



NIOSH Reference: TN-25765
Mfr. Reference: LWS904

National Institute for Occupational
Safety and Health (NIOSH)
National Personal Protective
Technology Laboratory (NPPTL)
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May 13, 2022

Peter Maguire
Vice President
Lighthouse Worldwide Solutions
300 W Antelope Road
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Dear Peter Maguire:

The National Institute for Occupational Safety and Health (NIOSH) has reviewed your request accepted March 23, 2022. This request was for Surgical N95 approval of model/part number SHIELD-95-MEDI air-purifying filtering facepiece respirator for protection against particulates at a N95 filter efficiency level. The model SHIELD-95-MEDI is identical to the existing N95 filtering facepiece H210 approved under TC-84A-9376. The complete respirator configuration is detailed on assembly matrix, file name *714252843-1AM17.xls*, revision 17, dated: March 17, 2022.

This request is granted. Approvals are granted only for documentation written in the English language. It is the approval holder's responsibility to correctly translate materials desired to be produced in languages other than English. Approval number **TC-84A-9431** has been assigned. This respirator is approved and cleared for use in healthcare settings under the FDA/NIOSH MOU 225-18-006. This Surgical N95 provides protection against particulates at the N95 filter efficiency level and conforms to recognized standards for flammability, fluid resistance and biocompatibility.

The approval label is included as an attachment to this letter. The abbreviated label has been accepted as submitted. The full and abbreviated approval label must be presented to users as approved by NIOSH, including the cautions and limitations as listed. The approved assembly consists 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).

Production and labeling of the approved respirator configuration can begin following receipt of this letter (the time and date of the approval action are recorded by my electronic signature, below).

This certificate of approval is not an endorsement of the respirator by NIOSH, and such endorsement shall not be stated or implied in advertisements or other publicity. However, you may publicize the fact that this respirator has met the requirements of Title 42, *Code of Federal Regulations*, Part 84 (42 CFR 84). You may market the device, subject to the general controls provisions of the Federal Food, Drug and Cosmetic Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

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