

ILS Laboratories

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Tesa 10mg - 10mg

Tested for: Summit Biotech USA
<http://www.summitbiotechusa.com/>

COA #: COA-2026-954734
Lot Number: TESA-BLCK08
Labeled Content: 10mg

Method: Full QC Panel

Analysis Date: 07/26/2026
Appearance: Good
Date Received: 07/08/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: UQZG8277

Identity	Peptide Purity	 Fentanyl Free
Tesa 10mg	99.76%	

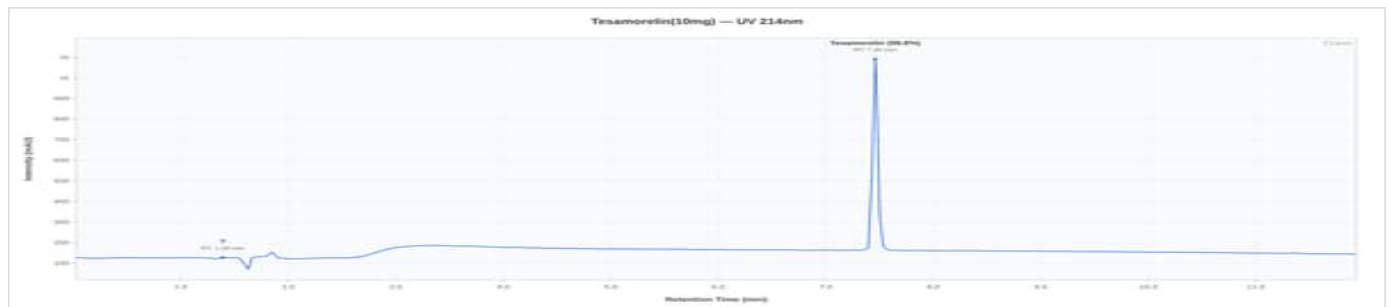
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.76%	%	PASS
Net Peptide Content	Report Only	10.43	mg	N/A
Identity (HPLC-RTM)	Tesamorelin	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



Tesa 10mg 10mg - TESA-BLCK08

HPLC Chromatogram



Tesa 10mg 10mg - TESA-BLCK08: 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
Lead (Pb)	NMT 1.0 ppm	0.409	PASS
Mercury (Hg)	NMT 1.5 ppm	Not Detected	PASS
Chromium (Cr)	NMT 10.0 ppm	Not Detected	PASS



Dr. Greg Kalyuzhny

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Lab Director
7/26/2026

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Verify: portal.ils-lab.com/verify/9IPRASGuu-OA_74D
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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	0.055 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: 08/01/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Tesa 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 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.




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