



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 04/21/2022 17:24  
**Last Report Issued** 06/20/2017 14:22

**88888**  
Clinical Example Report  
Attn: Example Reports  
200 Welsh Road  
Horsham, PA 19044

**Patient Name** 2171SP-POS  
**Patient ID** 2171SP-POS  
**Chain** 17001125  
**DOB** Not Given  
**Sex** Not Given  
**Workorder** 17001125  
**Received** 06/13/2017

**Lab ID** 17001125-001  
**Matrix** Serum or Plasma  
**Patient Name** 2171SP-POS  
**Patient ID** 2171SP-POS  
**Container Type** Clear vial

**Collect Dt/Tm** Not Given  
**Source** Not Given

**Approx Vol/Weight** Not Given

**Receipt Notes** None Entered

Analysis and Comments	Result	Units	Reporting Limit	Notes
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**2171SP Gold, Serum/Plasma**

Analysis by Inductively Coupled Plasma/Optical Emission Spectrometry (ICP/OES)

Gold	500	mcg/L	100	ELEVATED
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Normally: Less than 0.1 mcg/L

Steady-state concentrations following 50 mg weekly intramuscular gold sodium thiomalate injections: Approximately 3000 - 5000 mcg/L

Steady-state concentrations following 6 mg daily oral auranofin: Approximately 500 - 700 mcg/L

Specimens for elemental testing should be collected in certified metal-free containers. Elevated results for elemental testing may be caused by environmental contamination at the time of specimen collection and should be interpreted accordingly. It is recommended that unexpected elevated results be verified by testing another specimen in a trace metal free container.

This test was developed and its performance characteristics determined by NMS Labs. It has not been cleared or approved by the US Food and Drug Administration.