

ILS Laboratories
 8222 Vickers St, Suite 106, San Diego, CA 92111
 (619) 329-3999 | ils-lab.com

Tesamorelin

Tested for: Voltera Sciences
 www.volterasciences.com


COA #: **COA-2026-412739**
 Lot Number: **TSM10.05.28.26**
 Accession #: **ACC-2026-3697**

Method: **Full QC Panel**
 Analysis Date: **06/05/2026**
 Appearance: **Good**
 Date Received: **06/01/2026**

PASS



Scan to verify
 authenticity at ils-lab.com
 Access Code: GTD8AM7C

Identity	Peptide Purity	99.78%		Fentanyl Free
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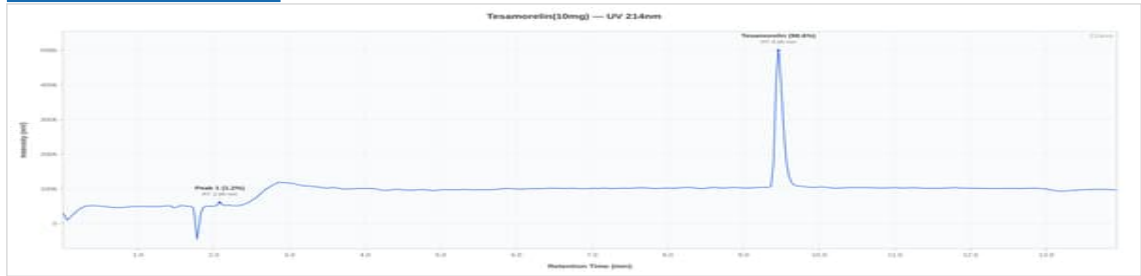


Tesamorelin - TSM10.05.28.26

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.78%	%	PASS
Net Peptide Content	Report Only	10.67	mg	N/A
Identity (HPLC-RTM)	Tesamorelin	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



Tesamorelin - TSM10.05.28.26: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	<i>Not Detected</i>	PASS
Cadmium (Cd)	NMT 0.5 ppm	<i>Not Detected</i>	PASS
Chromium (Cr)	NMT 10 ppm	<i>Not Detected</i>	PASS
Mercury (Hg)	NMT 1.5 ppm	<i>Not Detected</i>	PASS
Lead (Pb)	NMT 1 ppm	<i>Not Detected</i>	PASS

Sterility Testing (PCR)

Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS




Dr. Greg Kalyuzhny
 Lab Director
 6/7/2026

COA #: **COA-2026-412739**
 Access Code: **GTD8AM7C**
 Verify: portal.ils-lab.com/verify/KrWjuKgnvFQZdyls
 Issued: 6/7/2026

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	0.08 EU/mL	Reported

About this result: Endotoxin is reported as a quantitative value. Acceptable limits vary by product type and matrix, so no universal pass/fail threshold applies to RUO products. This result is below commonly referenced endotoxin thresholds.

Notes & Methodology

1. Date Tested: 06/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Tesamorelin by HPLC. Identification by chromatographic retention time comparison with a reference standard.
3. Elemental impurities analyzed by ICP-MS per USP <233> methodology. Acceptance criteria are internal laboratory quality screening limits for research-use materials and do not represent evaluation against any specific pharmacopeial monograph or product specification.
4. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
5. Peptide purity determined by RP-HPLC area normalization at 214 nm. Value represents the percentage of the target peptide relative to all peptide-related peaks. Non-peptide process-related impurities, if detected, are excluded from the calculation.



Dr. Greg Kalyuzhny
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Issued: 6/7/2026

Client: Voltera

Certified: 05/14/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	Tesamorelin	Form	Lyophilized powder
Batch	TESA.05.02.2026	Labeled Qty	10 mg
Mol. Formula	C22H366N72O67S	CAS Number	901758-09-6
Cap Color	Purple	Crimp Color	Blue

TEST RESULTS

	REFERENCE STANDARD	RESULT
Batch Avg Purity	(>98%)	99.866% <input checked="" type="checkbox"/>
Batch Avg Net Content	(10mg ± 10%)	11.64 mg <input checked="" type="checkbox"/>
Identity Confirmation (LC-MS)	(Tesamorelin)	Tesamorelin <input checked="" type="checkbox"/>
Endotoxin Safety Screen	(≤0.5 EU/mL)	PASS <input checked="" type="checkbox"/>
Microbial Sterility Screen	(No Growth)	No Growth <input checked="" type="checkbox"/>

HEAVY METAL SCREENING

Analyte	Result	Status
Arsenic (As)	Negative	<input checked="" type="checkbox"/>
Cadmium (Cd)	Negative	<input checked="" type="checkbox"/>
Lead (Pb)	Negative	<input checked="" type="checkbox"/>
Mercury (Hg)	Negative	<input checked="" type="checkbox"/>

BATCH CONFORMITY RESULTS

Vial	Purity (%)	Net Content (mg)
Vial 1	99.896	11.73
Vial 2	99.836	11.56
Batch Average	99.866%	11.64 mg

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm

