:	Quantitative High Performance Liquid Chromatography- Tandem Mass Spectrometry		
Performed:	Sun-Sat		
Reported:	1-6 days		
Note:	Thawing and refreezing of the specimen and exposure to light will result in decreased vitamin C concentration.		
CPT Codes:	82180		
New York DOH Approval Status:	This test is New York DOH approved.		
Interpretive Data:			

Effective Date: May 3, 2024

Intravenous vitamin C (IVC) administration produces millimolar plasma ascorbate (vitamin C) concentrations. Therapeutic concentrations average 15 mmol/L and range from 1-30 mmol/L. The maximum plasma concentration achieved by oral supplementation of vitamin C is approximately 250



Vitamin C concentration is reported as micromoles per liter (). To convert concentration to millimoles per liter (mmol/L), multiply the result by 0.001.

Reference Interval:

Test Number	Components	Reference Interval
	Vitamin C, Plasma	23-114 u mol/L

Effective Date: May 3, 2024

HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.