弓立（厦门）医疗用品有限公司
SPRO MEDICAL PRODUCTS (XIAMEN) CO.,LTD.

EU DECLARATION OF CONFORMITY

No: DOC

1. PPE (product, type, batch or serial number): GL001A/GL001

2. Name and address of the manufacturer: SPRO MEDICAL PRODUCTS(XIAMEN)CO.,LTD
West of 1-5th Floor, No.139 Factory Bldg,
TongAn Garden,TongAn Industry Area Xiamen
City, Fujian Province China

3. This declaration of conformity is issued under the sole responsibility of the manufacturer.

4. Object of the declaration (identification of PPE allowing traceability; where necessary for the identification of the PPE, a colour image of sufficient clarity may be included):
Disposable Respirator

5. The object of the declaration described in point 4 is in conformity with the relevant Union harmonisation legislation: Regulation (EU) 2016/425

6. References to the relevant harmonised standards:
EN 149:2001+A1:2009 Respiratory protective devices – Filtering half masks to protect against particles – Requirements, testing, marking.

7. The notified body INSPEC (NB:0194) performed the EU type-examination (Module B) and issued the EU type-examination certificate: PPE20161886

8. The PPE is subject to the conformity assessment procedure: conformity to type based on quality assurance of the production process (Module D) under surveillance of the notified body INSPEC (NB:0194).

9. Signed for and on behalf of:

Company: SPRO MEDICAL PRODUCTS(XIAMEN)CO.,LTD

General Manager: WONG SUET YI

Signature

Place & Date: Xiamen City ,Fujian Province,
CHINA 03st September, 2020

CE