BMO-200B

Patient Monitor

Suitable for ICU/CCU, Postoperative observatory, Emergency room, General ward, etc



ADVANTAGES AT A GLANCE

- 12.1" color TFT LCD screen with multi channel waveforms display
- Rechargeable high energy built in Li Battery
- Easy to learn, easy to use, simplified menus and delicated functions keys
- Easy connecting with central monitoring system 96 hours data storage
- Suitable for adult, pediatrics and neonates
- Monitors end-tidal CO2 and inspired CO2 accuracy at high respiration rates
- Gas monitoring with cutting edge water filter tube





Vital signs show perfectly

Suitable for ICU/CCU, Postoperative observatory, Emergency room, General ward, etc



80 120 /80 96





Standard Interface

Max 8 waveform display simultaneously, max 6 leads ECG waveforms display

Trend graphs

Big Font interface







Durable probes connector with high quality

Build in Thermal Recorder

2 Advanced

Our monitor could be used for Adult, Pediatric and Neonatal.

Our monitor has been welcomed by the doctors because of its stability in service reliability in operation and simplicity in maintenance.

BMV Medical, win the reputation by providing high quality products.

Oata Storage – stable and fast

- 400 groups NIBP measurements
- 72 hours Trend data and Trend graphs
- 200 groups Alarm events reviewing
- 12 hours Wave forms reviewing
- Supporting for Arrhythmia analysis and drug dose calculation

Alarming – more safe and reliable

3 level Audible and visual alarming, Dual alarm light for physiological and technical alarming



5 Standard Parameter: ECG, RESP, NIBP, SpO2, PR, TEMP Optional: 2-IBP, EtCO2, Touch screen, Build in thermal printer

www.bmv.cc



















SYS-50 Syringe Pump

Simple and Comfortable

3.5" Touch screen operation

Large and clear user interface, providing medical staff with a pleasant visual experience and facilitating real-time observation of the infusion status.

IV set management

A standard fitted tube clamp will prevent iv-sets entanglement and enhance infusion control.

Innovative and Safe

- certified by EN1789 standard for ambulances and medical transportation vehicles
- Syringe slider protection
- Internal battery up to 20 hours continuous power supply.

Information interconnection

- Optional built-in WiFi module.
- Can connected to HIS/CIS via WLAN or LAN.
- Can connected to MP-900 infusion central monitoring system to implement centralized monitoring of all infusions.

Smart and Convenient

- 8 infusion modes: Adapted for various clinical departments and a variety of special drugs.
- Continuous titration: Changing the flow rate without stopping the infusion.



Simplified Smart

Specifications

Dimension:	362(L) X 151(H) X 130(W)mm		
Weight:	1.8kg(including battery)		
Screen:	3.5 inch LCD touch screen		
Classification:	Type CF,Class I,IP24		
Syringe types:	All standard syringes of 5ml, 10ml, 20ml, 30ml, 50ml, 60ml		

Basic parameter

Infusion mode:
Rate, Time, Weight, Sequence, Trapezia, Loading dose, Micro, TIVA

Infusion rate range:
0.10-2000.0(ml/h), minimum increments 0.01ml/h

VTBI range:
0.1-9999ml

Accumulated volume:
0.0-999999.99ml

Accuracy:
≤2%

KVO rate:
0.1-5ml/h

Bolus rate:
0.10-2000.0(ml/h)

11 levels selectable, 225mmHg- 975mmHg

Special function			
Recent therapy:	20 recent therapies are recorded and can be used for rapid infusion		
Rate change during infusion:	Infusion rate can be changed during infusion without stop.		
Auto screen lock:	Available.		
Repeated Alarm:	The Alarm sounds again in 2 minutes, if the alarm is not resolved but set mute only.		
Storage:	Maximum 2000 records can be stored and reviewed.		
Sound volume:	11 levels selectable		
Power supply switching:	When AC/DC power supply is cut off, the infusion automatically switch to internal battery sup		

Power supply AC power: 100-240, 50/60 Hz, power consumption 45VA External DC power: DC 12 V, 1.5A Internal battery: Lithium battery 11.1V 1500mAh or Lithium battery 11.34V 2900mAh

Battery VS Charging time:					
Configuration	Fast charging time(Power off)	st charging time(Power off) Normal charging time(Power on)			
One 1500mAh@11.1V battery	3 hours up to 95% percetage, less than 4 hours to full	6 hours to full	Not less than 8 hours		
Two 1500mAh@11.1V batteries	6 hours up to 95% percetage, less than 8 hours to full	12hours to full	Not less than 15 hours		
Two 2900mAh@11.34V batteries	12 hours up to 95% percetage, less than 16 hours to full	etage, less than 16 hours to full 24hours to full			
NOTE:Infusion at 5ml/h with new battery under room temperature					

Connectivity	
WiFi/Wired:	Available
Data interface:	USB

Sare conditions			
	Temperature	Humidity	Pressure altitude
Operating conditions:	5°C to 40°C	15% to 95% RH, non-condensing	70.0 kPa-106.0 kPa
Storage and Shipping conditions:	-20°C to +55°C	10% to 95% RH. non-condensing	22.0 kPa-107.4 kPa

