

EC Declaration of Conformity

Manufacturer:

Wuhan Greentek Pty Ltd.
Room 03-2, Floor 3, Building 3, Phase III,
International Enterprise Center, Special
No.1, Guanggu Avenue, East Lake High-
tech Development Zone, Wuhan, China.
Tel: 86-027-88185488
E-mail: greentek@gtsensor.com

whose single Authorized Representative:

Luxus Lebenswelt GmbH
Kochstr.1, 47877, Willich, Germany
DIMDI: DE/0000047791
Lin Sun
Tel: 0049- 1715605732
E-mail: info.m@luxuslw.de

We, the manufacturer, herewith declare that the products

Name of device: EEG Electrode Cap; Medical EEG Electrode Cap; EEG Electrode Headset; EEG-Recording Cap; EEG Starter Kit; Gelfree Electrode Cap; Semi-Dry EEG electrode Cap

Name of device: EEG Caps; Electrode Caps

UMDN CODE: 17554

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class I according to Annex IX of the Directive 93/42/EEC. It bears the mark



following the procedure relating to the EC Declaration of Conformity set out in Annex VII of Directive 93/42/EEC.

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration.

The above mentioned declaration of conformity is exclusively under the responsibility of

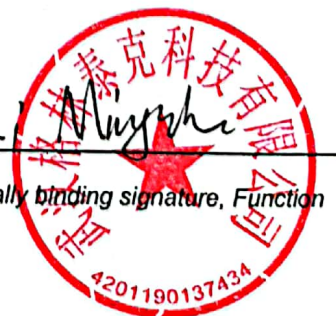
Wuhan Greentek Pty Ltd.

Room 03-2, Floor 3, Building 3, Phase III, International Enterprise Center, Special No.1, Guanggu Avenue, East Lake High-tech Development Zone, Wuhan, China.

January 15, 2021 Wuhan, China

Place, date

Li Mingsha
Legally binding signature, Function



EC Declaration of Conformity