

VENTILATOR
Y-30T - CHINA



VENTILATOR
725 BI level - CHINA



VENTILATOR 830 BI level - CHINA



COVID 19 TEST KIT - INDIAN



VENTILATOR
Savel 500 - INDIA



VENTILATOR
Savel 500e - INDIA



COVID 19 TEST KIT - CHINA



K N 95 Mask



N 95 Mask



K N 95 Mask - 2



3 PLY Mask - INDIA



Sanitizer (Gel+ Liquid)



DISPOSABLE PROTECTIVE
CLOTHING FOR MEDICAL
USE - INDIAN



DISPOSABLE PROTECTIVE
CLOTHING FOR MEDICAL
USE - CHINA



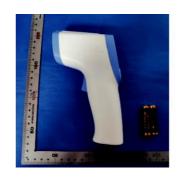
PPE KIT - 1

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PPE KIT - 2







MEDICAL INFRARED
THERMOMETER - CHINA



SECOND GENERATION MASK - CHINA



ISOLATER CHAMBER
- CHINA



SOLE CLEANING
MACHINE - CHINA



FOWLER BEDS
- CHINA



DISPOSABLE
MEDICAL MASK
- CHINA



SMART HELMET
- CHINA



DISINFECTION BOOTH - CHINA



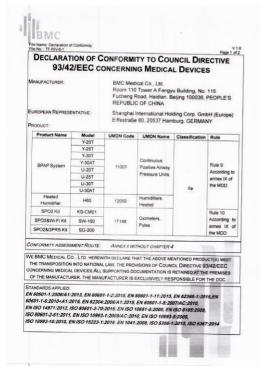
RAPID THERMOMETRY SAFETY DOOR - CHINA

VENTILATOR - Y-30T - CHINA













VENTILATOR - TECHNICAL PARAMETERS OF 725 BI-LEVEL PAP DEVICE - SHIPMENT FROM CHINA



1. Scope of application:

For adult patients with ARDS, hypoventilation syndrome, for bi-level non-invasive ventilation in a hospital or home

2. Themain function:

- 1. Different treatment options can be provided according to patients' needs, from CPAP to S, T, S / T, VGPS five modes
- Adjustable trigger sensitivity.
 Real-time monitoring and recording of blood oxygen saturation.
- 4. Alarmfunction.
- 5. Record function (SD card).
- 6. Maximum leak compensation function. 60L/min.
- 7. With trolley, easy to move in the hospital.

3. Thekey technical parameters:

1. Working mode: CPAP, S, T, ST, VGPS.

2. Ventilation parameters:

- 2.1 Inspiratory pressure: 4-25 cmH2O (0.5 cmH2O interval)
- 2.2 Expiratory pressure: 4-25 cmH2O (0.5 cmH2O interval)
- 2.3 Treatment pressure: 4-20 cmH2O (in CPAP mode, 0.5 cmH2O interval)
- 2.4 Delayed boost: 0-45 minutes (interval of 5 minutes)
- 2.5 Breathing frequency: 550 times / minute (interval 1 time / minute)
 2.6 Breathing ratio: 10 70% (interval of 1%)
- 2.7 Rise time: 1-5 steps (1 step interval, 1 step rises fastest, 5 steps rises most slowly)
- 2.8 Inhalation sensitivity: 1-5 steps (1 step interval, 1st step triggers fastest, 5th step 1st step triggers slowest)
- 2.9 Exhalation sensitivity: 1-5 (1 interval, the 1st trigges the fastest, 5th and 1st the
- 2.10 Tidal volume: 50-1500ml (effective in VGPS mode, interval 50ml)

3. VGPS (same with AVAPS, canset Target tidal volume) mode

- 3.1 Maximum IPAP value adjustable range: 4-25 cmH2O
- 3.2 Adjustable range of minimum IPAP value: 4-25 cmH2O
- 3.3 EPAP value adjustable range: 4-25 cmH2O
- 3.4 Target Tidal Volume: 50-1500ml

4. Display range of monitoring setting parameters: 4.1 Respiratory waveform can be displayed;

- 4.2 Current mode: CPAP, S, T, ST, VGPS
- 4.3 Inspiratory pressure: 4-25 cmH2O
- 4.4 Expiratory pressure: 4-25 cmH2O
- 4.5 Breathing frequency: 5-50 times / minute
- 4.6 Breathing ratio: 10-70%
- 4.7 Air leakag e: 20-99.9lpm 4.8 Tidal volume: 50-1500ml
- 4.9 minute ventilation: 0-50ipm
- 4.10 Blood oxygen saturation: 70-100%
- 4.11 Pulse rate: 25-250bpm
- 4.12 Delay boost: 0-45 minutes

5. Alarm items:

- 5.1 High pressure sound prompt: Turn on / off sound prompt
- 5.2 Lowminute ventilation sound prompt: Turn on / off sound prompt, can be set to
- 1-10L (interval 1L)
- 5.3 Air leakage sound prompt: on / off
- 5.4 Power off sound prompt: on / off
- 5.5 Choking voice prompt: On / off, choking time can be set to 10-40 seconds (intervaof 5 seconds)
- 5.6 Low Tidal Volume Sound Prompt: On / Off, Set Ti dal Volume Value Range 50-500ml ((Interval 50ml))

- 6.1 Total Weight: about 20kg including the trolley and the device
- 6.2 Operating temperature: 5-35 °C
- 6.3 Relative humidity: 10%-90%
- 6.4 Power supply: input is AC 100 -240V, 50-60Hz, 1.8A Max; output: DC 24V, 3.33A.



VENTILATOR - TECHNICAL PARAMETERS OF 830 BI-LEVEL PAP DEVICE - SHIPMENT FROM CHINA



1. Scope of application:

For adult patients with ARDS, hypoventilation syndrome, for bi-level non-invasive ventilation in a hospital or home.

