

METAPHARMACEUTICAL

N DE LOTE:

0030923

F&A PHARMA

ESKAY IODINE
PRIVATE LIMITED

05/09/2023

FINISHED PRODUCT: CERTIFICATE OF ANALYSIS

DEPARTMENT : QUALITY CONTROL

Ref.SOP No QCS/017

Format No. : QC/017/F/02

Name of Product	Povidone Iodine EP	Date of Manufacture	May. - 2023
Batch Number	28048E23	Date of Expiry	Apr. - 2026
A.R. Number	QC/FP/0486/23	Batch Size	500 kg
Date of Analysis	27/05/2023	Date of Sampling	25/05/2023
Date of Release	27/05/2023	Sample Quantity	150 gm
Mfg. Lic. No.	G/25/2008		

Sr. No.	Test	Observation	Specification
01.	Appearance	A reddish brown amorphous powder.	A yellowish brown or reddish brown amorphous powder.
02.	Solubility	Complies	Soluble in water and in alcohol (96%); practically insoluble in acetone.
03.	Identification		
	By IR	Complies	To be comply when compared with iodinated Povidone CRS.
	By Chemical	Complies	An intense blue colour is produced.
04	pH of 10% w/v Solution	1.95	1.5 - 5.0
05	Iodide on dry basis	3.50	Not More Than 6.0 %
06.	Loss On Drying @ 105°C.	2.70	Not More Than 8.0 %
07.	Sulphate Ash	0.005	Not More Than 0.1 %
08.	Assay on dried basis (Available Iodine)	11.72	Between 9.0 % to 12.0 %

Remarks: The product complies with EP Specification.

Prepared By

Checked By

Approved By

Signature

Date

(Sanjana Purohit - Officer -QC)

(Prakash Chauhan-Sr.Executive -QC) (Raman Kaushik - GM Quality)



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