Manufactured Exclusively for: FRTIL

Certificate of Analysis

Product Code: 93761	Lot Number: T225A	Date of Manufacture: Mar 2024		
Master Batch Record Name: FRTIL Beef Liver (150) Cap Bottling MBR				
Product Label Name: FRTIL Beef Liver (150) Caps				
Net Quantity of Contents: 150 Capsules	uantity of Contents: 150 Capsules Serving Size: 5 Capsules Product stability has not been as			

 $\begin{tabular}{ll} \hline Ingredient & rounded to the whole unit where ≥ 0.5 \\ \hline Liver (Bovine) (New Zealand) & 600mg \\ \hline AVERAGE FINISHED CAPSULE WEIGHT & 747 MG \\ \hline \end{tabular}$

Other Ingredients: capsule (gelatin, 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.

Consult a physician before use if you have hemochromatosis. Accidental overdose of iron-containing products is a leading cause of fatal poisoning in children under 6. Keep this product out of reach of children. In case of accidental overdose, call a doctor or poison control center immediately.

Assay	Method	Specification	Results
Identification	NIR (HPLC, IR, or FTIR)	Positive for Beef Liver	Positive as indicated
Appearance	Organoleptic	(00) Bovine Capsule	(00) Bovine Capsule
E. coli	AOAC 101101	USP Absent in 10g	USP Absent in 10g
Arsenic (As)	ICP-MS FPM 10.01	< 4ppm	As 0.009ppm
Cadmium (Cd)	ICP-MS FPM 10.01	< 2ppm	Cd 0.144ppm
Methylmercury (Hg)	ICP-MS FPM 10.01	< 2ppm	Hg 0.005ppm
Lead (Pb)	ICP-MS FPM 10.01	< 5ppm	Pb 0.028ppm
Salmonella	AOAC-RI 960801	USP Absent in 10g	USP Absent in 10g
Composition Analysis	Assayed by Weight	Conforms	Conforms

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.

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.

Quality Assurance Approval: Tana Wadigan 05/03/2024
Tana Madigan Date

Results per Capsule