2. The main function:

- 1. Different treatment options can be provided according to patients' needs, from CPAP to S, T, S / T, VGPS five modes
- 2. Adjustable trigger sensitivity.
- Real-time monitoring and recording of blood oxygen saturation.
- 4. alarm function.
- 5. Record function (SD card).
- 6. Maximum leak compensation function. 60L/min
- 7. With trolley, easy to move in the hospital

3. the key technical parameters:

- 1. Working mode: CPAP, S, T, ST, VGPS.
- 2. Ventilation parameters:
- 2.1 Inspiratory pressure: 4-30 cmH2O (0.5 cmH2O interval)
- 2.2 Expiratory pressure: 4-30 cmH2O (0.5 cmH2O interval)
- 2.3 Treatment pressure: 4-20 cmH2O (in CPAP mode, 0.5 cmH2O interval)
- 2.4 Delayed boost: 0-45 minutes (interval of 5 minutes)
- 2.5 Breathing frequency: 5-50 times / minute (interval 1 time / minute)
- 2.6 Breathing ratio: 10-70% (interval of 1%)
- 2.7 Rise time: 1-5 steps (1 step interval, 1 step rises fastest, 5 steps rises most slowly)
 2.8 Inhalation sensitivity: 1-5 steps (1 step interval, 1st step triggers fastest, 5th step
- 1st step triggers slowest)
- 2.9 Exhalation sensitivity: 1-5 (1 interval, the 1st trigger is the fastest, 5th and 1st the
- 2.10Tidal volume: 50-1500ml (effective in VGPS mode, interval 50ml)

3.VGPS (same with AVAPS, can set Target tidal volume)mode:

- 3.1 Maximum IPAP value adjustable range: 4-30 cmH2O
- 3.2 Adjustable range of minimum IPAP value: 4-30 cmH2O
- 3.3 EPAP value adjustable range: 4-30 cmH2O
- 3.4 Target Tidal Volume: 50-1500ml

4. Display range of monitoring setting parameters:

- 4.1 Respiratory waveform can be displayed; 4.2 Current mode: CPAP, S, T, ST, VGPS
- 4.3 Inspiratory pressure: 4-30 cmH2O 4.4 Expiratory pressure: 4-30 cmH2O
- 4.5 Breathing frequency: 5-50 times / minute
- 4.6 Breathing ratio: 10-70%
- 4.7 Air leakage: 20-99.9lpm
- 4.8 Tidal volume: 50-1500ml
- 4.9 minute ventilation: 0-50lpm
- 4.10 Blood oxygen saturation: 70-100%
- 4.11 Pulse rate: 25-250bpm
- 4.12 Delay boost: 0-45 minutes

5. Alarm Items:

- 5.1 High pressure sound prompt: Turn on / off sound prompt
- 5.2 Low minute ventilation sound prompt: Turn on /off sound prompt, can be set to
- 1-10L (interval 1L)
- 5.3 Air leakage sound prompt: on / off
- 5.4 Power off sound prompt: on / off
- 5.5 Choking voice prompt: On / off, choking time can be set to 10-40 seconds (interval
- 5.6 Low Tidal Volume Sound Prompt: On / Off, Set Tidal Volume Value Range 50-500m ((Interval 50ml))

6. Specifications

- 6.1 Weight: about 20kg including thetrolley and the device
- 6.2 Operating temperature: 5-35 °C
- 6.3 Relative humidity: 10%-90%
- 6.4 Power supply: input is AC 100-240V, 50-60Hz, 1.8A Max; output: DC 24V, 3.33A.



Baltimore, MD, US, 21217 +1 510 221-3232

corporate@natcor-atlantic.com www.natcor-atlantic.com

SAVEL 500 & 500e - INDIAN



Pneumatically Driven Pneumatically Controlled

- With IPPV, ASSIST/CONTROL AND MANUAL MODE
- Different parameters in different colour coding parameters in different colors,
- **OPERATING FRINDELY**
- Simple Hose Connections
- Compact and Portable
- Can be used at Emergency Department, Emergency Station, Ambulance, Hospital Casualty, Standby Ventilator for ICU and for Patient's Inter-Departmental Transfer
- MRI compatible model is also available. (Optional)
- Pneumatic Suction (Optional)

C € ISO 9001:2012 ISO 13485:2015

In any sudden emergency "SAVEL 500" in your hand to save a precious life.

In any serious emergency Artificial ventilation may be required. SAVEL 500 has been designed very operating friendly so that any paramedical staff can handle the primary situation in absence of Intensivisist or Doctors.

Modes of ventilation : IPPV Assisted and Manual

Parameter

Minute Volume : 3~20 L/min (used in 60%) 3~15 L/min (used in 100%)

Respiratory rate 10~60bpm I:E ratio : 1:1.5 and 1:2 FiO₂ : 60% and 100%

: Flow Range 1 to 15 LPM O₂ Flowmeter (on separate selection)

: Up to 300 mm of Hg Suction Monitoring

Pressure values

Alarm Paw Pawhigh, PawLow, O2 supply pressure down

Technical data

Gas supply

Maximum Insp. pressure Compliance

Noise

Dimensions (H xWxD)

Environment requirement

Temperature

Relative humidity

60 PSI

50cm of water 4mL/100Pa

65dB(A) 26 x12 x16 cm

: 3 kg

: -18°C~50°C (Operation) : -20°C~55°C (Storage) : 5%~95%, non-condensing (operation)

1. On/ Off

Proposed Ventilator Control Mode (CMV)Four Knob control System

2. Frequency
3. Tidal Volume
4. 40% and 100% Selection
5. Air way pressure Meter.

≤93% (Storage)



COVID 19 TEST KIT - INDIAN

Sub: Quotation for Supply of COVID-19 Kits

The world is fearing Coronavirus, but we have a solution for its early detection and isolation.

We are the providers of COVID-Self Help (Test at Home) Kits (CE Certified, EU Approved) - Listed on Indian Council of Medical Research (ICMR), India that comes with 98% - 99% accuracy rate. The test helps the users detect possible infection from coronavirus from their homes without going to laboratory. And labs can use it as an effective tool to assess the situation and take pro-active action. The Do-it-Yourself kit currently is getting widely adopted in India, Czech Republic, Japan, Switzerland, South Korea, Australia, LATAM and Italy. Apart from India, the kits are also EU approved - Netherlands, Ministry of Health & Sports, US FDA - pending.

The COVID-Test at Home Kits are available in three order types.

