







Certificate of Analysis

Product Name:	Crystel GOLD CRY 101	
Reference:	CRY/GOL/019	
Batch Number:	WO45264	
Date of Manufacture:	2018-05-29	
Expiry Date:	2020-05-29	

This is to certify that Crystel Gold solution above was tested on 2018-05-29 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	0.870 - 0.890	0.879
STERILITY	No growth	Conforms
IRRADIATION	Achieves minimum dose	Conforms
ENDOTOVIN	<0.25EU/ml	⟨0.1
ENDOTOXIN	Recovery 50-200%	134%

Completed By:

Quality Department: Canaleus

Date: 2018 - 08 - 10

Reviewed By

Date: 2018 -08 - 10

DRF1574 CRY/GOL/006/Issue4/Sept2012



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: CRY101 Gold

Lot No: WO45264 ICN: 0718012D

Subcontractor's Report Number: 2018133730

Lab Book Reference: ST03-179 Date Received: 16 JUL 2018 Date Completed: 08 AUG 2018

No. of samples: 10

Sterility Test - Membrane Filtration

Medium Result		
Tryptone Soy Broth	No Growth	
Fluid Thioglycollate Medium	No Growth	

Fluid Thioglycollate Medium	No Growth
Conclusions: The samples submitted comply with the test for s	sterility in accordance with the client's methodology.
Results Reported by: Laboratory Technician/Designee: L Ellis	Date: 08AUG 2019
Reviewed by: Quality Assurance Manager UK General Manager Technical Manager Laboratory Manager	Date: 09 Avg 2018
QA Approved by:	

alskyls	Date: 09 Aug 2018
Quality Assurance	

Revision History:

UK General Manager
Technical Manager

DCCF No:	Revision:	Description:	Originator:	Effective Date:
NA	1	Revision 1, no changes	L Ellis	08 AUG 2018

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ICN: 0718012D

Page 1 of 1 TSCTS044-1 Rev 3

REVISION NO: 1

DATE: 08 AUG 2018

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PYROQUANT DIAGNOSTIK GmbH
Opelstrasse 14
D-64546 Mörfelden-Walldorf
Germany
Tel: 0049 6105 96 10 0
Fax: 0049 6105 96 10 15



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

Sample ID: CRY101 Gold

Lot No: WO45264 ICN No: 0718012D Lab Book Reference: EL189-96-A
Date Received: 16 JUL 2018
Date Completed: 18 JUL 2018

Reagents

Pyrotell®-T: Lot Number 517-10-848-T. Pyrotell®-T was reconstituted with Pyrosol® Buffer and used to Assay all dilutions.

Pyrosol® Buffer: Lot Number 226-425.

LAL Reagent Water (LRW): Lot Number AC10244546. 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 159, The potency of the CSE is 14 EU/ng when measured against Reference Standard Endotoxin (RSE), lot number H0K354 (Food and Drug Administration, CBER) with Pyrotell®-T lot number 517-10-848-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 EL168-03. 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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Page 1 of 2

EL189-96-A REVISION NO:1 ICN: 0718012D DATE:18 JUL 2018 TSCTS015-3.1 Rev 18

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Sample Results:

Sample Identification	% Spike Recovery	EU/ml
WO45264	134	<0.1

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

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Approval:					
Techno Laboratory	ian: / Technician: L	Ellis	Date:	3062	018
Review E □ Laborato	oard	(Amers	Date:	19JUL2	2)18
QA App	roval:	iainart.	Date: 19	Jul 201	8
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References:					
Associates o	f Cape Cod SOF	CTS012, General Kinetic LAL Turbidimetri	ic Assay		
United State	s Pharmacopoe	ia <85> Bacterial Endotoxins Test			
European Ph	armacopoeia, S	Section 2.6.14.			
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Revision His	tory:				
DCCF No:	Revision:	Description:	Origin		Effective Date:
NA	1	Revision 2 No Changes	L Ellis	;	18JUL2018

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Page 2 of 2

EL189-96-A REVISION NO:1 ICN: 0718012D DATE:18 JUL 2018 TSCTS015-3.1 Rev 18



http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 08-Jul-2018

UK33S12133266-1-1

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

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

Order Information		
Account Number:	100708	
Synergy Health Sales Part Reference:	1108717	
Customer Reference Number:	707546	
Product Description:	ALCOHOL 1 LTR TRIGGER SPRAY DV4778	
	20-45kGy	
Validation Reference:	4778	
Quantity Received:	157	
Customer Minimum Specification kGy:	20.0	
Customer Maximum Specification kGy:	45.0	
Customer Unit Lot/Batch Number:	BN: W045264, 3 PLTS	
I	rradiation Data	
Date and Time of Irradiation:	08-Jul-2018 20:30	
Reference Dose Range kGy:	33.9 - 34.2	
Calculated Minimum Dose kGy: 27.5		
Calculated Maximum Dose kGy: 39.4		

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