Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL

**Axiom Analytics Labs LLC**
Sacramento, California, USA

## AX-2026-3426-KRNM

GL-3 RT (Retatrutide) · Lot GL-RT30MG-04-202610-001
Issued Oct 22, 2026 PT · v1
Verify: A577-3AB5-618F

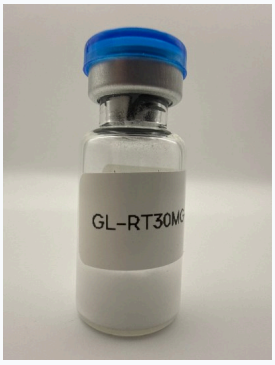
SCAN TO VERIFY

**PASS — ALL SPECIFICATIONS MET**

4 of 4 layers complete

| L1 | L2 | L3 | L4 |
|---|---|---|---|
| Identity | Purity | Net Peptide Content | Sterility |

### Sample & Vial Validation




Vial A: as received & analyzed

MORE VIEWS OF VIAL A



Vial A · right

| | | | |
|---|---|---|---|
| Brand | **Guardian Labs** | Product | **GL-3 RT (Retatrutide)** |
| Lot / Batch | **GL-RT30MG-04-202610-001** | Label Claim | **30 mg** |
| Matrix | **lyophilized powder** | Sample Count | **1** |
| Vial Size | **3mL** | Cap Color | **Translucent Blue** |
| Crimp Color | **Silver / Aluminum** | Contents Color | **White** |
| Seal Intact | **Yes** | Condition | **sealed, intact** |
| Received | **Oct 21, 2026 PT** | Sampling | **client submitted** |

### Test Results

| Test | Specification | Result | |
|---|---|---|---|
| **Identity**<br>research screen | every vial within ±1 Da of 4731.421 Da | `4731.417 Da` | PASS |
| **Purity**<br>research screen | ≥ 99.25 area % | `99.97%` | PASS |
| **Net Peptide Content**<br>research screen | every vial within ±10% of 30 mg | `Vial A: 31.93 mg` | PASS |
| **Sterility · Rapid PCR**<br>research screen | Rapid PCR screen | `Target DNA not 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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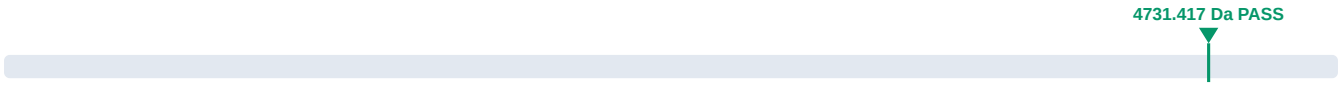
## Layer Results

### 1 Identity
LC-MS (ESI+)

**4731.417 Da** — PASS
every vial within ±1 Da of 4731.421 Da

| | | | |
|---|---|---|---|
| Theoretical mass | **4731.421 Da** | Acceptance | **within ±1 Da** |
| Vial A | **4731.417 Da · PASS** | | |




### 2 Purity
RP-HPLC 214nm (USP <621>)

**99.97%** — PASS
≥ 99.25 area %

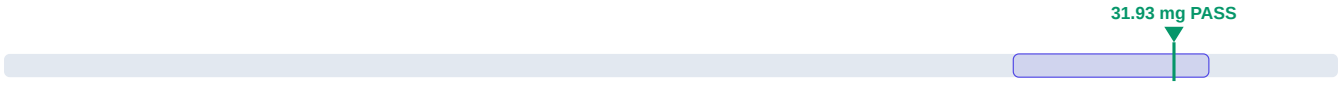
| | | | |
|---|---|---|---|
| Acceptance | **≥ 99.25%** | Vial A | **99.97% · PASS** |
| Criterion source | **Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31** | | |




### 3 Net Peptide Content
Quantitative HPLC vs reference std

**Vial A: 31.93 mg** — PASS
every vial within ±10% of 30 mg

| | | | |
|---|---|---|---|
| Target | **30 mg (±10%)** | Vial A | **31.93 mg · PASS** |




### 4 Sterility · Rapid PCR
Rapid Microbial DNA Screen (qPCR target panel)

**Target DNA not detected** — PASS
Rapid PCR screen

| | |
|---|---|
| Target panel | **Pan-bacterial 16S rRNA gene (VIC), Pan-fungal 18S rRNA gene (FAM)** |

`Run controls valid.`

## 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>.

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## Digital Signatures

| | | | |
|---|---|---|---|
| Analyst | **Alana Reeves** | Analyst Signature | **ed25519:25913afe1d30d73c756580e7...** |
| Technical Reviewer | **Michael Niles** | Reviewer Signature | **ed25519:97ed0c558fc94291e4ac4aaa...** |
| Laboratory Director | **Daniel Brennan** | Director Signature | **ed25519:ad39fd59a46d94221348dc18...** |

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-2026-3426-KRNM.

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

SHA-384: 970dcd41b83017f64bd0a21ab54cd238172dd7d7351571e921856e9693ecfd22...

Seal: ed25519:706e5e0d703bdcf4980937c92b678ac0989f56a3...

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