



PASS — ALL SPECIFICATIONS MET

5 of 5 layers complete

L1 Identity	L2 Purity	L3 Net Peptide Content	L4 Sterility	L8 Fentanyl Screen
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Sample & Vial Validation



Vial A: as received & analyzed

Brand	APEXX BIOLABS LLC	Product	Adamax 1032
Lot / Batch	T-ADA-0018	Label Claim	10 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		

MORE VIEWS OF VIAL A



Test Results

Test	Specification	Result	
Identity research screen	every vial within ±1 Da of 1032.228 Da	1032.112 Da	PASS
Purity research screen	≥ 99.25 area %	99.92%	PASS
Net Peptide Content research screen	every vial within ±10% of 10 mg	Vial A: 10.29 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



SCAN TO VERIFY

Axiom Analytics Laboratories
2108 N St Ste N, Sacramento, CA 95816 US
AX-2026-684-FC4X
Adamax 1032 · Lot T-ADA-0018
Issued Oct 16, 2026 PT · v1
Verify: C530-9A85-BFD4

Layer Results

1 Identity LC-MS (ESI+)		1032.112 Da every vial within ±1 Da of 1032.228 Da PASS	
Theoretical mass	1032.228 Da	Acceptance	within ±1 Da
Vial A	1032.112 Da · PASS		1032.112 Da PASS

2 Purity RP-HPLC 214nm (USP 621)		99.92% ≥ 99.25 area % PASS	
Acceptance	≥ 99.25%	Vial A	99.92% · PASS
Criterion source	Axiom Analytics SPEC-PURITY-001, Revision 1, effective 2026-08-31		
			99.92% PASS limit 99.25%

3 Net Peptide Content Quantitative HPLC vs reference std		Vial A: 10.29 mg every vial within ±10% of 10 mg PASS	
Target	10 mg (±10%)	Vial A	10.29 mg · PASS
Confirmed container volume	Unknown	Content concentration	Unknown
			10.29 mg PASS

4 Sterility · Rapid PCR Rapid Microbial DNA Screen (qPCR target panel)		Target DNA not detected Rapid PCR screen PASS	
Result	Target DNA not detected	Acceptance	Rapid PCR screen



SCAN TO VERIFY

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

Reporting Scope & Limitations

[Sterility • Rapid PCR] DNA-target research screen only. It detects microbial DNA from viable and non-viable organisms alike, and is neither viable-CFU enumeration under USP <61> nor a batch sterility determination under USP <71>.

[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:b542f9132fd7b702e4f8bfd4...
Technical Reviewer	Michael Niles	Reviewer Signature	ed25519:1e51eedbe0bb27e2676cbcd8...
Laboratory Director	Daniel Brennan	Director Signature	ed25519:69e7739a30a4c479de6a00b7...

Where physical instrument analysis is performed, it takes place within Axiom Analytics' laboratory network, allocated at Axiom's discretion based on method, capacity, and order volume. Axiom Analytics performs sample accession, chain of custody, analytical oversight, and data review, and issues and is responsible for this certificate. Where analysis is subcontracted, the method is performed within the partner laboratory's published ISO/IEC 17025 scope of accreditation for that method, not merely at a laboratory holding ISO/IEC 17025 accreditation. Axiom Analytics does not itself currently hold ISO/IEC 17025 accreditation. The method-validation status of each reported result is identified beside that result. Partner laboratory identities are not disclosed: where samples are analyzed under double-blind conditions, they are identified only by an internal lot ID.

Each signature is an ed25519 signature over the SHA-384 hash of this report. Scan the QR on the letterhead or verify independently at <https://axiomanalyticslab.com/verify/AX-2026-684-FC4X>.

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SHA-384: 9c16a2e0a20ea0b06552fffa67df535703d56d45b92dc64f1f43ad757f624b32...

Seal: ed25519:9e0d20b5f89501276cf0c66fcdfa43aeb4e663b7...

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.

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