Declaration of Conformity CE

Manufacturer:
Zhejiang Bangli Medical Products Co.,Ltd
No118,Yuegui South Road,City West New District,321300 Yongkang City, Zhejiang Province, China

whose single Authorized Representative:
ZOG EAR INDUSTRIES Co.,Ltd
145-157ST JOHN STREET,
LONDON ,EGLAND,ECN 4PW

We, the manufacturer, herewith declare that the products

CAPSICUM PLASTER
POLYETHILENE PLASTER
CORN REMOVAL PLASTER
ZINC OXIDE
SURGICAL TAPE
WOUND PLASTER
NON WOVEN

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to **Class I , rule1** according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE

The product concerned has been manufactured under a quality management system according to AnnexVII 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 Product Safety GmbH
Am Grauen Stein D-51105 Köln
Certificate No.:DD 600975010001
Issue date: 26.11.2014
Expiry date: 25.11.2019

following the procedure relating to the EC Declaration of Conformity set out in Annex VII coupled with Annex V of Directive 93/42/EEC.
Application of the abovementioned Annexes and the intervention by the Notified Body is limited to: the aspects of manufacture concerned with securing and maintaining sterile conditions.
The above mentioned declaration of conformity is exclusively under the responsibility of

Zhejiang Bangli Medical Products Co.LTD

浙江邦立医药用品有限公司
ZHEJIANG BANGLI MEDICAL PRODUCTS CO.,LTD