

## **DECLARATION OF CONFORMITY**

Manufacturer's Name:

Maxtec

Address:

2305 South 1070 West

Salt Lake City, Utah 84119

**USA** 

European Representative:

**QNET BV** 

Kantstraat 19

NL-5076 NP Haaren The Netherlands

Product:

MaxBlend Family Blender

Model(s):

MaxBlend2 & MaxBlend Lite (Including Pole Mount Bracket Accessories

RP05P07, RP05P09, R100P22 and R100P26)

Classification & GMDN:

Class IIb Mixer, Gas, Breathing 36327

Classification criteria:

Clause 3.1 Rule 9 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:

September 13, 2016

This declaration is considered valid from May 5, 2019 to December 18, 2021

Signature:

annif Java

Date: 2 May

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:

ISO 60601-1:2005 (3rd edition)

IEC 60601-1-2:214

IEC 60601-1-8:2007

IEC 60601-1-6:2010

CAN/CSA C22.2 NO. 60601-1:14

ISO 80601-2-55:2011

EN ISO 10993-1:2009

EN ISO 14971:2012

EN 62366:2008

ISO 15223-1:2016

EN 1014:2008

MIL-STD-810G (31 Oct. 2008)

ISO 11195:1995

ISO 15002:2008