Intertek

EC Certificate

FULL QUALITY ASSURANCE SYSTEM
Directive 93/42/EEC on Medical Devices, Annex II (3)

Certificate Number
41311197

Initial Certification Date
June 20, 2003

Certificate Issue Date
June 20, 2008

Certificate Expiry Date
June 20, 2013

The certification is subject to the organization maintaining their system in compliance with the regulations stated in this certificate, allowing regular assessments and following the contracted requirements of the Notified Body.

Intertek Semko AB is a Notified Body according to Directive 93/42/EEC on medical devices, with identification number 0413.

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We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the Swedish 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, and the result entitles the organization to use the CE 0413 marking on those products listed below.

Organization:

INVIVO Corporation

12501 Research Parkway, Orlando FL 32826, USA

Product Category:

- Patient monitoring devices

For further identification of the products covered, see the MDD product list/product schedule.

Marie Olsson, Certification Manager MDD
Intertek Semko AB, Kista, Sweden