

97	690	12-35	566

CERTIFICATE OF COMPLIANCE

PRODUCT DESCRIPTION : EXCIPIAL [®] - LIPOCREME 50g BATCH NUMBER : 9769012							
MANUFACTURING SITE : LABORATOIRES GALDERMA, ZI MONTDESIR, ALBY SUR CHERAN, 74540, FRANCE							
AUTHORIZATION NUMBER : MM 18/037 - February 16, 2018							
IMPORTING COUNTRY (DESTINATION) : FRANCE							
FORMULA NUMBER : 0769	MANUFACTURING DATE (MM/DD/YYYY) : 05/08/2019						
PRODUCT CODE : 023566	SHELF LIFE : 36 MONTHS						
		EXPIRY DATE (MM/YYYY) : 05/2022					
REFERENCE SPECIFICATION : MT.09.DOC.1433.R	ANALYSIS TYPE : Full						
ANALYSIS NUMBER : B-20190520-00022							
RELEASED UNITS QUANTITY : 13440 Units							
MEAN FILL (>= Nominal value of label claim AC.02.SOP.5005)	50.6 g						
TESTS	METHODS	SPECIFICATIONS	RESULTS				
Macroscopic appearance	Visual examination						
Form		Semi solid emulsion	Complies				
Color Other Decretics		White	Complies				
Other Properties		Smooth, supple	Complies				
Odor	Smell Examination	Characteristic (perfumed)	Complies				
Structure	Visual examination (Microscope)	No crystal visible, shiny structure <= 50µm visible	corresponds				
Total aerobic microbial count (TAMC)	Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05*	NMT 100 CFU/g	< 100 CFU/g				
Total combined yeast/mould count (TYMC)	Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05*		< 10 CFU/g				
Specific microorganisms	Ph. Eur 2.6.13/USP 1111,62/JP G4,4.05*						
Pseudomonas aeruginosa Staphylococcus aureus		Absence per gram Absence per gram	Absence per gram Absence per gram				
*All pharmacopools mothods are carried out according to the su	urrant adition						

*All pharmacopoeia methods are carried out according to the current edition.

Labvantage 0601.012.01

🖧 GALDERMA



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TESTS	METHODS	SPECIFICATIONS	RESULTS

NOTES : N/A

Certification :

I hereby certified that above information is authentic and accurate.

This batch has been manufactured (including packaging and quality control), at the above mentioned site in full compliance with the GMP requirements of the local regulatory authorities and with the specification in the marketing authorization of the importing country.

The batch processing, packaging and analysis records were reviewed and found to be in compliance with the GMP. All major processes (manufacturing and analytical) have been validated. All deviation or changes have been properly investigated, evaluated and approved by authorized personal.

This document has been electronically signed by a qualified person :

Aurélie GARREAU Date : June 10, 2019 🖧 GALDERMA