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CERTIFICATE OF ANALYSIS

Date: 13/12/2018

Product Name	CLEARKLENS DE VH29		
Product Code			
Batch Number	TNP 18266, 45966		
Production Date	22 109 1 2018		
Expiration Date	Exe 22 L03 L2020		

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	Lower - Upper		Results Clear Colonies	
Appearance	Visual				
Specific Gravity (20°C)	DM004	0.880	0.888	0.	888

On behalf of Diversey site Quality Manager	Name :	T. Stavon Movalariota
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

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



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 09-Oct-2018

UK33S12178073-2-1

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: 1106483 **Customer Reference Number:** P031851 **Product Description: DIVER DE BLK2 BOTTLE DV4648 25-45kgy Validation Reference:** 4648 **Quantity Received:** 160 Customer Minimum Specification kGy: 25.0 Customer Maximum Specification kGv: 45.0 Customer Unit Lot/Batch Number: FMP18265 45966, 3 PLTS **Irradiation Data** Date and Time of tradiation: 09-Oct-2018 01:03 Reference Dose Range kGy: 35.9 - 37.0 Calculated Minimum Dose kGy: 28.4 Cajculated Maximum Dose kGy: 41.9

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



http://www.sterls-ast.com

Certificate of Irradiation

Date Issued: 02-Oct-2018

UK33S12172913-3-1

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: 1106483 **Customer Reference Number:** P031825 **Product Description: DIVER DE BLK2 BOTTLE DV4648 25-45kGy** Validation Reference: 4648 **Quantity Received:** K4K Customer Minimum Specification kGy: 25.0 **Customer Maximum Specification kGv:** 45.0 **Customer Unit Lot/Batch Number:** FMP18265 45966, INC 5 SMPLS, 10 PLTS irradiation Data Date and Time of Irradiation: 02-Oct-2018 03:17 Reference Dose Range kGy: 34.4 - 34.5 Calculated Minimum Dose kGv: 27.2 Calculated Maximum Dose kGy: 39.0

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



Wickham Laboratories Ltd Hoeford Point, Barwell Lane, Gosport Hampshire PO13 0AU England Telephone: +44(0)1329 226600

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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 LA1 4XS Date Received: 14 Dec 2018

Date Tested: 04 Jan 2019

Date Test Completed: 18 Jan 2019

Purchase Order: 32300

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0046349/1

Test Required:

Sterllity by Membrane Filtration Steritest

Date Received:

14/12/2018

Test Article: Sample Code: Clearklens DE FMP18265 45966

Batch Ref: 9689

Qty Received:

20 x 900mL Bottles + 3 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	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.

Minne

CBOL MRSB

Date: 21 Jan 2019 15:32:51

Mrs C Moore

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0046349 Print Number: P0056514

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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA1 4XS Date Received: 14

14 Dec 2018

Date Tested:

18 Dec 2018

Date Test Completed:

18 Dec 2018

Purchase Order:

32300

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

r: 0046349/2

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL

Date Received:

14/12/2018

Test Article: Sample Code:

Clearklens DE FMP18265 45966

Batch Ref:

9689

Qty Received:

1 x 900mL Bottle

Test	Method item	Result	Specification
Bacterial Endotoxin by Kinetic QCI Test	MM110/00	<0.25 EU/mL	
Spiked Recovery	MM110/01	73 %	50-200
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	FM01-02	

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

Lasson

Date: 19 Dec 2018 14:56:31

Victoria Watson

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Laboratory Manager – Microbiology

Certificate of Analysis - O\$MM Consignment: 0046349 Print Number: P0055889

