CERTIFICATE OF COMPLIANCE

Becton, Dickinson and Company
BD Medical Surgical Systems
Route 7 and Grace Way
Canaan CT 06018-2415 US

Product Name : SYRINGE 1ML LL
1 mL BD Luer-Lok" Syringe sterile, single use polycarbonate
Catalog Number : 309628 **Manufacture Date**: 2018/01/01
Batch Number : 8001740
Expiration Date : 2022/12/31

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 21 CFR 820. Medical devices are listed with FDA per 21 CFR 807. Manufacturing sites are registered with FDA per 21 CFR 807. The devices satisfy FDA pre-market notification requirements per 21 CFR 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 Gamma. Sterilization cycle development/validation is performed to 10-6 SAL in accordance with current ISO 11137 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 release. The released devices meet applicable BD product specifications.

PRODUCT SPECIFIC SPECIFICATIONS

This product complies with the following BD Specification: SP100001

BD MANUFACTURING SITE

BD Manufacturing site:
BD Medical
Route 7 and Grace Way
Canaan, CT 06018, USA
Primary Sterilization site:
BD Medical
1852 10th Avenue
Columbus, NE 68601, USA

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Legal Manufacturer:
Becton, Dickinson and Company
1 Becton Drive
Franklin Lakes, NJ 07417, USA
EU Authorized Representative:
Becton Dickinson Distribution Center
Laagstraat 57B-9140
Temse, Belgium
CE Certificate Number: 252.231

Paul Strzepa
Quality Manager, BD Canaan