

Diversey Europe Operations BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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

## **CERTIFICATE OF ANALYSIS**

Date:

19 December 2022

Product Name	CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26 first off 7514852		
roduct Code			
Batch Number	FMP22353	55552	
Production Date	19/12/2022		
Expiration Date	EXP 19/12/2024		

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		Results
Appearance	Visual	Clear Colou	ırless Liquid	Clear Colourless Liquid
pH (neat solution)	DM001	9.0	12.5	11.43
Specific Gravity (20°C)	DM004	1.004	1.020	1.006
Potential Chlorine Dioxide		100	180	135

On behalf of Diversey site Quality Manger	Name:	Edyta Rodrígues Justyna Staron
	Position	Quality Control Inspector

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



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## **CERTIFICATE OF ANALYSIS**

Date:

19 December 2022

Product Name	CLEARKLENS BI-SPORE BASE SOLUTION VH26 first off 7514852		
Product Code			
Batch Number	FMP22353	55552	
Production Date	19/12/2022		
Expiration Date	EXP 19/12/2024		

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		Results
Appearance	Visual	Clear Color	ırless Liquid	Clear Colourless Liquid
pH (neat solution)	DM001	1.5	2.5	1.9
Specific Gravity (20°C)	DM004	1.010	1.030	1.019

On behalf of Diversey site  Quality Manger	Name:	Edyta Rodrígues Justyna Staron
	Position	Quality Control Inspector

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

## STERIS: Gamma Certificate Of Processing

Prepared for:

**FLEXIBLE MEDICAL PACKAGING LTD (8245)** 

Gamma Process Run ID

2173-33392A

**Product Code** 

Lot Number

Quantity UOM

**BI-SP B3 Components DV5857** 

55636

Case

Validation Reference Number:

5857

Processing Run Start Date: 10-Dec-2022 3:12 PM

Processing Run End Date: 10-Dec-2022 9:14 PM

Specified Dose Range (kGy):

25.0 - 40.0

Calculated Min Dose (kGy):

28.7

Reference Dose Range (kGy):

26.1 - 35.8

Calculated Max Dose (kGy):

34.3

PO Number: 404108

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

### Gamma Process Run Approval authorized by STERIS

Date/Time E-Signed: 11-Dec-2022 7:11 AM

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

Items irradiated under WO55636 will be used in finished batch Fmp22353 55552

Olga Kirchner

Document ID: 130213

# PHARMACEUTICAL ANALYSIS REPORT



Flexible Medical Packaging

Unit 8 Hightown White Cross Industrial Estate Lancaster LA14XS

FAO:

Mrs.Olga Kirchner

Report of Tests on:

Bi-Spore Activator

Sample Description:

Sample Code: Fmp22353 55552

Lucideon Sample Number: UK2367-630

Lucideon Report Number: UK2367-630/MFEP

Issue Number:

1

Date Logged:

06-Jan-2023

Order Number:

404243

**Date Reported:** 

26-Jan-2023

Date(s) of Test(s):

11-Jan-2023 to 25-Jan-2023

**Sterility Testing** 

Membrane Filtration EP

Test Results:

The test results meet the EP/USP criteria: Yes

Result: Pass

Tests carried out in accordance with cGMP

**End of Test Report** 

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Mrs Natalie Boot

Senior Business Support Administrator

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## PHARMACEUTICAL ANALYSIS REPORT



Flexible Medical Packaging

Unit 8 Hightown White Cross Industrial Estate Lancaster LA14XS

FAO:

Mrs.Olga Kirchner

Report of Tests on:

Bi-Spore Base

Sample Description:

Sample Code: Fmp22353 55552

Lucideon Sample Number: UK2367-631

Lucideon Report Number: UK2367-631/MFEP

Issue Number:

Date Logged:

06-Jan-2023

Order Number:

404243

**Date Reported:** 

26-Jan-2023

Date(s) of Test(s):

11-Jan-2023 to 25-Jan-2023

**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** 



Mrs Natalie Boot

Senior Business Support Administrator