vol. 72 ■ no. 7

CONTINUING EDUCATION

Health Care Workers' Comfort Ratings for Elastomeric Half-Mask Respirators Versus N95® Filtering Facepiece Respirators During the COVID-19 Pandemic

Lisa Pompeii, PhD^{1,2,3} D, Janelle Rios, PhD³, Colleen S. Kraft, MD, MSc⁴, Marie Kasbaum, MPH², Elisa Benavides, MPH², Scott J. Patlovich, DrPH³, Luis Ostrosky-Zeichner, MD³, Adam Hornbeck, MSN, APRN, FNP-BC, FNP-C⁵, Caitlin McClain, MS, GSP⁵, Rohan D. Fernando, MS⁵, Margaret Sietsema, PhD⁵, and Morgan Lane, MPH⁴

Abstract: Background: Reusable elastomeric half-mask respirators (EHMR) are an alternative to address shortages of disposable respirators. While respirator discomfort has been noted as a barrier to adherence to wearing an N95 filtering facepiece respirator (FFR) among health care personnel (HCP), few have examined EHMR comfort while providing patient care, which was the purpose of this study. Method: Among a cohort of 183 HCP, we prospectively examined how HCP rated EHMR tolerability using the Respirator Comfort, Wearing Experience, and Function Instrument (R-COMFI) questionnaire at Study Week 2 and Week 10. At the completion of the study (Week-12), HCP compared EHMR comfort with their prior N95 FFR use. Overall R-COMFI scores and three subscales (comfort, wear experience, and function) were examined as well as individual item scores. Findings: The HCP reported an improved overall R-COMFI score (lower score more favorable, 30.0 vs. 28.7/47, respectively) from Week 2 to Week 10. Many individual item scores improved or remained low over this period, except difficulty communicating with patients and coworkers. The overall R-COMFI scores for the EHMR were more favorable than for the N95 FFR (33.7 vs. 37.4, respectively), with a large proportion of workers indicating their perception that EHMR fit better, provided better protection, and they preferred to wear it in pandemic conditions compared with the N95 FFR. Conclusion/Application to Practice: Findings suggest that the EHMR is a feasible respiratory protection device with respect to tolerance. EHMRs can be considered as a possible alternative to the N95 FFR in the health care setting. Future work is needed in the EHMR design to improve communication.

Keywords: respiratory conditions/asthma, respiratory protection/respirators, personal protective equipment, health care worker/homecare worker, COVID-19

Background

Respiratory protection safeguards health care personnel (HCP) from exposure to infectious aerosol transmissible pathogens and is a critical component of infection control strategies in health care settings (Occupational Safety and Health Administration, 2011). The primary respiratory protection used in medical settings is the NIOSH Approved® N95® filtering facepiece respirator (N95 FFR; Wizner et al., 2016); however, hospitals faced significant challenges during the COVID-19 pandemic caused by the SARS-CoV-2 virus with shortages (Ahmed et al., 2020; Auerbach et al., 2021), leaving HCP vulnerable to exposure to infectious pathogens. In recent years, the elastomeric half-mask respirator (EHMR) has been proposed as one alternative to the N95 FFR (Baracco et al., 2015; Centers for Disease Control and Prevention, 2020; de Perio et al., 2020). EHMRs are half-face, tight-fitting respirators that are made of synthetic or rubber material and are equipped with replaceable filters or cartridges, and they have the same assigned protection factor (APF) as the N95 FFR (Centers for Disease Control and Prevention, 2020). One advantage over the N95 FFR is that the EHMR can be repeatedly disinfected, cleaned and reused. To date, literature in this area remains sparse, with very few studies having examined EHMR use in health care (Ciconte & Danyluk, 2013; Hines et al., 2019a, 2019b). Prior studies suggest that discomfort was a barrier to HCP's adherence to wearing the N95 FFR (Locatelli et al., 2014),

DOI: 10.1177/21650799241238755. From ¹Cincinnati Children's Hospital Medical Center, ²Baylor College of Medicine, ³The University of Texas Health Science Center at Houston, ⁴Emory University, and ⁵Centers for Disease Control and Prevention. Address correspondence to: Lisa Pompeii, PhD, Cincinnati Children's Hospital Medical Center, 3333 Burnet Ave., Cincinnati, OH 45229, USA; email: lisa.pompeii@cchmc.org.

For reprints and permissions queries, please visit SAGE's Web site at http://www.sagepub.com/journalsPermissions.nav.

Copyright © 2024 The Author(s)



Applying Research to Occupational Health Practice

This prospective cohort study examined HCWs level of tolerance to wearing the EHMR during routine patient care, which we measured every 2 weeks, over 12 weeks, using the validated R-COMFI questionnaire. We also asked workers to compare their tolerance for the EHMR with their prior use of the N95 FFR. Overall, workers showed an improved tolerance for the EHMR between Week 2 and Week 10 of the study. Moreover, they rated the EHMR more favorably than their N95 FFR. These findings suggest that the EHMR may be a feasible respiratory protection device with respect to tolerance, which includes comfort, wear experience, and function. There are numerous factors that the occupational health nurse should consider when introducing the EHMR into the hospital setting, including the processes for fit testing and training workers, as well as training them on how it is used, disinfected, and stored. Our study only examined a single make and model of EHMR, while it would be beneficial to have various EHMR makes/models available to workers. There are numerous steps involved with introducing the EHMR into the hospital setting, and perhaps pilot testing the process prior to scaling the program hospital-wide would be advantageous.

with limited findings about how the EHMR has been rated (Hines et al., 2019b; Radonovich et al., 2009). The purpose of this study was to examine HCP's level of comfort while wearing the EHMR when providing patient care and to compare comfort ratings between the EHMR and N95 FFR.

