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EXPERIENCE THE DIFFERENCE
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KN95 FFP2 FACE MASK
5 Layer

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High-Closed KN95 Professional Protective Dust Masks for Slider PM2.5 Comfortable Elastic Earring

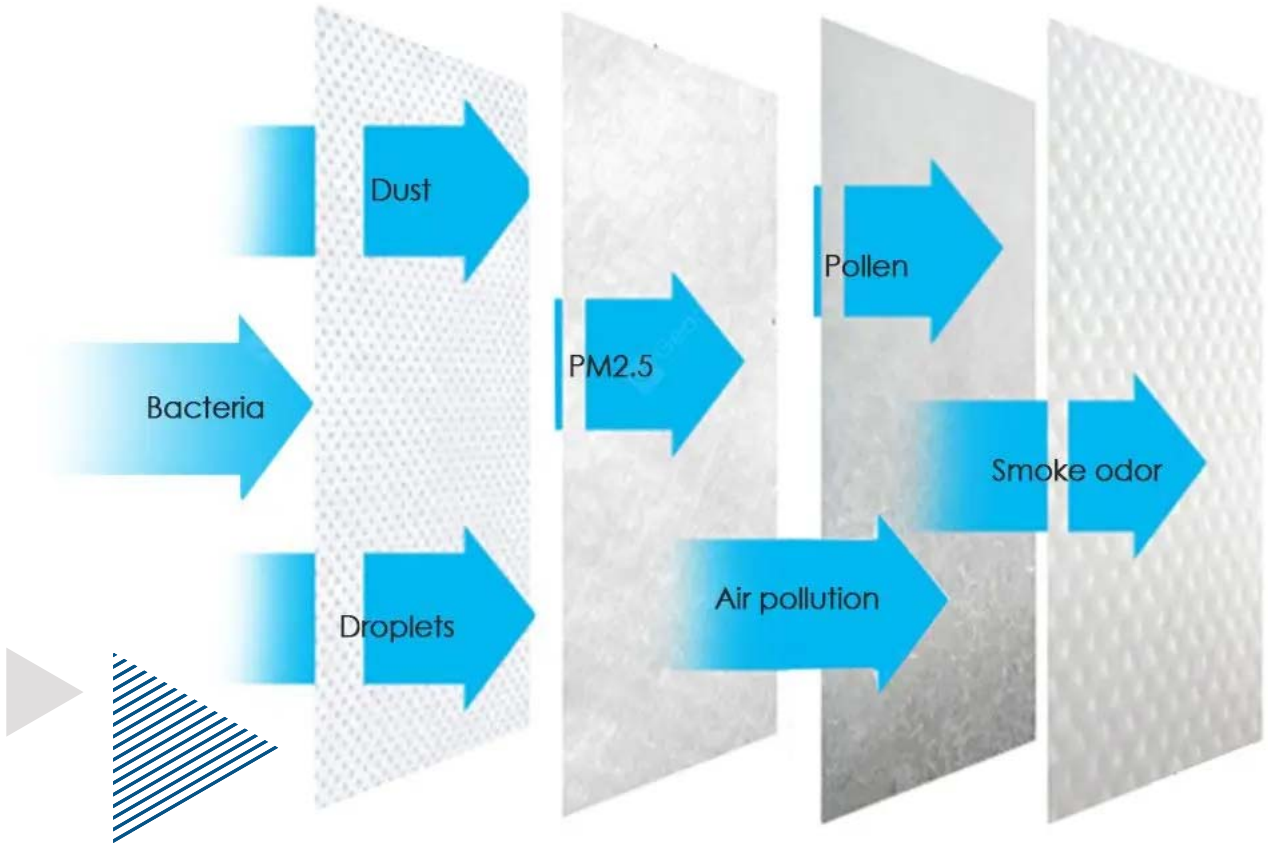
Main features:

Exclusive function:
Compared to the ordinary mask which is not water resistant, our KN95 mask effectively prevents infections caused by the splitting of fluid splashes from the patient's body or arterial intake (the size of the slit is between 1 and 5 μm).

More than 95% filtration efficiency:
Efficient removal of air pollution, dust, PM2.5 filters, fog, bacteria, odors, dust, pollen, etc., Material respiratory quality of the rejected being.

Tight
fit : Three-dimensional cut, fine points, KN95 mask for your face tightly, preventing splashes from getting scratched in your face.

Type of elastic earring:
High quality elastic earring, suitable for different face shapes, difficult to slide, allows you to take or remove easily and quickly.



Integrated adjustable metal bridge band Integrated adjustable metal
bridge bridge, simply press on it to adapt gently to the bridge of the nose to better seal your face.

Decrease gently:
Technology fabric, anti-bacteria and breathable, KN95 facial mask is comfortable to wear.

