Certificate of Analysis

ILS Laboratories
8222 Vickers St, Suite 106, San Diego, CA 92111
(619) 329-3999 | ils-lab.com

BPC-157 / TB-500 - 10mg

PASS

Tested for: Platinum Lion Peptides
America's Hometown Plymouth MA, 02360 | www.platinumlionpeptides.com

COA #:	**COA-2026-YJURPF**	Analysis Date:	**07/31/2026**
Lot Number:	**BPCTB55-1004**	Appearance:	**Good**
Accession #:	**ACC-2026-9456**	Sample Matrix:	**Lyophilized**
Labeled Content:	**10mg**	Volume:	**3mL**
		Date Received:	**07/10/2026**

Scan to verify authenticity at ils-lab.com
Access Code: FJZM6C2B

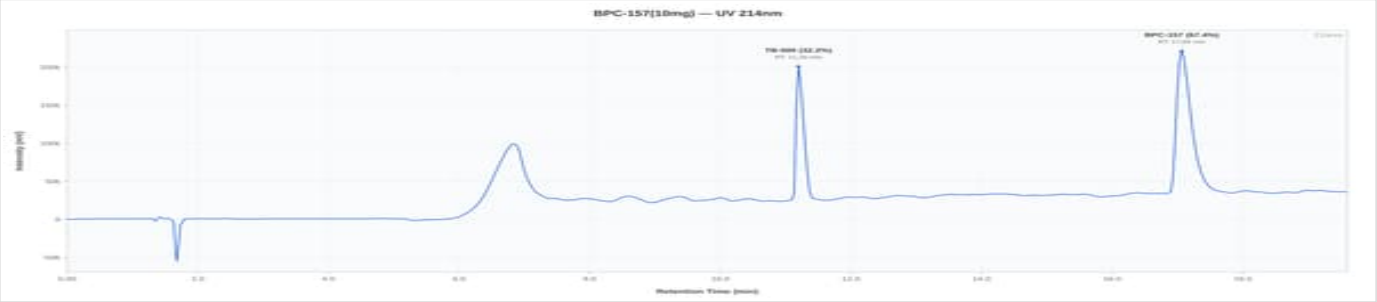
Method: **Full QC Panel**

Identity	Peptide Purity	
BPC-157 / TB-500	**99.68%**	**Fentanyl Free**

Blend Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.68%**	%	**PASS**
Identity (HPLC-RTM)	BPC-157 + TB-500	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Dedicated V0

Additional Tests

Analyte	Specification	Result	Unit	Status
Net Blend Peptide Content	Report Only	**10.61**	mg	N/A
-- BPC-157		*5.41*	*mg*	
-- TB-500		*5.20*	*mg*	

Dr. Greg Kalyuzhny
Lab Director
7/31/2026

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Verify: portal.ils-lab.com/verify/HdvIzH8Ueiw5m-dn
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HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.68%	10.61 mg	Confirmed	**PASS**
Conformity V1	99.68%	10.55 mg	Confirmed	**PASS**
Mean	**99.68%**	**10.58 mg**	—	—
Std Dev	—	**0.0300 mg**	—	—

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	.2629	**PASS**
Cadmium (Cd)	NMT 0.5 ppm	.365	**PASS**
Lead (Pb)	NMT 1.0 ppm	.12	**PASS**
Mercury (Hg)	NMT 1.5 ppm	.269	**PASS**
Chromium (Cr)	NMT 10.0 ppm	5.36	**PASS**

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	**0.165 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: 07/31/2026. Methods: Blend Purity, Identity & Quantitation (HPLC); Additional Tests.
2. The sample was confirmed to be BPC-157 / TB-500 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.
6. Per-component content calculated from total net peptide content using the manufacturer's stated formulation ratio (BPC-157, TB-500). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
7. Chromatogram shown is representative: Dedicated V0 (99.68% purity, closest to batch mean of 99.68%).

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