Methods

We conducted a 12-week prospective cohort study at two Level-1 trauma centers in Georgia (GA) and Texas (TX) from September through December 2021. Workers were fit-tested and trained to wear an EHMR, which they were asked to wear over a 12-week time period in lieu of their assigned N95 FFR. More specifically, they were trained to wear their EHMR when respiratory protection was warranted (e.g., patients with known aerosol transmissible diseases). Every 2 weeks, workers were asked to complete a survey about their perceived comfort and function with the respirator. Study details about EHMR wear time, respirator disinfection, use, and storage are reported elsewhere (Lane et al., 2024).

Elastomeric Respirator

The elastomeric respirator that was included in this study was the NIOSH Approved Honeywell RU8500 (Honeywell International Inc., Charlotte, NC) which participants were asked to use during routine patient care (e.g., feeding, bathing,

suctioning, and intubation). This model was chosen due to four characteristics (Figure 1): (a) the exhalation valve diversion cover that kept air from being exhaled into a patient's breathing zone; (b) filter covers that prevented potential liquid splashes from reaching the filter material; (c) the silicone body, as silicone is known to be easy to clean and disinfect; and (d) the speech diaphragm to facilitate clearer communication from the wearer. Because little was known about the EHMRs exhaled air during the start of the COVID-19 pandemic, the study hospitals required workers to place a surgical mask over the exhalation valve to ensure that patients were not inadvertently exposed to COVID-19 if a worker was infected (National Institute for Occupational Safety and Health [NIOSH], 2022). For purposes of feasibility, we chose to use smaller, pediatric masks, which workers were trained on how to place.

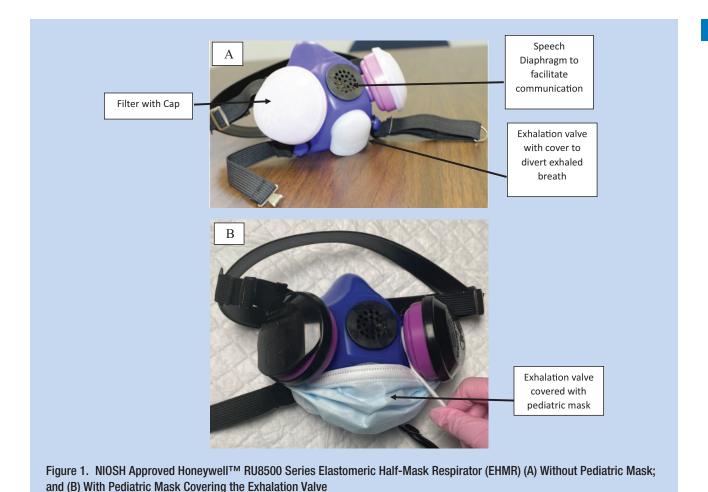
Subjects

Workers were eligible to participate if they were: (a) at least 18 years of age, (b) their jobs involved direct patient care or in-person counseling, (c) had been fit-tested to wear a N95 FFR in the prior two years, and (d) answered "yes" when asked if they regularly wore a respirator or mask to care for patients in the prior year. Workers who reported having anosmia or ageusia during recruitment were excluded from the study given the assumption that they would not be able to detect Bitrex™ (Dräger US, Houston, TX), the qualitative fit testing solution which was used to fit test workers prior to respirator use. They were also excluded if they had facial hair (e.g., beard or mustache) or a facial injury or adornment that would interfere with the seal of the respirator. At the TX hospital, recruitment was targeted to HCP who worked on inpatient units that were more likely to have patients with aerosol transmissible diseases such as COVID-19 or tuberculosis (e.g., emergency department and intensive care unit). The GA hospital restricted its recruitment to nurses and certified nursing assistants that provide care on one general medical ward and one intensive care unit. Workers were incentivized up to US\$450 for their participation in the seven study surveys. This study received human subjects' approval from the Institutional Review Board at The University of Texas-Health Science Center in Houston, Baylor College of Medicine, and Emory University.

Recruitment

Recruitment at the TX hospital was conducted virtually due to hospital-based COVID-19 restrictions that prevented study staff from entering the hospital. Study investigators presented an online study overview presentation to eight inpatient unit leaders who were then asked to email study invitations and flyers to their frontline workers. The invitations included a link to a REDCap (Harris et al., 2019) study site page in which workers had access to an online video about the study, an inclusion criteria questionnaire, and an online consent form. Workers who met the eligibility criteria and completed the consent form were

vol. 72 ■ no. 7 WORKPLACE HEALTH & SAFETY



then asked to sign up for a 60-minute time slot for fit testing and training. During the fit testing and training session, they watched a 10-minute video about EHMR use, what to expect over the course of the study, and the request to complete biweekly surveys. They were then fit-tested using qualitative methods and received face-to-face training on how to disinfect the EHMR and how to place the pediatric mask over the exhalation valve.

