

Particulate Respirator KN95 (GB2626)



Product Name	Particulate Respirator KN95
Performance Standard	GB2626-2006
Type	Respiratory protective product
Model	RZ95B
Protection Grade	KN95
Component	Mask body, nose clip and ear loops.
Function and application	Used for protection against virus. It can protect you from dust, droplets, saliva and bacteria.
Instructions	Hold the mask by the ear loops. Place a loop around each ear and pinch the nose clip to the shape of your nose.
Caution	<ol style="list-style-type: none">1.Do not use when expired.2.Do not wash or disinfect, avoid repeated use and destroy after use.3.Do not use if the package is damaged.4.Not suitable for people sensitive to non-woven fabric.5.Please refer to the instructions before use.
Storage	Stored at room temperature with good ventilation, non-corrosive environment and no more than 80% relative humidity. Avoid high temperature.
Expiration date	36 months in the correct storage environment.



Particulate Respirator KN95 (GB2626)



Quality Assurance: CE Certificate and FDA Medical Device Registration

Verification No.: CPAH20031022153

GTS

VERIFICATION OF COMPLIANCE

Applicant: Anhui RYZUR Medical Equipment Manufacturing Co., Ltd.

Address of Applicant: RYZUR Medical Industrial Park, West of Susong Road, South of Guanhai Road, Hefei Economic and Technological Development Zone, Anhui Province

Product Name: KN95 particulate respirator

Model No.: KN95B

Test Standard(s): EN 149: 2001+A1:2009

Test report(s): TPAH20031022153

Date of issue: March 17th, 2020

Date of Expiry: March 16th, 2025

In accordance with the following Applicable Directives:
The Personal Protective Equipment Regulation (EU)2016/425

Conclusion
The referred test report(s) show that the product complies with standard(s) recognized as giving presumption of compliance with the essential requirements.
The CE marking as shown below can be affixed on the product after preparation of necessary technical documentation.
Other relevant Directives have to be observed.

Approved by: Hermann Wolter

For and on behalf of
Global Testing Services Co., Ltd

Global Testing Services Co., Ltd.
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FDA The 3rd Party Certificate of
FDA Medical Device Registration

Note:
This file is Not being issued by FDA. GTS, as the 3rd party, produce it, intended to facilitate customer display & transmit information. The following contents, FDA registered Facility/Owner/Operator&FDA listing- Medical Device, are excerpted from database at www.fda.gov.

Establishment:
Anhui Ryzur Medical Equipment Manufacturing Co., Ltd
Ryzur Medical Industrial Park, South of Guanhai Road and West of Susong Road, Economic&Technological Development Zone, Hefei, Anhui, China 230601
Registration Number / FEI Number*:
* First Establishment Identifier (FEI) should be used for identification of entities within the imports message set

Status: Active
Date of Registration Status: 2020

Owner/Operator
Anhui Ryzur Medical Equipment Manufacturing Co., Ltd
Ryzur Medical Industrial Park, South of Guanhai Road and West of Susong Road, Economic&Technological Development Zone, Hefei, Anhui, China 230601
Owner/Operator Number: 10062907

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Devices Listing Information

Proprietary Name	Product Codes	Device Class	Listing Number	Establishment Operations
Face Mask	LYU	1	D37****	Manufacturer

Please careful protect your Listing Number.

Professional FDA Registration Services, by Shanghai Global Testing Services Co., Ltd.
More details on the website: <http://www.gts-lab.com>
Need help? Contact us, GTS, at +86(021) 33637866 & info@gts-lab.com FDA
CERTIFICATE NUM: GTS22155