



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 07 04 51392 004

Manufacturer: Proxy Biomedical Ltd.
Coilleach
Spiddal
Co Galway
IRELAND

Facility(ies): Proxy Biomedical Ltd.
Coilleach, Spiddal, Co Galway, IRELAND

Product Category(ies): Surgical Mesh and
Surgical Mesh Delivery Systems

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: 75900019

Valid until: 2009-07-14



Date, 2007-06-12

Reiner Krumme

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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