Legendary Europe

Covid Supplies

0048729373945 00966568654916

Ulica,Aleje Jerozolimskie 89/43 02-001 Warszawa - Polska

www.LegendaryEurope.eu

Work together against COVID-19

Our Five Guidelines:

1) Best selected manufactures.
2) FDA approval.
3) Inspect the goods.
4) Ex-factory price plus minimal administrative expenses.
5) Insured Shipment.

Work together against COVID-19

- In an effort to support the anti-pandemic work, we integrate high-quality supply chain resources to provide medical supplies.
- If you need the products on the list, please contact us in time and we will deliver them as soon as possible.
- If you need products other than those on the list, please inform us in time and we will provide you with the best quality products in time.

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No.	Product	Supply/Week	Certificates
1	N95 #1	1 million pcs	CE, FDA
2	N95 #2		CE, FDA
3	KN95 #1	1.4 million pcs	CE, FDA
4	KN95 #2	1.4 million pcs	CE, FDA
5	KN95 #3	1.4 million pcs	FDA,CE (expected by next Monday)
6	Disposable medical mask #1	1 million pcs	CE, ISO13485, FDA
7	Disposable Medical Mask #2	1 million pcs	CE, FDA
8	Disposable medical masks #3	7 million pcs	CE, FDA
9	Reusable Copper Infused Protective Masks	1.4 million pcs	FDA
10	Superbio IgM/IGG Antibody Fast Detection Kit	1.4 million pcs	CE, FDA
11	NewScen IgM/IGG Antibody Fast Detection Kit	5 million pcs	CE, FDA
12	BGI IgM/IGG Antibody Fast Detection Kit		
13	Medical Disposable Protective Gown-Sterile Type	/	FDA
14	Medical Disposable Protective Gown-Non Sterile Type	/	FDA
15	Medical goggles	0.5 million pcs	CE,FDA
16	Invasive ventilator	1500 units	CE
17	Non-Invasive ventilator yuwell Bi-level device	/	CE
18	Infrared Forehead Thermometer #1	/	FDA
19	Infrared Forehead Thermometer #2		CE, FDA
20	Thermographic Measurement (Hikvision)	/	/
21	MediDefense Penetrexx Surface Protection	/	FDA
22	mPulse Hand Sanitizer	/	FDA

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N95 #1

Certificate	CE, FDA
Supply/ Week	1 million pcs
Deliver Time	Arrive within 7 days

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N95 #2

Certificate of Conformity

NO.: 10062813

This is certifies that:

Applicant	:	ZHUHAI QINGCHUNYU BIOTECHNOLOGY CO., LTD
Address	:	4th floor, building 2, No.18, Honghui fifth road, Hongqi Town, Jinwan District Zhuhai, Guangdong, 519090, CHINA
Manufacturer	:	ZHUHAI QINGCHUNYU BIOTECHNOLOGY CO., LTD
Address	:	4th floor, building 2, No.18, Honghui fifth road, Hongqi Town, Jinwan District Zhuhai, Guangdong, 519090, CHINA
Device Listing	:	D374136 (Respirator, surgical)
EUT	:	Particulate respirator
Models	:	N95, FFP3(the product photo see the annex)
Test Standard	:	Food and Drug Administration Regulation

FDA

Certificate	CE, FDA
Supply/ Week	/
Deliver Time	/

KN95 #1

CERTIFICATION OF FDA REGISTRATION

FDA

CERTIFICATION OF FDA REGISTRATION

Certificate	CE, FDA
Supply/ Week	1 million pcs
Deliver Time	Arrive within 7 days

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KN95 #2

Certificate	CE
Supply/ Week	7 million pcs
Deliver Time	/

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KN95 #3

Certificate	FDA,CE (expected by next Monday
Supply/ Week	1.4 million pcs
Deliver Time	/

Fiscal Year 2020 USA FDA
CERTIFICATION OF REGISTRATION

This certifies that:

TimeBase Medical Science and Technology(Jiangsu) Co.,Ltd.
37 Building, No. 801 Jiankang Avenue, Taizhou Economic Development Zone, Taizhou City, Jiangsu Province

has completed the FDA Establishment Registration (as manufacturer) and Device Listing with the US Food & Drug Administration, through

Salude USA, LLC.

