











**Quantum**  
Commodities

MEDICAL SUPPLIES







# Medical Devices

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00001	SHANG HAI		(CT) whole body x-ray computed tomography spiral scanner	United Imaging 528	FDA NO. K183143 	0.28 unit/ton	part 1: 2424*1158*2324mm, net weight 1735kg gross weight 2000kg part 2: 2971*938*1234mm net weight 481kg gross weight 734kg part 3: 948*748*1450mm net weight 260kg gross weight 330kg part 4: 2058*988*1059mm net weight 125kg gross weight 223kg part 5: 1508*1058*1169mm net weight 120kg gross weight 205kg part 6: 855*845*930mm net weight 82kg gross weight 89.5kg total net weight: 2803kg total gross weight: 3581.5	In stock / Air transport	yes
00002	SHANG HAI		FlatDR medical diagnostic x-ray machine (DR)	United Imaging 266i	Chinese NMPA (National Medical Products Administration) attached: 沪械注准 20182300076  Same brand passes FDA	0.91 unit/ton	part 1: 2424*1158*2324mm, net weight 1735kg gross weight 2000kg part 2: 2971*938*1234mm net weight 481kg gross weight 734kg part 3: 948*748*1450mm net weight 260kg gross weight 330kg part 4: 2058*988*1059mm net weight 125kg gross weight 223kg part 5: 1508*1058*1169mm net weight 120kg gross weight 205kg part 6: 855*845*930mm net weight 82kg gross weight 89.5kg total net weight: 2803kg total gross weight: 3581.5	In stock / Air transport	yes







# Medical Devices

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00004	SHEN ZHEN		ECG machine	EDAN SE-1200 Express	<p>FDA NO. K160876</p> 	100 units/ton	5kg/Net weight unit(without battery and paper) 10kg/Gross weight unit 420*330*105mmv	In stock / Air transport	yes
00008	SHEN ZHEN		B-ultrasound	Sonoscape/E2	<p>FDA NO. K170303</p> 	77 units/ton	6.5kg/Net weight unit 13kg/Gross weight unit 378*352*114mm	In stock / Air transport	yes







# Ventilators

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00005	NAN JING		Invasive Ventilator	PA700b	CE Certification attached 	12 units/ton	77kg (with PA air compressor attached) net weight 83kg (gross weight) 1240*670*600mm	In stock / Air transport	yes
00006	JIANG SU		Non-invasive Ventilator	Yuwell YH830	Emergency Use Authorizations (EUA) products, or more information see FDA official site <a href="https://www.fda.gov/media/136702/download">https://www.fda.gov/media/136702/download</a> CE certification attached 	33.3 units/ton	25kg/unit (net weight) 30kg/unit (gross weight) 630*630*1050mm	In stock / Air transport	yes
00007	JIANG SU		Sputum suction	Yuwell/7E-A	Same brand passes FDA EUA CE certification attached 	196 units/ton	227 units/ton 4.4kg/unit (net weight) 5.1kg/unit (gross weight) 280*196*285mm	In stock / Air transport	yes



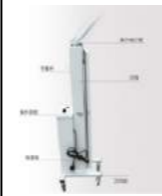

## Monitoring Devices

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00003	CN SHEN- ZHEN		Multifunctional ECG monitor	Midray/ Epm10	FDA NO. K191769 	188.6 units/ton	4kg/unit (net weight) 5.3kg/unit (gross weight) 269*252*159mm	In stock / Air transport	yes
00009	CN SHENZHEN		Central monitoring system	Mindray/ BeneVision	FDA NO. K183238 	N/A	N/A	In stock / Air transport	yes
00010	U.S.A		Defibrillator monitor	Zoll M series	FDA NO. K180482 	163.9 units/ton	5.23kg/unit (net weight with battery) 6.1kg/unit (gross weight) 350*320*450mm	In stock / Air transport	yes







# Hospital/Emergency Room Equipment

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00011	CN SHAN-DONG		ICU multifunctional rescue bed	Mingtai/Z1	CE Certification attached 	8.1 units/ton	110kg/unit (net weight) 123kg/unit (gross weight) 2000*720*630mm	In stock / Air transport	yes
00012	SHAN DONG		Operating table	Ming-tai/MT2000	CE Certificatio attached 	3.2 units/ton	210kg/unit (net weight) 308kg/unit (gross weight) 1550*850*950mm	In stock / Air transport	yes
00013	SHENZ HEN		Double headed shadowless lamp	Comen/L6	MHRA (The U.K.) certification attached 	4.8 units/ton	185kg (net weight) 205kg (gross weight) 1465*1095*960mm	In stock / Air transport	yes

# Hospital/Emergency Room Equipment

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00014	SHAN DONG	 A7 AS型挂床床架冲孔钢板床架	Ward manual double rocking nursing bed	Lekang/A7	CE Certification attached 	12.7 units/ton	64kg/unit (net weight with handles, wheels and mattress) 79kg (gross weight) 2080*823*690mm	In stock / Air transport	yes
00015	SHAN DONG		UV disinfection truck	Mingtai/M-DV1000	CE Certification attached 	232.5 units /ton	3.4kg/unit (net weight) 4.3kg/unit (gross weight) 310*260*843mm	In stock / Air transport	yes

## Miscellaneous

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00016	JIANGSU		Digital film printer	Hu.q/HQ-460DY	CE Certificator attached 	14.3 units/ton	1 / Box 70kg/unit (gross weight) 840*710*610mm	In stock / Air transport	yes
00017	JIANGSU		Digital film	Hu.q/HQ-460DY	CE Certification attached 	40 cases/ton (14"*17") 55.5 cases/ton (11"*14") 71.4 cases/ton (10"*12") 90.9 cases/ton (8"*10")	100 sheets/box 5 boxes/case 470*240*570mm per case 25kg per case (14"*17",350*430mm ) 18kg per case (11"*14",280*350mm) 14kg per case(10"*12",250*300mm) 11kg per case(8"*10",200*250mm)	In stock / Air transport	yes
00018	SHANDONG		Autoclave	Shinva/ MOST-T80	CE Certification attached 	5 units/ton	5 units/ton 180 kg/unit (net weight) 200 kg/unit (gross weight) 1180*890*735mm	In stock / Air transport	yes



For detailed certifications, see  
sheet "certifications"

