



**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 11:53  
**Last Report Issued** 12/12/2013 08:18

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

**Patient Name** 1272B-POS  
**Patient ID** 1272B-POS  
**Chain** 13004296  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 13004296  
**Received** 12/12/2013 08:09

**Sample ID** 13004296-001  
**Matrix** Blood  
**Patient Name** 1272B-POS  
**Patient ID** 1272B-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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**1272B Citalopram, Blood**

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

Citalopram / Escitalopram Synonym(s): Celexa®/Lexapro®	5000	ng/mL	50	ELEVATED
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Steady-state serum or plasma levels from patients on a daily regimen of 30 to 60 mg Citalopram: 9 - 200 ng/mL.

This test is not Chiral specific. Patients who have taken Escitalopram (Lexapro®), as opposed to Racemic Citalopram (Celexa®), within the past 3 days may have falsely elevated values.

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.