Aura Pharmaceutical Group

A Hong Kong Private Company | Companies Ordinance (Cap. 622)

CERTIFICATE OF ANALYSIS

Analytical Test Report

AL-COA-2026-08015

TASK NUMBER	PRODUCT SKU	BATCH / LOT	DISPOSITION
AL-8015	NJ500	AL-NJ500-2026-05	PASS — Released

SAMPLE & CLIENT

Client (ordered by)	**Aura Labs Asia**	**Manufacturer**	Aura Labs (GMP contract manufacture)
Sample	**NAD+ - 500mg**	**Batch**	AL-NJ500-2026-05
Product SKU	NJ500	**Form**	Lyophilized peptide powder
Dosage / strength	500mg	**Expiry (sealed)**	14 MAY 2028

TIMELINE

Testing ordered	07 APR 2026	**Sample received**	27 APR 2026
Analysis conducted	22 APR 2026	**Report issued**	24 JUL 2026

SAMPLE DESCRIPTION

Sealed research vial(s) of lyophilized peptide powder, labeled BPC-157 10mg. Visual inspection: white to off-white cake/powder. Pictures not included.

TESTS REQUESTED

- Identity confirmation by mass spectrometry (MS)
- Purity determination by reverse-phase HPLC (UV 220 nm)
- Net peptide content / assayed quantity
- Assessment of a peptide vial for research-use release

RESULTS

ANALYTICAL SUMMARY — Identity ESI-MS · Purity HPLC

NAD+ — **500 mg**

HPLC PURITY	IDENTITY	APPEARANCE
99.250%	Confirmed — NAD+	White lyophilized powder

COMMENTS

Sample meets Aura Labs release specification (≥99.0% HPLC purity; identity confirmed by MS). Lot released for research use only.

Quality Assurance

Authorized Signatory — Quality Control

Aura Labs Quality Control · Analysis conducted › 22 APR 2026

This certificate relates only to the sample tested and to the batch identified herein.

AL-MS20-17028A10

Scan QR or open auralabs.asia/verify

RUO. For laboratory and research use only (RUO). Not for human or veterinary use, diagnosis, treatment, or consumption. Not a drug, food, or cosmetic. Handle in accordance with institutional laboratory safety protocols.

COMPOUND IDENTIFIERS

Compound	NAD+	**CAS**	137525-51-0
Molecular formula	C62H98N16022	**MW (Da)**	1419.53
Sequence	Gly-Glu-Pro-Pro-Pro-Gly-Lys-Pro-Ala-Asp-Asp-Ala-Gly-Leu-Val		

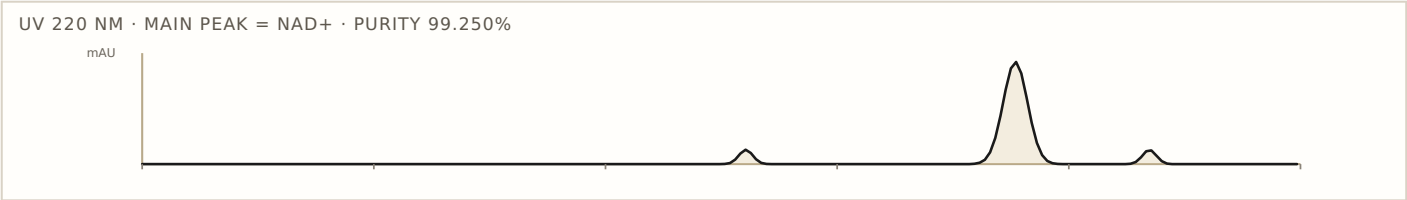
IDENTITY — MASS SPECTROMETRY

Method	ESI-MS
Theoretical mass	1419.53 Da
Observed mass	1419.48 Da
Mass error	-35.2 ppm
Conclusion	**Confirmed — NAD+**

HPLC METHOD PARAMETERS

METHOD RP-HPLC	COLUMN C18, 4.6 × 250 mm, 5 µm
MOBILE PHASE A 0.1% TFA in water	MOBILE PHASE B 0.1% TFA in acetonitrile
GRADIENT 5–65% B over 30 min	FLOW RATE 1.0 mL/min
DETECTION UV 220 nm	INJECTION / TEMP / RUN 20 µL · 30 °C · 35 min

REPRESENTATIVE CHROMATOGRAM (SCHEMATIC)



PEAK INTEGRATION TABLE

PEAK #	RT (MIN)	AREA	AREA %	ASSIGNMENT
1	8.42	12,480	0.12	Impurity
2	12.18	10,342,110	99.70	BPC-157 (main)
3	14.05	18,650	0.18	Impurity

SPECIFICATION VS RESULT

ATTRIBUTE	SPECIFICATION	RESULT	STATUS
Identity	Matches labeled compound	Confirmed — NAD+	**PASS**
Purity (HPLC)	≥ 99.0%	99.250%	**PASS**
Appearance	White to off-white lyophilized powder	White lyophilized powder	**PASS**
Assayed quantity	Consistent with labeled strength	NAD+: 500 mg	**PASS**

STORAGE & HANDLING

Lyophilized	Lyophilized: -20 °C, protected from light and moisture
After reconstitution	2–8 °C after reconstitution; use within the stability window for research protocols
Sealed expiry	14 MAY 2028

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