



CERTIFICATE OF ANALYSIS

COA Number: RUO-26-5C1057CE

SAMPLE SUMMARY

ISSUED TO	DECLARED IDENTITY	ISSUE DATE	
No Limit Labs	Retatrutide	Sep 29, 2026	
RUO SAMPLE ID	DECLARED LOT NUMBER	DECLARED CONTENT	DECLARED FORM
RUO-2026-CDD41A8E-30	RT-20-080326	20 mg / vial	Lyophilized powder

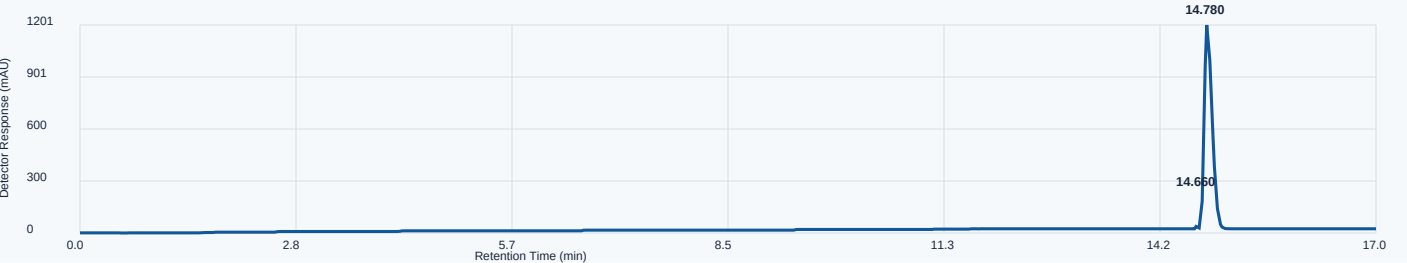
ANALYTICAL COMPOSITION AND IDENTITY

PURITY RESULT		VIAL 1	VIAL 2	AVERAGE	STATUS
PERCENTAGE PURITY		99.98%	99.94%	99.96%	PASS
COMPONENT	TARGET	VIAL 1	VIAL 2	AVERAGE	IDENTITY / STATUS
Retatrutide	20.00 mg	20.79 mg	20.75 mg	20.77 mg	CONFIRMED / PASS

COMPREHENSIVE 10-TEST ANALYTICAL PANEL

ANALYSIS	METHOD	REPORTED RESULT	ASSESSMENT
Molecular Identity Confirmation	LC-MS/HRMS	Identity confirmed	PASS
Chromatographic Purity	RP-HPLC-UV/DAD	99.98% / 99.94% (average 99.96%)	PASS
Net Peptide Content	Quantitative assay	20.79 / 20.75 mg (average 20.77 mg)	PASS
Vial-to-Vial Conformity Assessment	Multi-vial quantitative study	2/2 vials within criteria	PASS
Microbial Analysis (PCR)	Validated PCR-based microbial assay	No Detectable Microbial DNA	PASS
Bacterial Endotoxin	LAL - USP <85>	<=0.05 EU/mL	PASS
Heavy Metals	ICP-MS	Below reporting limit	PASS
Residual solvents	Headspace GC	Not detected	PASS
Reconstitution and Clarity	Visual assessment after reconstitution	Clear, colorless; no visible particles	PASS
Fentanyl Screen	Validated targeted LC-MS/MS	Not detected	PASS

RP-HPLC-UV CHROMATOGRAPHIC EVIDENCE



CONTROLLED REPORT STATUS
Evidence-backed panel recorded by RUO Eagle

VERIFY REPORT
Scan to confirm status and issued PDF



Results apply only to the submitted specimens and analyses shown, including the two identified vials; they do not establish lot-wide uniformity. Research use only. Verification authenticates RUO Eagle issuance; it does not establish clinical suitability.