

www.cambridge.org/dmp

Oren Mayer PhD^{1,2}, Tiffany Pfundt PharmD³, Gamola Z. Fortenberry PhD⁴, Brian H. Harcourt PhD¹ and William A. Bower MD¹

Report From the Field

Cite this article: Mayer O, Pfundt T, Fortenberry GZ, Harcourt BH, Bower WA (2022) Lessons learned from a COVID-19 biohazard spill during swabbing at a quarantine facility. *Disaster Med Public Health Prep* **16**: 1279–1281. doi: <https://doi.org/10.1017/dmp.2020.436>.

First published online: 5 November 2020

Keywords:

SARS-CoV-2; COVID-19; cruise ships; quarantine facility; biohazard mitigation; and biocontainment

Corresponding author:

Oren Mayer,
Email: etx9@cdc.gov.

¹Centers for Disease Control and Prevention, Atlanta, Georgia, USA; ²Laboratory Leadership Service assigned to NCEZID, Centers for Disease Control and Prevention, Atlanta, Georgia, USA; ³Food and Drug Administration, Silver Spring, Maryland, USA and ⁴US Department of Agriculture, Food Safety and Inspection Service, Washington DC

Abstract

The need for increased testing for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the virus that causes coronavirus disease 2019 (COVID-19), has resulted in an increase of testing facilities outside of traditional clinical settings and sample handling by individuals without appropriate biohazard and biocontainment training. During the repatriation and quarantine of passengers from the *Grand Princess* cruise ship at a US military base, biocontainment of a potentially infectious sample from a passenger was compromised. This study describes the steps taken to contain the spill, decontaminate the area, and discusses the needs for adequate training in a biohazard response.

Adequate testing capacity is fundamental to controlling the coronavirus disease 2019 (COVID-19) pandemic.¹ Demand for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) sampling in nontraditional healthcare settings presents unique challenges for the safety of staff collecting specimens.² To meet this demand, the scope of those performing sampling activities continues to expand to individuals without appropriate background in biohazard mitigation or biosafety training.^{3–6} Without proper training, this intersection of field response, infection prevention and control, and worker safety serves as a weak point to both the safety and quality of the response. An example of this occurred during sample collection from quarantined individuals from the *Grand Princess* cruise ship at a US military base. The resulting biohazard incident drove deeper discussion into proper sample handling, broader implications for improving the safety and practices of those working with COVID-19 samples in the field, and the importance of including those with biohazard and biocontainment training in these situations.

Narrative

Grand Princess passengers arrived for quarantine at the base from March 9–12, 2020. Testing for SARS-CoV-2 was offered to passengers regardless of presence of symptoms. In total, 383 of 873 passengers agreed to provide nasopharyngeal swab (NP) specimens. A quality management system (QMS) was instituted to ensure samples were handled with an emphasis on safety and sample integrity.

As part of the QMS, all samples were triple-contained; NP specimens were placed in a sealed sample tube (primary container, PC) inside a biocontainment bag (secondary container, SC) before being wrapped with other SCs inside an absorbent waterproof under pad (AWU) inside a 95-kPa sealed bag (tertiary container, TC). SCs were placed in an AWU-lined sample cooler with ice packs for transport to the shipment sample processing center (SSPC). The SSPC was a converted hotel kitchenette consisting of a table for data entry, sample processing space on the floor, 4 minifridges to store samples at 4°C until shipment, and a connected living room serving as the staff resting and staging area. In the SSPC, the sample cooler was handed off to a processing officer who performed quality control (QC). QC consisted of (1) removing each sample bag from the cooler; (2) inspecting sample tubes for container integrity, proper labeling, and accuracy of identifying information; and (3) placing each sample with other checked samples in the TC. The only personal protective equipment (PPE) used when performing QC was nitrile gloves. In this case, the processing officer was a non-Centers for Disease Control and Prevention (CDC) responder with basic PPE training and no background in sample biocontainment and biosafety. A CDC Laboratory Subject Matter Expert (LSME) was on-site overseeing sample processing and PPE usage.

During QC, the processing officer removed a sample bag from the cooler, stood, and saw a drop of colored liquid fall into the cooler while noting that approximately 0.5 mL of viral transport media was inside the SC. The processing officer immediately placed the SC containing the spill back into the cooler, and without moving, notified the LSME. The LSME blocked entry into the area and had the processing officer remain in place while checking the officer's

© Society for Disaster Medicine and Public Health, Inc. 2020. This is an Open Access article, distributed under the terms of the Creative Commons Attribution licence (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted re-use, distribution, and reproduction in any medium, provided the original work is properly cited.

