

DECLARATION OF CONFORMITY

Manufacturer's Name: Maxtec
Address: 2305 South 1070 West
Salt Lake City, Utah 84119
USA

European Representative: QNET BV
Kantstraat19
NL-5076 NP Haaren
The Netherlands

Product: MAXO2+ Series Oxygen Analyzers

Model(s): MAXO2+ A, MAXO2+ AE, and Handi+

Classification & GMDN: IIa Analyzer, Gas, Oxygen 35219

Classification criteria: Clause 3.2 Rule 10 of Annex IX of MDD

We herewith declare that the above mentioned products meet the provisions of the following EC Council Directives and Standards. All supporting documents are retained under the premises of the manufacturer and the European Representative.

Directives: General application directives: Medical Device Directive, COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 per Annex II as amended by 2007/47/EC of 5 September 2007

Notified Body: TÜV SÜD Product Service
RIDLERSTRASSE 65, D-80339 MUNICH, Germany
Number 0123

EC Certificate No.: G1 16 10 45041 020

Date CE mark was affixed: 21 June 2004, - MAXO2+ / 03 December 1998 – Handi+

This declaration is considered valid from November 1, 2018 to December 18, 2021.

Signature:  Date: 11/01/2018

Name: Tammy Lavery
Position: Director of Regulatory and Quality

Applied Standards

The referenced list of harmonized standards for which documented evidence of compliance can be provided includes:

EN ISO 14971:2012
IEC 60601-1:2005 (EN 60601-1:2006)
IEC 60601-1-2:2004 (EN 60601-1-2:2004)
EN 62366:2008
EN 1041:2008
ISO 15223-1:2016