



DN LABORATORY

A Govt. Approves Testing Laboratory Under "the Drugs & Cosmetics Act 1940 And Rules " There Under
Approved By: FDA Haryana Ayush Haryana: An ISO 9001 : 2015 & GLP Certified Lab.
 Address: Ind. Area, Phase-I, Phanchkula, Haryana.
 Mob: +91 9538239428, email id: dnlabpk@gmail.com, manisha256@gmail.com

Certificate of Analysis

Form-47, 160 (A) Report of test or Analysis by Approved Institution

Sample Name	Aumeto CholestShield Capsules			Received Date	22-07-2025
Customer Information	AUMETO, 201T ACADEMY ST, GREENVILLE, SC 29601			Ref. No.	NIL
Supplied By	NS			Report No.	DNL/534/25-26/317G/R
Batch Size	NS	LOT No.	AT-XXVT7N14	Sample Qty.	90 Capsules
Mfg. Date	07-2025	Exp. Date	22-07-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	23-07-2025	Date of Completion	29-07-2025	Protocol ID	IHS

RESULT OF ANALYSIS

Description	Grey powder filled in 0 sized transparent hard gelatin capsule		
Average Fill weight	: 500 mg		
Disintegration Time	: 10-11 min (NMT – 30 min)		
Assay	: Each Serving Two Capsule (on an average fill) contains: -		
Composition	Claim	Observed	Method
Citrus Bergamot	500 mg	500.15 mg	HPLC
Black Chokeberry Fruit	200 mg	199.92 mg	HPLC
Olive Leaf	50 mg	49.87 mg	HPLC
Niacin (As Nicotinic Acid)	50 mg	50.03 mg	HPLC
Hawthorn Berry, Turmeric, Ginger, Fenugreek, Artichoke Leaf, Holy basil, Garlic, Cinnamon, Apple, Black Pepper	200 mg	200.11 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characterstics	Grey Powder	Complies	Organoleptic
Odor & Taste	Characterstics	Conform	Organoleptic
Solubility	Soluble in Alcohol	Complies	Turbidity Meter
Assay	99%	99.8%	HPLC
Specific Rotation	(-4) – (+3)	+1.7	Polarimeter
Refractive Index	1.458-1.464 at 20°C	1.461	Refractometer
Specific Gravity	0.877-0.890 at 20°C	0.884	Densimeter
Optical Rotation	+5- +34°C	+31	Polarimeter
Residue on Ignition	NMT 1%	0.45%	Muffle Furnace
Particle Size	90 % pass 80 mesh	Conforms	CP 2015
Loss on drying	NMT 5.0%	3.47%	USP
Ash Content	NMT 5.0%	3.19 %	USP
Residue of Solvents	NMT 0.5%	Conforms	NLS-QCS-1007
Moisture	MNT 5 %	3.67 %	AOAC
pH 5 % in water	5-7 at 25°C	5.30	USP <791>
Uniformity of Weight	5.0-7.0%	5.60 %	Ph. Eur. Method 2.9.5
Bulk Density	0.35-0.50 g/mL	0.40 g/mL	Bulk Density Apparatus
Tapped Density	0.40-0.80 g/mL	0.50 g/mL	Tapped Density Apparatus
Heavy Metals:			
Lead	NMT - 2 ppm	0.68 ppm	ISO 17294 ICP-MS
Arsenic	NMT - 1 ppm	0.43 ppm	ISO 17294 ICP-MS
Cadmium	NMT - 1 ppm	0.29 ppm	ISO 17294 ICP-MS
Mercury	NMT - 0.1 ppm	0.05 ppm	EN 13806 AAS
Microbial Examination:			
Yeast & Mould	NMT 300 cfu/g Max	10 cfu/g	GB 4789.15
E. Coli	Absent/g	Absent	GB 4789.3
Salmonella	Absent/g	Absent	GB 4789.3.2016
Staphylococcus Aureus	Absent/g	Absent	GB 4789.3.2016
Pseudomonas Aeruginosa	Absent/g	Absent	GB 4789.3.2016

29-07-2025

Date of Completion

Signature of Person-in- charge of testing



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OPINION : In the opinion of the undersigned the above sample is of standard quality is of standard quality as defined in the Act and Rules made there under for the reasons given below

The sample confirms to IP ☐ BP ☐ USP ☐ BIS ☐ ISO ☐ AYUR ☐ test specification with respect to above test only.

29-07-2025

Date of Completion

Signature of Person-in- charge of testing

Note :

1. This report is not to be reproduced wholly or in part and cannot be used as evidence in the court of law and should not be used in any advertising media without our special permission in writing.
2. Samples (s) not drawn by us, unless otherwise stated.
3. Total liability of our analytical division is limited to the invoiced amount.
4. Sample will be destroyed after one month from date of issue of test certificate unless otherwise specified.
Results given in reports are released to sample tested.
- 5.