Owner/Operator Number: 10063090

Device Listings:

Listing Number	Product Code	Device Name ➢ Proprietary Name	Premarket Submission Type
D075897	OEA	**Disposable General Medical Gown(Non-Surgical)**	510(k) Exempt

Salude

Salude LLC will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. Salude LLC makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whom such benefit is intended. This certificate does not denote endorsement or approval of the certificate holder's device or establishment by the U.S. Food and Drug Administration. Salude LLC assumes no liability to any person or entity in connection with the foregoing.

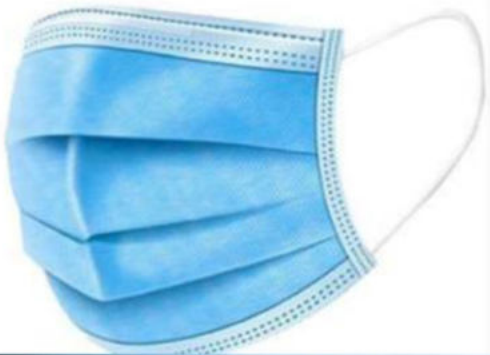
Pursuant to 21 CFR 807.20, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Salude, LLC is not affiliated with the U.S. Food and Drug Administration.

Salude USA, LLC

FDA

Executive VP
Issued: Mar. 19, 2020
Cert. No. 20200319003
Expiration Date: Dec. 31, 2020

Disposable Medical Mask #1



Certificate	CE, ISO13485, FDA
Supply/ Week	1 million pcs
Deliver Time	/

Applicant Name	Bengbu Huohe Pharmaceutical CO.,LTD
Address	NO.189 Xinghua Road Bengbu City
Manufacturer	Bengbu Huohe Pharmaceutical CO.,LTD
Address	NO.189 Xinghua Road Bengbu City
Product name	Disposable Mask
Model / Serial No.	HH/QB2020-01, HH/QB2020-02, HH/QB2020-03, HH/QB2020-04 17.5cm×9.5cm, 14.5cm×9.5cm, 12.0cm×7.0cm
Standard EN Norms	EN 149:2001+A1:2009
Applicable Directives	R 2016/425 (Regulation on Personal Protective Equipment)

NOTE: 烦请确认以上企业及产品信息，如无误，请签字盖章后扫描或者传真，谢谢
以上信息为欧盟 CE 认证证书正本所体现信息，正本签发后不能修改，请熟知！

企业签字(盖章)：


中华人民共和国
药品GMP证书
CERTIFICATE OF GOOD MANUFACTURING PRACTICES FOR PHARMACEUTICAL PRODUCTS
PEOPLE'S REPUBLIC OF CHINA
Manufacturer:
Address: No. 189 Xinghua Road Bengbu City
Scope of Inspection:
This is to certify that the above-mentioned manufacturer complies with the requirements of Chinese Good Manufacturing Practices for Pharmaceutical Products.
This certificate remains valid until
Issued By
Date for Issuing
国家药品监督管理局制
NATIONAL MEDICAL PRODUCTS ADMINISTRATION


药品生产许可证

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Disposable Medical Mask #2

Certificate	FDA, CE
Supply/ Week	1 million pcs
Deliver Time	/

Disposable Medical Mask #3

2020

CERTIFICATE OF REGISTRATION

This certifies that:

ZHANGJIAGANG ZHIYI MEDICAL HEALTH PRODUCTS CO., LTD
Renmin Road Leyu Town
Zhangjiagang City Jiangsu, CHINA 215622

is registered with the U.S. Food and Drug Administration for FY 2020 pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations:

Establishment Registration:	3008192718
DUNS No.:	56-297-8777
Device Classification Name:	MASK, SURGICAL
Product Code:	FXX
Regulation Number:	878.4040
Official Correspondent and U.S. Agent:	Registrar Corp 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0377 • Fax: +1-757-224-0179

Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until the end of the year stated above, unless said registration is terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. Registrar Corp assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.38, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding."