At the GA site, HCP employed on a general medical ward and an intensive care unit were invited to participate including nurses, certified nursing assistants, and patient care assistants. HCP received information about the study at daily unit huddles and through email. Charge nurses were recruited as "super users" and received additional training so that they could be the on-unit experts on the use of the EHMRs. Super users reminded participants to wear the EHMR and were available to answer questions and let the study team know if more supplies were needed. Upon enrollment, HCP filled out a screening survey and consent form online and then scheduled themselves to attend a fit testing session through the study hospital's employee health unit. During fit testing appointments, Emory's study team conducted a small training session with participants on the parts of the respirator, the placement of a pediatric mask over the

exhalation valve, and the surveys they would be asked to complete as part of the study. Super users were trained to conduct an in-service training for the participants on their unit which included protocols for disinfection and storage. Participants were also sent a link to a video describing the different aspects of EHMR use.

Data Collection

Participants were asked to complete pre- and post-surveys (baseline and Week 12) and five biweekly surveys during Weeks 2 through 10 of the study. For purposes of examining the level of comfort with wearing the EHMR, participants were asked to complete the validated Respirator Comfort, Wearing Experience, and Function Instrument (R-COMFI; LaVela et al., 2017).

For these analyses, we examined their responses to this survey pertaining to EHMR tolerability at study Weeks 2 and 10, in which workers were asked to rank their tolerability in the prior 2 weeks of EHMR use. In addition, at Week 12 workers were asked to compare the EHMR comfort over the entire study period with their comfort of prior N95 FFR use before the onset of the study.

The R-COMFI questionnaire includes 21 questions that measure respirator tolerability including subscales of discomfort (10 items), general wearing experience (6 items), and function (5 items). For respirator discomfort, workers were asked to rate on a three-point Likert-type scare (0 = none of the time, 1 =some of the time, 2 = all of the time) about physical discomfort such facial irritation, itching, and pinching as well as experiences with nausea and headache while wearing the respirator (score range 0 to 20; higher score indicates greater discomfort). For general wearing experience, the same threepoint Likert-type scale was used to rate workers' experiences with physical factors such as dizziness, claustrophobia, shortness of breath, and dry/itchy eyes (score range 0 to 12, higher score indicates more difficult wear experience). The function scale measured workers' ability to verbally communicate, their hearing and vision, as well as general interference with patient care duties while wearing the respirator, on a four-point Likert-type scale (0 = strongly)disagree, 1 = disagree, 2 = agree, and 3 = strongly agree, score range 0 to 15, higher score indicated greater interference). We added four additional questions to the survey in which workers were asked to rate their difficulty with getting straps over their hair, frustration with wearing, and whether patients or coworkers reacted negatively to the respirator.

At the 12-week post-study period, participants were asked their level of agreement on a four-point Likert-type scale about their experiences with wearing the EHMR versus the N95 FFR, including questions as to which respirator they felt provided a better fit, better respiratory protection, and their preference of which respirator they would prefer to wear in current and future pandemic and non-pandemic conditions.

Open-ended questions were asked in each of the biweekly surveys, where participants were asked to indicate any new or continuous challenges with wearing the EHMR in the previous 2 weeks as well as issues that had improved. In the post-survey, participants were also asked to describe what they liked least and most about the EHMR.

Demographic details were also collected including age category, race, Hispanic/Latino ethnicity (yes/no), self-described sex, job title, work department, and types of prior respirator use.

Data Analysis

Descriptive statistics were employed to describe the occupational characteristics of study participants and their history of respirator use. For purposes of comparing changes in the R-COMFI measures across time periods, we restricted our analyses to individuals who completed both Weeks 2 and 10 surveys. For examination of individual survey items, categories were collapsed to compare some/all of the time (some/all) versus none of the time (none) for the Discomfort and General Wear items, and strongly agree/agree versus strongly disagree/disagree for the Function items. To examine an overall R-COMFI score, the three subscales were summed, including Discomfort (score range: 0-20) and General Wear (score range:

0-12), and Function (score range: 0-15; LaVela et al., 2017; Radonovich et al., 2019). The total sum of the R-COMFI score could range from 0 to 47, with a lower score equaling better respirator tolerability. All participants were included in the analysis at Week 12 where we compared R-COMFI measures between the EHMR and the N95 FFR. Differences between categorical measures were examined using the Mantel-Haenszel chi-square test, while differences in R-COMFI scores were measured using the paired *t* test. Open-ended data regarding respirator comfort and use were categorized into similar themes by two study investigators and reviewed by a third to ensure the accuracy of coding. We sought to describe themes that were more commonly experienced. SAS 9.4 was used to conduct all quantitative data analysis.

