

**ESKAG PHARMA PVT. LTD.**

Manufacturing Address: Plot No. 31, 32, 33 & 34, Sector 6A, I.I.E. Sidcul, Ranipur,
Haridwar, Uttarakhand 249403, India
Corporate Address: AG-112, Sector-II, Suite No.804 & 805, Baishakhi, Salt Lake City, Kolkata,
West Bengal 700 091, India
Tel: +91 33 40251500. Email: eskag@eskag.in

CERTIFICATE OF ANALYSIS**Certificate of Analysis**

COMPLIES / PASSES

SAMPLE INFORMATION

Name of Product:	Liposomal Curcumin Capsule (100 mg)	Date of Sampling:	21/04/2026
Generic Name:	Liposomal Curcumin Capsule	Date of Release:	28/04/2026
Batch No.:	CCC32601	STP No.:	STP/FP/IIIF/004-00
Batch Size:	15,000 NOS	Specification No.:	SP/FP/IIIF/004-00
Manufacturing Date:	APR. / 2026	Sample Quantity:	100 Capsules
Expiry Date:	SEP. / 2027	A.R. No.:	FP/IIIF/260002

RESULTS

Parameter	Specification	Result	Status
Physical & Identification			
Description	White colour cap & White colour body hard gelatin capsule containing yellow colour granular powder.	White colour cap & White colour body hard gelatin capsule containing yellow colour granular powder.	PASS
Identification (HPLC)	The retention time of the major peak for liposomal curcumin obtained from test preparation should correspond to that of standard preparation as described under the assay.	Complies	PASS
Physical Parameters			
Disintegration Time	NMT 30 minutes	8 min 55 sec	PASS
Average Net Content per Capsule	255.0 mg $\pm$ 10.0% w/w	250.14 mg	PASS
Uniformity of Net Weight	$\pm$ 10.00% w/w	+4.14%, -6.05% w/w	PASS
Assay			
Each Hard gelatin capsule contains Liposomal Curcumin 100 mg	90.0 mg to 110.0 mg	105.9 mg	PASS
Microbial Enumeration Tests			
Total Bacterial Count	NMT 1000 CFU/gm	40 CFU/gm	PASS
Total Fungal Count	NMT 100 CFU/gm	Nil	PASS
Tests for Specified Micro-organisms			
<i>E. coli</i>	Should be Absent / gm	Absent / gm	PASS

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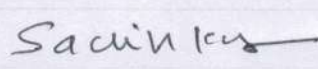
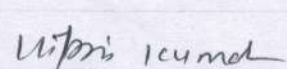
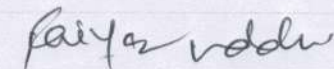
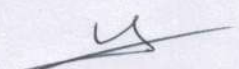
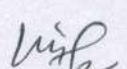
Overall Result: The sample COMPLIES as per above Specification.

CONCLUSION

The Liposomal Curcumin Capsule (Batch No. CCC32601) was analyzed in accordance with the standard test procedures stipulated in STP/FP/IIIF/004-00. All parameters tested, including physical description, identification by HPLC, disintegration time, assay, microbial enumeration, and tests for specified micro-organisms, were found to be within the prescribed acceptance criteria.

The present product is assessed as complying with the requirements in the course of the tests carried out.

AUTHORISATION

Analyzed By	Checked By	Approved By
Name	Name	Name
		
Signature	Signature	Signature
		
Date	Date	Date
28/04/2026	28/04/2026	28/04/2026



We confirm that the results shown in this certificate refer only to the investigated sample. Conditions beyond our cognizance (inappropriate packing, transport, etc.) may affect the results. Reproduction of this inspection report is only permitted in its entirety. Permission to reproduce extracts must be obtained from our company. Future changes to the legal basis may lead to a reassessment.