EC Certificate

TÜVRheinland®

Production Quality Assurance
Directive 93/42/EEC on Medical Devices, Annex V

Registration No.: DD 2070881-1

Digitally signed by Jighili Tatiana
Date: 2022.06.30 17:01:03 EEST
Reason: MoldSign Signature
Location: Moldova

Manufacturer: M & G Products Co., Ltd.
No. 968-970 Mingzhuwan,
Yangzhong
212200 Jiangsu
P.R. China

Products: Electronic Thermometers, Sterile Infusion sets for Single Use, Sterile Syringes for Single Use, Transfusion Sets, Hypodermic Needle for Single Use, Three way Stopcock, Sterile Heparin Caps for Single Use, Sterile Dental Needles for Single Use, I.V. Cannula for Single Use, Infusion Set with Burette, Extension Tubes, Syringe for Insulin, Scalp Vein set for Single use, I.V.flow regulator for Single use, Sterile Nelaton Catheters, Insulin Pen Needles, Sterile and Non-sterile Oxygen Masks, Sterile and Non-sterile Nasal Oxygen Cannulas, Sterile and Non-sterile Nebulizer Masks, Sterile and non-sterile venture masks, Sterile and non-sterile non Rebreathing Masks, Sterile and Non-sterile Nebulizers, Suction Tubing With Yankauer Handle, Needle Free Connectors, Tracheal Tubes, Reinforced Endotracheal Tubes, Sterile Surgical Blades, Disposable Scalpel with Plastic Handles, Lap Sponges;
Aspects of manufacture concerned with securing and maintaining sterile conditions of Sterile Suction Catheters, Sterile Feeding Tubes, Disposable Stomach Tubes,

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.

Report No.: 15080731 011

Effective date: 2021-04-30

Expiry date: 2024-05-26

Issue date: 2021-04-30

TÜV Rheinland LGA Products GmbH Zertifizierstelle TUVRheinland

Herbert Zhong
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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

EC Certificate

TÜVRheinland®

Production Quality Assurance
Directive 93/42/EEC on Medical Devices, Annex V

Registration No.: DD 2070881-1

Manufacturer:
M & G Products Co., Ltd.
No. 968-970 Mingzhuwan,
Yangzhong
212200 Jiangsu
P.R. China

Disposable Umbilical Cord Clamps, Disposable Mucus Extractors, Disposable Rectal Tubes, Disposable I.D. Bracelets, Urinary Collection Bags, Sterile Gauze Sponges, Sterile Bandages, Sterile Gauze Rolls, Sterile Cohesive Bandages, Sterile Wound Plaster/Strips, Sterile Dressings Adhesives

Replaces Approval, Registration No.: DD 60148986 0001

Report No.: 15080731 011

Effective date: 2021-04-30

Expiry date: 2024-05-26

Issue date: 2021-04-30

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle TÜVRheinland®

Herbert Zhong
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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

EC Certificate

TÜVRheinland®

Production Quality Assurance
Directive 93/42/EEC on Medical Devices, Annex V

Registration No.: DD 2070881-1

Manufacturer:
M & G Products Co., Ltd.
No. 968-970 Mingzhuwan,
Yangzhong
212200 Jiangsu
P.R. China

The scope of certification includes the following manufacturing sites:

No.	Location	Product groups manufactured
/01	M & G Products Co., Ltd. Binjiangdadao, Huangshantao, Lianhe Xinba Town, Yangzhong 212200 Jiangsu P.R. China	Same as the above mentioned Products

Report No.: 15080731 011

Effective date: 2021-04-30

Expiry date: 2024-05-26

Issue date: 2021-04-30

TÜV Rheinland LGA Products GmbH Zertifizierstelle TUVRheinland®

Herbert Zhong
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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