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-24L-K / DreamCan**
**Products Colour/Type: White, Non-reusable, Molded**
**Product Classification: FFP3 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: 0370-4263-PPE/B (Issue Date: 27/08/2020)**

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:0370-4442-PPE/C2 (Issue Date: 18/09/2020)**

Both Module B & Module C2 mentioned above are issued by:
LGAI Technological Center, S.A. (APPLUS), Campus UAB – Ronda de la Font del Carme, s/n, 08193, Bellaterra, Barcelona, Notified Body No. 0370

**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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