SUZHOU KD INTELLIGENT DEVICE CO., LTD.
NO.36 Gugang Road, Chengxiang Town, Taicang City, Jiangsu P.R. of China
TEL: +86-512-53110088 FAX: +86-512-53110099

Declaration of Conformity

Manufacturer : **Suzhou KD Intelligent Device Co., Ltd. located in No.36 Gugang Road, Taicang City, Jlangsu Province, China. Actor ID /SRN :CN-MF-000016513**

European Representative : Distributeur France Acekare
7 rue de Mireport 33310 LORMONT - 09 80 80 8515
SRN: FR-AR-000011114

Product Name: **Power wheel chair**

Type: PL001-1002, PL001-4001, PL001-2004-1, PL001-2009, PL001-2010, PL001-2009-2, PL001-2010-2,PL001-2003K, PL001-3005, PL001-2004, PL001-5001, PL001-5001-2, PL001-6001, PL001-6002 ,PL001-7001, PL001-7002, PL001-7003, PL001-7005, PL001-8001,PL001-8002, PL001-6003,PL001-6004

UMDNS Code:16214

Medical Device Classification : **Class I**

Model No.:	
PL001-1002	Basic UDI –DI:697057522100288
PL001-4001	Basic UDI –DI:69705752240018T
PL001-2004-1	Basic UDI –DI:6970575222004-1PV
PL001-2009	Basic UDI –DI:69705752220098V
PL001-2010	Basic UDI –DI:69705752220108E
PL001-2009-2	Basic UDI –DI:6970575222009-2QQ
PL001-2010-2	Basic UDI –DI:6970575222010-2PJ
PL001-2003K	Basic UDI –DI:6970575222003KZ9
PL001-3005	Basic UDI –DI:69705752230058U
PL001-2004	Basic UDI –DI:69705752220048K
PL001-5001	Basic UDI –DI:697057522500192
PL001-5001-2	Basic UDI –DI:6970575225001-2QP
PL001-6001	Basic UDI –DI:697057522600199
PL001-6002	Basic UDI –DI:69705752260029B
PL001-7001	Basic UDI –DI:69705752270019G
PL001-7002	Basic UDI –DI:69705752270029J
PL001-7003	Basic UDI –DI:69705752270039L
PL001-7005	Basic UDI –DI:69705752270059Q
PL001-8001	Basic UDI –DI:69705752280019P

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PL001-8002	Basic UDI –DI:69705752280029R
PL001-6003	Basic UDI –DI: 69705752260039D
PL001-6004	Basic UDI –DI: 69705752260049F

Conformity Assessment Route: **ANNEX XIII of Regulation (EU) 2017/745 : CLASSIFICATION RULES，NON-INVASIVE DEVICES： Rule1**

We herewith declare that the above mentioned products meet the provisions of the following EC Council regulations and Standards. All supporting documentations are retained under the premises of the manufacturer.

REGULATIONS

General applicable regulations:

Regulation (EU) 2017/745 of the European Parliament and the Council of 5 April 2017 on Medical devices (MDR)

EN 12184:2009; ISO7176; ISO9999:2011, EN60601-1-2：2016

Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS) and amendments.

Taicang,Suzhou,China

Jan,09,2023

Signature: yiwen zhang

Name: **Yiwen Zhang**

Position: **General Manager**

EC Declaration of Conformity
SZKD CE-01(A/0)