EU DECLARATION OF CONFORMITY

Product Identification:
Fluke Biomedical IDA-5 - Infusion Device Analyzer

First declared: 2009

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

Special Conditions:
None

Manufacturer
Fluke Biomedical
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[Signature]
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Product Compliance Manager, Fluke Corporation
CE-94 Rev. 002 - This declaration supersedes all previous declarations for this product.

30 June 2016