EC Certificate Full Quality Assurance System: Certificate GB10/80184

The management system of

Medbone-Medical Devices, Lda.

Rua Fonte da Carreira n° 350 B-17,
2645-467 CASCAIS, 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

Sterile synthetic bone substitute.

Substitutos ósseos sintéticos estéreis.

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 04 December 2013 until 24 April 2018 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 11 February 2016
Issue 3. Certified since 24 April 2010

Certification is based on reports numbered GB/PI 222625

Authorised by

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