


# Declaration of Conformity

Applications of Council Directive: 93/42/EEC Medical Devices Directive	 0413
Standards to which conformity is declared	EN 60601-1, IEC 60601-2-30, EN 60601-1-2 (EMC), EN 1060-1, EN 1060-3, “Non-Invasive Sphygmomanometers – General Requirements & Supplementary Requirements for Electro-Mechanical BP Measuring Systems”, AAMI SP10
Manufacturer’s name and address	SunTech Medical, Inc. 507 Airport Boulevard, Suite 117 Morrisville, NC 27560-8200 USA
Importer’s name and address	SunTech Medical, Ltd. Europe Oakfield Industrial Estate Stanton Harcourt Rd. Eynsham, Oxfordshire OX29 4TS England
Type of Equipment	Bravo™ Non-Invasive Ambulatory Blood Pressure Monitor and Report Generator Software Class IIa
Model Number	222B

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



DECEMBER 9, 2004

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

Date