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CERTIFICATE OF ANALYSIS

Date: 13 April 2026

Product Name	CLEARKLENS CLEANSINALD SC VH9
Product Code	101103552
Batch Number	FMP26103 63818
Production Date	13/04/2026
Expiration Date	EXP 13/04/2028

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 Lower**	**- Upper**	**Results**
Appearance	Visual	Clear Slightly Yellow Liquid		Clear Slightly Yellow Liquid
pH (neat solution)	DM001	12.0	13.0	12.9
Specific Gravity (20°C)	DM004	1.040	1.060	1.052
Cationic Content	DM020	14.25	15.75	15.20

On behalf of Diversey site Quality Manger	Name:	Justyna Staron Aneta Wiktorowicz
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

COA Template	*Version : 04*	*Date of issuing : March 11th 2019*

STERIS: Certificate Of Processing

Prepared For **FLEXIBLE MEDICAL PACKAGING LTD (8245)**

Gamma Process Run ID **2173-63042A**

Product Code	Lot Number	Quantity	UOM
C/SINALD BOTTLES/CAPS/BAGS DV4894	63824	29	Case

Validation Reference Number: 4894

Processing Run Start Date **21-Mar-2026 3:12 PM**

Processing Run End Date **21-Mar-2026 9:53 PM**

Specified Dose Range (kGy)	**25.0 - 40.0**	**Calculated Min Dose (kGy)**	**28.7**
Reference Dose Range (kGy)	**27.4 - 36.5**	**Calculated Max Dose (kGy)**	**35.3**

PO Number: 412685

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

Gamma Process Run Approval authorized by STERIS

DateTime Esigned 22-Mar-2026 7:48 AM

Operating facilities are in compliance with applicable local/state and federal regulations providing services under a certified quality system which meets the following requirements (if applicable): FDA QMSR, ISO 9001, ISO 13485 current certified version, national pharmaceutical GMP (EU GMP, Commission Directive 2017/1572, Veterinary Regulation (EU) 2019/6, etc.) and is in alignment with ISO 11137 current certified version. STERIS AST certifies that these processed items received the indicated doses within the precision and accuracy of the dosimetry system used. The product has been processed as per the requirements of the Customer Agreement and/or Customer Specification.

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 WO63824 will be used in finished batch Fmp20103 63818.

Olga Kirchner

Document ID: 232961

PHARMACEUTICAL ANALYSIS REPORT

LUCIDEON
providing material solutions

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: Fmp26103 63818

Lucideon Sample Number: N261605 - 1

Lucideon Report Number: N261605 - 1/MFEP **Issue Number:** 1

Date Logged: 20-Apr-2026 **Order Number:** 412869

Date Reported: 12-May-2026 **Date(s) of Test(s):** 28-Apr-2026 **to** 12-May-2026

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

Signed by:
Natalie Boot
Signer Name: Natalie Boot
Signing Reason: I have reviewed this document
Signing Time: 12 May 2026 | 16:21 BST
90369E9048134123B65D5E8FBD375BF3

Mrs Natalie Boot
Senior Business Support Administrator

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Certificate Of Completion

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Certified Delivered	Security Checked	12-May-26 | 16:19
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Completed	Security Checked	12-May-26 | 16:22

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