# Declaration of conformity

In compliance with the Council Directive 93/42/EEC of 14 June 1993 about medical devices, the company

**ORMED GmbH**
**Merzhauser Strasse 112**
**D-79100 Freiburg - Germany**

**declares that the products of the product line**

**ARTROMOT®** (see Annex)

**fulfill the requirements of the Council Directive 93/42/EEC of 14 June 1993, Annex II, as well as the essential requirements of Annex I.**

**With reference to Rule 9 of the Directive 93/42/EEC, the product is a device of risk class IIa.**

CE
0297

**Freiburg, 30 August 2011**

**- QA Management Representative -**

This certificate is valid until expiry of the certificate referred to.
(the certificate can be downloaded from:
https://de.dqs-ul.com/kunden/kundendatenbank.html)

**Annex:**

ARTROMOT®-S2*PRO*
ARTROMOT®-S3
ARTROMOT®-S3 Comfort
ARTROMOT® ACTIVE-K
ARTROMOT®-K1
ARTROMOT®-K2
ARTROMOT®-K2PRO
ARTROMOT®-K2PRO Chip
ARTROMOT®-K3
ARTROMOT®-K4
ARTROMOT®-SP2
ARTROMOT®-SP3
ARTROMOT®-E2
ARTROMOT®-E2 compact