Certificate of Analysis

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

R3 - 30mg

Tested for: RevPep
Revolutionpeptide.com

PASS

Scan to verify authenticity at ils-lab.com
Access Code: DAVAE8XV

COA #:	**COA-2026-955312**	Analysis Date:	**06/10/2026**
Lot Number:	**R3001P**	Appearance:	**Good**
Labeled Content:	**30mg**	Volume:	**3mL**
Volume:	**3mL**	Date Received:	**06/03/2026**

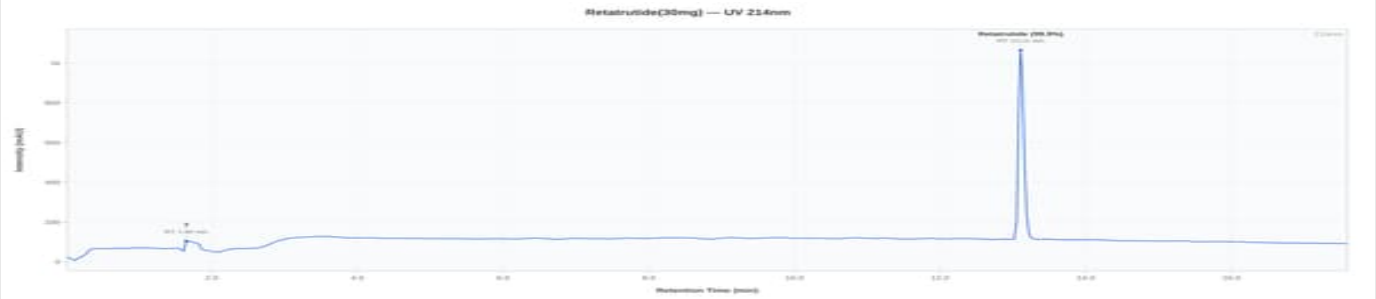
Method: **Full QC Panel**

Identity	Peptide Purity
R3	99.94%

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.94%**	%	PASS
Net Peptide Content	Report Only	**32.07**	mg	N/A
Identity (HPLC-RTM)	Retatrutide	**Confirmed**	-	PASS

HPLC Chromatogram



R3 30mg - R3001P: 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 ppm	0.0553 ppm	PASS
Mercury (Hg)	NMT 1.5 ppm	0.1073 ppm	PASS
Chromium (Cr)	NMT 10 ppm	0.0793 ppm	PASS

Sterility Testing (PCR)

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

Dr. Greg Kalyuzhny
Lab Director
8/8/2026

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Verify: portal.ils-lab.com/verify/ui17dZdf0eMQyDQ_
Issued: 8/8/2026

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: 08/08/2026. Methods: Full QC Panel.
2. The sample was confirmed to be R3 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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