Product	Silver	Gold	Diamond
Product Description	SARS-CoV-2 IgM Method	SARS-CoV-2 gG Method	SARSCoV- 2 igM + IgG (Combo)
Where you can use?	Test at Home	Test at Home	Test at Home
Ingredients	1 piece gM test-ki:	1 piece IgG :est-kit	1 piece of IgM – IgG test-kit
Use Case	- Fear of coronavirus infection Detects infection with high accuracy after 3 days of possible contact	- schibking symptoms like fever, dry cough, diarrehea, difficulty in breathing exc Detects infection with high accuracy after 10 42 days of possible contact	Detects infection with high accuracy Can be used at an point of time. All in one solution for everybody

Accreditation & Government Approvals	Approved in India, EU, Japan, Australia, South Korea, Switzerland &	Approved in India, EU, Japan, Australia, South Korea, Switzerland &	Approved in India, EU, Japan, Australia, South Korea, Switzerland &
FOB PRICE Country Specific Prices, only	China	China	China
Accuracy Rate (Tested through Clinical Trials*)		racy rate is over 98% 88% for Gold and Dia 99%	

NOTE: A USER CAN GET THE RESULTS OF ITS TEST IN LESS THAN 15 MINUTES FROM THE CONVENIENCE OF HIS/HER HOME/. NO WORRY OF INFECTING OTHERS.

WHY QUALITY MATTERS IN THIS SITUATION?

" In addition to this. Other companies are using Colloidal Gold Method. We are using Latex method, which is called Lateral Flow Immunoassay Systems. This makes the brand deliver high sensitivity and extremely high accuracy rate"

UK - NEWS (COVID-TEST RESULTS IN 5 MINUTES WITH NEW LATEX METHOD)

Transparent Price Points

	Distributor / V	Vholesale Price	206	Dispatch Schedule
Minimum Order Quantity	Silver	Gold	Diamond	Days
200000 - 1 Million				3-7
Orders above 1 Million		Ask the price		On priority

Payment Terms: FOB, 100% Pre-Payment only
*Additionally: freight on actuals

Weight & Dimensions

Master Box (Size)	Total Kits	Total Weight	Remarks
42.9 * 34.8 * 30.5 cm	600 Kits/each	8.5 KG	

a manual, 20 alcohol swabs Open it carefully and you will find

Total Kits Weight of each Small Box Master Box) (Size) 17 x 14 x 6 c.m 0.25 KG 20 Kits/each

Ingredients of the Kit

1. Test cassette; 2. Specimen buffer 3. Sample pipette; 4. Test instruction. 5. Lancet (optional).

Under this brand, Importer and distributor of antibodies kits, medical equipment and reagents. We have manufacturing sites in China, Israel, USA and Japan and we are the suppliers to governments in Italy, Netherlands, Switzerland, South Korea, Australia, LATAM, Japan etc.

- WE ARE SUPPLIERS TO GOVERNMENT BODIES IN 100+ COUNTRIES
 CE CERTIFIED PRODUCTS
 OWNS MANUFACTURING LINES IN CHINA, ISRAEL, USA AND JAPAN
 APPROVED IN JAPAN, EUROPEAN UNION, SWISS, SOUTH KOREA, AUSTRALIA, CHINA AND MORE
 700+ PRODUCTS PORTFOLIO INCLUDING N95 MASKS, HAZMAT SUITS ETC.

SARS-COV-2 ANTIBODY TEST (LATERAL FLOW METHOD), **HOME TESTING KIT - SHIPMENT FROM CHINA**













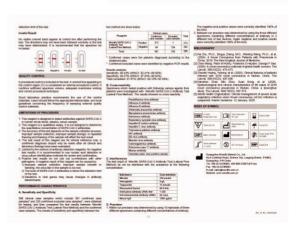


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K N95 MASK - SHIPMENT FROM CHINA







N-95 MASK - SHIPMENT FROM CHINA













K N-95 MASK (2) - SHIPMENT FROM CHINA

















3 PLY MASK - INDIA









 • 1826 Pennsylvania Ave, #1658
 Baltimore, MD, US, 21217
 • +1 510 221-3232

corporate@natcor-atlantic.com

www.natcor-atlantic.com

SANITIZERS (Gel + Liquid)























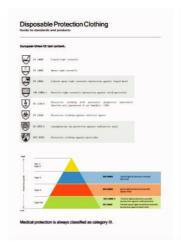
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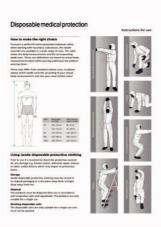
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DISPOSABLE PROTECTIVE CLOTHING FOR MEDICAL USE - INDIAN





















DISPOSABLE PROTECTIVE CLOTHING FOR MEDICAL USE - SHIPMENT FROM CHINA

















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from the bottom up Pull the cuff up and arrange the cuff position Pull the zipper up to the top and adjust the seal of the cap
LERIFICACIONAL Method of removing solution dothing
(1) 無效性的工 (2) 先上性不适构证的 (3) 从上性不适构证的 (4) 数下水、污染油的温度下放致人能扩展物理 1. Uncly the Zippor 2. pull the hat up and back so that the head is off the hat and the sleeves are off 3. from the box down, take off and roll
Trom the top down, take on and row The contamination is taken off the inside and put into the clinical waste bag

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贮存在通风干燥、无腐蚀锥气体的环境中。这类火器以及景燃物。产品运输过程中应防止 激 運、制防包装。

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DISPOSABLE PROTECTIVE CLOTHING FOR MEDICAL USE - SHIPMENT FROM CHINA



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	EN 14605:2005+A1		
Clause	Requirement - Test	Result - Remark	Verdic
4	Requirements		
4.1	Materials		
	Character protective clothing materials shall be tested according to the requirements of Tistle 1 and in accordance with the test mathods specified in EN 14325. A performance level of at least 1 shall be obtained for all requirements.		,
	Chemical protective clothing materials shall not be known to cause skin irritation or have any adverse offect to health lose also EN 340 2000.4 (2)		P
	Pire to testing, all chemical protective clothing materials table to closmed, The materialist set in instruction is indicate that derawing is allowed Martafacture's instructions with regard to number of cleaning cycles, cleaning procedure and presible reapplication of insertiments shall be observed. In maintenan number of cleaning cycles is indicated, materials that undergo the cycles is indicated, materials that undergo the cycles is indicated.		P
	All test specimens shall be conditioned at (20 a 2) **C and (65 ± 5) **C relative humsity for at least 24 h and testing shall start within 5 min after removing the specimen from the conditioning atmosphere.		Р
	MOTES Chemical protection clothing motions (for which a test method in Table 1 does not provide a clear in assurement and profit, should be marked him applicable in the test report and in the instructions for use. The mason why the test could not be completed should be indicated, i.e., where the aboutloy of the upschemic provincts to determine an end-port in the purchase resistance best,		P
	NOTE 2 Materials should be as tight and setfex ble as possible in order 15 ensure weaver comfact as well as providing setfective potention. Indicatal supporting are only one element for the determination of weaver confers of preceding coloring. Design feetiness of the chitming may even have a more important influence on weaver comfact than making properties.		P
	NOTE 3 if resistance to heat and flame is required, the chemical protective clothing should be tested and marked according to the appropriate standard.		P
4.2	Seams, John and assemblages		
	Seams, joins and assemblages shall be tested and classified according to the requirements of Table 2 and the corresponding dauses of EN 14005.		Р
4.3	Performance requirements for whole suits (Typus		
4.3.1	Gmeral		
	Characal protective clothing shall fulfil the relevant requirements of EN 340. The defining shall be made to that the water to an investment and is as confortable as possible, consisted with the protection affected by the garment, as can be verified by the "berein movements" test, described in 3.4.1		P

