

ILS Laboratories

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

GLOW - 70mg

Tested for: Regenerative Research

<https://www.regenerativeresearch.co/>

COA #: COA-2026-CXBIL
Lot Number: GW26-03
Accession #: ACC-2026-9667
Labeled Content: 70mg

Analysis Date: 07/31/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 07/13/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: 2ER9MSPN

Method: Full QC Panel

Identity
GLOW

Peptide Purity
99.88%



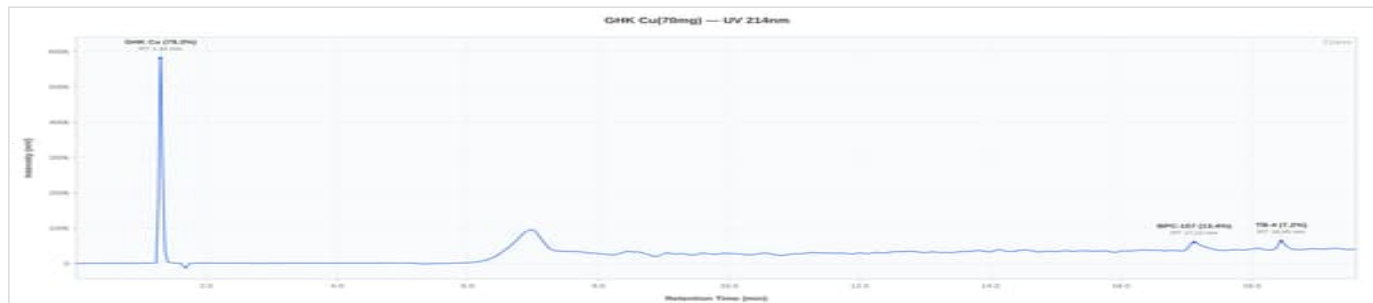
Blend Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.88%	%	PASS
Net Blend Peptide Content	Report Only	72.6	mg	N/A
-- GHK-Cu		52.27	mg	
-- BPC-157		10.31	mg	
-- TB-4		10.02	mg	
Identity (HPLC-RTM)	GHK-CU + BPC-157 + TB-4	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



GLOW 70mg - GW26-03

HPLC Chromatogram



Representative chromatogram, Conformity V1

HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.88%	72.6 mg	Confirmed	PASS
Conformity V1	99.87%	73.96 mg	Confirmed	PASS
Conformity V2	99.87%	71.94 mg	Confirmed	PASS
Mean	99.87%	72.83 mg	—	—
Std Dev	0.0047%	0.8410 mg	—	—



Dr. Greg Kalyuzhny

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Lab Director
7/31/2026

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Verify: <portal.ils-lab.com/verify/6cgpjTYI9Wv7n0S>
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Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	.963	PASS
Cadmium (Cd)	NMT 0.5 ppm	.336	PASS
Lead (Pb)	NMT 1.0 ppm	.582	PASS
Mercury (Hg)	NMT 1.5 ppm	.2247	PASS
Chromium (Cr)	NMT 10.0 ppm	6.32	PASS

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	0.124 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. This result is below commonly referenced endotoxin thresholds.

Notes & Methodology

1. Date Tested: 07/31/2026. Methods: Blend Purity, Identity & Quantitation (HPLC).
2. The sample was confirmed to be GLOW 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.
6. Per-component content calculated from total net peptide content using the manufacturer's stated formulation ratio (GHK-Cu, BPC-157, TB-4). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
7. Chromatogram shown is representative: Conformity V1 (99.87% purity, closest to batch mean of 99.87%).




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