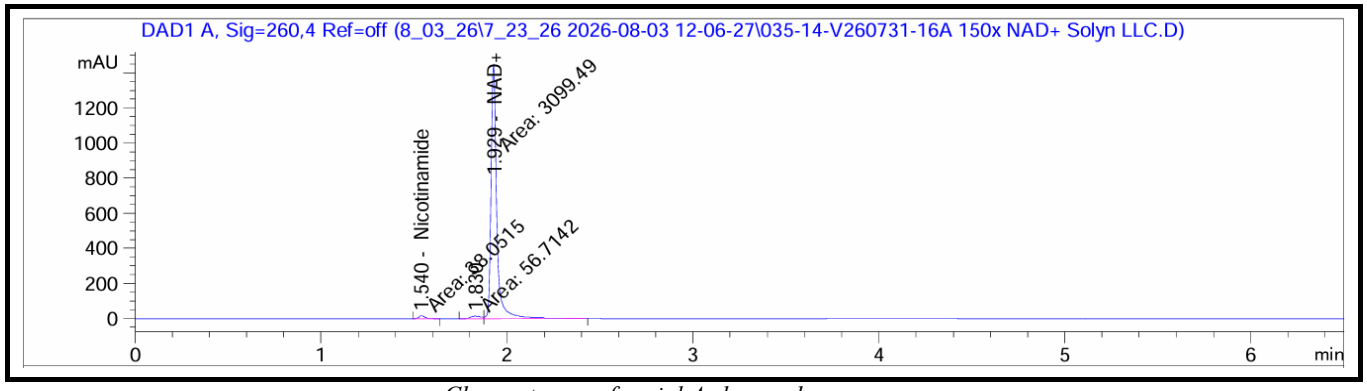
Vanguard Laboratory
2635 Parkmont Ln
Olympia, Wa 98502
360-967-7010

Certificate of Analysis

NAD+ 500 mg

Report To:
Solyn LLC

Compound: NAD+
Amount: 500 mg
Laboratory ID: V260731-16 001
Date Reported: 8/16/2026
Lot Number: 2607-213



Chromatogram for vial A shown above.

Analysis	Method	Result (per Vial)		
Identity: Retention Time	HPLC-UV/VIS	Pass - Matches Ref. Standard		
Identity: Spectral Analysis	HPLC-UV/VIS	Pass - Matches Ref. Standard		
Chromatographic Purity (Total)	HPLC-UV/VIS	>99.80	>99.80	>99.80
Chromatographic Purity (NAD+)	HPLC-UV/VIS	97.03%	97.23%	96.48%
Chromatographic Purity (Nicotinamide)	HPLC-UV/VIS	2.97%	2.77%	3.52%
Quantity: NAD+	HPLC-UV/VIS	729 mg	723 mg	748 mg
Fentanyl Screening	HPLC-UV/VIS	Pass - Not Detected		



ISO 17025:2017 ACCREDITED LABORATORY

Report By: Dustin Newman, Laboratory Director on 8/16/2026

Approved By: Tori Johnson, Operations Manager on 8/16/2026

Please consult A2LA Certificate #6377.01 for a list of accredited tests. Samples were received in acceptable condition. The result(s) in this report relate only to the portion of the sample(s) tested. All analyses were performed consistent with the Vanguard Laboratory Quality Management System. Vanguard Laboratory and its staff did not observe or participate in the sample selection process, and cannot confirm the authenticity of the sample or its representativeness of the associated lot/batch.

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Analyte	Method	LOQ (ppm)	Result (per Vial)	
Chromium (Cr)	ICP-MS	0.05	ND	ND
Arsenic (As)	ICP-MS	0.01	ND	ND
Cadmium (Cd)	ICP-MS	0.01	ND	ND
Lead (Pb)	ICP-MS	0.02	ND	ND
Mercury (Hg)	ICP-MS	0.005	ND	ND

Run ID:

Analysis	Method	Result (per Vial)	
Endotoxins	LAL	**A**	Pass - <5.00 EU/mg*
		B	Pass - <5.00 EU/mg*

*Pass/Fail criteria based on the USP/FDA threshold of 5 EU/kg (350 EU total for a 70 kg adult)

Analysis	Method	Result (per Vial)	
Sterility	USP <71>	**A**	Pass - No Growth
		B	Pass - No Growth
		C	Pass - No Growth
		D	Pass - No Growth

Report By: Dustin Newman, Laboratory Director on 8/16/2026

Approved By: Tori Johnson, Operations Manager on 8/16/2026

ND: Non-Detect
LOD: Limit of Quantification

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Certificate of Analysis

NAD+ 500 mg

Report To:
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Compound: NAD+
Amount: 500 mg
Laboratory ID: V260731-16 001
Date Reported: 8/16/2026
Lot Number: 2607-213

Residual Solvents Analysis			
Analyte	**Method**	**LOQ (ppm)**	**Result (ppm)**
Acetone	GC-MS	3.7	ND
Benzene	GC-MS	0.02	ND
Butanes	GC-MS	3.7	ND
Chloroform	GC-MS	0.02	ND
Cyclohexane	GC-MS	2.9	ND
Dichloromethane	GC-MS	0.45	ND
Ethanol	GC-MS	3.7	ND
Ethyl Acetate	GC-MS	3.7	ND
Heptane	GC-MS	3.7	ND
Hexanes	GC-MS	0.21	ND
Isopropanol	GC-MS	3.7	ND
Methanol	GC-MS	2.2	ND
Pentanes	GC-MS	3.7	ND
Propane	GC-MS	3.7	ND
Toluene	GC-MS	0.66	ND
Total Xylenes	GC-MS	1.6	ND

Analysis	**Method**	**Result (per Vial)**	
Container Closure Integrity	Dye Ingress Method	**A**	Pass
		B	Pass
Solubility Analysis	Fully Soluble in 2 mL Water	**A**	Pass
Trifluoroacetic Acid (TFA)	Ion Chromotography	ND - <0.10%	

Report By: Dustin Newman, Laboratory Director on 8/16/2026

Approved By: Tori Johnson, Operations Manager on 8/16/2026

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