CE

FLUKE®
Biomedical

EU DECLARATION OF CONFORMITY

Product Identification:

First declared: **2015**

Fluke Biomedical Incu II - Incubation 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)
Directive 2014/53/EU, Radio Equipment (RED)
Directive 2006/66/EC, Batteries

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
EN 301 489-1 V1.9.2 Radio equipment and services
EN 301 489-17 V2.1.1 Broadband Data transmission systems
EN 62133: 2013 Safety requirements for portable sealed secondary cells
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Special Conditions:

None

Manufacturer

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06 July 2016

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Product Compliance Manager, Fluke Corporation
CE-105 Rev. r002 - This declaration supersedes all previous declarations for this product.