EU DECLARATION OF CONFORMITY

Manufacturer: **Merivaara Corp.**
Puustellintie 2
15150 Lahti, FINLAND

This Declaration of conformity is issued under the sole responsibility of the manufacturer.

Objects of the Declaration:

Trade name of Electrical Medical Device:	**Product code(s):**
Practico	100030000, 145000
Promerix	100060000
Merimote	100060860
OpenOR™	200040100, 200040101, 200040102, 200040103 200041000, 200041001, 200042000 200040010, 200040020, 200043000
Q-Flow	520210, 520220, 520222, 520230, 520211 (incl. Camera option)

The objects of the declaration described above are in conformity of Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the Restriction of the use of certain Hazardous Substances (RoHS) in electrical and electronic equipment. Conformity is declared based on harmonized standard:

EN 50581:2012 Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances

Signed for and on behalf of Merivaara Corp.:

Lahti 21st of August 2019

Jari Kalja, Chief Executive Officer

Digitally signed by Ceaicovschi Tudor
Date: 2021.11.03 14:18:31 EET
Reason: MoldSign Signature
Location: Moldova