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CERTIFICATE OF ANALYSIS

Date: 19.06.2019

Product Name	CLEARKLENS CLEANSINALD SS VH9S		
Product Code			
Batch Number	TMP 19169 47378		
Production Date	18.06.2019		
Expiration Date	EXP 8/12/2020		

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	Clear Slightly Yellow Liquid		clear Slightly
pH (neat solution)	DM001	9.0	11.4	0.9
Specific Gravity (20°C)	DM004	0.990	1.010	0,998

On behalf of Diversey site Quality Manager	Position	Quality Control Inspector
	Name:	Staron Product

This document being issued electronically does not bear a signature

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

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-3030A

Product CodeLot NumberQuantityUOMC/SINALD SS 900ml Bottle DV4673WO473796Case

Validation Reference Number: 4673

Processing Run Start Date: 08-May-19 03:42 AM
Processing Run End Date: 08-May-19 09:30 AM

Specified Dose Range (kGy): 25.0 - 40.0 Calculated Min Dose (kGy): 29.1 Reference Dose Range (kGy): 27.3- 36.6 Calculated Max Dose (kGy): 35.1

Gamma Process Run Approval authorized by STERIS

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 WO47379 will be used in finished batch code Fmp19169 47378.

Olga Kirchner

Document ID: 20468

Last Revised in Rel 2.0.0.0

Rel Date: 08/13/2018

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mail@wickhamlabs.co.uk www.wickhamlabs.co.uk

Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received: 21 Jun 2019

Date Tested:

16 Jul 2019

Date Test Completed:

30 Aug 2019

Purchase Order:

33320

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0049322/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

21/06/2019

Test Article:

ClearKlens Cleansinald SS

Sample Code:

FMP19169 47378

Batch Ret:

10302

Qty Received:

20 x 900mL Bottles + 1 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
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	See external comments
Product Standard Data Sheet	FG047/psd	FM06-02

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.

BSG, CBIOL MRSB

Date: 09 Aug 2019 10:12:43

Mrs P. Pham-Lengoc

Documentation Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0049322 Print Number: P0061139 Page 1 of 1

