

CERTIFICATE OF ANALYSIS / BATCH RELEASE & IDENTITY VERIFICATION

BPC–157 10mg

Synthetic pentadecapeptide (**body-protection compound / BPC–157**) · sterile-filled lyophilised powder, released against independent third-party HPLC–UV, ICP–MS heavy-metal, endotoxin and sterility analysis.

PRODUCT	LOT · BATCH	TESTING LABORATORY	REPORT DATE
BPC-157 10mg White lyophilised powder	EL-BPC10-0626 Single production lot	Independent Third-party laboratory	4 June 2026

HPLC–UV PURITY 99.09% Main-peak area · reported $\pm 0.18\%$ • MEETS RELEASE SPEC	IDENTITY HPLC assay & retention BPC–157 Confirmed
	NET PEPTIDE CONTENT HPLC–UV/VIS assay 11.34mg ≥ 10mg
	BACTERIAL ENDOTOXIN LAL · see page 2 Pass 7.70 EU/mg

01 PURITY · IDENTITY · CONTENT

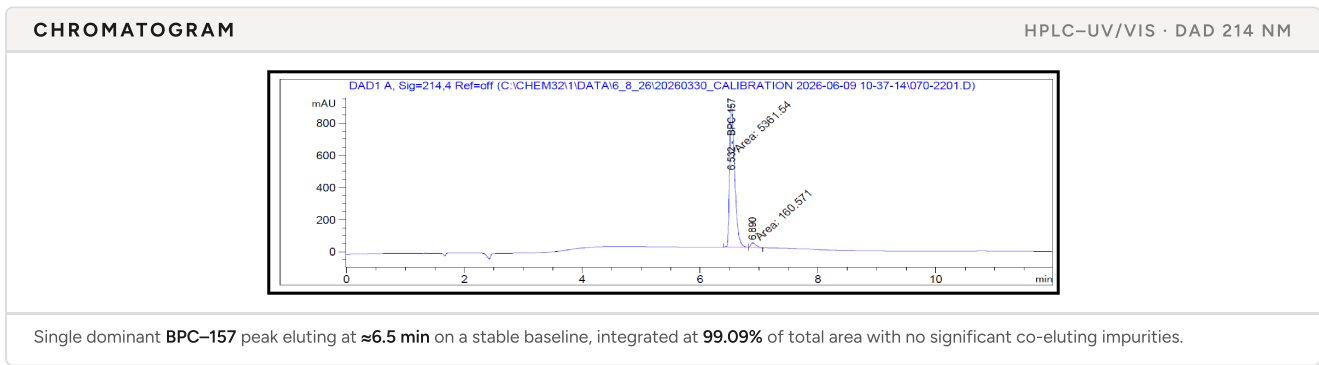
Analytical Results

TEST	METHOD	RESULT	RELEASE SPECIFICATION	STATUS
Identity Main-peak assignment	HPLC–UV/VIS	BPC–157 Single dominant peak · ≈ 6.5 min	Single dominant peak	PASS
Chromatographic Purity Main-peak area %	HPLC–UV/VIS	99.09% Reported $\pm 0.18\%$	$\geq 98.0\%$	PASS
Net Peptide Content Assay vs. label claim	HPLC–UV/VIS	11.34mg	≥ 10.0 mg (label claim)	PASS

Methods. Content and purity by high-performance liquid chromatography with UV/VIS detection (HPLC–UV/VIS) against a reference standard; chromatographic purity reported as main-peak area %. Identity is assigned from the single dominant HPLC peak and its assayed content (no orthogonal mass-spectrometric confirmation was performed for this batch). **Status** is assessed against acceptance criteria set by Eira London for research-grade (RUO) release. Elemental-impurity, endotoxin and sterility results are reported on **page 2**.

02 INSTRUMENT TRACE

Chromatogram



Released for research use. This batch conforms to Eira London's research-grade release specification on the basis of the independent analytical data above and the contaminant, endotoxin and sterility data on page 2. Certificate refers only to the lot and sample(s) tested.

Eira LondonAUTHORISED — QUALITY & COMPLIANCE
Batch released 4 June 2026**eira london**

MODERN HEALTH SCIENCE, REFINED

eiralondon.com

Batch COA available on request

Quote lot EL-BPC10-0626

Research Use Only. The material described is supplied strictly for laboratory research use and is **not** approved for human or veterinary use, nor for diagnostic, therapeutic or clinical application. Analysis was performed by an independent third-party laboratory under standard laboratory conditions; results are specific to the sample(s) provided and should be interpreted by qualified professionals within the scope of the intended research. Accuracy may be influenced by sample integrity, handling and other experimental variables. This certificate does not constitute a medical or clinical claim.

CONTINUED FROM PAGE 1 / LOT EL-BPC10-0626 · 4 JUNE 2026

Safety & Contaminant Screen

Independent third-party ICP-MS elemental-impurity, **bacterial endotoxin (LAL)** and **sterility (USP <71>)** analysis for **BPC-157 10mg**, batch EL-BPC10-0626.

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03 ELEMENTAL IMPURITIES

Heavy Metals · ICP-MS

ANALYTE	METHOD	LOQ (PPM)	RESULT (PPM)	STATUS
Arsenic (As) Elemental impurity	ICP-MS	0.01	ND	PASS
Cadmium (Cd) Elemental impurity	ICP-MS	0.01	ND	PASS
Lead (Pb) Elemental impurity	ICP-MS	0.02	ND	PASS
Mercury (Hg) Elemental impurity	ICP-MS	0.005	ND	PASS

Methods. Elemental impurities by inductively-coupled-plasma mass spectrometry (ICP-MS), reported in parts per million (ppm). Results accepted against USP <232>/<233> elemental-impurity limits; all four analytes were **non-detect**. **ND** = non-detect · **LOQ** = limit of quantification.

04 MICROBIAL · ENDOTOXIN

Sterility & Endotoxin

TEST	METHOD	RESULT	RELEASE SPECIFICATION	STATUS
Bacterial Endotoxin Pyrogen screen	LAL	7.70 EU/mg	≤ 5 EU/kg (≤350 EU/dose)	PASS
Sterility Direct inoculation	USP <71>	No growth detected	Sterile / no growth	PASS

Methods. Bacterial endotoxin by Limulus Amebocyte Lysate (LAL) assay; sterility by USP <71> direct inoculation. Endotoxin pass/fail is based on the USP/FDA threshold of 5 EU/kg (350 EU total for a 70 kg adult). **Status** is assessed against acceptance criteria set by Eira London for research-grade (RUO) release.

Released for research use. Read together with page 1 (purity, identity and content), the contaminant, endotoxin and sterility data above complete the batch-release record for lot EL-BPC10-0626. Certificate refers only to the lot and sample(s) tested.

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Batch released 4 June 2026

