

PRODUCT CERTIFICATE

Product: Contec Sterile ProChlor
Product Code: SBT102PC
Product Description: Sterile Stabilised Hypochlorous Acid in purified water 1L Trigger Spray
Batch Number: 250403069
Manufacture Date: APR / 2025
Expiry Date: APR / 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	2221
pH at 20°C :	3.0 – 6.0	3.72

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: 0000943851, 0000943852, 0000943853

Sterility test result: No evidence of microbial growth

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

Name:	1: Kobey Robson	2: Nicola Hunter
Position:	1: Quality Assurance Officer	2: Quality Assurance Officer
Date:	1: 09 JUL 25	2: 09 JUL 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
ZI du Prat RP 3707
56037 VANNES
France

China
Contec Cleanroom Technology (Suzhou) Co. Ltd
No. 17 Longyun Road
Suzhou 215024
China

www.conteclnc.com
[Infoeu@contecinc.com](mailto:infoeu@contecinc.com)

STERIS: Gamma Certificate Of Processing

Prepared For CONTEC CLEANROOM UK LTD (8250)
Gamma Process Run ID 2173-56571A

Checked 22MAY25

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
PRIMARY PACKAGING 1L SWIRL DV4688	250403069	64	Case

Customer Item ID: PREPSBT102PC
Validation Reference Number: 4688

Processing Run Start Date 16-May-2025 2:16 AM

Processing Run End Date 16-May-2025 9:26 AM

Specified Dose Range (kGy)	25.0 - 45.0	Calculated Min Dose (kGy)	31.1
Reference Dose Range (kGy)	26.2 - 41.6	Calculated Max Dose (kGy)	36.2

PO Number: 149676

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

Gamma Process Run Approval authorized by STERIS

DateTime Esigned 16-May-2025 1:07 PM

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: 208931

N/A

Last Revised in Rel 2.0.0.0

Rel Date: 13-Aug-2018

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