EC DECLARATION OF CONFORMITY

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

LATEX TOURNIQUET,LATEX FREE TOURNIQUET,LATEX FINGER COT
Classsification: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