



NIOSH Reference: TN-23625 Mfr. Reference: SFI1058/1059a Centers for Disease Control and Prevention (CDC)

National Institute for Occupational Safety and Health (NIOSH) National Personal Protective Technology Laboratory (NPPTL) 626 Cochrans Mill Road

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April 9, 2020

Tang Yingzhong Suzhou Fangtian Industries Co., Ltd. Xiehe Building, No. 158 Jinmen Road Suzhou City, Jiangsu CHINA

Dear Tang Yingzhong:

The National Institute for Occupational Safety and Health (NIOSH) has reviewed your request accepted January 8, 2020. This request was for extensions of approvals TC-84A-7863 and TC-84A-7866 to add private label versions of Suzhou Fangtian N95 air-purifying filtering facepiece respirators to ARMOR S.C.C. of Ecuador. The approval numbers, Suzhou Fangtian respirator model numbers, assembly matrix file names and ARMOR S.C.C. private-label model numbers are shown in the following table:

Approval Number Suzhou Fangtian Model	Assembly Matrix File Name	ARMOR S.C.C. Model
TC-84A-7863	FT-N058 AMc.xls	1058
Model: FT-N058	Revision C, Dated: 12-11-2019	
TC-84A-7866	FT-N059 AMe.xls	1059
Model: FT-N059	Revision E, Dated: 12-16-2019	

This request is granted. Extension of approvals are granted only for documentation written in the English language. It is the manufacturer's responsibility to correctly translate materials desired to be produced in languages other than English.

The final respirator approval labels are included as attachments to this letter. The abbreviated labels have been accepted as submitted. The cautions and limitations which apply to these approvals are on the approval label. Only those assemblies affected by this request, or where new approval numbers are assigned, apply to this approval action. Production approval labels cannot include information on unapproved configurations.

The manufacturer is responsible for properly packaging, labeling, and controlling the respirators produced under these private label approvals. At a minimum, the items that must be controlled are the approved user instructions, all approval labeling, all approved packaging, use claims, marketing materials, and the respirator design and construction details. Any change to these

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NIOSH-approved respirators or approval documentation without prior notification and approval is a violation of this approval and renders this certification as invalid.

No changes may be made to any respirators and accompanying documentation without prior written approval of NIOSH. Requests for changes must be submitted to NIOSH and a modification of this approval must be granted before changes are made.

Sincerely,

Jeffrey Peterson Chief, Conformity Verification and Standards Development Branch

Enclosures