







# **Certificate of Analysis**

Product Name:	Crystel Silver CRY 104
Reference:	CRY/SIL/019
Batch Number:	P625100
Date of Manufacture:	30-AUG-16
Expiry Date:	AUG-2018

This is to certify that Crystel Silver solution above was tested on 30-AUG-16 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	0.878
STERILITY	No growth	Conforms
IRRADIATION	Achieves minimum dose	Conforms
ENDOTOVIN	<0.25EU/ml	⟨0.1
ENDOTOXIN	Recovery 50-200%	119%

Completed By:

Quality Department:

Date: 8 Dec 2016

Reviewed By:

Date: 08 12 16

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:P625100 ICN: 1116018F

Subcontractor's Report Number: 2016259131

Lab Book Reference: ST03-41 Date Received: 14 NOV 2016 Date Completed: 01 DEC 2016

No. of samples: 10

Sterility Test - Membrane Filtration

Sterility lest - Membrane Filtration	
Medium	Result
Tryptone Soy Broth	No growth
Fluid Thioglycollate Medium	No growth

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Date: 02 06C 2016
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## Revision History:

DCCE No:	Revision:	Description:	Orig!nator:	Effective Date:
N/A	1	Revision 1, no changes	C Ray	01 DEC 2016

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ST03-41 REVISION NO: 1 ICN: 1116018F DATE: 01 DEC 2016 TSCT5044-1 Kev 2

UK Dearon Park Meorgate Road Knowsky Everpool L33 7RX TeE +44 (0)151 547 7444 Ees +44 (0)151 547 7100

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USA 124 Bernard E. Stint Jean Drive-East Falmouth MA 02536-4445 USA Tel: 001 508 540 3444 Usy: 001 508 540 8680 Germany PYROQUANT DIAGNOSTIK GmbH Opelstrase 11 D-64546 Mortelden-Walldort Germany 'Rd: 0049-6165-96-10-0 Ecc. 0049-6105-96-10-15

Web: www.neeiuk.co.uk



Specialists in Endotoxin and Glucon Detection

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

Sample ID: CRY104 SILVER

Lot No: P625100 ICN No: 1116018F Lab Book Reference: EL186-73 (B)
Date Received: 14 NOV 2016
Date Completed: 17 NOV 2016

#### 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 226-391.

LAL Reagent Water (LRW): Lot Number AAH207245. 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; 3G-1166.

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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EL186-73 (B) REVISION NO: 1 ICN: 1116918F DATE: 17 NOV 2016 TSCTS015-3.1 Rev 17

UK Deacon Park Mrougate Road Knowsky Liverpool L33 7RX Tel: +14 (0)151 547 7444 Eax: +44 (0)151 547 7400 USA 124 Bernard J., Saint, Jean Drive East Edinouth VIA 02536-1445 USA Tel. 001 508 540 3444 East 001 508 540 8680 Germany PYROQUANT DIAGNOSTIK GmbH Opelsnasse 14 Di64546 Mortelden-Walldorf Germany 1cl 0049 6105 96 10 0 Lay 0049 6105 96 10 15



### Sample Results:

I	Sample Identification	% Spike Recovery	EU/ml
	CRY104 SILVER P625100	119	<0.1

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

oval:	
Techncian: Laboratory Technician: Charlotte Ray	Date: 13 Nov 2016
Reviewed By:	Date: 11. NO. 17.016
☐ Quality Assurance idanager ☐ UK General Manager ☐ Technical Manager ☐ Laboratory Manager QA Approval:	Date: 18 NOV 2016
□ Quality Assurance Manager □ UK General Manager 교·Fechnical Manager	
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## **Revision History:**

DCCF No:	Revision:	Description:	Originator	Effective Date:
N/A	1	Revision 1: No Changes	C RAY	17 NOV 2016

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EL186-73 (3) REVISION NO: 1

ICN: 1116018F **DATE: 17 NOV 2016**  Page 2 of 2

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http://www.synergyhealthplc.com

# **Certificate of Irradiation**

Date Issued: 31-0ct-2016 UK33S11738288-1-1

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

Or	der Information
Account Number: Synergy Health Sales Part Reference: Customer Reference Number: Product Description:	100708 1108717 703438 ALCOHOL 1 LTR TRIGGER SPRAY DV4778
Validation Reference: Quantity Received: Customer Minimum Specification kGy: Customer Maximum Specification kGy: Customer Unit Lot/Batch Number:	20-45kGy 4778 153 20.0 45.0 CRY 104, P625100, 3PLTS
<u> </u>	rradiation Data
Date and Time of Irradiation: Reference Dose Range kGy: Calculated Minimum Dose kGy: Calculated Maximum Dose kGy:	28-Oct-2016 23:38 33.9 - 35.5 27.5 40.9