







Certificate of Analysis

Product Name:	Crystel Silver CRY 104	
Reference:	CRY/SIL/019	
Batch Number:	P706900	
Date of Manufacture:	08-MAR-17	
Expiry Date:	MAR-2019	

This is to certify that Crystel Silver solution above was tested on o8-MAR-17 and found to comply with all requirements as set out in the specifications and method of analysis.

PROPERTIES	SPECIFICATION	RESULTS
APPEARANCE	Clear and free from suspended matter	Conforms
ODOUR	Alcohol Like	Conforms
SPECIFIC GRAVITY	0.875 - 0.895	0.882
STERILITY	No growth	Conforms
IRRADIATION	Achieves minimum dose	Conforms
ENDOTOXIN	<0.25EU/ml	⟨0.1
ENDUTUAIN	Recovery 50-200%	138%

Completed By:	Reviewed By:
Quality Department: COMO News	X10pleC
Date: 25 April 2017	Date: 25 APR 17

DRF1574 CRY/SIL/007/Issue5/Sept 2012





Sterility Testing Analytical Report

Please note the results below are not generated at Associates of Cape Cod Int. Inc., but reported from a subcontractor's final report.

Sample ID: CRY104 Silver

Lot No: P706900 ICN: 0317023F

Subcontractor's Report Number: 2017062834

Lab Book Reference: ST03-71 Date Received: 29 Mar 2017 Date Completed: 21 Apr 2017 No. of samples: 10

Sterility Test - Membrane Filtration	
Medium	Result
Tryptone Soy Broth	No growth
Fluid Thioglycollate Medium	No growth
Fluid Thiogrycollate Fledram	

Tryptone Soy Broth	No growth
Fluid Thioglycollate Medium	No growth
Conclusions:	
	erility in accordance with the client's methodology.)
(The samples submitted comply with the test for st	erinty in accordance with the eneme means 37,7
- water many and a discount	

CONTROL PROPERTY SERVICE A SHIP HE AS A SHIP OF SERVICE SERVIC	
Results Reported by:	
Laboratory Tecnnician: Asnleigh Middleton	Date: 214pr2017
Reviewed by: Quality Assurance Manager UK General Manager	Date: 21 APR 2017
☐ Technical Manager ☐ Laboratory Manager	p.
QA Approved by:	0.5
Attens	Date: 216927297
Quality Assurance Menager	9.50

Revision History:

Technical Manager

DCCE NO:	Revision:	Description:	Originator:	Effective Date:
N/A	1	Revision 1, no changes	A Middleton	21 Apr 2017

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ST03-71 REVISION NO: 1

ICN: 0317023F DATE: 21 Apr 2017 TSCTS044-1 Rev 2

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Specialists in Endotoxin and Glucan Detection

LIMULUS AMEBOCYTE LYSATE (LAL) ANALYTICAL REPORT Release Test - Kinetic Turbidimetric Method (SOPCTS012)

Sample ID: CRY104 Silver

Lot No: P706900 ICN No: 0317023F **Lab Book Reference**: EL185-148 **Date Received**: 29 Mar 2017 **Date Completed**: 06 Apr 2017

Reagents

Pyrotell®-T: Lot Number 516-05-784-T. Pyrotell®-T was reconstituted with Pyrosol® Buffer and used to Assay all dilutions.

Pyrosol® Buffer: Lot Number 226-398.

LAL Reagent Water (LRW): Lot Number AAK207310. LRW was used to reconstitute endotoxin, prepare dilutions of endotoxin and sample(s), and serve as the negative control.

Instrumentation: Pyros® Kinetix Incubating Kinetic Tube Reader; S/N; 1E-1055.

Control Standard Endotoxin (CSE): Lot number 152, Escherichia coli O113:H10. The potency of the CSE is 20 EU/ng when measured against Reference Standard Endotoxin (RSE), lot number H0K354 (Food and Drug Administration, CBER) with Pyrotell®-T lot number 516-05-784-T.

The sensitivity of the assay is the lowest concentration of CSE used to construct the calibration curve. A series of 10-fold dilutions between 0.001 and 1.0 EU/mL was tested in duplicate and the results were used to construct the standard curve.

Preparation and Testing of Sample: The samples were diluted with LRW and analysed at a 1:100 dilution, in accordance with validation EL168-06. Positive product controls were tested, in parallel, on sample dilutions spiked with an Endotoxin concentration of 0.1 EU/ml. The sample to lysate ratio was 4:1.

Results: The Release Test is performed according to a validated method and is used to release finished products. The test can also be performed for release of raw/in-process materials. The absolute value for the correlation coefficient for the regression of log onset time versus log endotoxin concentration was greater than or equal to 0.980. The negative control (LRW) was tested in duplicate and contained less than 0.001 EU/mL. Inhibition or enhancement of the bacterial Endotoxin test by the sample is absent if the concentration of the Endotoxin assayed in the Positive Product Controls are within 50% to 200% of the nominal concentration.

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EL185-148 ICN; 0317023F TSCTS015-3.1 Rev 17
REVISION NO: 1 DATE: 06 Apr 2017

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Sample Results:

Sample Identification	% Spike Recovery	EU/ml
CRY104 Silver Lot: P706900	138	< 0.1

All samples analysed passed the specification of < 0.25 EU/ml.

Approval:	
Techncian: Sophie Hughes	Date: 06 APR 2013
Reviewed By:	Date: 1/AP/2017
☐ Quality Assurance Manager ☐ UK General Manager ☐ Technical Manager ☐ Laboratory Manager	
QA Approval:	Date: 12 AP/ 2017
☑ Quality Assurance Manager ☐ UK General Manager ☐ Technical Manager	

References:

Associates of Cape Cod SOPCTS012, General Kinetic LAL Turbidimetric Assay
United States Pharmacopoela <85> Bacterial Endotoxins Test
European Pharmacopoela, Section 2.6.14.

Revision History:

	£ .	Date:
N/A 1 Revision 1, no changes.	S. Hughes	06 Apr 2017

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EL185-148 REVISION NO: 1

ICN: 0317023F DATE: 06 Apr 2017 Page 2 of 2

TSCTS015-3.1 Rev 17



http://www.synergyhealthplc.com

Certificate of Irradiation

Date Issued: 19-Mar-2017

UK33S11834458-1-1

This is to certify that AST Daventry Synergy Health PLC has where appropriate delivered an irradiation process in accordance with:

EN ISO 11137-1:2015 Sterilisation of Health Care Products EN ISO 9001:2008 Quality Management System EN ISO 13485:2012 Quality System - Medical Devices

The Tristel Company Ltd
Unit 4C Lynx Bus Park, Fordham Rd
Newmarket
Suffolk CB8 7NY
UNITED KINGDOM

0	Order Information		
Account Number:	100708		
Synergy Health Sales Part Reference:	1108717		
Customer Reference Number:	704407		
Product Description:	ALCOHOL 1 LTR TRIGGER SPRAY DV4778		
	20-45kGy		
Validation Reference:	4778		
Quantity Received:	80		
Customer Minimum Specification kGy:	20.0		
Customer Maximum Specification kGy:	45.0		
Customer Unit Lot/Batch Number:	P706900, 2 plts		
I	rradiation Data		
Date and Time of Irradiation:	19-Mar-2017 09:20		
Reference Dose Range kGy:	33.7 - 34.3		
Calculated Minimum Dose kGy:	27.3		
Calculated Maximum Dose kGy:	39.5		