



EC Certificate Full Quality Assurance System: Certificate U019819943433

The management system of
Sechrist Industries Inc.

4225 East La Palma, Anaheim, CA, 92807, United States

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC
on medical devices, Annex II (excluding Section 4)

For the following products

Gas Mixers for respiratory care and heart bypass oxygenation equipment, Manoplace Hyperbaric Chamber.

Where the above scope includes class II medical devices, a valid CE Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 18 December 2010 until 14 May 2024
and remains valid subject to satisfactory surveillance audits,
Issue 1, Certified since 18 May 1998
and first certified by SGS Belgium NV since 18 December 2010

Certification is based on reports numbered WRMSC 08209

Authorized by

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