

ILS Laboratories
8222 Vickers St, Suite 106, San Diego, CA 92111
(619) 329-3999 | ils-lab.com

Semaglutide - 10mg

Tested for: Elevate

PASS

COA #: **COA-2026-954740**
Lot Number: **SM10-2026-001**
Accession #: **ACC-2026-13751**
Labeled Content: **10mg**

Analysis Date: **08/01/2026**
Appearance: **Good**
Sample Matrix: **Lyophilized**
Volume: **3mL**
Date Received: **08/01/2026**



Scan to verify
authenticity at ils-lab.com
Access Code: QHVTG3R

Method: **Full QC Panel**

Identity	Peptide Purity	
Semaglutide	99.51%	 Fentanyl Free

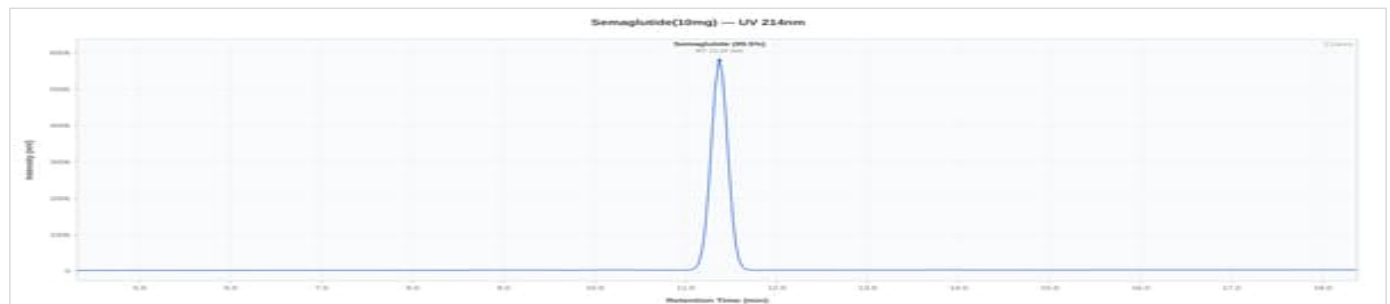
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.51%	%	PASS
Net Peptide Content	Report Only	10.37	mg	N/A
Identity (HPLC-RTM)	Semaglutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



Semaglutide 10mg - SM10-2026-001

HPLC Chromatogram



Semaglutide 10mg - SM10-2026-001: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	<i>Not Detected</i>	PASS
Cadmium (Cd)	NMT 0.5 ppm	<i>Not Detected</i>	PASS
Lead (Pb)	NMT 1.0 ppm	<i>Not Detected</i>	PASS
Mercury (Hg)	NMT 1.5 ppm	<i>Not Detected</i>	PASS
Chromium (Cr)	NMT 10.0 ppm	<i>Not Detected</i>	PASS



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Dr. Greg Kalyuzhny
Lab Director
8/1/2026

COA #: **COA-2026-954740**
Access Code: **QHVTG3R**
Verify: <portal.ils-lab.com/verify/QazdVX2JgD5HfPmc>
Issued: 8/1/2026

Sterility Testing (PCR)

Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	NMT 0.05 EU/mL	Reported

About this result: Endotoxin is reported as a quantitative value. Acceptable limits vary by product type and matrix, so no universal pass/fail threshold applies to RUO products.

Notes & Methodology

- 1. Date Tested: 08/01/2026. Methods: Full QC Panel.
- 2. The sample was confirmed to be Semaglutide by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- 3. Elemental impurities analyzed by ICP-MS per USP <233> methodology. Acceptance criteria are internal laboratory quality screening limits for research-use materials and do not represent evaluation against any specific pharmacopeial monograph or product specification.
- 4. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
- 5. Peptide purity determined by RP-HPLC area normalization at 214 nm. Value represents the percentage of the target peptide relative to all peptide-related peaks. Non-peptide process-related impurities, if detected, are excluded from the calculation.




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