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CERTIFICATE OF ANALYSIS

Date: 17 01 2019

Product Name	CLEARKLENS TEGO 2000SC VH25
Product Code	100961418
Batch Number	EM018344 46751
Production Date	10 12 120 18
Expiration Date	EXP (0)1212021

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 Clear Slightly Yellow Liquid		Results clear Stiphfly Yellow Liquis
Appearance	Visual			
Specific Gravity (20°C)	DM004	0.993	1.003	0.998
pH (neat solution)	DM001	7.5	8.5	8.0

Quality Manager	Position	Quality Control Inspector
On behalf of Diversey site	Name:	Navaburda Partyest

This document being issued electronically does not bear a signature

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



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Ms D Henderson

Flexible Medical Packaging Limited

Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

18 Jan 2019

Date Tested:

18 Feb 2019

Date Test Completed:

11 Mar 2019

Purchase Order:

PO32429

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0046783/2

Test Required:

Sterility by Membrane Filtration Steritest

Date Received: **Test Article:** Sample Code:

18/01/2019 Tego 2000SC

FMP18344 46751

621190

Qty Received:

Batch Ref:

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 TSB at 20-25°C after 4 - 7 days	MM107/07	No growth in two broths
Growth of Sub-culture in THY at 30-35°C after 4 - 7 days	MM107/08	No growth in two broths
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.0 2.6.1, USP 41 <71> & JP XVII 4.06
Product Standard Data Sheet	FG047/psd	FM08-02

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

BSc. CBlot MRSB

Date: 19 Mar 2019 10:05:15

Mrs P. Pham-Lengoc

Documentation Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0046783 Print Number: P0057960







STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-1048A

Product Code Lot Number Quantity MOU FMP18344 46751 180 Case

TEGO 2000SC DV4724

TEGO 2000SC DV4724 FMP18344 46751 INC 2 SAMPLE 2 Case

4724

Validation Reference Number: 4724

PO Number: PO32313

Processing Run Start Date: 01-Jan-19 09:56 PM Processing Run End Date: 02-Jan-19 05:46 AM

Validation Reference Number:

Specified Dose Range (kGy): 25.0 - 45.0 Calculated Min Dose (kGy): 27.5 Reference Dose Range (kGy): 29.4-40.5 Calculated Max Dose (kGv): 36.3

Signature Manifest

Reviewed and E-Signed By: Mark Morse (QA Officer)

Date/Time E-Signed: 2019-01-02 12:25 PM

Document Content Revision: 1

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 NN118RB**

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Document ID: 10748

Last Revised in Rel 2.0.0.0 Rel Date: 08/13/2018 Page 1 of 1