



Certificate of Analysis

All tests passed. This certificate is verified and authentic.

TESTED FOR

KDA COMPOUNDS @
kdacollection.com

Product Information

Product
TSM 10mg - 10mg

Lot Number
KDA-TES-2762

Appearance
Good

Test Type
Full QC Panel

Date Received
06/16/2026

Analysis Confirmed
06/23/2026



Confirmed Fentanyl Free

Immunoassay, 50 ng/mL cutoff

Test Results — Full QC Panel

Analyte	Limit	Result	Unit	Status
Peptide Purity (HPLC)	≥ 95.0%	99.48%	%	PASS
Net Peptide Content	Report Only	10.44	mg	N/A
Identity (ID)	Tesamorelin	Confirmed	-	PASS

⊠ Elemental Impurities — ICP-MS per USP <233>

Element	Limit (ppm)	Result (ppm)	Status
Arsenic (As)	≤ 1.5 ppm	0.1594 ppm	PASS
Cadmium (Cd)	≤ 0.5 ppm	0.1228 ppm	PASS
Chromium (Cr)	≤ 10 ppm	< LOQ	PASS
Mercury (Hg)	≤ 1.5 ppm	0.1612 ppm	PASS
Lead (Pb)	≤ 1 ppm	0.0696 ppm	PASS

Overall: PASS

🧫 Sterility Testing (PCR)

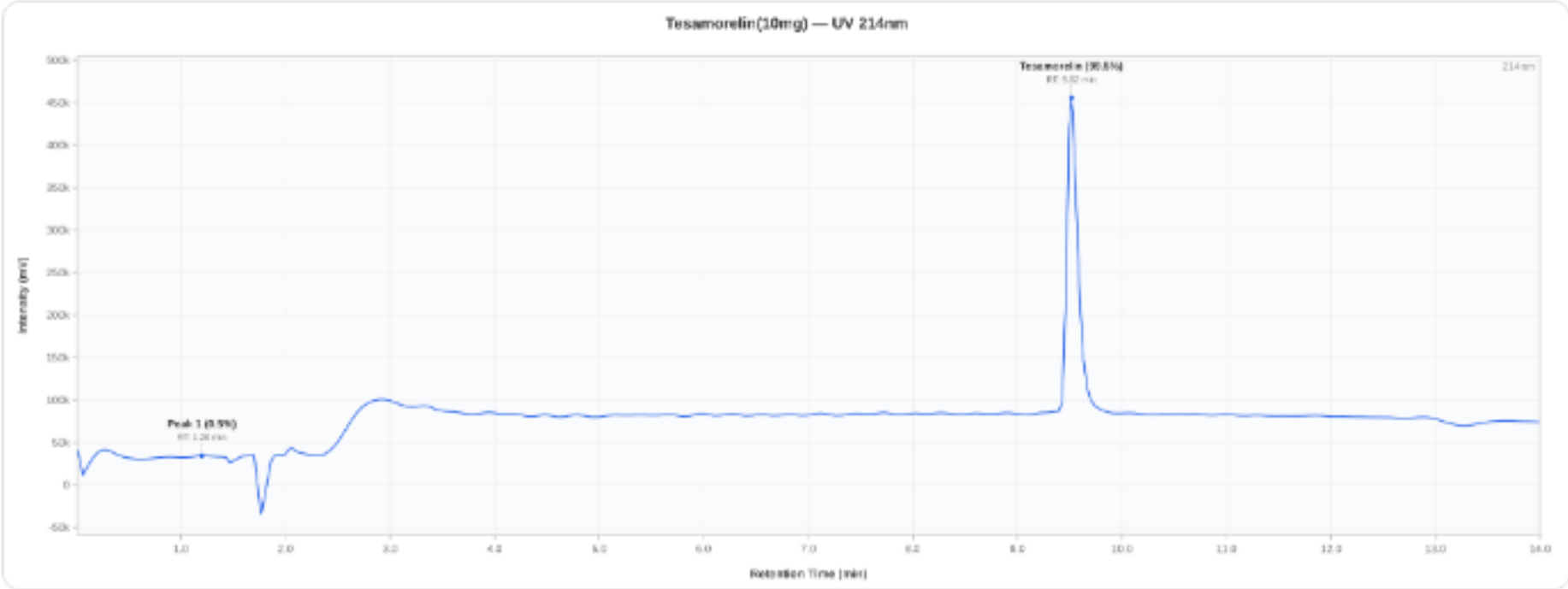
Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS

💧 Endotoxin Testing — USP <85>

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	≤ 0.05 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.

HPLC Chromatogram



CERTIFICATE CHANGE HISTORY

ILS QA

Sep 5, 2026, 1:29 PM PDT

Certain non-result date fields on this certificate were corrected to align with the original analysis record following data migration. Analytical results were not changed.

NOTES & METHODOLOGY

1. Date Tested: 06/23/2026. Methods: Full QC Panel.
2. The sample was confirmed to be TSM 10mg 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 chromogenic method. Acceptance criteria per client specification.
5. Sterility testing performed by PCR (Polymerase Chain Reaction) method.
6. 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.
7. Accreditation Scope: ILS Laboratories maintains ISO/IEC 17025 accreditation for activities listed in its published PJLA scope. Peptide testing reported on this certificate is performed for Research Use Only.
8. This certificate reports analytical results for the sample(s) submitted to ILS Laboratories for testing. All sample identifications, product names, brand names, lot/batch numbers, and labeled content are provided by the submitting party and have not been independently verified. Results reflect only the analytical testing of the specific sample(s) provided and do not constitute an evaluation or endorsement of the manufacturer's processes, quality systems, or overall production. ILS Laboratories makes no representations regarding the intended use, safety, efficacy, regulatory status, or intellectual property ownership of any materials tested.