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## Demo Report

**Report Issued** 04/21/2022 17:40
**Last Report Issued** 01/14/2019 06:54

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

**Patient Name** 2504SP-POS
**Patient ID** 2504SP-POS
**Chain** 19000128
**DOB** Not Given
**Sex** Not Given
**Workorder** 19000128

**Received** 01/14/2019

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**Lab ID** 19000128-001
**Matrix** Serum or Plasma
**Patient Name** 2504SP-POS
**Patient ID** 2504SP-POS
**Container Type** Clear vial

**Collect Dt/Tm** Not Given
**Source** Not Given

**Approx Vol/Weight** Not Given

**Receipt Notes** Not frozen as required

| Analysis and Comments | Result | Units | Reporting Limit | Notes |
|---|---|---|---|---|
| **2504SP Levodopa, Serum/Plasma** | | | | |
| Analysis by High Performance Liquid Chromatography/Tandem Mass Spectrometry (LC-MS/MS) | | | | |
| Levodopa | 5.0 | mcg/mL | 0.020 | |
| The target plasma concentration of levodopa in Parkinsonian patients is 2 +/- 0.5 mcg/mL. The average peak plasma levodopa concentration following a single oral dose of Sinemet® containing 200 mg levodopa was 1.2 mcg/mL at 0.5 hours for normal release and 3.3 mcg/mL at 2 hours for controlled release. At steady-state, the average trough plasma levodopa concentration following oral Sinemet® containing 200 mg levodopa was 0.07 mcg/mL for normal release and 0.16 mcg/mL for controlled release. | | | | |

Sample receipt condition is inappropriate for test code: Not frozen as required

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.