MINGTAI



ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ ZERTIFIKAT ◆ CERTIFICATE



ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ ZERTIFIKAT ◆ CERTIFICATE

证书附件—证书号: EU143518

产品名称	类型/型号
	AT-6732E/F, AT-85001A, AT-85001B, AT-85001E, AT-85001F, AT-85002A, AT-85002B, AT-85002A, AT-85002B, AT-75002A, AT-75002B, AT-85002E, AT-85002F, AT-80002E, AT-80002F, AT-75002E, AT-75002F, AT-87003A, AT-87003B, AT-79003A, AT-79003B, AT-62005A, AT-62512A, AT-62512B, AT-67312A, AT-67312B, AT-62512E, AT-62512F, AT-67312E, AT-67312F, AT-67516A, AT-67516B, AT-74616A, AT-74616B, AT-67516E, AT-67516F, AT-74616E, AT-74616F, AT-62516A, AT-62516B, AT-62516E, AT-62516F, AT-82021B, AT-61025A, AT-61025B, AT-66025A, AT-66025B, AT-61025E, AT-61025F, AT-66025E, AT-66025F, AT-98028, AT-87031A, AT-87031B, AT-87031E, AT-87031F, AT-79031A, AT-79031B, AT-72031A, AT-72031B, AT-79031E, AT-79031F, AT-72031E, AT-72031F, AT-66002A1, AT-62032A, AT-115056J, AT-115056S, AT-61056J1, AT-61056S1, AT-61056J2, AT-61056S2, AT-61056J3, AT-61056S3, AT-61056J5, AT-61056S5, AT-61056J6, AT-61056S6, AT-83071A, AT-83071B, AT-75071A, AT-75071B, AT-83071E, AT-83071F, AT-79071E, AT-79071F, AT-62582A, AT-62582B, AT-67382A, AT-67382B, AT-62582E, AT-62582F, AT-67382E, AT-67382F, AT-78060B, DZ-62512B, DZ-67516A/B, DZ-74616A/B, DZ-83071A/B, E-NT-85001CS3/DN3, CT-85001CS/D5, DZ-62512F1, Z1, Z2, Z3, Z4, Z5, Z6, Z7, Z8, Z9, Z21, Z22, Z23, Z24, Z25, Z26, Z27, Z28, Z29
转运车/对接车	D1, D2, D3, D4, D5, D6, D7, D8, D9, D10, D11, D12, D13, D14, D15, D16, D17, D18, D19, D20
床车	

附件结束



For detailed certifications, see  
sheet "certifications"

**YUWELL**

ZERTIFIKAT ♦ CERTIFICATE ♦ 認證證書 ♦ CERTIFICADO ♦ CERTIFICAT

  
Product Service

**EC Certificate**  
Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)  
No. G2 15 08 55329 013

**Manufacturer:** Jiangsu Yuyue Medical Equipment & Supply Co., Ltd.  
Yunyang Industrial Park  
212300 Danyang, Jiangsu  
PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:** Shanghai International Holding Corp. GmbH (Europe)  
Elifstraße 80  
20537 Hamburg  
GERMANY

**Product Category(ies):** Electric Suction Apparatus,  
Oxygen Concentrator,  
Air Compressive Nebulizer  
Electronic Blood Pressure Monitor

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

**Report No.:** SH15392EXT01

**Valid from:** 2015-11-11  
**Valid until:** 2020-11-10

**Date:** 2015-09-17

  
Hans-Heiner Junker





TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123  
Page 1 of 2

TÜV SÜD Product Service GmbH · Zertifizierstelle · Riederstraße 65 · 80339 München · Germany 

ZERTIFIKAT ♦ CERTIFICATE ♦ 認證證書 ♦ CERTIFICADO ♦ CERTIFICAT

  
Product Service

**EC Certificate**  
Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)  
No. G2 15 08 55329 013

**Facility(ies):** Jiangsu Yuyue Medical Equipment & Supply Co., Ltd.  
Yunyang Industrial Park, 212300 Danyang, Jiangsu,  
PEOPLE'S REPUBLIC OF CHINA

Page 2 of 2

TÜV SÜD Product Service GmbH · Zertifizierstelle · Riederstraße 65 · 80339 München · Germany 

## For detailed certifications, see sheet "certifications"

### COMEN

 Medicines & Healthcare products  
Regulatory Agency

 **MHRA**

registration. As the updated information does not affect your regulatory obligations or the information published on our Public Access Registration Database (PARDB).

Thank you for registering the following generic groups of devices:

**Class I Devices:**  
*Dental Lights*  
*Lamps (Ophthalmic Examination)*  
*Traction And Surgical Immobilisation Devices*  
*Medical Examination Luminaires*  
*Operating Tables And Accessories*  
*Surgical Instrument Accessories*  
*Autoclave Accessories (E.G. Trays And Tray Lifters, Shelves, Racks)*  
*Airway Devices/Monitoring Equipment And Accessories*  
*Electrosurgical Accessories(e.g. Transient Invasive Electrodes, Footswitches)*

**Custom Made Devices:**  
*None*

**Products Covered By Article 12:**  
*None*

**Confidentiality**

Please note that in accordance with Directive 2007/47/EC as of 21st March 2010 information on the registration of persons responsible for placing devices on the market will no longer be treated as confidential and the Competent Authority will provide third parties with information on the name and address of manufacturers and authorised representatives and their devices that have been registered. However the names of individuals, their telephone numbers and email addresses will remain confidential unless you have chosen to trade using personal details. This change only applies to medical devices and does not affect In Vitro Diagnostic devices registration, which remain confidential under Article 19 of the In Vitro Diagnostic Directive 98/79/EC.

**If your company name or that of a manufacturer that you represent is based on an individual's personal name it will be published unless you inform the MHRA that you would like the company name to remain confidential.**

**Likewise, if your company address or that of a manufacturer that you represent is the personal home address of an individual it will be published unless you inform the MHRA that you would like the company address to remain confidential.**

Should you have any queries regarding your registration please do not hesitate in contacting us.

Yours sincerely

  
 Malcolm Ridgway  
 Data Integrity Support Officer

Regrepmdd Vers 3 June 2015

 Medicines & Healthcare products  
Regulatory Agency

 **MHRA**

**Our Ref: CA017365**

My Peter Wei  
 Lotus Global Co Ltd  
 1 Fourseasons Terrace  
 West Drayton  
 Middlesex  
 UB7 9GG

**MHRA**  
 10 South Colonnade  
 Canary Wharf  
 London  
 E14 4PU  
 United Kingdom

**28 August 2019**

Dear Mr Wei

**MEDICAL DEVICES REGULATIONS 2002: REGULATION 19**  
**Registration of Persons Placing General Medical Devices on the Market**

Thank you for informing the Competent Authority of the details of **Manufacturers Name:- Shenzhen Comen Medical Instruments Co.,Ltd** located at **Manufacturers Address:- No 2 of FIYTA Timepiece Building, Nanhuan Avenue, Gongming sub-district, Guangming New District, Shenzhen Guangdong China 518106** for whom you are acting as the authorised representative and for supplying the medical device information.

**Your registration has been recorded based on your declaration that you have determined that the device(s) fall within the definition of "medical device", and that you have classified it/them as falling within Regulation 19 taking into account the intended purpose(s) and mode(s) of action. In accepting your registration, I should make clear that the Competent Authority does not examine each individual notification and therefore cannot and does not necessarily endorse these determinations. Neither does this letter represent any form of accreditation, certification or approval by the UK Competent Authority.**

Your registration is based upon your declaration on the RG2 form and means that:

**For Manufacturers of Class I medical devices, Assemblers, and Sterilisers**  
 You should now be operating under the Medical Devices Directive and the above Regulations for the products you asked us to register, by fully complying with the essential requirements, CE marking those products or labelling them as such.

