

CERTIFICATE OF ANALYSIS

Sermorelin Acetate - 5 mg

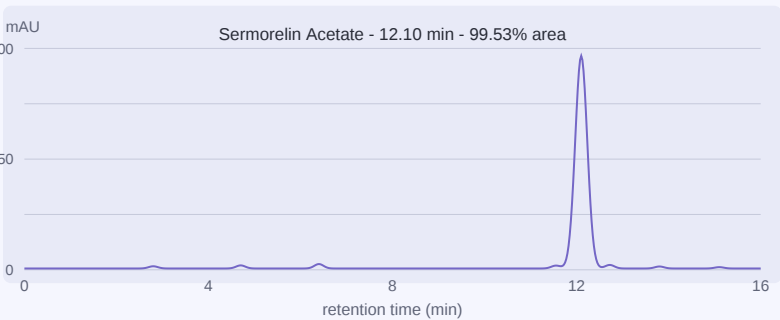
GHRH(1-29) amide acetate - synthetic peptide - lyophilised

|             |                    |             |                   |          |                          |
|-------------|--------------------|-------------|-------------------|----------|--------------------------|
| Material    | Sermorelin Acetate | Client      | novumpeptides.com | Received | 2026-07-15 at 2-8 C      |
| Batch / lot | PEP-32-7ET         | Label claim | 5 mg              | Analysed | 2026-07-16 to 2026-07-17 |

TEST RESULTS

| ATTRIBUTE                   | RESULT   | SPECIFICATION     | METHOD                         | VERDICT  |
|-----------------------------|----------|-------------------|--------------------------------|----------|
| Identity                    | Conforms | Matches reference | MA-MS-014 / LC-HRMS            | Conforms |
| Purity (area %)             | 99.53%   | >= 99.00%         | MA-UHPLC-031 / UV 214          | Pass     |
| Largest single impurity     | 0.14%    | <= 0.50%          | MA-UHPLC-031 / UV 214          | Pass     |
| Total related substances    | 0.47%    | <= 1.00%          | MA-UHPLC-031 / UV 214          | Pass     |
| Sermorelin Acetate per vial | 5.00 mg  | 4.50-5.50 mg      | MA-ASSAY-011 / UHPLC-UV        | Pass     |
| Water content               | 3.87%    | <= 8.00%          | MA-KF-004 / Karl Fischer       | Pass     |
| Residual acetonitrile       | 158 ppm  | <= 410 ppm        | MA-GC-006 / HS-GC-FID          | Pass     |
| Residual TFA                | 0.10%    | <= 1.00%          | MA-IC-003 / ion chromatography | Pass     |

RP-UHPLC processed trace, UV detection



Record F3KU7ET-PEP-01; duplicate independent preparations. System suitability met retention-time and resolution criteria; blank response was below threshold.

Related species > 0.05%

|                        |       |
|------------------------|-------|
| deletion sequence      | 0.14% |
| insertion sequence     | 0.10% |
| diastereomeric variant | 0.08% |
| N-terminal variant     | 0.06% |
| C-terminal variant     | 0.04% |
| unresolved peaks       | 0.05% |

Related species were assigned by orthogonal reference comparison and mass-spectrometric or chromatographic evidence. Mean area percentages; reporting threshold 0.05%.

Conclusion

Identity and quality attributes met the stated criteria. Mean Sermorelin Acetate content was 5.00 mg/vial and chromatographic purity was 99.53%.

Identity was established by LC-HRMS against a qualified reference; content and related substances were assessed independently.

Analytical review

Ines Abadi I. Abadi

Quality review

Marcus Oyelaran

Approval date: 2026-07-18 M. Oyelaran

Decision rule: guarded acceptance; reported outcomes account for method uncertainty and remain within the stated project specifications.