EC Declaration of Conformity

Manufacturer:

Wenzhou Bokang Instruments Co., Ltd.
Haining Road, Haibin, Longwan, Wenzhou 325024, China
Tel:0086-577-86876969

whose single Authorized Representative:

Shanghai International Trading Corp. GmbH (Hamburg)
Add: Eiffestrasse 80, 20537 Hamburg，Germany
Tel: 0049-40-2513175
Fax: 0049-40-255726

We, the manufacturer, herewith declare that the products

Electronic-sphygmomanometer

*(*BK6001,BK6002, BK6021,BK6022,BK6023,BK6031，BK6032，BK6003
BK1016，BK1018，BK1016B,BK1016L)
UMDNS-Code: 16157*; GMDN-Code/Preferred Terms:* 45617，47489

meet the provisions of Directive 93/42/EEC which apply to them.
The medical device has been assigned to class **II**a according to Annex IX of the Directive 93/42/EEC.
It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to Annex V of Directive 93/42/EEC.
Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: DD 60128509 0001
Issue date: 16.04.2018
Expiry date: 27.04.2023

following the procedure relating to the EC Declaration of Conformity set out in Annex V of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: **Wenzhou Bokang Instruments Co., Ltd.**
Address: **Haining Road, Haibin, Longwan, Wenzhou 325024, China**

WENZHOU, CHINA 2018.04.16
Place, date

温州博康医疗科技有限公司
WENZHOU BOKANG INSTRUMENTS CO.,LTD.

President
Legally binding signature, Function