

EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

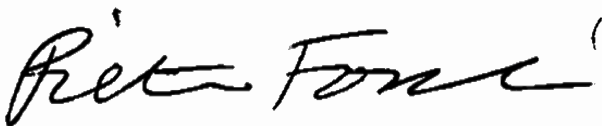
No. CE 538571
Issued To: **Medicom MTD Ltd**
99 Petrovskaya Str
Taganrog, Rostov Region
347900
Russian Federation

In respect of:

The manufacture of equipment for EEG/EMG/EP studies, long term Video EEG Monitoring, Cerebral Function Monitoring, PSG/Sleep Diagnostics, neurophysiological, psychophysiological and psychological studies, and equipment for biofeedback training and rehabilitation.

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

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



Pietro Foschi - Strategic Delivery Director

First Issued: **03 September 2008**

Date: **05 January 2015**

Expiry Date: **02 September 2018**

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Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

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List of Significant Subcontractors

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

Certificate No: **CE 538571**
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Taganrog, Rostov Region
347900
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Subcontractor:

Service(s) supplied

Polmed.de
Vertretung und Repräsentanz
Dipl.-oec Boguslaw Karcz
Steinäcker 5
Aichwald
73773
Germany

EU Representative

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EC Certificate - Production Quality Assurance Certificate History

Certificate No: **CE 538571**
 Date: **05 January 2015**
 Issued To: **Medicom MTD Ltd**
99 Petrovskaya Str
Taganrog, Rostov Region
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Date	Reference Number	Action
03 September 2008	7218012	First Issue
21 February 2011	7604149	Extention to scope changed from The manufacture of EEG equipment and Biofeedback psychophysiological rehabilitation devices to The manufacture of equipment for EEG/EP studies, long term Video EEG Monitoring, Cerebral Function Monitoring, PSG/Sleep Diagnostics, neurophysiological, psychophysiological and psychological studies, and equipment for biofeedback training and rehabilitation.
28 Jun 2013	7985407	Certificate renewal and addition of Polmed.de as EU representative.
05 January 2015	8269950	Extension to scope to include equipment for EMG studies

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