**For Manufacturers of Custom-made devices and Custom Made Active Implantable**  
 You should be ready to claim compliance with the Directive and Regulations and should be manufacturing custom-made devices in accordance with their requirements.

**If you stop placing devices on the market or if you are not complying with the Regulations you should inform us so that we can amend our records. You should be aware that it is an offence to place on the market CE marked devices that do not comply with the regulations.**

The information you provided has been recorded against the reference number shown at the top of this letter, which we ask you to quote in all future correspondence and communications.

**Please inform us of the following chargeable changes:**

- the company information e.g. name and address
- additional generic groups of devices (not individual products within an existing generic group)
- Change of authorised representative

Please also use the Devices Online Registration Database (DORS) to tell us of the following changes e.g. removal/discontinuation of a device from your registration record, change of contact person, postcode, telephone number and/or email address, for which payment of our statutory fee does not apply. Though, you are required to provide these non-chargeable changes in writing we will not provide an updated letter of

For detailed certifications, see  
sheet "certifications"

## LEKANG

File No: CE-TCF-001



**EC Declaration of Conformity**

Regarding Medical Device Directive(93/42/EEC)

Including Directive 2007/47/EC

**Manufacturer**  
 Manufacturer: Qufu LeKang Medical Technology Co., Ltd.  
 Address: North Xishou Road, Kongzi Avenue, Qufu City, Shandong Province

**Product**

Product Name	Type
Shadowless	LKZF-700/500, LKZF-700/700, LKZF-500/500, LKZF-700, LKZF-500, LKZF-700L, L
Operating Lamp	LKZF-500L, LKZF-200L, LKLED-700/500, LKLED-700/700, LKLED-500/500, LKLED-700, LKLED-500, LKLED-700L, LKLED-500L, LKLED-200/200, LKLED-200, LKLED-200L, LKLED-5+5, LKLED-5+3, LKLED-3+3, LKLED-5, LKLED-3, LKLED-3L, LKLED-3L
Operating Table	LKDS-I, LKDS-II, LKDS-III, LKDS-IV, LKDS-V, LKDS-VI
Medical Beds	LK-A1, LK-A2, LK-A3, LK-A4, LK-A5, LK-A6, LK-A7, LK-A8, LK-A9, LK-A10, LK-B1, LK-B2, LK-B3, LK-B4, LK-B5, LK-B6, LK-B7, LK-B8, LK-B9, LK-B10

Classification: I  
 Rule: According to Rule 12

We confirm the above products can meet the requirement of Medical Device Directive and the following harmonized standards:

EN ISO 14971:2012  
 EN ISO 15223-1:2016  
 EN 1041:2006 +A1:2013  
 EN 60601-1:2006/A1:2013  
 EN 60601-1-2:2015  
 EN 60601-2-41:2009  
 EN 60601-2-48:2011

Signature:   
 Date: 2017.11.22





# For detailed certifications, see sheet "certifications"

## United Imaging



January 3, 2019

Shangri United Imaging Healthcare Co., Ltd.  
Shumei Wang  
QM & RA VP  
No. 2258 Chengbei Rd., Jiading Industrial District  
SHANGHAI, 201807 CHINA

Re: K183143  
Trade/Device Name: uCT 520, uCT 528  
Regulation Number: 21 CFR 892.1750  
Regulation Name: Computed Tomography X-Ray System  
Regulatory Class: Class II  
Product Code: JAK  
Dated: November 14, 2018  
Received: November 15, 2018

Dear Shumei Wang:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/e2ltools/cfpnm/pnm.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the [Federal Register](#).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's

U.S. Food & Drug Administration  
10953 New Hampshire Avenue  
Silver Spring, MD 20903  
[www.fda.gov](https://www.fda.gov)

国产医疗器械产品（注册） <span style="float:right">返回</span>	
注册证编号	沪械注准20182300076
注册人名称	上海联影医疗科技有限公司
注册人住所	上海市嘉定区城北路2258号
生产地址	上海市嘉定区城北路2258号
产品名称	数字化医用X射线摄影系统
管理类别	第二类
型号规格	uDR 266i (配置1、配置2)
结构及组成/主要组成成分	产品由系统柜组件（包括高压发生器和电源分配单元）、X射线管组件、限束器、控制台、机架、移动床、无线数字平板探测器、图像处理系统、显示器、滤线栅、拼接支架组成。
适用范围/预期用途	产品供医疗机构通过X射线对人体进行摄影获得X射线图像，用于医学影像诊断。
产品储存条件及有效期 /	
附件	
其他内容	
备注	审批部门 上海市食品药品监督管理局
	批准日期 2018-02-27
	有效期至 2023-02-26
变更情况	
	注 网站发布的医疗器械产品注册和产品备案信息供公众查询,如网站公布的信息与原纸质批件不一致,请联系相应部门纠错。进口和国产三类医疗器械数据问题请发送邮件至国家药监局医疗器械数据纠错邮箱: qixiejucuo@nmpa.gov.cn【邮件主题请注明“医疗器械数据问题”, 邮件正文中请填写以下全部信息: 1.医疗器械注册号/备案号; 2.类型(注册、变更、延续等); 3.问题描述(500字以内); 4.企业名称(全称); 5.统一社会信用代码; 6.联系人姓名; 7.联系电话(手机和座机); 8.联系邮箱】。或致电国家药监局医疗器械数据纠错电话010-88331951(此电话为我局医疗器械数据纠错联系电话,并非相应产品/企业业务咨询电话)。国产一类和二类医疗器械数据问题请联系企业所在地省局,由省局通过数据共享平台进行纠错和维护。

## For detailed certifications, see sheet "certifications"

### EDAN



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

February 15, 2017

Food and Drug Administration  
10903 New Hampshire Avenue  
Document Control Center - WO66-G609  
Silver Spring, MD 20993-0002

Edan Instruments, Inc.  
% Doug Worth  
Sr. Dir. US RA/QA  
Edan Medical  
1200 Crossman Ave. Suite 200  
Sunnyvale, California 94089

Re: K160876

Trade/Device Name: Electrocardiograph, Models:  
SE-12, SE-12 Express, SE-1200, and SE-1200 Express  
Regulation Number: 21 CFR 870.2340  
Regulation Name: Electrocardiograph  
Regulatory Class: Class II  
Product Code: DPS  
Dated: January 5, 2017  
Received: January 9, 2017

Dear Doug Worth:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the [Federal Register](#).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in

For detailed certifications, see  
sheet "certifications"

## SONOSCAPE



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
10903 New Hampshire Avenue  
Document Control Center - W036-G609  
Silver Spring, MD 20993-0002

Sonoscape Medical Corp.  
% Ms. Toki Wu  
Regulatory Affairs Manager  
4/E, 5/E, 8/F, 9/F & 10/F, Yizhe Building, Yuquan Road, Nanshan  
Shenzhen, Guangdong 518051  
CHINA

May 19, 2017

Re: K170303  
Trade/Device Name: E5 Series Digital Color Doppler Ultrasound System  
Regulation Number: 21 CFR 892.1550  
Regulation Name: Ultrasonic pulsed doppler imaging system  
Regulatory Class: II  
Product Code: IYN, IYO, ITX  
Dated: February 20, 2017  
Received: March 1, 2017

Dear Ms. Wu:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.



