

# EC CERTIFICATE

## FULL QUALITY ASSURANCE SYSTEM APPROVAL CERTIFICATE

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

No. 41315225

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: AMVEX Corporation  
121 Granton Drive, Unit 21  
Richmond Hill, ON L4B 3N4  
Canada

Product category: - Flow meters  
- Hose assemblies  
- Suction regulators  
- Gas regulators

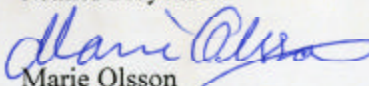
Date of expiry: 24 November 2010

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

Stockholm  
24 November 2005

The original certificate issued on  
24 November 2005

**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.

**Intertek** ETL SEMKO