



Test Report/ Certificate of Analysis

Date of Report May 21, 2026
To: Azoth
Santa Monica, CA
From: AxisPharm, LLC
Certificate ID Number: 21May2026-01

Results reported by: Lab Technician
Ning Yang

Review/Approved by: 
QA Director
Dawei Yue

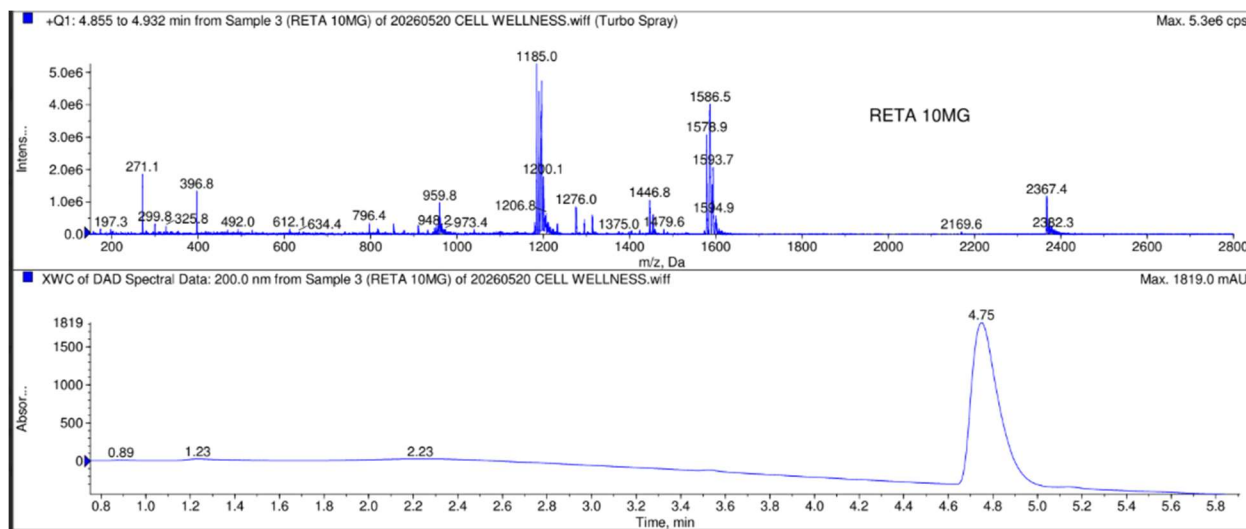
Sample Received: Date: 05/19/2026
Receiving Condition: ☒ Acceptable ☐ Other: _____
Storage Condition: ☐ Room temperature ☒ -20°C ☐ -80°C
☐ Other: _____

Sample Name: Retatrutide 10mg; Brand Name: Azoth ; Lot#: RET10VAS0526; MFD: 05/2026; BBD: 05/2028

Method(s): LCMS and LAL Endotoxin Assay

Procedure(s): Laboratory Activity Date: 05/08/2026

Results:



AxisPharm, LLC

Results only apply to tested Research Use Only samples within Test Report as received. Not for use in humans or animals. Analytical data apply only to the batch tested and do not indicate suitability for diagnostic or therapeutic use. Any unauthorized alteration, reproduction, or relabeling of this COA renders it invalid.



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Sample Name	Net Weight Measured	MW (g/mol)	Detected	UV Purity (208nm) (%)
Retatrutide 10mg	10.4mg	4731.33	2367.4[M+2H], 1586.5[M+3H]	>99.3

Endotoxin Results:

Sample Name	Endotoxin Testing Value (EU/mg)
Retatrutide 10mg	0.053

Salts and sugars that serves as fillers in injectables are not considered as impurities.

Conclusion: The product comes with Retatrutide 10.4mg at purity >99.3%.

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