For detailed certifications, see sheet "certifications"

**PUAO(PA700B)**



## For detailed certifications, see sheet "certifications"

### MINDRAY



October 31, 2019

Shenzhen Mindray Bio-medical Electronics Co., LTD.  
Yanhong Bai  
Manager Regulatory Affairs, Technical Regulation Department  
Mindray Building, Keji 12th Road South,  
Hi-tech Industrial Park, Nanshan  
Shenzhen, 518057 Cn

Re: K191769  
Trade/Device Name: ePM Series Patient monitors  
Regulation Number: 21 CFR 870.1025  
Regulation Name: Arrhythmia Detector And Alarm (Including ST-Segment Measurement And Alarm)  
Regulatory Class: Class II  
Product Code: MHX, DSL, MLD, DRT, DXN, DSK, FLL, DQA  
Dated: September 30, 2019  
Received: October 1, 2019

Dear Yanhong Bai:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pmn.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

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Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal

U.S. Food & Drug Administration  
10903 New Hampshire Avenue  
Silver Spring, MD 20993  
[www.fda.gov](http://www.fda.gov)



March 13, 2019

Shenzhen Mindray Bio-medical Electronics Co., LTD.  
Yanhong Bai  
Manager Regulatory Affairs, Technical Regulation Department  
Mindray Building, Keji 12th Road South  
Hi-tech Industrial Park, Nanshan  
Shenzhen, 518057 Cn

Re: K183238  
Trade/Device Name: BeneVision Central Monitoring System  
Regulation Number: 21 CFR 870.2300  
Regulation Name: Cardiac Monitor (Including Cardiachometer And Rate Alarm)  
Regulatory Class: Class II  
Product Code: MSX, DRQ, MHX, DRT, DXN, DQA, DSB  
Dated: February 7, 2019  
Received: February 8, 2019

Dear Yanhong Bai:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pmn.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the [Federal Register](#).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal

## For detailed certifications, see sheet "certifications"

### ZOLL



November 30, 2018

ZOLL Medical Corporation  
Pooja Dalvi  
Regulatory Affairs Specialist  
269 & 271 Mill Road  
Chelmsford, Massachusetts 01824-4105

Re: K180482

Trade/Device Name: ZOLL Propaq M  
Regulation Number: 21 CFR 870.1025  
Regulation Name: Arrhythmia detector and alarm (including ST-segment measurement and alarm)  
Regulatory Class: Class II  
Product Code: MHX, DRT, DXN, DSK, CCK, DQA, DPS, FLL  
Dated: October 22, 2018  
Received: October 23, 2018











Dear Pooja Dalvi:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database located at <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pmn.cfm> identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.











If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the [Federal Register](#).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part





# Protective Gear

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00019	HE NAN		COATS FOR Operating room	Scrubs. Surgical gown Medium size. Fabric construction non-woven, disposable SMMS type with loop closure. Fluid resistant bodysuits, sleeves with reinforced elasticized cuffs from 5 to 10 cm width, minimum gown length 120 cm			50 /box 60*40*45cm	1000000 / 7 day	1000000
00020	HE NAN		SHOES COVER	Operating rooms use, TNT, Large size with anti-slip to the traction, disposable			26/5000 8 / bag 200 / box 60*40*40cm 7 kg	1000000 / 7 day	1000000
00021	HE NAN		SURGICAL HAT	Made of non-woven fabric. With fold-over rubber, fluid resistant. Disposable.			20 / bag 1000 / box 60*40*40cm 7 kg	1000000 / 7 day	1000000
00022	JIANG SU		RUBBER BOOTS	White rubber boot with insole with treatment non-slip, in natural rubber, double PVC material lined by inside in rubberized fabric, sanitized vulcanized in one piece resistant to alcohol and all kinds of chemicals, long spout (40 cm high) Numbers 36 to 44	certificate will be added soon		36-44 Size	100 / day	100
00023	JIANG SU		DISPOSABLE CARPET	Disposable Sheet Carpet, Film Overlapping Sheet Block polyethylene with antibacterial adhesive layer with last sheet adhesive for fixing to the ground. Blue in colour and measures 66x113cm. 30 sheets per carpet / 10/8 carpets per box	certificate will be added soon		65*115cm	10 / day	10
00024	ZHE JIANG		Protective suit	Disposable Waterproof Hooded Jumpsuit: Protective Clothing Category III; Tyvek Classic Xpert LA - DuPont. Size M, G OR GG			1piece / bag 50 / box	1000000 / 7day	1000000



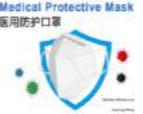

# Gloves

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00025	SHAN DONG		GLOVE FORRECOGNITION	Medium and Large Size, Latex resistant -long spout. 240 mm + -5mm unbreakable in the fit test, with absorbent powder. With flange			100 pieces/box, 10 boxes/box. 2100 ourter carton; 46100 ourter carton		1000000
00026			SURGICAL GLOVE	N ° 6- ½ and 7 in pairs, latex, thickness 0.2 to 0.3 mm, non-slip, Optimum sensitivity, unbreakable in the shim test with absorbent powder. 270 mm length, variation + - 10mm to be measured from the end top of middle finger to cuff edge.			100 pieces/box, 10 boxes/box. 2100 ourter carton; 46100 ourter carton		1000000
00027	SHAN DONG		NITRILE GLOVES	Nitrile Gloves, non-sterile, ambidextrous, dust-free, 30 cm which They feature a beaded cuff. Size M					100
00028	SHAN DONG		GLOVE FORRECOGNITION	Ambidextrous, dust-free 30 cm nitrile gloves featuring a beaded cuff. Size S			100 pieces/box, 10 boxes/box. 2100 ourter carton; 46100 ourter carton		100
00029	SHAN DONG		GLOVE FORRECOGNITION	Ambidextrous, dust-free 30 cm nitrile gloves featuring beaded cuff. Size L			100 pieces/box, 10 boxes/box. 2100 ourter carton; 46100 ourter carton		50











## Eye Protection

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00030	TIAN JIN		FACE SHIELD	Made of transparent plastic /polycarbonate, it must provide good visibility, both at that carries it like the patient.		EN166	200 / box 68*38.3*41.5cm 13.87KG	1000000 /3day	1000000
00031	SHEN ZHEN		GOGGLES	Provides good sealing with face skin, PVC frame flexible to easily adapt to all face contours with even pressure, close your eyes and surrounding areas, should accommodate users with prescription glasses. Lenses must be Clear fog and scratch resistant plastic, adjustable band not to loosen during clinical activity, indirect ventilation to avoid fogging, reusable. EU STANDARD DIRECTIVE 86/686 / EEC. EN 166/2002 ; ANSI / ISEA Z87.1-2010			26/5000 8 / bag 200 / box 60*40*40cm 7 kg	1000000 /7 day	1000000

