Test Definition: FISVU

Isavuconazole (CRESEMBA) LC-MS/MS Plasma

Overview

Method Name
Reversed-phase UPLC-MS/MS

NY State Available
Yes

Specimen

Specimen Type
Plasma

Specimen Required
*****Must submit one specimen per order. Specimens cannot be shared between multiple orders.*****

Collection Container/Tube: Lavender top (EDTA) (plasma gel/PST are **not acceptable**).
Submission Container/Tube: Sterile, screw-top tube
Specimen Volume: 1 mL Plasma
Collection Instructions:
1. Within 30 minutes of collection, centrifuge and aliquot 1 mL of plasma into a sterile, screw-top tube.
2. Send ambient

Specimen Minimum Volume
Plasma: 0.5 mL

Reject Due To

Gross hemolysis	Reject
Gross lipemia	Reject
Sodium heparin	Reject
ACD	Reject

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Plasma	Ambient (preferred)	14 days	
	Refrigerated	14 days	
	Frozen	30 days	

Clinical & Interpretive

Clinical Information
Refer to www.eurofins-viracor.com/test-menu/

Reference Values
The analytical measuring range is 0.1-10 mcg/mL. The therapeutic range has not yet been established.

Performance

PDF Report
No

Day(s) Performed
Monday through Saturday

Report Available
3 to 5 days

Performing Laboratory Location
Eurofins Viracor

Fees & Codes

Fees
- Authorized users can sign in to Test Prices for detailed fee information.
- Clients without access to Test Prices can contact Customer Service 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact Customer Service.

Test Classification
This test was developed and its performance characteristics determined by Eurofins Viracor. It has not been cleared or approved by the U.S. Food and Drug Administration.

CPT Code Information
80299

LOINC® Information

Test ID	Test Order Name	Order LOINC® Value
FISVU	Isavuconazole, Plasma	Not Provided

MAYO CLINIC LABORATORIES

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Isavuconazole (CRESEMBA) LC-MS/MS Plasma

Result ID	Test Result Name	Result LOINC® Value
FISVU	Isavuconazole, Plasma	10989-2