



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 03/30/2020 11:54  
**Last Report Issued** 02/02/2012 08:16

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

**Patient Name** 1275SP-POS  
**Patient ID** 1275SP-POS  
**Chain** 12000242  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 12000242  
**Received** 02/02/2012 08:13

**Sample ID** 12000242-001  
**Matrix** Serum or Plasma  
**Patient Name** 1275SP-POS  
**Patient ID** 1275SP-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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**1275SP Clonidine, Serum/Plasma**

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

Clonidine	5.0	ng/mL	0.050	ELEVATED
Synonym(s): Kapvay®; Duraclon; Catapres® Immediate-release, oral: 0.50 - 2.0 ng/mL, 2 hours after administration Sustained-release, patch: 0.20 - 2.0 ng/mL, at steady-state Sustained-release, oral: 0.20 - 0.27 ng/mL, 6.8 +/- 3.6 hours after a 0.1 mg single dose in healthy fed adults; children receive higher doses on a mg/kg basis.				

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.