

To: [5.1.2e]@nfu.nl [5.1.2e]@nfu.nl; [5.1.2e]@pinnedbyk.com [5.1.2e]@pinnedbyk.com]
From: coronabeschermingsmiddelen
Sent: Mon 3/23/2020 2:50:54 PM
Subject: FW: FFP 2 en FFP3 masks N95 + CE classificatie (voldoende voorraad)
Received: Mon 3/23/2020 2:50:54 PM
[image001.jpg](#)
[MEDICAL CE PDA N95-1.pdf](#)
[MEDICAL CE PDA N95.pdf](#)

Beste heer,

Nogmaals dank voor uw bericht.

De inkoop van beschermingsmiddelen wordt sinds kort centraal gecoördineerd. Daarom stuur ik met deze e-mail uw bericht door naar dit Landelijk Consortium Hulpmiddelen.

Graag wil ik u vragen om uw verdere correspondentie te richten aan dit consortium. Het mailadres is: [5.1.2e]@nfu.nl

Ik wil u hartelijk danken voor het meedenken.

Met vriendelijke groet,



Directie Geneesmiddelen en Medische Technologie

Team Corona

Ministerie van Volksgezondheid, Welzijn en Sport

Parnassusplein 5 | 2511 VX | Den Haag

Postbus 20350 | 2500 EJ | Den Haag |

Van: [5.1.2e] | Pinned by K <[5.1.2e]@pinnedbyk.com>
Verzonden: maandag 23 maart 2020 09:12
Aan: coronabeschermingsmiddelen <coronabeschermingsmiddelen@minvws.nl>
Onderwerp: FFP 2 en FFP3 masks N95 + CE classificatie (voldoende voorraad)

Geachte heer, mevrouw,

Onze fabrikant in China heeft op dit moment voldoende voorraad inzake mondkapjes met N95 classificatie en er worden elke dag [5.1.1c] stuks geproduceerd aangezien de vraag hoog is zijn de mondkapjes ook weer snel uitverkocht.

- *levertermijn betreft circa 9 dagen*

- *certificering zie bijlage*

- *prijs FFP 2 betreft [5.1.1c] (grotere afnames zal de prijs dalen)*

- *prijs FFP 3 betreft [5.1.1c] (grotere afnames zal de prijs dalen)*

- de huidige voorraad betreft op dit moment FFP2 5.1.1c stuks en FFP3 5.1.1c stuks

Ook levert onze fabrikant andere producten die men nu nodig heeft zie bijlage hiervoor kan ik een prijs aanvragen mocht dit nodig zijn.

Mochten er verder vragen zijn dan verneem ik deze graag.

Kind Regards,

5.1.2e

Head Office
Pinned by K & Overdose the Label
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by
K

Member of:





Attestation of Conformity

No. ICR Polska/M [REDACTED]



**Name and address
of Registered Manufacturer:** [REDACTED]

Product name: Disposable medical face mask

Product type/model: A1, A2, A3

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:

- Shenzhen MONLKA Technology Co., Ltd

No. of test report: MNK20200311043R

Issue date: 16.03.2020

Expiration date: 15.03.2025

The mutual obligations and rights of the certification are regulated by the contract No. ICR Polska/2020-2040.

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: Rafal Kalinowski

Warsaw, 16. 03. 2020.

ICR Polska Co. Ltd.

ul. Plac Przymierza 6, 03-944 Warszawa
www.icrpolska.com, e-mail: icrpolska@icrqa.com



Attestation of Conformity

No. ICR Polska/M [REDACTED]



Name and address
of Registered Manufacturer:

[REDACTED]

Product name: Disposable medical face mask

Product type/model: A1, A2, A3

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:

- Shenzhen MONLKA Technology Co., Ltd

No. of test report: MNK20200311043R

Issue date: 16.03.2020

Expiration date: 15.03.2025

The mutual obligations and rights of the certification are regulated by the contract No. ICR Polska/2020-2040.

