

# EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

**No.****CE 02000****Issued To:**

**Teledyne Analytical Instruments  
A business unit of Teledyne  
Instruments, Inc  
16830 Chestnut Street  
City of Industry  
California  
91748  
USA**

**In respect of:**

**The design and manufacture of oxygen sensors**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):



Gary E Slack, Senior Vice President Medical Devices

**First Issued: 1998-07-09****Date: 2021-02-03****Expiry Date: 2023-07-08**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

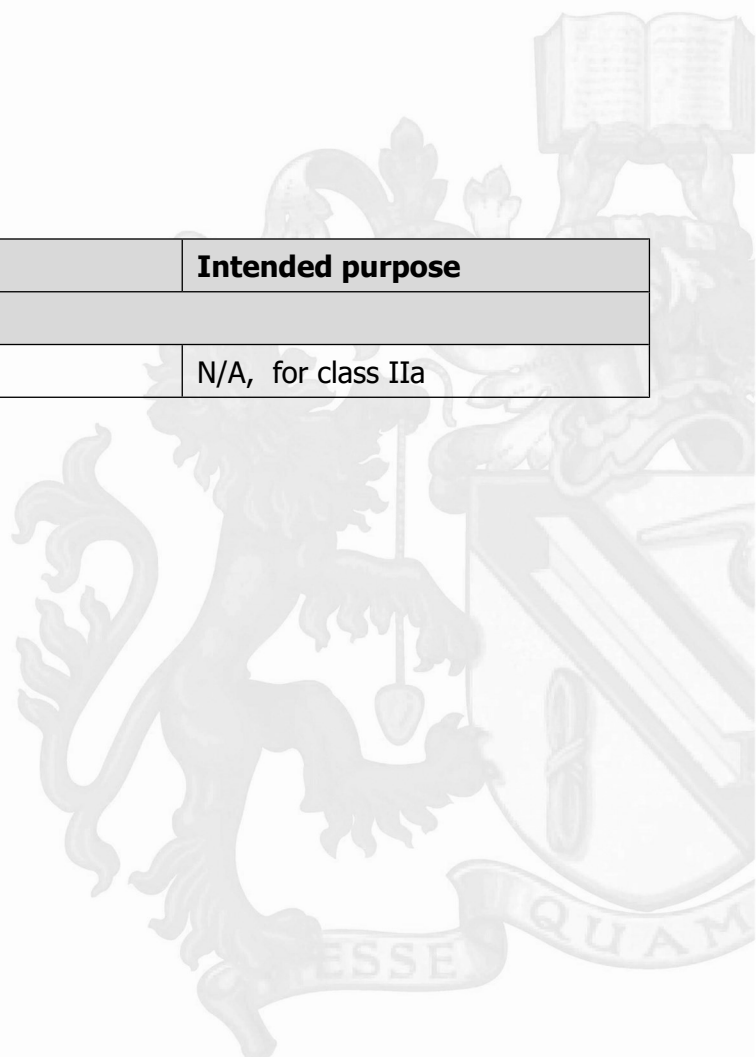
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## Supplementary Information to CE 02000

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**USA**

NBOG code(s)	Device description	Intended purpose
<b>Class IIa</b>		
MD 1102	Oxygen Sensors	N/A, for class IIa

First Issued: **1998-07-09**Date: **2021-02-03**Expiry Date: **2023-07-08**

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