Clause	Requirement -Test	Result - Remark	Ventic
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	d) the year of manufacture, and also the month of		
	manufacture if the expected shelf-life of the cluthing		
	is less than 24 months. This information maybe		
	marked on every commercial packaging until netead	1	
	of being marked on every term of clothing.	1	
	e) the manufacturer's type, identification ormodel		
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	NOTE Consideration should be green to suitable		- 1
	additional marking	1	
6	information supplied by the manufacturer		_
•	This information shall accompany every item of		-
	chemical protection clothing or at least every		,
	commercial packaging unit. The purpose is to		
	guarantee that the wearer is confronted with these		
	nstructions.		
	The information shall be at least in the official		
	language(s) of the country or region of destination		- 5
	They shall be clear, legible, unambiguous and, if		
	helpful. Bustrations, part numbers, marking etc. shall		- 1
	be added if appropriate, warnings shall be given		- 1
	against problems likely to be encountered.	1	
	The instructions together with the information on the		P.
	marking shall contain at least the following		350
	arternation:		
	a) the name, Yademark of other means of		- 1
	stentification of the manufacturer and/or fea-		- 1
	suthposed representative established in the		- 1
	European Union or the country where the product is		- 1
	placed on the market:		- 1
	to the reference number of this document and the	l .	- 1
	identification as "Type 3" or "Type 4" limited use or		- 1
	reusable full-body chemical protective clothing, gras-	1	- 1
	Type PR Di or PR (4) for partial body protection.		- 1
	c) if applicable, a statement to opecify additional		- 1
	personal protective aggioment with which the suit		- 1
	shall be worn, and how to attach or connect them to		- 1
	achieve the claimed performance classification. This		- 1
	datement shall be precise enough to kelp the user to		- 1
	select the appropriate enument, e.g. a bond model	1	- 1
	YY or equivalent, or respiratory protection including a		
	full face mask, etc	1	- 1
	d) the manufacturer's type, identification or model		- 1
	number		- 1
	e) the size range (as defined in EN 340):		- 1
	() a list of characely and characel products		- 1
	Including the names and approximate		- 1
	concentrations of the components) to which the		- 1
	protective clothing has been to sted and the		



TEST REPORT

Prepared For:	
Trade Mark	NA.
Product Name :	Disposable Protective Clothing For Medical Use
Model(s)	NA
Prepared By	Shenzhen CCT Testing Technology Co., Ltd. On Floor, Area I. Building 1, Instruids Science and Pethnology Procession Paid. Tell 400 155-335 7075 Fix: 400 755-336 5971 Fix: 400 755-336 5971 Fix: 400 755-336 5971 Fix: 400 755-336 5971
Test Date	Mar. 04, 2020 - Mar. 13, 2020
Date of Report	Mar. 13, 2020
Report No.:	CCT20090401GRS

This test report is limited to the above client company and the product model only, be duplicated without prior written consent of Shanzhan CCT Testing Technology	

Clause	Requirement - Test	Result - Remark	Verde
CHARL	Printing of the Printing of th	PRESENT PROPERTY.	70100
	MOTE I Wearer comfort can be judged in wearhale of the suit with but persons experienced in the type of work and environments for which the suits are elected as onitionise clothing.		P
	Chemical potactive clothing Type 3 and Type 4 shall full the requirements specified in 4.3-4 (Table 3), when combined with additional protective equipment, i.e. for protection of hands, feet, face, head anothe respectory that, according to the manufacturer's instructions and when bested as a complete sout.		P
	The requirements of his clause apply to the gammes as whole including component parts in a gammes, boots, hoods or respiration) that are not irregard to gammes, consistent that pare and search larger attaching these components are included within the scope of this document, whereas or takes for the components are obtained for the components are obtained for the components are obtained to the components are obtained to the components of the Components are obtained to the Components.		P
	MOTE 2 Parkel body protection covers only specific areas of the body, leaving others exposed to the hazard. Because of this only limited trading of this type of clusters in appropriate and this product standard is adding		P
6.32	Pre-conditioning		
	Fino to testing, the themical protective do thing shall be cleaned, if the manufacture is instructions indicate that cleaning is allowed. Manufacturer's instructions with regard to number of cleaning cytics, cleaning procedures and possible reagilization of treatments shall be obtained if no maniferant number of cleaning cycles is indicated, the clothing shall underso the cleaning cycle.		,
4.33	Conditioning		
	All chemical protective dothing shall be conditioned for at least 24 h at the same conditions as used for the task.		Р
4.3.4	Resistance to penetration by liquids		
4.3.4.1	Governi and preliminary testing	X.	
	Type 3 chemical protective dothing shall be tested against penatration by liquids by means of a jettest in accordance with 4.3.4.3.		P
	Type 4 chemical protective dofting shall be tested against penetration by liquids by means of a spray test in accordance with 4.3.4.2.		P
	Partial body protection items Type PB [4] shall notibe fested agent frees others. Dearns, Jones and accombigged of Type PB [5] distring shall behead to the jet test (EN ISO 17491-3) (see also Table 2, feetings as		P
	Prior to testing each suit in accordance with ENISO 17491-3, or ENISO 17491-4, a practical test shallbe camed out by a human feet subject. Histore than one size of chamical protective suit is manufactured, the set subject will be affected to select the appropriate.		,

	EN 14605 2005+A1	2009	
Jause	Requirement -Test	Result - Remark	Verdic
	performance were described a personal model personal model of control and any of the second and any of the sec		

