



Becton Dickinson Medical (S) Pte Ltd  
30 Tuas Avenue 2  
Singapore 639461 SG

# CERTIFICATE OF COMPLIANCE

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**Product Name** : SYRINGE 3ML LL W/NDL ECLIPSE 22X1-1/2 RB  
**Catalog Number** : 305783  
**Batch Number** : 8110129  
**Expiration Date** : 2023/04/30

## **REGULATORY COMPLIANCE AND QUALITY SYSTEM**

BD Products comply with the regulatory requirements of the region in which these are sold 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 development/validation is performed to 10<sup>-6</sup> SAL in accordance with current ISO 11135 guidelines.

## **BIOCOMPATABILITY**

This product has been evaluated in accordance with ISO 10993 "Biological Evaluation of Medical Devices", 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



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release. The released devices meet applicable BD product specification(s).

## **PRODUCT SPECIFIC SPECIFICATIONS**

This product complies with the following BD Specification(s): 10005001

## **BD MANUFACTURING SITE**

BD Manufacturing site:  
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: 252.231

Z3009\_CLST\_TUAS003 Rev. 02 Effective 09 April 2018

Alex Budiman  
Quality Assurance Director  
Signature Date: 2018/05/24