

## 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

Prinsessegracht 20 2514 AP The Hague The Netherlands

European Representative SRN: NL-AR-000000116

Basic UDI DI: Pending Assignment

Product Name(s): BlenderBuddy Family (BlenderBuddy, BlenderBuddy2)

Intended Purpose: The BlenderBuddy and BlenderBuddy2 are intended for use in conjunction

with a variety of air/oxygen blenders.

Risk Class & GMDN: Class IIa 44225 – Oxygen/air breathing gas mixer, hospital (Accessory)

Classification Criteria: Clause 1.2 Rule 2 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

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

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EC Certificate No.:	G1 045041 00 25 Rev. 00	
Date CE mark was affixed:	06 May 2009	
This Declaration is considered	valid from the date of the si	gnature below to May 26, 2024
Signature:	Date: _	12/28/2021
Name: Sidra Hankins Position: VP of QA/RA		
This Declaration supersedes an	y preceding Declaration of	Conformity for the BlenderBuddy Family
Applied Common Specification	and Standards	
The referenced list of Commor compliance can be provided in	1	ized standards for which documented evidence of

EN ISO 13485:2016 (ISO 13485:2016) BS EN ISO 14971:2019 (ISO 14971:2019) BS EN ISO 20417:2021 ISO 15223-1:2021