٦	Technical Specification
Ventilation Modes	
	VCV(A/C) PCV(A/C) PRVC SIMV(VCV)+PSV
	SIMV(PCV)+PSV SIMV(PRVC)+PSV SPONT/CPAP+PSV
	BIVENT+PSV NIV/CPAP NIV-T NIV-S/T
Parameters	
Tidal Volume:	20~2000 ml
Respiration Rate:	1~80 bpm
• Tinsp:	0.2~9 s
Tslope:	0~2 s
Tpause:	0~4 s
I:E Ratio:	1:10~4:1
• FiO <sub>2</sub> :	21%~100%
Trigger Sensitivity:	Pressure (-20~0 cmH <sub>2</sub> O, above PEEP)
	Flow (0.5~20 LPM)
• PEEP:	0~35 cmH₂O
Psupport:	0~70 cmH₂O
• Pinsp:	5~70 cmH <sub>2</sub> O
Special Procedures	
The state of the s	Apnea Ventilation Smart Suction Manual Breath
	Insp/ Exp Hold ETCO <sub>2</sub> Measurement
	Nebulization Waveform Freeze
Monitoring	
Pressure Value:	Ppeak, Pplat, Pmean, Pmin, PEEP
Volume / Flow Value:	Vti, Vte, MV, MVspont
Time Value:	ftotal,fspont, I:E
Real Time Curves:	Pressure-Time, Flow-Time, Volume-Time waveforms
	Pressure-Volume, Volume-Flow, Flow-Pressure loops
Gas Monitoring:	FiO <sub>2</sub> , ETCO <sub>2</sub>
Calculated Values:	Compliance(C)
	Resistance(R)
	MVleak
	RSBI
	WOB
	PEEPi
Alarm	
	Paw high / low MVe high / low Circuit disconnned
	FiO <sub>2</sub> high / low Inspiration / Expiratory tidal volume low
	High Respiration Rate Apnea AC Failure Nebulizer On
	Low Battery Air /O <sub>2</sub> supply down High / Low PEEP
	Leakage out of range Occlusion
Technical Data	
• Screen:	12" TFT color touch screen (detachable)
Supply Gas:	O <sub>2</sub> , 0.28~0.6 MPa
Power Supply:	AC100~240 V, 50 Hz/60 Hz
Communication Interface:	RS-232 Port, Nurse call Port, Ethernet Port
Dimension (WxDxH):	322 mm x 375 mm x 366 mm (Main Unit)
ee (e)	547 mm x 675 mm x 950 mm (Cart)
Weight:	12.5 kg (Main Unit)
	25 kg (Cart)
	==g (50.1)

 $Remark: Above \ configurations \ include \ standard \ and \ optional. \ Please \ check \ price \ with \ your \ Aeonmed \ sales \ representative.$ 



0: No.4 Hangfeng Road, Fengtai Science Park. Fengtai District, Beijing, China (100070) cience Park: No. 10, Chaobai Street, Yanjiao Development Zone, Sanhe City, Habri Poving China (DRS201)

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rmation contained in this leaflet is correct at the date of publication. The policy of Beijing Aeonmed Co., Ltd. is one of continued improvements to its products cause of this policy Beijing Aeonmed Co., Ltd. reserves the right to make any changes, which may affect the information in this leaflet without giving prior notices

out giving prior notices. AVT70-1708



### Superior Mobile ICU ventilator

- Comprehensive ICU ventilator including BIVENT and PRVC
- · Compact, big capacity battery, no air compressor, intra-hospital mobility
- Flexible device configuration: equipped on a trolley, bed or ceiling pendant

### Cost Effective Solution

- Unique metal-based, autoclavable, heated exhalation valve
- Built-in flow sensor, non-consumable design
- Upgradeable ventilation system software, with an available USB port















#### An Optimal Combination of Invasive and Noninvasive Ventilator

- As noninvasive ventilation is used increasingly in a wide range of clinical situations, we offer a dual solution
- VG70 combines the advantages of a flexible noninvasive ventilator with a full-featured invasive ventilator for the ICU

## Optimal patient-ventilator synchrony, increase patient comfort

- The Unique Leak Compensation System Keep precise control on the tidal volume of each breath delivered to the patient by adjusting compensation dosage automatically
- Advanced Trigger Technique Enhance sensitivity, avoid spurious triggering

Auto-detect and Adjust Leak Compensation Automatically Adapt to Patient's Breathing Pattern

