

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
November 18, 2013

Certificate Expiry Date
November 17, 2018

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, LVFS 2003:11, 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.

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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Organization:

Bovie Medical Corporation

5115 Ulmerton Road, Clearwater, FL 33760, USA

Product Category:

- Cauteries
- Disposable nerve locators
- Surgical light
- Electrosurgery equipment including accessories
- Ophthalmology burrs and power handles

November 15, 2013

Signed date



Mats Premfors, Certification Authority MDD
Intertek Semko AB, Kista, Sweden