EC- DECLARATION OF CONFORMITY

Following the EC Directive concerning medical devices 93/42/EEC, annex VII.

I, the undersigned, agent of the following manufacturer:

Haelvoet nv

Leon Bekaertstraat 8	**Tel.:**	**+32 (0) 51 48 66 95**
8770 Ingelmunster	**Fax:**	**+32 (0) 51 48 73 19**
België	**Email:**	**info@haelvoet.be**

Declare hereby that the following product:

Vico H-L Stretcher

No.: 09770

Medical device class I (non-invasive device)

when installed, maintained and used in accordance with the manual, the rules of good craftsmanship, and the intended purpose complies with all necessary safety requirements and other relevant provisions of annex I of:

Medical Devices directive 93/42/EEC

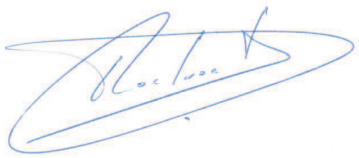
The following norms have been applied to indicate the conformity:

EN ISO 14971 Application of risk management to medical devices.

The above-mentioned product has been designed, produced and checked in accordance with the quality management system of ISO 9001:2000.

Ingelmunster, 28/09/2010

Signature:



Haelvoet Vincent
Managing director