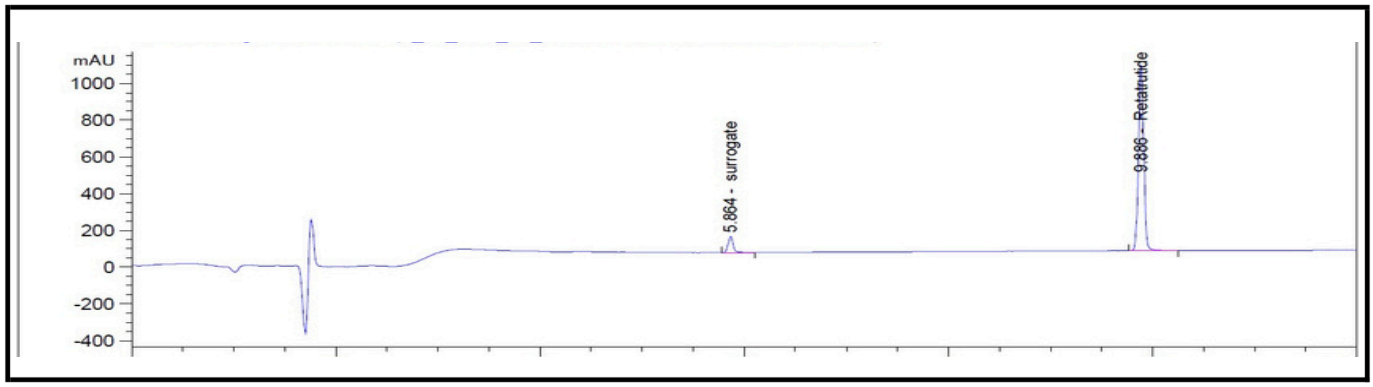
Kalbio Biotechnology
20503 Crescent Bay
Dr Lake Forest, Ca
92630 United States

# Certificate of Analysis

GLP3-RT 20 mg

Report For: Luxora

Compound: GLP3-RT
Quantity:20 mg
Batch# 80426-RETA20
Lot Number:335-RETA20
Manufactured Date: 08/06/2026
Expiration:06/06/2028




| Analysis | Method | Result (per Vial) | | |
|---|---|---|---|---|
| Chromatographic Purity | HPLC-UV/VIS | >99.80% | >99.80% | >99.80% |
| Quantity | HPLC-UV/VIS | 21.56 mg | 19.84 mg | 20.06 mg |

Report By: Daniel Myers, Laboratory Director on 08/21/2026

Approved By: Esteban Solis, Operations Manager on 08/21/2026

ISO 17025:2017 ACCREDITED LABORATORY

Please consult 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 Kalbio Biotechnology Quality Control. Kalbio 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.

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Batch# 80426-RETA20
Lot Number:335-RETA20
Manufactured Date: 08/06/2026
Expiration:06/06/2028

| Analyte | Method | LOQ (ppm) | Result (ppm) |
|---|---|---|---|
| Arsenic (As) | ICP-MS | 0.01 | ND |
| Cadmium (Cd) | ICP-MS | 0.01 | ND |
| Lead (Pb) | ICP-MS | 0.01 | ND |
| Mercury (Hg) | ICP-MS | 0.005 | ND |

RunID: 260224

| Analysis | Method | Result |
|---|---|---|
| Endotoxins | LAL | Pass - <5.00 EU/mg* |
| BioBurden | USP <61> | Pass - No GrowthDetected |

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

Report By: Daniel Myers, Laboratory Director on 08/21/2026

Approved By: Esteban Solis, Operations Manager on 08/21/2026

*ND: Non-Detect*
*LOD: Limit of Quantification*

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