You have been given an N95 respirator that has been decontaminated for multiple-user reuse by healthcare personnel (i.e., healthcare personnel may receive a different respirator following decontamination than the one they had previously used) in a healthcare setting to help prevent healthcare personnel exposure to pathogenic biological airborne particulates during the COVID-19 pandemic.

This Fact Sheet informs you of the significant known and potential risks and benefits of the emergency use of decontaminated, compatible N95 respirators. These compatible N95 respirators have been decontaminated using Stryker Sustainability Solutions’ (SSS) VHP N95 Respirator Decontamination System (hereafter referred to as “decontaminated N95 respirators” and “SSS Decontamination System,” respectively, throughout this Fact Sheet).

Decontaminated N95 respirators that have been decontaminated using the SSS Decontamination System are authorized for multiple-user reuse by healthcare personnel in a healthcare setting during the COVID-19 pandemic.

Whether or not you use a surgical mask, respirator, or face shield, always follow infection control measures: wash hands, cover cough and sneezes; and stay home if you may be sick.

What are the symptoms of COVID-19?

Many patients with confirmed COVID-19 have developed fever and/or symptoms of acute respiratory illness (e.g., cough, difficulty breathing). However, limited information is currently available to characterize the full spectrum of clinical illness associated with COVID-19. Based on what is known about the virus that causes COVID-19, signs and symptoms may appear any time from 2 to 14 days after exposure to the virus.

Public health officials have identified cases of COVID-19 infection throughout the world, including the United States, which may pose risks for public health. Please check the CDC webpage for the most up to date information.

What do I need to know about the emergency use of decontaminated N95 respirators?

- SSS Decontamination System has been authorized for emergency use to decontaminate compatible N95 respirators for multiple-user reuse by healthcare personnel during the COVID-19 pandemic to prevent wearer exposure to pathogenic biological airborne particulates.
- Successful testing on decontaminated N95 respirators demonstrated acceptable performance through 3 decontamination cycles for sporicidal activity, filtration efficiency, breathability, form fit testing, and strap integrity testing.
- Healthcare personnel should remember that respirators only offer safety when they fit properly. Tight-fitting respirators must form a tight seal around the wearer’s face to avoid exposure to hazardous substances.

Preparing compatible N95 respirators for decontamination:
- Remove the compatible N95 respirator per institutional procedures.
- Inspect respirators after each use prior to collection for decontamination.
- Discard if decontaminated three (3) times or if visibly soiled (e.g., blood) or damaged – do not use and do not send for decontamination.
- Place N95 respirators in a designated hamper stand for subsequent decontamination per your healthcare facility’s procedures.

Use of decontaminated N95 respirators:
- Decontaminated N95 respirators are not sterile.
- Discard if decontaminated three (3) times or if visibly soiled (e.g., blood) or damaged.
- Cellulose-based materials are incompatible with the SSS Decontamination System.
- Report problems with decontaminated N95 respirators to your healthcare facility.

Monitor healthcare personnel for signs and symptoms of potential infection with SARS-CoV-2

Report Adverse events, including problems with test performance or results, to MedWatch by submitting the online FDA Form 3500 (https://www.accessdata.fda.gov/scripts/medwatch/index.cfm?action=reporting.home) or by calling 1-800-FDA-1088.
or other respiratory infection for up to and including 14 days after last contact with the SARS-CoV-2 virus and related material, and promptly report such information to your healthcare facility.

- Report damage or discoloration observed upon receipt of the decontaminated N95 respirators, and potential exposure of healthcare personnel from breaks in or other damage to or degradation of the decontaminated N95 respirators to your healthcare facility.

Use appropriate personal protective equipment (PPE) when caring for individuals suspected of having COVID-19 as outlined in the CDC webpages, including *Interim Infection Prevention and Control Recommendations for Patients with Confirmed Coronavirus Disease 2019 (COVID-19) or Persons Under Investigation for COVID-19 in Healthcare Settings, Infection Control, and FAQ about PPE.*

Current information on COVID-19 for healthcare personnel is available at CDC’s webpage, *Information for Healthcare Professionals* (see links provided in “Where can I go for updates and more information” section).

What are the known and potential benefits and risks of using decontaminated N95 respirators?

Potential benefits include:

- May help prevent exposure to airborne pathogens, and therefore reduce the risk of infection or illness
- May extend the usability of compatible N95 respirators by allowing for decontamination and reuse

Potential risks include:

- Failure of filtration efficiency
- Reduced breathability
- Strap failure and ineffective face-fit
- Reused respirators may not have been effectively decontaminated of SARS-CoV-2 or other pathogens
- Cross-contamination from ineffective decontamination

Overview of SSS Decontamination System

The Stryker Sustainability Solutions Decontamination System is a self-contained system that uses vaporized hydrogen peroxide (VHP) for decontamination of compatible N95 respirators that are contaminated or potentially contaminated with SARS-CoV-2 or other pathogenic microorganisms. N95 respirators containing cellulose-based materials are incompatible with the SSS decontamination process.

Each decontamination cycle in the SSS Decontamination System consists of injecting VHP into the decontamination chamber until achieving a saturated atmosphere; maintaining the VHP exposure for a 120-minute dwell time; and allowing the VHP to off-gas to a level of 1 ppm prior to post decontamination processing. A minimum of six calibrated chemical indicators are dispersed throughout the system to indicate a successful decontamination cycle. This decontamination system enables the reuse of compatible N95 respirators that would otherwise be disposed of after a single use. However, respirators that are visibly soiled must be discarded and not reused or decontaminated.

What is an EUA?

The United States FDA has made the emergency use of SSS Decontamination System to decontaminate compatible N95 respirators available under an emergency access mechanism called an Emergency Use Authorization (EUA). The EUA is supported by the Secretary of Health and Human Service’s (HHS’s) declaration that circumstances exist to justify the emergency use of medical devices, including alternative products used as medical devices, due to insufficient supply during the COVID-19 pandemic.

The SSS Decontamination System has been made available under an EUA and has not undergone the same type of review as an FDA-approved or cleared device. FDA may issue an EUA when certain criteria are met, which includes that there are no adequate, approved, available alternatives, and based on the totality of scientific evidence available, it is reasonable to believe that the SSS Decontamination System may be effective at preventing healthcare personnel exposure to
FACT SHEET FOR HEALTHCARE PERSONNEL

Stryker Sustainability Solutions Decontamination System for Decontaminating Compatible N95 Respirators

May 27, 2020

pathogenic biological airborne particulates during periods of insufficient respirator supply during the COVID-19 pandemic by decontaminating, for a maximum of 3 decontamination cycles per respirator, compatible N95 respirators that are contaminated or potentially contaminated with SARS-CoV-2 or other pathogenic microorganisms.

The EUA for the SSS Decontamination System is in effect for the duration of the COVID-19 declaration justifying emergency use of medical devices, unless terminated or revoked (after which the products may no longer be used).

Where can I go for updates and more information?

CDC websites:
General: https://www.cdc.gov/COVID19

FDA websites:
General: www.fda.gov/novelcoronavirus
EUAs: https://www.fda.gov/medical-devices/emergencysituations-medical-devices/emergency-use-authorizations

Report Adverse events, including problems with test performance or results, to MedWatch by submitting the online FDA Form 3500 (https://www.accessdata.fda.gov/scripts/medwatch/index.cfm?action=reporting.home) or by calling 1-800-FDA-1088