

## UE DECLARATION OF CONFORMITY

This is a declaration made in compliance with the general safety and performance requirements set out in Annex I of MDR 2017/745

This declaration of conformity is issued under the sole responsibility of the manufacturer:

OT Bioelettronica S.R.L

Object of the declaration, cl. I rule 1 Annex VIII:

### EMG DRY ELECTRODE ARRAYS

Basic UDI: 805697785ELECARRAY009C6

SA 16/10	Ref.: OT0022
SA 16/5	Ref.: OT0023
SA 16/2.5	Ref.: OT0024
DAP05MM1501	Ref.: OT0250
DAM05MM1601	Ref.: OT0251

The object of the declaration above fulfills the general safety and performance requirements of Annex I of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC. The object above is in compliance with Directive 2011/65/EC of the European Parliament and of the Council of 8 June 2011, on the restriction of certain hazardous substances in electrical and electronic equipment, and Directive 2015/863 of 31 March 2015 laying down modification of Annex II of Directive 2011/65/EC, ISO 10993-1:2008 Biological evaluation of medical devices and ISO 14971:2019 Medical devices - application of risk management to medical devices.

Turin, 04/11/2024

Andrea Bottin (CEO)

