

## **EC** Certificate Directive 93/42/EEC Annex V **Production Quality Assurance** Medical Devices

Registration No.: DD 60133273 0001

Report No.:

15085900 005

Manufacturer:

**Xiantao Xingrong Protective** 

Products Co., Ltd.

No. 46, East of Pengchang Road,

433018 Xiantao, Hubei

China

**Products:** 

Aspects of manufacture concerned with securing and maintaining sterile conditions of Face Masks, Surgical

Gowns, Non-woven Caps, Non-woven Shoe Covers,

Plastic Shoe Covers, Coveralls

Replaces Approval, Registration No.: DD 60104282 0001

Notified Body

Zhong

**Expiry Date:** 

2023-12-05

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

**Effective Date:** 

2018-12-06

Date:

2018-12-06

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.