Declaration of Conformity

Manufacturer: SunTech Medical, Inc.
507 Airport Boulevard, Suite 117
Morrisville, NC 27650-8200

EU Rep: SunTech Medical, Ltd.
Oakfield Industrial Estate
Stanton Harcourt Road
Eynsham, Oxon OX29 4TS
United Kingdom

Device: 247 Non-invasive Blood Pressure device with optional Temperature and Pulse Oximetry

Product Class: Class IIa
Assessment Procedure: Annex II
Notified Body: Intertek Semko AB
Box 1103, SE-164 22 Kista, Sweden

Product Marking: 

We hereby declare that the above mentioned product, including system components and accessories, complies with LVFS 2003:11 transposing European Medical Devices Directive 93/42/EEC and the following standards and normative documents:

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12. FDA 21CFR801.5, Medical devices; adequate directions for use.
13. AMMI SP10:2008, Electronic or Automated Sphygmomanometers

Valid on and after: August 16, 2011

[Signature]

Name: Chuck Setzer
Position: Quality and Regulatory Affairs Manager
Company: SunTech Medical, Inc.