

EC Certificate - Production Quality Assurance

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

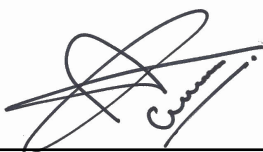
No. CE 538571
Issued To: **Medicom MTD Ltd**
68 Frunze Str.
Taganrog, Rostov Region
347900
Russian Federation

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 2797):



Albert Roossien, Regulatory Lead

First Issued: **2008-09-03**

Date: **2019-02-08**

Expiry Date: **2023-09-02**

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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.

Information and Contact: BSI, Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands Tel: + 31 20 346 0780

BSI Group The Netherlands B.V. registered in The Netherlands under 33264284.

A member of BSI Group of Companies.

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Supplementary Information to CE 538571

Issued To:

**Medicom MTD Ltd
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Number	Device Name
Class IIa	
NBOG Code	Device or Device subcategory
MD1301	Electroencephalograph-recorder computerized portable «ENCEPHALAN-EEGR-19/26»
MD1301	Cerebral Function Monitor "Encephalan-CFM"
MD1301	Sleep Signals Recorder "ApnOx"
MD1301	Neuromyoanalyzer NMA-4-01 "Neuromyan"
MD1301	Objective psychological analysis and testing system "Egoscop"
MD1103	Psychophysiological telemetric system "Rehacor-T"

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List of Significant Subcontractors

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

Certificate No: **CE 538571**
Date: **2019-02-08**
Issued To: **Medicom MTD Ltd**
68 Frunze Str.
Taganrog, Rostov Region
347900
Russian Federation

Subcontractor:

Service(s) supplied

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

EU Representative

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Certificate History

Certificate No: **CE 538571**
 Date: **2019-02-08**
 Issued To: **Medicom MTD Ltd**
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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.
26 October 2017	8799763	Change of address from Medicom MTD Ltd, 99 Petrovskaya Str, Taganrog, Rostov Region, 347900, Russian Federation to Medicom MTD Ltd, 68 Frunze Str, Taganrog, Rostov Region, 347900, Russian Federation.
24 August 2018	9643016	Certificate Renewal.
Current	7780277	Traceable to NB 0086.

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