# CERTIFICATE OF COMPLIANCE

Becton Dickinson Medical (S) Pte Ltd
30 TUAS AVENUE 2
SINGAPORE SG 639461 SG



**Product Name** : NEEDLE ECLIPSE 25X1 RB
25 G BD" Eclipse 1 in. single use sterile
**Catalog Number** : 305761 **Manufacture Date**: 2019/10/02
**Batch Number** : 9276555
**Expiration Date** : 2024/09/30

## REGULATORY COMPLIANCE AND QUALITY SYSTEM

BD Products comply with the regulatory requirements of the region in which these are sol d and manufactured.
BD Products sold in the US comply with the current FDA Quality System Regulation 21CFR820. Medical devices are listed with FDA per 21CFR807. Manufacturing sites are registered with FDA per 21CFR807. The devices satisfy FDA pre-market notification requirements per 21CFR 807.
BD Products which are CE marked comply with Medical Devices Directive 93/42/EEC and are manufactured within production facilities that comply with the international standard ISO 13485: Quality Systems - Medical Devices - Requirements For regulatory purposes.

## STERILITY

All products which are labeled as "sterile" and released for sale by BD are certified to be sterile as long as the package is unopened and undamaged.
This product is primarily sterilized via Ethylene Oxide. Sterilization cycle develo pment/validation is performed to 10-6 SAL in accordance with current ISO 11135 guidelines.

## BIOCOMPATABILITY

This product has been evaluated in accordance with ISO 10993 "Biological Evaluation of Medical De vices", and complies with all relevant sections.

## PYROGENICITY

All products which are labeled as non-pyrogenic and released for sale by BD have been tested per United States Pharmacopeia (USP) chapter 85 - Bacterial Endotoxins Test and meets limits as specified in chapter 161- Medical Devices- Bacterial Endotoxin and Pyrogen Tests.

## QUALITY CONTROL TESTING AND RELEASE

Representative production samples are collected and inspected in accordance with current applicable product specifications. Inspection records are reviewed and signed off by qualified personnel for product release. The released devices meet applicabl e BD product specification(s).

## PRODUCT SPECIFIC SPECIFICATIONS

This product complies with the following BD Specification(s): SP100123 and SP100129

## BD MANUFACTURING SITE

BD Manufacturing site:

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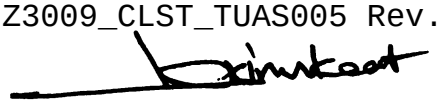
Becton Dickinson Medical (S) Pte Ltd
30 Tuas Ave 2, Singapore 639461 SG
Legal Manufacturer:
Becton Dickinson and Company
1 Becton Drive,Franklin Lakes,NJ 07417 USA
EU Authorized Representative:
Becton Dickinson Distribution Center NV
Laagstraat, 57 Temse 9140 Belgium
CE Certificate Number: 252.232
Z3009_CLST_TUAS005 Rev. 02 Effective 09 April 2018



Law Innkeat
Quality Assurance Management
Signature Date: 2019/10/31