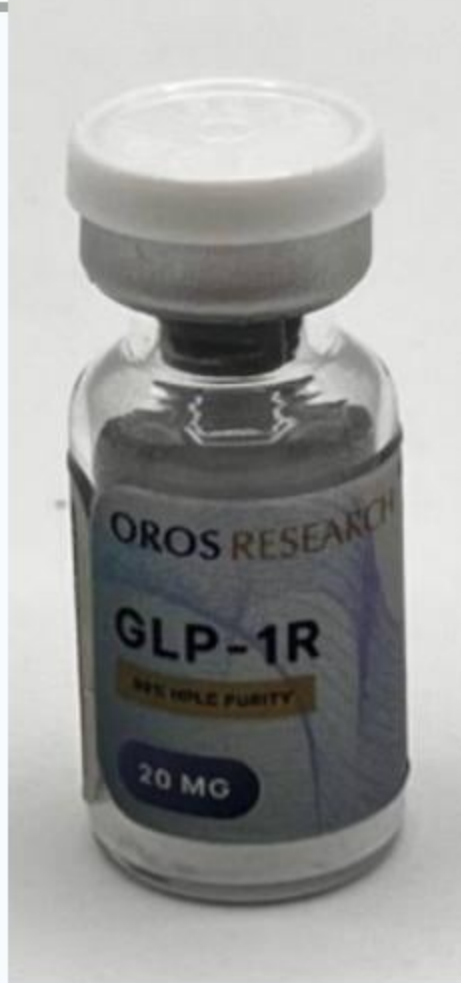


Summary of Results

| Analysis Type        | Method         | Date Tested | Status   |  |
|----------------------|----------------|-------------|----------|-------------------------------------------------------------------------------------|
| Peptides             | HPLC           | 04/09/2026  | Complete |                                                                                     |
| Bacterial Endotoxins | USP <85>       | 04/09/2026  | Complete |                                                                                     |
| Heavy Metals         | ICP-MS         | 04/07/2026  | Pass     |                                                                                     |
| Microbiological      | Petrifilm/qPCR | 04/08/2026  | Pass     |                                                                                     |
|                      |                |             |          |                                                                                     |
|                      |                |             |          |                                                                                     |
|                      |                |             |          |                                                                                     |

Serving Size: N/A

Peptide Analysis

| Analyte                                                                                                                 | Result       | Specification | LOD                        |
|-------------------------------------------------------------------------------------------------------------------------|--------------|---------------|----------------------------|
| Retatrutide                                                                                                             | 20.86mg/vial | 20mg/vial     | 0.1%                       |
| Identity of the analyte was confirmed by retention time concordance with a certified reference standard, per USP <621>. |              |               |                            |
| Chromatographic Purity                                                                                                  | >99%         | >99%          | 0.1%                       |
| Bacterial Endotoxins<br>(Gel-Clot: 1:10 Dilution)                                                                       | <1.25 EU/ml  | <5 EU/ml      | $\lambda =$<br>0.125 EU/mL |
|                                                                                                                         |              |               |                            |
|                                                                                                                         |              |               |                            |
|                                                                                                                         |              |               |                            |
|                                                                                                                         |              |               |                            |
|                                                                                                                         |              |               |                            |
|                                                                                                                         |              |               |                            |

NOTES: Injectable peptides may include acceptable salts, sugars, or buffering agents added to improve solubility and stability. These excipients are non-chromophoric, typically not detected by HPLC, and are not considered impurities. Two vials were tested for variability, and the average result is reported above.

LOD = LIMIT OF DETECTION; LOQ = LIMIT OF QUANTIFICATION

The sample analyzed was inspected and is free from visual mold, mildew, and foreign matter. The testing procedures, equipment calibration, and maintenance are all in accordance with ISO/IEC 17025:2017 standards. The presented report is only applicable to the sample specified above and may not be applied to any similar or identical products. Reports are prohibited from being reproduced with alterations of any kind.

Heavy Metal Analysis (USP <233>)

| Analyte | Result<br>(mcg/vial) | LOQ (ppm) | LOD (ppm) | Limit<br>(mcg/day) | Pass/Fail |
|---------|----------------------|-----------|-----------|--------------------|-----------|
| Arsenic | 0.003                | 0.010     | 0.005     | 0.5                | Pass      |
| Cadmium | <LOQ                 | 0.010     | 0.005     | 0.1                | Pass      |
| Lead    | 0.006                | 0.010     | 0.005     | 0.5                | Pass      |
| Mercury | <LOQ                 | 0.010     | 0.005     | 0.1                | Pass      |

Microbiological Analysis (USP <61>, USP <62>)

| Microbe                              | Result      | Limit       | Pass/Fail |
|--------------------------------------|-------------|-------------|-----------|
| Total Aerobic Plate                  | <10 CFU/g   | <100 CFU/g  | Pass      |
| Coliform                             | <10 CFU/g   | <10 CFU/g   | Pass      |
| Bile-Tolerant Gram-Negative Bacteria | <10 CFU/g   | <10 CFU/g   | Pass      |
| Yeast and Mold                       | <10 CFU/g   | <10 CFU/g   | Pass      |
| Escherichia Coli                     | Negative/1g | Negative/1g | Pass      |
| Salmonella                           | Negative/1g | Negative/1g | Pass      |
| Staphylococcus Aureus                | Negative/1g | Negative/1g | Pass      |

NOTES:

CFU = Colony Forming Unit  
NS = Not Specified  
NT = Not Tested

LOQ = Limit of Quantification  
LOD = Limit of Detection

The sample analyzed was inspected and is free from visual mold, mildew, and foreign matter. The testing procedures, equipment calibration, and maintenance are all in accordance with ISO/IEC 17025:2017 standards. The presented report is only applicable to the sample specified above and may not be applied to any similar or identical products. Reports are prohibited from being reproduced with alterations of any kind.