



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	CJC-1295 (No DAC)/Ipamorelin
Lot / Batch	926-Cl-10	Label Claim	10 mg
Matrix	lyophilized powder	Sample Count	1
Vial Size	3mL	Cap Color	Royal Blue
Crimp Color	Silver / Aluminum	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	2 of 2 components confirmed	PASS
Purity	each declared component at or above its own locked purity criterion	2 of 2 components within criterion	PASS
Net Peptide Content	each declared component within its own locked content criterion, at the declared ratio	10.41 mg of 10 mg declared (+4.1%), 2 of 2 components within criterion	PASS
Sterility · Rapid PCR	Rapid PCR screen	Target DNA not detected	PASS

Layer Results

1

Identity

LC-MS/MS (ESI+)

2 of 2 components confirmed

each +/- 1 Da of its component mass

PASS

Component	Its criterion	Observed	Status
CJC-1295 no DAC	3367.954 Da · acceptance ± 1 Da · average	3367.954 Da	PASS
Ipamorelin	711.868 Da · acceptance ± 1 Da · average	711.868 Da	PASS



2 Purity

 RP-HPLC 214nm (USP 621)

2 of 2 components within criterion
 each component at or above its own locked purity criterion

PASS

Component	Its criterion	Observed	Status
CJC-1295 no DAC	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.79%	PASS
Ipamorelin	Expected Purity ≥ 99.25 area % · Acceptance NLT 99.25 area % · Units area % · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.89%	PASS

3 Net Peptide Content

 Quantitative HPLC vs reference std

10.41 mg of 10 mg declared (+4.1%), 2 of 2 components within criterion
 each component within its own locked content criterion, at the declared ratio

PASS

Component	Its criterion	Observed	Status
CJC-1295 no DAC	1 part · 4.5–5.5 mg · Declared blend recipe on the controlled Axiom catalog record for CJC-1295 (No DAC)/Ipamorelin (component ratio 1); acceptance ±10% per Axiom SPEC-CONTENT-001	5.18 mg	PASS
Ipamorelin	1 part · 4.5–5.5 mg · Declared blend recipe on the controlled Axiom catalog record for CJC-1295 (No DAC)/Ipamorelin (component ratio 1); acceptance ±10% per Axiom SPEC-CONTENT-001	5.23 mg	PASS

 Observed ratio **CJC-1295 no DAC 1.00 : Ipamorelin 1.01** against declared **CJC-1295 no DAC 1.00 : Ipamorelin 1.00** · source Declared blend recipe on the controlled Axiom catalog record for CJC-1295 (No DAC)/Ipamorelin (component ratio 1); 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
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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>.

Digital Signatures

Analyst	Alana Reeves	Analyst Signature	ed25519:7c6d7ea6df536bbd46221c7c...
Technical Reviewer	Michael Niles	Reviewer Signature	ed25519:3c7e5e9b8b92ab4d4851a2e1...
Laboratory Director	Daniel Brennan	Director Signature	ed25519:1a207432a3a9210d8867d346...

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.

Each signature is an ed25519 signature over the SHA-384 hash of this report. Scan the QR on the letterhead or verify independently at <https://axiomanalyticslab.com/verify/AX-2026-4672-Q322>.



Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL



Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US

AX-2026-4672-Q322

CJC-1295 (No DAC)/Ipamorelin · Lot 926-Cl-10
Issued Sep 6, 2026 · v1
Verify: 28B4-A2C8-E2EF



SCAN TO VERIFY

✓ **CRYPTOGRAPHICALLY SEALED (ed25519 + SHA-384)**

SHA-384: 7d57330c4a8646567e627f8ba2e33ab05834e0b75e1936b3fcb667a24bcf01fd...

Seal: ed25519:08e950674b481cce5f6da0fcf224e0f54f864bfa...

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.

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