



Shangrila 510S

Portable Ventilator

TECHNICAL
DATA

AEO MED
Reliable Quality Thoughtful Service

Shangrila 510S is a universally used model. Not only for emergency ventilation, but also it is suitable for ideal transport ventilation. The circumstances include on-site emergency treatment, pre-hospital transport, inter-hospital transport, and even intra-hospital transport.



Patient Type

Adult, Child, Infant (>3.5 kg)

Ventilation Modes

A/C-V with Sigh	A/C-P
SIMV-V	SIMV-P
SPONT/PSV	CPAP
MANUAL	

Enhancements

Apnea back-up ventilation	Manual breath
Alarm silence	

Parameter Setting

Tidal volume (Vt)	0-2000 mL
Pinsp	5-50 cmH ₂ O
Psupp	0-50 cmH ₂ O
CPAP (NIV)	0-30 cmH ₂ O
PEEP	0-30 cmH ₂ O
Pressure trigger	-20-0 cmH ₂ O
Flow trigger	2-30 L/min
Frequency (f)	1-120 bpm (A/C) 1-40 bpm (SIMV)
I:E ratio	4:1-1:10
Pause time (Tpause)	0-5 s
O ₂ concentration	40-100 %
Inspiration flow	Max. 90 L/min

Monitoring Data

MV	VT	Frequency (f)
FiO ₂	Ppeak	PEEP

Waveform Display

P-T	F-T
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Alarm Setting

High MV	OFF, 1-25 L
Low MV	OFF, 0-24 L
High FiO ₂	OFF, 50-100 %
Low FiO ₂	OFF, 35-99 %
High Paw	6-80 cmH ₂ O
Low Paw	0-40 cmH ₂ O
AC power failure	Gas supply down
O ₂ deficiency	MV high
Low battery	No VT
Apnea	CP high

Language

Chinese	English
Spanish	Italian
Turkish	Other languages

Environmental Requirements

Temperature

Operating condition	-18-50 °C
Storage condition	-20-60 °C

Relative humidity

Operating condition	15-95 %
Storage condition	10-95 %

Barometric pressure

Operating condition	70-110 KPa
Storage condition	50-110 KPa

Electrical

AC voltage	100-240 V, 50/60 Hz
DC voltage	12 V
One battery run time	Minimum 4.5 hours
Two batteries run time	Minimum 10 hours

Gas Supply

Supply pressure (O2)	250-600 kPa
Flow	Max. 180 L/min

Physical Specifications

Total Weight	8 kg
Dimension (H*W*D):	168* 156 *300 mm
LCD screen	5" TFT
Communication Ports	RS232



Product Service

Certificate

No. Q5 065725 0022 Rev. 02

Holder of Certificate: **Beijing Aeonmed Co., Ltd.**
Room 405
Basement 1 to 4th Floor of 901 Unit
Building 9, No.26 Outer Ring West Road
Fengtai District
100070 Beijing
PEOPLE'S REPUBLIC OF CHINA

Certification Mark:



Scope of Certificate: **Design and Development, Production, Distribution, Installation and Servicing of Anaesthetic Workstation, Vaporizer, Ventilator, Medical Air Compressor, Infusion Pump, Ceiling Pendent, Operating Table, Surgical Light, Multi-Parameter Patient Monitor, Syringe Pump, Patient Warming System, Videoscope System.**

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.: BJ1985904

Valid from: 2020-03-23

Valid until: 2022-12-31

Date, 2020-03-23

Christoph Dicks
Head of Certification/Notified Body

Certificate

No. Q5 065725 0022 Rev. 02

Applied Standard(s):

EN ISO 13485:2016
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies):

Beijing Aeonmed Co., Ltd.
Room 405, Basement 1 to 4th Floor of 901 Unit, Building 9, No.26
Outer Ring West Road, Fengtai District, 100070 Beijing, PEOPLE'S
REPUBLIC OF CHINA

Beijing Aeonmed Co., Ltd.
No. 10 Chaobai Street, Yingbin Road West, Yanjiao Development
Zone, 065201 Langfang City, Hebei Province, PEOPLE'S
REPUBLIC OF CHINA



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



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 065725 0019 Rev. 04

Manufacturer:

Beijing Aeonmed Co., Ltd.

Room 405

Basement 1 to 4th Floor of 901 Unit

Building 9, No.26 Outer Ring West Road

Fengtai District

100070 Beijing

PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Anaesthetic Workstation, Vaporizer,
Ventilator, Medical Air Compressor,
Infusion Pump, Ceiling Pendant,
Multi-Parameter Patient Monitor,
Videoscope System, Patient Warming System.**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10657250019Rev.04

Report No.:

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2021-05-21

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Date,

2021-05-21

Christoph Dicks

Head of Certification/Notified Body