

Diversey Europe BV

Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

Tel. +31 (0)30 247 6427

CERTIFICATE OF ANALYSIS

Date: 22.08.2015

Product Name	CLEARKLENS DE VH29		
Product Code	7515783		
Batch Number	FMP 15197 38577		
Production Date	16/07/2015		
Expiration Date	16/07/2017		

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	Lim Lower ~	its Upper	Results
Appearance	Visual	Clear Co	lourless	clear Colovile
Specific Gravity (20°C)	JDM004	0.880	0.888	0.883

On behalf of Diversey site	Name:	Tartyisla, Alcoha
Quality Manager	Position	Quality Control Inspector

This document being issued electronical	ly does not bear a signature	
COA 7515783	Version: 03	Date of issuing: November 28th 2011



http://www.synergyhealthplc.com

Certificate of Irradiation

Date Issued: 09-Aug-2015

UK32S11435608-1-1

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

EN ISO 11137-1:2006 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 100432 **Account Number: Synergy Health Sales Part Reference:** 1002733 P024822 **Customer Reference Number:** JD DE 900ML BOTTLES 25-45kGy **Product Description:** Validation Reference: 4.1756 701 **Quantity Received:** 25.0 **Customer Minimum Specification kGy: Customer Maximum Specification kGy:** 45.0 697 X B/N FMP15197 38577 + 4 SAMPLES TUES Customer Unit Lot/Batch Number: 28TH JULY 2015 13 PLTS LOT 1 **irradiation Data** Date and Time of Irradiation: 08-Aug-2015 22:39 37.7 - 40.5 Reference Dose Range kGy: 28.3 Calculated Minimum Dose kGy: Calculated Maximum Dose kGy: 43.9



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

Telephone: +44(0)1329 226600 Fax: +44(0)1329 226688

mail@wickhamlabs.co.uk www.wickhamlabs.co.uk

Ms D Henderson

Date Received: Flexible Medical Packaging Limited

18 Aug 2015

Unit 8 Hightown

20 Aug 2015 Date Tested:

White Cross Industrial Estate

Lancaster LA14XS

Date Test Completed:

20 Aug 2015

Purchase Order:

PO24935

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0024273/2

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL: Ph Eur and USP

Date Received:

18/08/2015

Test Article:

ClearKlens DE VH29

Sample Code:

6773

Batch Ref: Qty Received: FMP15197 38577 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	
Product Standard Data Sheet	FG047/psd	FM01	

Comments

Test carried out according to current Pharmacopoeias.

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

WYZ

Date: 21 Aug 2015 16:32:24

Mr K.A. Barker

Business Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0024273 Print Number: P0029170 Page 1 of 1





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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA1 4XS Date Received:

18 Aug 2015

Date Tested:

21 Aug 2015

Date Test Completed:

04 Sep 2015

Purchase Order:

PO24935

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0024273/1

Test Required:

Sterility Test by Membrane Filtration (Steritest) Method I: Ph Eur and USP

Date Received:

18/08/2015

Test Article:

ClearKlens DE

Sample Code:

100

Batch Ref: Qty Received: FMP15197 38577 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
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.

Mary

Date: 07 Sep 2015 15:33:36

Mrs C Moore

Laboratory Manager - Pharmaceutical Microbiology