Sri Krishna Pharma
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CERTIFICATE OF ANALYSIS

Q.C DEPARTMENT,
DATE: 28.09.2024

Product	**PARACETAMOL IP**		
Batch No.	**8289/24-25**	**Manufacturing Date**	**September, 2024**
Batch Qty.	**2255.120 Kg**	**Expiry/ Retest Date**	**August, 2029**
A.R. No.	**FPC/PF3/0316/24-25**	**Date of Analysis**	**25.09.2024**

S.No.	TESTS	SPECIFICATIONS	RESULTS
1	**Description**	White crystals or white crystalline powder	White crystalline powder
2	**Solubility**	Freely soluble in ethanol (95%), sparingly soluble in water, very slightly soluble in Dichloromethane and in ether	Complies
3	**Identification**		
	B) By UV absorbance at about 249 nm	0.427 to 0.453	0.449
	C) Colour test	A violet colour develops which does not turn red.	Complies
	D) Test for Acetyl Groups	A blue color develops at the junction of two drops after 1 or 2 min. The colour intensifies and persists for a short time.	Complies
4	**Heavy Metals**	Not more than 10 ppm	Less than 10 ppm
5	**Sulphated Ash**	Not more than 0.10%	0.04%
6	**Loss on Drying**	Not more than 0.50%	0.15%
7	**Assay (On dried basis)**	Not less than 99.0% and Not more than 101.0% of $C_8H_9NO_2$	99.7%
8	**Related Substances (By HPLC)**		
	A) 4-Chloroacetanilide	Not more than 10 ppm	BDL(<0.5 ppm)
	B) 4-Aminophenol	Not more than 50 ppm	BDL(<1.15 ppm)
	C) 4-Nitrophenol	Not more than 0.05%	BDL(<0.0009%)
	D) Any other impurity	Not more than 0.05%	0.01%
	E) Total of other impurities	Not more than 0.1%	0.01%

CONCLUSION: The Material Complies/ Does not Complies to the above Specifications of IP 2022.

Remarks: BDL- Below Detection Limit, BDRL- Below Disregard Limit.

PREPARED BY QC DATE: 28.09.24

CHECKED BY QA DATE: 28.09.2024

APPROVED BY Sr. MANAGER -QC/DESIGNATE DATE: 28/09/2024

ISSUED TO: M/s. BCM CORPORATION.

REPORT NO.:8289(A)

QTY.: 2250 Kg

ISSUED DATE: 28.09.2024

QC/GEN/016-F-01/00

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