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CERTIFICATE OF ANALYSIS

Date: 28. 11. 2018

Product Name	CLEARKLENS CLEANSINALD SC VH9		
Product Code	7516429		
Batch Number	PHP 18332 46827		
Production Date	28/11/2013		
Expiration Date	Exp. 28/11/2020		

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	_	ntly Yellow Juid	clean slightly Yellow Liquist
pH (neat solution)	DM001	12.0	13.0	12.7
Specific Gravity (20°C)	DM004	1.040	1.060	1.048

On behalf of Diversey site Quality Manager	Name:	o. Linconne, Party usb		
	Position	Quality Control Inspector		

This document being issued electronically does not bear a signature

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



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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

07 Dec 2018

Date Tested:

20 Dec 2018

Date Test Completed:

08 Jan 2019

Purchase Order:

32264

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0046229/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

07/12/2018

Test Article: Sample Code:

Cleansinald SC FMP18332 45827

Batch Ref: 620529

Qty Received:

20 x 25mL Bottles + 5 Spare

Test	Method Item	Result
Sterility Test by Membrane Filtration (Steritest) Method	MM107/00	Pass
Growth in Tryptone Soya Broth at 20- 25°C after 14 days	MM107/01	No growth in one broth
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in one broth
Growth of Sub-culture in TSB at 20-25°C after 4 - 7 days	MM107/07	No growth in one broth
Growth of Sub-culture in THY at 30-35°C after 4 - 7 days	MM107/08	No growth in one broth
Volume Tested	MM107/05	20 x 25 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.0 2.6.1, USP 41 <71> & JP XVII 4.06
Product Standard Data Sheet	FG047/psd	FM06-01

Approval is provided by Electronic Signature. Their name and position is shown below.

CBIO! MRSB

Date: 11 Jan 2019 15:14:58

Mrs C Moore

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0046229 Print Number: P0056321 Page 1 of 1





http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 21-Oct-2018

UK33S12181434-3-2

This is to certify that AST Daventry Synergy Health PLC, a STERIS Company has where appropriate delivered an irradiation process in accordance with the current certified standards:

EN ISO 11137-1 Sterilisation of Health Care Products EN ISO 13485 Quality System - Medical Devices

Flexible Medical Packaging Ltd Unit 8, White Cross Ind Estate Hightown Lancaster Lancashire LA1 4XS UNITED KINGDOM

All in accordance with current Technical Agreement

Order information **Account Number:** 100432 Synergy Health Sales Part Reference: 1111497 **Customer Reference Number:** P031884 **Product Description:** C/SINALD BOTTLES/CAPS/BAGS DV4894 25-40KGS Validation Reference: 4894 **Quantity Received:** 16 **Customer Minimum Specification kGy:** 25.0 **Customer Maximum Specification kGy:** 40.0 **Customer Unit Lot/Batch Number:** W045828, 2 PLTS Irradiation Data Date and Time of Irradiation: 21-Oct-2018 13:45 Reference Dose Range kGy: 30.2 - 30.4 Calculated Minimum Dose kGy: 27.5 Calculated Maximum Dose kGy: 33.4

Items irradiated under WO45828, will be used in finished product batch FMP18332 45827

C Pascoe

Irradiation Release Authorised By Synergy Health Sterllisation UK, a STERIS Company