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0377 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lyndon
Executive Director
Registrar Corp
Dated: November 28, 2019

检测结果

RESULTS OF TEST

说 明

Certificate	FDA, CE
Supply/ Week	7 million pcs
Deliver Time	/

Reusable Cooper Infused Protective Masks

COPPER INFUSED
MACHINE WASHABLE
INHIBIT BACTERIA AND FUNGAL GROWTH
IMPROVE HYGIENE
REDUCE FACIAL TOUCHING

Certificate	FDA
Supply/ Week	1.4 million pcs
Deliver Time	/

MANUFACTURING SOLUTIONS CENTER

301 Conover Station SE; Conover, NC 28613

Irad Seese
Textile Based Delivery
301 Conover Station SE
Conover, NC 28613

Received Date: March 17, 2020
Completed Date: March 20, 2020

REFERENCE: TS 157967
Style: 850015085046
Color: White
Description: Face Mask
Copper Nylon

Sample Type: Knitted Fabric
Sample Form: Face Mask
Size: N/A

TEST RESULTS:

Antibacterial Finishes:

Assessment of Antibacterial Finishes on Textile Materials - AATCC 100-2012

This test is accredited under the laboratory's ISO/IEC 17025 accreditation issued by the ANSI-ASQ National Accreditation Board. Refer to certificate and scope of accreditation L2238

Testing Results:

Tested after 1x wash/dry cycle	Results: cfu/sample		
	Zero Contact Time	24hr Contact Time	Percent Reduction
Staphylococcus aureus ATCC 6538	1.30E+05	1.00E+03	99.23%
Klebsiella pneumoniae ATCC 4352	5.50E+05	1.00E+03	99.82%

Calculate % reduction to formula 1) 100 (B-A)/B = R; section 10.2

Testing Information:

- Staphylococcus aureus ATCC 6538
- Klebsiella pneumoniae ATCC 4352
- Growth media: Tryptic Soy Broth
- Sample size # layers: 2
- Sterilization: none
- Neutralizer: 100ml Letheen Broth w. Tween
- Target inoc. Level: (1.0-2.0) x 10^5 CFU/ml
- Inoculum carrier: 5% (1/20x) Nutrient broth w. 0.05% Triton X100
- Inoculum size: 1.0ml +/- 0.1ml
- Contact time: 18 - 24 h
- Temperature: 37 +/- 2° C
- Samples were tested after 1x wash/dry cycle
- Samples were prepared and enumerated using automatic equipment; Tempo, BioMerieux.
- Cultures stored at 5° +/-2°C.

MANUFACTURING SOLUTIONS CENTER

301 Conover Station SE; Conover, NC 28613

Testing Information:

- AATCC Laboratory Procedure 1 -2018, "Home Laundering: Machine Washing"
  - Home Laundered 1 times using AATCC Laboratory Procedure 1-2018
  - Table II A, B & C - Alternate Laundering Parameters
  - Laundry Test Conditions: load size = 1.8kg (4lbs.). Launder Right Side Out
  - Top Loading Machine Wash: Normal Cycle (IV) Hot: 49+/-3C ((120+/-5F)
  - using 50±1ml AATCC High Efficiency (HE)Standard Reference Liquid Detergent
  - (Aii) Tumble Dry Permanent Press Table V - Standard Drying Conditions
  - Type 3: 50% cotton/ 50% polyester bleached plain weave Table VII - Ballast Parameters

Report prepared:

Cheryl Nettesheim
Antimicrobial Testing
MSC – Testing Lab
(828) 327-7000 ext. 4512

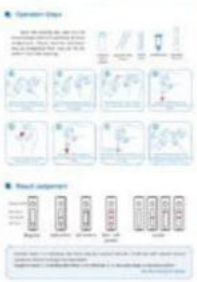
cnettesheim@manufacturingsolutionscenter.org
www.manufacturingsolutionscenter.org

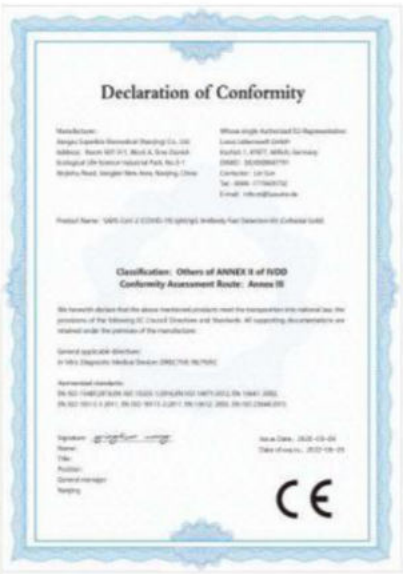
The test result are based on the submitted sample(s) only. MSC/HTC's liability shall not exceed the fees paid for the testing reflected on this report. It is the customer's responsibility to ensure that they comply with all U.S. federal, state/ local laws & regulations. These tests are not to be used for marketing purposes without the express written consent of the Director of the Manufacturing Solutions Center at Catawba Valley Community College. The test report shall not be reproduced except in full, without written approval from MSC/HTC. All results will be kept confidential.

