



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	Ultra Herbs NAD+ Boosting Nicotinamide Riboside Capsules			Received Date	05-09-2025
Customer Information	Ultra Herbs, 2711 W Whiteside, St # Springfield MO 65807			Ref. No.	NIL
Supplied By	NS			Report No.	DNL/09/25-26/148D/E
Batch Size	NS	LOT No.	UH-XXVT9N15	Sample Qty.	90 Capsules
Mfg. Date	09-2025	Exp. Date	05-09-2028	Fssai. Lic. No.	10822999000390
Date of Analysis	06-09-2025	Date of Completion	12-09-2025	Protocol ID	IHS

RESULT OF ANALYSIS

Description	Brown fine powder filled in transparent hard gelatine capsule			
Average Fill weight	: 500 mg			
Disintegration Time	: 18-20 min (NMT – 30 min)			
Assay	: Each serving capsule (on an average fill) contains: -			
Composition	Claim	Observed	Method	
Nicotinamide Riboside	300 mg	300.18 mg	HPLC	
Trans-Resveratrol	100 mg	99.95 mg	HPLC	
(From Polygonum Cuspidatum Root)				
Turmeric (Curcuma Longa) Rhizome	30 mg	30.10 mg	HPLC	
Quercetin (Sophora japonica, flower)	20 mg	19.80 mg	HPLC	
Rhodiola Rosea	20 mg	20.12 mg	HPLC	
Milk Thistle (Silybum marianum) Seed	20 mg	19.90 mg	HPLC	
Black Pepper (Piper nigrum fruit)	10 mg	9.88 mg	HPLC	
Odor & Taste	Characteristic	Conforms	Organoleptic	
Identification	Conform	Complies	IR Spectrum	
Assay	NLT 98%	99.7%	U.V.	
Loss on drying	NMT 0.5%	0.3%	Muffle Furnace	
Moisture	NMT 5.0%	2.5%	AOAC	
Particle Size	NLT 95 % Should pass through 80 mesh	Conforms	Eur. Ph.<2.9.12>	
Bulk Density	Between 20-40 g/100ml	34g/100ml	Eur. Ph.<2.9.34>	
Tapped Density	Between 40-60 g/100ml	57g/100ml	Eur. Ph.<2.9.34>	
pH of 1 % Suspension	2-4 at 25°C	3.2	USP <791>	
Ash Content	NMT 5 % w/w	2.7 %	Muffle Furnace	
Heavy Metals:				
Total Heavy Metals	10 ppm Max	Conforms	USP<231> ICP-MS	
Lead	NMT - 2 ppm	0.39 ppm	USP<231> ICP-MS	
Arsenic	NMT - 1 ppm	0.27 ppm	USP<231> ICP-MS	
Cadmium	NMT - 0.5 ppm	0.03 ppm	USP<231> ICP-MS	
Mercury	NMT - 0.1 ppm	0.01 ppm	USP<231> ICP-MS	
Microbial Examination:				
Total Aerobic Bacteria	NMT 1000 cfu/gm	150 cfu/g	USP<61>	
Yeast & Mould	NMT 100 cfu/g Max	20 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>	

12-09-2025

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.

12-09-2025

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.