

Disposable Isolation coveralls

– Safety Protection Refused to be Hurt –

Antibacterial rate was
more than 99.99%

Multi-layer Isolation, Free Breathing

FIGHT THE VIRUS
NOT LOVE



WuHan must Win

China must Win



COMFORT



PERFECT



BREATHABLE



NATURAL

Quality Guarantee Trustworthy

PRODUCT DETAILS

Fine cutting Professional protection



Name : Disposable Isolation coveralls

Recommended Industry: Typical applications may include:Apartment gate~street office~community~city traffic crossroads~company~Medical outpatient department~sick room~inspection chamber ~lab room etc.

Feature: Disposable (special+safety). Double Protection(Taped Seams+Velcro Design make it more easy to operate). Triple Barrier(help provide barrier protection against liquid splashes, hazardous dusts, and certain biological contaminants).Four kinds of comfortable(light and permeable,overhang design+slim fits ,Anti-static ,non-irritant)

Product Style: Coveralls **Color:** White

Material: PP non-woven fabric(37gsm)+PE film(28gsm)

Quality Guarantee Trustworthy

COVERALLS SIZE DETAILS

unit:cm

Model	Lenght	Chest	Sleeve length	Sleeve opening	Ankle opening
160	165	120	84	18	24
165	169	125	86	18	24
170	173	130	90	18	24
175	178	135	93	18	24
180	181	140	96	18	24
185	188	145	99	18	24
Error	±2	±2	±2	±2	±2

Quality Guarantee Trustworthy

DETAILS SHOW



01

Three panel hood design

Protective hood design; elastic cap mouth; Effectively block dust and biological contaminants

02

Zipper design

Simple and easy to wear and take off



03

Elastic cuff design

Comfortable and easy to work.

04

Elastic waist design

Elastic waist design, more comfortable to wear, but also to meet the needs of different staff figure.



05

Coveralls design

Coveralls design provide additional protection, breathable and waterproof, no cross infection.



06

Elastic waist and ankles

Elastic waist and ankles for convenience and freedom of movement.

Quality Guarantee Trustworthy

PHOTO OF MODELS SHOW

Front view



Back View



Inspection Report Safety Protection

CERTIFICATE OF REGISTRATION

DAKES
Product Quality Assurance System
No. 02022436

Certificate
No. Q6 065447 0001 Rev. 00

Holder of Certificate: Anhui Med Medical Tech
Jia Bao Indus
246500 Anhui
PEOPLE'S REP

Facility(ies): Anhui MedPure
Jia Bao Indus
PEOPLE'S REP

Certification Mark: 

Scope of Certificate: Production and Sterile Surgical Sterile Surgical Sterile Surgical

Applied Standard(s): EN ISO 13485:2 Medical device Requirements to ISO 13485:2016 EN ISO 138

The Certification Body of TÜV SUD Product Service Group has established and is maintaining a quality system which meets the requirements of the listed standard.

Report No.: 0161121001

Valid from: 2018-10-30

Valid until: 2021-10-30

Date: 2018-10-30

Page 1 of 1
TUV SUD Product Service GmbH - Certification Body - PE

CERTIFICATE OF REGISTRATION

EC Certificate
Production Quality Assurance System
Directive 93/40/EEC on Medical Devices (MDD)
(Devices in class I or sterile conditions, otherwise)
No. G25 065447 0002 Rev. 00

Manufacturer: Anhui Med Medical Tech
Jia Bao Indus
246500 Anhui
PEOPLE'S REP

Facility(ies): Anhui MedPure
Jia Bao Indus
PEOPLE'S REP

Product Category(ies): Sterile Sun Sterile Sun Sterile Sun

The Certification Body of TÜV SUD Product Service Group has established and is maintaining a quality system which meets the requirements of this Directive. It is subject to the requirements of this Directive. It is subject to the requirements of this Directive. It is subject to the requirements of this Directive.

Report No.: 0118113120

Valid from: 2018-10-30

Valid until: 2023-10-30

Date: 2018-10-30

Page 1 of 1
TUV SUD Product Service GmbH is Notified By
TUV SUD Product Service GmbH - Certification Body



**QUALITY MANAGEMENT SYSTEM
CERTIFICATE**

Certificate No. 00118Q312058R053400
We hereby certify that
Anhui MedPure Medical Technology Co., Ltd.
Unified Social Credit Code: 91340811078251917

D4-B-2 Building, Jiatiao Industrial Park, Anqing City, Anhui Province, P.R.China

by reason of its
Quality Management System
has been awarded this certificate for compliance with the standard
GB/T 19001-2016 / ISO 9001:2015
The Quality Management System Applies in the following area:
Manufacturing Non-Woven Medical Supplies, Like Surgical Gown, Surgical Packs (Coverall, Drape and Spatula) and So On.


Certified since: **November 26, 2018** Valid from: **November 26, 2018** Valid until: **November 25, 2021**
After a surveillance cycle, the certificate is valid only when used together with an acceptance Notice of Surveillance Audit issued by CQC.
Please access www.cqc.com.cn for checking validity of the certificate.

This certificate and its relevant information can query in the website of Certification and Accreditation Administration of the People's Republic of China (www.cca.gov.cn).

CHINA QUALITY CERTIFICATION CENTRE
Section 4, No. 144, Nanshanlihe South Fourth Ring Road, XitaiWoo Road, Beijing 100870, China
http://www.cqc.com.cn

Q 0133759 2015/8



**Fiscal Year 2018
IFICATION OF REGISTRATION**

Furthest Medical Technology Co., Ltd
ingmagna company, Laoling Industrial Park,
ng, Anhui



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

FDA SUNGO TECHNICAL SERVICE INC., CHICAGO,
6050 W EASTWOOD AVE APT 201, CHICAGO,
ILLINOIS 60636, USA
Telephone: +1-455-857-7779 / E-mail: USA.FDA@sungogroup.com

ator Number: 10056827
gify: See annex

If Service Inc. will confirm that such registration remains effective upon request and to re-verify until the end of the calendar year stated above, unless said registration is terminated or this certificate is/are withdrawn. Service Inc. makes no other warranties, nor does this certificate make any representations or warranties to any other than the named certificate holder. For whose sole benefit it is issued. The device endorsement or approval of the certificate holder's device or establishment and Drug Administration. SUNGO Technical Service Inc. assumes no liability to any communication with the foregoing.

28.07.2018: Registration of a device establishment or manufacturer of a registration is only upon device approval of the establishment or its products. Any registration approval of official approval because of registration or provision of a registration and continues accordingly. The U.S. Food and Drug Administration does not of registration, nor does the U.S. Food and Drug Administration recognize or maintain. SUNGO Technical Service Inc. is not affiliated with the U.S. Food and Drug

CE: Tel: 021-68871103 Email:Shingo02@163.com Website: www.sungogroup.com
8013rd Floor, No.1500 Century Avenue, Shanghai 200122, P.R.China