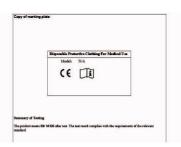
requirements (Type 4) cor	TEST REPORT EN 1466 2005-A1 2009 didning against liquid chemicals - performance for dobing with liquid-styr (Type 3) or sprey-light excitons, including term providing protection to if the body only (Types PB [3] and PB[4])
Reference No.	CCT28830481GRS
Date of issue	Mar 13, 2020
Contents	14 pages
Testing laboratory	
Name	Sheather CCT Testing Technology Co., Ltd.
Address	+
Testinglocation	
Client	
Name	
Address	
rea specification	
Standard	EN 14805 2005+A1 2009
Test procedure	CE
Procedure deviation	NA.
Non-standard test method	NA.
TEXTER	
Description	Disposable Protective Clothing For MedicalUse
Trademark	N/A
Model and/or type reference	N/A
Manufacturer	Address
	City, Province, China

Test case verificts	
Test case does not apply to the test object	N(A)
Test item does meet the requirement	P(ses)
Testitiem does not meet the requirement	F(all)
Testrig Date of recept of test term	Mar 04,7828
Date(s) of performance of test	From Mar. 04, 2020 to Mar. 13, 2020
General remarks	
This test report shall not be reproduced ex- laboratory.	cept in full without the written approval of the testing
laboratory.	
taboratory. The test results presented in this report relate	only to the item tested.
This test report shall not be reproduced ex- taboratory. The test results presented in this report relate "see remark ##" refers to a remark apprinted. "see appended table?" refers to a table appe	contyto the item tested. It to the report
taboratory. The test results presented in this report relate "(see remark #f" refers to a remark appended	only to the item tested to the report orded to the report.
taboratory. The test results presented in this report relate "(see remark #/f refers to a remark appended "(see appended table)" refers to a table appe	only to the item tested to the report orded to the report.
taboratory. The test results presented in this report relate "(see remark #/f refers to a remark appended "(see appended table)" refers to a table appe	only to the item tested to the report orded to the report.

Cause	Requiement - Test	Result - Remark	Verdic
Clause	Requiement - Lest	Heisut - Hernank	Vertic
4343	Resistance to penetration by liquids (critori)		
4.14.3			
	Three new subs, pre-conditioned in accordance with 4.3.2, shall be wated in accordance with ENISD 17491-3. If applicable, the subs shall be worn with the additional personal protection equipment specified in the marketaurer's snotucions.		
	All subs shall pass the test, i.e. the total stain area on any one undergament of each suit shall be less than or equal to three times the total calibrated stain area.		,
44	Visor		
4.4.1	General	£	
	Where a visor is litted as part of the sult, as distinct from a respiratory protective device joined to the sult, the visor shall comply with the requirements of 4.4.1 to 4.4.4.		P
	Where sold-fagging compounds are used or specified in the information supplied by the ensemblaching they shall not have an adverse sifect on the health of the we see, or on the properties of the protective garment.		P.
	NOTE if a visor is integrated in a bood, edequate respectory protection should be provided. The respectory protective daylor should meet the requirements of the relevant product standard and the compatibility between respiratory protective device and hood should be checked.		
4.4.2	Mechanical strength of visor		
	The visor shall not be visibly damaged in such a way so to be likely to affect the performance of the complete device, when stated in sicretiance with 7.5 of EN 1294 1:1998. This test shall be followed by a spray test (EN ISO 17491-4), or jet test (EN ISO 17491-4), or jet test (EN ISO 17491-4), as appropriate.		'
443	Field of vision		
	When corrying out the seven movements prior to the start of the spinar test, or the jet test as appropriate, (see 4.3.4.1 or 6.3.4.2), the field of vision shall be judged satisfactory by the test solyect(s).		P
···	Distertion of vision		
	The test subject(s) shall be able to read a sign with latters of 100 mm high and a proportional width from a distance of 6 m.		-
5	Marking		
	The chemical protective clothing shall be marked with at least the following information. Thermarking shall be charly visible and as durable as adequate for the life of the clothing.		Р
	a) the name, Yade mark or other means of identification of the manufacturer; b) the type of the chemical protective clothing, i.e.		P

	Requiement - Test	Result - Remark	Verd
Te	tile 1 — Test requirements for Type	3, Type 4, Type PS (3) and Type PS (4) clothin	9
	Clause in EN 14325 2004	Performancerequirement]
	44	admission resistance	
	4.5	flex cracking resistance	
	40	flex cracking resistance at -30 °C	
	4.7	tear resistance (trapezoidal)	
	4.9	tersile strength	
	4.10	purcture resistance	_
	4.11	resistance to permeation of Equita-	_
		seinted text	1
On	ly applicable to clothing intended for u	se at very low temperatures.	_

a and Type PB	[6] diething
Performance requirement	Reference
resistance to permeation of tracks*	EN 14325-2004, 4.11
resistance to perwitation by Equich*	EN ISO 17491-397 EN ISO 17491-4
sean strength	EN 14325 2004, 5.5
* Seams, pains and assendingers of Type PB (3) En (SO 17491-97). * Applicatio only to seams which are exposed in only seams relevant to the construction shall be lead if shall be obtained. * To be rectable which suit tests, i.e. I EIN ISO and IEN ISO 1981-IF I Wish level sarray sent for and IEN ISO 1981-IF I Wish level sarray sent for	nuse. For partial body protestion items considered and a performance level of a 17491-31 gettest) for Type 3 clothing



	EN 14605:2005+A1:2009		
lause	Requirement - Test	Result - Remark	Verdict
	use accepting to the manufacturer's information leafer. If applicable, the test subject that a bits were additional parameter applicant, as specified in the manufacturer's instruction. The test shall comprise their explorations, at moderate speed, of the feerom movements' sequence described.		P
	and the filtering non-minimal supposes. In the filtering non-minimal supposes. In the filtering non-minimal supposes th		***
	If the test subject is not able to parform one or several interements due to the fundance of the sub- or if the movements result in substantial damagato the suit, the suit shall be considered to have felled.		Р
	Subsequipped with a visor shall also pass the tests specified in 4.4 better further testing Falure will result in a disqualification for further testing and the subshall be considered to have failed.		P
1342	Resistance to penetration by liquids (spraytest)	•	
	Three new suits, pre-conditioned in accordance with 4.3.2, shall be belied in accordance with ENISO 17491-4. If applicable, the outs shall be worn with the additional personal protestive equipment specified in the mismatchure's instructions.		,
	All suits shall pass the test, i.e. the total stain area on any one undergoment of each out shall be less than any one undergoment of each suit shall be less than		P

