

Diversey Europe Operations BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date: 09 January 2024

Product Name	CLEA	CLEARKLENS TEGO 2000SS VH25S		
Product Code	7516427			
Batch Number	FMP23339	59140		
Production Date	05/12/2023			
Expiration Date	EXP 05/06/2025			

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	Limi Lower	ts - Upper	Results
Appearance	Visual	Clear Slightly	Yellow Liquid	Clear Slightly Yellow Liquid
pH (neat solution)	DM001	6.2	8.2	7.8
Specific Gravity (20°C)	DM004	0.980	1.020	0.996

On behalf of Diversey site	Name:	Edyta Rodrígues Ewelína Nowaczyk
Quality Manger	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

COA Template Version : 04 Date of issuing : March 11th 2019			
	COA Template	Version : 04	Date of issuing : March 11th 2019

STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID 2173-43075A

Product Code	Lot Number		<u>Quantity</u>	<u>UOM</u>
TEGO 2000SS BLK1 DV4767	59140 - STR0	01783	122	Case
Validation Reference Number:	4767			
TEGO 2000SS BLK1 DV4767	59140 - STR0	01783, 3 X SAMPLES	3	Case
Validation Reference Number:	4767			
Processing Run Start Date: 29-Dec-2023 5:26 AM				
Processing Run End Date: 29-Dec-2023 2:36 PM				
Specified Dose Range (kGy):	25.0 - 45.0	Calculated Min Dose	(kGy):	29.1
Reference Dose Range (kGy):	31.5 - 39.7	Calculated Max Dose	e (kGy):	42.9

PO Number: 407079

Product meets Customer specifications; zero nonconformities occurred during this irradiation run.

ſ	Signature Manifest			
I	Reviewed and E-Signed By: Sonal Sale (Quality Engineer)			
l	Date/Time E-Signed: 02-Jan-2024 9:33 AM			
I	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 NN11 8RB Phone: + 44(0) 1327 706 111

Document ID: 163763

PHARMACEUTICAL ANALYSIS REPORT



Flexible Medical Packaging Unit 8 Hightown White Cross Industrial Estate Lancaster LA1 4XS

Mrs. Olga Kirchner FAO:

Report of Tests on: Tego 20	000 SS			
Sample Description: Sample	Code: Fmp23339 5	59140		
Lucideon Sample Number: UK2494	-665			
Lucideon Report Number: UK2494	-665/MFEP	Issue Number:	1	
Date Logged: 10-Jan-	2024	Order Number:	407216	
Date Reported: 09-Feb-	-	Date(s) of Test(s):	25-Jan-2024 to	08-Feb-2024
		ility Testing		
	Membra	ine 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

DocuSigned by:

Tracy Wheeler

Signer Name: Tracy Wheeler Signing Reason: I have reviewed this document Signing Time: 09 February 2024 | 13:50 GMT

D85AB0246E184F31821E2EC1AFA6CA90

Tracy Wheeler

Business Support Administrator

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