GSTIN : 06AJZPM1523E1Z5

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-1, 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	Ultra Herbs L-Citrulline, L-Arginine Taurine Capsules			**Received Date**	04-09-2025
Customer Information	Ultra Herbs, 2711 W Whiteside, St # Springfield MO 65807			**Ref. No.**	NIL
Supplied By	NS			**Report No.**	DNL/09/25-26/78Q/R
Batch Size	NS	**LOT No.**	UH-XXVT9N08	**Sample Qty.**	180 Capsules
Mfg. Date	09-2025	**Exp. Date**	04-09-2028	**Fssai. Lic. No.**	10822999000390
Date of Analysis	05-09-2025	**Date of Completion**	11-09-2025	**Protocol ID**	IHS

RESULT OF ANALYSIS

Description : Brown powder filled in transparent hard gelatin capsule
Average Fill weight : 1250 mg
Disintegration Time : 10-11 min (NMT – 30 min)
Assay : Each Serving 2 Capsule (on an average fill) contains: -

Composition	Claim	Observed	Method
Vitamin D3 (as Cholecalciferol)	50 mcg	49.80 mcg	HPLC
Vitamin C	100 mg	99.95 mg	HPLC
L-Citrulline	1500 mg	1499.90 mg	HPLC
L-Arginine	200 mg	199.85 mg	HPLC
Taurine	200 mg	200.15 mg	HPLC
Creatine Monohydrate	200 mg	199.88 mg	HPLC
Ashwagandha Root (Withania Somnifera)	50 mg	49.98 mg	HPLC
Tongkat Ali Root	200 mg	200.10 mg	HPLC
Fadogia Agrestis Stem	50 mg	49.85 mg	HPLC
Black Pepper (Piper Nigrum) fruit	20 mg	20.15 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characterstics	Brown Powder	Complies	Organoleptic
Odor & Taste	Characterstics	Conform	Organoleptic
Solubility	Soluble in Ethanol	Complies	Turbidity Meter
L-Citrulline	NLT 98.5%	99.30%	HPLC
Refractive Index	1.425-1.510 at 20°C	1.479	Refractometer
Specific Optical Rotation	+17.5° ± 1°	+17.1°	Spectrophotometer
Specific Gravity	1.280-1.300 at 20°C	1.289	Densimeter
Particle Size	95% pass 80 mesh	Conforms	CP 2015
Loss on drying	NMT 0.3%	0.15%	USP
Residue on Ignition	NMT 0.1%	0.09%	USP<281>
Ash Content	NMT 0.5%	0.07 %	USP
Moisture	MNT 0.5 %	0.09 %	AOAC
pH 5 % in water	2.5-3 at 25°C	2.7	USP <791>
Bulk Density	40-60g/100mL	46g/100mL	USP <616>
Tapped Density	70-90 g/100mL	76g/100mL	USP <616>
Heavy Metals:			
Total Heavy Metals	10 ppm Max	Conforms	USP<233> ICP-MS
Lead	NMT - 1 ppm	0.34 ppm	USP <233> ICP-MS
Arsenic	NMT - 2 ppm	0.26 ppm	USP <233> ICP-MS
Cadmium	NMT – 0.5 ppm	0.15 ppm	USP <233> ICP-MS
Mercury	NMT - 0.1 ppm	0.05 ppm	USP <233> ICP-MS
Microbial Examination:			
Total Plate Count	NMT 1000 cfu/g	Complies	USP<2021>
Yeast & Mould	NMT 100 cfu/g	20 cfu/g	USP<2021>
E. Coli	Absent/g	Absent	USP<2021>
Salmonella	Absent/g	Absent	USP<2021>
Staphylococcus Aureus	Absent/g	Absent	USP<2021>
Pseudomonas Aeruginosa	Absent/g	Absent	USP<2021>

11-09-2025
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.

11-09-2025
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.