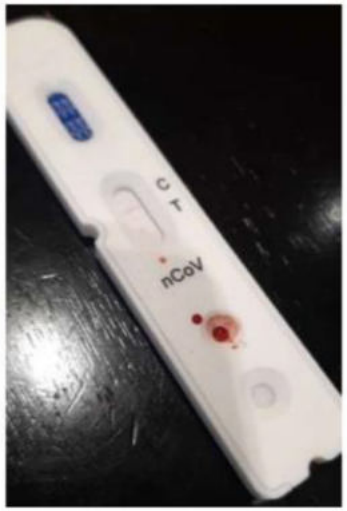
MADE IN USA

FDA CERTIFIED FACILITY

Superbio IgM/IGG Antibody Fast Detection Kit







Certificate	FDA, CE
Supply/ Week	1.4 million pcs
Deliver Time	Arrives within 7 days

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NewScen IgM/IGG Antibody Fast Detection Kit

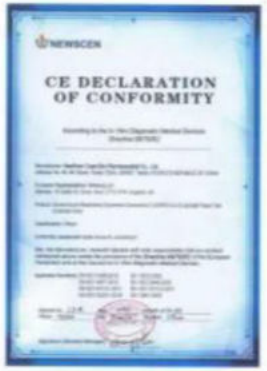
NewScen SARS-COV-2 Antibody IgM/IgG
Rapid Test Kit (Colloidal Gold)

CERTIFICATION

国际认证

ISO13485

ISO9001



EC DOC

NewScen	Diagnosed Sample				Total	
	Positive		Negative			
Positive	A	86	B	9	A+B	95
Negative	C	8	D	258	C+D	266
Total	A+C	94	B+D	267	ABCD	361

Sensitivity = A/(A+C)% = 91.49%
Specificity = D/(B+D)% = 96.63%
Total Accuracy = (A+D)/(ABCD)% = 95.29%

The evaluation is carried out on the sample from COVID-19 nucleic acid test (PCR) confirmed cases from CDC and classIII hospitals

Finger Tip Blood ✔

15 minute Result GET ✔

2-30℃ Storage ✔

24M Shelf-life ✔

FAST ! EASY ! ACCURATE!

Certificate	FDA, CE
Supply/ Week	5 million pcs
Deliver Time	Arrives within 7 days

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BGI IgM/IGG Antibody Fast Detection Kit

Test principle

The SARS-CoV-2 IgM/IgG rapid assay kit (Colloidal Gold) is to make a colloidal gold mark taking the advantage of the specific antigen of novel coronavirus, to coat taking the anti-human IgM and anti-human IgG monoclonal antibody as the detection line.

Positive result of IgM antibody indicating a recent infection

Positive result of IgG antibody signaling a longer or previous infection

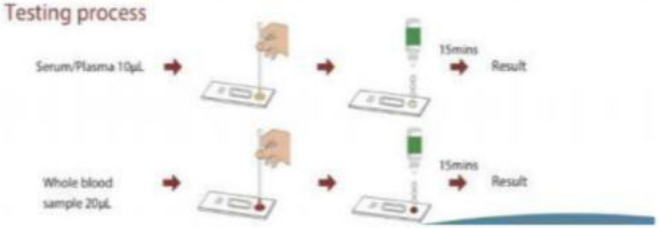
Sample Type

Serum / plasma / venous whole blood

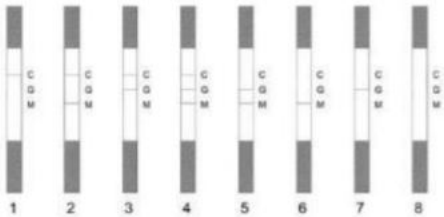
Advantage

SIMPLE	A simple and convenient one-step operation
FAST	Rapid test generates results in just 15 minutes
INTUITIVE	The result is readable with naked eyes, specific equipment is not required

Testing process



Results



"C" (Control line), "M"(IgM test line) , "G"(IgG test line).

