# EC – declaration of conformity

We,

**Joh. Stiegelmeyer GmbH & CO. KG**
**Ackerstraße 42**
**D - 32051 Herford,**

hereby declare under sole responsibility as the manufacturer that the product model named below:

**Care Bed Series**

**Venta**

with model versions **movo** and **forto**

in the version submitted complies with the regulations of the EC Directive 93/42/EEC for Medical Devices, last amended by Directive 2007/47/EC dated 5 September 2007.

It is categorised as a Class 1 active medical device.
The relevant technical documentation is kept by the manufacturer's safety representative.
To evaluate the conformity to the Directives, all applicable parts of the following standards were referred to:

**Harmonised standards**:

| | |
|---|---|
| EN 14971:2013-04 | Risk Analysis for Medical Products |
| EN 60601-1:2006 | Safety for Medical Electrical Equipment |
| EN 60601-1-2:2007 | Electromagnetic Compatibility |
| DIN EN 60601-1-6:2010 | Medical electrical equipment: Suitability for Intended Use |
| DIN EN 60601-2-52:2010 | Particular requirements for the safety including essential performance of medical beds |

**International standards:**

| | |
|---|---|
| IEC 60601-2-52:2009 | Medical electrical equipment: Particular requirements for the basic safety and essential performance of medical beds |

Herford, 2015-05-11

Georgios Kampisiulis Kemmler
(Management)

Ralf Wiedemann
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