



## EU Declaration of Conformity

### Product Identification

Product Name	Sensor Class
Oxygen Sensors	C1R, C2R, R13, R15, R17MED, R22MED, R23MED, R24MED, R26MED, R29MED, R29IMED, R30MED, R34MED, R36MED, J1, T1, T2, T4, T7, UFO130, UFO130-2.

### Manufacturer

Company Name	Address	Management Representative
Teledyne Analytical Instruments a business unit of Teledyne Instruments, Inc	16830 Chestnut Street City of Industry, CA 91748, USA	Vasu Narasimhan Director of Regulatory Affairs & Quality Assurance

### Authorized Representative

Company Name	Address	Contact Information
Viamed Limited	15 Station Road, Cross Hills Kieghley, W. Yorkshire BD20 7DT United Kingdom	Phone: +44 (0) 1535 634542 URL: <a href="http://www.viamed.co.uk">www.viamed.co.uk</a> E-mail: Jean@Viamed.co.uk

### Registration Information

Notified Body / ID #	CE Certificate Number	Date CE Marking first applied
BSI, UK / 0086	CE 02000	09 July 1998

### Conformity Assessment

Device Classification	Route of Compliance	Standards Applied
Class IIa (Annex IX, Rule 10)	Annex II, Section 3.2, of MDD 93/42/EEC, as amended by Directive 2007/47/EC	EN 60601-1, 3 <sup>rd</sup> Edition EN 60601-1-2 EN 60601-1-6 EN 14971:2007 EN ISO 21647

We, Teledyne Analytical Instruments, a business unit of Teledyne Instruments, Inc, declare that the above mentioned products meet the provision of the Council Directive 93/42/EEC, as amended by Council Directive 2007/47/EC, for Medical Devices. All supporting documents are retained at the premises of the manufacturer.

Signature: 

Name: Vasu Narasimhan

Title: Director of RA & QA

Date: October 8, 2013

Signature: 

Name: Stephen Broy

Title: Director of Engineering

Date: October 8, 2013