	Test results	Results Interpretation
1	Only the quality control (C) line appears	Negative
2	Both the quality control line (C) and the detection line M appears	Positive for the IgM antibody
3	Both the quality control line (C) and the detection line G appears	Positive for the IgG antibody
4	The quality control line (C) and both detection lines G and M appears	Positive for both the IgG and IgM antibodies
5	The quality control line (C) doesn't appear, no matter whether the detection line G and M appear	The test is invalid. New test should be made

50 test/kit, key composition of each kit:

Composition	Specification	Quantity
SARS-CoV-2 Detection reagent card	1pcs/bag	50 bags
Sample diluent solution	5mL/bottle	2 bottles

Certificate	FDA, CE
Supply/ Week	5 million pcs
Deliver Time	Arrives within 7 days

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Medical Disposable Protective Gown – Sterile Type

Certificate	FDA, CE
Supply/ Week	1.4 million pcs
Deliver Time	/

Fiscal Year 2020

CERTIFICATION OF FDA REGISTRATION

Listing No	Code	Device Name	Proprietary Name
D374151	FRL	FIBER, MEDICAL, ABSORBENT	cotton soft towels, compressed face towels, wet wipes, non-woven fabrics, disposable wipes
D374154	HCY	Shield, eye, ophthalmic (including sunlamp protective eyewear and post-myfolatic eyewear)	goggles
D374153	OKA	Non-surgical isolation gown	protective clothing
D374150	KHA	MASK, SCAVENGING	Masks, mask gaskets

Fiscal Year 2020

CERTIFICATION OF FDA REGISTRATION

This certifies that:

NINGBO SHISHA HOUSEWARE CO., LTD.
1F, No.2 Yongfeng Street, Lingfeng Village, Xiaowangmiao Street, Fenghua District, Ningbo, Zhejiang, 315800, CHINA

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, through

Shenzhen CCT Testing Technology Co., Ltd.

Owner/Operator Number: 10062816

Device Listing#: See Next page

Medical Disposable Protective Gown – Non-Sterile Type

Fiscal Year 2020

CERTIFICATION OF FDA REGISTRATION

Listing No	Code	Device Name	Proprietary Name
D374151	FRL	FIBER, MEDICAL, ABSORBENT	cotton soft towels, compressed face towels, wet wipes, non-woven fabrics, disposable wipes,
D374154	HOY	Shield, eye, ophthalmic (including sunlamp protective eyewear and post-mydriatic eyewear)	goggles,
D374151	OKA	Non-surgical isolation gown	protective clothing
D374150	KHA	MASK, SCAVENGING	Masks, mask gaskets

Fiscal Year 2020

CERTIFICATION OF FDA REGISTRATION

This certifies that:

NINGBO SHISHA HOUSEWARE CO., LTD.
1F, No.2 Yongfeng Street, Lingfeng Village, Xiaowangmiao Street, Fenghua District, Ningbo, Zhejiang, 315800, CHINA

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, through

Shenzhen CCT Testing Technology Co., Ltd.

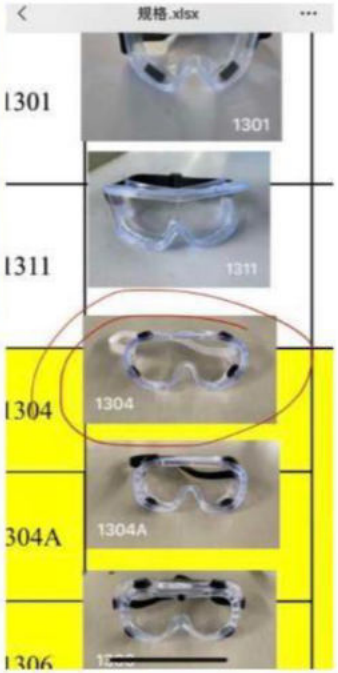
Owner/Operator Number: 10062816

Device Listing#: See Next page

Certificate	FDA, CE
Supply/ Week	1.4 million pcs
Deliver Time	/

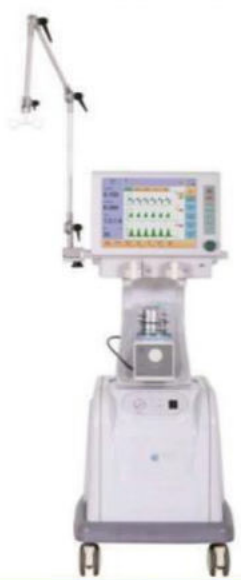
Medical Goggles

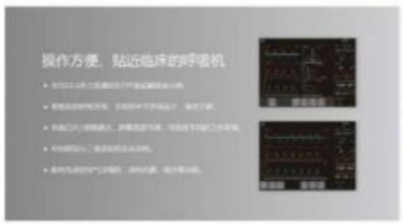
Certificate	FDA, CE
Supply/ Week	0.5 million pcs
Deliver Time	Arrives within 7 days



