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| **PART #:** | **REF #:** | **DESCRIPTION:** | **LOT #:** |
| **DF8088S** | **84011600** | **DISPOSABLE POLYSTYRENE STERILE BLUE FORCEPS WITH BLUNT TIP. FORCEPS ARE INDIVIDUALLY PEEL-PACKED.** | **358147** |
| **STERILIZATION DATE: 2024-08-26 (YYYY-MM-DD)** | **EXPIRATION DATE: 2029-07-31 (YYYY-MM-DD)** |

**CERTIFICATE OF STERILIZATION**

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

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

EN 556-1: 2001 Sterilization of medical devices – Requirements for medical devices to be designated “Sterile” – Part 1: Requirements for terminally sterilized medical devices

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:

ISO 13485:2016: Quality systems – Medical devices.

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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

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

 Requirements for medical devices to be designated “Sterile”

 Part 1: Requirement for terminally sterilized medical devices