Findings

Among the 208 workers recruited from both locations, 183 (88.0%) completed the study through Week 12, and of those, 139 (74.0%) completed both the Week 2 and Week 10 surveys. Participants were more likely to be aged 18 to 35 years (62.3%) and identify as female (74.3%; Table 1). A large

Table 1	Demographic Characteristics of Health Care
Personn	nel (n = 183)

Age	N (%)
18–25	38 (20.8)
26–35	76 (41.5)
36–45	31 (16.9)
45–55	21 (11.5)
>55	17 (9.3)
Self-described sex	
Male	44 (24)
Female	136 (74.3)
None of these	3 (1.6)
Race	
American Indian or Alaska Native	5 (2.7)
Asian	27 (14.8)
Black or African American	38 (20.8)
Native Hawaiian or Other Pacific Islander	1 (0.5)
White	108 (59)
More than one race	4 (2.2)

Table 1	(continued)	۱
Table I. (Comunica	,

Age	N (%)			
Hispanic ethnicity				
Yes	28 (15.3)			
No	155 (84.7)			
Education				
High school/GED/Tech/Certificate program	14 (7.6)			
Associate's degree	23 (12.6)			
Bachelor's degree	96 (52.5)			
Master's degree	34 (18.6)			
Doctoral (MD/PhD)	16 (8.7)			
Job title				
Patient Care Assistant/Patient Sitter/Lift Team Technician	17 (9.3)			
Nurse	95 (51.9)			
Physician/Physician Assistant/Nurse Practitioner/CRNA/Med Student	18 (9.8)			
Respiratory Therapist /Ultrasound Neurology	18 (9.8)			
Occupational Therapist/Assist/Physical Therapist/Asst./Speech Path	31 (16.9)			
Other	4 (2.2)			
Work department				
Emergency department/trauma	17 (9.3)			
Intensive care unit (ICU)	104 (56.8)			
Operating room	4 (2.2)			
Float (GM/ICU)	40 (21.8)			
General medicine (GM)	15 (8.2)			
Other	3 (1.6)			
Prior respirator or mask use (not mutually exclusive)				
Surgical/procedural mask ^a	176 (84.2)			
N95 FFR	155 (74.2)			
EHMR	4 (1.9)			
Powered Air Purifying Respirator (PAPR)	5 (2.4)			

^aSurgical/Procedural mask is not a form of respiratory protection.

proportion indicated their race as White (59.0%), Black (20.8%), or Asian (14.8%), while the majority were non-Hispanic ethnicity (84.7%). More than half (52.5%) reported having a bachelor's degree or higher (27.3%). Workgroups were predominantly nurses (51.9%), occupational/physical/speech therapists or assistants (16.9%), physicians/nurse practitioners or assistants (9.8%), or respiratory therapists (9.8%). A large proportion of workers were employed in intensive care units (ICU; 56.8%) or floated between hospital units (21.8%) followed by working in the emergency department (9.3%) or in general medicine units (8.2%).

EHMR Wear Week 2 Versus Week 10

Among participants who indicated wearing their EHMR in the prior 2 weeks at study Week 2 and Week 10, the overall R-COMFI score slightly improved from 30.0 to 28.7 (p < .01; Table 2).

When examining the mean R-COMFI subscales, the discomfort score slightly improved from Week 2 to Week 10 (mean score: 14.0 vs. 13.1, p < .01). For the individual categorical discomfort scale items, all but two items improved in Week 10 relative to Week 2, with two that were significantly improved including facial irritation (64.7% vs. 52.9%; p = .05) and sweat/moisture build-up (56.1% vs. 40.6%; p < .01), and two that were borderline significant, including tightness of straps (49.6% vs. 38.8%; p = .07), and lack of fresh air (28.1%) vs. 18.7%; p = .07). The other discomfort measures were not significantly different across time periods, with approximately 40% to 50% that indicated all/some of the time for nose pinching and facial heat/warmth. A much smaller percentage reported facial itching, pinching, nausea, and headache. The improved comfort scores over the 10-week period (Week 2 vs. 10) were reflected in participants' comments, such as one worker who referred to the EHMR as a "mask" when they stated, "I surprisingly liked the mask a lot more as I got more comfortable with it. I will probably continue to use it . . .," while another indicated "[The] comfort has gotten better-feels like [the] mask has molded to my facial features and sits perfectly, fitting got better."

For the total general wear experience mean score, no difference was observed between Weeks 2 and 10 (mean score: 6.5 vs. 6.5, p=.54). When examining the individual categorical general wear items, approximately one-third of participants reported general frustration with wearing the EHMR (30.9% vs. 25.9%, p=.35), noting that the EHMR interfered with wearing glasses/face shield/goggles (36.0% vs. 38.1%, p=.71) as well as experiencing difficulty getting straps over their hair, which was not significantly different across time periods (36.0% vs. 30.9%, p=.37). When participants were asked what they liked/disliked about the EHMR, one participant indicated "I didn't like my hair getting stuck in the straps" while others indicated "It was hard to keep my glasses in place on bridge of my nose," and "It took time to get used to wearing bifocals with the mask (i.e., EHMR). When I was able to adjust

Table 2 Continuous and Categorical Ratings of R-COMFI Among Health Care Personnel at Weeks 2 and 10 of Elastomeric (EHMR) Use $(n = 139)^a$

