Axiom ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL + 1 ADD-ON LAYER (L8)

Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US

SCAN TO VERIFY

AX-2026-1028-FEJH

Retatrutide 20 mg · Lot L-RET-018
Issued Oct 17, 2026 PT · v1
Verify: 1984-F842-585D

PASS — ALL SPECIFICATIONS MET

5 of 5 layers complete

L1	L2	L3	L4	L8
Identity	Purity	Net Peptide Content	Sterility	Fentanyl Screen

Sample & Vial Validation



Vial A: as received & analyzed

MORE VIEWS OF VIAL A



Vial A · right

Brand	**Research Vault Group**	Product	**Retatrutide 20 mg**
Lot / Batch	**L-RET-018**	Label Claim	**20 mg**
Matrix	**Lyophilized powder**	Sample Count	**1**
Vial Size	**3mL**	Cap Color	**Translucent Pink**
Crimp Color	**Silver / Aluminum**	Contents Color	**White**
Seal Intact	**Yes**	Condition	**sealed, intact**
Received	**Oct 9, 2026 PT**	Sampling	**client submitted**

Test Results

Test	Specification	Result	
Identity research screen	every vial within ±1 Da of 4731.421 Da	4731.478 Da	PASS
Purity research screen	≥ 99.25 area %	99.4%	PASS
Net Peptide Content research screen	every vial within ±10% of 20 mg	Vial A: 21.1 mg	PASS
Sterility · Rapid PCR research screen	Rapid PCR screen	Target DNA not detected	PASS
Fentanyl Screen research screen	no fentanyl or analog at or above 1 ng/mg	none detected	PASS

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.

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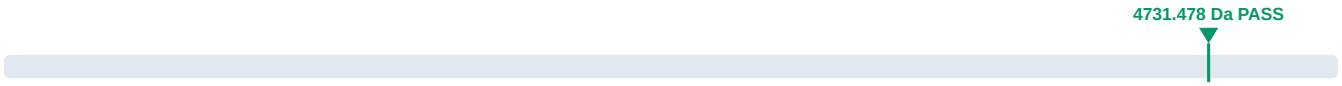
Layer Results

1 Identity

LC-MS (ESI+)

4731.478 Da — every vial within ±1 Da of 4731.421 Da — PASS

Theoretical mass	4731.421 Da	Acceptance	within ±1 Da
Vial A	4731.478 Da · PASS		



2 Purity

RP-HPLC 214nm (USP <621>)

99.4% — ≥ 99.25 area % — PASS

Acceptance	≥ 99.25%	Vial A	99.4% · PASS
Criterion source	Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31		

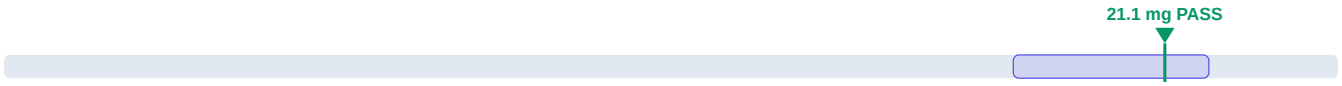


3 Net Peptide Content

Quantitative HPLC vs reference std

Vial A: 21.1 mg — every vial within ±10% of 20 mg — PASS

Target	20 mg (±10%)	Vial A	21.1 mg · PASS
Confirmed container volume	Unknown	Content concentration	Unknown



4 Sterility · Rapid PCR

Rapid Microbial DNA Screen (qPCR target panel)

Target DNA not detected — Rapid PCR screen — PASS

Result	Target DNA not detected	Acceptance	Rapid PCR screen
Target panel	Pan-bacterial 16S rRNA gene (VIC), Pan-fungal 18S rRNA gene (FAM)		

Run controls valid.

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8 Fentanyl Screen

LC-MS/MS targeted adulterant screen

none detected
no fentanyl or analog at or above 1 ng/mg

PASS

Result	**Not detected**	Detection limit	**1 ng/mg**
Analytes screened	**Fentanyl, Acetylfentanyl, Acrylfentanyl, Butyrylfentanyl, Carfentanil, Cyclopropylfentanyl, Furanylfentanyl, Methoxyacetylfentanyl, ortho-Fluorofentanyl, Valerylfentanyl, 4-ANPP (despropionyl fentanyl)**	Source container	**A**

Reporting Scope & Limitations

`[Sterility · Rapid PCR]` DNA-target research screen only. It detects microbial DNA from viable and non-viable organisms alike, and is neither viable-CFU enumeration under USP <61> nor a batch sterility determination under USP <71>.

`[Fentanyl Screen]` Targeted LC-MS/MS adulterant screen, reported as detected or not detected against its stated detection limit. A harm-reduction research screen, not an exhaustive adulterant panel.

Digital Signatures

Analyst	**Alana Reeves**	Analyst Signature	**ed25519:3d59fd14da67bf68e9d2088c...**
Technical Reviewer	**Michael Niles**	Reviewer Signature	**ed25519:2c15f824034bc2bb712b5ece...**
Laboratory Director	**Daniel Brennan**	Director Signature	**ed25519:323313eb4bc26c04063306d5...**

Where physical instrument analysis is performed, it takes place within Axiom Analytics' laboratory network, allocated at Axiom's discretion based on method, capacity, and order volume. Axiom Analytics performs sample accession, chain of custody, analytical oversight, and data review, and issues and is responsible for this certificate. Where analysis is subcontracted, the method is performed within the partner laboratory's published ISO/IEC 17025 scope of accreditation for that method, not merely at a laboratory holding ISO/IEC 17025 accreditation. Axiom Analytics does not itself currently hold ISO/IEC 17025 accreditation. The method-validation status of each reported result is identified beside that result. Partner laboratory identities are not disclosed: where samples are analyzed under double-blind conditions, they are identified only by an internal lot ID.

Each signature is an ed25519 signature over the SHA-384 hash of this report. Scan the QR on the letterhead or verify independently at https://axiomanalyticslab.com/verify/AX-2026-1028-FEJH.

✓ **CRYPTOGRAPHICALLY SEALED (ed25519 + SHA-384)**

`SHA-384: 22eeeddd464ac5229fda9b4d33d91313a13fb7bbc5f45d7a1551c2a868fb724c...`

`Seal: ed25519:44d6f6f52be0ac623ad7a22c72fe4e2baabb1cb3...`

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