Diversey Europe Operations BV
Maarssenbroeksedijk 2
3542 DN Utrecht
The Netherlands

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

CERTIFICATE OF ANALYSIS

Date: 12/10/2020

Product Name	**CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26**
Product Code	75 15 828
Batch Number	FMP 20279 50320
Production Date	05/10/2020
Expiration Date	EXP 05/10/2022

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	Limits Upper	Results
Appearance	Visual	Clear Colourless Liquid		Clear Colourless Liquid
pH (neat solution)	DM001	9.0	12.5	11.5
Specific Gravity (20°C)	DM004	1.004	1.020	1.005
Potential Chlorine Dioxide (ppm)		100	180	135

On behalf of Diversey site Quality Manager	Name :	Nawabuda Partyusho
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

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

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CERTIFICATE OF ANALYSIS

Date: 12/10/2020

Product Name	**CLEARKLENS BI-SPORE BASE SOLUTION VH26**
Product Code	7515828
Batch Number	FMP20279 50320
Production Date	05/10/2020
Expiration Date	EXP 05/10/2022

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	Limits Upper	Results
Appearance	Visual	Clear Colourless Liquid		Clear Colourless Liquid
pH (neat solution)	DM001	1.5	2.5	2.0
Specific Gravity (20°C)	DM004	1.010	1.030	1.016

On behalf of Diversey site Quality Manager	Name :	Navaneeda Pantyish
	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: 2173-13313A

Product Code	Lot Number	Quantity	UOM
BI-SP B3 COMPONENTS DV4725	WO50319	8	Case

Validation Reference Number: 4725

Processing Run Start Date: 07-Sep-20 12:47 AM

Processing Run End Date: 07-Sep-20 06:52 AM

Specified Dose Range (kGy):	**25.0 - 40.0**	**Calculated Min Dose (kGy):**	**27.5**
Reference Dose Range (kGy):	**30.2- 39.2**	**Calculated Max Dose (kGy):**	**35.3**

PO Number: 36309

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

Items Irradiated under WO50319 will be used in finished batch Fmp20279 50320.

Olga Kirchner

PHARMACEUTICAL ANALYSIS REPORT

LUCIDEON
insight creating advantage

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

FAO: Mrs.Olga Kirchner

Report of Tests on:	Bi-Spore Activator		
Sample Description:	Sample Code: Fmp20279 50320		
Lucideon Sample Number:	UK204167-29010		
Lucideon Report Number:	UK204167-29010/MFEP	**Issue Number:**	1
Date Logged:	14-Oct-2020	**Order Number:**	PO 36615
Date Reported:	09-Nov-2020	**Date(s) of Test(s):**	23-Oct-2020 **to** 06-Nov-2020

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

09NOV20

Mrs Andrea Saunders
Business Support Assistant

Lucideon Limited
Queens Road, Penkhull
Stoke-on-Trent
Staffordshire ST4 7LQ

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PHARMACEUTICAL ANALYSIS REPORT

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Hightown
White Cross Industrial Estate
Lancaster
LA1 4XS

FAO: Mrs.Olga Kirchner

Report of Tests on:	Bi-Spore Base		
Sample Description:	Sample Code: Fmp20279 50320		
Lucideon Sample Number:	UK204167-29011		
Lucideon Report Number:	UK204167-29011/MFEP	**Issue Number:**	1
Date Logged:	14-Oct-2020	**Order Number:**	PO 36615
Date Reported:	09-Nov-2020	**Date(s) of Test(s):**	23-Oct-2020 **to** 06-Nov-2020

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

09 NOV 20

Mrs Andrea Saunders
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