

Randy Cole
Regenerative Research
1475 S. Fillmore St
Denver, CO 80210



Project 135241
Lab # 127996
Date Rec'd 8/15/2025
Report Issued 8/25/2025

Project

Sample NAD+ 500 mg ND25-02

Certificate of Analysis

<u>Analyte</u>	<u>Result</u>	<u>LOQ</u>	<u>Units</u>	<u>Method</u>	<u>Date</u>	<u>% of Label</u>
Microbiology						
Endotoxins	ND	0.5	EU/mL	USP 85	8/22/2025	
Peptide Analysis						
Purity by area	99.95	0.5	%	HPLC-UV/MS	8/22/2025	
Purity by mass	500	0.5	mg	HPLC-UV/MS	8/22/2025	99.94
Spectral identification	Confirmed			HPLC-UV/MS	8/22/2025	

The data presented are from the analysis of the sample shown and meet Krause Analytical internal quality assurance criteria unless otherwise flagged.
Methods shown reference current Krause Analytical SOPs
ND - Not detected LOQ - limit of quantification
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Respectfully submitted,

Mark C. Krause
Laboratory Director

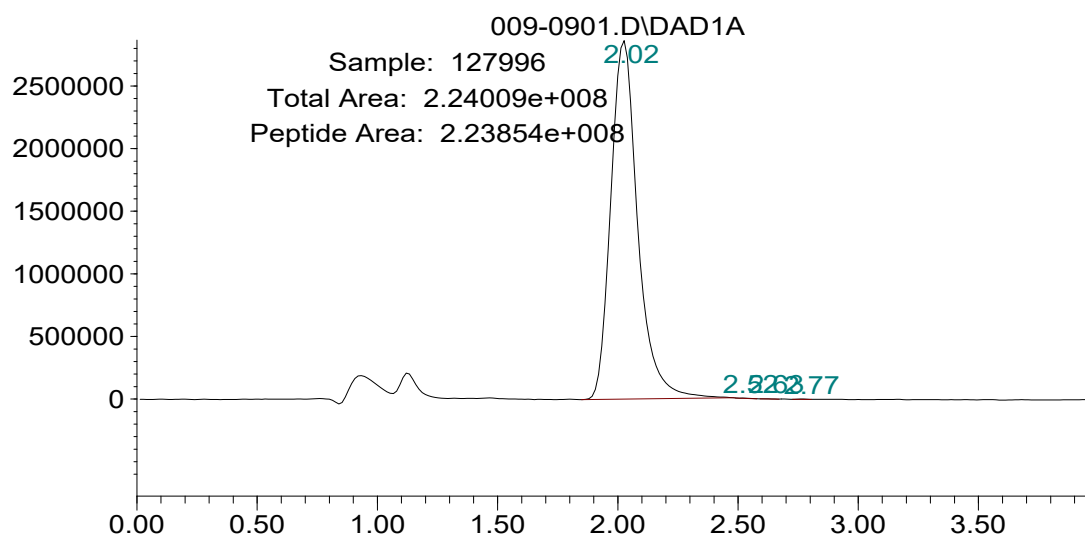


krause analytical

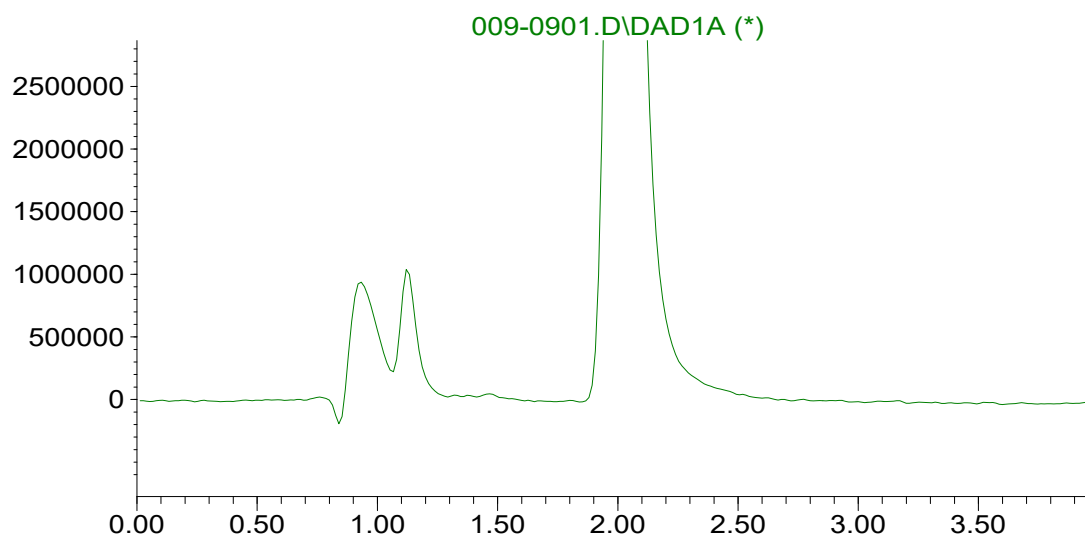
Lab #:127996 NAD+ 500 mg

Chromatogram

Response_

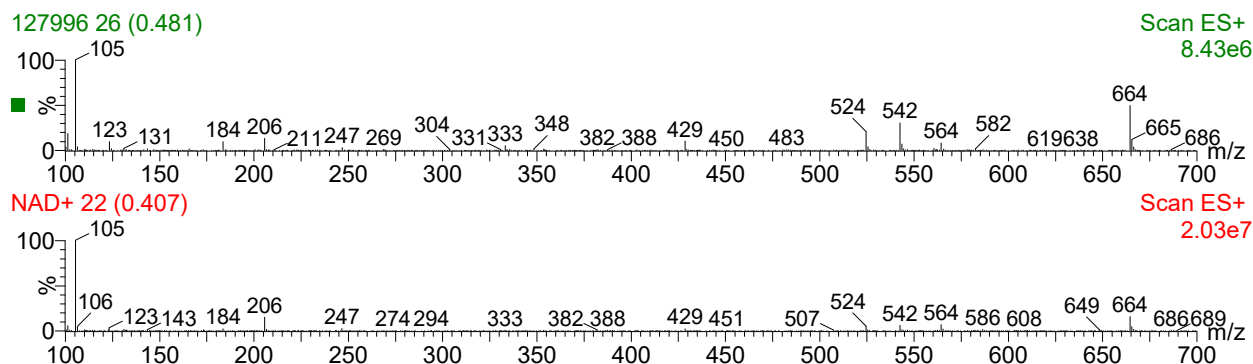


Time
Response_





Spectra



Method Summary

- 2 mL of purified water is added to the lyophilized powder in the vial, and the contents mixed to dissolve the lyophilized powder.
- An aliquot is taken from the vial and diluted to contain approximately 500 mg/L of the peptide.
- The diluted sample is analyzed by HPLC-UV-MS.
- The mass spectrum obtained is compared to an authentic standard of the peptide for identification.
- The total area of all of the peaks in the chromatogram is calculated, and the area of the peak of the peptide is divided by the total area to obtain the purity by area value, reported in percent.
- The area of the peptide is compared to the area of the peptide peak in the known standard to obtain a concentration in the solution. This concentration is used to calculate the total mass of peptide in the vial, which is compared to the stated mass (label claim) and reported as both total mass in the vial and as a percent of the label claim.