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CERTIFICATE OF ANALYSIS

Date: 15/02/2019

Product Name	CLEARKLENS DE VH29
Product Code	7515783
Batch Number	FMP19026 46760
Production Date	26/01/2019
Expiration Date	EXP 26/01/2021

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	
Appearance	Visual	Clear Colourless Liquid		Clear colourless liquid
Specific Gravity (20°C)	DM004	0.880	0.888	0.881

On behalf of Diversey site Quality Manager	Name :	J. Staver, P. J. J. J. J.
	Position	Quality Control Inspector

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

Contract Analytical Services

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Ms D Henderson
Flexible Medical Packaging Limited
Unit 8
Hightown
White Cross Industrial Estate
Lancaster
LA1 4XS

Date Received: 15 Feb 2019
Date Tested: 19 Feb 2019
Date Test Completed: 19 Feb 2019
Purchase Order: 32604

CERTIFICATE OF ANALYSIS

Laboratory Reference Number: 0047246/2
Test Required: Bacterial Endotoxin Testing by Kinetic QCL
Date Received: 15/02/2019
Test Article: Clearklens DE
Sample Code: FMP19026 46760
Batch Ref: 9937
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	97 %
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.

Date: 20 Feb 2019 12:20:35

Victoria Watson
Laboratory Manager – Microbiology



Wickham Laboratories

Contract Analytical Services

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Date Received: 15 Feb 2019
Date Tested: 12 Mar 2019
Date Test Completed: 26 Mar 2019
Purchase Order: 32604

CERTIFICATE OF ANALYSIS

Laboratory Reference Number: 0047246/1
Test Required: Sterility by Membrane Filtration Steritest
Date Received: 15/02/2019
Test Article: Clearlens DE
Sample Code: FMP19026 46760
Batch Ref: 9937
Qty Received: 20 x 900mL Bottles + 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 Thiolglycollate 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: 28 Mar 2019 08:18:38

Mrs P. Pham-Lengoc
Documentation Manager – Microbiology

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-1516A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
IPA BOTTLE DV4648	FMP19026 46760	166	Case
Validation Reference Number: 4648			

Processing Run Start Date: 09-Feb-19 05:51 PM

Processing Run End Date: 10-Feb-19 01:38 AM

Specified Dose Range (kGy):	25.0 - 45.0	Calculated Min Dose (kGy):	27.2
Reference Dose Range (kGy):	31.6- 39.8	Calculated Max Dose (kGy):	42.3

Other Information

FMP19026 46760, 3 PLTS

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

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-1445A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
DIVER DE BLK2 BOTTLE DV4648	FMP19026 46760	540	Case
Validation Reference Number: 4648			
DIVER DE BLK2 BOTTLE DV4648	SAMPLES	5	Case
Validation Reference Number: 4648			

Processing Run Start Date: 02-Feb-19 11:56 AM

Processing Run End Date: 02-Feb-19 09:27 PM

Specified Dose Range (kGy):	25.0 - 45.0	Calculated Min Dose (kGy):	26.8
Reference Dose Range (kGy):	31.6- 39.8	Calculated Max Dose (kGy):	39.6

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

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