Certificate of Analysis

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

GLOW - 70mg

Tested for: Bandlien

PASS

Scan to verify authenticity at ils-lab.com
Access Code: 3W88R9TD

COA #:	**COA-2026-956508**	Analysis Date:	**08/31/2026**
Lot Number:	**GLOW-26/07-01**	Appearance:	**Good**
Labeled Content:	**70mg**	Volume:	**3mL**
Sample Matrix:	**Lyophilized**	Date Received:	**07/17/2026**

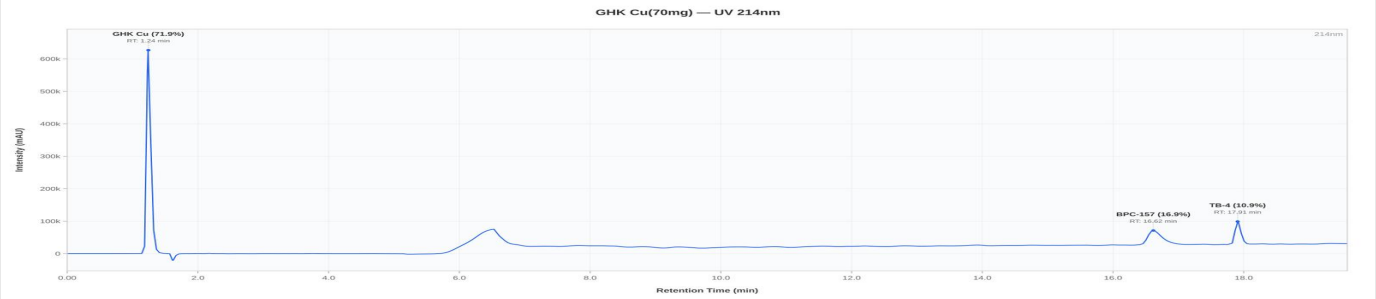
Method: **Full QC Panel**

Identity	Peptide Purity
GLOW	99.67%

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.67%**	%	PASS
Net Blend Peptide Content	Report Only	**72.49**	mg	N/A
-- GHK-Cu		*52.19*	*mg*	
-- BPC-157		*10.29*	*mg*	
-- TB-4		*10.00*	*mg*	
Identity (HPLC-RTM)	GHK-CU + BPC-157 + TB-4	**Confirmed**	-	PASS

HPLC Chromatogram



GLOW 70mg - GLOW-26/07-01: UV Chromatogram

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
Lead (Pb)	NMT 1.0 ppm	*Not Detected*	PASS
Mercury (Hg)	NMT 1.5 ppm	*Not Detected*	PASS
Chromium (Cr)	NMT 10.0 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	**0.071 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: 08/31/2026. Methods: Full QC Panel.
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 chromogenic 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. Accreditation Scope: ILS Laboratories maintains ISO/IEC 17025 accreditation for activities listed in its published PJLA scope. Peptide testing reported on this certificate is performed for Research Use Only.
8. This certificate reports analytical results for the sample(s) submitted to ILS Laboratories for testing. All sample identifications, product names, brand names, lot/batch numbers, and labeled content are provided by the submitting party and have not been independently verified. Results reflect only the analytical testing of the specific sample(s) provided and do not constitute an evaluation or endorsement of the manufacturer's processes, quality systems, or overall production. ILS Laboratories makes no representations regarding the intended use, safety, efficacy, regulatory status, or intellectual property ownership of any materials tested.

ILS LABS

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ils-lab.com

Dr. Greg Kalyuzhny

Lab Director

8/31/2026

COA #: **COA-2026-956508**

Access Code: **3W88R9TD**

Verify: portal.ils-lab.com/verify/_Z_MuUcp0WFAOAd2

Issued: 8/31/2026

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This certificate relates only to the sample(s) tested.