DET NORSKE VERITAS

CERTIFICATE OF CONFORMITY

Application of: Council Directive 93/42/EEC of 14 June 1993, issued as "Forskrift for Medisinsk Utstyr" by the Norwegian Ministry of Health and Social Affairs.

Certificate no.: 2002-RGC-MDD-0003

This is to certify that the technical documentation for the product:

VARIOUS SUPPORTS

Manufactured by

E-LIFE INTERNATIONAL CO., LTD.
LIVEN SPORTS MFG. (XIAMEN) CO., LTD.

at

4F-2, NO. 366, ZHONGHE RD., ZHONGHE CITY, TAIPEI COUNTY 235, TAIWAN

complies with the requirements applicable to it

The manufacturer's technical documentation, as required for **Class I** devices, has been reviewed and found to comply with the requirements in Annex VII, section 3.
Further identification and description of the products covered by this certificate are given in the Appendix.

18 January, 2006
for Det Norske Veritas Region Greater China

Limitations:

Any changes of the medical devices shall immediately be reported to Det Norske Veritas Region Greater China in order to examine whether this Certificate remains valid.

DET NORSKE VERITAS ★ 1864 ★

Dennis Lin
Technical Director,
DNV R.G.C. Product Certification

Any significant changes in design or construction of the product, or amendments to the Directive or Standards referenced above may render this receipt invalid. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC.