



NMS Labs

CONFIDENTIAL

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Demo Report

Report Issued 04/21/2022 17:43
Last Report Issued 01/23/2015 13:05

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

Patient Name 2527SP
Patient ID 2527SP
Chain 15000174
DOB Not Given
Sex Not Given
Workorder 15000174
Received 01/23/2015

Lab ID 15000174-001
Matrix Serum or Plasma
Patient Name 2527SP
Patient ID 2527SP
Container Type Clear vial

Collect Dt/Tm Not Given
Source Not Given

Approx Vol/Weight Not Given

Receipt Notes None Entered

Table with 5 columns: Analysis and Comments, Result, Units, Reporting Limit, Notes

2527SP Lacosamide, Serum/Plasma

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

Table row: Lacosamide, None Detected, mcg/mL, 0.50

Synonym(s): Vimpat®

Peak plasma concentrations are reached 1 to 2 hours after a single oral or intravenous dose with a half-life of 13 hours. Following a single 200 mg dose administered as a 30-minute infusion, a 60-minute infusion, or orally as a tablet to 24 male subjects, mean maximum plasma lacosamide concentrations were 5.95 +/- 1.49, 5.38 +/- 1.10 and 5.15 +/- 1.4 mcg/mL, respectively.

Mean plasma concentrations following maintenance doses:
200 mg/day: 4.99 +/- 2.51 mcg/mL;
400 mg/day: 9.35 +/- 4.22 mcg/mL;
600 mg/day: 12.46 +/- 5.60 mcg/mL.

NMS Labs derived data:
5th - 95th Percentile Data: 1.8 - 13.0 mcg/mL
Mean: 5.3 mcg/mL
(N = 14900)

Results for sample 15000174-001 are continued on next page



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Lab ID 15000174-001
Matrix Serum or Plasma
Patient Name 2527SP
Patient ID 2527SP

Collect Dt/Tm Not Given
Source Not Given

Analysis and Comments	Result	Units	Reporting Limit	Notes
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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.