Intertek

EC Certificate

PRODUCTION QUALITY ASSURANCE
Directive 93/42/EEC on Medical Devices, Annex V

Certificate Number
41312909

Initial Certification Date
February 11, 1999

Certificate Valid from
February 12, 2014

Certificate Expiry Date
February 11, 2019

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 production 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 V of the Directive 93/42/EEC on medical devices. We certify that the production quality system conforms with the relevant provisions of the aforementioned legislation, and the result entitles the organization to use the CE 0413 marking on those products listed below.

Organization:

Thought Technology Ltd.

8205 Montreal/Toronto Blvd. Suite 223, Montreal West, Quebec, H4X 1N1, Canada

Product Category:

- Equipment for biofeedback and for therapy of urinary and faecal incontinence, Class I with measuring function.
- Muscle stimulators, Class IIa

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

February 7, 2014
Signed date

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

2008-09-01