Multi-paramete Monitoring

## Safe Ventilation Through Whole Treatment Phase

### **Initial Treatment Phase**

- Noninvasive ventilation mode associated with decreased intubation rates, shortened patient stays, improved patient comfort, and a reduced risk of cross infection
- Preset patient's height and IBW. Reduce clinician's workload

## Stable Condition Phase

- PRVC and BIVENT employ lung-protective strategies, delivering intelligent ventilation
- Comprehensive lung mechanics monitoring include compliance, airway resistance, PEEPi and time constant
- Three waveforms & three loops with user-friendly display provide a continuous monitoring of the patient's condition

## Weaning Phase

- Various ventilation modes enhance the weaning process
- The unique trigger and leakage compensation system safeguards each and every patient breath resulting in smooth and comfortable breathing, avoiding extra workload on the patient and promoting recovery
- RSBI and WOB provide accurate reference for weaning

#### Rehab Phase

- Data export port provides connection to hospital monitors and Patient Data Management Systems
- · Provides pressure support for the patient when spontaneous breathing is present













截止2020-02-10 08:00 全国数据统计

较昨日+3073

较昨日+4008

较昨日+296

较昨日+97

较昨日+632

**40235** 确诊 **23589** 疑似

6484

909

3283

在 确 诊 患 者 病 例 中 <u>重 症 患 者</u> 达 到 6 4 8 4 例

哀悼武汉疫情死难者,致敬全国抗疫一线医护!

# 重症、危重症病人 呼吸支持治疗难点在哪里?

2月4日15时,国家卫建委召开新闻发布会,在回答香港无线台记者就香港一名39岁新型冠状病毒死亡中表示称,确实可能存在青壮年患病后病情急速发展,在短短的几个小时内急速恶化,且难以救治,最终死亡的情况。

对此,《新型冠状病毒感染的肺部炎症诊疗方案(试行第5版)》强调:

- 当患者接受标准氧疗后呼吸窘迫或低氧血症无法缓解 时,可考虑使用高流量鼻导管氧疗或无创通气。
- 若短时间内(1-2小时)内病情无改变甚至恶化,应当及时进行气管插管和有创机械通气。
- 同时对于重症ARDS患者,建议进行肺复张。

## 数据来源

丁香园.丁香医生新型冠状病毒感染肺炎疫情实时动态 国家卫健委2月4日15时新闻发布会

AEOMED 谊安 产品可靠 服务贴心

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传真: 010-83681616-8130

网站: www.aeonmed.com



## 从无创-有创-无创 提供全方位治疗手段

- 从轻症患者的无创通气治疗,再到重症患者或无创通气 不耐受患者的有创通气治疗,一台呼吸机全部解决。
- 减少医务人员因为更换机器带来的设备消毒等工作量,同时也可以避免更换机器带来的病人间交叉感染。

## ■独特的屏机分离技术 方便医护人员操作



- 显示器屏幕可以与气路平台完全分离,方便医护人员操作,同时在操作过程中可以增加与患者的距离,保证医护人员的安全。
- 可以做到真正做到专机专用,最大程度的降低院内交叉感染的几率。

# 摆脱气源依赖 转运 更安全



当患者需要进行相关影像学检查时,可以承担患者转运途中的呼吸支持工作。对于此次疫情需要转运评估治疗效果的患者可以做到安全快捷的转运。

# 具备肺复张工具 满足严重ARDS患者治疗需要

对于急性呼吸窘迫综合征的患者可以帮助其复张"湿区"的肺泡,同时单次有效的设计可以避免"干区"肺泡的过度膨胀,对临床医生治疗患者起到帮助作用。

# 呼吸力学测量 让患者肺"一目了然"

VG70呼吸机可以为患者进行相关肺功能测量,为评估患者治疗效果提供一定的帮助,保证医生能够更好的制定患者治疗方案及机械通气方案。



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#### 历经考验 精益求精

Shangrila590P融合智能的操作方式,带给客户更完美的操作体验;更舒适的通气模式,让患者获得更多的通气保护与支持;独具特色的免维护气路设计,更能有效降低客户使用和维护成本。

- 12.1"高清彩色触摸显示屏+飞梭旋钮双操作模式,轻松完成参数设置;
- 高精度的20~2500mL宽范围潮气量控制技术,带给成人与小 儿患者更多通气保障;
- · 一体化恒温主动式呼气阀技术, 非耗材设计, 杜绝隐性成本;
- 有创与无创通气类支持,实现完美的序贯通气;
- 智能的PRVC、BIVENT通气模式,提供有效肺保护策略,具有更好的人机同步协调性和通气舒适度;
- •丰富的临床实用工具:同步雾化、吸/呼气保持、智能增氧;
- 实用的波形工具、呼吸环、肺功能测量工具,时刻关注患者病情发展;
- \* 360度全景报警显示,提供实时监控与保护。

