

ILS Laboratories

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

GHK-Cu / TB-500 / BPC-157 / KPV - 80mg

Tested for: Neo Blue Logistics (Baja Peps)

PASS

COA #: COA-2026-COLJMT
Lot Number: Lot# BP - KL80 - 1001
Accession #: ACC-2026-12602
Labeled Content: 80mg

Analysis Date: 08/06/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 07/27/2026



Scan to verify
authenticity at ils-lab.com
Access Code: RKH3S6PW

Method: Full QC Panel

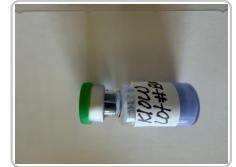
Identity
GHK-Cu / TB-500 / BPC-157 / KPV

Peptide Purity
99.84%



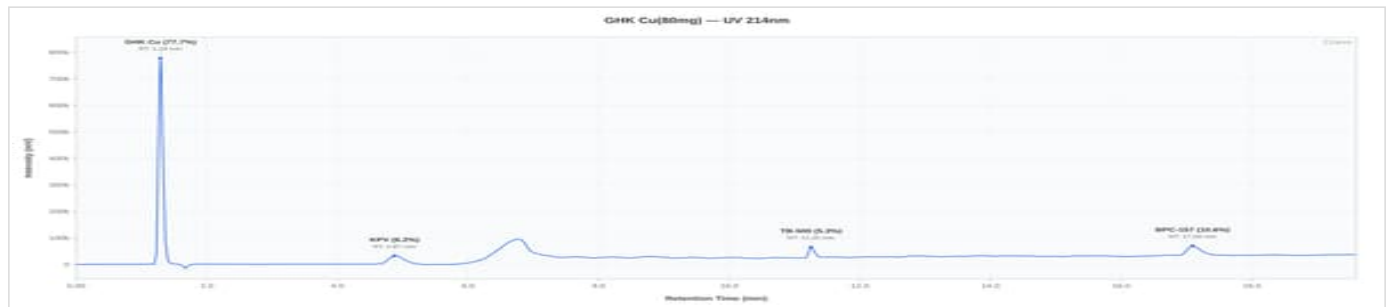
Blend Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.84%	%	PASS
Identity (HPLC-RTM)	GHK-CU + BPC-157 + TB-500 + KPV	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



GHK-Cu / TB-500 / BPC-157 / KPV
80mg - Lot# BP - KL80 - 1001

HPLC Chromatogram



Additional Tests

Analyte	Specification	Result	Unit	Status
Net Blend Peptide Content	Report Only	83.45	mg	N/A
-- GHK-Cu		48.11	mg	
-- BPC-157		12.03	mg	
-- TB-500		12.23	mg	
-- KPV		11.92	mg	

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.83%	84.17 mg	Confirmed	PASS
Conformity V1	99.84%	83.45 mg	Confirmed	PASS
Mean	99.84%	83.81 mg	—	—
Std Dev	0.0050%	0.3600 mg	—	—

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	0.0336	PASS
Cadmium (Cd)	NMT 0.5 ppm	0.015	PASS
Lead (Pb)	NMT 1.0 ppm	0.01816	PASS
Mercury (Hg)	NMT 1.5 ppm	0.00191	PASS
Chromium (Cr)	NMT 10.0 ppm	0.016	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

- Date Tested: 08/06/2026. Methods: Blend Purity, Identity & Quantitation (HPLC); Additional Tests.
- The sample was confirmed to be GHK-Cu / TB-500 / BPC-157 / KPV by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- 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.
- Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
- 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.
- Per-component content calculated from total net peptide content using the manufacturer's stated formulation ratio (GHK-Cu, BPC-157, TB-500, KPV). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
- Chromatogram shown is representative: Conformity V1 (99.84% purity, closest to batch mean of 99.84%).




Dr. Greg Kalyuzhny
Lab Director
8/6/2026

COA #: **COA-2026-COLJMT**
Access Code: **RKH3S6PW**
Verify: <portal.ils-lab.com/verify/LUCU0-zgpeNNmVFd>
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