

EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4 Full Quality Assurance System **Medical Devices**

Registration No.: HD 60112688 0001

Report No.:

15096014 001

Manufacturer:

Shanghai SA Medical & Plastic

Instruments Co., Ltd. No. 18, Hengxi Road Pujiang Industrial Zone

Minhang District Shanghai 201114

China

Products:

Spinal Needles, Epidural Needles, Nerve Block Needles, Bone Marrow Needles, Introducer Needles

Replaces Approval, Registration No.: HD 60087230 0001

Expiry Date:

2021-08-14

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 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 II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date:

2016-08-15

Date:

2016-08-08

TÜV Rheinland LGA Products GmbH - Tillystraße 90431 Nürnberg

Notified Bo

K. Ren

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



Certificate

The Certification Body of TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

Shanghai SA Medical & Plastic Instruments Co., Ltd. No. 18, Hengxi Road Pujiang Industrial Zone Minhang District Shanghai 201114 China

has established and applies a quality management system for the following scope:

Design and Development, Manufacture and Distribution of Needles for Medical Use (This certificate information can be searched on CNCA official website http://www.cnca.gov.cn)

Proof has been furnished that the requirements specified in

EN ISO 9001:2008

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date:

2016-08-08

Certificate Registration No.:

SY 60112690 0001

An audit was performed. Report No.: 15096015 001

This Certificate is valid until:

2018-09-14

Date 2016-08-08

Certification Body Rheinla K. Ren

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