"EU" DECLARATION OF CONFORMITY

Manufacturer: **Allmobility Trading S.R.L.**

Operative headquarters: Via G. Balla, 4 42124 Reggio Emilia (RE) – ITALY

Registered head office: Via Mentana, 150 43100 Parma (PR) – ITALY

Tel. +39 0522 941078

E-mail: info@allmobility.it

Web: www.allmobility.it

EUDAMED SRN IT-MF-000021947

Medical device: **"Allmobility BUFFALO™ 150" BOARD**

Class: **I** (Annex VIII, Rule 1, MDR 2017/745)

Basic UDI-DI: **805660018ALL03BUFCU**

Applicable Directives: 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

We hereby declare, under our sole responsibility, that the products listed above meet all applicable general safety and performance requirements of Annex I of Regulation (EU) 2017/745 concerning Medical Devices.

The Manufacturer undertakes to keep and make available to the competent authorities the technical documentation specified in Annex II of Regulation (EU) 2017/745, for a period of 10 years from the date on which the product was last manufactured.

REGGIO EMILIA – ITALY (IT) - 17 / 05 / 2021

The Legal Representative

(writer and signatory of this declaration)

(Maurizio Cassinadri)

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