4. EC Declaration of Conformity

according to the following guidelines:

Electromagnetic compatibility (EMC) (2004/108/EC)
RoHS substance bans (2011/65/EU)
WEEE Waste Electrical Equipment Disposal (2002/96/EG & 2008/34/EG)

The manufacturer / Distributor / Authorized representative

Mindfield Biosystems Ltd.
Hindenburgring 4
D-48599 Gronau
Germany

WEEE-Reg.-Nr. DE 24465971

hereby declares that the following product:

“Mindfield® eSense Respiration” in combination with the Mindfield® eSense Skin Response”

complies with the provisions of the Directives identified above, including their amendments in force at the time of the declaration.

The following harmonized standards have been applied:

DIN EN 60950-1 Information technology equipment – Safety – Part 1: General Requirements (2011-01)

DIN EN 55022 Information technology equipment – Radio disturbance characteristics (2008-05)

DIN EN 55024 Information technology equipment – Immunity characteristics (2011-09)

Place: Gronau
Date: 08. May 2019
Niko Rockensüß, Managing Director