CE

FLUKE®
Biomedical

EU DECLARATION OF CONFORMITY

Product Identification: First declared: **2009**

Fluke Biomedical 451B-RYR., 451DE-SE-RYR - Ion Chamber Survey Meter

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 61326-2-2: 2013 Particular EMC requirements for portable test, measuring and monitoring equipment

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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CE 66 Rev. r008

23 June 2016

Thomas M. Smith P.E.
Product Compliance Manager, Fluke Corporation
CE 66 Rev. r008 - This declaration supersedes all previous declarations for this product.