

Diversey Europe BV Maarasenbroeksedlijk 2 3542 DN Utrecht The Netherlands

Tel. +31 (0)30 247 6427 www.Diversey.com

CERTIFICATE OF ANALYSIS

Date: 28 11 2019

Product Name	CLEARKLENS CLEANSINALD SC VH9		
Product Code			
Batch Number	FMP 19 332 48103		
Production Date	28/11/2019		
Expiration Date	EXP 28 111 2021		

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	Limits		Results	
	Method	Lower	- Upper		
Appearance	Visual	Clear Slightly Yellow Liquid		Clear Slightly	
pH (neat solution)	DM001	12.0	13.0	12. Q	
Specific Gravity (20°C)	DM004	1.040	1.060	1.052	
Cationic content	DM020	14.25	15.75	14.51	

On hehalf of Director site	Name:	Navaharda T. Stawn
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

COA Template Version: 02 Date of issuing: November 24th 2017

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-5272A

Product CodeLot NumberQuantityUOMC/SINALD BOTTLES/CAPS/BAGSWO4810411 Case

DV4894

Validation Reference Number: 4894

Processing Run Start Date: 27-Sep-19 11:01 PM
Processing Run End Date: 28-Sep-19 04:49 AM

Specified Dose Range (kGy): 25.0 - 40.0 Calculated Min Dose (kGy): 30.1

Reference Dose Range (kGy): 27.4-36.5 Calculated Max Dose (kGy): 36.9

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 Irradiated under WO48104 will be used in finished batch Fmp19332 48103.

Olga Kirchner

Document ID: 29375

Rel Date: 08/13/2018

Last Revised in Rel 2.0.0.0

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 SC

Sample Description:

Sample Code: Fmp 19332 48103

Lucideon Sample Number: (196608)-50137

Lucideon Report Number: (196608)-50137/MFEP

Issue Number:

1

Date Logged:

05-Dec-2019

Order Number:

PO34403

Date Reported:

06-Jan-2020

Date(s) of Test(s):

07-Dec-2019 to 04-Jan-2020

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

BCOL 06-JON-20

Mrs Natalie Boot

Senior Business Support Assistant