

ILS Laboratories

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

KLOW - 80mg



Tested for: NextGen Peptides
www.NGpeptide.com

COA #: COA-2026-QHRTGB
Lot Number: klw06
Accession #: ACC-2026-4533
Labeled Content: 80mg

Analysis Date: 06/25/2026
Appearance: Good
Vial Size: 3mL
Date Received: 06/10/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: P66ZQ6Q5

Method: Full QC Panel

Identity

KLOW

Peptide Purity

99.84%

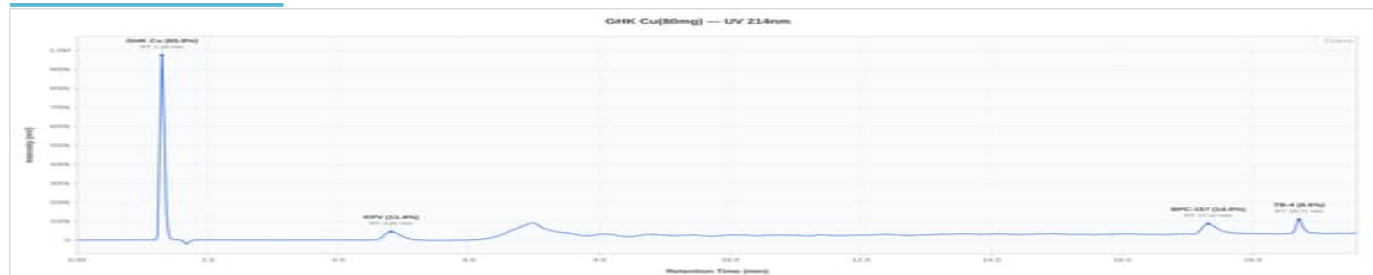


Fentanyl
Free

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.84%	%	PASS
Net Blend Peptide Content	Report Only	85.33	mg	N/A
-- GHK		53.46	mg	
-- BPC-157		10.59	mg	
-- TB-500		10.80	mg	
-- KPV		10.48	mg	
Identity (HPLC-RTM)	GHK + BPC-157 + TB-500 + KPV	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.84% purity, closest to batch mean of 99.84%)

HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.84%	85.33 mg	Confirmed	PASS
Conformity V1	99.84%	85.69 mg	Confirmed	PASS
Conformity V2	99.84%	84.68 mg	Confirmed	PASS
Mean	99.84%	85.23 mg	—	—
Std Dev	0.0000%	0.4180 mg	—	—



Dr. Greg Kalyuzhny

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Lab Director
6/25/2026

COA #: COA-2026-QHRTGB
Access Code: P66ZQ6Q5
Verify: portal.ils-lab.com/verify/7lnd2U-cKgD09Wj
Issued: 6/25/2026

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>)	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: 06/25/2026. Methods: Full QC Panel.
2. The sample was confirmed to be KLOW 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, BPC-157, TB-500, KPV). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
7. Chromatogram shown is representative: Dedicated V0 (99.84% purity, closest to batch mean of 99.84%).




Dr. Greg Kalyuzhny
Lab Director
6/25/2026

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