

PRODUCT CERTIFICATE

Product: Contec Sterile NeutraKlean
Product Code: SBC503NK
Product Description: Sterile Neutral Detergent in purified water (EP) 5L Capped
Batch Number: 250303029
Manufacture Date: Mar / 2025
Expiry Date: Mar / 2027

ANALYSIS

Test	Specification	Results
Colour:	Colourless	Complies
Clarity:	Clear	Complies
Filtration:	Filtered to 0.2 microns	Complies
SG at 20°C:	0.995 - 1.010	1.000
pH:	6.0 to 8.0	7.84

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.

IRRADIATION

Irradiation certificate number: 2173-55641A
Irradiation Dose (kGy): > 25 kGy > 33.1



We certify that the notified goods have undergone irradiation by exposure to γ (Gamma) irradiation with an exposure of not less than 25 kGy to conform to the European Pharmacopoeia.

Irradiation treatment applied was accordance with:

ISO 13485:2016 Quality Management System – Medical Devices
ISO 11137:2015 Sterilisation of Healthcare Products – Requirements for Validation & Routine Control – Radio-sterilisation

STERILITY

Sterility test number: 0000930940
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: 17 JUN 25 2: 17 JUN 25
Authorised Signature: 1:  2: 
For and on behalf of Contec Inc

STERIS: Gamma Certificate Of Processing

Prepared For CONTEC CLEANROOM UK LTD (8250)
Gamma Process Run ID 2173-55641A

<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
Neutraklean 5L DV6097	250303029	95	Case
Validation Reference Number: 6097			

Processing Run Start Date 12-Apr-2025 4:56 AM

Processing Run End Date 12-Apr-2025 2:17 PM

Specified Dose Range (kGy)	25.0 - 50.0	Calculated Min Dose (kGy)	33.1
Reference Dose Range (kGy)	31.3 - 46.4	Calculated Max Dose (kGy)	45.5

PO Number: 148665

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

Gamma Process Run Approval authorized by STERIS

DateTime Signed 13-Apr-2025 8:24 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

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