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CERTIFICATE OF ANALYSIS

Date: 20 July 2026

Product Name	CLEARKLENS CLEANSINALD SS VH9S		
Product Code	FMP26198		
Batch Number	100848254	64437	
Production Date	17/07/2026		
Expiration Date	EXP 17/01/2028		

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		Results
		Lower	Upper	
Appearance	Visual	Clear Slightly Yellow Liquid		Clear Slightly Yellow Liquid
pH (neat solution)	DM001	9.0	11.4	10.3
Specific Gravity (20°C)	DM004	0.990	1.010	0.996

On behalf of Diversey site Quality Manger	Name:	Edyta Rodrigues Justyna Staron
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

COA Template	Version : 04	Date of issuing : March 11th 2019
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STERIS: Certificate Of Processing

Prepared For FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID 2173-65371A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
C/SINALD SS 900ml Bottle DV4673	64439	13	Case

Validation Reference Number: 4673

Processing Run Start Date 04-Jul-2026 3:12 PM

Processing Run End Date 04-Jul-2026 10:53 PM

Specified Dose Range (kGy)	25.0 - 40.0	Calculated Min Dose (kGy)	30.4
Reference Dose Range (kGy)	27.3 - 36.6	Calculated Max Dose (kGy)	37.6

PO Number: 413232

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

Gamma Process Run Approval authorized by STERIS

DateTime Esigned 05-Jul-2026 7:41 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 WO64439 will be used in finished batch Fmp26198 64437.

Olga Kirchner

PHARMACEUTICAL ANALYSIS REPORT

Flexible Medical Packaging

Unit 8
Hightown
White Cross Industrial Estate
Lancaster
LA1 4XS

FAO: Mrs. Olga Kirchner

Report of Tests on: Cleansinald SS

Sample Description: Sample Code: Fmp26198 64437

Lucideon Sample Number: N263081 - 1

Lucideon Report Number: N263081 - 1/MFEP

Issue Number: 1

Date Logged: 23-Jul-2026

Order Number: 413417

Date Reported: 10-Aug-2026

Date(s) of Test(s): 25-Jul-2026 to 08-Aug-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: 10 August 2026 | 15:12 BST

90369E9048134123B65D5E8FBD375BF3

Mrs Natalie Boot

Senior Business Support Administrator

Page 1 of 1

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Certificate Pages: 2
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Status: Completed

Envelope Originator:
Natalie Boot
Queens Road
Stoke-on-Trent, Staffordshire ST4 7LQ
Natalie.Boot@uk.lucideon.com
IP Address: 86.188.245.148

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Status: Original
10-Aug-26 | 15:11
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Natalie.Boot@uk.lucideon.com

Location: DocuSign

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Signature

Signed by:

Signer Name: Natalie Boot
Signing Reason: I have reviewed this document
Signing Time: 10 August 2026 | 15:12 BST
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Sent: 10-Aug-26 | 15:12
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