Certificate of Analysis

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

GLOW - 70

8x Tested

PASS

Tested for: Axis Research Supply Inc

COA #:	**COA-2026-VPEXCC**	Analysis Date:	**10/17/2026**
Lot Number:	**AX26219-14**	Appearance:	**Good**
Labeled Content:	**70**	Date Received:	**10/08/2026**

Method: **Full QC Panel**

Scan to verify authenticity at ils-lab.com
Access Code: EMFTJB8C

Identity	Peptide Purity	
GLOW	**99.92%**	Fentanyl Free

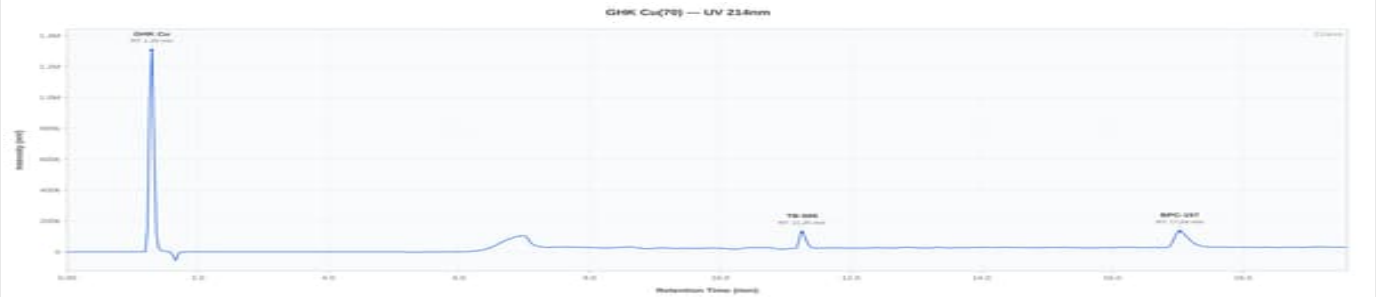


GLOW 70 - AX26219-14

Blend Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.92%**	%	**PASS**
Identity (HPLC-RTM)	GHK-CU + TB-500 + BPC-157	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Dedicated V0

Additional Tests

Analyte	Specification	Result	Unit	Status
Net Blend Peptide Content	Report Only	**74.48**	mg	N/A
-- GHK-CU		*52.99*	*mg*	
-- TB-500		*10.70*	*mg*	
-- BPC-157		*10.79*	*mg*	

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.92%	74.48 mg	Confirmed	**PASS**
Conformity V1	99.83%	73.69 mg	Confirmed	**PASS**
Mean	**99.88%**	**74.09 mg**	—	—
Std Dev	**0.0450%**	**0.3950 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>)	NMT 5.0 EU/mL	**NMT 0.05 EU/mL**	**PASS**

Acceptance criteria: NMT 5.0 EU/mL per client specification.

Notes & Methodology

1. Date Tested: 10/17/2026. Methods: Blend Purity, Identity & Quantitation (HPLC); Additional Tests.
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 determined by individual peak integration. Not all labeled components may be present.
7. Chromatogram shown is representative: Dedicated V0 (99.92% purity, closest to batch mean of 99.88%).
8. 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.
9. 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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Dr. Greg Kalyuzhny
Lab Director
10/17/2026

COA #: **COA-2026-VPEXCC**
Access Code: **EMFTJB8C**
Verify: portal.ils-lab.com/verify/EK6-s8q6mBGz6eDa
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