Accumark Labs

CERTIFICATE OF ANALYSIS

Paramount Peptides
https://paramountpeptides.com

SAMPLE NAME:	BPC-157 + TB500
SAMPLE CODE:	8S2R-RZDE
RECEIVED DATE:	09/21/2026
PUBLISHED DATE:	09/22/2026



DECLARED IDENTITY	BPC-157, TB500 (17-23 Fragment)
MATRIX TYPE:	PEPTIDE BLEND
SAMPLE NAME:	BPC-157 + TB500
SAMPLE SIZE:	40.0 mg
Nº LOT CODE:	10KWLV09082026

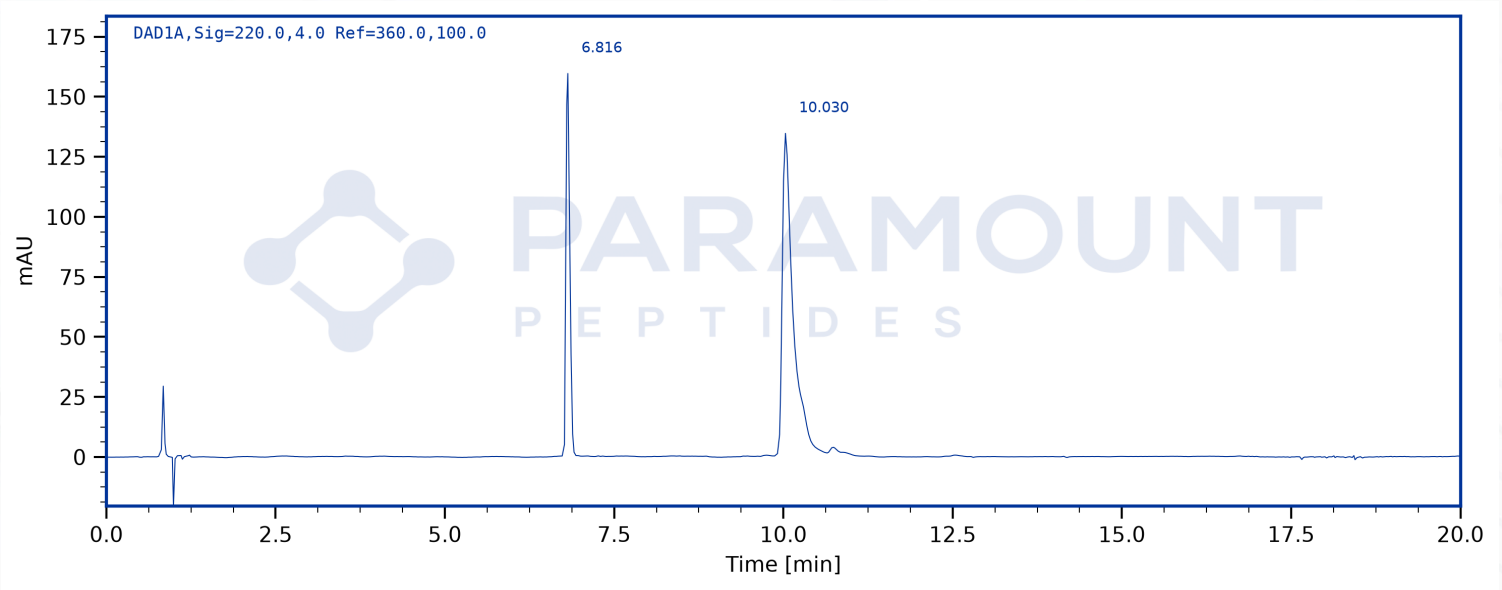
Variance
4 Vials

43.46 mg
Total Quantity

99.62%
Total Blend Purity

Sample	Test	Specification	Result	Status
Peptide Blend	IDENTITY	BPC-157, TB500 (17-23 Fragment)	BPC-157, TB500 (17-23 Fragment)	CONFORMS
Peptide Blend	QUANTITY	MEASURE	mean 43.46 mg · SD 0.44 · %RSD 1.02% · n=4	MEASURED
Peptide Blend	PURITY	> 98.0%	mean 99.62% · SD 0.05 · %RSD 0.05% · n=4	CONFORMS

HPLC Chromatogram Report



RESULTS REVIEWED BY:

Anthony Burke PhD
Lab Manager
09/22/2026

RESULTS CERTIFIED BY:

Dr. Levi Fried MD
Lab Director
09/22/2026

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Analyte	Test	Specification	Result	Status
BPC-157	IDENTITY	BPC-157	Conforms 4/4	Conforms 4/4
BPC-157	QUANTITY	MEASURE	mean 20.79 mg · SD 0.17 · %RSD 0.81% · n=4	MEASURED
BPC-157	PURITY	> 98.0%	mean 99.26% · SD 0.11 · %RSD 0.11% · n=4	CONFORMS
TB500 (17-23 Fragment)	IDENTITY	TB500 (17-23 Fragment)	Conforms 4/4	Conforms 4/4
TB500 (17-23 Fragment)	QUANTITY	MEASURE	mean 22.67 mg · SD 0.28 · %RSD 1.22% · n=4	MEASURED
TB500 (17-23 Fragment)	PURITY	> 98.0%	mean 99.95% · SD 0.02 · %RSD 0.02% · n=4	CONFORMS

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VIAL	TB500 (17-23 Fragment)		
	Identity	Quantity mg	Purity %
Vial 1	TB500 (17-23 Fragment)	22.884 mg	**99.931%**
Vial 4	TB500 (17-23 Fragment)	22.806 mg	**99.946%**
Vial 5	TB500 (17-23 Fragment)	22.265 mg	**99.931%**
Vial 6	TB500 (17-23 Fragment)	22.709 mg	**99.974%**

VIAL	BPC-157		
	Identity	Quantity mg	Purity %
Vial 1	BPC-157	20.876 mg	**99.229%**
Vial 4	BPC-157	20.872 mg	**99.21%**
Vial 5	BPC-157	20.536 mg	**99.425%**
Vial 6	BPC-157	20.876 mg	**99.194%**

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Endotoxin USP85 LAL

Test	Specification (EU/mL)	Result (EU/mL)	LOQ (EU/mL)	Verdict
Endotoxin USP85 LAL	≤5	< LOQ	2.5	**Conforms**

Method: Endotoxin testing performed using Limulus Amebocyte Lysate assay in accordance with USP <85> under validated laboratory conditions.

Rapid Sterility Screening (PCR)

Test	Specification	Result	Unit	Verdict
Rapid Sterility Screening (PCR)	Not Detected	Not Detected		**Conforms**

Method: Rapid sterility screening performed by real-time polymerase chain reaction (qPCR) targeting conserved microbial DNA sequences under validated laboratory conditions.

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09/22/2026

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