KOVERA LABS

CERTIFICATE OF ANALYSIS

REPORT # KVR-2026-85A68A

Client(s): Next Level Research Labs

Certified: 10/26/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	Klow	**Form**	Lyophilized Powder
Batch	06012026KO8	**Labeled Qty**	80 mg
Cap Color	Blue	**Crimp Color**	Silver

TEST RESULTS

	REFERENCE STANDARD	RESULT
Blend Avg Purity	(>98%)	99.577% ✓
Blend Avg Net Content	(~80mg)	84.79 mg ✓
Blend Identification (LC-MS)	(Klow)	Klow ✓
Endotoxin Safety Screen	(≤0.5 EU/mL)	PASS ✓

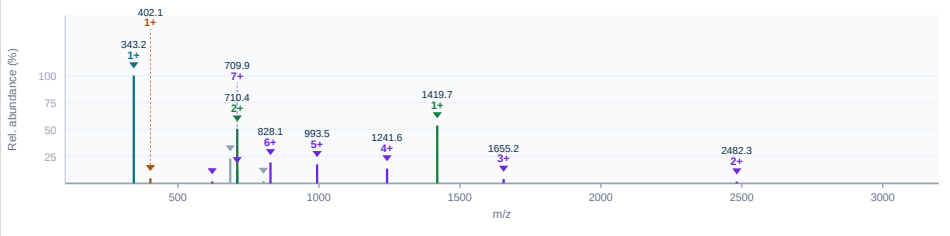
BLEND COMPONENT ANALYSIS

Component	Labeled	Measured
GHK-Cu	50 mg	53.23 mg
KPV	10 mg	9.74 mg
BPC-157	10 mg	10.08 mg
TB-500	10 mg	11.74 mg

MASS SPECTRUM

ESI + mode, MS1 centroid Charge states confirmed **11**

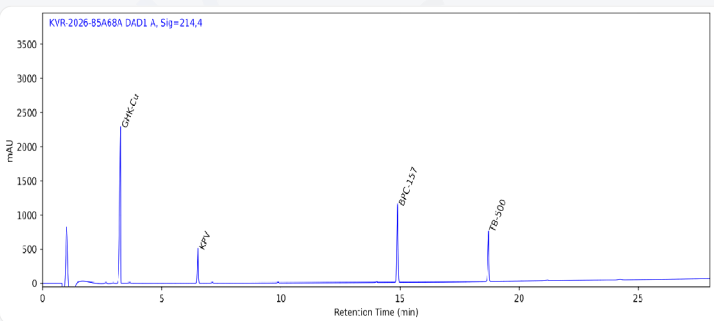
KPV 1 ion **GHK-Cu** 1 ion **TB-500** 7 ions **BPC-157** 2 ions



Each coloured stick is a confirmed charge state of the compound(s) above, drawn at the intensity measured for that ion. Height is relative to the strongest confirmed ion and is not proportional to concentration; content is reported in the Test Results above. The 2 grey stick(s) are the same compound's own in-source fragment and multimer ions, labelled by species: they are produced from the analyte in the ion source and are not separate substances in the vial.

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm





Lemar Arghandiwal
Lab Director

koveralabs.com/verify

Report#: KVR-2026-85A68A
Access Code: XPNL86C

KOVERA LABS

BACTERIAL ENDOTOXIN ANALYSIS

(Kinetic Chromogenic LAL, USP <85>)

Report Number: **KVR-2026-F19DE9**

CLIENT & SAMPLE INFORMATION

Client	Next Level Research Labs	**Certified Date**	10/26/2026
Product Name	Klow	**Strength**	80 mg
Lot / Batch	06012026KO8	**Condition**	Lyophilized

This Certificate of Analysis certifies that the sample listed herein was tested for bacterial endotoxin using a kinetic chromogenic LAL assay in accordance with USP <85> Bacterial Endotoxins Test.

TEST METHODOLOGY

Test Performed	Quantitative Bacterial Endotoxin Test	**Compendial Ref**	USP <85>
Assay Type	Kinetic Turbidimetric	**Detection λ**	660 nm
Detection Range	0.01 - 1.0 EU/mL	**Endotoxin Std**	E. coli O111:B4
Dilution Vol	2.0 mL	**Matrix**	LAL Reagent Water

QUANTITATIVE RESULTS

Parameter	Result
Endotoxin Level	< 0.04 EU/mL
Acceptance Limit	≤ 0.5 EU/mL
Sample CV (%)	2.8%
Spike CV (%)	2.8%
Spike Recovery (%)	89%
Final Determination	**PASS**

CONTROLS

Control	Expected	Observed	Status
Positive Control	Detectable	As expected	Pass
Negative Control	No signal	As expected	Pass

INTERPRETATION

Endotoxin content was quantitatively determined using a kinetic chromogenic LAL assay in accordance with USP <85>. Result reported as below quantitation threshold. The measured endotoxin level meets the specified acceptance criteria. The sample passes the endotoxin limit test.

AUTHORIZATION

REVIEWED BY

Lemar Arghandiwal

Lab Director