

Manufactured Exclusively for:
FRTIL

Certificate of Analysis

Product Code: 93762R1	Lot Number: T661A	Date of Manufacture: Jan 2025
Master Batch Record Name: FRTIL Acerola Cherry Vitamin C (90) Cap Bottling MBR		
Product Label Name: Acerola Cherry Vitamin C (90)		
Net Quantity of Contents: 90 Capsules	Serving Size: 3 Capsules	Product stability has not been assessed.

Ingredient	Results per Capsule rounded to the whole unit where ≥ 0.5
Calories	7
Total carbohydrate	1g
Total Sugars	1g
Vitamin C (as For Organic Acerola Cherry (Fruit) Freeze Dried Extract)	125mg
Calcium (as calcium for Organic Acerola Cherry (Fruit) Freeze Dried Extract)	1mg
Potassium (as potassium for Organic Acerola Cherry (Fruit) Freeze Dried Extract)	10mg
Organic Acerola Cherry (Fruit) Freeze Dried Extract (Fruit) (Malpighia glabra L.)	625mg
AVERAGE FINISHED CAPSULE WEIGHT	737 MG

Other Ingredients: capsule (hypromellose, purified water)

This product does not contain any of the 9 major allergens as defined by the Food Allergy Safety, Treatment, Education, and Research (FASTER) Act, signed into law April 23, 2021. The FASTER Act defines the 9 major food allergens as: milk, eggs, fish, Crustacean shellfish, tree nuts, peanuts, wheat, soybeans, and sesame.

Assay	Method	Specification	Results
Identification	NIR (HPLC, IR, or FTIR)	Positive for Acerola Cherry Vitamin C (90)	Positive as indicated
Potency	HPLC	Vitamin C 375mg per serving	Vitamin C 408mg/serving
Appearance	Organoleptic	(0) Vegi Capsule	(0) Vegi Capsule
E. coli	AOAC 101101	USP Absent in 10g	USP Absent in 10g
Total Aerobic Plate Count	AOAC 071203	AHPA $\leq 10^7$ CFU/g	AHPA $\leq 10^7$ CFU/g
Arsenic (As)	ICP-MS FPM 10.01	< 4ppm	As 0.005ppm
Cadmium (Cd)	ICP-MS FPM 10.01	< 2ppm	Cd 0.001ppm
Methylmercury (Hg)	ICP-MS FPM 10.01	< 2ppm	Hg 0.001ppm
Lead (Pb)	ICP-MS FPM 10.01	< 5ppm	Pb 0.041ppm
Enterobacteriaceae	AOAC 121901	AHPA $\leq 10^4$ CFU/g	AHPA $\leq 10^4$ CFU/g
Composition Analysis	Assayed by Weight	Conforms	Conforms
Coliform	USP	≤ 100 CFU/g	≤ 100 CFU/g
Pesticides Analysis	USP <561>	USP <561> Residual Panel as Specified	Pass

This Certificate Of Analysis is based on raw material assays and in house results checked by weight. Above stated ingredients are the inclusive list of all constituents contained in said lot and batch. We acknowledge that each dosage may have up to a 5% difference in stated results due to processing techniques, and said percentage falls within U.S. Guidelines.

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This lot was analyzed and released by our authorized Quality Control Department and was found to meet all intended specifications as given above and in 21 CFR Part 111 and the SOPs that govern those regulations in this facility for purity, potency, strength, and composition. As per First Priority Manufacturing SOPs finished batches are analyzed on a rotational testing basis, every finished product receives a minimum of two tests, as not all tests are completed on every batch some testing information on your COA may be historical data from testing gathered during the production of the product at hand.

Quality Assurance Approval: Tana Madigan 01/27/2025
Tana Madigan Date