Intertek

EC Certificate

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

Certificate Number
41312698

Initial Certification Date
November 2, 2003

Certificate Valid from
July 10, 2009

Certificate Expiry Date
November 17, 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:

Bovie Medical Corporation

5115 Ulmerton Road, Clearwater, FL 33760, USA

Product Category:

- Cauteries
- Disposable nerve locators
- Medical light
- Electrosurgery equipment including accessories
- Ophthalmology burrs and power handles
- Nail drills and power handle

July 10, 2009
Signed date

Hans Ericsson, Acting Certification Manager MDD
Intertek Semko AB, Kista, Sweden

2008-09-01