

EC Certificate - Full Quality Assurance System

Supplementary Information to CE 699333

Issued To:

**Arrow International LLC
 Subsidiary of Teleflex Incorporated
 3015 Carrington Mill Blvd.
 Morrisville
 North Carolina
 27560
 USA**

Number	Device Name	Intended purpose per IFU
Class III		
MD 0102	ARROWg+ard Blue and ARROWg+ard Blue Plus Central Venous Catheters (containing chlorhexidine), Sets and Kits	See CE 699338
MD 0102	Single-Lumen and Multi-Lumen Central Venous Catheter Sets and Kits	See CE 699342
MD 0102	Arrow Single and Multiple Lumen Peripherally Inserted Central Catheters (PICC)	See CE 699339
MD 0102	Hemodialysis Two-Lumen Catheters, Kits and Sets	See CE 699348
MD 0102	ARROWg+ard Blue® 2-Lumen Hemodialysis Catheters, Kits and Sets	See CE 699355
MD 0102	ARROWg+ard Blue ® Percutaneous Sheath Introducers, Kits and Sets	See CE 699356
MD 0106	Spring Wire Guide/ Guidewire	See CE 699363
MD 0100	Non-Heparin Coated ThermoDilution Catheters and Kits	See CE 699364

First Issued: **2020-06-11**

Date: **2020-12-01**

Expiry Date: **2024-05-26**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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