Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL

Axiom Analytics Labs LLC
Sacramento, California, USA

AX-2026-2553-J7VZ

BPC-157 · Lot GL-BPC10MG-04-202610-001
Issued Oct 22, 2026 PT · v1
Verify: B333-0C25-B9B7

SCAN TO VERIFY

PASS — ALL SPECIFICATIONS MET

4 of 4 layers complete

L1	L2	L3	L4
Identity	Purity	Net Peptide Content	Sterility

Sample & Vial Validation

Vial A: as received & analyzed

MORE VIEWS OF VIAL A

Vial A · right

Brand	**Guardian Labs**	Product	**BPC-157**
Lot / Batch	**GL-BPC10MG-04-202610-001**	Label Claim	**10 mg**
Matrix	**lyophilized powder**	Sample Count	**1**
Vial Size	**3mL**	Cap Color	**Black**
Crimp Color	**Silver / Aluminum**	Contents Color	**White**
Seal Intact	**Yes**	Condition	**sealed, intact**
Received	**Oct 21, 2026 PT**	Sampling	**client submitted**

Test Results

Test	Specification	Result	
Identity research screen	every vial within ±1 Da of 1419.556 Da	`1419.546 Da`	PASS
Purity research screen	≥ 99.25 area %	`99.95%`	PASS
Net Peptide Content research screen	every vial within ±10% of 10 mg	`Vial A: 10.39 mg`	PASS
Sterility · Rapid PCR research screen	Rapid PCR screen	`Target DNA not detected`	PASS

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.

Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL

Axiom Analytics Labs LLC
Sacramento, California, USA

AX-2026-2553-J7VZ

BPC-157 · Lot GL-BPC10MG-04-202610-001
Issued Oct 22, 2026 PT · v1
Verify: B333-0C25-B9B7

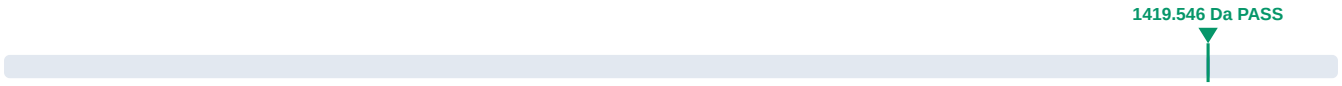
SCAN TO VERIFY

Layer Results

1 Identity
LC-MS (ESI+)

1419.546 Da
every vial within ±1 Da of 1419.556 Da
PASS

Theoretical mass	**1419.556 Da**	Acceptance	**within ±1 Da**
Vial A	**1419.546 Da · PASS**		



2 Purity
RP-HPLC 214nm (USP <621>)

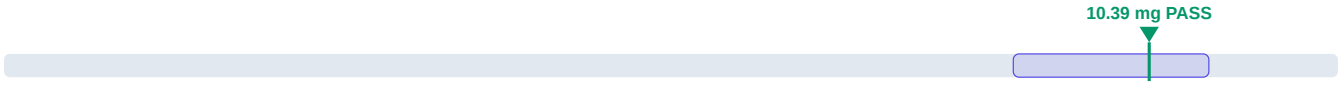
99.95%
≥ 99.25 area %
PASS

Acceptance	**≥ 99.25%**	Vial A	**99.95% · PASS**
Criterion source	**Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31**		

3 Net Peptide Content
Quantitative HPLC vs reference std

Vial A: 10.39 mg
every vial within ±10% of 10 mg
PASS

Target	**10 mg (±10%)**	Vial A	**10.39 mg · PASS**



4 Sterility · Rapid PCR
Rapid Microbial DNA Screen (qPCR target panel)

Target DNA not detected
Rapid PCR screen
PASS

Target panel	**Pan-bacterial 16S rRNA gene (VIC), Pan-fungal 18S rRNA gene (FAM)**

`Run controls valid.`

Reporting Scope & Limitations

`[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>.

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.

Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL

Axiom Analytics Labs LLC
Sacramento, California, USA

AX-2026-2553-J7VZ

BPC-157 · Lot GL-BPC10MG-04-202610-001
Issued Oct 22, 2026 PT · v1
Verify: B333-0C25-B9B7

SCAN TO VERIFY

Digital Signatures

Analyst	**Alana Reeves**	Analyst Signature	**ed25519:3e346715956526268a73efa0...**
Technical Reviewer	**Michael Niles**	Reviewer Signature	**ed25519:24a5436c932e166246f799ca...**
Laboratory Director	**Daniel Brennan**	Director Signature	**ed25519:4f8e24894ea807298df876a1...**

Where physical instrument analysis is performed, it takes place within Axiom Analytics' laboratory network, allocated at Axiom's discretion based on method, capacity, and order volume. Axiom Analytics performs sample accession, chain of custody, analytical oversight, and data review, and issues and is responsible for this certificate. 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: where samples are analyzed under double-blind conditions, they are 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-2026-2553-J7VZ.

✓ CRYPTOGRAPHICALLY SEALED (ed25519 + SHA-384)

SHA-384: 2ea6a915aebacfa76c0d58b1a7b0b1dc55341f1d99af668a9bed180c2d95e3f3...

Seal: ed25519:3f22322f9ab3fe14df108a7b7009496423d645c9...

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.

RESEARCH USE ONLY. All analytical testing, validation reports, and interactive Certificates of Analysis (COAs) provided by Axiom Analytics are intended strictly for Research Use Only (RUO) in in-vitro or laboratory animal model environments. Axiom Analytics does not analyze, validate, evaluate, or make any representations regarding the safety, efficacy, toxicity, or legality of any tested substance for human or animal consumption, veterinary use, clinical diagnostics, food additives, cosmetics, or household applications.

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.