

Declaration of Conformity

The manufacturer, or his authorized representative in the Community:

Company: **Profit Royal Pharmaceutical Limited**

Address: **RM 1211, 12/F, Sunbeam Centre, 27 Shing Yip Street, Kwun Tong, Kowloon, HONG KONG.**

Declares that our Surgical Masks described hereafter

Product Name: **NASK FFP2 Nanofiber SMART Mask (NASK 納米纖維智能殺菌口罩)**

Product Description: **Splash Resistance. Particle Filtration Efficiency (PFE).
Bacterial Filtration Efficiency (BFE). Differential Pressure (Breathability).
Flammability. Microbial Cleanliness (Bioburden). Biocompatibility.
Antimicrobial (Bacteria Killing). Viral Filtration Efficiency (VFE).**

Product Code: **M0011**

is in conformity with the provisions of **CE European Medical Device Regulation (MDR) EU 2017 / 745** (replaced Directive 93 / 42 / EEC) and, where such is the case, the national standard transposing harmonized standard

Standard: CE European EN 14683:2014 Medical face masks – Requirements and test methods

Risk Classification: Class I Medical Device

Functional level: Meeting TYPE II R

Tested by: US Nelson LABs (nelsonlabs.com) and HK SGS (sgs.com)

Signed: Henry Chow 

Position: Quality Manager

Date: 2020-11-03

