EC Declaration of Conformity

Manufacturer:

Shenzhen Dongdixin Technology Co., LTD.

No. 3 Building XiliBaimang Xusheng

Industrial Estate, 518108 Nanshan,

Shenzhen China

Tel: (86) 755 2765 7648

Fax: (86) 755 2765 2674

Authorized Representative:

Shanghai international holding Corp. GmbH (Europe)

Eiffestraße 80, 20537 Hamburg Germany

Tel: 0049-40-2513175

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DIMDI No.: DE/0000040627

We, the manufacturer, herewith declare that the products

Nerve and Muscle Stimulator

DX6605BO（EMT-4）, EMT-6, T-6

(including system components and accessories)

UMDNS-Code: **13762**

meet the provisions of Directive 93/42/EEC and amended by Directive 2007/47/EC which apply to them.

The medical device has been assigned to class IIa according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to Annex II.3 of Directive 93/42/EEC and amended by Directive 2007/47/EC.

Compliance of the designated product with the Directive 93/42/EEC and amended by Directive 2007/47/EC have been assessed and certified by the Notified Body

TÜV Rheinland LGA Product GmbH
Am Grauen Stein D-51105 Köln
Certificate No.: HD 60039057 0001
Issue date: 19 Jul., 2011
Expiry date: 21 Jun. 2016

following the procedure relating to the EC Declaration of Conformity set out in Annex II.3 of Directive 93/42/EEC and amended by Directive 2007/47/EC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: Shenzhen Dongdixin Technology Co., LTD.

Address: No. 3 Building XiliBaimang Xusheng Industrial Estate
518108 Nanshan, Shenzhen China

Shenzhen 2012.3.20

Place , date

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Legally binding signature, Function