



Certificate of Analysis

FULL QUALITY CONTROL PANEL · PEPTIDE ANALYTICAL REPORT

CERTIFICATE NO.

COA-2026-9444

Issued 07/02/2026 · Page 1 of 4

CLIENT		PREPARED FOR		SAMPLE & ACCESSION		ACC - 2026 - 7444			
Aion Limitless aionlimitless.com Client reference: on file Distribution: client confidential				ANALYTE / IDENTITY		CAS NUMBER		LOT NUMBER	
				BPC-157		137525-51-0		BP-2602	
				LABELED CONTENT		SAMPLE MATRIX		APPEARANCE	
				10 mg		Lyophilized		Conforms	
				MANUFACTURED		RETEST / EXPIRY		STORAGE	
				05/02/2026		05/2028		-20 C, dark	
				DATE RECEIVED		DATE ANALYZED		DISPOSITION	
				06/28/2026		07/02/2026		RELEASED	

PEPTIDE PURITY 98.10% Spec ≥ 95.0% ✓ PASS	IDENTITY BPC-157 HPLC-RTM + LC-MS ✓ PASS	NET PEPTIDE CONTENT 10.30 mg measured · label 10 mg REPORTED	BACTERIAL ENDOTOXIN 0.6 EU/vial · limit NMT 5 EU/mL ✓ PASS	DISPOSITION RELEASED Full QC Panel complete
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1 RELEASE STATEMENT

The submitted BPC-157 sample was evaluated under the Full Quality Control Panel covering identity, purity, net peptide content, composition, sterility, bacterial endotoxin, elemental impurities, and targeted adulterant screening. Identity was confirmed by HPLC retention-time matching with an LC-MS [M+2H]²⁺ ion at m/z 710.77 (neutral mass 1419.5 Da). Chromatographic purity of 98.10 percent comfortably exceeded the not less than 95.0 percent release criterion. Sterility returned No Growth, bacterial endotoxin was well within the applicable limit, and arsenic, cadmium, chromium, mercury, and lead were each Not Detected, with fentanyl Not Detected at the screening cutoff. On the basis that all tested attributes meet their respective specifications, the laboratory supports a disposition of RELEASED for the material as received.

Authorized Disposition

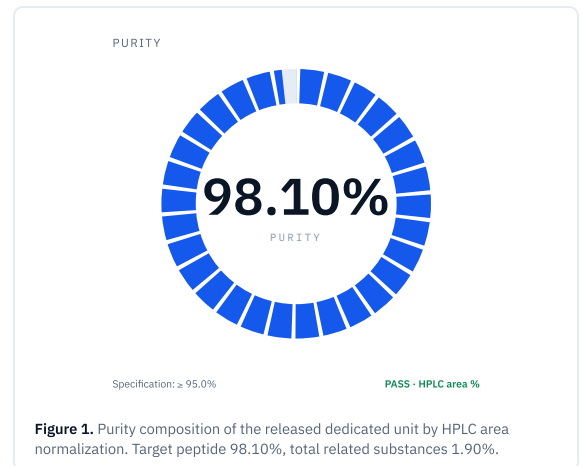
IDENTITY Confirmed ✓ PASS	PURITY 98.10% ✓ PASS	NET CONTENT 10.30 mg ✓ PASS	MOISTURE 5.20% w/w ✓ PASS	STERILITY No Growth ✓ PASS	ENDOTOXIN 0.6 EU/vial ✓ PASS	HEAVY METALS Not Detected ✓ PASS
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2 PURITY, QUANTITATION & IDENTITY

HPLC / HPLC-RTM

ANALYTE	SPECIFICATION	RESULT	UNIT	STATUS
Peptide Purity (HPLC)	≥ 95.0%	98.10	%	✓ PASS
Net Peptide Content	Report only	10.30	mg	REPORTED
Identity (HPLC-RTM)	BPC-157	Confirmed	—	✓ PASS
Fentanyl Screen	Immunoassay, 50 ng/mL	Not Detected	—	✓ PASS

Interpretation. The measured chromatographic purity of 98.10 percent exceeds the not less than 95.0 percent release criterion, indicating a homogeneous principal peak with a low related-substance burden. Net peptide content of 10.30 mg is the measured mass recovered from the unit and is reported against a nominal 10 mg label claim, confirming the vial delivers at least the labeled mass of active peptide.



