Certificate of Analysis

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

GLOW 50/10/10MG - 50/10/10mg

Tested for: NEXT LEVEL COMPOUNDS

PASS

COA #:	**COA-2026-413053**	Analysis Date:	**06/23/2026**
Lot Number:	**GLO501010-1001**	Appearance:	**Good**
Labeled Content:	**50/10/10mg**	Sample Matrix:	**Lyophilized**
Sample Matrix:	**Lyophilized**	Vial Size:	**3mL**
		Date Received:	**06/17/2026**

Scan to verify authenticity at ils-lab.com
Access Code: 9EFTBYV5

Method: **Full QC Panel**

Identity	Peptide Purity
GLOW 50/10/10MG	**99.95%**



GLOW 50/10/10MG 50/10/10mg - GLO501010-1001

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.95%**	%	**PASS**
Net Blend Peptide Content	Report Only	**74.42**	mg	N/A
-- GHK-Cu		*53.58*	*mg*	
-- BPC-157		*10.57*	*mg*	
-- TB-4		*10.27*	*mg*	
Identity (HPLC-RTM)	GHK-CU + BPC-157 + TB-4	**Confirmed**	-	**PASS**
Identity (HPLC-RTM)	GHK-CU + BPC-157 + TB-4	**Confirmed**	-	**PASS**

HPLC Chromatogram



GLOW 50/10/10MG 50/10/10mg - GLO501010-1001: 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**
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**

Dr. Greg Kalyuzhny
Lab Director
6/23/2026

COA #: COA-2026-413053
Access Code: **9EFTBYV5**
Verify: portal.ils-lab.com/verify/j13kIvkV7l3fPXMe
Issued: 6/23/2026

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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.

Notes & Methodology

1. Date Tested: 06/23/2026. Methods: Full QC Panel.
2. The sample was confirmed to be GLOW 50/10/10MG 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.

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