Axiom ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL + 1 ADD-ON LAYER (L8)

Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US

AX-2026-4578-2GMP

PRO-Nad+ · Lot T-NAD-0068
Issued Nov 12, 2026 PT
Verify: A612-0E9E-3CD9

SCAN TO VERIFY

PASS — ALL SPECIFICATIONS MET

5 of 5 layers complete

L1	L2	L3	L4	L8
Identity	Purity	Net Content	Sterility	Fentanyl Screen

Sample & Vial Validation



Photograph associated with this sample

MORE VIEWS OF VIAL A



Vial A · right

Brand	**Precise Research Only**	Product	**PRO-Nad+**
Lot / Batch	**T-NAD-0068**	Label Claim	**1000 mg**
Matrix	**lyophilized powder**	Sample Count	**1**
Cap Color	**Black**	Crimp Color	**Silver / Aluminum**
Contents Color	**White**	Seal Intact	**Yes**
Condition	**sealed, intact**	Received	**Sep 24, 2026 PT**
Sampling	**client submitted**		

Test Results

Test	Specification	Result	
Identity research screen	every vial within ±1 Da of 663.43 Da	663.236 Da	PASS
Purity validated in house	Expected ≥ 99.25 · Acceptance ≥ 99.25 · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31	99.72%	PASS
Net Content research screen	every vial within ±10% of 1000.00 mg	Vial A: 1069.22 mg	PASS
Sterility · Rapid PCR research screen	Rapid PCR screen	Target DNA not detected	PASS
Fentanyl Screen research screen	no fentanyl or analog at or above 1 ng/mg	none detected	PASS

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.

Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US

AX-2026-4578-2GMP

PRO-Nad+ · Lot T-NAD-0068
Issued Nov 12, 2026 PT
Verify: A612-0E9E-3CD9

SCAN TO VERIFY

Layer Results

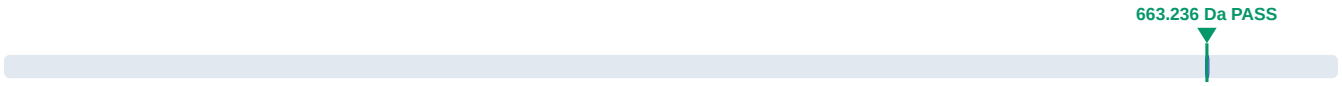
1 Identity

LC-MS (ESI+)

663.236 Da
every vial within ±1 Da of 663.43 Da

PASS

Theoretical mass	**663.43 Da**	Acceptance	**within ±1 Da**
Vial A	**663.236 Da · PASS**		



2 Purity

HILIC-CAD homogeneity (universal detection)

99.72%
Expected ≥ 99.25 · Acceptance ≥ 99.25 · Source Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31

PASS

Acceptance	**≥ 99.25%**	Vial A	**99.72% · PASS**
Criterion source	**Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31**		



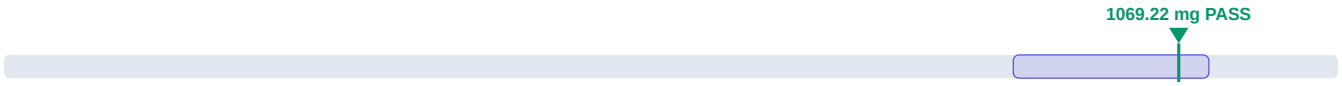
3 Net Content

Quantitative HPLC vs reference std

Vial A: 1069.22 mg
every vial within ±10% of 1000.00 mg

PASS

Target	**1000.00 mg (±10%)**	Vial A	**1069.22 mg · PASS**



4 Sterility · Rapid PCR

Rapid Microbial DNA Screen (qPCR target panel)

Target DNA not detected
Rapid PCR screen

PASS

8 Fentanyl Screen

LC-MS/MS targeted adulterant screen

none detected
no fentanyl or analog at or above 1 ng/mg

PASS

Result	**Not detected**	Detection limit	**1 ng/mg**
Analytes screened	**Fentanyl, Acetylfentanyl, Acrylfentanyl, Butyrylfentanyl, Carfentanil, Cyclopropylfentanyl, Furanylfentanyl, Methoxyacetylfentanyl, ortho-Fluorofentanyl, Valerylfentanyl, 4-ANPP (despropionyl fentanyl)**	Source container	**A**

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.

Axiom
ANALYTICS

CERTIFICATE OF ANALYSIS · 4X ESSENTIAL PANEL + 1 ADD-ON LAYER (L8)

Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US

AX-2026-4578-2GMP

PRO-Nad+ · Lot T-NAD-0068
Issued Nov 12, 2026 PT
Verify: A612-0E9E-3CD9

SCAN TO VERIFY

Reporting Scope & Limitations

`[Sterility · Rapid PCR]` Rapid PCR screen for microbial DNA. It detects DNA from live and dead organisms and is not a USP <71> sterility test or a USP <61> count. Full USP <71> sterility testing is available as an add-on.

`[Fentanyl Screen]` Targeted LC-MS/MS adulterant screen, reported as detected or not detected against its stated detection limit. A harm-reduction research screen, not an exhaustive adulterant panel.

Digital Signatures

Analyst	**Alana Reeves**	Analyst Signature	**ed25519:152ba1d043da79d463f08201...**
Technical Reviewer	**Michael Niles**	Reviewer Signature	**ed25519:0b2610ab08ce74cd0801d967...**
Laboratory Director	**Daniel Brennan**	Director Signature	**ed25519:f7d8b7ad2aad2cec04530639...**

Testing is performed within Axiom's laboratory network under Axiom's methods. Every result is reviewed and released by Axiom, which issues and is responsible for this certificate. Axiom Analytics does not hold ISO/IEC 17025 accreditation. Samples are identified to the testing laboratory by an internal lot ID only. The method status of each result is shown beside it.

Each signature is an Ed25519 signature over the SHA-384 hash of this report. Scan the QR code or verify it yourself at https://axiomanalyticslab.com/verify/AX-2026-4578-2GMP.

✓ CRYPTOGRAPHICALLY SEALED (ed25519 + SHA-384)

`SHA-384: cfa2ad05936609ddc8e7a5698a3c987379ea82728bb610107fb20ac74268c8ea...`

`Seal: ed25519:77419f5c5db3b58b85414606e351c89d87cd0ad2...`

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.

RESEARCH USE ONLY. All analytical testing, validation reports, and interactive Certificates of Analysis (COAs) provided by Axiom Analytics are intended strictly for Research Use Only (RUO) in in-vitro or laboratory animal model environments. Axiom Analytics does not analyze, validate, evaluate, or make any representations regarding the safety, efficacy, toxicity, or legality of any tested substance for human or animal consumption, veterinary use, clinical diagnostics, food additives, cosmetics, or household applications.

Results relate only to the item or portion tested. Unless sampling was performed or witnessed by Axiom Analytics, the laboratory does not attest that the submitted sample represents the wider lot or batch. A PASS designation means only that the reported result met the specification shown. It does not establish safety, efficacy, regulatory approval, legality, or suitability for human or animal use.