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CERTIFICATE OF ANALYSIS

Date: 03.04.2018

Product Name	CLEARKLENS CLEANSINALD SS VH9S
Product Code	100848254
Batch Number	FMP 18 183 / 45474
Production Date	02104/2018
Expiration Date	Exp. 02/01/2020

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	Liı	nits	Results
		Lower -	- Upper	
Appearance	Visual	_	ntly Yellow Juid	clean slightly Yellow Liquid
pH (neat solution)	DM001	9.0	11.4	10.7
Specific Gravity (20°C)	DM004	0.990	1.010	1.000

O. 1. 1. 16 . 6 D.	Name:	apa kinenna, ilavaluda
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

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	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:

05 Jul 2018

Date Tested:

11 Jul 2018

Date Test Completed:

25 Jul 2018

Purchase Order:

31329

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0043553/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

05/07/2018

9433

Test Article:

ClearKlens Cleansinald SS

Sample Code:

FMP18183 45474

Batch Ref: **Qty Received:**

20 x 900mL Bottles

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
Volume Tested	MM107/05	20 x 50 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

Comments

The testing procedure was performed in compliance with the current Pharmacopoeias however, the volume tested does not comply with the requirements of the current Pharmacopoeias.

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

CB6 M858

Date: 26 Jul 2018 13:24:16

Mrs C Moore

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0043553 Print Number: P0052319 Page 1 of 1





http://www.steris-ast.com

Certificate of Irradiation

Date issued: 02-Jun-2018

UK33S12110989-3-1

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:	1105380	
Customer Reference Number:	P031109	
Product Description:	C/SINALD SS 900ml Bottle DV4673 25-40kGy	
Validation Reference:	4673	
Quantity Received:	6	
Customer Minimum Specification kGy:	25.0	
Customer Maximum Specification kGy:	40.0	
Customer Unit Lot/Batch Number:	W045567, 1 PLT	
li li	rradiation Data	
Date and Time of Irradiation:	01-Jun-2018 23:42	
Reference Dose Range kGy:	32.3 - 32.6	
Calculated Minimum Dose kGy:	29.6	
Calculated Maximum Dose kGy:	35.6	

Items irradiated under WO45567, will be used in finished product batch FMP18183 45474

C. Pascoe

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

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