

ILS Laboratories

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(619) 329-3999 | ils-lab.com

GLOW GHK-CU + BPC-157 + TB-500 - 70mg

Tested for: LifeGen Labs
LifeGenLabs.com

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: FCHC38NB

COA #: COA-2026-PZYQUH
Lot Number: 4052
Accession #: ACC-2026-10149
Labeled Content: 70mg

Analysis Date: 08/14/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 07/16/2026

Method: Full QC Panel

Identity

GLOW GHK-CU + BPC-157 + TB-500

Peptide Purity

99.80%



Fentanyl
Free

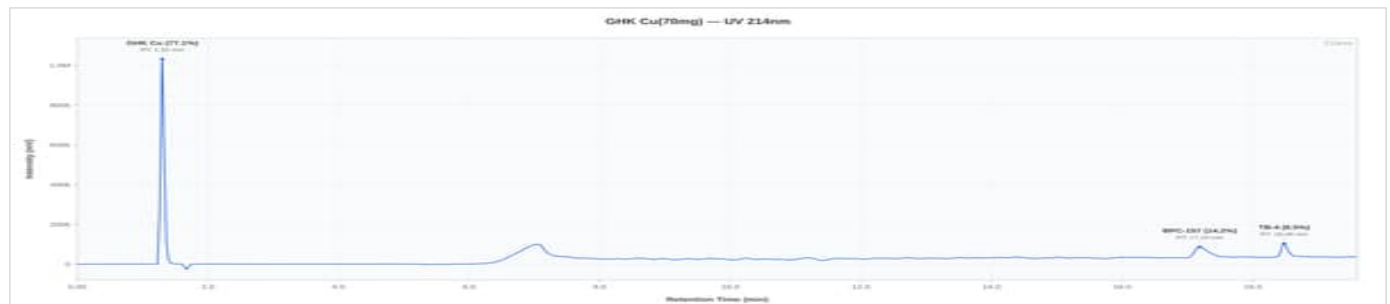
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.80%	%	PASS
Net Blend Peptide Content	Report Only	74.93	mg	N/A
-- BPC-157		38.21	mg	
-- TB-500		36.72	mg	
Identity (HPLC-RTM)	BPC-157 + TB-500	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



GLOW GHK-CU + BPC-157 + TB-500
70mg - 4052

HPLC Chromatogram



GLOW GHK-CU + BPC-157 + TB-500 70mg - 4052: 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



Dr. Greg Kalyuzhny

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Lab Director
8/14/2026

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Verify: <portal.ils-lab.com/verify/Oo6VqnR784xzZGwn>
Issued: 8/14/2026

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>)	NMT .5 EU/mL	0.146 EU/mL	PASS

Acceptance criteria: NMT .5 EU/mL per client specification.

Notes & Methodology

1. Date Tested: 08/14/2026. Methods: Full QC Panel.
2. The sample was confirmed to be GLOW GHK-CU + BPC-157 + TB-500 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 (BPC-157, TB-500). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
7. 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.




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