	EHMR Wee	k 2 (<i>N</i> = 139) ^c	EHMR Week	10 (<i>N</i> = 139)°	
R-COMFI Scores ^b	M	(SD)	М	(SD)	P value
Discomfort Score	14.0 (3.1)		13.1 (3.4)		<.01
General Wear Experience Score	6.	5 (0.95)	6.	5 (1.8)	.54
Function Score	9.	5 (2.5)	9.	1 (2.9)	.04
Overall Tolerance Score	30.	0 (5.2)	28.	28.7 (6.5)	
	Some/All	None	Some/All	None	
Individual R-COMFI Items	N (%)	N (%)	N (%)	N (%)	
Discomfort					
Tightness of straps	69 (49.6%)	70 (50.4%)	54 (38.8%)	85 (61.2%)	.07
Facial irritation (leaves marks/indents)	90 (64.7%)	49 (35.3%)	73 (52.9%)	65 (47.1%)	.05
Facial itching	30 (21.6%)	109 (78.4%)	25 (18%)	114 (82%)	.45
Facial pinching	32 (23%)	107 (77%)	28 (20.1%)	111 (79.9%)	.56
Nose, nose-bridge (pinching, redness)	61 (43.9%)	78 (56.1%)	54 (39.4%)	83 (60.6%)	.45
Facial heat/warmth	63 (45.3%)	76 (54.7%)	52 (37.4%)	87 (62.6%)	.18
Sweat/moisture build-up	78 (56.1%)	61 (43.9%)	56 (40.6%)	82 (59.4%)	<.01
Lack of fresh air	39 (28.1%)	100 (71.9%)	26 (18.7%)	113 (81.3%)	.07
Nausea	5 (3.6%)	134 (96.4%)	8 (5.9%)	128 (94.1%)	.37
Headache	14 (10.3%)	122 (89.7%)	18 (12.9%)	121 (87.1%)	.49
General wear experience					·
Dizziness	4 (2.9%)	134 (97.1%)	5 (3.6%)	133 (96.4%)	.76
Loss of energy/ tiredness/ fatigue	11(7.9%)	128 (92.1%)	15 (10.8%)	124 (89.2%)	.41
Claustrophobia	11 (8%)	127 (92%)	14 (10.1%)	125 (89.9%)	.54
Shortness of breath	14 (10.1%)	125 (89.9%)	15 (10.8%)	124 (89.2%)	.84
Difficulty of breathing	15 (10.9%)	123 (89.1%)	15 (10.8%)	124 (98.2%)	.98
Dry or itchy eyes	9 (6.5%)	130 (93.5%)	7 (5.1%)	131 (94.9%)	.62
Frustration with wearing ^d	43 (30.9%)	96 (69.1%)	36 (25.9%)	103 (74.1%)	.35
Interference with wearing glasses, goggles, face shield ^d	50 (36.0%)	89 (64%)	53 (38.1%)	86 (61.9%)	.71
Difficulties getting straps over hair (hair pulling) ^d	50 (36.0%)	89 (64%)	43 (30.9%)	96 (69.1%)	.37

(continued)

Table 2. (continued)

	Strongly agree/ agree	Strongly disagree/ disagree	Strongly agree/ agree	Strongly disagree/ disagree	
Function					
Mask ^e affected concentration (always adjusting the mask)	21 (15%)	119 (98%)	18 (12.9%)	121 (87.1%)	.62
Difficulty verbally communicating to others (unintelligible, muffled speech)	84 (60%)	56 (40%)	73 (52.5%)	66 (47.5%)	.21
Difficulty hearing others	20 (14.3%)	120 (85.7%)	14 (10.1%)	125 (89.9%)	.37
Mask ^e obstructed my vision	11 (7.9%)	129 (92.1%)	8 (5.8%)	129 (94.2%)	.51
Mask ^e interfered with my patient care duties (quicker to leave room, less interaction)	15 (10.7%)	125 (89.3%)	18 (12.9%)	121 (87.1%)	.56
Patients reacted negatively to the respirator ^a	20 (14.3%)	120 (85.7%)	15 (10.9%)	123 (89.1%)	.39
Coworkers reacted negatively to the respirator ^a	11 (7.9%)	129 (92.1%)	8 (5.8%)	131 (94.2%)	.49

^aMust have worn respirator in Week 2 and Week 10. ^b R-COMFI scores do not include questions that were not in the original R-COMFI questionnaire. ^c <2% missing responses for some cells. ^d Questions not included in the original R-COMFI questionnaire. ^e Mask in this survey refers to the elastomeric half-mask respirator.

my glasses with the straps there were very few problems." Only a small proportion of participants (<10%) reported dizziness, loss of energy/fatigue, shortness of breath, or dry/itchy eyes in either time period.

Differences in the EHMR mean function score between Weeks 2 and 10 were modestly improved (mean score: 9.5 vs 9.1, p = .04). The individual categorical function items revealed that more than half of the participants during Week 2 (60.0%) had difficulty communicating to others despite this EHMR having a speech diaphragm, which persisted through to Week 10 (52.5%; p = .21). When participants were asked to indicate what they did not like about the EHMR, there was consensus that communicating with patients was difficult, with one worker indicating, "My main dislike was that it was difficult for patients and peers to hear me with the respirator on," and another indicating ". . . I felt like I was always yelling." When participants were asked how they would improve the design of the respirator, some indicated that the respirator needed an "improved voice box" or a "microphone." Participants rated the other function items more favorably with only a small proportion indicating difficulty hearing others (Week 2,14.3% vs. Week 10, 10.1%, p = .37), visual obstruction due to mask (7.9% vs. 5.8%, p = .51), adversely affecting concentration (15.0% vs.12.9%, p = .62), and interference with patient care (10.7% vs. 12.9%, p = .56). A small proportion indicated that patients reacted negatively to the EHMR (14.3%

vs. 10.9%, p=.39), and fewer (7.9% vs. 5.8%, p=.49) indicated that participants reacted negatively, with some stating, "It does freak out some older, confused patients" and "People react to it as if it's a bit sci-fi."

