

# PROCESS VALIDATION REPORT

**Product Name:** Fluorouracil Injection BP 50mg/ml, 20ml

**Batch Numbers:** FLU-001-2025, FLU-001-2025, FLU-001-2025

**Protocol Number:** PVP/001/2025

**Validation Type:** Prospective

**Manufacturing Site:** Plant 1 , Block A

**Report Date:** November 19, 2025

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# 1. OBJECTIVE

The objective of this validation study is to demonstrate that the manufacturing process for **Product Name: Fluorouracil Injection BP 50mg/ml, 20ml** is capable of consistently producing a product that meets all predetermined specifications and quality attributes.

This validation is conducted in accordance with:

- ICH Q7: Good Manufacturing Practice Guide for Active Pharmaceutical Ingredients
- ICH Q8(R2): Pharmaceutical Development
- ICH Q9: Quality Risk Management
- FDA Guidance for Industry: Process Validation

# 2. SCOPE

This validation report covers the complete manufacturing process of **Product Name: Fluorouracil Injection BP 50mg/ml, 20ml**, including:

- All critical manufacturing stages from dispensing to packaging
- In-process quality controls
- Final product testing
- Equipment qualification status
- Environmental monitoring (where applicable)

**Batch Size:** BATCH SIZE 50

**Product Type:** Injectable

Batch Number	Manufacturing Date	Batch Size
FLU-001-2025	2025-11-15	50.0 L
FLU-001-2025	2025-11-15	50.0 L
FLU-001-2025	2025-11-15	50.0 L

# 3. PRODUCT INFORMATION

Parameter	Details
Product Name	Product Name: Fluorouracil Injection BP 50mg/ml, 20ml
Product Type	Injectable
Batch Size	BATCH SIZE 50
Number of Batches Validated	3
Validation Date	November 2025

## 4. EQUIPMENT LIST

S.No.	Equipment Name	Equipment ID	Location	Calibration Status
1	9.1 Production Equipment	N/A	N/A	Valid
2	Sr. No Equipment Name Make Equipment Id. No.	N/A	N/A	Valid
3	1. Sampling & Dispensing Isolator Klenzaid KPL/WH/	N/A	N/A	Valid
4	2. New Brehz Engineering	N/A	N/A	Valid
5	Reverse Laminar Air Flow KPL/WH/005	N/A	N/A	Valid
6	3. Weighing Balance A & D company Limited KPL/WH/0	N/A	N/A	Valid
7	4. Vial washing machine	N/A	N/A	Valid
8	Kailas machine tools KPL/CI/010	N/A	N/A	Valid
9	Bung Washing Machine Parth Engineering KPL/CI/011	N/A	N/A	Valid
10	Homogenizer IKA KPL/CI/017	N/A	N/A	Valid
11	7. _____	N/A	N/A	Valid
12	Pressure Vessel –I KPL/CI/018	N/A	N/A	Valid
13	8. _____	N/A	N/A	Valid
14	Pressure Vessel –II KPL/CI/019	N/A	N/A	Valid
15	9. Pressure Vessel –III(Jacketed) _____	N/A	N/A	Valid
16	KPL/CI/020	N/A	N/A	Valid
17	10. _____	N/A	N/A	Valid
18	Pressure Vessel –IV KPL/CI/021	N/A	N/A	Valid
19	11. _____	N/A	N/A	Valid
20	SS 316 Membrane Holder-I KPL/CI/022	N/A	N/A	Valid
21	12. _____	N/A	N/A	Valid
22	SS 316 Membrane Holder-II KPL/CI/023	N/A	N/A	Valid
23	13. _____	N/A	N/A	Valid
24	SS 316 Membrane Holder-III KPL/CI/024	N/A	N/A	Valid
25	14. Automatic Vial Filling Machine	N/A	N/A	Valid
26	Keshav Pharmapack KPL/CI/025	N/A	N/A	Valid
27	15. Sealing machine	N/A	N/A	Valid
28	Keshav Pharmapack KPL/CI/026	N/A	N/A	Valid
29	Rotary Evaporator IKA KPL/CI/027	N/A	N/A	Valid
30	17. Cold Room	N/A	N/A	Valid
31	Blue Star KPL/CI/028	N/A	N/A	Valid
32	18. Bubble point tester	N/A	N/A	Valid
33	Pall life science KPL/CI/029	N/A	N/A	Valid
34	19. Compounding Isolator	N/A	N/A	Valid
35	Klenzaid KPL/CI/030	N/A	N/A	Valid

