

LABORATORY REPORT

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108 **EXAMPLE, REPORT W**

WX0000003826 F 12/05/1988 34 Y

Referral Testing

Collected: 09/22/2023 12:14 Received: 09/22/2023 12:14

<u>Test Name</u> <u>Result</u> <u>Flag Ref-Ranges</u> <u>Units</u> <u>Site</u>

Leflunomide as Metab (Teriflunomide), S/P 30000 ng/mL NMRL

Reporting Limit: 500 ng/mL Synonym(s): Leflunomide Metabolite Leflunomide is used for the management of the signs and symptoms of rheumatoid arthritis. Teriflunomide is an active metabolite of Leflunomide and is indicated for the treatment of patients with relapsing forms of multiple sclerosis.

Mean steady-state trough plasma concentrations of teriflunomide from patients on daily regimens of 10 or 25 mg of leflunomide were 18000 and 63000 ng/mL, respectively, after 24 days. Recommended doses of teriflunomide and leflunomide result in a similar range of teriflunomide plasma concentrations. Based on this, the expected therapeutic range is between 18000 ng/mL and 63000 ng/mL.

Following completion of an elimination regimen, plasma concentrations should be determined twice at least 14 days apart to verify that concentrations are less than 20 $\,\mathrm{ng/mL}$.

THIS TEST IS NOT MEANT TO MONITOR THE ELIMINATION OF TERIFLUNOMIDE IN WOMEN OF CHILDBEARING POTENTIAL WHO DISCONTINUE LEFLUNOMIDE. The drug carries a black box warning for hepatotoxicity and pregnancy.

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

Testing performed at NMS Labs, Inc. 200 Welsh Road Horsham, PA 19044-2208 CLIA 39D0197898

Reported Date: 2023.09.22 12:15