



EC CERTIFICATE

MedApps Inc.

7975 N Hayden Road, Suite A-203
Scottsdale AZ 85258
USA

Full Quality Assurance System Approval Certificate

Annex II, section 3 of Council Directive 93/42/EEC concerning medical devices

Scope of Certificate:

Design and manufacture of remote patient monitoring equipment

Device Classification:

Class IIa

Device Descriptions:

Remote Patient Monitoring System

Model Type:

MedApps 2.0

File Number A17343

Cycle Start Date 23 December 2009

Certificate No. 616.111012

Effective Date 12 October 2011

Expiry Date 16 December 2012

Authorised by

Barry A. Fitch

Certification Manager

For and on Behalf of UL International (UK) Ltd

We hereby declare that an examination of the full quality assurance system has been carried out per report SR7415652, following the requirements of the national legislation to which the undersigned is subject, transposing Annex II (with the exemption of section 4) of Council Directive 93/42/EEC on Medical Devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive and is subject to periodic surveillance as required by 93/42/EEC, Annex II, Section 5. For Class III devices where they are covered by this certificate, an EC Design Examination certificate according to 93/42/EEC, Annex II, Section 4 is required. This certificate is issued with 0 attachments listing model numbers and 0 addendums listing additional locations covered by this certificate.

Notified Body

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00-NB-F0413 1.0

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