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, Panchkula, 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	Liposomal NMN Trans-Resveratrol Capsules			**Received Date**	02-01-2026
Customer Information	Reverse Clinics by Threads Physio Orthosport LLP, F-10/4, DLF Phase 1, Sector-27, Gurugram, Haryana 122001			**Ref. No.**	NIL
Supplied By	NS			**Report No.**	DNL/01/26-27/18B/C
Batch Size	NS	**LOT No.**	CW-091	**Sample Qty.**	60 Capsules
Mfg. Date	01-2026	**Exp. Date**	02-01-2028	**Fssai. Lic. No.**	20825005004121
Date of Analysis	03-01-2026	**Date of Completion**	09-01-2026	**Protocol ID**	IHS

RESULT OF ANALYSIS

Description : Yellow Powder filled in transparent hard gelatin capsule
Average Fill weight : 590 mg
Disintegration Time : 10-11 min (NMT – 30 min)
Assay : Each Serving Two Capsule (on an average fill) contains: -

Composition	Claim	Observed	Method
Liposomal NMN	500 mg	500.10 mg	HPLC
Trans-Resveratrol	150 mg	149.95 mg	HPLC
Quercetin Phytosome	150 mg	150.15 mg	HPLC
Coenzyme Q10	100 mg	99.85 mcg	HPLC
Ashwagandha KSM-66	100 mg	100.20 mg	HPLC
Green Tea Extract (EGCG)	60 mg	59.90 mg	HPLC
Grape Seed Extract	60 mg	60.10 mcg	HPLC
Panax Ginseng	50 mg	49.90 mg	HPLC
Black Pepper	5 mg	4.95 mg	HPLC
Identification	Conform	Complies	IR Spectrum
Characterstics	Yellow Powder	Complies	Organoleptic
Odor & Taste	Characterstics	Conform	Organoleptic
Solubility	Soluble in Water	Complies	Turbidity Meter
Assay	NLT 98%	99.5%	HPLC
Refractive Index @ 25 C	1.44-1.48 at 20°C	1.45	Refractometer
Specific Gravity	1.735-1.825 at 20°C	1.810	Densimeter
Particle Size	95% pass 80 mesh	Conforms	CP 2015
Loss on drying	NMT 1.0%	0.34%	USP
Ash Content	NMT 1.0%	0.58 %	USP
Moisture	MNT 0.5 %	0.27 %	AOAC
pH 5 % in water	3-4 at 25°C	3.25	USP <791>
Acid Insoluble Ash	NMT 0.3%	0.07%	Muffle Furnace
Residual Solvent	NMT 50 ppm	Conforms	Eur. Ph.<2.4.24>
Bulk Density	30-50g/100mL	43g/100mL	Bulk Density Apparatus
Tapped Density	60-80 g/100mL	71g/100mL	Tapped Density Apparatus
Heavy Metals:			
Total Heavy Metals	10 ppm Max	Conforms	USP<231> ICP-MS
Lead	NMT - 1 ppm	0.35 ppm	USP <231> ICP-MS
Arsenic	NMT – 0.5 ppm	0.27 ppm	USP <231> ICP-MS
Cadmium	NMT – 0.2 ppm	0.11 ppm	USP <231> ICP-MS
Mercury	NMT - 0.1 ppm	0.03 ppm	USP <231> ICP-MS
Microbial Examination:			
Total Plate Count	NMT 2000 cfu/g	Complies	USP<61>
Yeast & Mould	NMT 50 cfu/g	10 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>

09-01-2026
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.

09-01-2026
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.