ILS Laboratories

Verified Certificate

Certificate of Analysis

All tests passed. This certificate is verified and authentic.

TESTED FOR

Panda Peptides
pandapeptides.com

Product Information

Product Lipo-C with B12 216mgmL - 216mg	Lot Number PP2605-029
Appearance Good	Test Type Full QC Panel
Date Received 09/08/2026	Analysis Confirmed 09/16/2026

Confirmed Fentanyl Free
Immunoassay, 50 ng/mL cutoff

Test Results — Full QC Panel

Analyte	Limit	Result	Unit	Status
Peptide Purity (HPLC)	≥ 95.0%	99.82%	%	PASS
Net Peptide Content	As Labeled	216	mg	N/A
Identity (ID)	Lipo-C	Confirmed	-	PASS
Arsenic (As)	≤ 1.5 ppm	Not Detected	ppm	PASS
Cadmium (Cd)	≤ 0.5 ppm	Not Detected	ppm	PASS
Lead (Pb)	≤ 1.0 ppm	Not Detected	ppm	PASS
Mercury (Hg)	≤ 1.5 ppm	Not Detected	ppm	PASS
Chromium (Cr)	≤ 10.0 ppm	Not Detected	ppm	PASS

Sterility Testing (PCR)

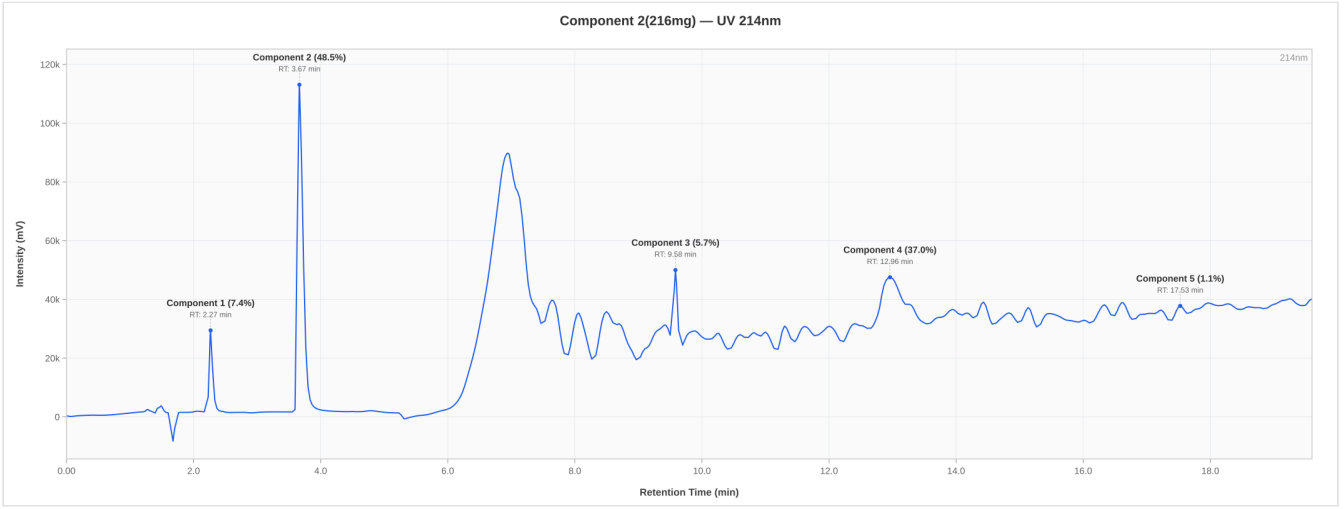
Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS

Endotoxin Testing — USP ⟨85⟩

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	≤ 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.

HPLC Chromatogram



Representative chromatogram, Conformity V1 (closest to batch mean purity)

Sample	Purity	Net Content	Identification	Result
Dedicated V0	99.82%	216 mg	Confirmed	PASS
★ Conformity V1	99.47%	216 mg	Confirmed	PASS
Conformity V2	97.58%	216 mg	Confirmed	PASS

STATISTICAL SUMMARY

Purity (HPLC)	Net Peptide Content	Endotoxin (USP <85>)
Mean: 98.96 %	Mean: 216.00 mg	Mean: 0.02 EU/mL
Std Dev: 0.9839 %	Std Dev: 0.0000 mg	Std Dev: 0.0000 EU/mL

★ Representative sample: Conformity V1 (closest to batch mean purity). The chromatogram above corresponds to this sample.

NOTES & METHODOLOGY

1. Date Tested: 09/16/2026. Methods: Full QC Panel.

2. The sample was confirmed to be Lipo-C with B12 216mgmL 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. Sterility testing performed by PCR (Polymerase Chain Reaction) method.

6. 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.

7. 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.

8. 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.

Dr. Greg Kalyuzhny
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09/16/2026

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