



# Certificate of Conformance

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

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**Product Name** : TUBE LIHEP PLH 13X100 6.0 PLBL GN  
**Catalog Number** : 367886 **Manufacture Date:** 2019/12/02  
**Batch Number** : 9315437  
**Expiration Date** : 2021/04/30

## CERTIFICATE OF STERILITY

### STERILITY CLAIM:

All products which are labeled as either "Sterile" or "Sterile Interior" and released for sale by BD Diagnostics Preanalytical Systems are certified to be sterile as long as the product package or product is unopened and undamaged. For those products labeled "Sterile Interior" only the product interior is sterile.

### 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 CMDCAS and ISO 13485:2003 certified.

**This product conforms to product specification VS57886.**

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

Jennifer Jackson  
Quality Assurance Manager