

NPPTL COVID-19 Response: International Respirator Assessment

Manufacturer: Guangdong Golden Leaves Technology Development Co., Ltd.

Model Tested: 8862 KN95

Date Tested: May 27, 2020

These findings pertain to the Guangdong Golden Leaves Technology Development Co., Ltd., model 8862 KN95. The packaging and labeling for this product indicate that it meets GB2626-2006 (the Chinese standard for Respiratory Protective Equipment – Non-Powered Air-Purifying Particle Respirator).

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 99.50% and 99.00%, 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: May 27, 2020

Report Prepared: May 27, 2020

Manufacturer: Guangdong Golden Leaves Technology Development Co., Ltd.

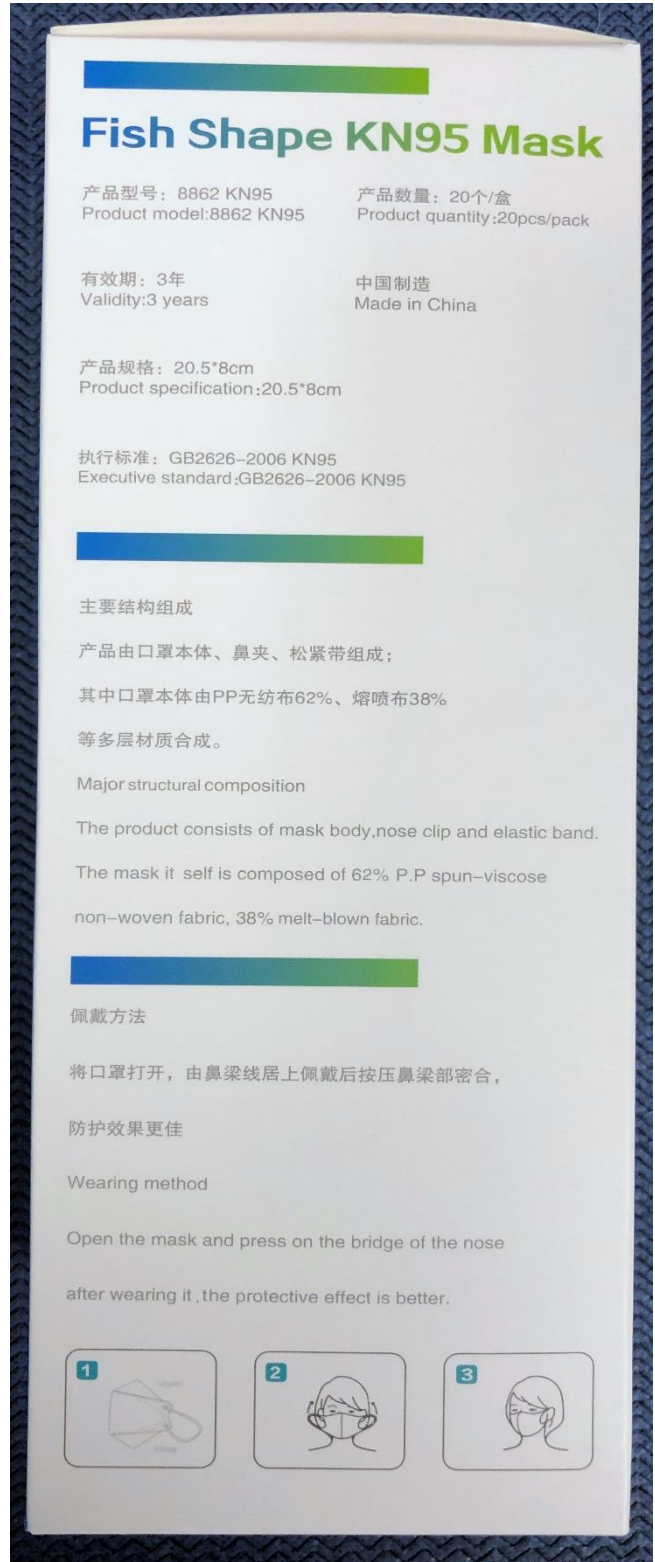
Item Tested: 8862 KN95

Country of Certification: China (GB2626-2006)

