GSTIN : 06AJZPM1523E1Z5

DNL

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@gmai.com, manisha256@gmail.com

Certificate of Analysis

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

Sample Name	Aumeto Tribulus Capsules			**Received Date**	20-07-2025
Customer Information	AUMETO, 201T ACADEMY ST, GREENVILLE, SC 29601			**Ref. No.**	NIL
Supplied By	NS			**Report No.**	DNL/355/25-26/215L/D
Batch Size	NS	**LOT No.**	AT-XXVT7N12	**Sample Qty.**	150 Capsules
Mfg. Date	07-2025	**Exp. Date**	20-07-2028	**Fssai. Lic. No.**	10822999000390
Date of Analysis	21-07-2025	**Date of Completion**	27-07-2025	**Protocol ID**	IHS

RESULT OF ANALYSIS

Description : Brown powder filled in 0 sized transparent hard gelatin capsule
Average Fill weight : 550 mg
Disintegration Time : 10-11 min (NMT – 30 min)
Assay : Each Serving Two Capsule (on an average fill) contains: -

Composition	Claim	Observed	Method
Tribulus Terrestris	600 mg	600.10 mg	HPLC
Ashwagandha Root	100 mg	100.03 mg	HPLC
Korean Red Panax Ginseng	100 mg	99.87 mg	HPLC
Coenzyme Q10, Cordyceps, Yohimbe Bark, Maca Root, Black Pepper	300 mg	299.90 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characterstics	Brown Powder	Complies	Organoleptic
Odor & Taste	Characterstics	Conform	Organoleptic
Solubility	Soluble in Water	Complies	Turbidity Meter
Assay			
Saponins	NTL 95%	96.78%	UV-VIS
Protodioscin	NTL 60%	61.82%	UV-VIS
Withanolides	NTL 8%	8.65%	HPLC
Specific Rotation	+23 - +33	+27	Polarimeter
Residue on Ignition	NMT 1%	0.3%	Muffle Furnace
Particle Size	98 % 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	None	Conforms	NLS-QCS-1007
Moisture	MNT 10 %	6.83 %	AOAC
pH 5 % in water	4-7 at 25°C	5.52	USP <791>
Uniformity of Weight	5.0-7.5%	6.5 %	Ph. Eur. Method 2.9.5
Bulk Density	0.45-0.55 g/mL	0.50 g/mL	Bulk Density Apparatus
Tapped Density	0.40-1.0 g/mL	0.67 g/mL	Tapped Density Apparatus
Heavy Metals:			
Lead	NMT - 2 ppm	0.85 ppm	ICP-MS
Arsenic	NMT - 2 ppm	0.63 ppm	ICP-MS
Cadmium	NMT - 1 ppm	0.39 ppm	ICP-MS
Mercury	NMT - 0.5 ppm	0.23 ppm	ICP-MS
Microbial Examination:			
Yeast & Mould	NMT 300 cfu/g Max	10 cfu/g	USP 2021.2022
E. Coli	Absent/g	Absent	USP 2021.2022
Salmonella	Absent/g	Absent	USP 2021.2022
Staphylococcus Aureus	Absent/g	Absent	USP 2021.2022
Pseudomonas Aeruginosa	Absent/g	Absent	USP 2021.2022

27-07-2025
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.

27-07-2025
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.