

Client: Example Client ABC123

123 Test Drive

Salt Lake City, UT 84108 UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB 4/1/1955 Gender: Female

Patient Identifiers: 01234567890ABCD, 012345

Visit Number (FIN): 01234567890ABCD **Collection Date:** 00/00/0000 00:00

Immunoglobulins, Serum Cryoprecipitins (Reflex for 2002063 CRYGB QNT and 2002403 CRYO TYPING only. NOT orderable by clients.)

ARUP test code 2002125

Immunoglobulin G, Cryoprecipitate

16 mg/dL

Н

(Ref Interval: 0-0)

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.

Immunoglobulin A, Cryoprecipitate

See Note mg/dL

(Ref Interval: None Detected)

The cryoprecipitate IgA concentration is less than the analyzer sensitivity for this analyte. IgA cryoprecipitate is less than 7

mg/dL.

Immunoglobulin M, Cryoprecipitate

16 mg/dL

Н

(Ref Interval: 0-0)

Immunofixation Electrophoresis, Serum

ARUP test code 2012572

Immunofix Electrophoresis Serum

See Note

The cryoglobluin is characterized as a Type II cryoglobulin with

monoclonal IgM kappa and polyclonal IgG.

INTERPRETIVE INFORMATION: Immunofix Electrophoresis, Serum

This information should be correlated with the results of serum protein electrophoresis, quantitative immunoglobulins and other clinical and laboratory information.

EER Immunofix Electrophoresis Serum

See Note

Authorized individuals can access the ARUP Enhanced Report using the following link:

H=High, L=Low, *=Abnormal, C=Critical

4848



Cryoglobulin, Qualitative with Reflex to IFE Typing and Quantitative IgA, IgG, and IgM

ARUP test code 2002403

Cryoglobulin, Qualitative

POS 24Hour * (Ref Interval: NEG 72Hour)

Cryoglobulin detected at 24 hrs.

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.

VERIFIED/REPORTED DATES				
Procedure	Accession	Collected	Received	Verified/Reported
Immunoglobulin G, Cryoprecipitate	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Immunoglobulin A, Cryoprecipitate	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Immunoglobulin M, Cryoprecipitate	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Immunofix Electrophoresis Serum	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Cryoglobulin, Qualitative	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
EER Immunofix Electrophoresis Serum	23-215-122043	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

Patient: Patient, Example
ARUP Accession: 23-215-122043
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
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