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CERTIFICATE OF ANALYSIS

Date: 15 June 2022

Product Name	CLEARKLENS CLEANSINALD SC VH9		
Product Code	7516429		
Batch Number	FMP22166	55390	
Production Date	15/06/2022		
Expiration Date	EXP 15/06/2024		

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	12.0	13.0	13.0
Specific Gravity (20°C)	DM004	1.040	1.060	1.050
Cationic Content	DM020	14.25	15.75	15.18

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

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

LUCIDEON

insight creating advantage

Flexible Medical Packaging

Unit 8
Hightown
White Cross Industrial Estate
Lancaster
LA1 4XS

FAO: Mrs.Olga Kirchner

Report of Tests on: Cleansinald SC

Sample Description: Sample Code: Fmp22166 55390

Lucideon Sample Number: UK222549-16274

Lucideon Report Number: UK222549-16274/MFEP **Issue Number:** 1

Date Logged: 22-Jun-2022 **Order Number:** 402814

Date Reported: 13-Jul-2022 **Date(s) of Test(s):** 24-Jun-2022 to 08-Jul-2022

Sterility Testing

Membrane Filtration EP

Test Results:

The test results meet the EP/USP criteria: Yes

Result: Pass

Tests carried out in accordance with cGMP

End of Test Report

NCR001 13-Jul-22

Mrs Natalie Boot

Senior Business Support Administrator

Page 1 of 1

This report is issued in accordance with the Conditions of Business of Lucideon Limited and relates only to the sample(s) tested. No responsibility is taken for the accuracy of the sampling unless this is done under our own supervision. This report shall not be reproduced in part without the written approval of Lucideon Limited, nor used in any way as to lead to misrepresentation of the results or their implications.

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STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID 2173-28640A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
C/SINALD BOTTLES/CAPS/BAGS DV4894	54518	16	Case

Validation Reference Number: 4894

Processing Run Start Date: 28-May-2022 3:05 PM

Processing Run End Date: 28-May-2022 9:12 PM

Specified Dose Range (kGy):	25.0 - 40.0	Calculated Min Dose (kGy):	29.1
Reference Dose Range (kGy):	27.4 - 36.5	Calculated Max Dose (kGy):	36.5

PO Number: 402391

Product meets Customer specifications; zero nonconformities occurred during this irradiation run.

Gamma Process Run Approval authorized by STERIS

Date/Time E-Signed: 29-May-2022 3:05 PM

□

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 WO54518 will be used in finished batch Fmp22166 55390.

Olga Kirchner