CE

FLUKE®
Biomedical

EU DECLARATION OF CONFORMITY

Product Identification:

First declared: **2003**

Fluke Biomedical Dale 301 - Rigid Endoscope Tester

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

Standards Used:

EN 50581: 2012 Restriction of the use of hazardous substances

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Special Conditions:

None

Manufacturer

Fluke Biomedical
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CE-103 Rev r002 - This declaration supersedes all previous declarations for this product.