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CERTIFICATE OF ANALYSIS

Date: 12.06.2019

Product Name	CLEARKLENS DE VH29	
Product Code	7515783	
Batch Number	TMP19122 41006	
Production Date	02.05.2019	
Expiration Date	EXP OF OR	

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 Lower - Upper		Results
Appearance	Visual	Clear Col		clear colouries
Specific Gravity (20°C)	DM004	0.880	0.888	0.885

	Name:	J. Slavon, Pourtyus
On behalf of Diversey site Ouality Manager	Position	Quality Control Inspector

This document being issued electronically does not bear a signature		Date of issuing: November 24 th 2017
COA Template Version : 02	Date of issuing: November 24 201	
	46131011 1 0E	

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-3392A

UOM Quantity **Lot Number Product Code**

Case 705 FMP19122 47006 DIVER DE BLK2 BOTTLE DV4648

Validation Reference Number: 4648

Case FMP19122 47006, INC 5 SMPLS DIVER DE BLK2 BOTTLE DV4648

Validation Reference Number: 4648

Processing Run Start Date: 05-Jun-19 04:00 AM Processing Run End Date: 05-Jun-19 02:48 PM

Calculated Min Dose (kGy): 26.5 25.0 - 45.0 Specified Dose Range (kGy):

40.6 Calculated Max Dose (kGy): 31.6-39.8 Reference Dose Range (kGy):

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

Document ID: 22327

Rel Date: 08/13/2018 Last Revised in Rel 2.0.0.0

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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA1 4XS

Date Received:

13 Jun 2019

Date Tested:

14 Jun 2019

Date Test Completed:

14 Jun 2019

Purchase Order:

33269

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0049167/1

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL

Date Received:

13/06/2019

Test Article:

Clearklens DE FMP19122 47006

Sample Code: Batch Ref:

10229

Qtv 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	98 %
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 42 <85> & JPXVII 4.01
Product Standard Data Sheet	FG047/psd	FM01-02

Comments

Corrected Certificate of Analysis to LR0049167/1 authorised 18/06/19 due to batch error on paperwork supplied by client.

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

Date: 07 Aug 2019 16:24:34

Victoria Watson

Laboratory Manager – Microbiology

Certificate of Analysis - OSMM Consignment: 0049167 Print Number: P0061103 Page 1 of 1





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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

13 Jun 2019 Date Received:

12 Jul 2019 Date Tested:

26 Jul 2019 Date Test Completed:

33269 **Purchase Order:**

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0049167/2

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

13/06/2019

Test Article: Sample Code: Clearklens DE FMP19122 47006

Batch Ref:

10229

Qty Received:

20 x 900mL Bottle + 2 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	MM107/02	No growth in one broth
at 30-35°C after 14 days Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP	MQ005/07	See external comments
& JP Product Standard Data Sheet	FG047/psd	FM03-02

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.

Certificate of Analysis for LR0049167/2, authorised 06/08/19 due incorrect batch reference provided by client.

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

Victoria Watson Laboratory Manager – Microbiology

Certificate of Analysis - OSMM Consignment: 0049167 Print Number: P0061119 Page 1 of 1



Date: 08 Aug 2019 15:32:53