





EC Certificate

Full Quality Assurance System according to Medical Devices Directive 93/42/EEC Annex-II Section 3

Certificate Number: 1984-MDD-14-314

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive.

Organization:

DAEYANG MEDICAL CO. LTD.

147 Donghwakongdan-ro Donghwa-ri, Munmak-eup, Wonju-si, Gangwon-do, Korea

Product: High frequency therapy device **Models:** Back 3SE, Back 3S, Back 1S, Back 1E, Mabel6

The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa Meyer for details. This certificate is valid until 04 December 2019.

Report Number: M.4378.01

Kiwa Meyer Certification Services Inc. is Notified Body under Council Directive 93/42/EEC, concerning medical devices with identification number: 1984

05 December 2014, Istanbul, Turkey

Head of Notified Body

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