

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