

CERTIFICATE OF ANALYSIS

Product Name:	Nattokinase Vege Delayed Release Capsules				
Product#	DM012	Manufactured for: Dr. Murray Superfood & Natural Products, LLC			
Lot#:	260396	Address: 4823 N, 35 th Place, Phoenix, AZ 85018			
Date Manufactured:	3/2026	Sample Receipt Date: 3/9/2026			
Sample Receipt Condition:	Ambient Temperature		Login Date: 3/9/2026		
Sample Testing Date:	3/10/2026	Sampling: Sample results apply as received			
Product Appearance:	#1 clear/clear oblong acid resistant vege capsules filled with off white to beige powder. Result: Passed				
Weight Variation:	Theoretical Weight: 335 mg Result: 360.72 mg	Specification: 302-369 mg Method: USP <2091>			
Disintegration Time:	Specification: NLT 30 mins.	Result: 40 mins.		Method: USP <2040>	
Reference:	ATDS# 0712-00/26				
DIETARY INGREDIENTS					
Ingredient Name	LC/cap	Results	% of LC	Spec	Method
Nattokinase (20,000 FU/gm)	2,000 FU	2,000 FU	100.00	NLT 100%	**
OTHER INGREDIENTS					
Microcrystalline cellulose, hypromellose (capsule) and monolaurin.					
HEAVY METAL					
Heavy Metal	Specification	Results	Method	QP# 151.03	
Lead:	≤ 0.5 mcg	0.024 mcg	ICP-MS		
Arsenic:	≤ 10 mcg	0.003 mcg	ICP-MS		
Cadmium:	≤ 4.1 mcg	ND	ICP-MS		
Mercury:	≤ 3 mcg	ND	ICP-MS		
MICROBIOLOGY					
Micro Study# MB0059755	Specification	Result	Method		
Total Bacteria Count:	<10,000 CFU/g	<10 CFU/g	USP <2021>		
Total Yeast & Mold Count:	<1,000 CFU/g	<10 CFU/g	USP <2021>		
E. Coli:	Negative/10g	ND/10g	USP <2022>		
Salmonella:	Negative/10g	ND/10g	USP <2022>		
S. Aureus:	Negative/10g	ND/10g	USP <2022>		
Prepared by: <i>Salma Jahan</i>				Date: 3/30/2026	
Reviewed by: <i>Rajivinder Thakur</i>				Date: 3/30/2026	
Approved by: <i>Santosh Kumar</i>				Date: 3/30/2026	
Report Number: 260396.0	Report Status: Final				

** In Accordance with 21 CFR 111.75(d) (1) the following finished dietary ingredients product specifications are exempt from directly finished batch testing requirements as set forth in paragraph 21 CFR 111.75(c) (1). Also confirmed by proper raw material identification, verified by production process controls and QA batch record review and approval to ensure finished product meets all approved MMR in-process specifications. Ref: SOP No. QC-3.

These results apply only to the items tested. This Certificate of Analysis shall not be reproduced, except in its entirety, without the written approval.

-End of Report-