

NPPTL COVID-19 Response: International Respirator Assessment

Manufacturer: UNKNOWN

Model Tested: UNKNOWN

Date Tested: April 16, 2020

These findings pertain to the respirator submitted for testing. The packaging and respirator have no markings. The manufacturer and model are unknown. The form submitted indicates that this product is a KN95. The form also indicates the respirators were obtained from K & K Supply.

Ten respirators were submitted for evaluation. The samples were tested using a modified version of NIOSH Standard Test Procedure (STP) TEB-APR-STP-0059. This modified assessment plan can be found [here](#).

No certificate of approval was provided with the samples received; therefore, the authenticity of the claims cannot be validated.

The maximum and minimum filter efficiency was 99.60% and 98.14%, respectively. All ten respirators measured more than 95%.

While the above-listed product classification has similar performance requirements to NIOSH-approved devices, NIOSH does not have knowledge about the sustained manufacturer quality system and product quality control for these products. NIOSH also does not have knowledge about the product's handling and exposures after leaving its manufacturer's control.

In addition, this product is an ear loop design. Currently, there are no NIOSH-approved products with ear loops; NIOSH-approved N95s have head bands. Furthermore, limited assessment of ear loop designs, indicate difficulty achieving a proper fit. While filter efficiency shows how well the filter media performs, users must ensure a proper fit is achieved.

This assessment is not a part of the NIOSH respirator approval process and will in no way lead to or preclude NIOSH approval through the official approval process. This assessment was developed as an assessment of the filter efficiency for those respirator's represented as certified by an international certification authority, other than NIOSH, to support the availability of respiratory protection to US healthcare workers due to the respirator shortage associated with COVID-19. Only particulate filter efficiency was assessed.

The results provided in this letter are specific to the subset of samples that were provided to NPPTL for evaluation.

These results will be used to update the CDC guidance for [Crisis Capacity Strategies \(during known shortages\)](#).

Evaluation of International Respirators

Test: Modified TEB-APR-STP-0059

Date Tested: April 16, 2020

Report Prepared: April 17, 2020

Manufacturer: UNKNOWN

Item Tested: UNKNOWN

Country of Certification: China (GB2626-2006)

Pictures have been added to the end of this report.

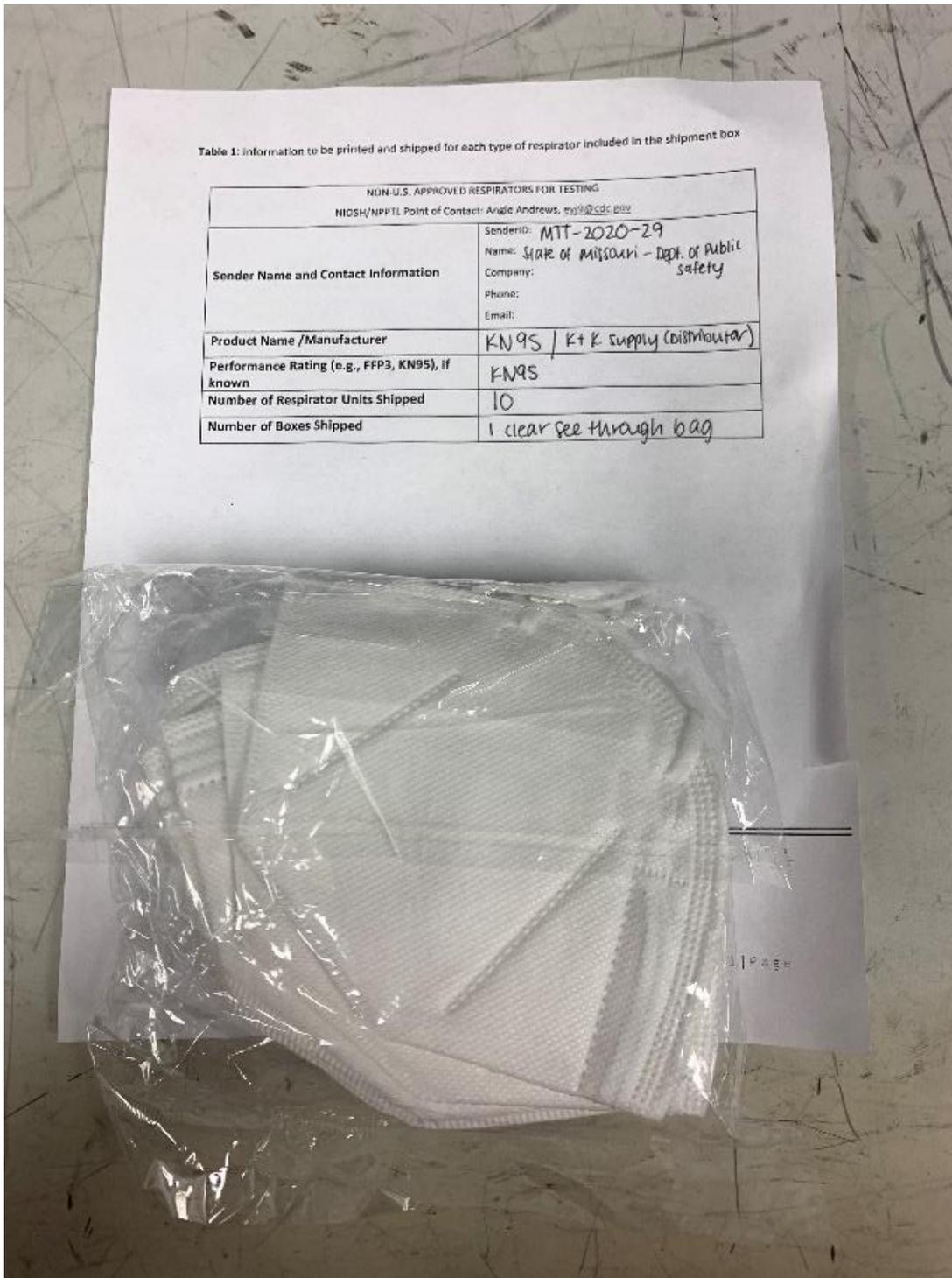
Filter	Flow Rate (Lpm)	Initial Filter Resistance (mmH ₂ O)	Initial Percent Leakage (%)	Maximum Percent Leakage (%)	Filter Efficiency
1	85	11.0	1.860	1.860	98.14
2	85	13.9	0.464	0.464	99.54
3	85	13.4	0.401	0.401	99.60
4	85	10.9	0.840	0.854	99.15
5	85	10.5	0.498	0.524	99.48
6	85	13.2	0.485	0.485	99.52
7	85	12.6	0.559	0.559	99.44
8	85	9.5	0.931	0.975	99.03
9	85	13.6	0.430	0.430	99.57
10	85	9.8	1.020	1.110	98.89
Minimum Filter Efficiency: 98.14			Maximum Filter Efficiency: 99.60		

