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CERTIFICATE OF ANALYSIS

Date: 20/09/2018

Product Name	CLEARKLENS DE VH29
Product Code	100862174
Batch Number	FMP 18245 45788
Production Date	0210912018
Expiration Date	EXP 02/09/12020

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	Lir	nits	Results
		Lower ·	 Upper 	
A	177 mu a 1	Clear Co	olourless	Clear Colour-
Appearance	Visual	Lic	uid	less Liguria
Specific Gravity (20°C)	DM004	0.880	0.888	0.884

0 1 1 16 CD' '4	Name:	Partyusto, Marchada
On behalf of Diversey site Quality Manager	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



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 18-Sep-2018

UK33S12166360-1-1

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 100432 **Account Number:** Synergy Health Sales Part Reference: 1106483 **Customer Reference Number:** P031711 **Product Description: DIVER DE BLK2 BOTTLE DV4648 25-45kGy** Validation Reference: 4648 541 **Ouantity Received: Customer Minimum Specification kGy:** 25.0 **Customer Maximum Specification kGy:** 45.0 FMP18245 45788, 10 PLTS **Customer Unit Lot/Batch Number: Irradiation Data** 17-Sep-2018 08:50 Date and Time of Irradiation: Reference Dose Range kGy: 36.3 - 36.8 28.7 Calculated Minimum Dose kGy: 41.6 Calculated Maximum Dose kGy:

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



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 09-Sep-2018

UK33S12162973-1-2

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

Oi	Order Information			
Account Number: Synergy Health Sales Part Reference: Customer Reference Number: Product Description: Validation Reference: Quantity Received: Customer Minlmum Specification kGy:	100432 1106483 P031693 DIVER DE BLK2 BOTTLE DV4648 25-45kGy 4648 167 25.0			
	==:			
Customer Unit Lot/Batch Number:	FMP18245 45788, INC 5 SMPLS, 3 PLTS			
lı	rradiation Data			
Date and Time of Irradiation:	09-Sep-2018 15:55			
Reference Dose Range kGy:	35.2 - 36.2			
Calculated Minimum Dose kGy:	27.8			
Calculated Maximum Dose kGy:	41.0			

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



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

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mail@wickhamlabs.co.uk

Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS Date Received:

21 Sep 2018

Date Tested:

27 Sep 2018

Date Test Completed:

11 Oct 2018

Purchase Order:

PO31766

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0044872/3

Test Required:

Sterility by Membrane Filtration Steritest

Date Received: Test Article: 21/09/2018

Clearklens DE FMP18245 45788

Sample Code: Batch Ref:

9568

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	FM03-01

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.

BSc. CBiol. MRSB

Date: 12 Oct 2018 12:43:41

Mrs P. Pham-Lengoc

Documentation Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0044872 Print Number: P0054208 Page 1 of 1





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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

21 Sep 2018

Date Tested:

24 Sep 2018

Date Test Completed:

24 Sep 2018

Purchase Order:

PO31766

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0044872/4

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL

Date Received: Test Article:

21/09/2018

Clearkiens DE FMP18245 45788

Sample Code: **Batch Ref:**

9568

Qty Received:

1 x 900mL Bottles

Test	Method Item	Result
Bacterial Endotoxin by Kinetic QCL Test	MM110/00	<0.25 EU/mL
Spiked Recovery	MM110/01	73 %
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 9.3 2.6.14, USP 41 <85> & JPXVII 4.01
Product Standard Data Sheet	FG047/psd	FM02-01

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

BSc. CBiol, MRSB

Date: 25 Sep 2018 10:43:38

Mrs P. Pham-Lengoc

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0044872 Print Number: P0053806

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