Certificate of CE-Registration

This is to certify that, in accordance with the Medical Device Directive 93/42/EEC, Medical Device Safety Service GmbH (MDSS) agrees to perform all duties and responsibilities as the Authorized Representative for:

Flotec, Inc.
Mr. John Pichon
7625 West New York Street
Indianapolis, IN 46214
UNITED STATES OF AMERICA

as stipulated and demanded by the aforementioned Directive. The German Competent Authority has allocated the medical devices of the Manufacturer registration numbers as foreseen in:

Annex A dated December 20, 2017

The Manufacturer has provided MDSS with the appropriate Declaration(s) of Conformity confirming that the medical devices fulfill the applicable requirements of Directive 93/42/EEC. In compliance with German law, a safety officer has been appointed for Germany.

2017-12-20

Joy Grimm
Senior Consultant
MDSS GmbH

Annex A dated December 20, 2017
Manufacturer: Flotec, Inc.

UMDNS Code Description Notified Medical Device Product Name & Catalogue Number	UMDNS Code	Risk Class	Cat. Code	Registration Number	NB No. / NB Certificate No.	NB Cert. Valid Until YYYY-MM-DD
Flowmeters, Gas	11-748	IIb	02	DE/CA09/0170/F07/002	0843/717.171028	2022-10-27
Flowmeters						
FXXX-XXXXX Flowmeters						
Regulators, High-Pressure Gas	13-323	IIb	02	DE/CA09/0170/F07/001	0843/717.171028	2022-10-27
Regulators; Ingage Regulators; Mini Regulators						
RXXXX-XXXXX Regulators *DXXXX-XXXXX Ingage Regulators* *MXXXX-XXXXX Mini Regulators*						