- The test method utilized in this assessment is not the NIOSH standard test procedure that is used for certification of respirators. Respirators assessed to this modified test plan do not meet the requirements of STP-0059, and therefore cannot be considered equivalent to N95 respirators that were tested to STP-0059.
- Respirators tested may not be representative of all respirators with the same certification mark. NIOSH has no control over suppliers and distributors of respirators certified by other national or international parties.
- This assessment is not a confirmation that it conforms with any or all of its specifications in accordance with its certification mark.
- This assessment was not a part of the NIOSH approval program. These results do not imply nor preclude a future approval through the NIOSH respirator approval program.

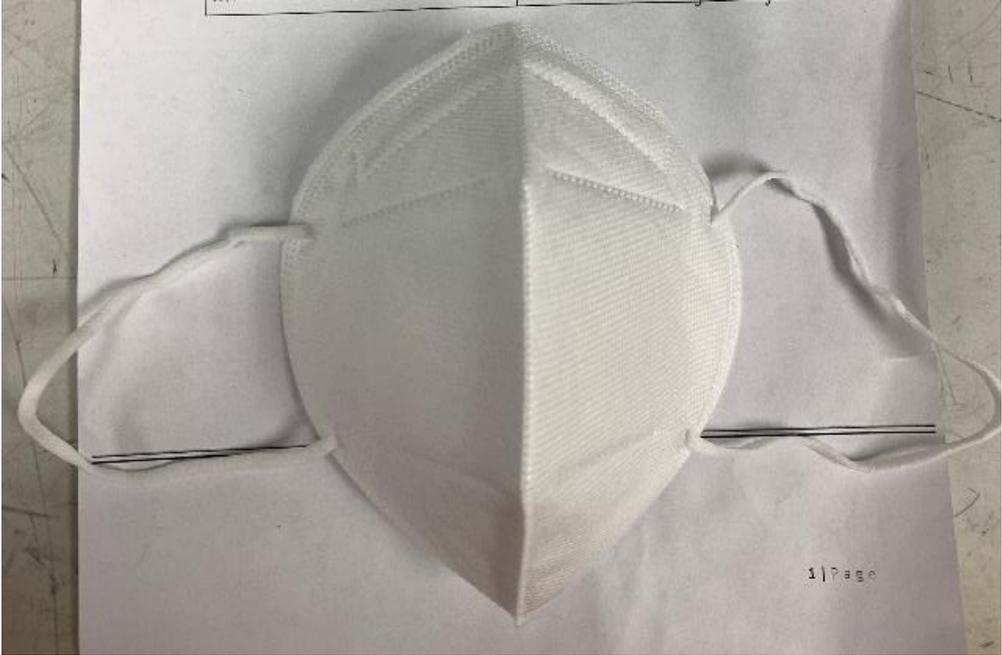
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Table 1: Information to be printed and shipped for each type of respirator included in the shipment box

NON-U.S. APPROVED RESPIRATORS FOR TESTING	
NIOSH/NPPTL Point of Contact: Avigle Andrews, avi@cdc.gov	
Sender Name and Contact Information	Sender ID: MIT-2020-29 Name: STATE of Missouri - Dept. of Public Safety Company: Phone: Email:
Product Name /Manufacturer	KN95 / K+K Supply (dismount)
Performance Rating (e.g., FFP3, KN95), if known	KN95
Number of Respirator Units Shipped	10
Number of Boxes Shipped	1 clear see through bag



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