

M/S IVM PHARMACIA

Plot No. 05, Industrial Township, Bhatoli- Kalan, Baddi (H.P.) -173205 CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

	QUALITY CONTROL D	EPARTMENT	
PRODUCT NAME	PARACETAMOL INFUSION IP (1000 MG/100 ML)		IK24120
GENERIC NAME	PARACETAMOL INFUSION IP (1000 MG/100 ML)	A. R. NO.	QC/FP/24/0501
MFG. DATE	03/2024	PRODUCT CODE	K
EXP .DATE	02/2026	BATCH SIZE	1500 LTR
SAMPLE SIZE	27 X 100 ML	PACK SIZE	100 ML
DATE TEST STARTED	01/04/2024	DATE TEST COMPLETED	16/04/2024
MFG. LIC. NO.	LVP/18/09	SPECIFICATION NO.	FPS/F010-03

S.NO.	TEST	OBSERVATION	SPECIFICATION	
1	Description	A clear colorless solution	A clear colorless solution.	
2	Identification	Principle peak in assay solution is corresponds to principle peak in reference solution	In the assay, the principle peak in the chromatogram obtained with test solution corresponds to that in the chromatogram obtained with reference solution.	
3	pH	5.73	4.5 to 6.5	
4	Light absorption	0.0011	The absorbance of infusion at 500 nm is no more than 0.04.	
5	Related Substances	ND ND 0.027%	4 Amino phenolNMT 0.1% 4 ChloroacetanilideNMT 10 ppm Any other secondary peaksNMT 0.25%	
6	Extractable volume	102 ml	Not less than nominal volume	
7	Bacterial Endotoxins Test	Less than 2.0EU/ml	Not more than 2.0 EU/ml of Paracetamol	
8	Sterility	Sterile	No growth observed during incubation period of 14 days.	
9	Particulate Contamination	26.67 particles/container 6.67 particles/container	≥ 10 µm Not more than 6000 particles/ container. ≥ 25 µm Not more than 600 particles/ container.	
10	Assay Each 100ml contains: Paracetamol IP1000mg Mannitol IP5.0%w/v	1011.45 mg/100ml (101.1%) 5.05 %w/v (101.1 %)	900.0mg to 1100.0mg/100 ml (90.0 % to 110.0 % of label claim) 4.50 % w/v to 5.50 % w/v (90.0 % to 110.0 % of label claim)	

Remark: The above sample complies / does not comply as per fH/ IP/BP/USP specification or external lab

analysis report. ND: Not Detected

	ANALYSED BY	CHECKED BY	APPROVED BY
Signature/ Date	Rakhi 24	Cond 16/10/1/24	Cony 16/04/24
18.	Officer-QC	Executive-QC	Quality Head

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