

ILS Laboratories

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

BPC & TB Blend - 20mg (10/10)



Tested for: NOVA Labs
www.nova-biolabs.com

COA #: COA-2026-MSW45K
Lot Number: 0627
Accession #: ACC-2026-13806
Labeled Content: 20mg (10/10)

Analysis Date: 08/10/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 08/02/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: UFJN3X4B

Method: Full QC Panel

Identity	Peptide Purity	Fentanyl Free
BPC & TB Blend	99.82%	

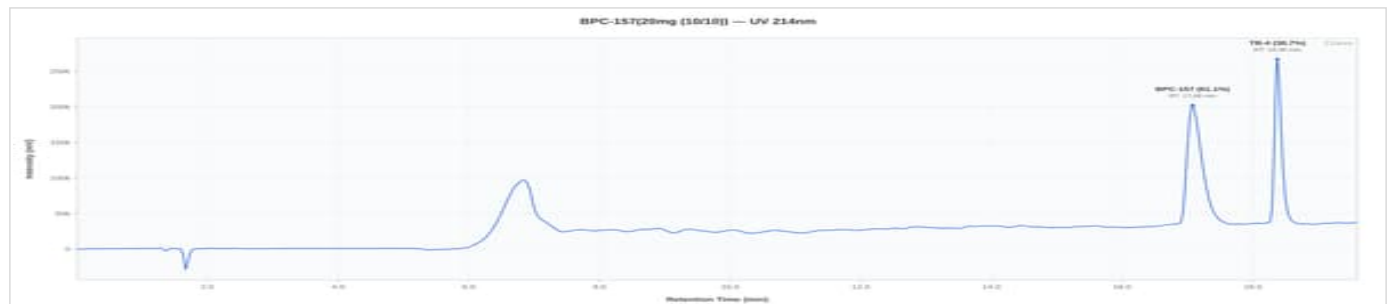
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.82%	%	PASS
Net Blend Peptide Content	Report Only	21.16	mg	N/A
-- BPC-157		10.79	mg	
-- TB-500		10.37	mg	
Identity (HPLC-RTM)	BPC-157 + TB-500	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



BPC & TB Blend 20mg (10/10) - 0627

HPLC Chromatogram



BPC & TB Blend 20mg (10/10) - 0627: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	Not Detected	PASS
Cadmium (Cd)	NMT 0.5 ppm	Not Detected	PASS
Lead (Pb)	NMT 1.0 ppm	Not Detected	PASS
Mercury (Hg)	NMT 1.5 ppm	Not Detected	PASS
Chromium (Cr)	NMT 10.0 ppm	Not Detected	PASS



Dr. Greg Kalyuzhny

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Lab Director
8/10/2026

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Verify: portal.ils-lab.com/verify/Zv1tyQpQPJyozM1I
Issued: 8/10/2026

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: 08/10/2026. Methods: Full QC Panel.
2. The sample was confirmed to be BPC & TB Blend 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.




Dr. Greg Kalyuzhny
Lab Director
8/10/2026

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Verify: <portal.ils-lab.com/verify/Zv1tyQpQPJyozM1I>
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