DEPARTMENT OF HEALTH & HUMAN SERVICES

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
Pittsburgh, PA 15236-0070
Phone: 412-386-4000
Fax: 412-386-4051

NIOSH Reference: TN-24359
Mfr. Reference: SMP0227PL1

December 4, 2020

[redacted]

Dear [redacted]

The National Institute for Occupational Safety and Health (NIOSH) has reviewed your request accepted October 30, 2020. This request was for extension of approval TC-84A-9252, to private label [redacted] N95 models TN02-27 (medium) and TN02-28 (small) air-purifying filtering-facepiece respirators. The complete respirator configurations are detailed on assembly matrix, file name *TN0227AMb.xls*, revision b, dated: 10/16/2020. The private label companies and private label part numbers are noted below.

1. Unicharm Corporation, of Tokyo, Japan. The Unicharm private label part numbers are SNF-M01 Medium and SNF-S01 Small.
2. [redacted]

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 approval labels are included as attachments to this letter. The abbreviated labels have been accepted as submitted. The cautions and limitations are listed on the approval label or labels. 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 approved assemblies consist of the parts as listed on the approval label and the assembly matrix. Parts are to be marked with the numbers indicated on the approval label in a legible and permanent manner (marking cannot be removed without evidence of its previous presence).

The approval holder 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 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 A. Peterson -S
Digitally signed by Jeffrey A. Peterson -S
Date: 2020.12.04 11:29:22 -05'00'

Jeffrey Peterson
Chief, Conformity Verification and
Standards Development Branch

Enclosures