

# EC CERTIFICATE

## FULL QUALITY ASSURANCE SYSTEM APPROVAL CERTIFICATE

(Annex II of the Directive 93/42/EEC on Medical Devices)

No. 41315235

We hereby declare that an examination of the under mentioned full 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 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 legislation.

Manufacturer: Neurologica Corporation  
14 Electronics Avenue  
Danvers, MA 01923  
USA

Product category: Computed Tomography Scanners

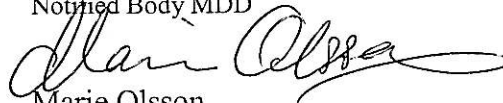
Date of expiry: 6 October 2011

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

Stockholm  
6 October 2006

The original certificate issued on  
6 October 2006

**Intertek Semko AB**  
Notified Body MDD

  
Marie Olsson  
Certification Manager MDD

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