

**EUROCAT Institute for  
Certification and Testing**

Quarat<sup>®</sup> Center \* Wittichstraße 2  
64295 Darmstadt, Germany

herewith confirms, that the company



**Ad Rem Technology**

**162, rue du Faubourg Saint-Honoré  
FR-75008 Paris, France**

has established a

**quality management system**

for the

**design/development,  
manufacturing, sales and  
servicing support for neuro  
muscular electro stimulators.**

An audit has given evidence that the  
quality system is in compliance with the  
requirements of

**DIN EN ISO 9001 : 2008.**

Based on the periodical surveillance the  
certificate is valid until 23 March 2012.

Report No.: BQM09040TK01

**Certificate No.: CQ090832-9**



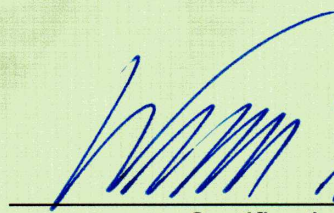


TGA TRÄGERGEMEINSCHAFT  
FÜR AKKREDITIERUNG  
GERMAN ASSOCIATION FOR  
ACCREDITATION GMBH



Accreditation granted by TGA  
Represented in DAR  
TGA-ZM-24-97-00

Darmstadt, 24 March 2009



Certification Body

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**Quality management system**

for the

**design/development,  
manufacturing, sales and  
servicing support for neuro  
muscular electro stimulators.**

An audit has given evidence that the  
quality system is in compliance with the  
requirements of

**DIN EN ISO 13485 : 2007**

**(EN ISO 13485 : 2003 + AC : 2007).**

Based on the periodical surveillance the  
certificate is valid until 13 February 2013.

Report No.: BQM09040TK01

**Certificate No.: CQ090832-13**



Accredited by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln  
und Medizinprodukten  
ZLG-ZQ-684.99.05-46

Darmstadt, 24 March 2009



Certification Body

## 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<sup>®</sup>Center, Wittichstraße 2  
64295 Darmstadt, Germany  
Identification No. 0535

that the manufacturer



**Ad Rem Technology**

**162, rue du Faubourg Saint-Honoré  
FR-75008 Paris, France**

concerning the medical device  
**VEINOPLUS, Model 2.1**

(UMDNS: 13-762, Class IIa)

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.

Based on the periodical surveillance the  
certificate is valid until 13 February 2013.

Report No.: BQM08018TK01

Certificate No.: **CQ080451-V**

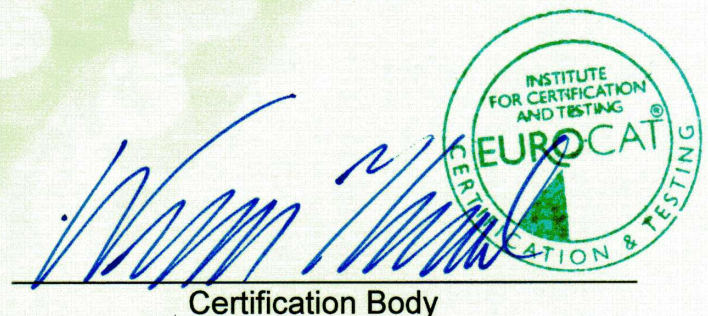
**CE 0535**

Accreditation granted by  
Zentralstelle der Länder  
für Sicherheitstechnik

**ZLS**

ZLS-ZQ-302/06

Darmstadt, 14 February 2008



\_\_\_\_\_  
Certification Body

The seal is circular with the text "INSTITUTE FOR CERTIFICATION AND TESTING" around the top and "EUROCAT" in the center. Below "EUROCAT" is the text "CERTIFICATION & TESTING".