# Mask

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00032	HE NAN		surgical mask (non-sterilized)	Surgical mask, highly resistant to fluids, allows good breathability, the inner and outer faces must be clearly identified. Hypoallergenic with 3-fold filter attached to the ear. Fluid resistance at a minimum pressure of 120 mmhg according to ASTM F1862-07, ISO 22609 or equivalent. Filtration efficiency: ASTM F2101, EN 14683 ANNEX B or EQUIVALENT		YY 0469-2011 iSO 9001 ISO 13485	2000 / Box 52.5*49*56.5cm 11.5KG	1000000 / 5day	1000000
00033	SHENYANG		GOGGLES	N95 or FFP2 mask or higher Good breathability. NIOSH Compliant from the USA or FFP2 according to EN 149		GB19083-2010	500 / box 46.5 * 43.3 * 36.3 cm 4.5 KG	1000000 / 15 day	1000000

# Oxygen Delivery / Equipment








Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00034	SHEN ZHEN		BALL OR CYLINDER FOR 6M3 OXYGEN	00034-00051 medical gases are provided by Shenzhen Huateng Medical Engineering Equipment Co., Ltd	   			3500 / 7day	3500
00035	SHEN ZHEN		AIR MEDICINAL GENERATOR SYSTEM			20 / 7day	20		
00036	SHEN ZHEN		MEDICINAL MEDICAL VACUUM GENERATOR SYSTEM			300 / 7 day	300		
00037	SHEN ZHEN		VISUAL AND ACOUSTIC ALARM PANEL FOR MEDICAL GAS			3500 / 7day	1000		
00038	SHEN ZHEN		MEDICINAL OXYGEN REGULATOR WITH HUMIDIFIER			10 / day	3500		
00039	SHEN ZHEN		1,300 CC MINIMUM SUCTION BOTTLE			1000 / 7day	1000		







# Oxygen Delivery / Equipment

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00040	SHEN ZHEN		BALL OR CYLINDER FOR OXYGEN 1M3 ALUMINUM		00034-00051 medical gases are provided by Shenzhen Huateng Medical Engineering Equipment Co., Ltd			700 / 7day	700
00041	SHEN ZHEN		CONSOLE OUTLET FOR MEDICINAL GAS					3000 / 7day	3000
00042	SHEN ZHEN		VACUUM REGULATOR WITH TRAP OHMEDA TYPE				400 / 7day	400	
00043	SHEN ZHEN		VACUUM REGULATOR WITH TRAP OHMEDA TYPE				10 / 7day	10	
00044	SHEN ZHEN		OXYGEN MANIFOLD 5 + 5				10 / day	10	
00045	SHEN ZHEN		FLEXOMETER FOR OXYGEN CONNECTION OHMEDA TYPE				1200 / 7day	1200	







# Oxygen Delivery / Equipment

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00046	SHEN ZHEN		FLUXOMETERS FOR OXYGEN DISS INTERNATIONAL SYSTEM	00034-00051 medical gases are provided by Shenzhen Huateng Medical Engineering Equipment Co., Ltd	 			1300 / 7day	1300
00047	SHEN ZHEN		MEDICAL GAS PIPE			3000 / 7day	3000		
00048	SHEN ZHEN		OXYGEN HUMIDIFIER			500 / 7day	500		
00049	SHEN ZHEN		CONNECTION FOR MEDICINAL HOSES			600 / 7day	600		
00050	SHEN ZHEN		SECTORIZING BALL VALVE			200 / 7day	600		
00051	SHEN ZHEN		VACUUM REGULATOR WITH TRAP DISS I NTERNATIONAL SYSTEM			600 / 7day	600		

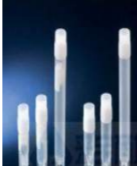





# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00052	Axygen/ Hangzhou		PCR TUBES	PCR Tube Strips for REAL TIME PCR EQUIPMENT, Strip8 tubes per strip compatible with Real Time PCR equipment Available in the LCSP (Applied Biosystem 7500 Thermal Cycler)	certificate will be added soon		1250pcs/Box	100box/day	160
00047	SHEN ZHEN		LID FOR MICROPLATE	Cap for microplate, CAP STRIP FOR PCR EQUIPMENT PLATES IN REAL TIME. Cap Strip, 8 caps per strip	certificate will be added soon		1250pcs/Box	100box/day	80
00048	SHEN ZHEN		RNA EXTRACTION KITS	Extraction Kit for RNA with support of silica gel from plasma, serum, cell-free fluids, and cell culture supernatant. Must contain columns, collection tubes, wash buffer and carrier RNA. It should not require heating anywhere in the process of extraction, do not use B mercapto-ethanol anywhere in the processing. It must contain all the reagents necessary for the extraction procedure. Insert required to verify technical specifications accompanying the offer. Kit for 250 determinations as a minimum				2000 kit / day	8000







# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00055	Hunan		RNA EXTRACTION KITS	Extraction kit for total nucleic acids with extraction equipment automated that can take from 1 to 96 samples at a time, with 4 pipette channels, with liquid level detector in tubes and plates, clot detector and drip detector. With security cabin with UV light and with opening lock during operation. Able to make the master mix for the polymerase chain reaction			24test/kit	2000 kit/day	3000
00056	Benjing		PACKING FOR SAMPLES TRANSPORT	Triple packaging to transport infectious samples in the form refrigerated, according to international standards (IATA).			11L	50/day	50
00057	Thermo Fisher/ Shanghai		MICRO CENTRIFUGE TUBE	Capacity 1.5 ml, with cap attached to the safelock system tube, for PCR, free of DNase, RNase and autoclavable pyrogens, with good user experience in the LCSP, through "Proof of use Satisfactory" issued by the Quality Management Unit of the LCSP and send sample			5000pcs/Box	50box/day	500











## Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00058	Thermo Fisher/ Shanghai		Cryovials POLYPROPYLENE, THREAD LID	STERILE CRYOVIALS, 2 ml, with external screw cap, polypropylene, autoclavable and sterile, send sample			600set/Box	100box/day	5000
00059	Changchun		INTAKING SAMPLES EQUIPMENT	EQUIPMENT FOR SAMPLE TAKING, Containing 2 ml. As minimum, liquid transport medium for viruses. For sample transport viral, with two polyester swabs or other synthetic material (not cotton) and rigid plastic stick. The tube must have a conical bottom, Sterile, without activated carbon			600set/Box	100box/day	5000
00060	Hunan		DETECTION OF COVID-19 BY PCR IN REAL TIME	in the management of equipment all the time required and in case of entry of new officials on campus. Submit a written program annual maintenance for the duration of the contract, which it must be included in the offer folder. The provider shall provide a written report of maintenance and / or repairs made to the equipment, so that the user registers the observations regarding operation and compliance with preventive and corrective maintenance. Notification time and response, to any inconvenience presented, should be immediately and within laboratory working hours and with staff qualified company. The supplier must put into operation the five laboratories within a maximum period of 20 (twenty) days once signed the contract			24test/kit	2000kit/day	10000a


