## 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

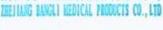
Expiry date: 25.11.2019

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 #立該药用品有限公司



重功意

EC Declaration of Conformity ECDC-001 ver: 1