MEDCAPTAIN()

of a condition

Occlusion level:

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD

Add: 12th Floor, Baiwang Research Building, No.5158 Shahe West Road, Xili, Nanshan, Shenzhen, P.R. China

Tel: +86 755 26953369 Fax: +86 755 26001651 E-mail: info@medcaptain.com Website: www.medcaptain.com



DISTRIBUTOR:



EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 14 09 81250 002

Manufacturer:

BMV Technology Co.,Ltd

4/F, BMV of Yinjin Building, Block 71,Baoan Centre District, Shenzhen 518010,Guangdong, PEOPLE'S REPUBULIC OF CHINA

EC-Representative:

Wellkang Ltd

Suite B, 29 Harley Street

LONDON W1G 9QR

UNITED KINGDOM

Product Category(ies):

Patient Monitors

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

748496336

Valid from:

2018-10-30 2020-10-29

Valid until:

H.- St.

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 2

Date,

2018-10-30



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 14 09 81250 002

Facility(ies):

BMV Technology Co.,Ltd

4/F, BMV of Yinjin Building, Block 71, Baoan Centre District,

Shenzhen 518010, Guangdong, PEOPLE'S REPUBULIC OF CHINA



Page 2 of 2





EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 086916 0017 Rev. 00

Manufacturer: MEDCAPTAIN MEDICAL

TECHNOLOGY CO., LTD.

12th Floor, Baiwang Research Building No.5158 Shahe West Road

Xili. Nanshan

518055 Shenzhen, Guangdong PEOPLE'S REPUBLIC OF CHINA

EC-Representative: Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80, 20537 Hamburg, GERMANY

Product Category(ies): Syringe Pump, Infusion Pump, Infusion Workstation, Sterile

Syringes for single use, Sterile Infusion sets for single use,

DVT Preventive Pump

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: GZ18117EXT01

 Valid from:
 2019-02-17

 Valid until:
 2024-02-16

Date, 2019-01-30

Stefan Preiß

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III



EC Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 086916 0017 Rev. 00

Facility(ies):

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD. 12th Floor, Baiwang Research Building, No.5158 Shahe West Road, Xili, Nanshan, 518055 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD. 1-4 Floor, 11th Factory Building, Nangang 1st Industrial Park, Songbai Road, Xili, Nanshan, 518055 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD. Room No. 401, Block B, Weike Science and Technology Building, Dongming Avenue, Guangming Subdistrict, Guangming New District, 518107 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

TÜV®



CERTIFICATE

No. Q1N 14 10 81250 003

Holder of Certificate:

BMV Technology Co., Ltd.

4/F, BMV of Yinjin Building, Block 71, Baoan Centre District, Shenzhen 518010, Guangdong. PEOPLE'S REPUBULIC OF CHINA

Facility(ies):

BMV Technology Co., Ltd.

4/F, BMV of Yinjin Building, Block 71, Baoan Centre District, 518010, Shenzhen, Guangdong, PEOPLE'S

REPUBLIC OF CHNA

Certification Mark:



Scope of Certificate:

Design and Development, Production and Distribution

of Patient Monitors

Applied

Standard(s):

EN ISO 13485:2012 + AC:2012 Medical devices - Quality management systems -

Requirements for regulatory purposes (ISO 13485:2003 - Car 1:2009)

DIN EN ISO 13485:2012

The Certification Body of TÜV SÜD Product Service Company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf

Report No .:

748496336

Valid from:

2018-10-30

Valid until:

2020-10-29

Date, 2018-10-30

Hans-Heiner Junker

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CERTIFICATE

No. Q1N 16 12 86916 007

Holder of Certificate:

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.

12th Floor, Baiwang Research Building

No.5158 Shahe West Road

Xili, Nanshan

518055 Shenzhen, Guangdong PEOPLE'S REPUBLIC OF CHINA



Certification Mark:



Scope of Certificate:

Design and Development, Production and Distribution of Syringe Pump, Infusion Pump, Infusion Workstation, Infusion Central Monitoring System, Sterile Syringes for single use, Sterile Infusion sets for single use and DVT Preventive Pump

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.:

GZ1711701

Valid from:

2017-03-17

Valid until:

2020-02-16

Date. 2017-03-17

Stefan Preiß

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CERTIFICATE No. Q1N 16 12 86916 007

Applied Standard(s):

EN ISO 13485:2012 + AC:2012

Medical devices - Quality management systems -

Requirements for regulatory purposes (ISO 13485:2003 + Cor. 1:2009)

DIN EN ISO 13485:2012

Facility(ies):

MEDCAPTAIN MEDICAL TECHNOLOGY CO., LTD.

1-4 Floor, 11th Factory Building, Nangang 1st Industrial Park, Songbai Road, Xili, Nanshan, 518055 Shenzhen, Guangdong,

PEOPLE'S REPUBLIC OF CHINA

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12th Floor, Baiwang Research Building, No.5158 Shahe West Road, Xili, Nanshan, 518055 Shenzhen, Guangdong, PEOPLE'S

REPUBLIC OF CHINA



