

PART #: REF #: DESCRIPTION: LOT #:

DISPOSABLE POLYSTYRENE STERILE BLUE FORCEPS
WITH BLUNT TIP. FORCEPS ARE INDIVIDUALLY PEELPACKED.

343834

CERTIFICATE OF STERILIZATION

EXPIRATION DATE: 02/2026

The validation and routine control of the sterilization process are carried out in accordance with the following standards:

BS EN ISO 11135: 2014 Sterilization of health care products – Ethylene oxide – Part 1: Requirements

for development, validation and routine control of a sterilizations process for

medical devices

BS EN 556-1: 2001 Sterilization of medical devices – Requirements for medical devices to be

designated "Sterile" - Part 1: Requirements for terminally sterilized medical

devices

BS EN ISO 10993-7: 2008 Biological evaluation of medical devices – Part 7: Ethylene oxide sterilization

residuals

Routine control: The criteria for product release from sterilization

- Conformity with the specified physical parameters for the sterilization cycle

- No growth of the biological indicators

Provided appropriately stored, products remain sterile for at least 5 years from the sterilization date.

CERTIFICATE OF CONFORMITY

Has been developed, manufactured, inspected and sterilized in accordance with the requirements of:

21 CFR part 820: Quality system regulation.

ISO 13485:2016: Quality systems – Medical devices.

93/42/EEC: Council Directive concerning medical devices

EN 550-1994: Sterilization of medical devices –

Validation and routine control of ethylene oxide sterilization

EN 556-1, 2001: Sterilization of medical devices –

Requirements for medical devices to be designated "Sterile" Part 1: Requirement for terminally sterilized medical devices