SDMPH
SOCIETY FOR DISASTER MEDICINE & PUBLIC HEALTH

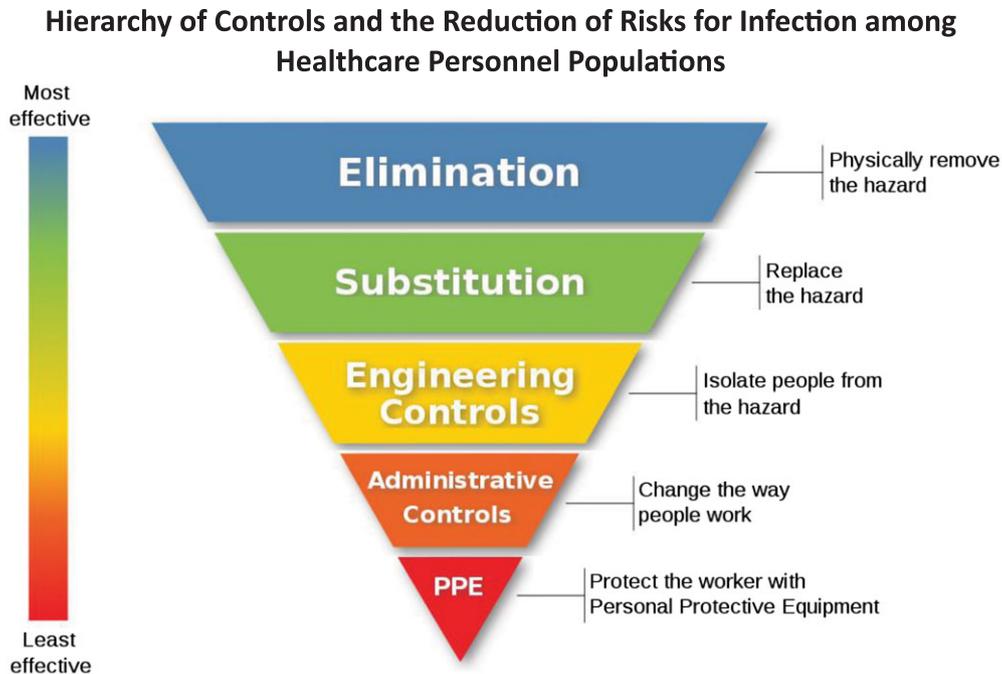


Figure 1. Hierarchy of controls and the reduction of risks for infection among healthcare personnel populations. Each level of the pyramid is associated with processes or functions that directly or indirectly protect staff from exposure or infection by infectious agents. When applied to sample acquisition for disease diagnostics (such as swabbing for SARS-CoV-2), while elimination or substitution strategies are most effective at protecting health care workers, neither are possible in these scenarios so maximizing engineering controls, administrative controls, and PPE must be emphasized.

clothing, shoes, and surrounding floor area for indication of spillage. Confirming no visible spillage, the processing officer removed their gloves and placed them into the contaminated cooler, washed their hands, and secured the area by physically blocking entry. The LSME contacted the CDC Infection Control Team Lead (ICTL) to discuss exposure risk and cleanup steps. By phone, the LSME and ICTL performed a risk assessment considering the location and volume of spill, staff present, and decontamination capability. The following cleanup plan was immediately implemented:

1. The LSME asked all nonessential personnel to leave the SSPC. Two team members volunteered to remain to assist the LSME.
2. The LSME and both assistants donned in order N95 respirators, face shields, gowns, and double gloved with nitrile gloves.
3. Cleanup roles:
 - a. The LSME decontaminated the potentially contaminated items and work areas. All decontamination was performed by wiping with Sani-cloth bleach disinfecting wipes⁷ (PDI, NJ).
 - b. The assistants stood directly on the other side of a half-wall separating the SSPC and staff staging area, handing items to the LSME and receiving waste for disposal.
4. The LSME removed the SCs from the larger TC and placed them back into the cooler, then decontaminated and discarded the used TC. The LSME then removed individual SCs for integrity inspection, decontamination, and placement on a clean AWU. When the suspect bag was removed, the LSME noted the PC appeared closed with the lid attached. However, the SC was not properly sealed, facilitating the external leak. The LSME placed the compromised tube and bag into a new sealable bag and discarded them per CDC guidelines.
5. In this same manner, the LSME removed, decontaminated, and placed all the contents of the cooler into a clean area for

subsequent processing. No other samples leaked, but several had open SCs, which were sealed before placement in the clean area.

6. The LSME decontaminated the inside and outside of the cooler and moved it to the clean area.
7. The LSME decontaminated the working area.
8. The LSME and the 2 assistants doffed PPE in order gloves, gowns, face shields, and then N95 respirators and disposed of as per CDC guidelines.
9. Clean nitrile gloves were donned by 1 assistant, and the remaining samples were processed for shipment. All samples were kept on ice packs during cleanup and processing; temperature integrity was not considered compromised.

Risk mitigation strategies were initiated as a result of this incident, including: (1) when removing samples from the cooler, ensuring these be held closely over the opening of the cooler with QC inspection performed before moving the sample to the TC; (2) re-training sample collection staff to ensure the SCs are adequately sealed before transport; and (3) storing disinfecting wipes and additional PPE (gloves, N95 respirators, gowns, and face shields) in the sample processing area. Real-time biosafety and biocontainment training conducted by an LSME in the field before sample handling was deemed essential to remove any confusion and discomfort felt by staff when handling a potential infectious spill.

Discussion

In the National Institute for Occupational Safety and Health (NIOSH) Hierarchy of Controls to reduce occupational hazards (Figure 1),^{8,9} the goal is to use the highest level possible of risk reduction. PPE is considered the least impactful level to reduce risk of infection, yet, before the incident, was the primary mitigation method used. The response plan for this incident effectively used

