



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 Apple Cider Vinegar Capsules			Received Date	21-07-2025
Customer Information	AUMETO, 201T ACADEMY ST, GREENVILLE, SC 29601			Ref. No.	NIL
Supplied By	NS			Report No.	DNL/534/25-26/323T/H
Batch Size	NS	LOT No.	AT-XXVT7N16	Sample Qty.	150 Capsules
Mfg. Date	07-2025	Exp. Date	21-07-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	22-07-2025	Date of Completion	28-07-2025	Protocol ID	IHS

RESULT OF ANALYSIS

Description	Brown powder filled in 0 sized transparent hard gelatin capsule		
Average Fill weight	: 505 mg		
Disintegration Time	: 10-11 min	(NMT – 30 min)	
Assay	: Each Serving Two Capsule (on an average fill) contains: -		
Composition	Claim	Observed	Method
Vitamin D3 (as Cholecalciferol)	125 mcg	124.98 mcg	HPLC
Apple Cider Vinegar	300 mg	300.07 mcg	HPLC
Berberine HCL (Berberis Aristala, Root)	200 mg	199.85 mg	HPLC
Turmeric (Curcuma longa, root)	100 mg	99.94 mg	HPLC
Ceylon Cinnamon (Cinnamomum verum, bark)	200 mg	200.07 mg	HPLC
Gymnema Sylvestre (leaf), Bitter Melon (Momordica charantia, fruit), Fenugreek (seed), Ginger (Zingiber officinale, rhizome), Holy basil (Ocimum sanctum, stem)	200 mg	200.16 mg	HPLC
Black Pepper (Piper nigrum) (fruit)	10 mg	10.05 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characteristics	Brown Powder	Complies	Organoleptic
Odor & Taste	Characteristics	Conform	Organoleptic
Solubility	Soluble in PEG	Complies	Turbidity Meter
Total Acidity (Acetic Acid)	NLT 10%	10.23%	Titration
Refractive Index	0.90-2.0 at 20°C	1.05	Refractometer
Specific Gravity	1.001-1.083 at 20°C	1.039	Densimeter
Particle Size	95% pass 80 mesh	Conforms	CP 2015
Loss on drying	NMT 5.0%	3.25%	USP
Ash Content	NMT 3.5%	2.66 %	USP
Moisture	MNT 5 %	3.54 %	AOAC
pH 5 % in water	3-7 at 25°C	5.61	USP <791>
Bulk Density	50-60g/100mL	55g/100mL	Bulk Density Apparatus
Tapped Density	90-130 g/100mL	108g/100mL	Tapped Density Apparatus
Heavy Metals:			
Lead	NMT - 2 ppm	0.73 ppm	ISO 17294 ICP-MS
Arsenic	NMT - 2 ppm	0.57 ppm	ISO 17294 ICP-MS
Cadmium	NMT - 1 ppm	0.24 ppm	ISO 17294 ICP-MS
Mercury	NMT - 0.1 ppm	0.03 ppm	EN 13806 AAS
Microbial Examination:			
Yeast & Mould	NMT 100 cfu/g Max	20 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

28-07-2025

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 ☒ HP ☐ BP ☐ USP ☐ BIS ☐ ISO ☐ AYUR ☐ test specification with respect to above test only.

28-07-2025

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.
5. Results given in reports are released to sample tested.