VERIFIED VIALS

VV-COA-NAD-001
Document reference

19 October 2026
Date issued

Certificate of Analysis

NAD+ · Research Grade Peptide — Purity & Identity (HPLC + LC-MS)

Issued by Verified Vials on the basis of testing carried out by an independent third-party laboratory, PeptideVerify.

01 PRODUCT

PRODUCT
NAD+ · Research Grade Peptide

INTENDED USE
Laboratory research use only

SUPPLIED BY
Verified Vials, Belfast, Northern Ireland

SAMPLE NAME ON LABORATORY RECORD
NAD+

BATCH NUMBERS & EXPIRY
500 mg NAD500-260626-A · 1000 mg NAD1000-260626-A · Expiry 26 June 2029

02 RESULT

PURITY — HPLC (UV)

98.10 %

3 peaks detected · minor peaks included in the purity figure

✓ IDENTITY CONFIRMED

LC-MS identity check: the detected ion pattern matches the laboratory's reference for NAD+.

Purity reflects UV-detectable compounds only; non-UV excipients are not measured by this method.

03 TEST DETAILS

LABORATORY
PeptideVerify (independent of the supplier)

LABORATORY CERTIFICATE
PV-C9D61B-KAZS-73C

LABORATORY CERTIFICATE ISSUED
17 June 2026

TEST TYPE
HPLC + LC-MS

METHOD
HPLC + ESI-MS

ANALYSIS DATE
16 June 2026

INSTRUMENT
Shimadzu LCMS-8040

SAMPLE AS RECEIVED
White lyophilised powder

CONDITION ON RECEIPT
As Expected

04 IDENTITY & SYSTEM SUITABILITY

PARAMETER	OBSERVED	ACCEPTANCE / REFERENCE	OUTCOME
MS identity match (ESI-MS)	Ion pattern matches reference	Lab reference: NAD+	CONFIRMED
Detected molecular weight	663.2 Da	Expected MW: 663.4 Da	—
Main peak retention time	6.635 min	Total peaks detected: 3	—
Resolution (Rs)	0.87	not applicable	N/A · HILIC
Theoretical plates (N)	202.00	not applicable	N/A · HILIC

Tailing factor (T) was reported by the laboratory as not applicable for this sample (limit ≤ 3.5). The laboratory notes that minor peaks are common in synthetic peptides, typically synthesis-related by-products such as incomplete sequences or minor structural variants, and are included in the purity calculation. HILIC method: the laboratory separates this polar compound by hydrophilic-interaction chromatography, where peaks are inherently broad; reverse-phase plate-count and resolution limits do not apply, and identity is established by LC-MS intact mass. Sample received as a white lyophilised powder, condition as expected. Purity is the HPLC purity reported by the laboratory and reflects UV-detectable compounds only; non-UV excipients (mannitol, sugars, salts) are not measured.

05 VERIFICATION & AUTHORISATION

peptideverify.co.uk/verify/PV-C9D61B-KAZS-73C

The analysis was carried out by PeptideVerify. Their certificate PV-C9D61B-KAZS-73C is the source record for every figure on this page and can be checked at the address above or by scanning the code.

This document is issued by Verified Vials and summarises that laboratory record. It is not a PeptideVerify document.

AUTHORISED BY
Quality Control
Verified Vials
19 October 2026

SIGNATURE

For laboratory research use only. Not for human or veterinary use. This certificate relates to the sample as received by the laboratory and summarises the laboratory's reported findings for that sample. It is not a guarantee of other units or batches, is not a safety assessment, and is not medical advice. Where this summary and the laboratory record differ, the laboratory record prevails.