



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 03/30/2020 11:07  
**Last Report Issued** 11/14/2011 06:09

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

**Patient Name** 0013SP  
**Patient ID** 0013SP  
**Chain** 11003488  
**Age** Not Given **DOB** Not Given  
**Gender** Not Given  
**Workorder** 11003488  
**Received** 11/14/2011 06:05

**Sample ID** 11003488-001  
**Matrix** Serum or Plasma  
**Patient Name** 0013SP  
**Patient ID** 0013SP  
**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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**0013SP Acamprosate, Serum/Plasma**

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

Acamprosate	None Detected	ng/mL	50	
Synonym(s): Campral; N-acetylhomotaurine; calcium acetylhomotaurinate				

Acamprosate is a synthetic psychotropic drug used in the treatment of alcohol dependence. It has been shown to reduce cumulative days drinking and alcohol-induced cravings. Upon ingestion, acamprosate immediately dissociates into N-acetyl homotaurine; this is the compound that is measured and reported. Steady-state acamprosate concentrations following 2 x 333 mg tablets three times daily ranged from 370 +/- 145 to 644 +/- 386 ng/mL and were achieved 3.5 +/- 0.5 to 9.0 +/- 1.9 hours after dose.

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.