



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 Turkesterone Capsules			Received Date	10-08-2025
Customer Information	AUMETO, 201T ACADEMY ST, GREENVILLE, SC 29601			Ref. No.	NIL
Supplied By	NS			Report No.	DNL/247/25-26/487R/G
Batch Size	NS	LOT No.	AT-XXVT8N15	Sample Qty.	90 Capsules
Mfg. Date	08-2025	Exp. Date	10-08-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	11-08-2025	Date of Completion	17-08-2025	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		
Assay	: Each Serving Capsule (on an average fill) contains: -		
Composition	Claim	Observed	Method
Ajuga Turkestanica (whole herb)	350 mg	350.07 mg	HPLC
Ecdysterone 98%	50 mg	49.96 mg	HPLC
L-Arginine	50 mg	50.02 mg	HPLC
Vitamin D3 (as Cholecalciferol)	50 mg	49.98 mg	HPLC
Tongkat Ali, Fadogia Agrestis, Ashwagandha, Fenugreek, Maca Root, Panax Ginseng, Mucuna Pruriens, Ginkgo Biloba, Black Pepper	50 mg	49.85 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characteristics	Brown Powder	Complies	Organoleptic
Odor & Taste	Characteristics	Conform	Organoleptic
Solubility	Soluble in DMSO	Complies	Turbidity Meter
Assay Turkesterone	NLT 2 %	2.23%	HPLC
Irradiation Status	Non-Irradiated	Irradiation Free	ESR Spectroscopy
Particle Size	95% pass 80 mesh	Conforms	USP<786>
Loss on drying	NMT 5%	2.85%	USP<731>
Ash Content	NMT 5 %	2.34 %	USP<281>
Solvent Residues	NMT 0.07%	Conforms	USP-GC
pH 5 % in water	5.5-7.5 at 25°C	5.8	USP <791>
Bulk Density	45-55g/100ml	49.2g/100ml	Bulk Density Apparatus
Moisture	MNT 8 %	4.02 %	AOAC
Heavy Metals:			
Lead	NMT - 2 ppm	0.05 ppm	USP <231> ICP-MS
Arsenic	NMT - 2 ppm	0.02 ppm	USP <231> ICP-MS
Cadmium	NMT - 2 ppm	0.01 ppm	USP <231> ICP-MS
Mercury	NMT - 0.5 ppm	0.02 ppm	USP <231> ICP-MS
Microbial Examination:			
Total Plate Count	NMT 1000 cfu/g	Complies	USP <61>
Yeast & Mould	NMT 100 cfu/g	15 cfu/g	USP <61>
E. Coli	Absent/g	Absent	USP <61>
Salmonella	Absent/g	Absent	USP <61>
Staphylococcus Aureus	Absent/g	Absent	USP <61>
Pseudomonas Aeruginosa	Absent/g	Absent	USP <61>

17-08-2025

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.

17-08-2025

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.