

## PRODUCT CERTIFICATE

Product: Contec Sterile ProChlor  
Product Code: SBT102PC  
Product Description: Sterile Stabilised Hypochlorous Acid in purified water 1L Trigger Spray  
Batch Number: 250303039  
Manufacture Date: MAR / 2025  
Expiry Date: MAR / 2027

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### ANALYSIS

Test	Specification	Results
Colour:	Colourless	Complies
Clarity:	Clear	Complies
Filtration:	Filtered to 0.2 microns	Complies
SG at 20°C:	0.990 – 1.010	1.001
Available chlorine:	>1000ppm	2179
pH at 20°C :	3.0 – 6.0	3.74

Manufactured product via a Quality System certified to ISO 9001:2015, tested in accordance with documented quality procedures and approved when required specifications are met.

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### STERILITY

Sterility test number: 0000935197, 0000935198, 0000935199

Sterility test result: No evidence of microbial growth

Test method as described in the current edition of the European Pharmacopoeia.

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Name:	1: Kobey Robson	2: Nicola Hunter
Position:	1: Quality Assurance Officer	2: Quality Assurance Officer
Date:	1: 19 JUN 25	2: 19 Jun 25
Authorised Signature:	1: 	2: 

For and on behalf of Contec Inc

COA09 Rev 4

**Manufactured by:**  
Contec Cleanroom (UK) Ltd  
Unit 6A Wansbeck Business Park  
Ashington  
UK

**America**  
Contec Inc  
P.O.Box 530  
Spartanburg SC  
USA

**Europe**  
Contec Inc  
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France

**China**  
Contec Cleanroom Technology (Suzhou) Co. Ltd  
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# STERIS: Gamma Certificate Of Processing

Prepared For                    CONTEC CLEANROOM UK LTD (8250)  
Gamma Process Run ID      2173-55643A

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<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
PRIMARY PACKAGING 1L SWIRL DV4688	250303039	64	Case
Validation Reference Number: 4688			

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Processing Run Start Date 17-Apr-2025 10:35 PM

Processing Run End Date 18-Apr-2025 5:07 AM

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Specified Dose Range (kGy)	25.0 - 45.0	Calculated Min Dose (kGy)	30.3
Reference Dose Range (kGy)	26.2 - 41.6	Calculated Max Dose (kGy)	36.1

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PO Number: 148867

Product meets Customer specifications; zero nonconformities occurred during this irradiation run.

**Gamma Process Run Approval authorized by STERIS**

DateTime Esigned 18-Apr-2025 6:47 AM

Operating facilities are in compliance with applicable state and federal regulations providing services under a certified quality system which meets the following requirements (if applicable): FDA QSR (will be updated to QMSR), ISO 9001, EN/ISO 13485 current certified version, national pharmaceutical GMP and is in alignment with EN ISO 11137 current certified version. STERIS AST certifies that these processed items received the indicated doses within the precision and accuracy of the dosimetry system used.

## Processing Location

Synergy Health Sterilisation UK  
Limited, a STERIS Company  
Brunel Close  
Drayton Fields Industrial Estate  
Daventry  
Northants  
NN11 8RB  
Phone: + 44(0) 1327 706 111



Document ID: 206111

N/A

Last Revised in Rel 2.0.0.0

Rel Date: 13-Aug-2018

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