



PASS — ALL SPECIFICATIONS MET

4 of 4 layers complete

■ L1
Identity

■ L2
Purity

■ L3
Net Peptide Content

■ L4
Sterility

Sample & Vial Validation



Vial A: as received & analyzed

Client	AncientCity Biotech	Product	Klow
Lot / Batch	926-Kl-80	Label Claim	80 mg
Matrix	lyophilized powder	Sample Count	1
Vial Size	3mL	Cap Color	Blue
Crimp Color	Silver / Aluminum	Contents Color	Blue
Seal Intact	Yes	Condition	sealed, intact
Received	Sep 2, 2026	Sampling	client submitted

Test Results

Test	Reference standard	Result	
Identity	every declared component within ± 1 Da of its own theoretical mass	4 of 4 components confirmed	PASS
Purity	each declared component at or above its own locked purity criterion	4 of 4 components within criterion	PASS
Net Peptide Content	each declared component within its own locked content criterion, at the declared ratio	82.7 mg of 80 mg declared (+3.4%), 4 of 4 components within criterion	PASS
Sterility · Rapid PCR	Rapid PCR screen	Target DNA not detected	PASS

Layer Results

1

Identity

HILIC-ESI-MS intact GHK-Cu full-scan/isotope identity + LC-MS/MS (ESI+) accurate mass

4 of 4 components confirmed

each +/- 1 Da of its component mass

PASS

Component	Its criterion	Observed	Status
GHK-Cu	402.92 Da · acceptance ± 1 Da · average	402.92 Da	PASS
BPC-157	1419.556 Da · acceptance ± 1 Da · average	1419.556 Da	PASS
TB-500 (Thymosin B4)	4963.506 Da · acceptance ± 1 Da · average	4963.506 Da	PASS
KPV	342.44 Da · acceptance ± 1 Da · average	342.44 Da	PASS



2

Purity

HILIC-CAD homogeneity (universal detection) + RP-HPLC 214 nm area% (USP <621>)

4 of 4 components within criterion

each component at or above its own locked purity criterion

PASS

Component	Its criterion	Observed	Status
GHK-Cu	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.96%	PASS
BPC-157	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.97%	PASS
TB-500 (Thymosin B4)	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.99%	PASS
KPV	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.94%	PASS

3

Net Peptide Content

Quantitative HPLC vs reference std

82.7 mg of 80 mg declared (+3.4%), 4 of 4 components within criterion

each component within its own locked content criterion, at the declared ratio

PASS

Component	Its criterion	Observed	Status
GHK-Cu	5 parts · 45–55 mg · Declared blend recipe on the controlled Axiom catalog record for Klow (component ratio 5); acceptance ±10% per Axiom SPEC-CONTENT-001	51.32 mg	PASS
BPC-157	1 part · 9–11 mg · Declared blend recipe on the controlled Axiom catalog record for Klow (component ratio 1); acceptance ±10% per Axiom SPEC-CONTENT-001	10.57 mg	PASS
TB-500 (Thymosin B4)	1 part · 9–11 mg · Declared blend recipe on the controlled Axiom catalog record for Klow (component ratio 1); acceptance ±10% per Axiom SPEC-CONTENT-001	10.63 mg	PASS
KPV	1 part · 9–11 mg · Declared blend recipe on the controlled Axiom catalog record for Klow (component ratio 1); acceptance ±10% per Axiom SPEC-CONTENT-001	10.18 mg	PASS

Observed ratio **GHK-Cu 4.86 : BPC-157 1.00 : TB-500 (Thymosin B4) 1.01 : KPV 0.96** against declared **GHK-Cu 5.00 : BPC-157 1.00 : TB-500 (Thymosin B4) 1.00 : KPV 1.00** · source Declared blend recipe on the controlled Axiom catalog record for Klow (component ratio 5); acceptance ±10% per Axiom SPEC-CONTENT-001

4

Sterility

Rapid Microbial DNA Screen — Qualification-Bound Target Panel

Sterility · Rapid PCR: No Growth

No microbial growth or target DNA detected

PASS

Sterility · Rapid PCR

Result

No Growth

Specification

Rapid PCR screen

PASS

Reporting Scope & Limitations

[Purity] The chromatographic trace was not retained for this certificate. The purity value and its acceptance limit are unaffected.

[Identity] The mass spectrum was not retained for this certificate. Identity is reported from the recorded accurate mass against its theoretical value.

[Sterility · Rapid PCR] DNA-target research screen only. It detects microbial DNA from viable and non-viable organisms alike, and is neither viable-CFU enumeration under USP <61> nor a batch sterility determination under USP <71>.



SCAN TO VERIFY

Digital Signatures

Analyst	Alana Reeves	Analyst Signature	ed25519:02970c648042af6d35707613...
Technical Reviewer	Michael Niles	Reviewer Signature	ed25519:77bb2ab5aba8be01e5d6b9cb...
Laboratory Director	Daniel Brennan	Director Signature	ed25519:c044dd25f0f49bc8d9e7ccd1...

Physical instrument analysis is performed at Axiom Analytics' discretion (based on method, capacity, and order volume), either in Axiom's own laboratory under its documented quality system or by an independent partner laboratory engaged under confidentiality. Where analysis is subcontracted, the method is performed within the partner laboratory's published ISO/IEC 17025 scope of accreditation for that method, not merely at a laboratory holding ISO/IEC 17025 accreditation. Axiom Analytics does not itself currently hold ISO/IEC 17025 accreditation. The method-validation status of each reported result is identified beside that result. Partner laboratory identities are not disclosed: samples are analyzed under double-blind conditions, identified only by an internal lot ID.

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SHA-384: 8e1466ff1a122bb83aeb1bc3690d23143ee640d1678bd7f0a84a43610215d5bd...

Seal: ed25519:b89e1eca11cabf1cd2ef7840b6fc740c00b27e31...

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