

## DECLARATION OF CONFORMITY

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

SRN: US-MF-000003961

European Representative: EMERGO EUROPE  
Westervoortsedijk 60,  
6827 AT Arnhem  
The Netherlands

European Representative SRN: NL-AR-000000116

Basic UDI DI: 0853061006MAXO2ME76

Product Name(s): MaxO2ME

Intended Purpose: The MaxO2 ME oxygen monitor is intended for continuous monitoring of the concentration of oxygen being delivered to patients ranging from newborns to adults. It can be used in the hospital and sub-acute settings.

*The MaxO2 ME is not a life supporting device.*

Risk Class & GMDN: Class IIb 35219 – Analyzer, Gas, Oxygen

Classification Criteria: Clause 3.1 Rule 9 of Annex IX of MDD

This EU declaration of conformity is issued under the sole responsibility of the manufacturer. We herewith declare that the above mentioned products meet the provisions of the following EC Council Regulations, Common Specifications and Standards. All supporting documents are retained under the premises of the manufacturer and the European Representative.

Regulations: General Application Directives: Medical Device Directive, Council Directive 93/42/EEC of 14 June 1993 per Annex II Excluding Section 4 as amended by 2007/47/EC of September 2007

RoHs (EU) 2015/863

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

Number 0123

Conformity Assessment: Full Quality Assurance per Annex II MDD 93/42/EEC Excluding Section 4

EC Certificate No.: G1 045041 00 25 Rev. 00

Date CE mark was affixed: 14 May 2016

This Declaration is considered valid from the date of the signature below until May 26, 2024

Issued by Signature:  Date: May 08, 2023Name: Anvitha Anand Rao  
Position: Regulatory and Quality Product Engineering SpecialistApproved by Signature:  Date: May 8, 2023Name: Sidra Hankins  
Position: VP of QA/RA

This Declaration supersedes any preceding Declaration of Conformity for the MaxO2ME Family.

#### Applied Common Specification and Standards

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

EN ISO 13485:2016 (ISO 13485:2016)  
EN ISO 14971:2019 (ISO 14971:2019)  
EN ISO 20417:2021  
ISO 15223-1:2021  
EN 60601-1:2006/A1:2013 (IEC 60601-1:2005/A1:2012)  
ISO 80601-2-55:2011  
EN ISO 10993-1:2009 (ISO 10993-1:2009)  
IEC 60601-1-8:2006/AMD1:2012  
ISTA 2A (2011)



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EN 60601-1-2:2015 (IEC 60601-1-2:2014)  
EN 62304:2006