

Diversey Europe Operations BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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

CERTIFICATE OF ANALYSIS

Date: 16.01.2018

Product Name	CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26		
Product Code	7515828		
Batch Number	_FMP 18009 44375		
Production Date	09/01/2018		
Expiration Date	EXP 09 1011 2020		

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	Lim	uits	Results
		Lower -	Upper	
Appearance	Visual	Clear Colour	less Liquid	Clear Colour -
pH (neat solution)	DM001	9.0	12.5	12-4
Specific Gravity (20°C)	DM004	1.004	1.020	1.008
Potential Chlorine Dioxide (ppm)		100	180	138.375ppm

On behalf of Diversey site Quality Manager	Name:	4. Staron Maralandie
	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: 16.01.2018

Product Name	CLEARKLENS BI-SPORE BASE SOLUTION VH26
Product Code	7515828
Batch Number	FMP 18009 44375
Production Date	09/01/2018
Expiration Date	EXP 09/01/2020

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 Co Lic	olourless juid	Clear Colonless
pH (neat solution)	DM001	1.5	2.5	2.2
Specific Gravity (20°C)	DM004	1.010	1.030	1.016

O. 1. 1. 16 - 6Diit-	Name:	Varabuda J. Staron
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

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



Wickham Laboratories Ltd Hoeford Point, Barwell Lane, Gosport Hampshire PO13 0AU England

Telephone:

+44(0)1329 226600

+44(0)1329 226688

mail@wickhamlabs.co.uk www.wickhamlabs.co.uk

Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 **Hightown**

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

23 Jan 2018

Date Tested:

31 Jan 2018

Date Test Completed:

14 Feb 2018

Purchase Order:

PO30443

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0040415/2

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

23/01/2018

Test Article:

Bi-Spore Activator FMP18009 44375

Sample Code: **Batch Ref:**

595594

Qty Received:

20 x 100mL Bottles

Test	Method Item	Result
Sterility Test by Membrane Filtration (Steritest) Method	MM107/00	Pass
Growth in Tryptone Soya Broth at 20- 25°C after 14 days	MM107/01	No growth in two broths
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in two broths
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.0 2.6.1, USP 40 <71> & JP XVII 4.06
Product Standard Data Sheet	FG047/psd	FM04

Approval is provided by Electronic Signature. Their name and position is shown below.

BSc. CBiol. MRSB

Date: 28 Feb 2018 17:09:31

Mrs S.L. Wood

Page 1 of 1

Senior Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0040415 Print Number: P0049205





Wickham Laboratories Ltd Hoeford Point, Barwell Lane, Gosport Hampshire PO13 0AU England

Telephone:

+44(0) 1329 226600

Fax:

+44(0) 1329 226688

mail@wickhamlabs.co.uk www.wickhamlabs.co.uk

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Lancaster LA14XS

Date Received:

23 Jan 2018

Date Tested:

31 Jan 2018

Date Test Completed:

14 Feb 2018

Purchase Order:

PO30443

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0040415/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

23/01/2018

Test Article:

Bi-Spore Base

Sample Code:

FMP18009 44375

Batch Ref: 587518

Qty Received:

20 x 100mL Bottles

Test	Method Item	Result
Sterility Test by Membrane Filtration (Steritest) Method	MM107/00	Pass
Growth in Tryptone Soya Broth at 20- 25°C after 14 days	MM107/01	No growth in two broths
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in two broths
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.0 2.6.1, USP 40 <71> & JP XVII 4.06
Product Standard Data Sheet	FG047/psd	FM05

Approval is provided by Electronic Signature. Their name and position is shown below.

BSc. CBiol, MRSS

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Page 1 of 1

Senior Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0040415 Print Number: P0049205







http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 22-Dec-2017

UK33S12030190-1-1

This is to certify that AST Daventry Synergy Health PLC, a STERIS Company has where appropriate delivered an irradiation process in accordance with the current certified standards:

EN ISO 11137-1 Sterilisation of Health Care Products EN ISO 13485 Quality System - Medical Devices

Flexible Medical Packaging Ltd Unit 8, White Cross Ind Estate Hightown Lancaster Lancashire LA1 4XS UNITED KINGDOM

All in accordance with current Technical Agreement

Account Number: 100432 Synergy Health Sales Part Reference: 1106825 **Customer Reference Number:** P030170 **Product Description:** BISP B3 COMPONENTS DV4725 25-40kGy Validation Reference: 4725 **Ouantity Received:** 25.0 **Customer Minimum Specification kGy: Customer Maximum Specification kGy:** 40.0 Customer Unit Lot/Batch Number: W044378, 1 plt

Order Information

Irradiation Data				
Date and Time of Irradiation: 22-Dec-2017 17:46				
Reference Dose Range kGy:	30.2 - 30.3			
Calculated Minimum Dose kGy:	25.0			
Calculated Maximum Dose kGy:	30.9			

Items irradiated under WO44378, will be used in finished product batch FMP18009 44375

C Pascoe

Irradiation Release Authorised By Synergy Health Sterilisation UK, a STERIS Company