



**NMS Labs**

**CONFIDENTIAL**

200 Welsh Road, Horsham, PA 19044-2208  
Phone: (215) 657-4900 Fax: (215) 657-2972  
e-mail: nms@nmslabs.com

Robert A. Middleberg, PhD, F-ABFT, DABCC-TC, Laboratory Director

**Demo Report**

**Report Issued** 03/30/2020 12:25  
**Last Report Issued** 01/30/2013 14:52

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

**Patient Name** 2330SP-POS  
**Patient ID** 2330SP-POS  
**Chain** 13000611  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 13000611  
**Received** 01/30/2013 12:38

**Sample ID** 13000611-001  
**Matrix** Serum or Plasma  
**Patient Name** 2330SP-POS  
**Patient ID** 2330SP-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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**2330SP Hydrochlorothiazide, Serum/Plasma**

Analysis by High Performance Liquid Chromatography/  
Tandem Mass Spectrometry (LC-MS/MS)

Hydrochlorothiazide Synonym(s): Microzide®	500	ng/mL	25	ELEVATED
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Daily administration of oral hydrochlorothiazide produced the following plasma concentrations in hypertensive patients:

25 mg = 17 +/-8 ng/mL trough  
and 76 +/- 26 ng/mL 5 hours post dose

75 mg = 34 +/- 17 ng/mL trough  
and 200 +/- 93 ng/mL 5 hours post dose.

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.