

Manufacturer

Tyeo Healtheare Group LP Nelleor Puritan Bennett Division Pleasanton, CA 94588 USA

European Authorized Representative

Tyeo Elealtheare UK LTD 154 Farcham Road Gosport PO13 BAS 11 K

European Declaration of Conformity to the Medical Device Directive, 93/42/EEC

Product Name: WarmTouch 5900 Patient Warming Unit

Product Model Number: 5900

Description: The WarmTouch 5900 Patient Warming Unit is a device which is part of a system used in the warming of patients for the treatment and prevention of hypothermia. The Model 5900 blows warmed air through a flexible hose into the disposable.

WarmTouch CareDrape and CareQuilt Blankets.

Classification/Rule: IIb / Rule 9

Conformity Assessment Route: Annex II

Declaration:

- Neileor Puritan Bennett deckares under our sole responsibility that the above product(s) to which this declaration relates, and which bear(s) the CE Marking, is ture) in conformity with the applicable requirements of EC Directive 93/42/BEC of 14 June 1993, concerning medical devices.
- As required by Annex II, Clause 3 of the above Directive, this Declaration is supported by EC Certificate
 and Quality System Certificate approval statement issued by TÜV Product Service.

Place:

Pleasanion, CA USA

Date:

02 June, 2003

Signature Name:

Sham Tarekere

Director, Design Assurance.