



**NMS Labs**

**CONFIDENTIAL**

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**Demo Report**

**Report Issued** 05/05/2022 10:34  
**Last Report Issued** 03/17/2022 08:07

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

**Patient Name** 4783B  
**Patient ID** 4783B  
**Chain** 22000663  
**DOB** Not Given  
**Sex** Not Given  
**Workorder** 22000663  
**Received** 02/14/2022

**Lab ID** 22000663-001  
**Matrix** Blood  
**Patient Name** 4783B  
**Patient ID** 4783B  
**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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**4783B Voclosporin, Blood**

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

Voclosporin	None Detected	ng/mL	1.0	
Synonym(s): Lupkynis®				

Voclosporin is a calcineurin inhibitor approved for the treatment of adults with lupus nephritis. Published Phase 3 clinical trial data indicated voclosporin in combination with mycophenolate mofetil (MMF) and low-dose steroids led to a clinically and statistically superior complete renal response rate versus MMF and low-dose steroids alone, with a comparable safety profile. Peak whole blood concentrations are reported to average 32 +/- 9.9 ng/mL and 950 +/- 250 ng/mL approximately 1-2 hours following single oral doses of voclosporin at 0.25 mg/kg and 4.5 mg/kg, respectively.

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.