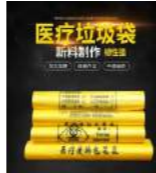



# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00061	Changchun		VIRUS TRANSPORT SOURCE GENEXPERT	<p>Sterile. Kit for taking and transporting viral samples, with swab and sterile means of transport. For the transportation of nasal samples, nasopharyngeal, ocular, dermal or mucous lesions.</p> <p>Sterile. Kit for taking and transporting viral samples, with swab and sterile means of transport. For the transportation of nasal samples, nasopharyngeal, ocular, dermal or mucous lesions.</p>			600set/Box	100box/day	2000
00062	Hunan		GENEXPERT	Kit for the determination of Sars-Cov-2 (OBS: price suggested by the OPS) US \$ 19.80			24test/kit	2000kit/day	50000
00063	Hunan		RESPIRATORY REAGENT FOR VIRUS	<p>Reagent kit for Coronavirus</p> <p>CoVD-2019 detection by Rt-PCR in real time. Compatible with thermal cycler equipment for time PCR real 96 sample plate format, with adapter for test strips tubes, LightCycler 480 II, available in service. Must include tubes for sampling, equipment for automated extraction of acids nucleic, with extraction kits, tips and tubes free of DNase and RNase. Preventive and corrective maintenance, as well as all supplies equipment consumables will be borne by the supplier.</p>			24test/kit	2000kit/day	384

# Laboratory Supplies



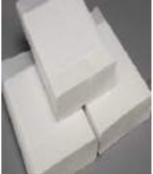





Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00064	3M/USA		STERILITY BIOLOGICAL CONTROL	Sterility Biological steam indicator. Self-contained ampoules with Geobacillus stearothermophilus to control autoclave spores maturity in two years			50pcs/kit	20kit/day	20
00065	3M/USA		STERILITY BIOLOGICAL CONTROL	Sterility Biological steam indicator. Self-contained ampoules with Geobacillus stearothermophilus to control autoclave spores maturity in two years			25test/kit (include 25 pcs of +- contrast reagent)	20kit/day	20
00066	3M/USA		STERILITY BIOLOGICAL CONTROL	Sterility Biological steam indicator. Self-contained ampoules with Geobacillus stearothermophilus to control autoclave spores maturity in two years			24test/kit (include 24 pcs of +- contrast reagent)	20kit/day	20
00067	3M/USA		STERILITY BIOLOGICAL CONTROL	Sterility Biological steam indicator. Self-contained ampoules with Geobacillus stearothermophilus to control autoclave spores maturity in two years			24test/kit (include 24pcs of +- contrast reagent)	20kit/day	20
00068	Zhuhai		ALCOHOL IN FOAM	Foam alcohol refill for dispenser			100grain/ 1Bottle)	1000Bottle/day	40

# Laboratory Supplies











Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00069	Shenzhen		MICROFIBER REPLACEMENT	Microfiber cloth replacement 40 cm	certificate will be added soon		400PCS/pack	1000pack/day	20
00070	Shanghai		PATHOLOGIAL WASTE BAG	Autoclavable polypropylene pathological waste bags. Measurements not exceeding 25 x 25 cm	certificate will be added soon		32cm*38cm, 100pcs/pack	1000pack/day	40
00071	Shanghai		PATHOLOGIAL WASTE BAG	Screen-printed autoclave bags with the danger symbol, transparent. 60 x 75 cm. 64-liter capacity Polypropylene.	certificate will be added soon		60*70cm, 100pcs/pack)	1000pack/day	100
00072	Tianjin		COOKING POT	High pot, with lid, capacity of 12 liters. 30 cm. diameter; stainless steel. triple bottom (stainless steel + aluminium + stainless steel) for autoclave	certificate will be added soon		9L	20	20
00073	Tianjin		COOKING POT	High pot, with lid, capacity of 9 litres. 30 cm. diameter stainless steel, triple bottom (stainless steel + aluminium + stainless steel) for autoclave	certificate will be added soon		12L	20	20













# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00074	Hebei		ABSORBENT PAD	Disposable absorbent pad with waterproof undercoat. 43 x 61 cm	certificate will be added soon		50pcs/pack	1000pack/day	8
00075	Beijing		SANITIZING TOWEL	Autoclavable polypropylene pathological waste bags. Measurements not exceeding 25 x 25 cm	certificate will be added soon		32cm*38cm, 100pcs/pack	1000pack/day	40
00076			PAPER TOWEL FOR KITCHEN DOUBLE LEAVES	Paper towel interspersed 22 x 21 cm minimum for laboratory use. 350 to 400 sheets x package			4000pcs/Box	100box/day	14000
00077	Thermo Fisher/ Shanghai		MICROCENTRIFUGA PIPES	Capacity 1.5 ml, with cap attached to the safelock system tube, for PCR, free of DNase, RNase and autoclavable pyrogens, with good user experience in the LCSP, through "Proof of use Satisfactory issued by the Quality Management Unit of the LCSP and send sample			5000pcs/Box	100box/day	120
00078	Thermo Fisher/ Shanghai		POLYPROPYLENE Cryovials THREAD LID	STERILE CRYOVIALS, 2 ml, with external screw cap, polypropylene, autoclavable and sterile, samples if possible			500pcs/Box	100box/day	20


# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00079	Thermo Fisher/ Shanghai		AUTOMATIC PIPETTE TIPS	Universal tips. Sterile, thin-walled, with filter for automatic pipettes from 200 to 1000 microliters. Free of DNase, RNase and pyrogens. Arranged in 96-tip racks, pipette compatible available at LCSP. Human, Eppendorf, Ranin			50pack/Box	100pack/day	400
00080	Thermo Fisher/ Shanghai		AUTOMATIC PIPETTE TIPS	Universal tips. Sterile, thin-walled, with filter for automatic pipettes from 10 to 200 microliters. Free of DNase, RNase and pyrogens. Arranged in 96-tip racks, pipette compatible available at LCSP. Human, Eppendorf, Ranin			50pack/Box	100pack/day	400
00081	Thermo Fisher/ Shanghai		AUTOMATIC PIPETTE TIPS	Sterile thin-walled automatic pipet tip, with filter for automatic pipettors from 2 to 20 microliters. Free of DNase, RNase and pyrogens. Arranged in 96-tip racks, pipette compatible Available from L.C.S.P. Eppendorf or Human brands			50pack/Box	100pack/day	400
00082	Eppendorf /Shanghai		VARIABLE VOLUME AUTOMATIC PIPETTE	Variable Volume Automatic Pipette, Multichannel, 8 Channels, adjustable from 0.5 to 10 microliters, with individual support.			1pcs	30	8
00083	Eppendorf /Shanghai		VARIABLE VOLUME AUTOMATIC PIPETTE	Automatic pipette with variable volume, GRADUABLE from 5 to 50 microliters. With quality certificate of origin. For biology molecular.			1pcs	30	8









