



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 03/30/2020 11:28  
**Last Report Issued** 01/31/2013 09:12

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

**Patient Name** 0796SP-POS  
**Patient ID** 0796SP-POS  
**Chain** 13000645  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 13000645  
**Received** 01/31/2013 09:10

**Sample ID** 13000645-001  
**Matrix** Serum or Plasma  
**Patient Name** 0796SP-POS  
**Patient ID** 0796SP-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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**0796SP Bumetanide, Serum/Plasma**

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

Bumetanide	500	ng/mL	25	ELEVATED
Synonym(s): Bumex®				
Peak plasma concentrations were 97 +/- 15 ng/mL in eight healthy subjects following 2 mg and 180 +/- 100 ng/mL in four healthy subjects following 5 mg oral bumetanide.				

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.