Certificate of Analysis

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

BPC-157 - 10mg

Tested for: YOUR COMPANY NAME
Your website here

EXAMPLE

BRANDING PREVIEW

Not valid for batch verification

COA #:	EXAMPLE-COA	Analysis Date:	Example date
Lot Number:	YOUR-LOT	Appearance:	**Good**
Accession #:	EXAMPLE-SAMPLE	Sample Matrix:	**Lyophilized**
Labeled Content:	**10mg**	Volume:	**3mL**
		Date Received:	Example date

Method: **Full QC Panel**

Identity	Peptide Purity	
BPC-157	**99.88%**	Fentanyl Free

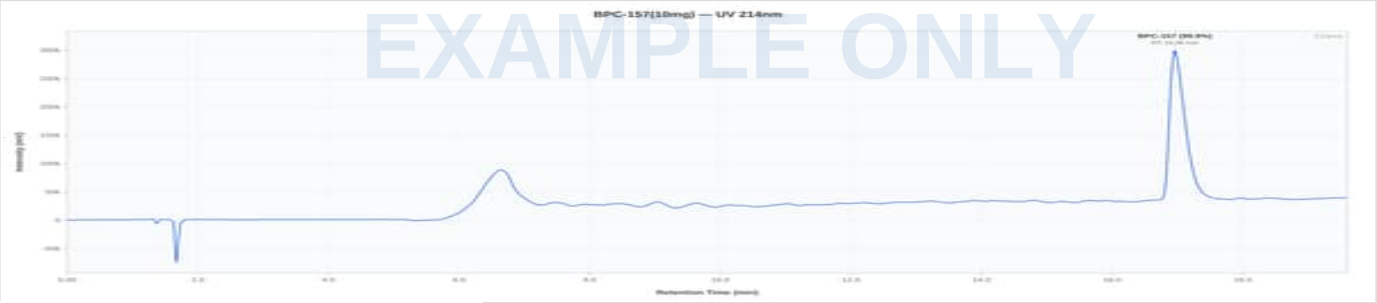
YOUR LABEL

YOUR VIAL PICTURE HERE

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.88%**	%	**PASS**
Net Peptide Content	Report Only	**10.57**	mg	N/A
Identity (HPLC-RTM)	BPC-157	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



BPC-157 10mg - YOUR LOT: 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 ppm	*Not Detected*	**PASS**
Mercury (Hg)	NMT 1.5 ppm	*Not Detected*	**PASS**
Chromium (Cr)	NMT 10 ppm	*Not Detected*	**PASS**

EXAMPLE ONLY - NOT A VALID CERTIFICATE

Source-report values illustrate the format. They are not results for your product or batch.

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>)	NMT 0.25 EU/mL	**0.077 EU/mL**	**PASS**
Acceptance criteria: NMT 0.25 EU/mL per client specification.			

EXAMPLE ONLY - NOT A VALID CERTIFICATE

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EXAMPLE ONLY

Notes & Methodology

1. Date Tested: Example date. Methods: Full QC Panel.
2. The sample was confirmed to be BPC-157 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 chromogenic 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. Accreditation Scope: ILS Laboratories maintains ISO/IEC 17025 accreditation for activities listed in its published PJLA scope. Peptide testing reported on this certificate is performed for Research Use Only.
7. This certificate reports analytical results for the sample(s) submitted to ILS Laboratories for testing. All sample identifications, product names, brand names, lot/batch numbers, and labeled content are provided by the submitting party and have not been independently verified. Results reflect only the analytical testing of the specific sample(s) provided and do not constitute an evaluation or endorsement of the manufacturer's processes, quality systems, or overall production. ILS Laboratories makes no representations regarding the intended use, safety, efficacy, regulatory status, or intellectual property ownership of any materials tested.

YOUR COMPANY NAME / YOUR VIAL

Final report, signature and verification are issued by the laboratory after testing.

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This certificate relates only to the sample(s) tested.