

Client: Modern Aminos

Certified: 04/17/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	SEMAX	Form	Lyophilized powder
Batch	BX-P-J3R8	Labeled Qty	10 mg
Mol. Formula	C37H51N9O10S	CAS Number	80714-61-0
Cap Color	White	Crimp Color	Sliver

TEST RESULTS

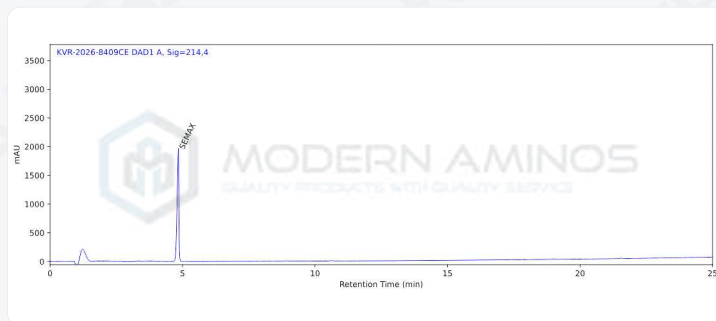
	REFERENCE STANDARD	RESULT
Batch Avg Purity	(>98%)	99.980% <input checked="" type="checkbox"/>
Batch Avg Net Content	(10mg ± 10%)	10.51 mg <input checked="" type="checkbox"/>
Identity Confirmation (LC-MS)	(SEMAX)	SEMAX <input checked="" type="checkbox"/>
Bacterial Endotoxins (LAL)	(≤ 0.50 EU/mL)	PASS <input checked="" type="checkbox"/>

BATCH CONFORMITY RESULTS

Vial	Purity (%)	Net Content (mg)
Vial 1	99.971	10.54
Vial 2	99.982	10.46
Vial 3	99.988	10.53
Batch Average	99.980%	10.51 mg

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm


 Omar Arghandinal
 Lab Director


koveralabs.com/verify

 Report#: KVR-2026-8409CE
 Access Code: JJT2Y1X

CLIENT & SAMPLE INFORMATION

Client	Modern Aminos	Analysis Date	04/17/2026
Product Name	Semax	Strength	10 mg
Lot / Batch	BX-P-J3R8	Condition	Lyophilized

This Certificate of Analysis certifies that the sample listed herein was tested for bacterial endotoxin using a kinetic chromogenic LAL assay in accordance with USP <85> Bacterial Endotoxins Test.

TEST METHODOLOGY

Test Performed	Quantitative Bacterial Endotoxin Test	Compendial Ref	USP <85>
Assay Type	Chromogenic (Kinetic)	Detection λ	405 nm
Detection Range	0.01 – 1.0 EU/mL	Endotoxin Std	E. coli O111:B4
Dilution Vol	2.0 mL	Matrix	LAL Reagent Water

QUANTITATIVE RESULTS

Parameter	Result
Endotoxin Level	< 0.04 EU/mL
Acceptance Limit	≤ 0.5 EU/mL
Sample CV (%)	2.8%
Spike CV (%)	3.5%
Spike Recovery (%)	98%
Final Determination	PASS

CONTROLS

Control	Expected	Observed	Status
Positive Control	Detectable	As expected	Pass
Negative Control	No signal	As expected	Pass

INTERPRETATION

Endotoxin content was quantitatively determined using a kinetic chromogenic LAL assay in accordance with USP <85>. Result reported as below quantitation threshold. The measured endotoxin level meets the specified acceptance criteria. The sample passes the endotoxin limit test.

AUTHORIZATION

REVIEWED BY
Lemar Arghandiwal
Lab Director

