

EC Certificate Full Quality Assurance System: Certificate ES19/86919

The management system of

Air Liquide Medicinal, S.A.

Rua Dr Antonio Loureiro Borges, N 4-3 Arquiparque Miraflores,
1495-131 Algés, Portugal

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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 16 December 2019 until 16 February 2023
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 16 February 2009
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered ES/MAD 231815

This is a multi-site certification.

Additional site details are listed on subsequent pages

Authorised by

SGS Belgium NV, Notified Body 1639

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LPMDS007 - Certificate CE1639 Annex II-4-EN rev. 02

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Certificate ES19/86919 continued

Air Liquide Medicinal, S.A.

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

Issue 1

Detailed scope

**Fixed Medical Gases Distribution Systems
with or without anaesthetic gas exhaust system.**

***Sistema de Distribuição de Gases Medicinais Fixo
com ou sem sistema de exaustão de gases anestésicos***

Where the above scope includes class III medical device(s), a valid EC 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.

Additional facilities

Avenida Vasco da Gama, no 7375, 4430 - 755 Avintes, Portugal