



EU DECLARATION OF CONFORMITY

Product Identification:

Fluke Biomedical 601ProXL - International Electrical Safety Analyzer

First declared: 2012

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

EN 61010-2-030: 2010 Particular requirements for testing and measuring circuits

Special Conditions:

None

Manufacturer

Fluke Biomedical
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CD-16 Rev. r005 - This declaration supersedes all previous declarations for this product.

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