

Diversey Europe Operations BV Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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

CERTIFICATE OF ANALYSIS

Date: 13.03.18

Product Name	CLEARKLENS BI-SPORE BASE SOLUTION VH26
Product Code	7514852
Batch Number	FMP18064 44681
Production Date	05/03/2018
Expiration Date	EXP 05/03/2020

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	Lir	nits	Results
		Lower -	- Upper	
Amagranas	Visual	Clear Co	olourless	Clear Courless
Appearance	VISUAI	Lic	quid	liquid
pH (neat solution)	DM001	1.5	2.5	16
Specific Gravity (20°C)	DM004	1.010	1.030	1.020

On habelf of Diversors site	Name:	Marsharda 7 Storen
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

This document being issued electronical	lly does not bear a signature		
COA Template	Version: 02	Date of issuing: November 24 th	2017



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CERTIFICATE OF ANALYSIS

Date: 13,03.18

Product Name	CLEARKLENS BI-SPORE ACTIVATOR SOLUTION VH26
Product Code	7514852
Batch Number	FMP18064 44681
Production Date	05/03/2018
Expiration Date	EXP 05/03/2020

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	Li	mits	Results	
		Lower	- Upper		
Appearance	Visual	Clear Colo	urless Liquid	Clear Colourless	uelid
pH (neat solution)	DM001	9.0	12.5	11.6	6
Specific Gravity (20°C)	DM004	1.004	1.020	1.004	
Potential Chlorine Dioxide (ppm)		100	180	148.5ppm	

On habelf of Diverse with	Name:	4. Staron Navabuda
On behalf of Diversey site Quality Manager	Position	Quality Control Inspector

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

Telephone: Fax: +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 LA 1 4XS **Date Received:**

15 Mar 2018

Date Tested:

22 Mar 2018

Date Test Completed:

05 Apr 2018

Purchase Order:

30669

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0041484/2

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

15/03/2018

Test Article: Sample Code: Bi-Spore (Base) FMP 18064 44681

Batch Ref:

595593

Qty Received:

20 x 100mL Bottles Base (2 Spares)

Test	Method Item	Resulf
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.

Margo

CBiol MRSB

Date: 06 Apr 2018 16:16:49

Mrs C Moore

Page 1 of 1

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0041484 Print Number: P0050053





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

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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

15 Mar 2018

Date Tested:

22 Mar 2018

Date Test Completed:

05 Apr 2018

Purchase Order:

30669

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0041484/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

15/03/2018

Test Article: Sample Code: Bi-Spore (Activator) FMP 18064 44681

595594

Qty Received:

Batch Ref:

20 x 100mL Bottles Activor (2 Spares)

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.

2377200

CBio MRSB

Date: 06 Apr 2018 16:16:49

Mrs C Moore

Page 1 of 1

Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0041484 Print Number: P0050053





http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 11-Feb-2018

UK33S12057025-2-1

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

Order Information		
Account Number:	100432	
Synergy Health Sales Part Reference:	1106825	
Customer Reference Number:	P030455	
Product Description:	BI-SP B3 COMPONENTS DV4725 25-40kGy	
Validation Reference:	4725	
Quantity Received:	15	
Customer Minimum Specification kGy:	25.0	
Customer Maximum Specification kGy:	40.0	
Customer Unit Lot/Batch Number:	W044680, 2 pits	
I	rradiation Data	
Date and Time of Irradiation:	11-Feb-2018 07:03	
Reference Dose Range kGy:	32.9 - 33.3	
Calculated Minimum Dose kGy:	27.2	
Calculated Maximum Dose kGy:	34.0	

Items irradiated under WO44680, are to be used in finished product batch FMP18064 44681.

C Pascoe

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