GSTIN: 06AJZPM1523E1Z5



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 C	lutathione, Res	veratrol NAC Capsules	Received Date	01-09-2025
Customer Information	Ultra Herbs, 2711 W Whiteside, St. # Springfield MO 65807			Ref. No. Report No.	NIL DNL/09/25-26/18B/C
Supplied By					
Batch Size	NS	LOT No.	UH-XXVT9N04	Sample Qty.	150 Capsules
Mfg. Date	09-2025	Exp. Date	01-09-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	02-09-2025	Date of Completion	08-09-2025	Protocol ID	IHS

		RESULT OF ANALYSIS			
Description	Brown powder filled	in transparent hard gelatin ca	psule		
Average Fill weight	: 530 mg				
Disintegration Time : 10-11 min		(NMT - 30 min)			
Assay	: Each Serving Capsu	le (on an average fill) contain	IS: -		
Composition		Claim	Observed	Method	
Vitamin C		50 mg	50.10 mg	HPLC	
Glutathione		400 mg	400.15 mg	HPLC	
Trans-Resveratrol		20 mg	19.85 mcg	HPLC	
NAC (N-Acetyl-L-Cysteine)		10 mg	9.80 mg	HPLC	
Organic-Collagen (Collagen	Type I. III)	20 mg	19.90 mg	HPLC	
(As Hydrolyzed Fish Collage					
Hyaluronic Acid (as Sodium)		20 mg	19.95 mg	HPLC	
Black Pepper (Piper Nigrum)		10 mg	10.15 mg	HPLC	
Identification	11.00.0	Conform	Complies	IR Spectrum	
Characterstics		Brown Powder	Complies	Organoleptic	
Odor & Taste		Characterstics	Conform	Organoleptic	
Solubility		Soluble in Water	Complies	Turbidity Meter	
Glutathione		NLT 98%	99.47%	HPLC	
Refractive Index	M.	1.492-1.600 at 20°C	1.571	Refractometer	
Specific Gravity		1.35-1.50 at 20°C	1.44	Densimeter	
Optical Rotation		-15.5 to -17.5 °	-16.8°	Polarimeter	
Particle Size		98% pass 80 mesh	Conforms	CP 2015	
Loss on drying		NMT 0.5%	0.15%	USP	
Residue on Ignition		NMT 0.1%	0.02%	USP<281>	
Residue of Solvents •		NMT 5000 ppm	Conforms	GC-USP<467>	
Ash Content		NMT 0.1%	0.05 %	USP	
Moisture		MNT 5 %	2.65 %	AOAC	
pH 5 % in water		4-5 at 25°C	4.30	USP < 791>	
Bulk Density		30-50g/100mL	41g/100mL	USP <616>	
Tapped Density	46	40-60 g/100mL	54g/100mL	USP <616>	
Heavy Metals:		10 00 g 100m2	0.8		
Total Heavy Metals		10 ppm Max	Conforms	USP<233> ICP-MS	
Lead		NMT - 3 ppm	0.64 ppm	USP <233> ICP-MS	
Arsenic		NMT - 1 ppm	0.55 ppm	USP <233> ICP-MS	
Cadmium		NMT - 1 ppm	0.27 ppm	USP <233> ICP-MS	
		NMT - 0.1 ppm	0.02 ppm	USP <233> ICP-MS	
Mercury Microbial Examination:		Tivii - 0.1 ppm	o.oz ppin		
Total Plate Count		NMT 1000 cfu/g	Complies	USP<2021>	
Yeast & Mould		NMT 1000 cfu/g	22 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>	
08-09-2025		1 LOSCHU B	A		
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Date of Completion		Sig	gnature or Fe	ison-in- charge of tes	ung

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DN LABORATORY

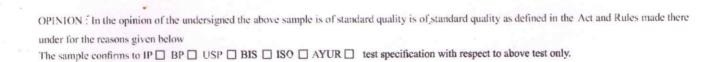
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08-09-2025 Date of Completion

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.

 Samples (s) not drawn by us, unless otherwise stated.
- Total liability of our analytical division is limited to the invoiced amount.
- 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.