EC Certificate

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

TÜVRheinland®

Registration No.: HD 60101875 0001

Report No.: 15042093 004

Manufacturer:
Shanghai Angel Electronic
Equipment Co., Ltd.
4/F, Bldg. 5, No. 1135
Bao Tou Rd., Yang Pu District
Shanghai 200438
China

Products:
Infusion Pumps, Syringe Pumps, Patient-controlled Analgesia
Infusion Pumps, Enteral Nutrition Pumps

Replaces Approval, Registration No.: HD 60038763 0001

Expiry Date: 2016-06-10

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: 2015-05-22

Date: 2015-05-22

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Notified Body

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.