

EC CERTIFICATE

FULL QUALITY ASSURANCE SYSTEM APPROVAL CERTIFICATE

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

No. 41313430

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: KanMed AB
Gårdsfogdev 18B
SE-168 66 Bromma
Sweden

Product category: Patient Warming Systems

Date of expiry: 19 June 2010

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

Stockholm
6 October 2005

Intertek Semko AB
Notified Body MDD

The original certificate issued on
19 June 2000


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.

Intertek ETL SEMKO

2004-05-14