

Diversey Europe Operations BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

Tel. +31 (0)30 247 6427 www.Diversey.com

CERTIFICATE OF ANALYSIS

Date: [9 | 11 | 2019

Product Name	CLEARKLENS TEGO 2000SC VH25		
Product Code			
Batch Number	EMB 10303 /10101		
Production Date	30/10/2019		
Expiration Date	EVP 3010 13033		

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
Appearance	Visual			Clear Stigut
Specific Gravity (20°C)	DM004	0.993	1.003	1.001
pH (neat solution)	DM001	7.5	8.5	80

On behalf of Diversey site Quality Manager	Position	Quality Control Inspector	
	Name:	Party 86	lavabuola

This document being issued electronically does not bear a signature

COA Town I. A.			
COA Template	Version: 02		
		Date of issuing: November 24 2017	
		Date of issuiria , November 24 2017	

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-6119A

Product Code Lot Number Quantity **UOM TEGO 2000SC DV4724** FMP19303 48491, INC 1 SAMPLE 100 Case

Validation Reference Number: 4724

Processing Run Start Date: 11-Nov-19 01:21 AM Processing Run End Date: 11-Nov-19 08:59 AM

Specified Dose Range (kGy): 25.0 - 45.0 Calculated Min Dose (kGy): 30.3 40.3 Reference Dose Range (kGy): 29.4-40.5 Calculated Max Dose (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: 32902

Rel Date: 08/13/2018

Last Revised in Rel 2.0.0.0

Page 1 of 1

PHARMACEUTICAL ANALYSIS REPORT



Flexible Medical Packaging

Unit 8 Hightown White Cross Industrial Estate Lancaster LA14XS

FAO:

Mrs.Olga Kirchner

Report of Tests on:

Tego 2000 SC

Sample Description:

Sample Code: Fmp19303 48491

Lucideon Sample Number: (196384)-48692

Lucideon Report Number: (196384)-48692/MFEP

Issue Number:

1

Date Logged:

22-Nov-2019

Order Number:

PO34324

Date Reported:

16-Dec-2019

Date(s) of Test(s):

27-Nov-2019 to 15-Dec-2019

Sterility Testing Membrane Filtration EP

Test Results:

Result: Pass

The test results meet the EP/USP criteria: Yes

Tests carried out in accordance with cGMP

End of Test Report

MOBOOK 10-DEC-19

Mrs Natalie Boot

of the results or their implications.

Senior Business Support Assistant

This report is issued in accordance with the Conditions of Business of Lucideon Limited and relates only to the sample(s) tested. No responsibility is taken for the accuracy of the sampling unless this is done under our own supervision. This report shall not be reproduced in part without the written approval of Lucideon Limited, nor used in any way as to lead to misrepresentation