

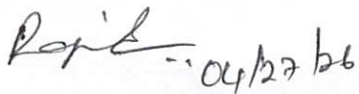


Certificate of Analysis

Product: NAD+ (5mL) 200mg/mL	
Compounding Date: 04/16/2026	Lot ID: LG52243282
Beyond Use Date: 07/15/2026	Product ID: 305471994
BUD Length: 90 Days	Quantity Yield: 250 Vials

Physical Appearance:	5 ml of clear solution in amber vial
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Test	Acceptance Criteria	Results	Compliance Statement
Endotoxin Test USP <85>	350 EU/mL	<2.50 EU/ml	PASS
Sterility-ATP Bioluminescence Test	No Evidence of growth	No Evidence of growth	PASS
Potency	90.0 - 110.0 %	N/A	N/A
Hood Surface Sampling	NMT 3 cfu/Plate	< 3 cfu/Plate	PASS
Glove Fingertip and Thumb Testing	NMT 3 cfu/Plate	< 3 cfu/Plate	PASS

Released By (Quality)	Approved By (Pharmacist)
 04/27/26	 4/27/26

Notes:



PHARMETRIC LABORATORY

CERTIFICATE OF ANALYSIS (CoA)

Pharmetric Laboratory
7265 Ulmerton Road
Largo, FL, 33771

Issue #:	0	SAMPLE #:	S424060
CoA RELEASE DATE:	22 Apr 2026		

CLIENT:	Vios Compounding Pharmacy	CONTACT:	Victor Hill
CLIENT ADDRESS:	31035 Schoolcraft Rd, Livonia, MI, 48150		
DESCRIPTION:	Nicotinamide Adenine Dinucleotide. Oxidized (NAD+) 200 mg (5ml)	LOT #:	LG52243282
STRENGTH:	200mg/mL	CONTAINER:	5.0 ml Vial
STORAGE CONDITIONS:	Refrigerated (2°C to 8°C)	BATCH SIZE:	200
DATE(S) RECEIVED:	17 Apr 2026	EXP.:	Not Provided
		QUANTITY RECEIVED:	11

TEST DATE	TEST	QTY	ACCEPTANCE CRITERIA	RESULTS	COMPLIANCE STATEMENT
20 Apr 2026	USP <85> Endotoxin Test	1	350 EU/mL	< 2.50 EU/mL	PASS
22 Apr 2026	Sterility-ATP Bioluminescence Test	10	No Evidence of growth	No Evidence of growth	PASS

STERILITY TEST (BATCH & SIZE COMPLIANCE)	USP 71 COMPLIANT	METHOD SUITABILITY ON FILE	Yes
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Samples accepted by Pharmetric Lab are assumed to be prepared by the customer within the guidelines of USP <71> regarding sample size and number of units sampled. Pharmetric conducts sterility testing using bioluminescence technology. Per USP <797> "A method not described in the USP may be used if verification results demonstrate that the alternative method is at least effective and reliable as the USP membrane filtration method as described in USP<71>." Sterility Testing performed by Pharmetric Lab using bioluminescence technology was validated as equal to compendial test by direct comparison. Validation documents are on file at Pharmetric Lab, LLC.

TEST DATE refers to start date. Please note some tests are carried out over multiple days.

The results on this CoA relate only to the item(s) tested and apply to the sample as received. This Report shall not be reproduced except in full without approval of the laboratory.



ACCREDITED TESTING LABORATORY. ACCREDITATION NUMBER 5386.01

COMMENTS	QUALITY SIGNATURE
	Yovany Gomez Quality Assurance Associate 1 4/22/2026 2:58 PM