Annex ZA (Informative) Relationship between the European Standard and to bits with if Requirements of BJ Directive Basic Standard and to bits of bit	Clause	Requirement - Test		Result - Remark	Verdi
The Surgeon Standard by a less parameter look a model age to 10.5% by the Surgeon Commission and the Surgeon Street Trade Association is provide a memor of continuing in Extended Requirements of the New Agents of Continuing Street Street Street Standard Street					
the European First Took Association is provide a means of conforming to Executed Requirements of the New Appearsh Disector (ISSEMENE). Once this described in citized in the Official Journal of the European Communities under that Describe and has been implemented as a national streaming of the Management Communities under that Describe and has been implemented as a national streaming of the Management of the Section Communities of this standard given in Table ZA Colories, which is least of the second of the Section (ISSEMENT) and the	Re	lationship between this Euro		Essential Requirements	of EU Directive
been implemented as a national standard in at least one Member State, compliance with the clauses of this standard given in Tutols ZA confers, within the limits of the scope of this standard, a presumption of conformity	the Eu	rope an Free Trade Association			
	been it	nplemented as a national stand rd given in Tubble ZA confers, wi	and in at least one Member than the limits of the scope	State, compliance with the of this standard, a presumpt	clauses of this
	Halis	Conceptions Estatement		and and the same of	

Clause(s) of this standard	Clause(s) of EU Directive 99/606/EEC, Annex 6
4.1	1.21.1 Suitable conditions materials
4.1	1.3.2 Lightness and design strength
4.1	3.10.2 Protection significations are substances and infective agents - Protection against outsineous and ocular contact.
42	1.32 Lightness and design strength
42	3.10.2 Protection against dangerous substances and infective agents - Protection against cultureous and ocular contact
4.3.1	1.2.1 Absence of risks and other nassance factors
4.3.1	1213 Maximum permissible user impediment.
432	2.4 FPE subject to ageing
43.4.1	1.1.1 Ergonomics
4341	1.213 Maximum permissible user impedment
43.4.1	1.3.3 Compatibility of different classes of PPE designed for simultaneous use
4342	3.10.2 Protection against dangerous substances and infective agents. Protection against cutaneous and ocular cented.
43.43	3.10.2 Protection against dangerous substances and infective agents- Protection against cuterrous and ocular centect
4.4	2.3 PPE for the face, every and respiratory tract
1	2.12 FPE bearing one or more identification or recognition marks threefy or indirectly relating to health and safety.







PPE KIT-1 - SHIPMENT FROM CHINA

NO CODE	РНОТО	DESCRIPTION	COLOR	UNIT	QTY	REMARKS
1	1	Disposable Protection Coverall Material - PP Non Woven fabric (30 gsm) + Breathable Film (30 gsm) + Glue (3 gsm) Style - 3 panel Hood Coveralls, Zipper, Elastic Waist and Cuffs - Wrist / Ankle	White	Piece	1	Œ approval
2	其前	Disposable Protection foot cover	Blue	Pair	1	CE approval
		Disposable face Mask 3 Ply mask with bacterial filter	blue or white	Piece	1	Œ approval
•		Fully Closed Double-Side Anti-Fog Lens Medical Protective Safety Goggles	standard	Piece	1	CE approval
.		Disposable viny examination gloves.	standard	Pair	3	CE approval
Cost for C	one PPE Kit		0	_	1.00	
	NT TERMS: 100%	i Advance.				
	ERMS: FOB Chin					



PPE KIT-2 - SHIPMENT FROM CHINA

10	of 1	DESCRIPTION	COLOR	UNIT	QTY	REMARKS
1		Disposable Protection Coverall Material - PP Non Woven fabric (37 gsm.) + PE Film (28 gsm.) Style - 3 panel Hood Coveralls, Zipper, Elastic Waist and Cuffs - Wrist / Ankle	White	Piece	1	CE approval
2	共動	Disposable Protection foot cover	Blue	Pair	1	CE approval
3		Disposable face Mask	blue or white	Piece	1	CE approval
4		Fully Closed Double-Side Anti-Fog Lens Medical Protective Safety Goggles Certificate: CE	standard	Piece	1	CE approval
5	V	Disposable viny examination gloves.	standard	Pair	1	CE approval
Cost for O	ne PPE Kit	W.				
	NT TERMS: 100%	Advance.				
S 1660 47 1650	ERMS: FOB China	2393030310				
20170-0111-0	18 house charges and a	g days after receiving full payment				



ANTIBACTERIAL GOGGLES - SHIPMENT FROM CHINA











FACE SHIELD - SHIPMENT FROM CHINA







INSTRUCTIONS

- Wash and dry hands thoroughly before opening
 With the nose strip facing outward, cover mouth and nosewith

- With the nose strip facing outwars, core mounts
 the mask
 Pull elastic bands on both sides over the east to hold mask in
 place
 Adjust the elastic bands to tighten against the bridge of the nose
 Dispose of after removing

ATTENTION

- If you experience symptoms such as liching or rash when using the mask, piease remove it immediately and consult your doctors, should you swell a strange odor when using, please discontinue usage.

 The mask may change over time as you wear it; please use with caution; maximum use time is 4. hours.

 Please avoid juilling the elastic bands too tightly against the ears.

 This product is disposable, please do not wash or reuse.

 Please do not use near fire.

RISKS OF REUSE

- The Face Shield is designed for up to 4 hours of usage; please do not re-use.

Face Shield PRODUCT SPECIFICATIONS

- Masks may become contaminated following use, you should ensure that they are disposed properly. In order to reduce potential contamination, please wash your hands before putting on a mask. You should also avoid touching the mask with weeking it. If you touch the mask, please wash or disinfect your hands. Please remove and discard the mask should it become moist.

 When removing the mask, take hold of the ear bands and pull them around your ears. Fold up the mask, cover it up and discard in a waste receptacle. Wish your hands immediately after discarding the mask.

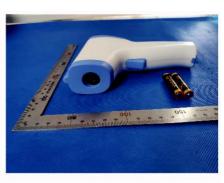


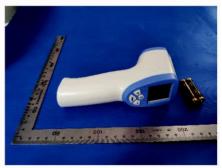




MEDICAL INFRARED THERMOMETER - SHIPMENT FROM CHINA







A:RoHS Directive 2011/85/EU
(*):With "reference to 1—3 82211 5:2013, determination of Creations by ICP-CIHS.
(2):With "sterence to 150 8221-4:2013, determination of Lead by ICP-CIES.
(3):With "alternace to 150 6222-4:2013, determination of Marcury by CP-CIES.
(4):With "alternace to 150 6222-4:2013, determination of Hexavalient Chromium by Ocioimeric Molicula. Singli UNIVERS.
(5):With "deternace to 1,0 82211 8:2015, determination of PILSe ord PIBULa by GIG MS.