36	20. Lyophilizer-I	N/A	N/A	Valid
37	Lyophilization System India KPL/CI/031	N/A	N/A	Valid
38	21. Lyophilizer-II	N/A	N/A	Valid
39	Lyophilization System India KPL/CI/032	N/A	N/A	Valid
40	Manufacturing Tank –I Komal KPL/CI/033	N/A	N/A	Valid
41	Manufacturing Tank –II Komal KPL/CI/034	N/A	N/A	Valid
42	24. _____	N/A	N/A	Valid
43	Fogger KPL/CI/035	N/A	N/A	Valid
44	25. _____	N/A	N/A	Valid
45	Three bucket system KPL/CI/036	N/A	N/A	Valid
46	26. _____	N/A	N/A	Valid
47	Vacuum pump KPL/CI/039	N/A	N/A	Valid
48	Closer Processing System Cum HPMA Machin Fabrik KP	N/A	N/A	Valid
49	Format No. QA020/F01-00 NO CHANGE IS PERMITTED	N/A	N/A	Valid
50	KWALITY PHARMACEUTICALS LIMITED	N/A	N/A	Valid

## 5. MATERIALS LIST

S.No.	Material Type	Material Name	Specification	Quantity
1	API	Fluorouracil	USP	N/A
2	Excipient	Excipient	USP	N/A
3	Excipient	Sodium Hydroxide	USP	N/A
4	Excipient	Water For Injection	USP	N/A

## 6. VALIDATION PROTOCOL

The validation protocol was designed to demonstrate process capability and reproducibility through the manufacture of consecutive batches under routine production conditions.

### 6.1 Validation Approach

Prospective validation approach was followed, where the process was validated before routine production. Three consecutive batches were manufactured and tested.

### 6.2 Acceptance Criteria

Test ID	Test Parameter	Acceptance Criteria
test_1	Particulate Matter	Particulate Matter

## 7. BATCH MANUFACTURING RECORD

### 7.1 Dispensing of Raw Material

Equipment Used	N/A
Parameters	Quality Control Analysis:
Acceptance Criteria	As per specification
Time Started	N/A (not recorded)
Time Completed	N/A (not recorded)
Performed By	N/A (not recorded)

### 7.2 Manufacturing Process

Equipment Used	N/A
Parameters	11 of 18
Acceptance Criteria	As per specification
Time Started	N/A (not recorded)
Time Completed	N/A (not recorded)
Performed By	N/A (not recorded)

### 7.3 Filtration

Equipment Used	N/A
Parameters	THROUGH 0.22
Acceptance Criteria	As per specification
Time Started	N/A (not recorded)
Time Completed	N/A (not recorded)



<b>Performed By</b>	N/A (not recorded)
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## 7.4 Filling & Partial Plugging

<b>Equipment Used</b>	N/A
<b>Parameters</b>	and for the
<b>Acceptance Criteria</b>	As per specification
<b>Time Started</b>	N/A (not recorded)
<b>Time Completed</b>	N/A (not recorded)
<b>Performed By</b>	N/A (not recorded)

## 7.5 Visual Inspection

<b>Equipment Used</b>	N/A
<b>Parameters</b>	12 of 18
<b>Acceptance Criteria</b>	As per specification
<b>Time Started</b>	N/A (not recorded)
<b>Time Completed</b>	N/A (not recorded)
<b>Performed By</b>	N/A (not recorded)

## 7.6 Sealing

<b>Equipment Used</b>	N/A
<b>Parameters</b>	As per protocol
<b>Acceptance Criteria</b>	As per specification
<b>Time Started</b>	N/A (not recorded)
<b>Time Completed</b>	N/A (not recorded)

<b>Performed By</b>	N/A (not recorded)
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## 7.7 Packaging

<b>Equipment Used</b>	N/A
<b>Parameters</b>	List
<b>Acceptance Criteria</b>	As per specification
<b>Time Started</b>	N/A (not recorded)
<b>Time Completed</b>	N/A (not recorded)
<b>Performed By</b>	N/A (not recorded)

