

HPLC Purity Analysis

BPC157/TB500 — Research Grade Verification

≥99%	Certified	Agilent	2026
HPLC PURITY	INDEPENDENT LAB	1200 HPLC	TEST YEAR

ANALYSIS REPORT — BPC157/TB500

BPC157/TB500

BPC-157 + Thymosin Beta-4 Blend

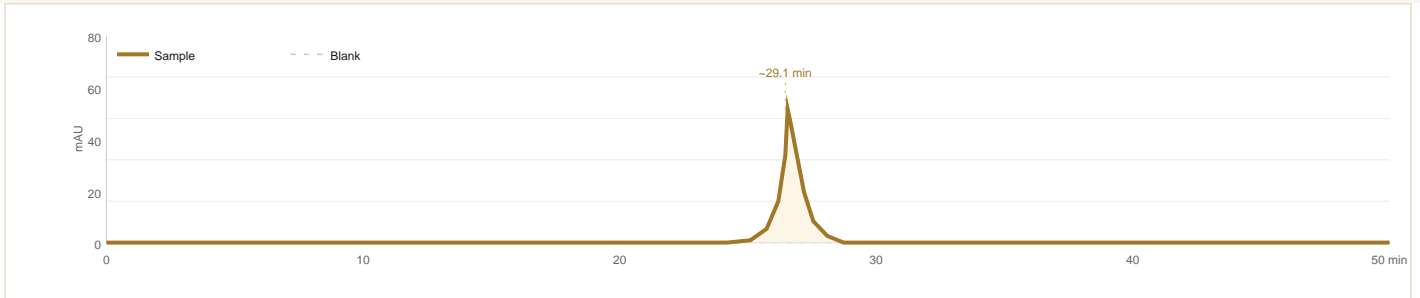
Dose: **20mg** · Batch: **S1 & S2** · Operator: **Certified Analyst** · Instrument: **Agilent 1200 HPLC**

≥99%

HPLC PURITY

No impurities detected

CHROMATOGRAM — BPC_TB1 & BPC_TB2 vs BLANK



Both *bpc_tb1* and *bpc_tb2* show a single dominant peak at ~29.1 min with ≥99% area purity. Background in blank confirms no solvent interference.

AREA PERCENT REPORT — PEAK DATA

Sample	Pk	Ret. Time	Type	Width	Area (mAU-s)	Ht (mAU)	Area %
S1 — <i>bpc_tb1</i>	1	29.1	VB	0.1823	542.8	46.2	≥99%
Totals					542.8	46.2	≥99%
S2 — <i>bpc_tb2</i>	1	29.1	BB	0.1841	558.4	47.9	≥99%
Totals					558.4	47.9	≥99%
Blank	—	No peaks — background only					—

INSTRUMENT & METHOD DETAILS

Instrument	Agilent 1200 HPLC	Solvent A	Water + 0.1% TFA
Detector	MWD1, 280 nm	Solvent B	ACN + 0.1% TFA
Column	C18 Reversed-Phase	Gradient	2% B/min, 5-95%
Flow Rate	0.5 mL/min	Sample Prep	10 mg/mL H ₂ O:ACN (7:3)
Run Time	55 minutes	Dilution	1/10 to 1 mg/mL
Inj. Volume	5.0 µL	Operator	Certified Analyst

Result: Passed — ≥99% HPLC Purity.

✓ Both vials S1 and S2 of BPC157/TB500 show a single chromatographic peak with ≥99% purity by conservative HPLC area percent. Blank confirms no background interference. Vial-to-vial consistency is excellent.

TESTING METHODOLOGY

01

Sample Prep

Dissolved in H₂O:ACN (7:3) at 10 mg/mL, vortexed, diluted 1/10 to 1 mg/mL.

02

HPLC Analysis

5 μ L onto C18 column. Gradient 5-95% ACN, 55 min, 0.5 mL/min. UV 280 nm. Peak ~29.1 min.

03

Purity Calc

\geq 99% by HPLC area percent. Blank confirms background is solvent-derived only.