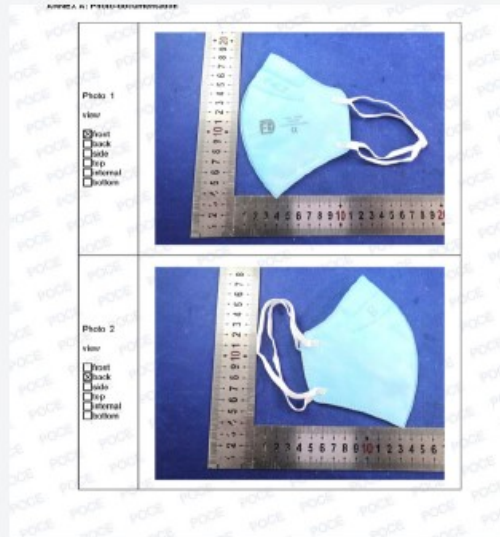


# PPE: FDA APPROVED KN95 MASK



KN95 masks go through the same testing for FFP2 as their N95 equivalent in the US. The FDA has cleared these masks as a substitute for N95s due to shortages and price issues. The testing for this mask was completed in China and passed compliance and CE certification. The factory has also been vetted thoroughly by our contacts overseas. There is no FDA cert, but this product is a viable option for anyone looking for shorter lead times and a reduced cost but the same quality of an N95.

## Product Specs

- 1,000 Minimum Order (Boxes of 840)
- Lead Time 5-7 days from PO
- Cost: \$4.75 each
- One Size Fits All , Med/Large (4.8")
- 100% Payment Required at Time of Order



440-343-3362

[jartiste@americanbus.com](mailto:jartiste@americanbus.com)

TEST REPORT	
EN 149	
Respiratory protective devices. Filtering half masks to protect against particles. Requirements, testing, marking	
Report Reference No.	POCE/2021/0000495
Completed by (signature)	Jiah Jiao
Approved by (signature)	Chen Chen
Date of Issue	Mar 10, 2021
Issuing Laboratory	SHENZHEN POCE TECHNOLOGY CO., LTD. H Building, Hengde Science and Technology Park, Tanggu, Shijiazhuang District, Shijiazhuang, China
Applicant's name	Shenzhen Dada Medical Technology Co., Ltd.
Address	Oct. 10th Floor, Building D1, Hengde Industrial Park, Shuang Community, Shijiazhuang City, Shijiazhuang District, Shijiazhuang, Shijiazhuang, China
Test specification	EN 149:2001+A1:2003
Non-standard test method	NA
Test item description	Particulate Filter
Trade Mark	POCE
Model/Type reference	EN149:2001, EN149:2001+A1:2003, EN149:2001+A1:2003, EN149:2001+A1:2003
Manufacturer	Shenzhen Dada Medical Technology Co., Ltd.
Address	Oct. 10th Floor, Building D1, Hengde Industrial Park, Shuang Community, Shijiazhuang City, Shijiazhuang District, Shijiazhuang, Shijiazhuang, China
Classification	FFP2 NR

Attestation of Conformity	
No. ICR Polska/MI002520	
CE	
Name and address of Registered Manufacturer	Shenzhen Dada Medical Technology Co., Ltd.
Product name	FFP2 NR
This Attestation certifies that the product meets the requirements of the following normative documents and within the scope of the manufacturer's responsibility with essential requirements of Directive 93/42/EEC	EN 149:2001+A1:2003
Reference to the standard	EN 149:2001+A1:2003
Conformity assessment procedure	Class I according to Rule 1 of Annex II of Directive 93/42/EEC
Classification	Class I according to Rule 1 of Annex II of Directive 93/42/EEC
Applied normative documents	EN 149:2001+A1:2003
Applied Quality Management System	EN ISO 9001:2015
This Attestation of Conformity will remain valid only if the Quality Management System Certificate remains valid and the surveillance audits are successful.	
The extension of validity has been carried out in accordance with the procedure ICR P-07-02.	
Validation has been carried out in accordance with the procedure ICR P-07-02.	
No. of test reports	1
Issue date	10.03.2021
Expiry date	10.03.2022
The issuer's obligations and rights of the certificate are regulated by the contract No. ICR P-07-02-001.	
This Attestation applies to products having the same technical specifications, intended use, the same design and construction as the product described in the contract.	
Signature	[Signature]
Director	Rafał Kucharski
Stamp	ICR Polska