CE

FLUKE®
Biomedical

EU DECLARATION OF CONFORMITY

Product Identification: First declared: **2013**

Fluke Biomedical Impuse 6000D, Impulse 7000DP - Defibrillator/Pacer Analyzers

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

Standards Used:

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

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Thomas M. Smith P.E.
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CE 13 Rev. r005 - This declaration supersedes all previous declarations for this product.

25 May 2016