DREAMCAN
深圳德睿康科技有限公司
Shenzhen DreamCan Technology Co.,Ltd

EU DECLARATION OF CONFORMITY

This Declaration of Conformity, issued under the sole responsibility of the manufacturer **Shenzhen DreamCan Technology Co., Ltd, 301/B3 Huaqiang Industrial Park, Qingfeng Ave., Baolong, Longgang District, Shenzhen, Guangdong, China** hereby declaring the following Personal Protective Equipment (PPE) & Medical Device (MDD)

Product Description: Particle Filtering Half Mask & Disposable Medical Mask
Product Model / Brand: DP-A-22L-K / DreamCan
Products Colour/Type: White, Non-reusable, Molded
Product Classification: FFP2 NR & TYPE IIR

is in conformity with the provisions of the following European Regulation:

PPE (Personal Protective Equipment) Regulation

The model is in conformity with the provisions of Regulation (EU) 2016/425, including fulfilment of the applicable essential health and safety requirements set out in Annex II, and with the National Standard transposing the harmonised European Standard Number (s):

EN149:2001+A1:2009

and is identical to the PPE which is the subject of EU type-examination (Module B of Regulation (EU) 2016/425) referenced on the certificate number:

Module B Certificate Number: CE-PC-200401-177-01-9B (Issue Date: 29/05/2020)

and is subject to the type based on internal production control plus supervised product checks at random intervals set out in Module C2 of Regulation (EU) 2016/425:

Module C2 Certificate Number: CE-PC-200325-106-FPC-C (Issue Date: 06/05/2020)

Both Module B & Module C2 mentioned above are issued by:
CCQS Certification Services Limited –Block 1 Blanchardstown Corporate Park,
Ballycoolin Road, Blanchardstown, Dublin15, D15 AKK1, Ireland, Notified Body No. 2834.

Medical Devices Directive

The model also is complied with the transposition into national law, the provision of Council Directive 93/42/EEC concerning medical devices; including, at 21 March 2010, the amendments by Council Directive 2007/47/EEC.

Relevant EC Directive: Medical Device Directive 93/42/EEC

and according to Annex IX rules of Medical Device Directive is classified as Class 1 device.
And is in conformity with the provisions of Annex VII and all other applicable provisions of
Council Directive 93/42/EEC (Medical Device) including the National Standard transposing the harmonised European Standard Number(s)

EN14683:2019+AC:2019 Type IIR

EU authorised Representative
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SIGNED BY: WALTER WANG
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