

FIRST OFF



Diversey Europe Operations BV
Maarssebroeksedijk 2
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The Netherlands

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www.Diversey.com

CERTIFICATE OF ANALYSIS

Date: 02/07/2020

Product Name	CLEARLENs BI-SPORE ACTIVATOR SOLUTION VH26
Product Code	7515828
Batch Number	FMP 20184, 49734
Production Date	02/07/2020
Expiration Date	Exp 02/07/2022

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		Results
		Lower	Upper	
Appearance	Visual	Clear Colourless Liquid		Clear Colourless Liquid
pH (neat solution)	DM001	9.0	12.5	11.4
Specific Gravity (20°C)	DM004	1.004	1.020	1.005
Potential Chlorine Dioxide (ppm)		100	180	131.625 ppm

On behalf of Diversey site Quality Manager	Name :	A. Bays, Ystou
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

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

Date: 02.07.2020

Product Name	CLEARLENs BI-SPORE BASE SOLUTION VH26
Product Code	7515828
Batch Number	FMP 20184, 49734
Production Date	02.07.2020
Expiration Date	EXP 02/07/2022

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		Results
		Lower	Upper	
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 :	J. Stanon, Uvabanda
	Position	Quality Control Inspector

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STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID: 2173-11116A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
BI-SP B3 COMPONENTS DV4725	WO49733	8	Case

Validation Reference Number: 4725

Processing Run Start Date: 08-Jun-20 12:10 PM

Processing Run End Date: 08-Jun-20 07:15 PM

Specified Dose Range (kGy):	25.0 - 40.0	Calculated Min Dose (kGy):	27.2
Reference Dose Range (kGy):	30.2- 39.2	Calculated Max Dose (kGy):	34.3

PO Number: PO35587

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 WO49733 will be used in finished batch Fmp20184 49734.

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: Fmp20184 49734

Lucideon Sample Number: UK202766-19921

Lucideon Report Number: UK202766-19921/MFEP **Issue Number:** 1

Date Logged: 08-Jul-2020 **Order Number:** PO 35872

Date Reported: 29-Jul-2020 **Date(s) of Test(s):** 09-Jul-2020 to 23-Jul-2020

Sterility Testing

Membrane Filtration EP

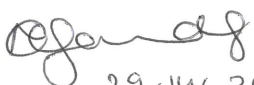
Test Results:

Result: Pass

The test results meet the EP/USP criteria: Yes

Tests carried out in accordance with cGMP

End of Test Report


29 JUL 20

Mrs Andrea Saunders
Business Support Assistant

Page 1 of 1

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Lucideon is the trading name of Lucideon Limited. Registered in England No. 1960455.

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enquiries@lucideon.com
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PHARMACEUTICAL ANALYSIS REPORT

Flexible Medical Packaging

Unit 8
Hightown
White Cross Industrial Estate
Lancaster
LA1 4XS

FAO: Mrs.Olga Kirchner

Report of Tests on: Bi-Spore Base

Sample Description: Sample Code: Fmp20184 49734

Lucideon Sample Number: UK202766-19922

Lucideon Report Number: UK202766-19922/MFEP **Issue Number:** 1

Date Logged: 08-Jul-2020 **Order Number:** PO 35872

Date Reported: 29-Jul-2020 **Date(s) of Test(s):** 09-Jul-2020 to 23-Jul-2020

Sterility Testing

Membrane Filtration EP


Test Results:

Result: Pass

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