

EC CERTIFICATE

PRODUCTION QUALITY ASSURANCE SYSTEM APPROVAL CERTIFICATE (Annex V of the Directive 93/42/EEC on Medical Devices)

No. 41312909

We hereby declare that an examination of the under mentioned production quality assurance system has been carried out following the requirements of the 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 assurance system conforms with the relevant provisions of the aforementioned legislation.

Manufacturer:

Thought Technology Ltd.
2180 Belgrave Ave.
Montreal, Quebec, H4A2L8
Canada

Product category:

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

Date of expiry:

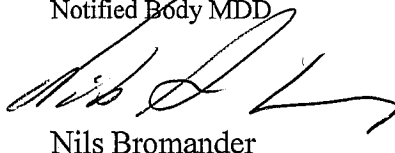
11 February 2009

The Certificate is valid for the devices which are stated in the present MDD – Product list

Stockholm
22 November 2004

Intertek Semko AB
Notified Body MDD

Original certificate issued
11 February 1999



Nils Bromander

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