



Product Service

# EC-CERTIFICATE

## Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 04 65725 008

**Manufacturer:****China Shikang Medical Equipment Group Limited**

Room 305, Yuehuyinzu

No.66 East Road

Nanzhan, Ningbo, CHINA

**EC-Representative:****Shanghai International Holding  
Corp. GmbH (Europe)**

Eiffestraße 80

20537 Hamburg

GERMANY

**Product  
Category(ies):****Anaesthetic Workstation, Vaporizer,  
Ventilator, Hummer Medical Air Compressor,  
Infusion pump, Ceiling Pendant, Operating Lamp,  
Operating Table, Autoclave, Infant Incubator, Hospital Bed**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

**Report No.:**

BJ1085907

**Valid until:**

2015-05-03

**Date,** 2010-05-18

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.