

Diversey Europe BV

Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date: 11.11.13

Product Name	CLEARKLENS TEGO 2000SC VH25	
Product Code	7513387	
Batch Number	FNP13290 35050	
Production Date	10/2013	
Expiration Date	10/2015	

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	Lin Lower -	nits Upper	Results
Appearance	Visual			deat Slightly
pH 100% (20°C)	JDM001	7.5	8.5	7.7
Specific Gravity (20°C)	JDM004	0.993	1.003	0.998

On behalf of Diversey site Quality Manager	Name:	Makeis	O. Linchner
	Position	Quality	Control Inspector

This document being issued electronically does not bear a signature

COA 7513387	Version: 03	Date of issuing: November 28th 2011



http://www.synergyhealthplc.com

Certificate of Irradiation

Date Issued: 07-Nov-2013 UK32S11069063-1-1

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 Account Number: 100432 Synergy Health Sales Part Reference: 1002722 **Customer Reference Number:** P021240 **Product Description:** DIVERSEY TEGO 2000SC 50ml 25-42kGy Validation Reference: 4.0716 **Quantity Received:** 105 Customer Minimum Specification kGy: 25.0 Customer Maximum Specification kGy: 42.0 Customer Unit Lot/Batch Number: 103 X B/No.FMP13290 35050 + 2 SAMPLES . 2 PLTS LOT 6 THURS 31.10.13 **Irradiation Data** Date and Time of Irradiation: 06-Nov-2013 19:44 Reference Dose Range kGy: 34.3 - 35.6 Calculated Minimum Dose kGy: 28.5 Calculated Maximum Dose kGy: 39.1



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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA1 4XS Date Received:

11 Nov 2013

Date Tested:

18 Nov 2013

Date Test Completed:

10 Dec 2013

Purchase Order:

21291

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

Test Required:

0013283/2

Sterility Test by Membrane Filtration (Steritest) Method I with Subculture:

PH Eur and USP

Date Received:

11/11/2013

Test Article: Sample Code: TEGO 2000sc Block 1 FMP 13290 35050

Sample Code:

452581, Exp. 2015/10

Qty Received:

20 x 50mL 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 two broths
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in two broths
Growth of Sub-culture in Tryptone Soya Broth at 30-35°C after 4 days	MM107/07	No growth in two broths
Growth of Sub-culture in Fluid hioglycollate at 20-25°C after 4 days	MM107/08	No growth in two broths
/olume Tested	MM107/05	20 x 50 mL
Product Standard Data Sheet	FG047/psd	FM08

Comments

As Steritest Sterility Test

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

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Date: 12 Dec 2013 12:07:59

Mrs C Moore

ISO 17025

Laboratory Manager - Pharmaceutical Microbiology

Consignment: 0013283

GLP GMP FDA

Certificate of Analysis - OSMM

Print Number: P0016007

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