Diversey Europe Operations BV

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CERTIFICATE OF ANALYSIS

Date: 15.07.2015

Product Name	CLEARKLENS CLEANSINALD SS VH9S		
Product Code	100848254		
Batch Number	FMP 15196 38578		
Production Date	15/07/2015		
Expiration Date	15/01/2017		

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	<u> </u>
Appearance	Visual	Clear Slig	htly Yellow	Clear Slightly
		Liq	uid	Yellow Liquid
pH (100% 20°C)	JDM001	9.0	11.4	10.8
Specific Gravity (20°C)	JDM004	0.990	1.010	0.998

On hahalf of Diversors site	Name:	Mature Partierdia
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

COA 7516432	Version: 05	Date of issuing: April 27th 2012



http://www.synergyhealthplc.com

Certificate of Irradiation

Date Issued: 07-Jul-2015

UK32S11419229-1-1

This is to certify that Bradford Synergy Health PLC has where appropriate delivered an irradiation process in accordance with:

EN ISO 11137-1:2006 Sterilisation of Health Care Products EN ISO 9001:2008 Quality Management System EN ISO 13485:2012 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 100432 **Account Number:** 1002720 **Synergy Health Sales Part Reference: Customer Reference Number:** P024640 **Product Description:** C/SINALD SS 900ml BOTTLE 25-40kGy 4.1909 Validation Reference: 5 **Quantity Received:** 25.0 **Customer Minimum Specification kGy:** 40 D **Customer Maximum Specification kGy:** B/N W038573 THURS 2ND JULY 2015 1 PLT LOT 4 **Customer Unit Lot/Batch Number: Irradiation Data** 07-Jul-2015 13:48 Date and Time of Irradiation: 35.1 - 35.5 Reference Dose Range kGy: Calculated Minimum Dose kGy: 30.7 38.0 Calculated Maximum Dose kGy:

Items irradiated under WO38573 will be used in finished product batch code FMP15196 38578

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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA1 4XS Date Received:

17 Jul 2015

Date Tested:

21 Jul 2015

Date Test Completed:

04 Aug 2015

Purchase Order:

24753

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0023709/1

Test Required:

Sterility Test by Membrane Filtration (Steritest) Method I: Ph Eur and USP

Date Received: Test Article: 17/07/2015 Cleansinald SS

Sample Code:

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Batch Ref: Qty Received: FMP15196 38578 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
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.

March

Date: 05 Aug 2015 14:30:54

Mrs C Moore

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0023709 Print Number: P0028806 Page 1 of 1

