PHARMASOL	IN-PROCESS RELEASE FORM	Procedure:	100-5-760
		DCO:	19-0322
		Rev: 2	Page: 1 of 2

Title:	**Clobetasol Propionate Foam, 0.05% (Concentrate)**			Effective Date: 08/29/19
Customer:	**Aucta Pharma**	Item #	**8112010P; 8112010E**	Dept. Quality Control

Prepared By:	Date:	Dept. Approval By:	Date:	QA Approval By:	Date:
	07/22/19		07/22/19		08/15/19

Batch# _____

TEST	METHOD	TEST SPECIFICATION	RESULTS	NOTEBOOK REFERENCE
1. Appearance (at 45°C)	Visual	Clear solution with no visible particles		
2. Identification by HPLC	100-3-283	The retention time (RT) of Clobetasol Propionate in the test sample solution shall correspond to the retention time of Clobetasol Propionate in the standard solution for the assay chromatogram.		
3. Identification by UV	100-3-283	UV spectrum matches that of standard.		
4. pH	Current USP<791>	5.0-7.0		
5. Assay by HPLC	100-3-283	Not less than 90.0 percent and not more than 110.0 percent of the labeled amount of Clobetasol Propionate		

Batch# _____

TEST	METHOD	TEST SPECIFICATION	RESULTS	NOTEBOOK REFERENCE
6. Ethanol Content	100-3-143	90.0-110.0%		

QC Tested by: _____ Date: _____

QC Reviewed by: _____ Date: _____

QA Reviewed by: _____ Date: _____