

ILS Laboratories

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AOD-9604 - 10mg

20 Tested for: S20 Research, LLC
AMINOS www.20aminos.com

COA #: COA-2026-953618
Lot Number: B10043
Accession #: ACC-2026-5911
Labeled Content: 10mg

Analysis Date: 07/02/2026
Appearance: Good
Sample Matrix: Lyophilized
Vial Size: 3mL
Date Received: 06/20/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: GY5GBCNM

Method: Full QC Panel

Identity
AOD-9604

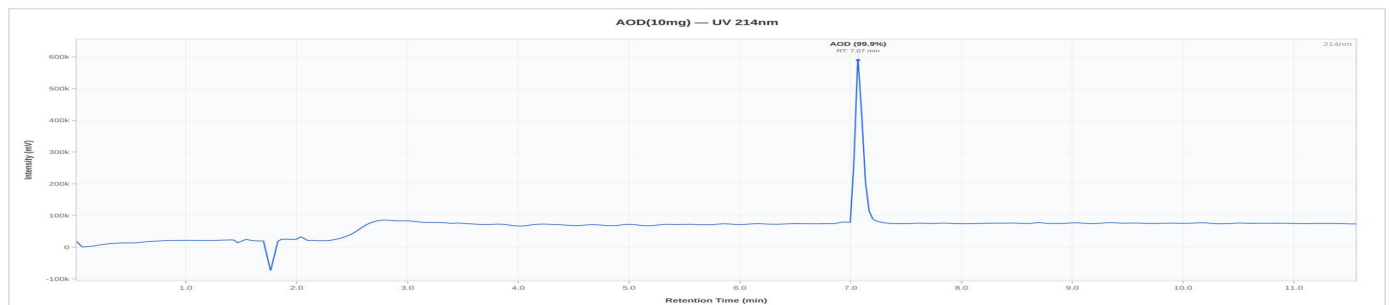
Peptide Purity
99.90%



Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.90%	%	PASS
Net Peptide Content	Report Only	10.73	mg	N/A
Identity (HPLC-RTM)	AOD	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



Representative chromatogram, Conformity V1

Heavy Metals (ICP-MS)

Analyte	Specification	Result	Unit	Status
Heavy Metals				
Arsenic (As)	<= 1.5 ppm	Not Detected	ppm	PASS
Cadmium (Cd)	<= 0.5 ppm	Not Detected	ppm	PASS
Lead (Pb)	<= 1.0 ppm	Not Detected	ppm	PASS
Mercury (Hg)	<= 1.5 ppm	Not Detected	ppm	PASS
Chromium (Cr)	<= 10.0 ppm	0.866	ppm	PASS



Dr. Greg Kalyuzhny

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Lab Director
7/9/2026

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Verify: portal.ils-lab.com/verify/3ekAl6zTdmF66pbE
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HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.89%	10.6 mg	Confirmed	PASS
Conformity V1	99.90%	10.73 mg	Confirmed	PASS
Mean	99.90%	10.66 mg	—	—
Std Dev	0.0050%	0.0650 mg	—	—

Sterility Testing (PCR)

Test	Specification	Result	Status
Sterility (PCR)	No Growth	No Growth	PASS

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	0.084 EU/mL	Reported

About this result: Endotoxin is reported as a quantitative value. Acceptable limits vary by product type and matrix, so no universal pass/fail threshold applies to RUO products. This result is below commonly referenced endotoxin thresholds.

Notes & Methodology

- 1. Date Tested: 07/09/2026. Methods: Purity & Quant (HPLC); Heavy Metals (ICP-MS).
- 2. The sample was confirmed to be AOD-9604 by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- 3. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
- 4. Peptide purity determined by RP-HPLC area normalization at 214 nm. Value represents the percentage of the target peptide relative to all peptide-related peaks. Non-peptide process-related impurities, if detected, are excluded from the calculation.
- 5. Chromatogram shown is representative: Conformity V1 (99.90% purity, closest to batch mean of 99.90%).




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