CERTIFICATE OF ANALYSIS

PuriLab Analytical

Retatrutide 40 mg

RT-2026-086 Lot PP-NOV26-RT-6835 Analyzed November 6, 2026



TIMELINE

ORDERED
October 25, 2026

RECEIVED
November 4, 2026

ANALYZED
November 6, 2026

SAMPLE

NAME
Retatrutide 40 mg

EXPECTED
Retatrutide

AMOUNT
1 vial, lyophilized powder

TESTING

CLIENT
PEP PLUG UK

TEST TYPE
Content & purity

METHOD
HPLC-UV

CONTENT, HOW MUCH IS IN IT

Not detected

Label claim: 40 mg

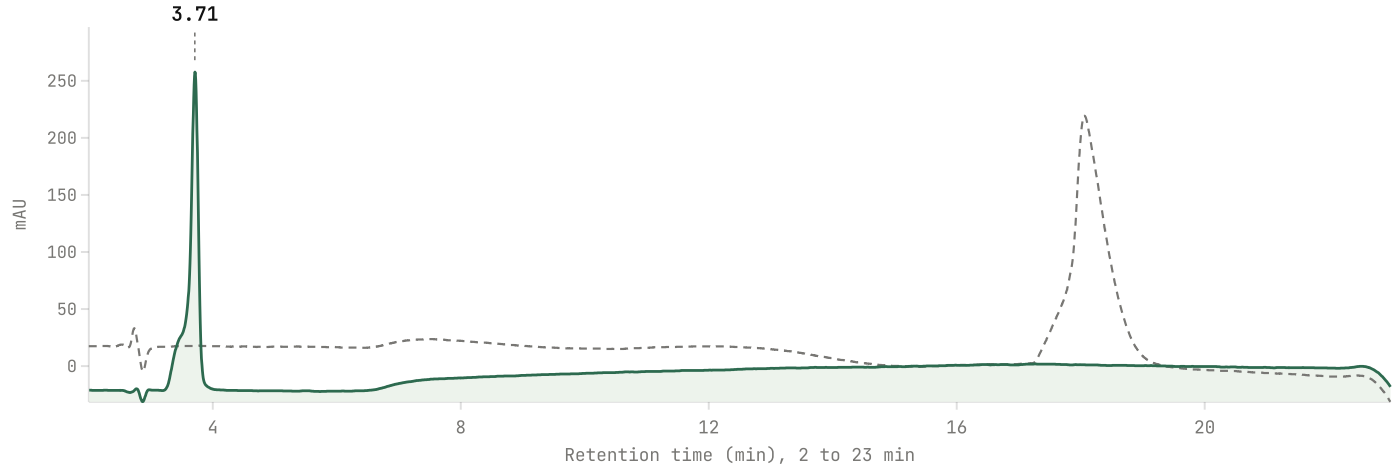
IDENTITY, WHAT IT IS

Not detected

No retatrutide detected

CHROMATOGRAM

HPLC-UV



● 3.71 min, Unidentified component --- Retatrutide reference, same method

ANALYST COMMENTS

White lyophilized powder, purple-cap vial, received sealed. Retatrutide was not found in the sample (detection limit about 0.2 mg per vial). The dashed trace is a retatrutide reference run on the same method, marking where retatrutide elutes. The only significant peak is an early eluting unidentified UV-absorbing component.

ANALYST SIGNATURE

Max Barrett
Lab Director, PuriLab Analytical
Analysis date: November 6, 2026

VERIFICATION CODE

RT-F8BH-FPV9

Verify at purilab.org/verify/RT-F8BH-FPV9

SCAN TO VERIFY

RT-F8BH-FPV9

Results apply solely to the sample as received and are not representative of any larger lot, batch, or production run. Findings are limited to the analytes stated above. PuriLab Analytical is an independent testing laboratory, is not CLIA certified or ISO 17025 accredited, does not sell or distribute any substance, and makes no representation as to the safety, legality, or fitness of any material for any use.