

EC Certificate Full Quality Assurance System: Certificate ES19/86854

The management system of

Albino Dias de Andrade, S.A.

Rotunda do Complexo Desportivo,
15 Seroa, 4595-069 Paços de Ferreira, 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 27 April 2021 until 24 May 2024
And remains valid subject to satisfactory surveillance audits.
Issue 4. Certified since 11 October 2013.

Certification is based on reports numbered ES/MAD 226345

Authorised by



Global Medical Devices Head of Notified Body

SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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Albino Dias de Andrade, S.A.

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

Issue 4

Detailed scope

**Non-sterile Gauze & non-woven dressings.
Sterile Gauze & non-woven dressings.
Gauze and non-woven dressings with X-ray thread,
Sterile and Non sterile**

**Compressas de gaze estéril e não estéril.
Compressas de tecidos e não-tecidos
estéreis e não estéreis.**

**Compressas de gaze, e de tecido não tecido,
estéril e não estéril com fios de contraste ao raio-x.**

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.