



Certificate of Conformance

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

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Product Name : DEVICE URIN TRANSFER STRAW
Catalog Number : 364966
Batch Number : 5093545

MANUFACTURING CLAIM:

BD Diagnostics Preanalytical Systems products are manufactured in accordance with the medical device regulations (21CFR820) and comply with Medical Device Reporting (MDR) Regulations (21CFR803). All products and manufacturing facilities comply with FDA registration and listing requirements (21CFR807). The released products satisfy BD Diagnostics Preanalytical Systems finished product specifications. The Broken Bow facility is also ISO 13485:2003 certified.

Broken Bow Quality Assurance
Release date:2015/05/12
Name:Erlinda Larsen

This certificate is produced and controlled electronically and is valid without handwritten signature.