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CERTIFICATE OF ANALYSIS

Date: 19 10 2018

Product Name	CLEARKLENS IPA VH1		
Product Code	7513400		
Batch Number	FHP 18274 45969		
Production Date	011012018		
Expiration Date	EXP 01 1 10 1 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 Liq		Gear Colour- less Liquid
Specific Gravity (20°C)	DM004	0.872	0.883	0.876

On behalf of Diversey site Quality Manager	Name:	Reityests, W. Marchiche
	Position	Quality Control Inspector

This document being issued electronically does not bear a signature

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



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 24-Oct-2018

UK33S12184414-1-1

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

Order Information **Account Number:** 100432 Synergy Health Sales Part Reference: 1104747 **Customer Reference Number:** P031918 **Product Description:** IPA BOTTLE DV4648 25-45kGv Validation Reference: 4648 **Ouantity Received:** 557 **Customer Minimum Specification kGy:** 25.0 **Customer Maximum Specification kGy:** 45.0 **Customer Unit Lot/Batch Number:** FMP18274 45969, 11 PLTS **Irradiation Data** Date and Time of Irradiation: 24-Oct-2018 04:05 Reference Dose Range kGy: 34.2 - 35.3 Calculated Minimum Dose kGy: 27.0 39.9 Calculated Maximum Dose kGy:

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



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 24-Oct-2018 UK33S12181434-1-1

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

o	rder information
Account Number:	100432
Synergy Health Sales Part Reference:	1104747
Customer Reference Number:	P031884
Product Description:	IPA BOTTLE DV4648 25-45kGy Validation
Reference:	4648
Quantity Received:	757
Customer Minimum Specification kGy:	25.0
Customer Maximum Specification kGy:	45.0
Customer Unit Lot/Batch Number:	FMP18274 45969, INC 1 SMPL, 14 PLTS
	rradiation Data
Date and Time of Irradiation:	24-0ct-2018 01:52
Reference Dose Range kGy:	34.6 - 35.8
Calculated Minimum Dose kGy:	27.3
Calculated Maximum Dose kGy:	40.5

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

A2



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 09-Oct-2018

UK33S12178073-1-2

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

0	rder Information
Account Number:	100432
Synergy Health Sales Part Reference:	1104747
Customer Reference Number:	P031851
Product Description:	IPA BOTTLE DV4648 25-45kGy
Validation Reference:	4648
Quantity Received:	114
Customer Minimum Specification kGy:	25.0
Customer Maximum Specification kGy:	45.0
Customer Unit Lot/Batch Number:	FMP18274 45969, INC 6 SAMPLES, 2 PLTS
ı	rradiation Data
Date and Time of Irradiation:	09-0ct-2018 00:13
Reference Dose Range kGy:	36.2 - 36.7
Calculated Minimum Dose kGy:	28.6
Calculated Maximum Dose kGy:	41.5

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



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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

17 Oct 2018

Date Tested:

19 Oct 2018

Date Test Completed:

19 Oct 2018

Purchase Order:

31899

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0045305/2

Test Required:

Bacterial Endotoxin Testing by Kinetic QCL

Date Received:

17/10/2018

Test Article:

Clearklens IPA FMP18274 45969

Sample Code: **Batch Ref:**

9688

Qty Received:

1 x 900mL Bottle

Test	Method Item	Result
Bacterial Endotoxin by Kinetic QCL Test	MM110/00	<0.2 EU/mL
Spiked Recovery	MM110/01	56 %
Dilution Tested	MM110/04	1/40
Endotoxin Spike Level	MM110/02	0.5 EU/mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.3 2.6.14, USP 41 <85> & JPXVII 4.01
Product Standard Data Sheet	FG047/psd	FM02-01

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

Date: 22 Oct 2018 10:25:29

Vicky Wilson

Laboratory Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0045305 Print Number: P0054425 Page 1 of 1





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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hiahtown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

17 Oct 2018

Date Tested:

24 Oct 2018

Date Test Completed:

07 Nov 2018

Purchase Order:

PO31899

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0045305/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received: Test Article:

17/10/2018 Clearklens IPA

FMP18274 45969

Sample Code: **Batch Ref:**

9688

Qty Received:

20 x 900mL Bottles (+ 3 Spare)

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 one broth
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in one broth
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	See external comments
Product Standard Data Sheet	FG047/psd	FM03-01

Comments

The testing procedure was performed in compliance with the current Pharmacopoeias however, the volume tested does not comply with the requirements of the current Pharmacopoeias.

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

BSc CBlot MRSB

Date: 08 Nov 2018 13:05:48

Mrs P. Pham-Lengoc

Documentation Manager - Microbiology

Certificate of Analysis - OSMM Consignment: 0045305 Print Number: P0054873 Page 1 of 1