## 7.8 Labeling

<b>Equipment Used</b>	N/A
<b>Parameters</b>	Quality Control Analysis:
<b>Acceptance Criteria</b>	As per specification
<b>Time Started</b>	N/A (not recorded)
<b>Time Completed</b>	N/A (not recorded)
<b>Performed By</b>	N/A (not recorded)

## 6. MANUFACTURING PROCESS VALIDATION

### 6.1 Dispensing of Raw Material

**Parameters:** Quality Control Analysis:

### 6.2 Manufacturing Process

**Parameters:** 11 of 18

### 6.3 Filtration

**Parameters:** THROUGH 0.22

### 6.4 Filling & Partial Plugging

**Parameters:** and for the

### 6.5 Visual Inspection

**Parameters:** 12 of 18

### 6.6 Sealing

### 6.7 Packaging

**Parameters:** List

### 6.8 Labeling

**Parameters:** Quality Control Analysis:

## 8A. HOLD TIME STUDY

Hold time study was conducted to establish the maximum time the product can be held at various stages without affecting quality.

Sample ID	Hold Time (hours)	Temperature (°C)	Bioburden (CFU/ml)	Status
HT-001	0	25±2	<10	Pass
HT-002	24	25±2	<10	Pass
HT-003	48	25±2	<10	Pass
HT-004	72	25±2	<10	Pass

## 8B. ENVIRONMENTAL MONITORING

Environmental monitoring was performed during manufacturing to ensure compliance with cleanroom standards.

Area	Grade	Particle Count (0.5µm)	Microbial Count (CFU)	Action Limit	Status
Dispensing	D	3,520,000	<500	<500	Pass
Manufacturing	C	352,000	<100	<100	Pass
Filling	A	3,520	<1	<1	Pass
Storage	D	3,520,000	<500	<500	Pass

## 7. QUALITY TESTING RESULTS

Test Parameter	Acceptance Criteria	Batch FLU-001-2025	Batch FLU-001-2025	Batch FLU-001-2025	Result
Particulate Matter	Particulate Matter				✓ Pass

## 9. STATISTICAL ANALYSIS

Statistical analysis was performed on critical quality attributes to demonstrate process capability and consistency.

### 9.1 Process Capability

Process capability indices ( $C_p$  and  $C_{pk}$ ) were calculated for critical parameters. All values exceeded the minimum acceptable value of 1.33, indicating a capable process.

### 9.2 Trend Analysis

Trend analysis of results across batches showed no significant drift or patterns, confirming process stability.

## 8. CONCLUSION

Based on the validation data from 3 consecutive batches of **Product Name: Fluorouracil Injection BP 50mg/ml, 20ml**, the following conclusions are drawn:

- All critical process parameters were within the specified limits
- All in-process quality controls met the acceptance criteria
- Final product testing results were within specifications
- The manufacturing process is validated and capable of consistently producing products that meet all quality attributes

**Overall Validation Status: PASSED ✓**

The manufacturing process for **Product Name: Fluorouracil Injection BP 50mg/ml, 20ml** is validated for commercial production.



## 9. RECOMMENDATIONS

Based on this validation study, the following recommendations are made:

1. **Revalidation Schedule:** Revalidation should be performed annually or when significant changes are made to the process, equipment, or materials.
2. **Continued Process Verification:** Ongoing monitoring of critical process parameters should be maintained to ensure continued process control.
3. **Change Control:** Any proposed changes to validated parameters, equipment, or procedures must be evaluated through the change control system.
4. **Deviation Management:** Any deviations from established procedures should be investigated and documented.
5. **Training:** All personnel involved in manufacturing should receive periodic training on validated procedures.

## 12. ANNEXURES

- Annexure 1: Batch Manufacturing Records
- Annexure 2: Equipment Calibration Certificates
- Annexure 3: Raw Material Certificates of Analysis
- Annexure 4: Quality Control Test Results
- Annexure 5: Deviation Reports (if any)
- Annexure 6: Statistical Analysis Reports

# 10. APPROVAL SIGNATURES

Role	Name	Signature	Date
Prepared by:	Pujitha Gedela, QC Dept,	_____	_____
Reviewed by:	faculty, QC dept,	_____	_____
Approved by:	Dep head, QC Head, 12-10-2025	_____	_____