Verification OF CONFORMITY

Company Name	[redacted]
Company Address	
Related Regulation	**: Directive Directive 93/42/EEC**
Standards	**: EN 14683:2019**
Product Name	**: Single-use medical face mask**
Classification:	**: Class I (Non-sterile)**
Model	**: Medical mask, Kn95 defensive mask, Gauze mask, Gas mask**

The certification is valid in accordance with the test report MTMR0320-MDD and when the product is manufactured in accordance with the tested sample.

In the Directive on Medical Devices, the medical device is classified in accordance with the provisions of Annex IX of the MDD as class I (low risk), class IIa or IIb (medium risk) or class III (high risk). The involvement of a Notified Body is not necessary for medical devices of class I
After inspection of the technical documentation issued by the customer, and in his request, we express our opinion that the product meets the technical requirement of the following directives:
Directive Directive 93/42/EEC

CE

The label of the CE Marking on the left side should be not less than 5mm height. CE Marking and EC Declaration of Conformity are duties for the manufacturer or its applicant who puts the product on the market. This one is responsible to start the CE marking and certification procedure as required by the legislation in force. Only forthe products which are compulsorily included into specific Directives or Regulations will be necessary to appoint a Notified Body.

Certificate NO. JZ86200323565

Issued Date: 23 Mar.. 2020

General Manager Kerol

SH JIANZHENGWANG LTD. JZ-CERT. SH OFFICE