


# Declaration of Conformity

<b>Applications of Council Directive: 93/42/EEC Medical Devices Directive</b>	 0413
Standards to which conformity is declared	IEC 60601-1, IEC 60601-2-30, IEC 60601-1-2 (EMC), IEC 60601-2-49, EN 1060-1, EN 1060-3, “Non-Invasive Sphygmomanometers – General Requirements & Supplementary Requirements for Electro-Mechanical BP Measuring Systems”, ISO 9919 “Pulse Oximeters for Medical Use”,
Manufacturer’s name and address	SunTech Medical, Inc. 507 Airport Boulevard, Suite 117 Morrisville, NC USA 27560-8200
European Authorized Representative	SunTech Medical, Ltd. Oakfield Industrial Estate Stanton Harcourt Road Eynsham, Oxon OX29 4TS United Kingdom
Type of Equipment	Cycle™ Class IIa Non-Invasive Blood Pressure Monitor System. System includes: cuffs, cables, power supply, and all accessories.
Model Number	1110, Cycle™ BP Monitor
Date of First Manufacture	December, 2005

I, the undersigned, hereby declare that the equipment specified above conforms to the above Directive(s) and Standard(s).



Name: Dayn McBee  
Position: Chief Executive Officer  
Company: SunTech Medical, Inc.

December 2, 2005  
Date