

EC DECLARATION OF CONFORMITY

**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
EN60601-1-2:2015
EN60601-1-6:2010
EN ISO 80601-2-55:2018
EN 14971:2019

MANUFACTURER'S NAME : TELEDYNE ANALYTICAL INSTRUMENTS
a business unit of Teledyne Instruments, Inc.

MANUFACTURER'S ADDRESS : 16830 Chestnut Street
City of Industry, CA 91748
U.S.A.

PRODUCT NAME : Oxygen Sensor

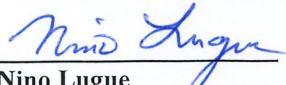
DEVICE CLASSIFICATION : Class IIa (Annex IX, Rule 10)

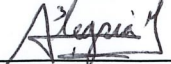
MODEL NUMBER : C1, C1R, C2R, R13, R15, R17MED, R22MED, R23MED,
R24MED, R26MED, R29MED, R29IMED, R30MED,
R34MED, R36MED, J1, T1, T2, T4, T7, UFO130,
UFO130-2

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Registration Information : NB: BSI Netherlands /# 2797
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: 
FULL NAME: Nino Luge
POSITION: Quality Analyst
Date: 09-16-22

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