



Attestation of Conformity

No. ICR Polska/



**Name and address
of Registered Manufacturer:**

Jiangxi yue 'an medical equipment co., LTD
3rd floor, no.15 workshop, wudu project area, industrial park,
xiushui county, jiujiang city, jiangxi province

Product name:

Respirator

Product type/model:

YA-KN95

Trade mark:

n/a

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.

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 14683:2005

**Applied Quality
Management System**

n/a

This AoC will remain valid only if Quality Management System Certificate remains valid.
The assessment process has been carried out in accordance with the program PC-P-07-07.
Evaluation has been carried out in accordance with test report made by:

- CMAT Testing and Certification Laboratories (Hua nan) Limited

No. of test report:

CMAT20200323050MDD

Issue date:

19.03.2020

Expiration date:

18.03.2025

The mutual obligations and rights of the certification are regulated by the contract
No. ICR Polska/2020-3125.

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



Director: Rafał Kalinowski

Warsaw, 19. 03. 2020.

ICR Polska Co. Ltd.

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