## 完善的参数监测和事件记录

参数监测

事件 记录

肺功能测量





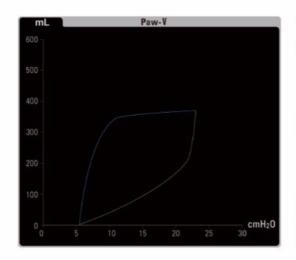


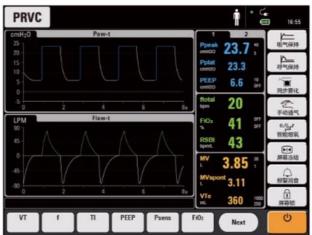
吸/呼 气保持 同步雾化

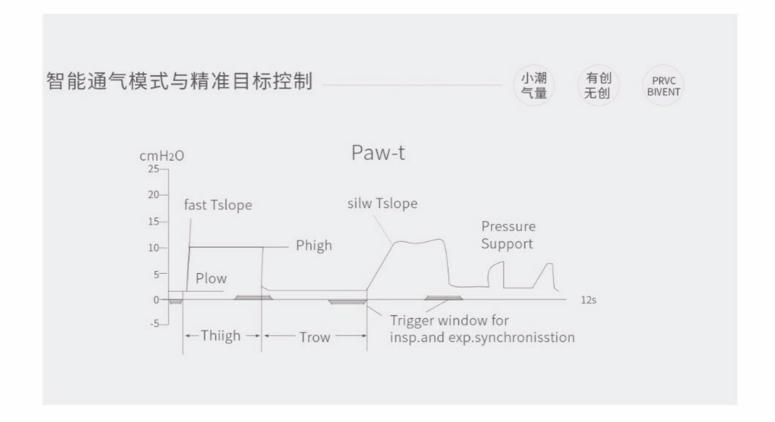
智能增氧

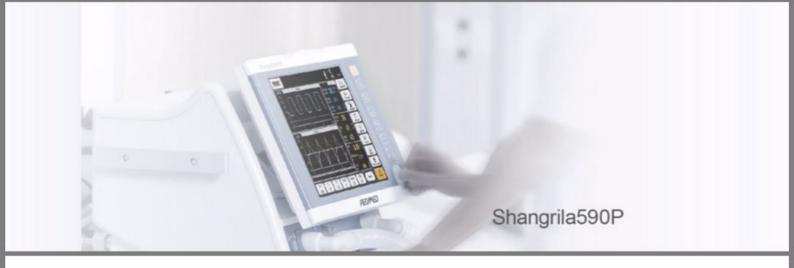
波形冻结

波形与 环形工具









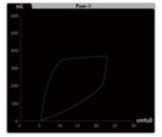


- The Shangrila590P ventilator ensures reliable, safe and high quality ventilation. 590P delivers excellent ventilation for both pediatric and adult patients
- 12.1" TFT touch color screen with navigation knob provide intuitive interface

 Detachable screen reduces the possibility of cross-infection

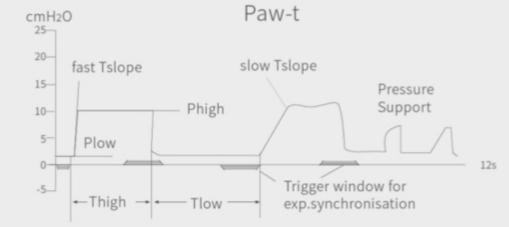


- Both Invasive & Non-invasive ventilation modes cover all types of patients need
- Supports external communication
   RS232 port enables external communication





- Built-in flow sensor, non-consumable, low maintenance cost
- Durable integrated exhalation valve, easy to sterilize







## **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. 01

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

EC-Representative: Shanghai International Holding Corp. GmbH

(Europe)

Eiffestraße 80, 20537 Hamburg, GERMANY

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. See also notes overleaf.

Report No.: BJ19859044

 Valid from:
 2019-07-16

 Valid until:
 2022-05-03

Date, 2019-07-16

Stefan Preiß

1. Punil

Head of Certification/Notified Body

Page 1 of 2

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



## **EC Certificate**

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

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REPUBLIC OF CHINA







