

DECLARATION OF CONFORMITY

COMPANY
REH4MAT Sławomir Wroński

DECLARES

That product:

REFERENCE NUMBER: **EB-SK/2RA**

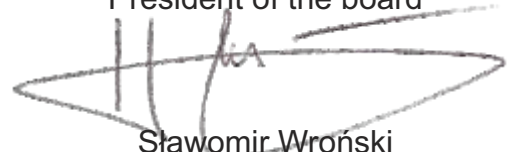
It conforms the requirements of the Council Directive 93/42/EEC of 14 June 1993 (**Annex VII**), as amended by Medical Device Directive 2007/47/EEC regarding medical products **belong to the 1 Class Of Medical Devices without measuring function, non-sterile** and associated with it norms:

- | | |
|------------------------|--|
| PN-EN 1041:2010 | - 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 13485:2012 | - Medical devices - Quality management systems - Requirements for regulatory purposes |
| PN-EN ISO 14971:2011 | - Medical products - Application of risk management to medical products |
| PN-EN ISO 15223-1:2012 | - Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General requirements |

and with the Act of Law of 20 May 2010 on Medical Devices
(Dz.U. 2010 nr 107 poz. 679)

The Company REH4MAT SŁAWOMIR WRÓŃSKI, 36-660 Głogów Małopolski, ul. Piaski 47
takes full responsibility for issuing the herby declaration.
Declaration is in compliance with the attachment No VII

President of the board



Sławomir Wroński