

EC DECLARATION OF CONFORMITY

We, as the Manufacturer, certifies that the following medical device:

LATEX TOURNIQUET, LATEX FREE TOURNIQUET, LATEX FINGER COT
Classification: Classified as class I according to Annex IX, rule 1 of the
Directive 93/42/EEC

meets all applicable requirements of the Medical Devices Directives

Directive Name / Number
93 / 42 / EEC and 47/2007/EC

The declaration is sole responsibility of the manufacturer

Name of manufacturer
Jiangsu High Hope International Group Sunshine I/E Corp.
No. 50, Zhonghua Road, Nanjing, China

The Authorized Representative within EU who has been empowered
to enter into commitments on our behalf:

Name of Representative in EU:
MedNet GmbH
Borkstrasse 10, 48163 Muenster, Germany

Date:
July 15, 2018

Signature: Zhu Baoyu
General Manager

江苏汇鸿国际集团盛世进出口有限公司
JIANGSU HIGH HOPE INTERNATIONAL GROUP
SUNSHINE IMPORT AND EXPORT CORPORATION

朱宝宇