3 REPRESENTATIVE CHROMATOGRAM

Reversed-phase HPLC-UV

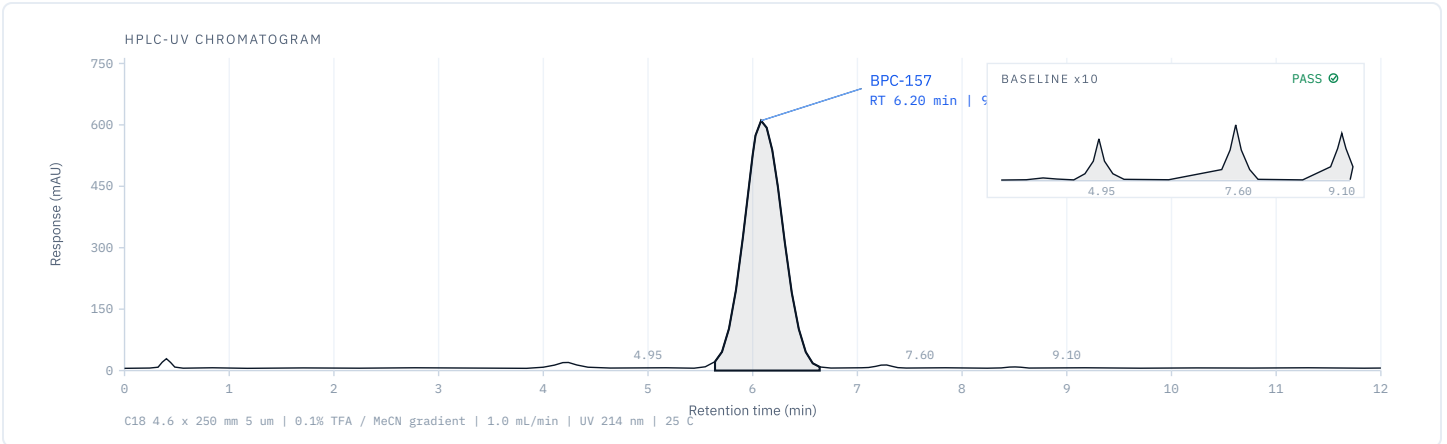


Figure 2. Representative reversed-phase HPLC-UV chromatogram (Conformity unit V1, 98.03% main-peak area, near the batch mean of 98.06%). Trace related substances are shown magnified 10x in the inset. A sharp, symmetric principal peak with a low related-substance burden is consistent with high chromatographic purity.

PEAK	RT (MIN)	AREA (MAU·S)	AREA %	PLATES (N)	TAILING
BPC-157 (principal)	6.20	4,901,500	98.03	9,120	1.05
Related substance 1 (RRT 0.80)	4.95	42,500	0.85	6,980	1.11
Related substance 2 (RRT 1.23)	7.60	34,000	0.68	6,610	1.09
Related substance 3 (RRT 1.47)	9.10	22,000	0.44	6,240	1.13
Total		5,000,000	100.00		

4 MASS CONFIRMATION

LC-MS, ESI positive



Figure 3. LC-MS mass confirmation (electrospray positive mode). Base peak [M+2H]2+ at m/z 710.77 with [M+H]+ at 1420.54, consistent with the 1419.5 Da mass of BPC-157 and confirming identity.

5 CONFORMITY TESTING

3 units · HPLC

SAMPLE	PURITY	NPC (MG)	IDENTITY	RESULT
V0 (dedicated)	98.10%	10.30	Confirmed	✓ PASS
V1	98.03%	10.24	Confirmed	✓ PASS
V2	98.06%	10.41	Confirmed	✓ PASS
Mean	98.06%	10.32	—	—
Std Dev	0.035%	0.0862	—	—

Interpretation. Across three independently tested units the purity standard deviation was 0.035 percent and content standard deviation 0.0862 mg, a low dispersion consistent with a homogeneous, well-controlled batch.



