Certificate of Analysis

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

PT-141 - 10mg

PASS

Tested for: Felix Chem LLC

COA #:	**COA-2026-5TZUM_**	Method:	**Full QC Panel**
Lot Number:	**052126-PT1-C**	Analysis Date:	**05/29/2026**
Accession #:	**ACC-2026-3109**	Appearance:	**Good**
Labeled Content:	**10mg**	Vial Size:	**3mL**
Sample Matrix:	**Lyophilized**	Date Received:	**05/22/2026**

Scan to verify authenticity at ils-lab.com

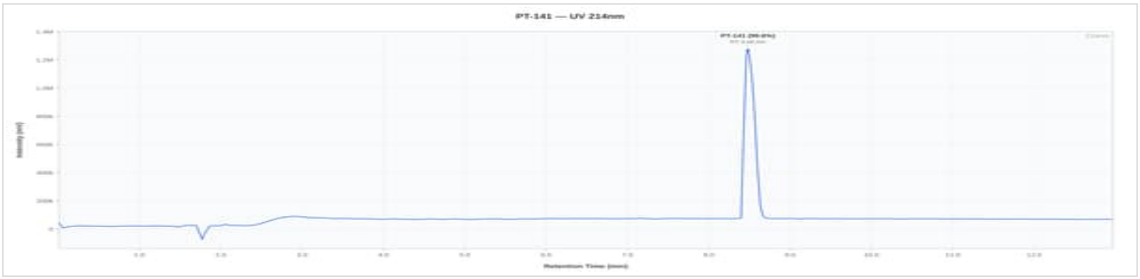
Identity	Peptide Purity	
PT-141	99.65%	Fentanyl Free

PT-141 10mg - 052126-PT1-C

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.65%**	%	**PASS**
Net Peptide Content	Report Only	**10.24**	mg	N/A
Identity (ID)	PT-141 10mg	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



PT-141 10mg - 052126-PT1-C: 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**

ilac-MRA PJLA Testing Accreditation No. 59423

Dr. Greg Kalyuzhny
Lab Director
5/31/2026

COA #: **COA-2026-5TZUM_**
Access Code: **G1SNYKBF**
Verify: portal.ils-lab.com/verify/mmotD4yVgXSl7Sy-
Issued: 5/31/2026

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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	**NMT 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.

Notes & Methodology

1. Date Tested: 05/29/2026. Methods: Full QC Panel.
2. The sample was confirmed to be PT-141 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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Lab Director
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