Electromagnetic disturbances – Requirements and tests	
To comply with its intended use, the limitations and restrictions regarding the electromagnetic environment in which the device may be used are specified below. The customer and/or user of the device should ensure that these requirements are respected.	
Statement for the operational environments	The device may be used in a domestic healthcare environment EXCEPT near active HF surgical equipment and the RF shielded room of an ME system for magnetic resonance imaging, where the intensity of EM disturbances is high.
Performance features of the ME system that have been determined to be essential to the performance of the device	Vibration of the motor contained in the device.
WARNINGS	Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally. Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this device could result in increased electromagnetic emissions or decreased electromagnetic immunity of this device and result in improper operation. Portable RF communications equipment (including peripherals such an antenna cables and external antennas) should be used no closer than 30cm (12 inches) to any part of the bruXane® 2go including cables specified by the manufacturer. Otherwise, degradation of the performance of this device could result.
A list of all cables, transducers and other accessories that are relevant for EMC compliance	Not applicable
Test	**Compliance**
RF emissions CISPR11	Group 1, Class B
Harmonic Emissions	Not applicable
Voltage Fluctuations/Flicker	Not applicable