Diversey Europe Operations BV

Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date: 29.02.2016

Product Name	CLEARKLENS DE VH29
Product Code	100862174
Batch Number	FMP 16040 39800
Production Date	09/02/2016
Expiration Date	EXP 09/02/2018

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	Lii	mits	Results
		Lower -	Upper	
Appearance	Visual	Clear C	Colourless	Clear Colour-
		Lie	quid	less Liquid
Specific Gravity (20°C)	JDM004	0.880	0.888	0.887
	<u> </u>			

On habelf of Diverse site	Name:	Partyisla, Ulling	
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector	

This	document	being	issued	electronical	lv does	not	bear a	signature

COA / 313/03 Version: 04 Date of issuing: April 2/ 2012	COA 7515783	Version: 04	Date of issuing: April 27th 2012
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http://www.synergyhealthplc.com

Certificate of Irradiation

Date Issued: 22-Feb-2016

UK32S11567235-1-1

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

EN ISO 11137-1:2015 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 **Account Number:** 100432 **Synergy Health Sales Part Reference:** 1002733 **Customer Reference Number:** P026002 **Product Description:** JD DE 900ML BOTTLES 25-45kGy Validation Reference: 4.1756 **Quantity Received:** 704 Customer Minimum Specification kGy: 25.0 **Customer Maximum Specification kGy:** 45.0 **Customer Unit Lot/Batch Number:** 700 X BATCH NO: FMP16040 39800 + 4 SAMPLES THURS 18TH FEB 2016 13 PLTS LOT 3 irradiation Data Date and Time of Irradiation: 22-Feb-2016 00:38 Reference Dose Range kGy: 36.3 - 37.3 Calculated Minimum Dose kGy: 27.2 Calculated Maximum Dose kGy: 40.4

Irradiation Release Authorised By Synergy Health plc



Wickham Laboratories Ltd Hoeford Point, Barwell Lane, Gosport Hampshire PO13 0AU England

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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:

29 Feb 2016

Date Tested:

01 Mar 2016

Date Test Completed:

01 Mar 2016

Purchase Order:

PO26114

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0027555/2

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL

Date Received: Test Article:

29/02/2016

Clearklens DE

Sample Code:

FMP16040 39800

Batch Ref:

7345

Qty Received:

1 x 900mL Bottle

Test	Method Item	Result
Bacterial Endotoxin by Kinetic QCL Test	MM110/00	<0.25 EU/mL
Spiked Recovery	MM110/01	121 %
Dilution Tested	MM110/04	1/50
Endotoxin Spike Level	MM110/02	0.5 EU/mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 8.0 2.6.14, USP 38 <85> & JPXVI 4.01
Product Standard Data Sheet	FG047/psd	FM01

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

Page 1 of 1

Date: 02 Mar 2016 15:06:53

Mrs P. Pham-Lengoc

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0027555 Print Number: P0033231





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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

29 Feb 2016

Date Tested:

04 Mar 2016

Date Test Completed:

18 Mar 2016

Purchase Order:

PO26114

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0027555/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

29/02/2016

Test Article:

Clearklens DE FMP16040 39800

Sample Code: Batch Ref:

7345

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	See external comments
Product Standard Data Sheet	FG047/psd	FM03

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.

ARTHOR

Date: 18 Mar 2016 15:45:49

Mrs C Moore

Page 1 of 1

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0027555 Print Number: P0033659

