Certificate of Analysis

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

NAD+ - 1000mg

PASS

Tested for: Crush Research
https://www.crushresearch.shop/

COA #:	**COA-2026-9_RBYQ**	Method:	**Full QC Panel**
Lot Number:	**CR-NAD-1000MG-2605-2**	Analysis Date:	**06/01/2026**
Accession #:	**ACC-2026-3014**	Appearance:	**Good**
Labeled Content:	**1000mg**	Vial Size:	**10mL**
Sample Matrix:	**Lyophilized**	Date Received:	**05/22/2026**

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

Identity	Purity	
NAD+	99.22%	Fentanyl Free

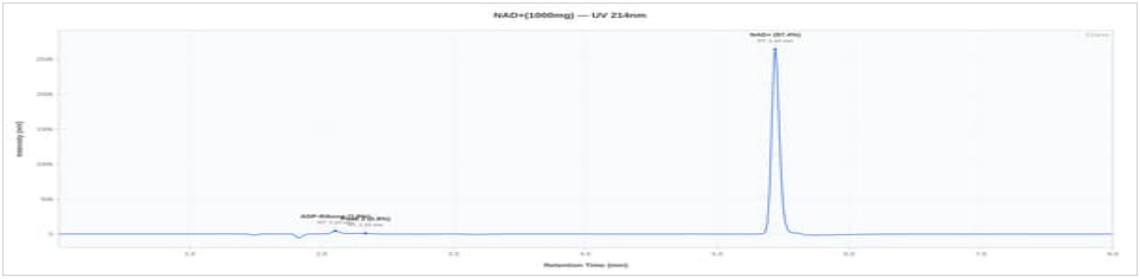


NAD+ 1000mg - CR-NAD-1000MG-2605-2

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Purity (HPLC)	>= 95.0%	**99.22%**	%	**PASS**
Net Peptide Content	Report Only	**1049.15**	mg	N/A
Identity (HPLC-RTM)	NAD+	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.17% purity, closest to batch mean of 99.15%)

Dr. Greg Kalyuzhny
Lab Director
6/1/2026

COA #: **COA-2026-9_RBYQ**
Access Code: **KNQCYMVR**
Verify: portal.ils-lab.com/verify/KPscL34f69Z0cKWp
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HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.17%	1044.86 mg	Confirmed	**PASS**
Conformity V1	99.22%	1049.15 mg	Confirmed	**PASS**
Conformity V2	99.07%	1015.82 mg	Confirmed	**PASS**
Mean	**99.15%**	**1036.61 mg**	—	—
Std Dev	**0.0624%**	**14.8047 mg**	—	—

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**
Chromium (Cr)	NMT 10 ppm	*Not Detected*	**PASS**
Mercury (Hg)	NMT 1.5 ppm	*Not Detected*	**PASS**
Lead (Pb)	NMT 1 ppm	*Not Detected*	**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	**NMT 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.

Dr. Greg Kalyuzhny
Lab Director
6/1/2026

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Accession #:	**ACC-2026-3014**	Appearance:	**Good**
Labeled Content:	**1000mg**	Vial Size:	**10mL**
Sample Matrix:	**Lyophilized**	Date Received:	**05/22/2026**

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Notes & Methodology

1. Date Tested: 06/01/2026. Methods: Purity & Quant (HPLC).
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.
6. Chromatogram shown is representative: Dedicated V0 (99.17% purity, closest to batch mean of 99.15%).

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Lab Director
6/1/2026

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