EC Certificate

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

TÜVRheinland®

Registration No.:	DD 60090679 0001
Report No.:	15064931 001

Manufacturer: Shanghai K&G International Co., Ltd.
Room 604, 1619 Dalian Road
200086 Shanghai
China

Products: Medical Devices

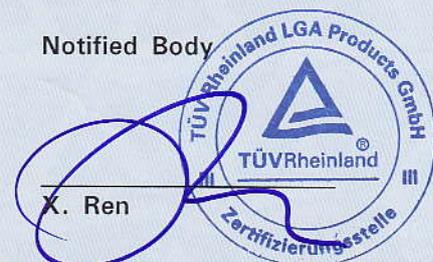(see attachment for products included)

Expiry Date: 2018-12-10

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.

		Notified Body
Effective Date:	2013-12-11	
Date:	2013-12-11	X. Ren

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.

TÜVRheinland ®

TÜV Rheinland

LGA Products GmbH

Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.:   DD 60090679 0001
Report No.:   15064931 001

Manufacturer: Shanghai K&G International Co., Ltd.
Room 604, 1619 Dalian Road
200086 Shanghai
China

Products:

Syringes, Sterile Hypodermic Needles for Single Use, Scalp
Vein Sets, Blood Transfusion Sets, Disposable Infusion Sets,
Sterile Surgical Gloves, Sterile Urethral Catheters for
Single Use, Sterile Surgical Blades, Disposable Scalpels
with Plastic Handle, Sterile Suction Tubes, Blood
Collection Needles, Stomach Tubes, Endotracheal Tubes
with and without cuff;

Aspects of manufacture concerned with securing and
maintaining sterile conditions of Urine Drainage Bags,
Vaginal Speculum

Notified Body

X. Ren

Date: 2013-12-11