DECLARATION OF CONFORMITY EC

No. 55/D/EN/4

MANUFACTURER

REH4MAT

PRODUCT

WEDGE

MODEL: KLW-01; KLA-Z; KLA-B; KLO-Z; KLO-B; WEDGE KLW-02; WEDGE KLW-03; WEDGE KLW-04; WEDGE KLW-05; WEDGE KLW-06; WEDGE KLW-07; WEDGE KLW-08; WEDGE KLW-09; WEDGE KLW-10; WEDGE KR-01; WEDGE KR-02; WEDGE KR-03; WEDGE KR-04; WEDGE KR-05; WEDGE KR-06; WEDGE KR-07; WEDGE KR-08; WEDGE KR-09; WEDGE KR-10

marked with the CE sign is Class I medical device, of rule 1, consistent with the requirements of the **Council Directive 93/42/EEC of 14 June 1993 on medical devices** and the **Law of 20 May 2010 on medical devices** as amended. Conformity assessment was carried out on the basis of the **VII Council Directive 93/42/EEC**.

The product meets the requirements of the harmonized standards

PN-EN 1041+A1:2013-12
Information supplied by the manufacturer of medical devices

PN-EN 12182:2012
Assistive products for persons with disability. General requirements and test methods

PN-EN ISO 13485:2016
Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

PN-EN ISO 14971:2012
Medical devices. Application of risk management to medical device

PN - EN ISO 15223-1:2017-02
Medical devices Symbols to be used with medical device
labels, labelling and information to be supplied - Part 1:General requirements

THIS DECLARATION HAS BEEN ISSUED ON THE RESPONSIBILITY
OF THE MANUFACTURER
REH4MAT, 36-060 Głogów Małopolski, ul. Piaski 47, Poland

QUALITY SYSTEM CERTIFICATION DNV·GL ISO 13485

COMPANY OWNER
Sławomir Wroński

Głogów Małopolski 27.09.2019