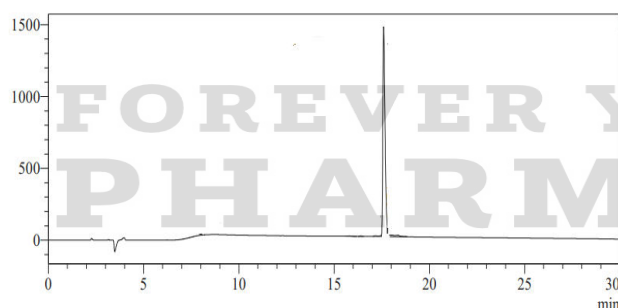


**CERTIFICATE OF ANALYSIS****SAMPLE INFORMATION**

Product Name	NAD+ 500 mg (Nicotinamide adenine dinucleotide)
Client Name/Lot No.	AvivaAmino
Sequence	C21H28N7O14P2
Dissolution condition	100% H2O
Length	13AA
Molecular Weight	664 g/mol

CHROMATOGRAM

mV



Peak #	Ret. Time	Area %
1	8.094	0.119
2	17.722	99.683
3	18.806	0.124
4	20.255	0.074

TEST RESULTS

	Specifications	Results
Strength	500.00 mg	548.61 mg
Appearance	White to off white lyophilized powder	Conforms
Purity	≥98.0%	99.7%
pH value	4.0-8.0	7.0
Impurity	Single Impurity ≤1.0%	0.1%
	Total Impurity ≤2.0%	0.3%

TEST PARAMETERS

Pump A	0.1% trifluoroacetic in 100% water
Pump B	0.1% trifluoroacetic in 100% acetonitrile
Total Flow	1.0ml/min
Wavelength	214nm
Analytical Column Type	Agilent ZORBAX StableBond 5µm C18 (4.6*250mm*5 µm)
Dissolution Method	100% H2O
Injection Volume	20uL

CONCLUSION

One 10ml amber vial contained a white lyophilized powder and has a blue cap with a silver crimp.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 99.7% NAD+ (548.61), and the rest are impurities of minor significance.

CERTIFIED BY:

Dane Fredericksen
Analytical Chemist
06/03/2025



****Verify the validity of test results by contacting support@foreveryoungpharmacy.com****