



BD Diagnostics Preanalytical Systems 150 South 1st Avenue Broken Bow NE 68822-2203 US

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 Product Name
 : TUBE URIN PLC 16X100 8.0 CONI UA YEL

 Catalog Number
 : 364980
 Manufacture Date: 2017/11/06

 Batch Number
 : 7279799

 Expiration Date
 : 2019/04/30

REGULATORY COMPLIANCE AND QUALITY SYSTEM

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 21CFR807.

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.

STERILITY

All products which are labeled as "sterile" and released for sale by BD are certified to be sterile. For those products labeled "sterile fluid path", only the fluid path is sterile. This product is sterilized via Gamma irradiation process. Sterilization cycle development/validation is performed to 10-3 SAL in accordance with current ISO 11137 guidelines.

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 specification(s).

PRIMARY STERILIZIATON SITE: BD Life Sciences Preanalytical Systems 150 South 1st Avenue Broken Bow, NE 68822-2203 USA

LEGAL MANUFACTURER: Becton, Dickinson and Company 1 Becton Drive Franklin Lakes, NJ 07417-1885 USA



Certificate of Conformance

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Jennifer Jackson Quality Assurance Manager