



**NMS Labs**

**CONFIDENTIAL**

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Robert A. Middleberg, PhD, F-ABFT, DABCC-TC, Laboratory Director

**Demo Report**

**Report Issued** 03/30/2020 12:12  
**Last Report Issued** 10/03/2011 14:08

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

**Patient Name** 1968TI-POS  
**Patient ID** 1968TI-POS  
**Chain** 11002854  
**Age** Not Given      **DOB** Not Given  
**Gender** Not Given  
**Workorder** 11002854  
**Received** 10/03/2011 14:01

**Sample ID** 11002854-001  
**Matrix** Tissue  
**Patient Name** 1968TI-POS  
**Patient ID** 1968TI-POS  
**Container Type** Clear vial

**Collect Dt/Tm** Not Given  
**Source** Adipose Tissue (Fat)

**Approx Vol/Weight** Not Given

**Receipt Notes**      None Entered

Tissue specimen required homogenization: 11002854-001

No testing performed on this sample.



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**Sample ID** 11002854-002  
**Matrix** Tissue  
**Patient Name** 1968TI-POS  
**Patient ID** 1968TI-POS  
**Container Type** Clear vial

**Collect Dt/Tm** Not Given  
**Source** Adipose Tissue (Fat)

**Approx Vol/Weight** Not Given

**Receipt Notes** Not frozen as required

NMS Labs generated homogenized Tissue sample: 11002854-002

Analysis and Comments	Result	Units	Reporting Limit	Notes
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**1968TI Eszopiclone / Zopiclone, Tissue**

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

Eszopiclone / Zopiclone	2000	ng/g	8.0	
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Synonym(s): Lunesta®, Imovane®

No reference data available.

This test is not chiral specific; therefore, Eszopiclone and/or Racemic Zopiclone (not approved in the US) may be present.

Sample receipt condition is inappropriate for test code: Not frozen as required

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.