



Verification

Of Conformity with European Directives

Product: Stretcher
Type: YXH-1, YXH-2, YXH-3, YXH-4, YXH-5
Applicant: Zhangjiagang Xiehe Medical Apparatus and Instruments Co., Ltd.
Address: Fuqian Development Zone, Yangshe Town, Zhangjiagang City, Jiangsu Province

The submitted sample of the above product has been tested for CE marking according to the following European Directives:
93/42/EEC Medical Device Directive (including 2007/47/EC)

Standard(s) used for showing compliance with the essential requirements in the specified directive(s):

Standard(s): **EN 1865:1999**
Test report(s): **No. 2007J20—60—108125**
Issued by: **National Center of Testing Technology Shanghai**
Dates: **2007.8.10**

The referred report(s) show that the product complies with standard(s) recognized as giving presumption of compliance with the essential requirements in the specified EU Directive(s).
The CE marking as shown below may be affixed on the product.

Certificate No. **No. 01122**
Date: **23 Aug 2007**
Reissue date: **5 Jan 2013**
Valid until: **4 Jan 2017**
NQA

General manager (Signature)



NQA is a trading division of NICEIC Group Ltd, Registration No. 02513162. Registered Office: Warwick House, Houghton Hall Park, Houghton Regis, Dunstable, Bedfordshire, LU5 5ZX.
This certificate is the property of NQA and must be returned on request.

HPRA Wholesalers authorisation No. W00426/00001

Tel: +353 (0) 1 835 2411

Email: sales@medguard.ie

Unit 28B Ashbourne Business Centre, Ashbourne, Co Meath, A84 WA49

Certificate of Registration