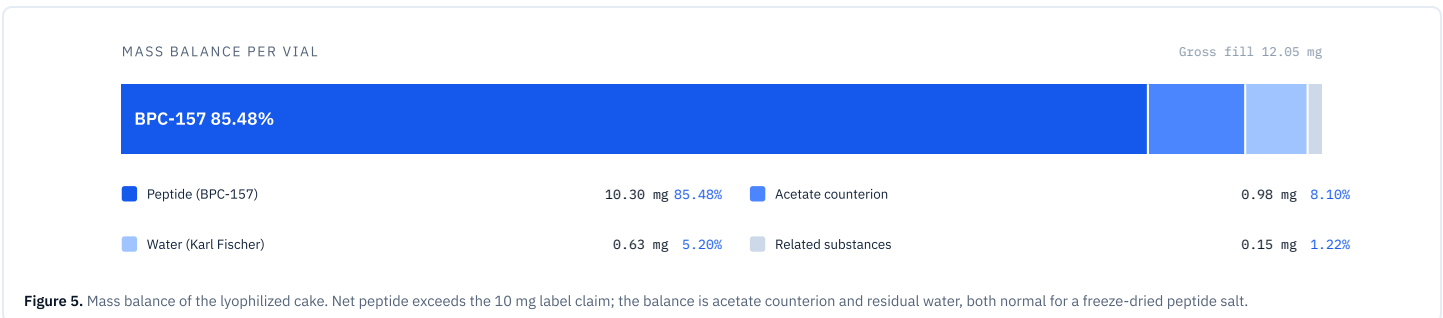
Figure 4. Purity conformity across tested units with mean and plus or minus 3 SD control band (SD 0.035%).

6 COMPOSITION, MOISTURE & COUNTERION

KF · IC · USP <467>

PARAMETER	METHOD	SPECIFICATION	RESULT	STATUS
Net Peptide Content	HPLC / AAA	Report only	10.30 mg	REPORTED
Water (Karl Fischer)	USP <921>	NMT 8.0% w/w	5.20%	✓ PASS
Counterion (acetate)	Ion chromatography	Report only	8.10% w/w	REPORTED
Residual Solvents (TFA)	USP <467>	Not Detected	Not Detected	✓ PASS
Gross Fill Weight	Gravimetric	Report only	12.05 mg	REPORTED

Interpretation. Karl Fischer titration returned 5.20 percent residual water and the acetate counterion accounts for 8.10 percent by weight, both typical for a freeze-dried peptide salt. The 12.05 mg gross fill reconciles peptide, counterion, water, and trace related substances.

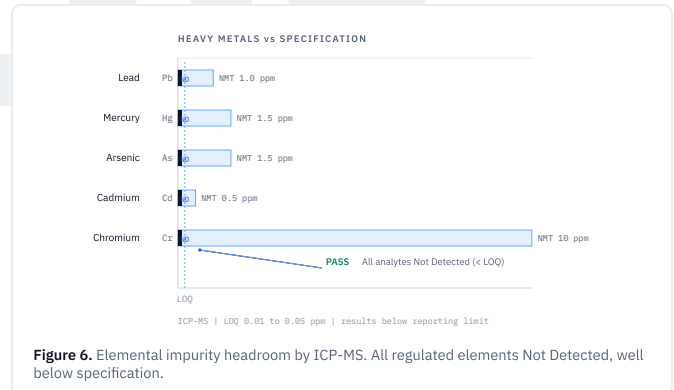


7 SAFETY & CONTAMINATION CONTROLS

ICP-MS · USP <71> · <85>

TEST	METHOD	SPECIFICATION	RESULT	STATUS
Arsenic (As)	ICP-MS	NMT 1.5 ppm	Not Detected	✓ PASS
Cadmium (Cd)	ICP-MS	NMT 0.5 ppm	Not Detected	✓ PASS
Chromium (Cr)	ICP-MS	NMT 10 ppm	Not Detected	✓ PASS
Mercury (Hg)	ICP-MS	NMT 1.5 ppm	Not Detected	✓ PASS
Lead (Pb)	ICP-MS	NMT 1 ppm	Not Detected	✓ PASS
Sterility	USP <71>	No Growth	No Growth	✓ PASS
Bioburden	USP <61>	< 10 CFU/mL	< 1 CFU/mL	✓ PASS
Bacterial Endotoxin	USP <85>	NMT 5 EU/mL	0.6 EU/vial	✓ PASS

Interpretation. By ICP-MS, arsenic, cadmium, chromium, mercury, and lead were each Not Detected below their specification thresholds. Sterility by membrane filtration returned No Growth with bioburden below the reporting limit, and bacterial endotoxin measured 0.6 EU/vial, at or below 0.60 EU/mL once reconstituted in 1 mL or more, within the not more than 5 EU/mL release limit.



