EC Certificate Full Quality Assurance System: Certificate ES19/86982

The management system of

ULTRA-CONTROLO - PROJECTOS INDUSTRIAIS, LDA

Parque Industrial da Quinta do Lavi, Armz 8 - Abrunheira,
2710-089 SINTRA, 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

Anaesthetic Gas Scavenging Systems ULTRASEG
Medical Vacuum Systems ULTRAVAC
Medical Compressed Air Systems ULTRAAR
Medical Compressed Air Systems MEDIAR
Medical Oxygen Supply Systems ULTRAOX

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.

This certificate is valid from 16 December 2019 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 07 October 2010
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered ES/MAD 227898

Authorised by

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