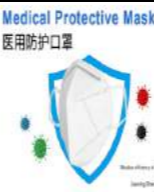

# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00084	Eppendorf /Shanghai		VARIABLE VOLUME AUTOMATIC PIPETTE	Automatic pipette with variable volume, GRADUABLE from 100 to 1000 microliters. With quality certificate of origin. For biology molecular.			1pcs	30	8
00085	Eppendorf /Shanghai		VARIABLE VOLUME AUTOMATIC PIPETTE	Automatic pipette with variable volume, GRADUABLE from 100 to 200 microliters. With quality certificate of origin. For biology molecular			1pcs	30	8
00086	Zhuhai		INFLUENZA DETERMINATION REAGENT	Rapid immunochromatographic test for the detection of Influenza Virus, and discrimination between type A and B, and subtyping of type AH1N1.			25test/kit	2000kit/day	6000
00087	changchun		EQUIPMENT FOR SAMPLING	Containing 2 ml. of liquid means to transport virus. For transport of viral samples, with two polyester swabs or other synthetic material (not cotton) and rigid plastic toothpick. The tube must have a conical bottom, sterile, without activated carbon			600set/Box	100box/day	10000
00088	ZHE JIANG		Quick-dry wash-free sanitizing Gel MSDS (100ML)	Medical				100 / BOX 38*38*22CM	20000

# Laboratory Supplies

Product Number	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Qty/Ton	Spec	Production ability / Delivery	Available live inventory? QUANTITY
00089	SHANG HAI		HYDROPHIL COTTON		certificate will be added soon		1 pack	10000pack/day	200

# Masks





No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00091	DONG GUAN		KN95 masks	KN95 Face Mask - FFP2 CE / FDA certified		GB2626-2006	1000个/Box 63.5*34*45.5cm 9KG	20 million / 37 days	2m per week
00092	GUANG ZHOU		Disposable surgical mask	Medical 3 ply Face Mask - CE / FDA		YY/T 0969-2013	50 / package 40 bales/carton 2000 pieces/carton 52*42*38 8.2KG	20 million / 30 days	3m per week
00093	HE NAN		surgical mask (non-sterilized)	Surgical 3 ply Face Mask - CE certified		YY/T 0969-2013	2000 / Box 52.5*49*56.5cm 11.5KG	1000000 / 5day	3m per week
00094	GUANG ZHOU		surgical mask (non-sterilized)	Surgical 3 ply Face Mask - CE certified		YY 0469-2011	50 / package 40 bales/carton 2000 pieces/carton 52*42*38 8.2KG	20 million / 30 days	3m per week
00095	SHEN YANG		N95 mask/respirator	N95 FFP2 CE / FDA certified		GB19083-2010	500 / box 46.5*43.3*36.3cm 4.5KG	2 million / 25 days	2m per week

## Gloves









No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00096	HE BEI		Disposable medical PVC gloves	PVC Vinyl Medical Gloves		ISO 13485 ISO 9001	100 / box 1000 / box 35*27*26cm 8KG	1 million / 3 days	1m 3 days
00097	HE BEI		Disposable nitrile gloves	Nytril Butyronitrile Medical Gloves		ISO 9001:2015	100 / box 1000 / box 35*27*25cm 5KG	1 million / 3 days	1m 3 days

## Eye Protection





\*M = million

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00098	TIAN JIN		FACE SHIELD	Made of transparent plastic/polycarbonate, it must provide good visibility, both at that carries it like the patient.		EN166	200 / box 68*38.3*41.5cm 13.87KG	100000 / day	600000 per week
00099	SHEN ZHEN		GOGGLES	Provides good sealing with face skin, PVC frame flexible to easily adapt to all face contours with even pressure, close your eyes and surrounding areas, should accommodate users with prescription glasses. Lenses must be Clear fog and scratch resistant plastic, adjustable band not to loosen during clinical activity, indirect ventilation to avoid fogging, reusable. EU STANDARD DIRECTIVE 86/686 / EEC, EN 166/2002: ANSI / ISEA Z87.1-2010		EN166:2001 EN ISO 4849:1981	200 / box 92.5*35.5*45.5cm 13.78KG	5-100000 / day	15000 per week

## Protective Clothing









No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00100	HE NAN		SHOES COVER	Operating rooms use, TNT, Large size with anti-slip to the traction, disposable		The national standard	50 /box 60*40*45cm	1000000 / 7 day	300000 per week
00101			SURGICAL HAT	Made of non-woven fabric. With fold-over rubber, fluid resistant. Disposable.		The national standard	20 / bag 1000 / box 60*40*40cm 7 kg		300000 per week
00102			COATS FOR OPERATING ROOM	Scrubs. Surgical gown Medium size. Fabric construction non-woven, disposable SMMS type with loop closure. Fluid resistant bodysuits, sleeves with reinforced elasticized cuffs from 5 to 10 cm width, minimum gown length 120 cm		The national standard	50 /box 60*40*45cm		100000 per week
00103	ZHE JIANG		PROTECTIVE SUIT	Sterile Hazmat suit Material: SMS / PP+PE Function: Anti-tear, Anti cross-contamination Without Cover Shoes Size: S M L XL XXL XXXL		GB19082 Moisture resistant grade 4 Blood penetrates level 3 EN14126	1 piece / bag 50 / box	30000/day	200000 per week

## Miscellaneous







No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00104	ZHE JIANG		Dry quickly without washing hands Sanitizing gel MSDS (500 ml)	500ml 75% Alcohol waterless gel		ISO22716:2007(E)	24 bottles/case 13KG 53*36*17.5cm	250,000 bottles /day	60000 per week Ethanol content 76%
00105	GUANG ZHOU		Forehead Temperature Gun	Infrared Thermometer Model No GL-B7F55 Tolerance: +/- 0.1 degree centigrade		YY/T0287-2017 ISO 13485:2016	50 boxes / boxes 64.5*33.5*39cm 12KG	4000 /week	4000 per week




## Ventilator Supplies

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note
00106	NING BO		Disposable use Invasive ventilator line	Invasive breathing circuit, FDA/CE approved				5000 per week	5000 per week
00107	NING BO		Disposable use Noninvasive ventilator line	None invasive breathing circuit, FDA/CE approved				5000/ week	5000 per week
00108	NING BO		Disposable use Invasive ventilator line	FDA/CE 1.6M EXPANDABLE TUBE / Y CONNECTORS WITH PORTS/ADDITIONAL 0.8M TUBE ( HMEF-optional +\$0.94COST)				5K / 2week 5K/ week	5k 2weeks then 5k per week
00109	JIANG XI		Disposable use Breathing filter	HEPA filter, HV ultrafiltration membrane, H87-W74 CE/FDA approved		ISO9001 ISO13485		20000 / week	20000 per week

