

ILS Laboratories

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(619) 329-3999 | ils-lab.com

NAD+ - 500mg

Tested for: RevPep
Revolutionpeptide.com

COA #: COA-2026-955329
Lot Number: NAD010
Labeled Content: 500mg
Volume: 10mL

Analysis Date: 08/05/2026
Appearance: Good
Volume: 10mL
Date Received: 07/10/2026

Method: Full QC Panel

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: NPLEBKGM

Identity

NAD+

Purity

99.46%

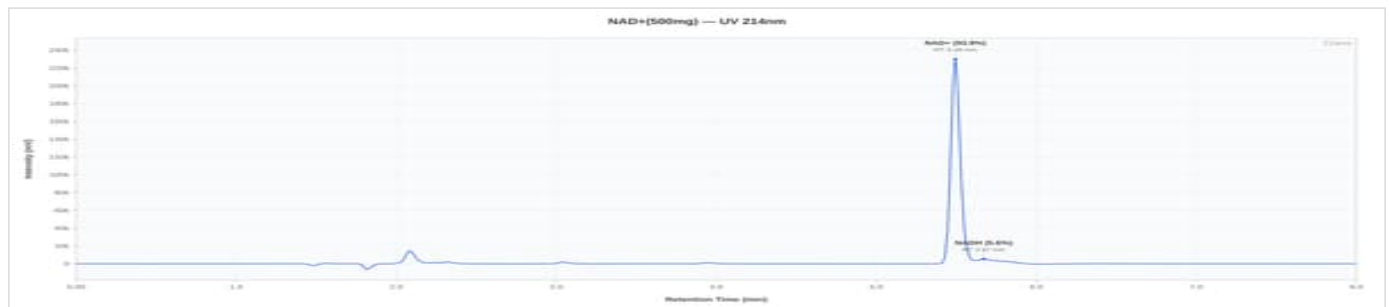


Fentanyl
Free

Full QC Panel

Analyte	Specification	Result	Unit	Status
Purity (HPLC)	>= 95.0%	99.46%	%	PASS
Net Peptide Content	Report Only	509.62	mg	N/A
Identity (HPLC-RTM)	NAD+	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



NAD+ 500mg - NAD010: 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



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
8/8/2026

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Access Code: NPLEBKGM
Verify: portal.ils-lab.com/verify/I-LAvqQu0INtybs
Issued: 8/8/2026

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	0.154 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.

Notes & Methodology

- 1. Date Tested: 08/08/2026. Methods: Full QC Panel.
- 2. The sample was confirmed to be NAD+ 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 turbidimetric method. Acceptance criteria per client specification.
- 5. Purity determined by RP-HPLC area normalization at 214 nm.




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