



Diversey Europe Operations BV
Maarssenbroeksedijk 2
3542 DN Utrecht
The Netherlands

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www.Diversey.com

CERTIFICATE OF ANALYSIS

Date: 28 March 2023

Product Name	CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26 first off		
Product Code	7514852		
Batch Number	FMP23087	56714	
Production Date	28/03/2023		
Expiration Date	EXP 28/03/2025		

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	Limits Lower - Upper		Results
Appearance	Visual	Clear Colourless Liquid		Clear Colourless Liquid
pH (neat solution)	DM001	9.0	12.5	10.98
Specific Gravity (20°C)	DM004	1.004	1.020	1.018
Potential Chlorine Dioxide		100	180	131.625

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

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

Date: 28 March 2023

Product Name	CLEARKLENS BI-SPORE BASE SOLUTION VH26 first off		
Product Code	7514852		
Batch Number	FMP23087	56714	
Production Date	28/03/2023		
Expiration Date	EXP 28/03/2025		

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	Limits		Results
		Lower	Upper	
Appearance	Visual	Clear Colourless 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 behalf of Diversey site Quality Manger	Name:	Justyna Staron Edyta Rodrigues
	Position	Quality Control Inspector

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STERIS: Gamma Certificate Of Processing

Prepared for: FLEXIBLE MEDICAL PACKAGING LTD (8245)

Gamma Process Run ID 2173-35885A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
BI-SP B3 Components DV5857	56737	24	Case

Validation Reference Number: 5857

Processing Run Start Date: 17-Mar-2023 3:43 AM

Processing Run End Date: 17-Mar-2023 10:05 AM

Specified Dose Range (kGy):	25.0 - 40.0	Calculated Min Dose (kGy):	30.2
Reference Dose Range (kGy):	26.1 - 35.8	Calculated Max Dose (kGy):	36.5

PO Number: 404819

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

Gamma Process Run Approval authorized by STERIS

Date/Time E-Signed: 17-Mar-2023 12:36 PM

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 WO56737 will be used in finished batch Fmp23087 56714.

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: Fmp23087 56714

Lucideon Sample Number: UK231282-8729

Lucideon Report Number: UK231282-8729/MFEP **Issue Number:** 1

Date Logged: 11-Apr-2023 **Order Number:** 405003

Date Reported: 03-May-2023 **Date(s) of Test(s):** 13-Apr-2023 to 27-Apr-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

NcBoot 03 May 23

Mrs Natalie Boot

Senior Business Support Administrator

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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PHARMACEUTICAL ANALYSIS REPORT

LUCIDEON

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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: Fmp23087 56714

Lucideon Sample Number: UK231282-8730

Lucideon Report Number: UK231282-8730/MFEP **Issue Number:** 1

Date Logged: 11-Apr-2023 **Order Number:** 405003

Date Reported: 03-May-2023 **Date(s) of Test(s):** 13-Apr-2023 to 27-Apr-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

Natalie Boot 03 May 23

Mrs Natalie Boot

Senior Business Support Administrator

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