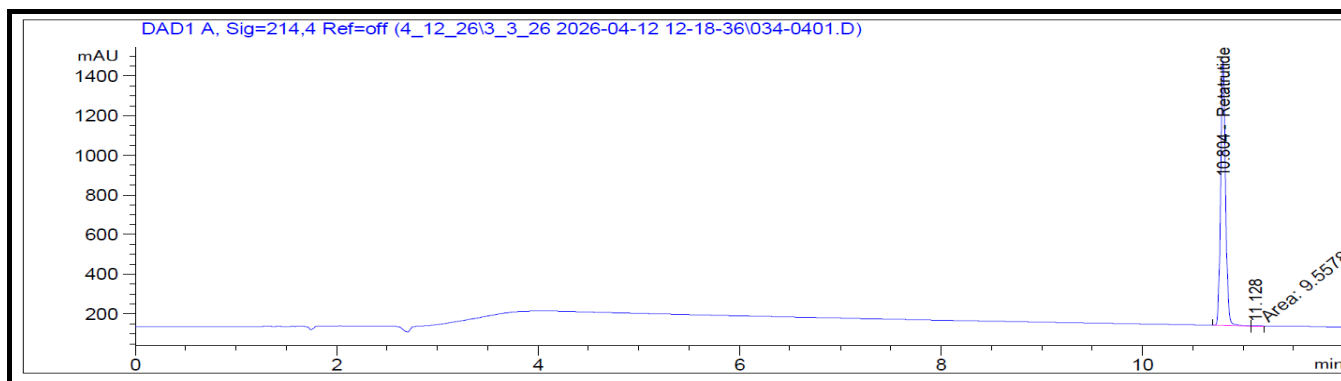


Certificate of Analysis

Retatrutide 10 mg

Report To:
AmiNexa

Compound: Retatrutide
Amount: 10 mg
Laboratory ID: V260402-3 001
Date Reported: 4/14/2026
Lot Number: White Cap



Analysis	Method	Result
Chromatographic Purity	HPLC-UV/VIS	99.79% + 0.18%
Assay	HPLC-UV/VIS	8.64 mg
Endotoxins	LAL	Pass - <5.00 EU/mg*

*Pass/Fail criteria based on the USP/FDA threshold of 5 EU/kg (350 EU total for a 70 kg adult)




Report By: Dustin Newman, Laboratory Director on 4/14/2026



Approved By: Tori Johnson, Operations Manager on 4/14/2026

Please consult A2LA Certificate #6377.01.01 for a list of accredited tests. Samples were received in acceptable condition. The result(s) in this report relate only to the portion of the sample(s) tested. All analyses were performed consistent with the Vanguard Laboratory Quality Management System. Vanguard Laboratory and its staff did not observe or participate in the sample selection process, and cannot confirm the authenticity of the sample or its representativeness of the associated lot/batch.