Certificate of Analysis

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

Retatrutide 10mg - 10mg

Tested for: Synthesis Peptides

PASS

COA #:	**COA-2026-954912**	Analysis Date:	**07/24/2026**
Lot Number:	**3G10726US**	Appearance:	**Good**
Accession #:	**ACC-2026-6959**	Sample Matrix:	**Lyophilized**
Labeled Content:	**10mg**	Volume:	**3mL**
		Date Received:	**06/29/2026**

Scan to verify authenticity at ils-lab.com
Access Code: YVU2D9FR

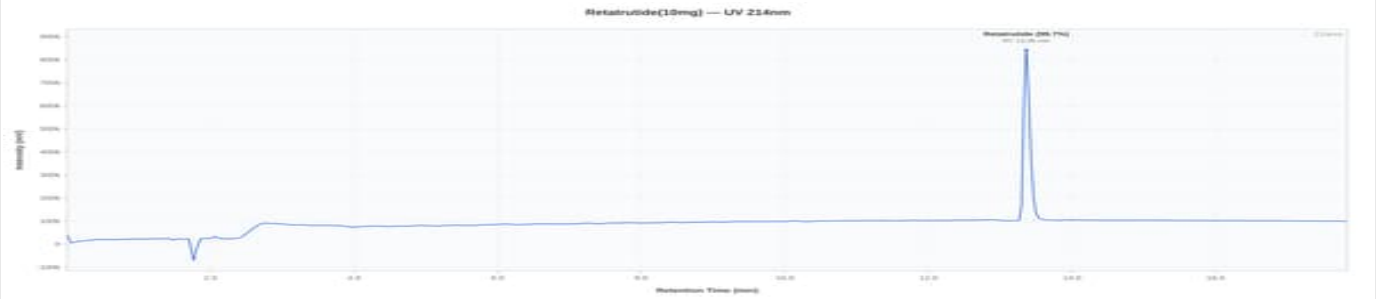
Method: **Full QC Panel**

Identity	Peptide Purity	
Retatrutide 10mg	**99.70%**	**Fentanyl Free**

Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.70%**	%	**PASS**
Net Peptide Content	Report Only	**10.32**	mg	N/A
Identity (HPLC-RTM)	Retatrutide	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Conformity V2

HPLC Conformity Testing Results (4 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.70%	10.32 mg	Confirmed	**PASS**
Conformity V1	99.69%	10.23 mg	Confirmed	**PASS**
Conformity V2	99.64%	10.4 mg	Confirmed	**PASS**
Conformity V3	99.37%	10.4 mg	Confirmed	**PASS**
Mean	**99.60%**	**10.34 mg**	—	—
Std Dev	**0.1347%**	**0.0701 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
8/5/2026

COA #: **COA-2026-954912**
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Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	.0485	**PASS**
Cadmium (Cd)	NMT 0.5 ppm	.0167	**PASS**
Lead (Pb)	NMT 1.0 ppm	0	**PASS**
Mercury (Hg)	NMT 1.5 ppm	0	**PASS**
Chromium (Cr)	NMT 10.0 ppm	.2247	**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 5 EU/mL	**NMT 0.05 EU/mL**	**PASS**

Acceptance criteria: NMT 5 EU/mL per client specification.

Notes & Methodology

1. Date Tested: 08/05/2026. Methods: Purity, Identity & Quantitation (HPLC).
2. The sample was confirmed to be Retatrutide 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.
6. Chromatogram shown is representative: Conformity V2 (99.64% purity, closest to batch mean of 99.60%).

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