Certificate of Analysis

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

GLOW - 70mg

5X TESTED

PASS

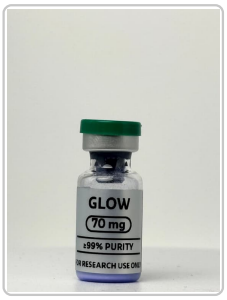
Tested for: HighRock Research Compounds
highrockresearchcompounds.com

COA #:	**COA-2026-XNG7MB**	Analysis Date:	**11/03/2026**
Lot Number:	**26-HR-08**	Cap Color:	**Green**
Labeled Content:	**70mg**	Date Received:	**10/26/2026**
Volume:	**3mL**		

Scan to verify authenticity at ils-lab.com
Access Code: 3R4CX9US

Method: **Blend Purity, Identity & Quantitation (HPLC), Endotoxin (USP <85>)**

Identity	Peptide Purity	
GLOW	**99.73%**	Fentanyl Free

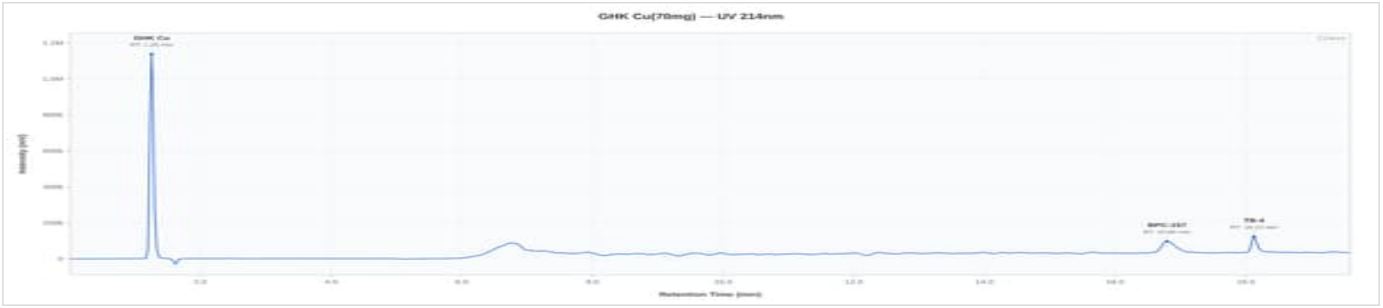


GLOW 70mg - 26-HR-08

Blend Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.73%**	%	PASS
Net Blend Peptide Content	Report Only	**71.82**	mg	N/A
-- BPC-157		*10.12*	*mg*	
-- TB-500		*10.21*	*mg*	
-- GHK-Cu		*51.49*	*mg*	
Identity (HPLC-RTM)	BPC-157 + TB-500 + GHK-CU	**Confirmed**	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	PASS

HPLC Chromatogram



GLOW 70mg - 26-HR-08: UV Chromatogram

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.

Notes & Methodology

1. Date Tested: 11/03/2026. 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. Endotoxin tested per USP <85> kinetic chromogenic method. Acceptance criteria per client specification.
4. 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.
5. Per-component content determined by individual peak integration. Not all labeled components may be present.
6. Quality System: ILS Laboratories operates an ISO/IEC 17025 quality management system. Peptide testing is performed for Research Use Only.
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.

Issued by ILS Laboratories
(619) 329-3999
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Dr. Greg Kalyuzhny
Lab Director
11/03/2026

COA #: **COA-2026-XNG7MB**
Access Code: **3R4CX9US**
Verify: portal.ils-lab.com/verify/x9QEElRjd2GLq3F3
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