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**CONFIDENTIAL**

## Demo Report

**Report Issued** 03/30/2020 12:17
**Last Report Issued** 04/05/2017 14:15

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

**Patient Name** 2110B-POS
**Patient ID** 2110B-POS
**Chain** 17000661
**Age** Not Given **DOB** Not Given
**Gender** Not Given
**Workorder** 17000661

**Received** 04/05/2017 13:30

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**Sample ID** 17000661-001
**Matrix** Blood
**Patient Name** 2110B-POS
**Patient ID** 2110B-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 |
|---|---|---|---|---|
| **2110B Fluphenazine, Blood** | | | | |
| Analysis by High Performance Liquid Chromatography/ Tandem Mass Spectrometry (LC-MS/MS) | | | | |
| Fluphenazine | 250 | ng/mL | 0.50 | ELEVATED |

Synonym(s): Permitil®; Prolixin®

Schizophrenic patients maintained with depot injections of fluphenazine decanoate at 12.5 to 50 mg every 1 to 2 weeks had plasma fluphenazine concentrations ranging from 1 to 17 ng/mL. Healthy subjects given single oral doses of 5 mg fluphenazine had peak plasma concentrations averaging 0.6 ng/mL (SEM +/- 0.1 ng/mL).

The blood to plasma ratio for fluphenazine is approximately 1.

Substance(s) known to interfere with the identity and/or quantity of the reported result: Trimeprazine

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.