

## EC Declaration of Conformity

Manufacturers Registered Name: DOOWONID Co., Ltd.

1-dong, 44, Techno 8-ro, Yuseong-gu, Daejeon, Korea

Tel: +82-42-721-3644

EC Representative: MedNet GmbH

Borkstrasse 10, 48163 Muenster, Germany T) +49-251-32266-61 F) +49-251-32266-22

## **Declares that the product**

**Brand Name**: Screw Driver **Device Name**: Ball Driver

**Model Name**: Total 5 models: 2type - Hex(3), Torx(2)

**GMDN Code**: 33968

Relevant EC Directives: Medical Device Directive 93/42/EEC amended by 2007/47/EC

**Classification**: Class I (Rule 1)

**Conformity Assessment Route**: Annex VI

We herewith declare that the above mentioned products meet the provisions of the Council Directive 93/42/EEC as amended by Directive 2007/47/EC for medical devices. All supporting documentation is retained under the premises of the manufacturer.

In conformity with the national standard transposing harmonised standards; See Attachment

Date: 2017. 02. 24

Doo-hyun Baek / C.E.O

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## [ List of Harmonized Standards ]

No.	Category	Title	Description
1	System	Directive 2007/47/EC	of 5 September 2007 concerning medical devices
		EN ISO 14971 : 2012	Medical devices - Application of risk management to medical devices
		MEDDEV2.12-1 Rev.8	Guidelines on a medical device vigilance system
2	ETC	MEDDEV2.4/1 Rev.9 June 2010	Guidelines relating to the application of the council directive 93/42/EEC on medical devices
		ISO 15223-1 : 2012	Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied
		EN 1041: 2008	Information supplied by the manufacturer with medical devices
3	Clinical investigation	ISO14155 : 2011	Clinical investigation of medical devices for human subjects – Good clinical practice
		MEDDEV2.7.1 Rev.3 December 2009	Evaluation of clinical data
		MEDDEV2.12-2 Rev.2 January 2012	Post market clinical follow up