EHMR Versus N95 FFR

When examining the overall and subscale *R*-COMFI scores at the end of the study period (Week 12) between the EHMR and N95 FFR, the overall score was more favorable for the EHMR (33.7) versus the N95 FFR (37.4, p < .01; Table 3). Similarly, two of the three subscales of discomfort (p < .01) and general wear experience (p = .03) were more favorable for the EHMR, while function was more favorable for the N95 FFR (p < .01).

When comparing individual comfort items across the R-COMFI subscales, most were rated more favorably for the EHMR relative to the N95 FFR. Among the discomfort items, all were rated more favorably for the EHMR, including facial itching (EHMR: 35.7% vs. N95 FFR: 61.9%; p < .01), lack of fresh air (50.3% vs. 78.7%, p < .01), nausea (8.8% vs. 16.5%, p = .03) and headache (21.8% vs. 38.2%, p < .01).

Similar patterns were observed for the general wear items, including shortness of breath (EHMR 20.2% vs. N95 FFR: 41.2%; p < .01), difficulty breathing (21.4% vs. 47.3%; p < .01), and dizziness (6.6% vs. 14.8%; p < .01). Although more than half (62.3%) of participants indicated frustration with wearing the

Table 3. Comparison of R-COMFI of Elastomeric Versus N95 Filtering Facepiece Respirators Among Health Care Personnel at Two U.S. Hospitals (n=183)

	EHMR Week	12 (<i>N</i> = 183) ^b	N95 Week	12 (<i>N</i> = 183) ^b	
R-COMFI scores ^a	M	(SD)	M	(SD)	P value
Discomfort Score	14.0 (3.5)		16.9 (4.0)		<.01
General Wear Experience Score	7.0) (1.7)	7.8	7.8 (2.1)	
Function Score	11.	0 (2.8)	10.	7 (2.8)	<.01
Overall Tolerance Score	33.	7 (6.7)	37.	4 (7.2)	<.01
	Some/All	None	Some/All	None	
Individual R-COMFI items	N (%)	N (%)	N (%)	N (%)	
Discomfort					_
Tightness of straps	118 (64.5%)	65 (35.5%)	145 (79.2%)	38 (20.8%)	<.01
Facial irritation (leaves marks/indents)	131 (71.6%)	52 (28.4%)	161 (88%)	22 (12%)	<.01
Facial itching	65 (35.7%)	117(64.3%)	112 (61.9%)	69 (38.1%)	<.01
Facial pinching	94 (51.4%)	89 (48.6%)	123 (67.2%)	60 (32.8%)	<.01
Nose, nose-bridge (pinching, redness)	104 (57.8%)	76 (42.2%)	157 (87.2%)	23 (12.8%)	<.01
Facial heat/warmth	115 (63.5%)	66 (36.5%)	144 (79.6%)	37 (20.4%)	<.01
Sweat/moisture build-up	126 (69.6%)	55 (30.4%)	149 (81.9%)	33 (18.1%)	<.01
Lack of fresh air	92 (50.3%)	91 (49.7%)	144 (78.7%)	39 (21.3%)	<.01
Nausea	16 (8.8%)	166 (91.2%)	30 (16.5%)	152 (83.5%)	.03
Headache	39 (21.8%)	140 (78.2%)	68 (38.2%)	110 (61.8%)	<.01
General wear experience					
Dizziness	12 (6.6%)	170 (93.4%)	27 (14.8%)	155 (85.2%)	.01
Loss of energy/tiredness/fatigue	36 (19.7%)	147 (80.3%)	52 (29.2%)	126 (70.8%)	.03
Claustrophobia	34 (18.7%)	148 (81.3%)	44 (24.2%)	138 (75.8%)	.2
Shortness of breath	37 (20.2%)	146 (79.8%)	75 (41.2%)	107 (58.8%)	<.01
Difficulty of breathing	39 (21.4%)	143 (78.6%)	86 (47.3%)	96 (52.7%)	<.01
Dry or itchy eyes	20 (11%)	162 (89%)	29 (16%)	152 (84%)	.16
Frustration with wearing ^c	114 (62.3%)	69 (37.7%)	141 (77%)	42 (23%)	<.01
Interference with wearing glasses, goggles, face shield ^c	105 (57.7%)	77 (42.3%)	104 (57.5%)	77 (42.5%)	.96
Difficulties getting straps over hair (hair pulling) ^c	98 (53.6%)	85 (46.4%)	98 (53.6%)	85 (46.4%)	.99

(continued)

Table 3. (continued)

