



maxtec

p r e m i e r o 2 s o l u t i o n s

DECLARATION OF CONFORMITY

Manufacturer's Name: Maxtec
Address: 6526 South Cottonwood Street
Salt Lake City, Utah 84107
USA

European Representative: QNET BV
Hommerterweg 286
6436 AM Amstenrade
The Netherlands

Product: Flowmeter

Model(s): Blender Buddy

Classification: Class I - Medical Device Accessory (Flowmeter – 11746)

Classification criteria: Clause 1.1 Rule 1 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

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

EC Certificate No.: Not Applicable, Product Self Certified

Date CE mark was affixed: 06 May 2009

This declaration is considered valid from September 13, 2011 to August 31, 2014

Signature:  **Date:** 

Name: Tammy Lavery, RAC
Position: Director of Regulatory & Quality/Maxtec Management Representative
6526 South Cottonwood Street
Salt Lake City, UT 84107
General Tel: 801-266-5300/800-748-5355 & Fax: 801-270-5590

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