



# CERTIFICATE OF COMPLIANCE

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

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**Product Name** : SYRINGE ORAL 3ML AMBER  
**Catalog Number** : 305210 **Manufacture Date** : 2018/09/24  
**Batch Number** : 8263592

### 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 21 CFR 820. Medical devices are listed with FDA per 21 CFR 807. Manufacturing sites are registered with FDA per 21 CFR 807. The devices satisfy FDA pre-market notification requirements per 21 CFR 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.

### BIOCOMPATABILITY

This product has been evaluated in accordance with ISO 10993 "Biological Evaluation of Medical Devices", and complies with all relevant sections.

### 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: SP100061

### BD MANUFACTURING SITE

BD Manufacturing 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

EU Authorized Representative:

Becton Dickinson Distribution Center  
Laagstraat 57 B-9140  
Temse, Belgium

CE Certificate Number: 252.308

Paul Strzepa  
Quality Manager, BD Canaan