

PRODUCT CERTIFICATE

Product: Contec Sterile ProChlor

Product Code: SBT102PC

Product Description: Sterile Stabilised Hypochlorous Acid in purified water 1L Trigger Spray

Batch Number: 250503087

Manufacture Date: May / 2025

Expiry Date: May / 2027

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	2122
pH at 20°C :	3.0 – 6.0	3.58

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.

STERILITY

Sterility test number: 0000947670, 0000947671, 0000947672

Sterility test result: No evidence of microbial growth

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

Name: 1: Nicola Hunter 2: Kobey Robson

Position: 1: Quality Assurance Officer 2: Quality Assurance Officer

Date: 1: 14 AUG 25 2: 14 AUG 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
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America
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STERIS: Gamma Certificate Of Processing

Prepared For CONTEC CLEANROOM UK LTD (8250)
Gamma Process Run ID 2173-57014A

Claxon 01Jun25

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
PRIMARY PACKAGING 1L SWIRL DV4688	250503087	64	Case

Validation Reference Number: 4688

Processing Run Start Date 31-May-2025 5:30 PM

Processing Run End Date 01-Jun-2025 12:17 AM

Specified Dose Range (kGy)	25.0 - 45.0	Calculated Min Dose (kGy)	30.2
Reference Dose Range (kGy)	26.2 - 41.6	Calculated Max Dose (kGy)	35.9

PO Number: 150088

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

Gamma Process Run Approval authorized by STERIS

DateTime Signed 01-Jun-2025 4:43 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
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Document ID: 210420

N/A

Last Revised in Rel 2.0.0.0

Rel Date: 13-Aug-2018

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