PAS

PREMIER ANALYTICAL SCIENCES

REPORT NO.

PAS-2026-1673

ISSUED 2026-11-20

PAGE 1 OF 1

REVISION 0 — CURRENT

CLIENT

4Ever Research

CHAIN

RECEIVED 2026-11-10

CONDITION — INTACT, SEALED

ANALYZED 2026-11-18

RELEASED 2026-11-20 · 12:03 PM PST

BASIS — PER VIAL

METHODS & LIMITS

a · Fluorometric — Fluorometric assay (peptide content) (SOP-13); LOQ 20 µg/mL

b · HPLC-UV / ESI-QTOF-MS — USP/NF 621 Chromatography (SOP-13); LOQ 1 µg/mL

ACCEPTANCE

Purity: 97 % or higher.

Content: 50 to 150 % of the label claim.

Identity: conforms to the reference standard.

VERIFY

https://premieranalytical.com/verify/PAS-2026-1673

CERTIFICATE OF ANALYSIS

INDEPENDENT ANALYTICAL LABORATORY · MISSION HILLS, CALIFORNIA

Retatrutide

CAS 2381089-83-2 · LABEL CLAIM 30 MG · CLIENT LOT RETA 30MG · CLIENT DESIGNATION "RETA 30MG" · 1 VIAL, NUMBERED · MATERIAL AS STATED BY CLIENT

1 OF 1 SAMPLES PASS Purity 99.06 %, content 69.8 % of the stated label claim across 1 sample, measured per vial by the methods keyed in the rail.

SAMPLE 01 · LAB NO. PAS-26-01698 · LOT RETA 30MG · RECEIVED INTACT, SEALED · DESIGNATION "RETA 30MG" **PASS**

PURITY	% OF LABEL	CONTENT	IDENTITY
99.06 %	69.8 %	20.93 mg	CONFORMS
HPLC-UV / ESI-QTOF-MS[b]	of 30 mg claimed	Fluorometric[a]	HPLC-UV / ESI-QTOF-MS[b]



PHOTO 1 — AS RECEIVED

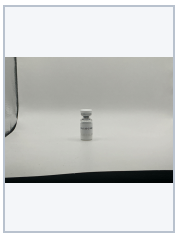


PHOTO 2 — LAB NO. AFFIXED

Released by — Michael Khoury, Principal Chemist

released electronically; recorded on the verification register

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REVISION 0 · STATUS: CURRENT

PAS-2026-1673 · SHA-256 BD81 6F6A DC47 6A4E 8E05 … · VERIFY: https://premieranalytical.com/verify/PAS-2026-1673 · PAGE 1 OF 1

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/// END OF REPORT ///