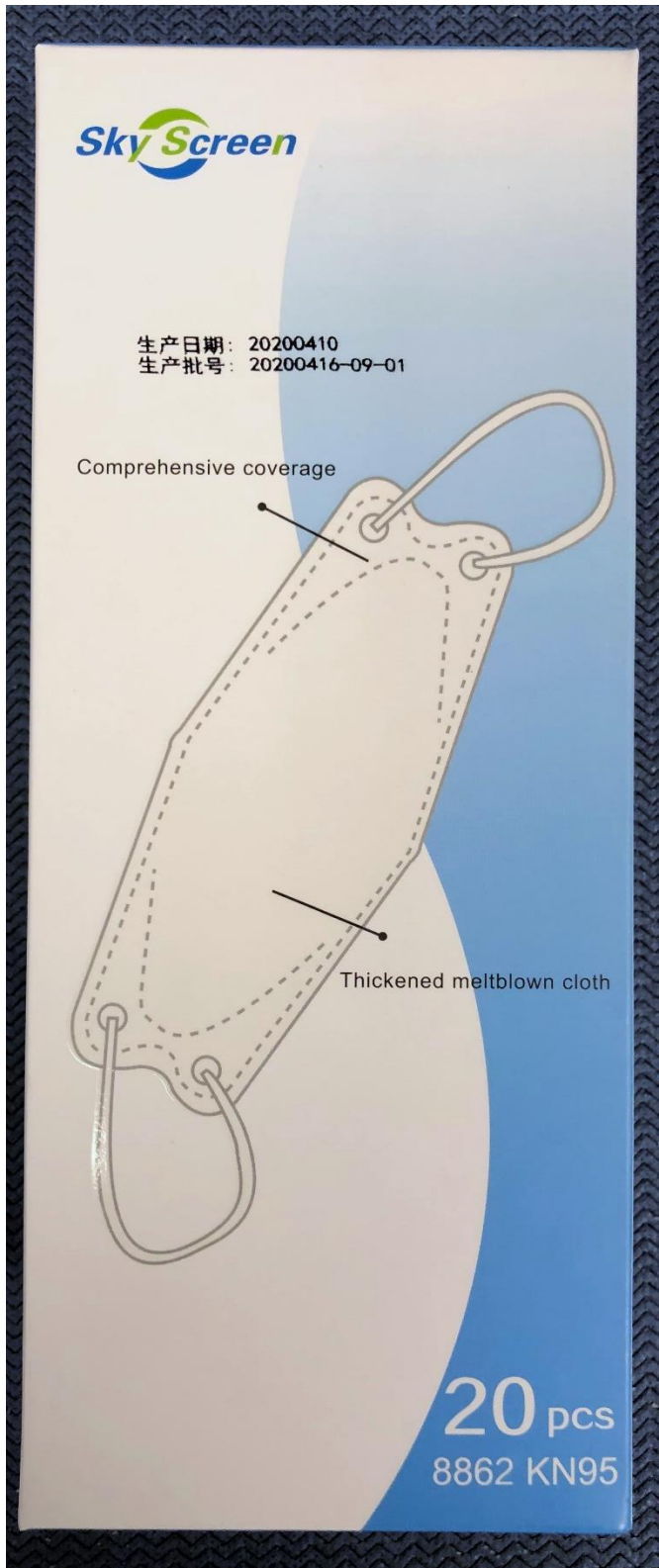
Pictures have been added to the end of this report.

Filter	Flow Rate (Lpm)	Initial Filter Resistance (mmH ₂ O)	Initial Percent Leakage (%)	Maximum Percent Leakage (%)	Filter Efficiency
1	85	10.1	0.69	0.69	99.31
2	85	10.3	0.60	0.60	99.40
3	85	9.8	0.60	0.60	99.40
4	85	10.8	0.63	0.63	99.37
5	85	10.1	0.78	0.78	99.22
6	85	10.3	0.73	0.73	99.27
7	85	10.6	0.50	0.50	99.50
8	85	12.7	1.00	1.00	99.00
9	85	10.2	0.72	0.72	99.28
10	85	10.2	0.81	0.81	99.19
Minimum Filter Efficiency: 99.00			Maximum Filter Efficiency: 99.50		

- 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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1. 不要在50℃以上的环境中
2. 不要在含氧量低于20%的环境下
3. 不要在有毒气体环境中使
4. 小孩, 孕妇以及老人不建
5. 如果包装损坏或者部件不
该口罩仅限于对非油性颗粒
6. 建议储存在干燥、通风、
储存温度在-20℃-30℃之
远离火源和易燃物品。
7. 本产品不适用于缺氧环
特殊行业或婴幼儿呼吸保
8. 不要在不通风、呼吸困
9. 心肺功能不正常者应慎
10. 如果口罩已经损坏或者
11. 如在使用过程中有任何
12. 不可重复使用, 口罩是
**请在使用前仔细阅读。
Cautions and Limitations
A. Do not use in an environm
B. Do not use in an environm
C. Do not use in a toxic gas
D. Not recommended for ch
E. If the package is damag
it is forbidden to use the m
It is limited to respiratory p
F. It is recommended to s
less than 80% and storage
-20℃ and 30℃. Keep awa
flammable materials.
G. This product is not app
as anoxic environment, un
and industrial dust protecti
H. Do not wear a mask if u
or during sleep.
I. People with abnormal h
masks with caution.
J. If a mask is damaged or
replace it in time.
K. In case of any discomf
stop to use immediately.
L. Do not reuse. The mask
**Pls read carefully befor
*注意使用前, 使用者必须
请保存这些说明以便参考
生产地址: 广东省东莞市
公司名称: 广东金叶科技
电话: 0769-87989800
*Pay attention before use,
Please save these intructio
Manufacturer:Guangdong G
Development Co.,Ltd
Production Address:5/F, b
Tangxia Town,Dongguan,
Contact:0769-87989800

