



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 03/30/2020 13:30  
**Last Report Issued** 01/20/2014 13:02

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

**Patient Name** 4367B  
**Patient ID** 4367B  
**Chain** 14000176  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 14000176  
**Received** 01/20/2014 12:33

**Sample ID** 14000176-001  
**Matrix** Blood  
**Patient Name** 4367B  
**Patient ID** 4367B  
**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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**4367B Teriflunomide (Pre-Pregnancy Monitoring), Blood**

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

Teriflunomide	None Detected	ng/mL	5.0	
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Synonym(s): Aubagio®

Teriflunomide is indicated for the treatment of patients with relapsing forms of multiple sclerosis.

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 ng/mL. The blood to plasma ratio is 0.5 to 0.7. The drug carries a black box warning for hepatotoxicity and teratogenicity.

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.