EC Certificate

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

Registration No.: DD 60145018 0001

Report No.: 16804044 007

Manufacturer:
Henan Kangdi Medical Devices
Co., Ltd.
No.4, 2nd Area, SME Pioneer Park
Industry Cluster Zone, Huaiyang County
Zhoukou City
466700 Henan
China

Products:
Heat Pads

(see attachment for site included)

Replace Approval, Registration No.:DD 60097951 0001

Expiry Date: 2024-05-27

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.

Effective Date: 2019-12-17

Date: 2019-12-17

Notified Body

Jing Zhang

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.