	Strongly agree/ agree	Strongly disagree/ disagree	Strongly agree/ agree	Strongly disagree/ disagree	
Function					
Mask ^d affected concentration (always adjusting the mask)	41 (22.4%)	142 (77.6%)	73 (40.1%)	109 (59.9%)	<.01
Difficulty verbally communicating to others (unintelligible, muffled speech)	139 (76%)	44 (24%)	97 (53.3%)	85 (46.7%)	<.01
Difficulty hearing others	41 (22.4%)	142 (77.6%)	30 (16.6%)	151 (83.4%)	.16
Mask ^d obstructed my vision	24 (13.2%)	158 (86.8%)	20 (11%)	162 (89%)	.52
Mask ^d interfered with my patient care duties (quicker to leave room, less interaction)	54 (29.5%)	129 (70.5%)	48 (26.4%)	134 (73.6%)	.51
Patients reacted negatively to the respirator ^c	42 (23.1%)	140 (76.9%)	15 (8.2%)	168 (91.8%)	<.01
Coworkers reacted negatively to the respirator ^c	30 (16.5%)	152 (83.5%)	14 (7.7%)	167 (92.3%)	.01

^aR-COMFI scores do not include questions that were not in the original R-COMFI questionnaire. ^b <2% missing responses for some cells. ^c Questions not included in the original R-COMFI questionnaire. ^d Mask in this survey refers to the elastomeric half-mask respirator.

EHMR, even more (77.0%, *p* < .01) rated the N95 FFR as frustrating to wear. When participants were asked what they liked about the EHMR, many indicated that they favored it over the N95 FFR, with one stating, "I like that there is a good, secure fit to my face and it's easy to test the positive/negative pressure to ensure a good fit. The material also felt better against my face, compared to the N95 FFR." Another participant indicated, "[It] made me feel extra secure during my patient interactions. I felt the fit was more customized to my face compared with the N95s FFRs provided by my department."

For the EHMR function measures, only one item was rated more favorably for the EHMR versus the N95 FFR, which included the respirator affecting the worker's concentration while working (EHMR 22.4% vs. N95 FFR 40.1%; p < .01). The EHMR was rated as being more difficult than the N95 FFR for verbally communicating (76.0% vs. 53.3%; p < .01). Workers also indicated that they perceived more negative reactions from patients (23.1% vs. 8.2%, p < .01) and coworkers (16.5% vs. 7.7%, p < .01) with the EHMR relative to the N95 FFR. No significant differences between the two devices were observed for difficulty hearing others, obstructed vision, or interference with patient care.

In the post-study period, compared to the N95 FFR, the majority of participants (88%) indicated that they felt the EHMR fit better, felt it provided greater protection from viruses and bacteria (88%), and they preferred to wear it while caring for

COVID-19 patients (74.9%; Table 4). These findings were reiterated in the open-ended questions where participants were asked what they liked about the EHMR, with many indicating that they felt more protected relative to the N95 FFR. For example, one worker indicated that "It had a great seal and I felt better protected related to mask [facepiece] leaks with N95 FFR," while others indicated that it "felt safe and more comfortable than a N95 FFR," it has a "better seal than [the] N95 FFR," and "I can breathe easier." With regard to wearing the EHMR during non-pandemic conditions, the majority of participants (69.9%) indicated that they felt comfortable doing so; however, half (53.6%) indicated that in nonpandemic conditions they would prefer to wear the N95 FFR over the EHMR. Some participants commented on their preference for N95 FFR in non-pandemic conditions, with one participant who indicated "for extended wear the elastomeric is much more comfortable, the N95 FFR is more convenient for intermittent use as it doesn't require disinfection and drying time."

When participants were asked why they chose to participate in this study, as well as aspects of the EHMR they liked, many indicated that they liked that the EHMR could be reused, with participants stating, "It minimizes waste," "I feel like I am creating less waste by using the [EHMR]," and "I do like that you do not have to throw it away and it is essentially sustainable." Because of respirator shortages, participants also liked that they were assigned their own respirator, with one worker indicating,

Table 4.	Health Care Personnel's Perce	ptions of the EHMR After	Three Months of Use	(N = 183)

Survey Items	Agree/ strongly agree	Disagree/ strongly disagree
The EHMR fits better than the N95 [FFR]	158 (86.3%)	25 (13.7%)
The EHMR provides more protection from viruses and bacteria than the N95 [FFR]	161 (88.0%)	22 (12.0%)
Prefers to wear the elastomeric over the N95 [FFR] during this pandemic while caring for COVID-19 patients	137 (74.9%)	46 (25.1%)
Prefers to wear the N95 [FFR] over EHMR in non-pandemic conditions	98 (53.6%)	85 (46.4%)
Feels comfortable continuing to wear the EHMR during the remainder of the pandemic	152 (83.1%)	31 (16.9%)
Feels comfortable wearing the EHMR while caring for patients in a non-pandemic environment	128 (69.9%)	55 (30.1%)
Feels comfortable recommending the EHMR to their colleagues	170 (92.9%)	13 (7.1%)

Note. EHMR = elastomeric half mask respirator. N95 [FFR] = N95 filtering face piece respirator.

"It was only mine, I was responsible for cleaning and caring for it, and I didn't worry that it was worn by others then disinfected." Others indicated, "I feel secure knowing I have an alternative mask [i.e., respirator] especially with supply shortage we've seen in the past" and "I liked having a personal respirator and not being reliant upon department provision of N95 FFR."

