PROCESS VALIDATION REPORT

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

Batch Numbers: FLU-001-2024, FLU-002-2024, FLU-003-2024

Report Date: November 12, 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

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

No equipment data available.

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. MANUFACTURING PROCESS VALIDATION

No stage data available.

7. QUALITY TESTING RESULTS

Test Parameter	Acceptance Criteria	Batch FLU-001-2024	Batch FLU-002-2024	Batch FLU-003-2024	Result
Particulate Matter	Particulate Matter	N/A	N/A	N/A	✗ Fail

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.

10. APPROVAL SIGNATURES

Role	Name	Signature	Date
Prepared by:	_____	_____	_____
Reviewed by:	_____	_____	_____
Approved by:	_____	_____	_____