Invasive Ventilator

Certificate	CE
Supply/ Week	1500 units
Deliver Time	Arrives within 14 days





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Non-Invasive Ventilator Yuwell Bi-level device

Certificate	CE
Supply/ Week	/
Deliver Time	/

YH730
Bi-Level
CE & CFDA APPROVED

1.Maximum inhale pressure 30cmH2O
2.With ST and target tidal volume mode settings
3.Fast rise time
4. Normal inhalation intensity, respiration rate is 54 times / min, and each exhalation triggers the ventilator perfectly. Easily keep in sync
5. Good synchronization with big flow

YH830
Bi-Level
CE & CFDA APPROVED

YH-830 mobile Bi-level for hospital

1.Maximum inhale pressure 30cmH2O, with separate humidifier, better humidification
2.With ST and target tidal volume mode settings
3.Fast rise time
4. Normal inhalation intensity, respiration rate is 54 times / min, and each exhalation triggers the ventilator perfectly. Easily keep in sync
5. Good synchronization with big flow

EC Certificate

No. G2 052486 0006 Rev. 01

Manufacturer: Suzhou Yuyue Medical Technology Co., Ltd.

Facility(ies):

Product Category(ies): I.V. Catheter for Single Use, Positive Airway Pressure Units, Humidifier

Certificate

Holder of Certificate: Beijing Eternity Electronic Technology Co.,Ltd.

Facility(ies):

Certification Mark:

Scope of Certificate: Design and Development, Production and Distribution of Anaesthetic machines, Ventilators, Emergency and Transport Ventilators, Infusion Pumps, Medical Air Compressors, BiPAP System (Nasal Continuous Positive Airway Pressure System), Syringe Pump

Applied Standard(s):

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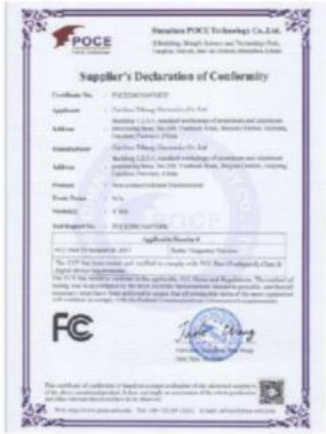
Infrared Forehead Thermometer #1

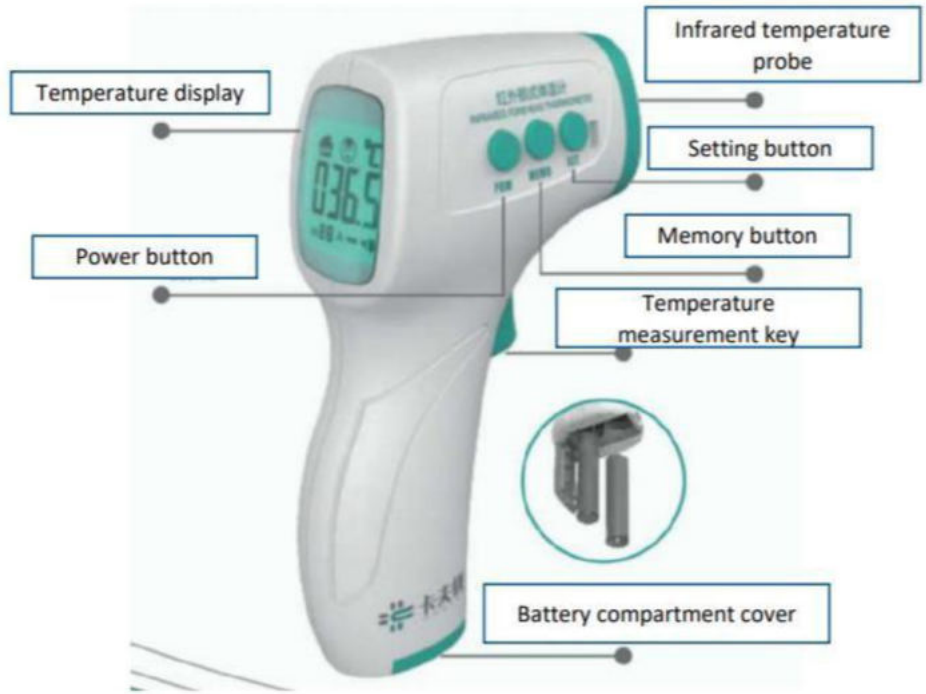


roduct structure	①	Infrared sensor	
	②	LCD	
	③	Backlight on / off button	
	④	Up button (view historical data)	
	⑤	Down button (view historical data)	
	⑥	Sound switch button	
	⑦	Measuring switch	
	⑧	Battery cover	
	⑨	Handle	
LCD Display Overview	①	Body temperature mode (BODY)	
	②	Digital reading	
	③	Storage location	
	④	Temperature ℃ / ℉	
	⑤	Read out of stored data	

