

The management system of

Micropace Pty Ltd

Unit 7/186-188 Canterbury Road,
Canterbury, NSW, 2193, Australia

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 21 March 2011 until 23 March 2012 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 23 March 2012
Issue 5. Certified since November 2003

Certification is based on reports numbered AUCOR 81

This is a multi-site certification.
Additional site details are listed on the subsequent page.

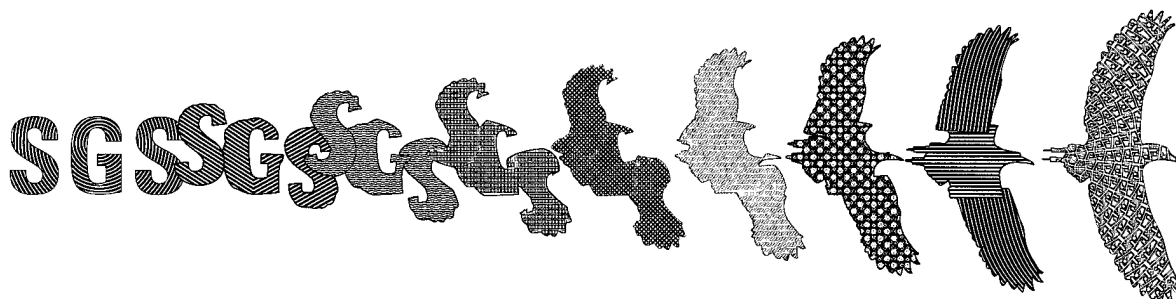
Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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SGS CE 01 0311 M2

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Micropace Pty Ltd

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

Issue 5

Detailed scope

Cardiac stimulator device.

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.

Additional facilities

Micropace EP Inc, 3205 West Warner Ave, Santa Ana CA, USA, 92704