# EC DECLARATION OF CONFORMITY

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| **APPLICATION OF MEDICAL DEVICE DIRECTIVE** | **: Annex II excluding section 4 93/42/EEC, as amended by Directive 2007/47/EC** |
| **STANDARDS TO WHICH CONFORMITY IS DECLARED** | **: EN 60601-1 3.1 Edition<br>EN60601-1-2:2015<br>EN60601-1-6:2010<br>EN ISO 80601-2-55:2018<br>EN 14971:2019** |
| **MANUFACTURER'S NAME** | **: TELEDYNE ANALYTICAL INSTRUMENTS a business unit of Teledyne Instruments, Inc.** |
| **MANUFACTURER'S ADDRESS** | **: 16830 Chestnut Street<br>City of Industry, CA 91748<br>U.S.A.** |
| **PRODUCT NAME** | **: Oxygen Sensor** |
| **DEVICE CLASSIFICATION** | **: Class IIa (Annex IX, Rule 10)** |
| **MODEL NUMBER** | **: C1, C1R, C2R, R13, R15, R17MED, R22MED, R23MED, R24MED, R26MED, R29MED, R29IMED, R30MED, R34MED, R36MED, J1, T1, T2, T4, T7, UFO130, UFO130-2** |
| **EU Authorized Representative** | **: Viamed GmbH-UG<br>An der Trave 15<br>23923 Selmsdorf, Germany<br>Ph: + 49 38823 548800<br>E-mail: Info@Viamed-EU.com** |
| **Registration Information** | **: NB: BSI Netherlands /# 2797<br>CE Certification # CE 02000** |

**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.**

SIGNATURE: Nino Lugue
FULL NAME: Nino Lugue
POSITION: Quality Analyst
Date: 09-16-22

SIGNATURE: Alegria
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POSITION: New Products Manager
Date: 9-16-22