	Technical Index	
lasic Parameters	Exact digits showing	0.1℃(0.1℉)
	Storage temperature	-20-55 ℃
	Ambient temperature for operation	5 ℃ ~ 40 ℃, the best temperature is 25 ℃
	Relative humidity	≤85%
	Power supply	DC 3V (2 AA batteries in series)
	Specification	160 * 100 * 40mm
	weight	100g
Measuring Range	Human body model temperature range	32.0-42.9 ℃
	Measuring distance range	5-15 cm (best measurement is 5 cm)
	Automatic shutdown time	20s
Measurement Accuracy	32~35.9℃（93.2~96.6℉）	±0.3℃(0.5℉)
	36~39℃（96.8~102.2℉）	±0.2℃(0.4℉)
	39~43℃（102.2~109.4℉）	±0.3℃(0.5℉)
Service life	40,000 times	/

Certificate	FDA, CE
Supply/ Week	/
Deliver Time	/

Infrared Forehead Thermometer #2



Certificate	FDA, CE
Supply/ Week	/
Deliver Time	Deliver within 24 hour if order less than 1000; deliver in 3-5 days if order amount between 10K-50K

Thermographic Measurement

DS-2TD2617B-3/6PA(B)

- Thermal : 160 × 120 ;
- Lens: 6mm ;
- Optical : 2688 × 1520 ;
- Optical lens: 4mm / 8mm ;
- Video mode : Bi-spectrum image fusion ;
- Accuracy : ±0.5℃ (± 0.3℃ with blackbody)
- Range : 30-45℃
- Audio alarm support

DS-2TD1217B-3/6PA(B)

- Thermal : 160 × 120 ;
- Lens: 6mm ;
- Optical : 2688 × 1520 ;
- Optical lens: 4mm / 8mm ;
- Video mode : Bi-spectrum image fusion ;
- Accuracy : ±0.5℃ (± 0.3℃ with blackbody)
- Range : 30-45℃
- Audio alarm support

Black Body

- Temperature resolution: 0.1℃
- Accuracy: ±0.1℃
- Temperature stability: ±0.1℃/h
- Effective emissivity: 0.97±0.02
- Operating temperature: 0~30℃

DS-2TD2636B-15/P

- Thermal : 384 × 288, lens: 15mm ;
- Optical : 2688 × 1520, lens: 6mm ;
- Accuracy : ±0.5℃ (± 0.3℃ with blackbody)
- Measurement Range : 30-45℃
- Working temperature : 10~35℃
- AI face detection, reduce false alarms.
- Simultaneous fever screening for multiple person (ca.10-15 persons).

NP-SG318LT-F

- 18detect zones
- Thermal imaging resolution : 160*120
- Temperature measurement accuracy : ±0.5℃
- Temperature measurement range : 30-45℃
- 7 inch LCD touch screen

- Face recognition & body temperature measurement
- Non-contact fast body temperature measurement (face detection)
- All in one Fast to deploy
- Voice reminder to broadcast real-time alarm
- Face capacity: 20,000
- Passing Speed: Up to 60 Persons/Min
- Measurement range: 30~45℃
- Accuracy: ±0.5℃.

DS-K5671-ZU+DS-2TD2617B-3/6PA(B)
DS-K3B601-L/Mpg-Dp65 or DS-K3B601-M/Mpg-InTtNDp65 or DS-K3B601-R/MpgTtN-Dp65

DS-2TP21B-6AVFW

- Thermal resolution : 160 × 120;
- Optical resolution : 640 × 480;
- Range : 30-45℃
- Touchable screen
- Accuracy : ±0.5℃ (± 0.3℃ with blackbody)
- Bi-spectrum fusion supported
- WIFI support
- Audio alarm support
- Automatic screenshot & upload

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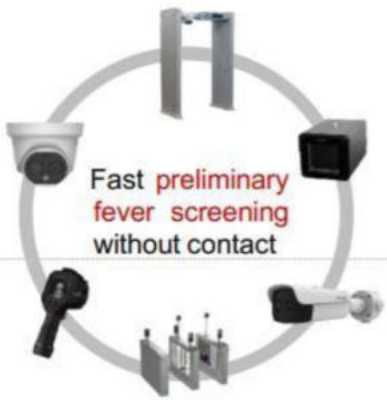
Thermographic Measurement



Locate potential fever person

Manual measurement by mercury, ear thermometer etc.

