RESULTS

CERTIFICATE OF COMPLIANCE

PRODUCT DESCRIPTION: EXCIPIAL® - HYDROCREME 50g

BATCH NUMBER: 9767004

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

METHODS

AUTHORIZATION NUMBER: MM 18/037 - February 16, 2018

IMPORTING COUNTRY (DESTINATION): FRANCE

FORMULA NUMBER: 0767 MANUFACTURING DATE (MM/DD/YYYY): 08/13/2019

PRODUCT CODE: 023565 SHELF LIFE: 36 MONTHS

EXPIRY DATE (MM/YYYY): 07/2022

SPECIFICATIONS

REFERENCE SPECIFICATION: MT.09.DOC.1473.R01 ANALYSIS TYPE: Full

ANALYSIS NUMBER: B-20190822-00056 RELEASED UNITS QUANTITY: 18240 Units

TESTS

MEAN FILL (>= Nominal value of label claim AC.02.SOP.5005) : 50.4 g

| <u></u> | IVIETTIODS | 31 EGII ICATIONS | |
|---|---|---------------------|------------------|
| Macroscopic appearance | Visual examination | | |
| Form | | Semisolid emulsion | Complies |
| Color | | White | Complies |
| Other Properties | | Homogeneous | Complies |
| Odor | Smell Examination | Almost odorless | Complies |
| Structure | Visual examination (Microscope) | No crystals visible | corresponds |
| рН | Ph. Eur 2.2.3 * | 3.5 - 6.5 | 5.6 |
| 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 | | Absence per gram | Absence per gram |
| Staphylococcus aureus | | Absence per gram | Absence per gram |
| | | | |

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

GALDERMA

9767004-3565

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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: September 09, 2019