Test Item(s)	Unit	Limited(ppm)	Result 1	MDL	
Cadmium (Cd)	mg/kg	100	N.D.	2	
Leac (Pb)	mg/kg	1000	N.D.	2	
Moreury (Hg)	mu-kg	1800	N.D.	2	
lexagalen: Chromium (CrVI)	mg/kg	1909	N.D.	2	
byAlkal no extraction	33355753		0.2753		
Sum of PBBs	mg/kg	1000	N.D.	-	
Monobrom abiphenyl	mg/kg		V.D.	5	
Ditricmotiphonyl	mg/kg		VID	5	
Tribremobiphonyl	mg/kg		N.D.	5	
Pentsbromobiphenyl	mg/kg		V.D.	5	
Hexabramab phenyl	mg/kg		AD	5	
Hoptabromobiphonyl	mg/kg	5+3	N.D.	5	
Optabromab oher yl	mg/kg		N.D.	5	
Nanabromooiphenyl	mg/kg		N.D.	h	
Decabromob phonyl	mu-kg		N.D.	5	
Sum of PBDEs	mg/kg	1000	N.D.		
Monabromabiphenyl ether	mg/kg	-	V.D.	h	
D bromobiphonyl office	nig-kg	-	N.D.	5	
Fribromobiphanyl aths e.her	mg/kg	-	N.D.	5	
Penta bromooiphenyl ether	mg/kg		N.D.	5	
Hexabramob attenyl ether	mg/kg		V.15.	. 5	
Heplabromobiphenyl ather	mg/kg	12.0	N.D.	5	
Ostabromobiohenyl ether	mg/kg		N.D.	5	
Nandarnmoorphenyl effrer	mg/kg		N.D.	5	
Decabramob alienglie.her	ing/kg		N.D.	5	
Pertabramoniphenyl ether	mg/kg	200.0	N.D.	5	

Filtration Weigh comple and place it in a conical flash Adjust the pH value

Sample cutting / preparation	
+	
Sample Measurement	
+	
Solvent extraction	
\	
Concentration/Dilution	
+	
Filtration	
+	
GC-MS	

DATA

Phthalates Testing Flow Chart

Test Hem(s)	Unit	Limited(ppm)	Result 1
Dibulyl Ph.he alo (DBP)	majkg	1000	ND.
Benzy buty Phthelate (SBP)	mg/kg	1000	ND
Bis (2 ethylhexyt) Prificalitie (DEHP)	m thyta	1000	1/12
Diisc bulyl Phthalate (DIBP)	majkg	1000	ND.

Vo.es
(*) Reference Information RoHS C rective 2011/absEJ Annex II amending Annex (EU)2015/863 and amending
Annex (EU)2017/21/32

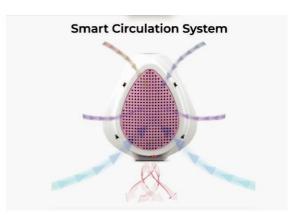
Bis (2-athy haryli) phthalate (DEHP), Buth) bencyl prithalara (BBP), Disc b, tyl Phitha are (DIBP) and Dibutyl
prithalara (DBP) are considered us a crisidy for risk crystopilors and substance restriction.

Remarks:
(1) 1 ing/kg = 1 ppm = 0.0001%
(2) MDL = Method Detector Limit
(3) ND = Not Detectod (< MDL)
(4) " " = No. Regulated

SECOND GENERATION MASK - SHIPMENT FROM CHINA

















PRODUCT FEATURES	MASK	
Colors	Classic White, Pink, Gray	
Filter Layers	5	
HEPA Compliant	Yes	
Purification efficiency	Upto 95.99%	
Fan Speed	3 Modes	
Charging Mode	USB Charging	
Charging Time	1.5 hours	
Battery Duration	4-6 hours	
Battery Capacity	680 mAh	
Dimensions	10.5*9.5*4.6cm	
Weight	70 grams	
Voltage	3.7V-5.2V	
Noise	28dB	

ISOLATER CHAMBER - SHIPMENT FROM CHINA



Oxygen supply connector

Technical features







- The isolator is mainly used for the isolation transportation of patients suspected or identified as respiratory infectious diseases in the environment of toxic gas (such as gas bomb pollution).
- The isolator frame is made of high quality and durable transparent material without pollution release, and sealed through zipper.
- The isolator adopts the principle of positive and negative pressure. The positive pressure principle can prevent the poisonous gas from entering the cabin. The negative pressure can prevent polluted air from leaking in the cabin.
- The difference pressure between inside of cabin pressure and outside of cabin pressure is not less than 20Pa.
- The filter tank can make external toxic gas or internal pollution gas highly efficient filtration to ensure that the filtered gas is clean.
- The filtration efficiency of the filter tank for aerosol particles with diameter of 0.3 m is not less than 99.99%.
- The fan with lithium battery and can be used continuously for not less than 5 hours. When the battery under voltage should be alarm function;
 The fan can use DC12V power supply.
- The cabin is equipped with 8 sealing covers, the medical staff need use rubber gloves to open the sealing covers, which is convenient for examination the medicine, food and patients.
- 9. The oxygen access valve can be connected on the sealing covers.

To an inches of the last of th	(printed tion	Fidence Store	Ser-ings:	Same Branch	and theory Parkey MSR		E.M	
1950001 3248		25-WSA	11 1	7 7 24	Queto E S	- 元ガルマー	□ W 有量	
NP-320	195×57×43cm	120×60×25cm	17Kg		1	123×40×54cm	18Kg	

Verification of Conformity Verification Number: 2020031025HA-V1 On the hause of the conformity of the proof of the state of the string product has been found to string end of the string product has been found to string end of the string product has been found to string end of the string product has been found to string end of the string product has been found to string end of the string



E SERIES - BELT BRUSH SOLE CLEANING MACHINE - SHIPMENT FROM CHINA

1838*768*140mm(L*W*H) zone handrails, Clean 1115+530mm Dry 400*520mm, 304, Induction switch: infrared, style wet, Water temperature 0-40°C, Mator voltage 180W. Pump power 200W, Current <60db. N.W: 100KG. Load capacity 300KG, Safety protection Guarantee one year except brush & dry pad







40°C, Water tank capacity 20L, Working voltage 220V, Motor power 180W, Pump power 200W, Current 1.5A, Noise <60db, N.W: 100KG, Load capacity 300KG, Safety protection. Short circuit/over current/overload voltage protection. Guarantee one year except brush & dry pad

Additional FUNCTION: When a user induce the temperature and mask and get an permission, access control will open to finish show sale class and disinfection.



