

ILS Laboratories

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GLP-3 RT 20mg - 20mg

Tested for: NextSelf

 www.nextselflabs.com

COA #: COA-2026-957557
Lot Number: NS-3RT-20MG
Labeled Content: 20mg
Volume: 3mL
Method: Purity, Identity & Quantitation (HPLC)

Analysis Date: 10/08/2026
Appearance: Good
Volume: 3mL
Date Received: 10/02/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: GB5TM76Y

Identity

GLP-3 RT 20mg

Peptide Purity

98.85%



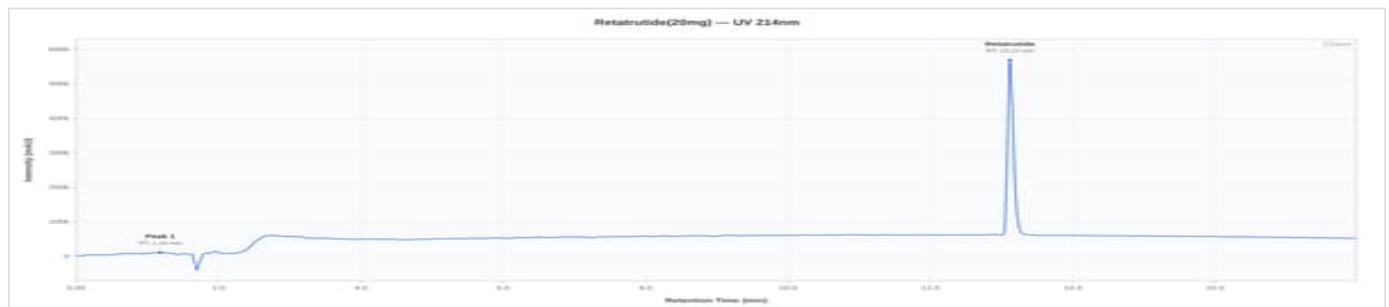
Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	98.85%	%	PASS
Net Peptide Content	Report Only	20.4	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



GLP-3 RT 20mg 20mg - NS-3RT-20MG

HPLC Chromatogram



GLP-3 RT 20mg 20mg - NS-3RT-20MG: UV Chromatogram

Notes & Methodology

- Date Tested: 10/08/2026. Methods: Purity, Identity & Quantitation (HPLC).
- The sample was confirmed to be GLP-3 RT 20mg by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- 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.
- 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.
- 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.