



**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 12:05  
**Last Report Issued** 05/13/2019 08:06

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

**Patient Name** 1777B-POS  
**Patient ID** 1777B-POS  
**Chain** 18001898  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 18001898  
**Received** 11/21/2018 08:54

**Sample ID** 18001898-001  
**Matrix** Blood  
**Patient Name** 1777B-POS  
**Patient ID** 1777B-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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**1777B Dipyridamole, Blood**

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

Dipyridamole	50	mcg/mL	0.050	
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Synonym(s): Persantine®

Steady-state trough plasma concentrations following a  
three times daily regimen of 50 mg:  
0.85 +/- 0.50 mcg/mL.  
The blood to plasma ratio is approximately 0.7.

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.