EC Certificate Full Quality Assurance System: Certificate CN19/41069

The management system of

Mespere LifeSciences Inc.

180 Frobisher Dr, Unit 1C, Waterloo, Ontario, N2V 2A2, Canada

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Non Invasive Central Venous Pressure Monitor (Model: Mespere-Venus-2000) for assessment of central venous blood pressure (CVP);

Non Invasive Venous Oximetry System (Model: Mespere VO 100) for assessment of venous blood oxygenation;

Non Invasive NeurOs Cerebral Oximetry System (Model: Mespere VO200) for assessment of Cerebral tissue oxygenation

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 16 December 2019 until 29 June 2023
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 29 June 2012
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered CN/SZX 14039

Authorised by

SGS Belgium NV, Notified Body 1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com