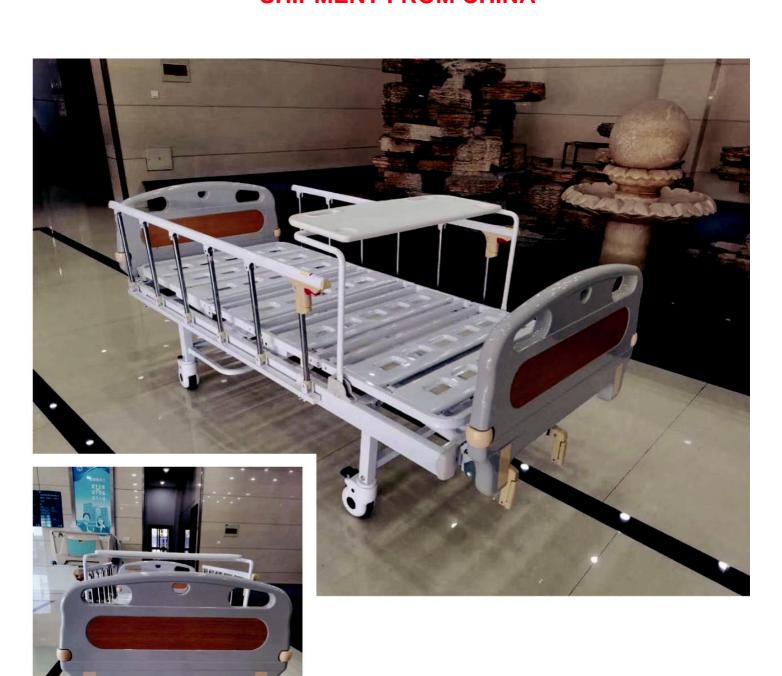




№ 1826 Pennsylvania Ave, #1658 Baltimore, MD, US, 21217

+1 510 221-3232
corporate@natcor-atlantic.com
www.natcor-atlantic.com

FOWLER BEDS - SHIPMENT FROM CHINA





DISPOSABLE MEDICAL MASK - NMT SHIPMENT FROM CHINA





























KC N901 SMART HELMET - SHIPMENT FROM CHINA

KC N901 Smart Helmet

Created with decades of ingenuity First choice for epidemic prevention Advanced Metamaterials
High-precision Temperature Screening
Maneuvering Non-interference Screening
Hi-tech Ultimate Experience



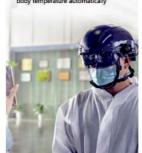
Five Powerful Functions



Smart Helmet for Unaware and Contactless **Temperature** Measurement

Efficient Helmet for Temperature Recording

Record personal info with their daily body temperature automatically



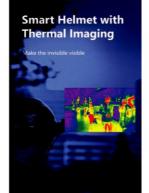
Powerful Helmet for Vehicle Screening

Rapid screening for vehicles



Powerful Helmet for Verification

Rapid face recognition and identity verification



Nine Modes



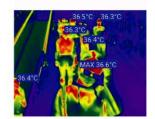
Single-person temperature measurement mode

The temperature of the single target in the center of the screen will be measured. The maximum temperature of different parts of the body is displayed on the AR module. The temperature above the normal range will trigger a audible and visual alarm.



Large-crowd temperature measurement mode

The temperature of the forehead, collar, arm, and other body parts exposed in the screen will be measured. The system will display the temperature if any part in the screen falls into the present temperature range. The alems will trigger when any part of the temperature goes above the threshold value.



QR code mode

Scan the QR code to automatically record personal temperature info into the database in real time, allowing paperless data logging.



QR code & temperature measurement mode

Scan the QR code to acquire the personal information first, and take a temperature measurement of the person within 3s. The personnel information and the corresponding temperature will be automatically recorded into database. This will implement paperless registration of the personnel information and the corresponding the paperless registration of the personnel information and the corresponding temperature.



License plate recognition mode

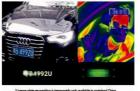
Recognize the vehicle license plate, identify and alert unregistered vehicles or suspect vehicles recorded in database.



License plate recognition & temperature

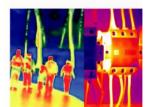
Besides plate identification mentioned before, the helmet can measure the temperature of the single target in the center of the screen. The maximum temperature of different parts of the body is displayed on the AR module, and the temperature above the normal range will trienge will be the screen that the screen that

measurement mode

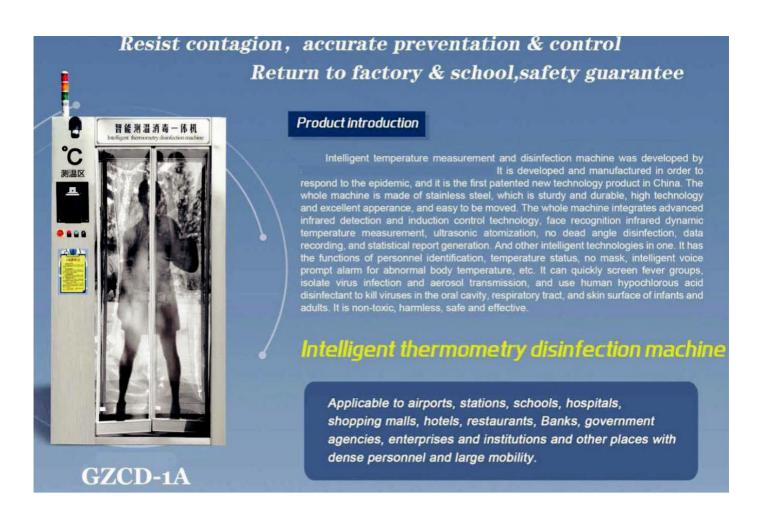


Night-vision /Facility inspection mode

Thermal imaging scanning of industrial facilities or establishments of night places, HVAC equipment, pipelines and electronic equipment, to assist finding target with abnormal temperature or searching for unauthorized personal.



DISINFECTION BOOTH - SHIPMENT FROM CHINA









RAPID THERMOMETRY SAFETY DOOR - SHIPMENT FROM CHINA







