

EC Declaration of Conformity

This declaration of conformity is issued under the sole responsibility of the manufacturer in accordance with the Council Directive 93/42/EEC in its currently valid version (Annex I) concerning medical devices.

Product: **Disposable Pulse Oximetry Sensor
WRAP**

Type: DW-2203-6 Nihon Kohden
DW-2203-6S Nihon Kohden
DW-2211-6 Nellcor
DW-2221-6 BCI
DW-2271-6 Mindray
DW-2411-6 HP/Philips
DW-3003-6 Flexmon

Classification: **Class IIb**
(RL 93/42/EEC,
Annex IX)

CE mark: 

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 Munich, Germany

Conformity Assessment Process: Annex II, section 3 of the Directive 93/42/EEC

Particular product standard applied: ISO 80601-2-61

Issued by: **EnviteC-Wismar GmbH
Alter Holzhafen 18
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Germany**

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Authorized Signature:



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