

ILS Laboratories

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

GLOW - 70mg

PASS

Tested for: Vanta

Vantapeps.com

COA #: COA-2026-OFICJF
Lot Number: GLW0001
Accession #: ACC-2026-1478
Concentration: 70mg
Sample Matrix: Lyophilized

Method: Full QC Panel
Analysis Date: 07/01/2026
Appearance: Good
Volume: 3mL
Received: 07/01/2026

Identity

GLOW

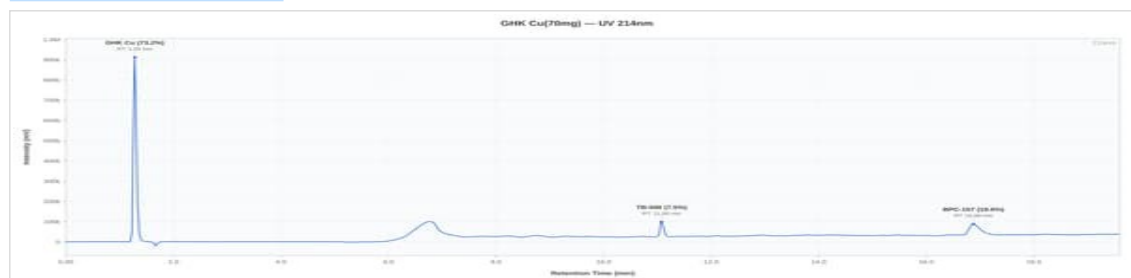
Peptide Purity

99.77%

Blend Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.77%	%	PASS
Net Blend Peptide Content	Report Only	73.27	mg	N/A
-- GHK-Cu		52.75	mg	
-- BPC-157		10.40	mg	
-- TB-4		10.11	mg	
Identity (ID)	GHK-CU + BPC-157 + TB-4	Confirmed	-	PASS

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.77% purity, closest to batch mean of 99.77%)



Certificate of Analysis

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HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.77%	73.27 mg	Confirmed	PASS
Conformity V1	99.76%	72.21 mg	Confirmed	PASS
Mean	99.77%	72.74 mg	—	—
Std Dev	0.0050%	0.5300 mg	—	—

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	Not Detected	PASS
Cadmium (Cd)	NMT 0.5 ppm	Not Detected	PASS
Chromium (Cr)	NMT 10 ppm	Not Detected	PASS
Mercury (Hg)	NMT 1.5 ppm	Not Detected	PASS
Lead (Pb)	NMT 1 ppm	Not Detected	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	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. This result is below commonly referenced endotoxin thresholds.



Dr. Greg Kalyuzhny
Lab Director



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COA #:	COA-2026-OFICJF	Method:	Full QC Panel
Lot Number:	GLW0001	Analysis Date:	07/01/2026
Accession #:	ACC-2026-1478	Appearance:	Good
Concentration:	70mg	Volume:	3mL
Sample Matrix:	Lyophilized	Received:	07/01/2026

Notes & Methodology

1. Methods: Blend Purity & Quant (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: Dedicated V0 (99.77% purity, closest to batch mean of 99.77%).



Dr. Greg Kalyuzhny
Lab Director