

EC-Certificate

(Production quality assurance system)
according to annex V, section 3
Medical Devices Directive 93/42/EEC

It is herewith confirmed by the
Notified Body

**EUROCAT Institute for
Certification and Testing**
Quarat[®] Center, Wittichstraße 2
64295 Darmstadt, Germany
Identification No. 0535

that the manufacturer
LD Technology LLC
100 N. Biscayne Blvd Suite 500
33132 Miami, USA
concerning the medical devices
(products/variants specified in enclosure)
fulfills the requirements according to
Annex V, section 3, Medical Devices
Directive 93/42/EEC. The manufacturer
has established a quality assurance
system for the production and final
inspection of specified devices.

The enclosure is part of this certificate
and contains 1 page.

Based on the periodical surveillance the
certificate is valid until 23 March 2014.

Report No.: BQM09046ES02
Certificate No.: **CQ090831-V**

CE 0535

Accreditation granted by
Zentralstelle der Länder
für Sicherheitstechnik

ZLS
ZLS-ZQ-302/06

Darmstadt, 24 March 2009


Certification Body



Enclosure of EC-Certificate

(Production quality assurance system)

according to annex V, section 3, Medical Devices Directive 93/42/EEC

Certificate No.: CQ090831-V

Manufacturer

LD Technology LLC

100 N. Biscayne Blvd Suite 500

33132 Miami, USA

Medical devices concerned:

Name of product	Variant	Item spec.	UMDNS	class
Electro Sensor Teck System (ES Teck System)	Model EIS-BC Model PEMS Model Complex			Ila
Impedance Tomography Mammography (ITM)	USB-1			Ila

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Darmstadt, 24 March 2009



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