8 FOR THE RECIPIENT

Plain-language summary

- ✓ **Purity 98.10 percent.** Almost everything in the vial is BPC-157. Only 1.90 percent is trace related substances, comfortably past the 95 percent release bar.
- ✓ **You get at least the labeled amount.** Measured net peptide content is 10.30 mg, at or above the 10 mg on the label, so the measured content meets or exceeds the label claim.
- ✓ **It is genuinely BPC-157.** HPLC retention time plus an LC-MS [M+2H]²⁺ ion at m/z 710.77 (neutral mass 1419.5 Da) confirm the identity, so this is not a substitute or a mislabel.
- ✓ **Sterility and endotoxin within specification.** Sterility shows No Growth, and bacterial endotoxin measured 0.6 EU/vial, which stays under the not more than 5 EU/mL limit at the tested reconstitution volume.
- ✓ **Screened for contamination.** Arsenic, cadmium, chromium, mercury, and lead were Not Detected, and the fentanyl adulterant screen was Not Detected.
- ✓ **Water and salt are normal.** The small amounts of water (5.20 percent) and acetate (8.10 percent) are expected for a freeze-dried peptide and are already accounted for in the net peptide figure.

Purity (HPLC)

Percent of the main peptide peak relative to all detected peaks.

Net peptide content

Measured mass of actual peptide, separate from water and salt.

NMT

Not more than; an upper acceptance limit.

Not Detected

Below the method limit of quantitation (LOQ).

EU/mL

Endotoxin units per milliliter, a measure of bacterial toxin.

RRT

Relative retention time; an impurity position versus the main peak.

Counterion

The salt partner of the peptide.

w/w

By weight, as a percent of total mass.

USP <71> / <85>

Standard chapters for sterility and bacterial endotoxin.

Karl Fischer

Titration method for measuring residual water (USP <921>).

Storage. Store the sealed lyophilized vial at -20 C, protected from light. Reconstituted solution is stable at 2 to 8 C; use within 30 days. **Reconstitution.** Add bacteriostatic or sterile water down the vial wall, swirl gently, do not shake. **Handling.** Research use only; not for human or veterinary use.

9 ANALYTICAL METHODS & NOTES

Method principles

HPLC Purity and Net Peptide Content

Reversed-phase HPLC with UV detection separates the principal peptide from related substances; purity is percent main-peak area and content is quantified against a calibrated reference. Acceptance: not less than the stated release limit.

LC-MS Mass Confirmation

Electrospray positive-mode mass spectrometry resolves the molecular ion and isotope or charge-state signature of the analyte.

Counterion by Ion Chromatography

Ion chromatography determines the counterion content by weight, characterizing the peptide salt form.

Heavy Metals by ICP-MS

Inductively coupled plasma mass spectrometry quantifies elemental impurities against per-element limits, with results below detection reported as Not Detected.

Bioburden by USP <61>

Total aerobic microbial and yeast or mold counts establish microbial load prior to release.

Identity by HPLC Retention-Time Matching

Sample retention time is compared to a qualified reference standard under identical conditions; identity is confirmed on co-elution within the established window.

Water Content by Karl Fischer

Coulometric Karl Fischer titration per USP <921> quantifies residual moisture in the lyophilized cake. Acceptance: not more than 8.0 percent w/w.

Residual Solvents

Headspace GC per USP <467> screens process solvents; trifluoroacetic acid is assessed against a not-detected threshold.

Sterility by USP <71>

Membrane filtration sterility test; No Growth is reported when no microbial recovery is observed across the incubation period.

Bacterial Endotoxin by USP <85>

Kinetic bacterial endotoxin test; acceptance is not more than 5 EU/mL per client specification.

DocuSigned by:



10101011 11111000 00111100 01000111

DS

FC

Finn C.

Laboratory Director

Date of authorization: 07/02/2026

DocuSigned by:



00011101 10001111 00000001 01000110

DS

BC

Brennan C.

Quality Assurance

Reviewed against acceptance criteria: 07/02/2026



Authenticate this certificate

Scan the code or verify online at titreonanalytical.com/verify/COA-2026-9444

Verification ID: TLA-9444-BPC157-3C9B



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