

Process Validation Interim / Final Report**Product Title:****PRODUCT CODE:**

	WRITTEN BY:	REVIEWED BY:
Name:		
Signature:		
Position:		
Date:		

Qualification Status
Qualification of _____ as per protocol no. _____ has been completed.
All deviations and additional protocol results for the batch are documented in this interim report. All acceptance criteria have been met according to protocol no. _____.
☐ The qualification for the use of _____ has been successfully completed.
☐ The qualification status of the use of _____ in the manufacture remains on-going until all qualification data has been compiled for this study and will be documented in a subsequent report.

REPORT COMPLETION APPROVAL:			
Name:			
Signature:			
Position:	Validation Manager	Quality Engineer	QA Team-Leader
Date:			

1. OBJECTIVE

2. VALIDATION STRATEGY

3. RESULTS

4. CONCLUSIONS AND RECOMMENDATIONS

Entered by:	Date:
Transcription Verified by:	Date:

Note. Adapted from gmpsop.com (2024)