Long-term fever screening at entrance with Walk through metal detector

Long-term fever screening at entrance with access control

Short-term fever screening at entrance with Thermal handheld camera

Short-term fever screening at entrance with Thermal bullet/turret camera

Long-term fever screening for crowd in open area

Patrol fever screening

SAFE
Non-contact, long distance and accurate, reduce cross-infection possibility

EFFICENT
Efficient, auto alarm triggering, reduce manpower resources investment.

ADAPTABLE
Multiple product types, easy to deploy, suitable for different scenarios.

TRACEABLE
Combining temperature, image and face recognition for easy management and query

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MediDefense Penetrexx Surface Protection

Calvert Memorial Hospital tested 60-day efficacy of Medi Defense Penetreuxx Surface Protection and the reduction of ATP scored achieved within a hospital environment commenced on Jun21, 2017 and continued for approximately 60 days.

Test Protocol

A Study in Efficacy of Medi Defense Penetrexx Surface Protection

Clearstream Technologies LLC performed a study by treating 5 high touch, heavy wear surfaces at Calvert/Frederick Memorial Hospitals, comparing the ATP counts of the surfaces treated with MediDefense Penetrexx against the ATP counts on the similar untreated surfaces. The data was collected using a standard ATP meter between regularly scheduled cleanings, two times weekly in the morning prior to cleaning and at the end of the day after daily use for a period of 60 days.

Surfaces treated with MediDefense Penetrexx yield a greater percent reduction in bio load than control surfaces when monitored in a side by side comparison without the interaction of established cleaning protocols in the reduction of bio loads on the monitored surfaces.

Certificate	FDA
Supply/ Week	/
Deliver Time	/

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Methicillin Resistant Staphylococcus aureus (MRSA) Strain

These data indicate that the fabric test, treated with the CT Siquat Technology, reduces the total population of MRSA bacteria >99.99% after one hour simulated dynamic contact.

CERTIFICATE OF ANALYSIS

These antibacterial tests were performed to demonstrate the effectiveness of the CT SiQuat Technology against the specific Methicillin Resistant *Staphylococcus aureus* (MRSA) strain. Test Methods conform to **ASTM E2149-01** guidelines (Standard Test Method for Determining the Antimicrobial Activity of Immobilized Antimicrobial Agents Under Dynamic Contact Conditions). Specific details of testing and materials is listed below the table.

These data indicate that the fabric tested, treated with the CT Siquat Technology, reduces the total population of MRSA bacteria >99.99 % after one hour simulated dynamic contact. Untreated fabric samples tested in parallel demonstrated no effectiveness at reducing the total MRSA population. These results indicate the antimicrobial effectiveness of the sample treated with the CT Siquat Technology against the resistant bacteria MRSA..

	Microbiological Analysis		
	Initial concentration	Final concentration	Percent Reduction
Untreated Fabric Sample	1.8 x 10s/ml	1.9x10s/ml	0%
Treated Fabric Sample	1.8x10s/ml	< 1.0 x 10 if ml	> 99.99%

ASTM E2149-01 StandaldTest Method for Detennining the Antimicrobial Activity of Immobilized Antimicrobial Agents Under Dynamic Contact Conditions. Total contact time: 1 hour Total Volume: 50 ml 0.3 mM KH.PO. +0.01% Q2-5211 Bacterial Strain: Clinical Isolate Methicilin Resistant *F. staphylococcus, aureus,* (MR SA) Description of sample tested: lg each **fabric** treated and untreated with AEM5772/5

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mPluse Hand Sanitizer

- The problem with base ingredients of hand sanitizer in US is that although they both provide an instantaneous kill; neither have any long - term efficacy.
- Adding preservatives like SiQ to these base formulas has the potential to add residual protection to the formula and provide for a sanitizer that would have additional protection properties after initial application.
- The added benefit of the SiQ formula to BAC products is the increased persistence on the skin, which will lead to a further reductions of microbial growth or persistence on the skin.

Certificate	FDA
Supply/ Week	/
Deliver Time	/

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THANK YOU

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