

ILS Laboratories

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

Tesamorelin - 10mg

Tested for: Virtus Peptides

virtus | virtuspeptides.com

COA #: COA-2026-954425
Lot Number: NS-20260601 | TSM10-WHITE
Labeled Content: 10mg

Analysis Date: 07/21/2026
Appearance: Good
Date Received: 07/07/2026

Method: Full QC Panel

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: EEJGQH3D

Identity	Peptide Purity	Fentanyl Free
Tesamorelin	99.31%	

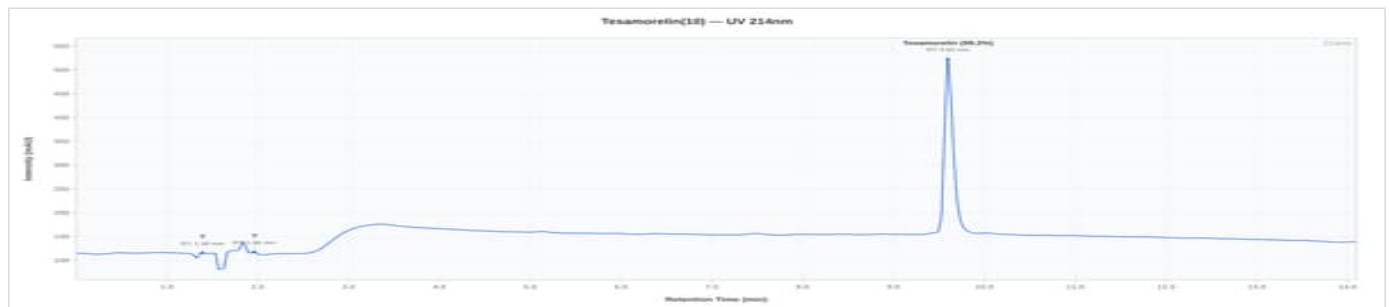
Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.31%	%	PASS
Net Peptide Content	Report Only	10.42	mg	N/A
Identity (HPLC-RTM)	Tesamorelin	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



Tesamorelin 10mg - NS-20260601 | TSM10-WHITE

HPLC Chromatogram



Tesamorelin 10mg - NS-20260601 | TSM10-WHITE: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

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

Sterility Testing (PCR)

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

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	NMT 0.5 EU/mL	0.142 EU/mL	PASS

Acceptance criteria: NMT 0.5 EU/mL per client specification.

Notes & Methodology

1. Date Tested: 07/28/2026. Methods: Purity, Identity & Quantitation (HPLC).
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
Lab Director
7/21/2026

COA #: **COA-2026-954425**
Access Code: **EEJGQH3D**
Verify: portal.ils-lab.com/verify/rQHviho_5ZmWdxzp
Issued: 7/21/2026