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EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **Guangdong Baihe Medical Technology Co., Ltd.
No.89, Taoyuan East Road, Nanhai District, Foshan City,
Guangdong, China**

We declare under our sole responsibility that

the medical device: **Disposable silicone foley catheters**

of class: **II b**

meets the provisions of the directive 93/42/EEC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure: / **Directive 93/42/EEC Annex II, excluding Section 4**

Registration No.: **HD 60149301 0001**

Notified Body: **TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197**

Foshan, 2020-10-10

John. Cen, Management Representative

2020.10.10
Place, date /

