

Diversey Europe BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date: 07/06/2021

Product Name	CLEARKLENS BI-SPORE BASE SOLUTION VH26
Product Code	7514852
Batch Number	FMP21147; 51718
Production Date	27/05/2021
Expiration Date	EXP 27/05/2023

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	Lir	Results	
		Lower -	- Upper	
Appearance	Visual	Clear Colou	rless Liquid	Clear Colourless Liquid
pH (neat solution)	DM001	1.5	2.5	1.9
Specific Gravity (20°C)	DM004	1.010	1.030	1.016

On helelf of Discourse 14	Name:	U.Haraburda A.Borys
On behalf of Diversey site Quality Manager	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: 27/05/2021

Product Name	CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26
Product Code	7514852
Batch Number	FMP21147; 51718
Production Date	27/05/2021
Expiration Date	EXP 27/05/2023

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	-	nits - Upper	Results
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	1,008
Potential Chlorine Dioxide (ppm)		100	180	155.25

On hehelf of Di	Name:	U.Haraburda A.Borys
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

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	The state of the s	
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: 2173-18772A

Product CodeLot NumberQuantityUOMBI-SP B3 COMPONENTS DV4725WO517198 Case

Validation Reference Number: 4725

Processing Run Start Date: 28-Apr-21 09:49 PM

Processing Run End Date: 29-Apr-21 03:37 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): 34.2

PO Number: 37882

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

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 WO51719 will be used in finished batch Fmp21147 51718.

Olga Kirchner

Rel Date: 08/13/2018

Document ID: 77409

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:

Bi-Spore Activator

Sample Description:

Sample Code: Fmp21147 51718

Lucideon Sample Number: UK212674-18306

Lucideon Report Number: UK212674-18306/MFEP

Issue Number:

Date Logged:

21-Jun-2021

Order Number:

PO 400105

Date Reported:

09-Jul-2021

Date(s) of Test(s):

23-Jun-2021 to 07-Jul-2021

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

NOBOOK 09-JUI-21

Mrs Natalie Boot

Business Support Administrator

This report is issued in accordance with the Conditions of Business of Lucideon Limited and relates only to the sample(s) tested.

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: Fmp21147 51718

Lucideon Sample Number: UK212674-18307

Lucideon Report Number: UK212674-18307/MFEP

Issue Number:

Date Logged:

21-Jun-2021

Order Number:

PO 400105

Date Reported:

09-Jul-2021

Date(s) of Test(s):

23-Jun-2021 to 07-Jul-2021

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

nBOOK 09-JUI-21

Mrs Natalie Boot

Business Support Administrator

This report is issued in accordance with the Conditions of Business of Lucideon Limited and relates only to the sample(s) tested.