

RESULTS

CERTIFICATE OF COMPLIANCE

PRODUCT DESCRIPTION: EXCIPIAL® - LIPOCREME 50g

BATCH NUMBER: 1769009

MANUFACTURING SITE: LABORATOIRES GALDERMA, ZI MONTDESIR, ALBY SUR CHERAN, 74540, FRANCE

METHODS

AUTHORIZATION NUMBER: M 20/018 - February 12, 2020

IMPORTING COUNTRY (DESTINATION): FRANCE

FORMULA NUMBER: 0769 MANUFACTURING DATE (MM/DD/YYYY): 05/07/2021

SHELF LIFE: 36 MONTHS PRODUCT CODE: 023566

EXPIRY DATE (MM/YYYY): 04/2024

SPECIFICATIONS

REFERENCE SPECIFICATION: MT.09.DOC.1433.R00.1 ANALYSIS TYPE: Full

ANALYSIS NUMBER: B-20210510-00032 RELEASED UNITS QUANTITY: 13608 Units

TFSTS

MEAN FILL (>= Nominal value of label claim AC.02.SOP.5005) : 50.3 g

INIE I HODS	SPECIFICATIONS	RESULTS
Visual examination		
	Semi solid emulsion	Complies
	White	Complies
	Smooth, supple	Complies
Smell Examination	Characteristic (perfumed)	Complies
Visual examination (Microscope)	No crystal visible, shiny structure $\leq 50 \mu m$ visible	corresponds
Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05*	NMT 100 CFU/g	< 100 CFU/g
Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05*	NMT 10 CFU/g	< 10 CFU/g
Ph. Eur 2.6.13/USP 1111,62/JP G4,4.05*		
	Absence per gram	Absence per gram
	Absence per gram	Absence per gram
	Visual examination Smell Examination Visual examination (Microscope) Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05* Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05* Ph. Eur 2.6.13/USP	Visual examination Semi solid emulsion White Smooth, supple Characteristic (perfumed) Visual examination (Microscope) Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05* Ph. Eur 2.6.12/USP 1111,61/JP G4,4.05* NMT 100 CFU/g NMT 10 CFU/g NMT 10 CFU/g Absence per gram

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

GALDERMA

1769009-3566

CERTIFICATE OF COMPLIANCE

PRODUCT DESCRIPTION: EXCIPIAL® - LIPOCREME 50g

BATCH NUMBER: 1769009

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 :

GOEHRY Calvin

Date: May 27, 2021