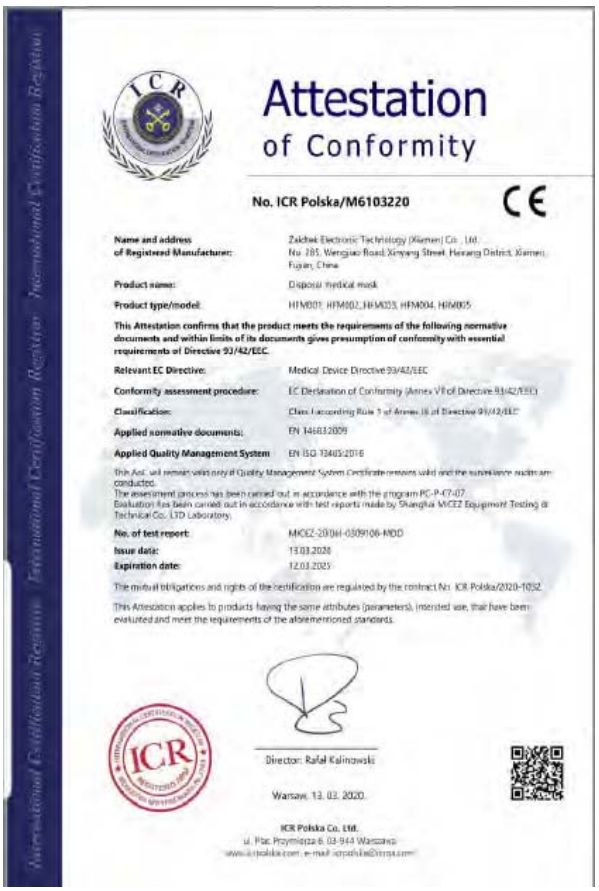
Wide application:
Suitable for personal care, laboratories, manufacturing industry, service industry and UV, etc.

Specification

Product weight: 0.0600 kg
Package:

Or

CE & FDA Mark Certification



Attestation of Conformity

No. ICR Polska/M6103220 CE

Name and address of Registered Manufacturer:	Zaidtek Electronic Technology (Xiamen) Co., Ltd. No. 285, Wengjiao Road, Xinyang Street, Haicang District, Xiamen, Fujian, China
Product name:	Disposal medical mask
Product type/model:	HFM001, HFM002, HFM003, HFM004, HFM005
Relevant EC Directive:	Medical Device Directive 93/42/EEC
Conformity assessment procedure:	EC Declaration of Conformity (Annex VII of Directive 93/42/EEC)
Classification:	Class I according Rule 1 of Annex IX of Directive 93/42/EEC
Applied normative documents:	EN 14683:2009
Applied Quality Management System:	EN ISO 13485:2016
No. of test report:	MCEZ-2020H-0309106-MDD
Issue date:	13.03.2020
Expiration date:	12.03.2025

This Attestation confirms that the product meets the requirements of the following normative documents and within limits of its documents gives presumption of conformity with essential requirements of Directive 93/42/EEC.

This AoC will remain valid only if Quality Management System Certificate remains valid and the surveillance audits are conducted.
The assessment process has been carried out in accordance with the program PC-P-47-07.
Evaluation has been carried out in accordance with test reports made by Shanghai MCEZ Equipment Testing & Technical Co., LTD Laboratory.

The mutual obligations and rights of the certification are regulated by the contract No. ICR Polska/2020-1052.

This Attestation applies to products having the same attributes (parameters), intended use, that have been evaluated and meet the requirements of the aforementioned standards.

Director: Rafał Kalinowski

Warsaw, 13. 03. 2020.

ICR Polska Co. Ltd.
ul. Plac Przymierza 8, 03-944 Warszawa
www.icrpolska.com, e-mail: icrpolska@icrpol.com

Fiscal Year 2020

CERTIFICATION OF REGISTRATION

This certifies that:

Zaidtek Electronic Technology (Xiamen) Co., Ltd.
No. 285, Wengjiao Road, Xinyang Street, Haicang District, Xiamen, Fujian, China

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

Shanghai JAT Enterprise Management Consultation Co., Ltd.

Owner Operator Number: 10063956

Device Listing#:

Listing No	Code	Device Name	Proprietary Name
D677450	KHA	MASK	Disposal medical mask HFM001, HFM002, HFM003, HFM004, HFM005

Device Listing#: See annex

JAT 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. JAT 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. JAT assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "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, JAT is not affiliated with the U.S. Food and Drug Administration.

JAT FDA

Chief engineer
Issued: 03/23/2020
Expiration Date:12/31/2020

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FOR MORE INFORMATION: contact@c4icommunication.com or +1 (302).981.1340