By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. **CE 661971**

Issued To: **Active Life Scientific, Inc.**
Suite 503
27 E Cota Street
Santa Barbara
California
93101
USA

In respect of:

Design, development, and manufacturing of OsteoProbe for diagnosing and monitoring material aspects of bone quality.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):

Stewart Brain, Head of Compliance & Risk - Medical Devices

First Issued: **2017-08-15** Date: **2017-08-15** Expiry Date: **2022-08-14**

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Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 661971**
Date:	**2017-08-15**
Issued To:	**Active Life Scientific, Inc.** **Suite 503** **27 E Cota Street** **Santa Barbara** **California** **93101** **USA**

Subcontractor:	**Service(s) supplied**
Emergo Europe Prinsessegracht 20 2514 AP The Hague The Netherlands	**EU Representative**

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EC Certificate - Full Quality Assurance System Certificate History

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Date	Reference Number	Action
Current	8608289	First Issue.