



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 Red Vine Leaf Capsules			Received Date	16-07-2025
Customer Information	AUMETO, 201T ACADEMY ST, GREENVILLE, SC 29601			Ref. No.	NIL
Supplied By	NS			Report No.	DNL/307/25-26/218T/R
Batch Size	NS	LOT No.	AT-XXVT7N11	Sample Qty.	90 Capsules
Mfg. Date	07-2025	Exp. Date	16-07-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	17-07-2025	Date of Completion	23-07-2025	Protocol ID	IHS

RESULT OF ANALYSIS

Description	Brown 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
Red Vine Leaf (Vitis Vinifera)	500 mg	500.12 mg	HPLC
Horse Chestnut (Aesculus Hippocastanum) (Seed)	200 mg	199.95 mg	HPLC
Beet Root (Beta Vulgaris)	100 mg	99.98 mg	HPLC
Ginger Root (Zingiber Officinale), Gotu Kota (Centella Asiatica) Leaf, Butcher's Broom Root (Ruscus Aculeatus), Bilberry (Vaccinium Myrtillus) Fruit, Hesperidin (from Citrus Sinensis), Fruit Peel, Grape Seed (Vitis Vinifera), L-Arginine, Diosmin (from Citrus Aurantium) Fruit Peel, Hawthorn Berry (Crataegus Spp.)	200 mg	199.93 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characteristics	Brown Powder	Complies	Organoleptic
Odor & Taste	Characteristics	Conform	Organoleptic
Solubility	Soluble in Water	Complies	Turbidity Meter
Polyphenols Flavonoids	4-7%	6.5%	UV
Residue on Ignition	NMT 30%	6.58%	Muffle Furnace
Particle Size	95 % pass 80 mesh	Conforms	USP39<786>
Loss on drying	NMT 5%	3.68%	Eur.Ph.7.0[2.8.17]
Ash Content	NMT 7%	6.15 %	Eur.Ph.7.0[2.4.16]
Moisture	MNT 6 %	4.85 %	AOAC
pH 5 % in water	4-8 at 25°C	5.69	USP <791>
Uniformity of Weight	5.0-7.50 %	6.3 %	Ph. Eur. Method 2.9.5
Heavy Metals:			
Lead	NMT - 1 ppm	0.2 ppm	ISO 17294-2 2016
Arsenic	NMT - 0.5 ppm	0.3 ppm	ISO 17294-2 2016
Cadmium	NMT - 0.5 ppm	0.10 ppm	ISO 17294-2 2016
Mercury	NMT - 0.5 ppm	0.03 ppm	ISO 17294-2 2016
Microbial Examination:			
Yeast & Mould	NMT 100 cfu/g Max	15 cfu/g	USP35<965>
E. Coli	Absent/g	Absent	USP35<965>
Salmonella	Absent/g	Absent	USP35<965>
Staphylococcus Aureus	Absent/g	Absent	USP35<965>
Pseudomonas Aeruginosa	Absent/g	Absent	USP35<965>

23-07-2025

Date of Completion

Signature of Person-in- charge of testing



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RESULT OF ANALYSIS

Specific Compound:	Observed	Specification	Method
Sildenafil	Not Detected	Below MDL	USP <2251>
Vardenafil	Not Detected	Below MDL	USP <2251>
Tadalafil	Not Detected	Below MDL	USP <2251>
Phenolphthalein	Not Detected	Below MDL	USP <2251>
Fluoxetine	Not Detected	Below MDL	USP <2251>
4-DHEA (4-androstene-3	Not Detected	Below MDL	USP <2251>
Desmethylcarbodenafil	Not Detected	Below MDL	USP <2251>
Sibutramine	Not Detected	Below MDL	USP <2251>
Desmethyisibutramine	Not Detected	Below MDL	USP <2251>
Didesmethyisibutramine	Not Detected	Below MDL	USP <2251>
Ostarine	Not Detected	Below MDL	USP <2251>
Ugandrol	Not Detected	Below MDL	USP <2251>

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.

23-07-2025

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.