# Air Purifiers

No.	Origin	Picture	Product	Description/ Specification	Certificate CE/FDA	Spec	Qty/Ton	Delivery	Note																																																																																																																		
00110	FU ZHOU		LK/KJH type Medical air purifying sterilizer	PM2.5、TVOC、 微生物PM2.5、 TVOC、 microorganism		ISO9001 ISO13485	1/Box 53*35*88.5cm 26KG	150 / week	In-stock, 150 per week																																																																																																																		
00111	FU ZHOU		LK/KJF - the -1150 plasma Air purifying and sterilizing machine	PM2.5、TVOC、 微生物PM2.5、 TVOC、 microorganism		ISO9001 ISO13485	1/Box 60.7*25.5*190cm	120 / week	In-stock, 120 per week																																																																																																																		
00112	CHENG DU		LK/KDSJ - Y1000 multi-function Air sterilizer	<table border="1" data-bbox="1438 852 1599 1015"> <tr> <th>项目</th> <th>规格/单位</th> <th>额定/标准</th> </tr> <tr> <td>额定风量 (m³/h)</td> <td>1000</td> <td>1000</td> </tr> <tr> <td>额定功率 (W)</td> <td>1000</td> <td>1000</td> </tr> <tr> <td>额定电压 (V)</td> <td>220</td> <td>220</td> </tr> <tr> <td>额定频率 (Hz)</td> <td>50</td> <td>50</td> </tr> <tr> <td>额定重量 (kg)</td> <td>10</td> <td>10</td> </tr> <tr> <td>额定尺寸 (mm)</td> <td>1000*1000*1000</td> <td>1000*1000*1000</td> </tr> <tr> <td>额定噪音 (dB)</td> <td>55</td> <td>55</td> </tr> <tr> <td>额定寿命 (h)</td> <td>10000</td> <td>10000</td> </tr> <tr> <td>额定温度 (°C)</td> <td>10-40</td> <td>10-40</td> </tr> <tr> <td>额定湿度 (%)</td> <td>10-90</td> <td>10-90</td> </tr> <tr> <td>额定海拔 (m)</td> <td>1000</td> <td>1000</td> </tr> <tr> <td>额定风速 (m/s)</td> <td>1.0</td> <td>1.0</td> </tr> <tr> <td>额定效率 (%)</td> <td>99.99</td> <td>99.99</td> </tr> <tr> <td>额定净化率 (%)</td> <td>99.99</td> <td>99.99</td> </tr> <tr> <td>额定净化速度 (m³/h)</td> <td>1000</td> <td>1000</td> </tr> <tr> <td>额定净化时间 (h)</td> <td>10000</td> <td>10000</td> </tr> <tr> <td>额定净化量 (m³)</td> <td>10000000</td> <td>10000000</td> </tr> <tr> <td>额定净化率 (ppm)</td> <td>0.1</td> <td>0.1</td> </tr> <tr> <td>额定净化率 (mg/m³)</td> <td>0.001</td> <td>0.001</td> </tr> <tr> <td>额定净化率 (μg/m³)</td> <td>0.0001</td> <td>0.0001</td> </tr> <tr> <td>额定净化率 (ng/m³)</td> <td>0.00001</td> <td>0.00001</td> </tr> <tr> <td>额定净化率 (pg/m³)</td> <td>0.000001</td> <td>0.000001</td> </tr> <tr> <td>额定净化率 (fg/m³)</td> <td>0.0000001</td> <td>0.0000001</td> </tr> <tr> <td>额定净化率 (ag/m³)</td> <td>0.00000001</td> <td>0.00000001</td> </tr> <tr> <td>额定净化率 (zg/m³)</td> <td>0.000000001</td> <td>0.000000001</td> </tr> <tr> <td>额定净化率 (kg/m³)</td> <td>0.0000000001</td> <td>0.0000000001</td> </tr> <tr> <td>额定净化率 (t/m³)</td> <td>0.00000000001</td> <td>0.00000000001</td> </tr> <tr> <td>额定净化率 (g/m³)</td> <td>0.000000000001</td> <td>0.000000000001</td> </tr> <tr> <td>额定净化率 (mg/m³)</td> <td>0.0000000000001</td> <td>0.0000000000001</td> </tr> <tr> <td>额定净化率 (μg/m³)</td> <td>0.00000000000001</td> <td>0.00000000000001</td> </tr> <tr> <td>额定净化率 (ng/m³)</td> <td>0.000000000000001</td> <td>0.000000000000001</td> </tr> <tr> <td>额定净化率 (pg/m³)</td> <td>0.0000000000000001</td> <td>0.0000000000000001</td> </tr> <tr> <td>额定净化率 (fg/m³)</td> <td>0.00000000000000001</td> <td>0.00000000000000001</td> </tr> <tr> <td>额定净化率 (ag/m³)</td> <td>0.000000000000000001</td> <td>0.000000000000000001</td> </tr> <tr> <td>额定净化率 (zg/m³)</td> <td>0.0000000000000000001</td> <td>0.0000000000000000001</td> </tr> <tr> <td>额定净化率 (kg/m³)</td> <td>0.00000000000000000001</td> <td>0.00000000000000000001</td> </tr> <tr> <td>额定净化率 (t/m³)</td> <td>0.000000000000000000001</td> <td>0.000000000000000000001</td> </tr> </table>	项目	规格/单位	额定/标准	额定风量 (m³/h)	1000	1000	额定功率 (W)	1000	1000	额定电压 (V)	220	220	额定频率 (Hz)	50	50	额定重量 (kg)	10	10	额定尺寸 (mm)	1000*1000*1000	1000*1000*1000	额定噪音 (dB)	55	55	额定寿命 (h)	10000	10000	额定温度 (°C)	10-40	10-40	额定湿度 (%)	10-90	10-90	额定海拔 (m)	1000	1000	额定风速 (m/s)	1.0	1.0	额定效率 (%)	99.99	99.99	额定净化率 (%)	99.99	99.99	额定净化速度 (m³/h)	1000	1000	额定净化时间 (h)	10000	10000	额定净化量 (m³)	10000000	10000000	额定净化率 (ppm)	0.1	0.1	额定净化率 (mg/m³)	0.001	0.001	额定净化率 (μg/m³)	0.0001	0.0001	额定净化率 (ng/m³)	0.00001	0.00001	额定净化率 (pg/m³)	0.000001	0.000001	额定净化率 (fg/m³)	0.0000001	0.0000001	额定净化率 (ag/m³)	0.00000001	0.00000001	额定净化率 (zg/m³)	0.000000001	0.000000001	额定净化率 (kg/m³)	0.0000000001	0.0000000001	额定净化率 (t/m³)	0.00000000001	0.00000000001	额定净化率 (g/m³)	0.000000000001	0.000000000001	额定净化率 (mg/m³)	0.0000000000001	0.0000000000001	额定净化率 (μg/m³)	0.00000000000001	0.00000000000001	额定净化率 (ng/m³)	0.000000000000001	0.000000000000001	额定净化率 (pg/m³)	0.0000000000000001	0.0000000000000001	额定净化率 (fg/m³)	0.00000000000000001	0.00000000000000001	额定净化率 (ag/m³)	0.000000000000000001	0.000000000000000001	额定净化率 (zg/m³)	0.0000000000000000001	0.0000000000000000001	额定净化率 (kg/m³)	0.00000000000000000001	0.00000000000000000001	额定净化率 (t/m³)	0.000000000000000000001	0.000000000000000000001		ISO9001 ISO13485	1/Box 61*48.6*94cm	200 / week	200 per week
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