

ILS Laboratories

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(619) 329-3999 | ils-lab.com

BPC-157 - 10mg



Tested for: Aeon Aminos
www.aeonaminos.com

COA #: COA-2026-XHWGSZ
Lot Number: AEON-BPC10-C1726
Accession #: ACC-2026-10642
Labeled Content: 10mg

Analysis Date: 08/07/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 07/17/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: F43PK8HQ

Method: Full QC Panel

Identity	Peptide Purity	Fentanyl Free
BPC-157	99.76%	

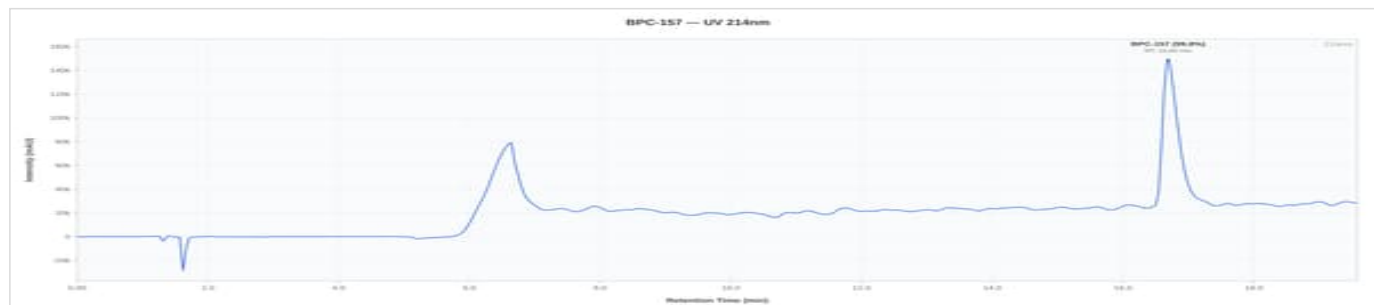
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.76%	%	PASS
Net Peptide Content	Report Only	10.59	mg	N/A
Identity (HPLC-RTM)	BPC-157	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



BPC-157 10mg - AEON-BPC10-C1726

HPLC Chromatogram



BPC-157 10mg - AEON-BPC10-C1726: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	0.0998	PASS
Cadmium (Cd)	NMT 0.5 ppm	0.01888	PASS
Lead (Pb)	NMT 1.0 ppm	0.0178	PASS
Mercury (Hg)	NMT 1.5 ppm	0.0828	PASS
Chromium (Cr)	NMT 10.0 ppm	0.078	PASS



Dr. Greg Kalyuzhny

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Lab Director
8/7/2026

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Verify: portal.ils-lab.com/verify/LD70y2yCU7MBn82b
Issued: 8/7/2026

Sterility Testing (PCR)

Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	NMT 0.05 EU/mL	Reported

About this result: Endotoxin is reported as a quantitative value. Acceptable limits vary by product type and matrix, so no universal pass/fail threshold applies to RUO products. This result is below commonly referenced endotoxin thresholds.

Notes & Methodology

1. Date Tested: 08/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be BPC-157 by HPLC. Identification by chromatographic retention time comparison with a reference standard.
3. Elemental impurities analyzed by ICP-MS per USP <233> methodology. Acceptance criteria are internal laboratory quality screening limits for research-use materials and do not represent evaluation against any specific pharmacopeial monograph or product specification.
4. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
5. Peptide purity determined by RP-HPLC area normalization at 214 nm. Value represents the percentage of the target peptide relative to all peptide-related peaks. Non-peptide process-related impurities, if detected, are excluded from the calculation.




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