

CERTIFICATE OF COMPLIANCE

Page: 1 of 2

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

<pre>Product Name : SYRINGE 5ML LL SP125 Catalog Number : 309646 Manufacture Date :2016/03/28 Batch Number : 6056715 Expiration Date : 2021/02/28 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 210FR80. 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. STERLITY 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 E-beam. 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- Transfusion ad Infusion Assemblies and Similar Medical Devices. 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 specifications. PRODUCT SPECIFIC SPECIFICATIONS This product complies with the following ED Specification: SP10001 BD MANUFACTURING SITE ED Manufacturing site: ED Medical, Route 7 and Grace Way,</pre>	
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. STERLITY 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 E-beam. Sterilization cycle development/validation is performed to 10-6 SAL in accordance with current ISO 11137 guidelines. BIOCOMPATABLITY This product has been evaluated in accordance with ISO 10993 "Biological Evaluation of Medical Devices", and complies with all relevant sections. PYROCENICITY 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- Transfusion and Infusion Assemblies and Similar Medical Devices. 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 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, Route 7 and Grace Way, Canaan, CT 06018, USA	Catalog Number : 309646 Manufacture Date : 2016/03/28 Batch Number : 6056715
by BD have been tested per United States Pharmacopeia (USP) chapter 85 - Bacterial Endotoxins Test and meets limits as specified in chapter 161- Transfusion and Infusion Assemblies and Similar Medical Devices. 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, Route 7 and Grace Way, Canaan, CT 06018, USA Legal Manufacturer: Becton, Dickinson and Company, 1 Becton Drive, Franklin Lakes, NJ 07417, USA	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 E-beam. 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
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, Route 7 and Grace Way, Canaan, CT 06018, USA Legal Manufacturer: Becton, Dickinson and Company, 1 Becton Drive, Franklin Lakes, NJ 07417, USA	by BD have been tested per United States Pharmacopeia (USP) chapter 85 - Bacterial Endotoxins Test and meets limits as specified in chapter 161- Transfusion and Infusion Assemblies and Similar Medical Devices. 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.
LITE HAME TIONS CHELTS CHO KEATRIOH DACE 03/11/2012	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, Route 7 and Grace Way, Canaan, CT 06018, USA Legal Manufacturer: Becton, Dickinson and Company, 1 Becton Drive,



CERTIFICATE OF COMPLIANCE

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

Page: 2 of 2

Product Name	:	SYRINGE	5ML	LL	SP125		
Catalog Number	:	309646			Manufacture	Date	:2016/03/28
Batch Number	:	6056715					
Expiration Date	:	2021/02/	28				

Paul Strzepa 📿 Quality Manager, BD Canaan