

1. GENERAL PRODUCT INFORMATION

Brand Name: [Company Name]

Product Name: [Enter Product Name]

Lot Number: [Enter Lot Number]

Description: [Product Count/Packaging]

Date of Certification: [MM/DD/YYYY]

Expiration Date: [MM/DD/YYYY]



Purity
Passed



Potency
Passed



Microbial Safety
Passed



Heavy Metals
Passed



Pesticide Residues
Passed



Residual Solvents
Passed

2. INGREDIENT VERIFICATION

Ingredient	Label Claim	Actual	Percent Label Claim Chart
Active Ingredient 1	1400 mg	1610 mg	115%
Label Ingredient 2	750 mg	825 mg	110%
Active Ingredient 3	500 mg	650 mg	130%
Active Ingredient 4	250 mg	300 mg	120%
Active Ingredient 5	100 mg	105 mg	105%
Active Ingredient 6	100 mg	125 mg	125%
Active Ingredient 7	50 mg	57.5 mg	115%
Active Ingredient 8	10 mg	11 mg	110%

0%

100%

3. MICROBIAL ANALYSIS

Test	Specification	Result	Method
Total Aerobic Count	[Specification]	[Result]	USP <61>
Total Yeast & Mold	[Specification]	[Result]	USP <61>
E. coli	[Specification]	[Result]	USP <62>
Salmonella spp.	[Specification]	[Result]	USP <62>
Staphylococcus aureus	[Specification]	[Result]	USP <62>
Pseudomonas aeruginosa	[Specification]	[Result]	USP <62>

4. HEAVY METALS ANALYSIS

Test	Specification	Result	Method
Total Aerobic Count	[Specification]	[Result]	USP
Total Yeast & Mold	[Specification]	[Result]	[Method]



Test	Specification	Result	Method
E. coli	[Specification]	[Result]	[Method]
Salmonella spp.	[Specification]	[Result]	[Method]

5. CHEMICAL ANALYSIS

Test	Specification	Result	Method
Residual Solvents	Under USP Limits	None Detected	USP <467>
Pesticide Residues	Under USP Limits	None Detected	USP <651>
Extraneous?	[Specification]	[Result]	[Method]

* USP refers to the current United States Pharmacopoeia.

CONCLUSION

This product meets label claims and USP safety standards.

Approval Signature: _____

Quality Control Director

Date: 3/1/2026