CE

FLUKE®
Biomedical

EU DECLARATION OF CONFORMITY

Product Identification:

First declared: **2009**

Fluke Biomedical IDA-5 - Infusion Device Analyzer

Statement of Conformity:

This Declaration of Conformity is issued under the sole responsibility of the manufacturer, based on sample product type-test, using ISO9001:2008 Quality Management System. The manufacturer hereby declares this product conforms to the following EU Directives:

Directive 2011/65/EU, RoHS
Directive 2014/30/EU, Electromagnetic Compatibility (EMC)
Directive 2014/35/EU, Low Voltage (LVD)

Standards Used:

EN 50581: 2012 Restriction of the use of hazardous substances
EN 61326-1: 2013 Electrical equipment for measurement, control and laboratory-EMC; General
EN 61010-1: 2010 Safety requirements for measurement, control and laboratory; General
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Special Conditions:

None

Manufacturer

Fluke Biomedical
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30 June 2016

Thomas M. Smith P.E.
Product Compliance Manager, Fluke Corporation
CE-94 Rev. 002 - This declaration supersedes all previous declarations for this product.