NON STERILE

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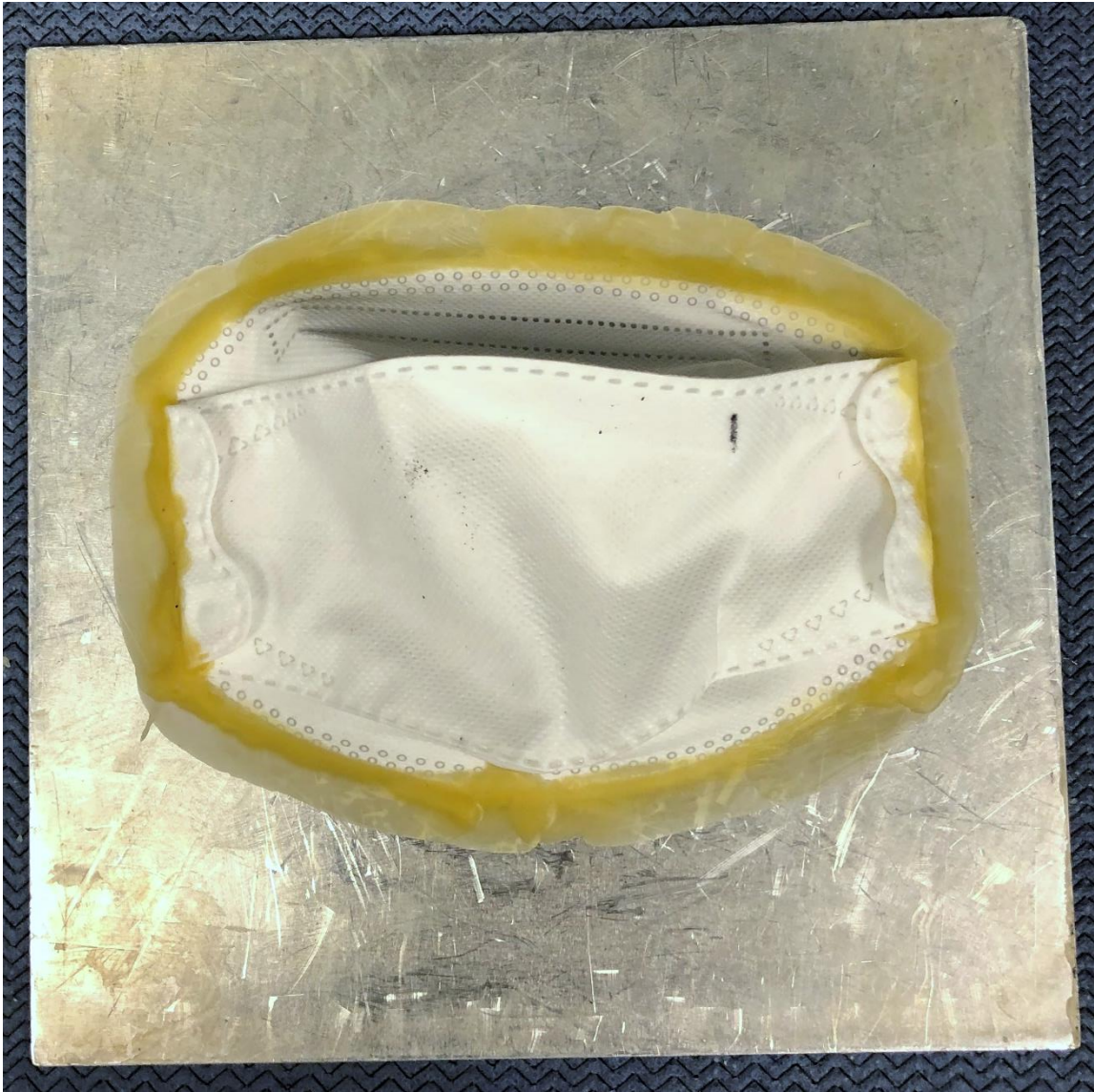
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