

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	5ML	AMBER
Catalog Number	:	305208			
Batch Number	:	7146857			

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. BIOCOMPATABILITY This product has been evaluated in accordance with ISO 10993 "Biological Evaluation of Medical Devices", and complies with all relevant sections. OUALITY 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 File name Z1003 CLST_IS_CN16 Revision Date 09/08/2015 Paul Strzepa 4

Quality Manager, BD Canaan