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, 16. 03. 2020.

ICR Polska Co. Ltd.

ul. Plac Przymierza 6, 03-944 Warszawa
www.icrpolska.com, e-mail: icrpolska@icrqa.com





Certificate

No. ICR Polska/P [REDACTED]

Name and address of certificate owner:



Name and address of manufacturer:



Product name: Disposable particulate respirator

Product types: FFP3, N95

This certificate confirms that the product meets the requirements of the following standards and within limits of its standards gives presumption of conformity with essential requirements of Regulation 2016/425

EN 149:2001+A1:2009

The certification process has been carried out in accordance with the program PC-P-07-07.

Evaluation has been carried out in accordance with test reports made by Shenzhen M Technology Co.,Ltd.

No. of test reports: MNK20200305023R

Certificate issue date: 16.03.2020

Expiration date: 15.03.2025

The mutual obligations and rights of the certification are regulated by the contract No. Polska/2020-3014.

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



Certificate

No. ICR Polska/P [REDACTED]

Name and address of certificate owner:



Name and address of manufacturer:



Product name: Disposable particulate respirator

Product types: FFP3, N95

This certificate confirms that the product meets the requirements of the following standards and within limits of its standards gives presumption of conformity with essential requirements of Regulation 2016/425

EN 149:2001+A1:2009

The certification process has been carried out in accordance with the program PC-P-07-07.

Evaluation has been carried out in accordance with test reports made by Shenzhen M Technology Co.,Ltd.

No. of test reports: MNK20200305023R

Certificate issue date: 16.03.2020

Expiration date: 15.03.2025

The mutual obligations and rights of the certification are regulated by the contract No. Polska/2020-3014.

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

防护服选抗呗奥丰生产的防护服

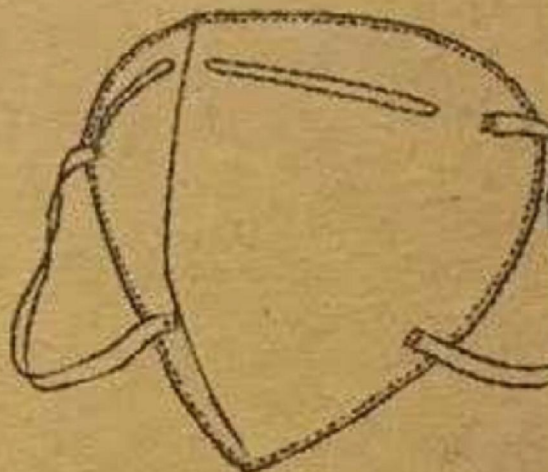
「百」海固防护服 型号齐全 现货供应「家现..

「百」防护服选恒泰,恒泰防护



Particulate respirator N95/ FFP3

- Filters bacteria and dust 95%
- Easy to carry flat fold design
- Hygienic and skin friendly
- Suitable for use in a sterile environment and throw



Use for:
 • Used in medical, dental, laboratory, household, industry, etc.
 • Effectively filter out bacteria and dust 95%
 Caution and instructions

1. This respirator does not supply oxygen, not for use in atmospheres containing less than 19.5% oxygen.
2. Not for use in atmospheres immediately dangerous to life or health.
3. Do not use for vapors, oil aerosols, asbestos, arsenic, cadmium, lead, H₂S, hydrogen cyanide, etc.
4. Do not exceed maximum use concentration as established by regulatory standards.
5. Failure to properly use and maintain this product could result in injury or death.
6. Never substitute, modify, add, or omit parts.



How to wear

1. Be sure the part is facing at the side of your nose after it is facing the face, and the seal is facing the face.
2. Press the seal against the side of the face, and pull the bottom back to the ear and hold it on the side of the face.
3. Press the hinges of both bands in middle of the nose strip and press upward according to the contour of the nose bridge.
4. Be sure both on your face when you wear, and check the tightness with your face.

5pcs

FDA CE
EN 14683:2005

STERILE EO

Made in China



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