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

| Product Name : Syringe 10ml LL |
|--|
| Catalog Number : 309604 |
| Batch Number : 7001704 |
| Expiration Date : 2021/12/31 |
| DECILLATION COMPLETANCE AND OTAL TTY CRETEM |
| 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 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 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 File name Z1003_CLST_IS_CN6 Revision Date 09/11/2015 |
| LITE HOUS COST TO CAN VEATOR DACE 03/11/2012 |
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CERTIFICATE OF COMPLIANCE

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Paul Strzepa 📿 Quality Manager, BD Canaan