TEM-295

Issue date: _____

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)