

Diversey Europe Operations EV Massesenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date:

21 January 2021

Product Name	CLEARKLENS CLEANSINALD SS VH9S				
Product Code	7516431				
Batch Number	FMP21019	51130			
Production Date	19/01/2021				
Expiration Date	EXP 19/07/2022				

This is to certify that the above batch of product has been tested and conforms to the manufacturing Quality Control specification for the product.

Test	Test Method	Limits		Results
		Lower	- Upper	
Appearance	Visual	Clear Slightly	Yellow Liquid	Clear Slightly Yellow Liquid
pH (neat solution)	DM001	9.0	11.4	10.9
Specific Gravity (20°C)	DM004	0.990	1.010	0.997

On behalf of Diversey site	Name:	Justyna Staron	Angelika Partynska
Quality Manger	Position	Quality Control Inspector	

This document being issued electronically does not bear a signature

COA Template Version : 02 Date of issuing : November 24th

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-15901A

Product CodeLot NumberQuantityUOMC/SINALD SS 900ml Bottle DV4673WO5112911 Case

Validation Reference Number: 4673

Processing Run Start Date: 19-Dec-20 02:00 AM

Processing Run End Date: 19-Dec-20 07:29 AM

Specified Dose Range (kGy): 25.0 - 40.0 Calculated Min Dose (kGy): 28.9

Reference Dose Range (kGy): 27.3 - 36.6 Calculated Max Dose (kGy): 36.6

PO Number: 37043

Gamma Process Run Approval authorized by STERIS

Operating facilities are in compliance with applicable regulations providing services under a certified quality system which meets the requirements of FDA QSR, EN/ISO 13485 current certified version, and is in alignment with EN ISO 11137 current certified version

Processing Location

Synergy Health Sterilisation UK Limited, a STERIS Company Brunel Close Drayton Fields Industrial Estate Daventry Northants NN11 8RB

Phone: + 44(0) 1327 706 111

Items Iraddiated under WO51129 will be used in finished batch Fmp21019 51130.

Olga Kirchner

Document ID: 67763

Last Revised in Rel 2.0.0.0

Rel Date: 08/13/2018 Page 1 of 1

PHARMACEUTICAL ANALYSIS REPORT



Flexible Medical Packaging

Unit 8 Hightown White Cross Industrial Estate Lancaster LA14XS

FAO:

Mrs.Olga Kirchner

Report of Tests on:

Cleansinald SS

Sample Description:

Sample Code: Fmp21019 51130

Lucideon Sample Number: UK21310-2065

Lucideon Report Number: UK21310-2065/MFEP

Issue Number:

Date Logged:

22-Jan-2021

Order Number:

PO 37271

Date Reported:

19-Feb-2021

Date(s) of Test(s):

04-Feb-2021 to 18-Feb-2021

Sterility Testing

Membrane Filtration EP

Test Results:

Result: Pass

The test results meet the EP/USP criteria: Yes

Tests carried out in accordance with cGMP

End of Test Report

0BOOT 19-Feb-21 Mrs Natalie Boot

This report is issued in accordance with the Conditions of Business of Lucideon Limited and relates only to the sample(s) tested.