Discussion

This study examined HCP's perceived levels of physical comfort and function with wearing the EHMR while conducting routine patient care. Overall, the EHMR received respectable R-COMFI scores among participants who wore the respirator consistently over a 10-week period, including improved scores for several of the comfort items between the two time periods. These findings are promising, especially given that we also observed significantly better comfort ratings for the EHMR relative to the N95 FFR. To date, only one study has examined differences in comfort between these two devices, in which the N95 FFR was rated more favorably among HCPs who only wore the N95 FFR compared with HCPs who only wore the EHMR (Hines et al., 2017, 2019b). In this study, participants were asked a single question about how much they liked their respirator with respect to comfort and they rated the N95 FFR slightly higher (3.42) than the EHMR (3.28) on a five-point scale.

Although participants rated many of the R-COMFI items more favorably over time, and more favorably compared with the N95 FFR, there were some poorly rated items that persisted including issues with communication, facial irritation, facial pinching, nose pinching, heat/warmth, and interference with wearing glasses. Difficulty communicating while wearing the EHMR was not an unexpected finding as it has been reported in prior studies (Ciconte & Danyluk, 2013; Hines et al., 2019b),

and this is recognized as an issue that needs to be addressed when designing future EHMRs specifically for use in health care settings (Radonovich et al., 2010). Prior multidisciplinary efforts to improve the design of the EHMR have proven successful, including efforts put forth by Project B.R.E.A.T.H.E., which is a partnership between federal agencies (e.g., NIOSH's NPPTL), academic researchers and respirator manufacturers (U.S. Department of Veterans Affairs et al., 2009). It is essential to engage respirator manufacturers to improve the long-standing communication limitations with the EHMR.

We tested the use of the EHMR during an ongoing pandemic, with a large proportion of participants who indicated that they perceived the EHMR to fit better and to provide better protection relative to the N95 FFR, which is similar to findings reported in other studies conducted during non-pandemic conditions (Ciconte & Danyluk, 2013; Hines et al., 2019b). Similar to Hines et al. (2019b), our study participants indicated that while the EHMR fit better and they perceived better protection, a large proportion also indicated that they would prefer to wear the N95 FFR in nonpandemic conditions (Hines et al., 2017, 2019b). Qualitative data from our study suggested that participants felt the N95 FFR was easier to use than the EHMR, especially when quick tasks needed to be conducted. Additional findings from our study (reported elsewhere; Lane et al., 2024) suggest that the process of doffing and disinfecting the respirator between patients was perceived as timeconsuming, perhaps contributing to a preference for the disposable N95 FFR when shorter periods of respiratory protection are warranted.

There are several limitations to this study that should be considered. We compared EHMR use during the study period to workers' N95 FFR use prior to the start of the study; however, we did not obtain detailed measures such as

frequency of use. We used specific inclusion criteria to ensure that we included workers that had prior experience wearing the N95 FFR including those that had been previously fit-tested for the N95 FFR. Differences that we observed in EHMR vs. N95 FFR measures may have been influenced by differences in frequency of N95 FFR that we did not measure. Other limitations to consider, especially among occupational health professionals that are planning on implementing the EHMR into their health care setting. First, we only tested one model of EHMR which was the Honeywell RU8500 series. There are various EHMRs (makes and models) that should be pilot tested and considered prior to implementation. This includes newer models that do not have an exhalation valve or have adaptors to filter exhaled breath through valves, removing the need to place a mask or face shield over the valve. Such changes in model design could remove certain communication barriers but also exacerbate others such as breathing. This issue has not been examined. Given the stated dislikes for the EHMR by study participants, such as difficulty communicating and facial irritation, more studies need to be conducted for purposes of examining participants' physical comfort while performing patient care tasks among a range EHMR makes and models. Future studies should also examine if and/or how duration of EHMR use over a work shift can be modified based on comfort, which our study did not examine. For purposes of examining if EHMR comfort changed over time, we restricted our longitudinal analysis to participants who reported using the EHMR at study Weeks 2 and 10; however, this most likely excluded participants that may have decreased their use because they found the respirator to be uncomfortable. This may explain why the 12-week R-COMFI findings for the EHMR were less favorable for the whole cohort compared to those in the restricted analysis. In addition, some participants may not have worn the N95 FFR since before the study period, which may have biased their rating to be more favorable for the EHMR relative to the N95 FFR.

Implications for Occupational Health Practice

There are numerous factors to consider when implementing the EHMR into the health care setting. This study focused on the individual workers' comfort while wearing the EHMR, their ability to use the EHMR, disinfect and store it, as well as use it while performing specific patient care tasks (reported elsewhere; Lane et al., 2024). However, organizational factors must also be considered. For example, in a prior study, we observed that it was feasible to rapidly fit test and train HCPs to wear the EHMR in a hospital setting that had an established Occupational Safety and Health Administration (OSHA)approved Respiratory Protection Program (OSHA, 2011; Pompeii et al., 2020). Furthermore, Hines and colleagues (2023) conducted a study examining an existing EHMR program where the hospital, rather than the worker, was responsible for EHMR disinfection and storage. While findings from these and other studies demonstrate that EHMR use in health care is feasible,

more research is needed to support the implementation of an EHMR program at the work unit and organizational levels.

Acknowledgments

The authors would like to recognize Kevin Strickland, Emily Haas, and Mihili Edirisooriya from the National Personal Protective Technology Laboratory for their thoughtful reviews of this document. We would also like to thank Dr. Lewis Radonovich for his earlier contributions to this work.

Disclaimer

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention. Mention of any company or product does not constitute endorsement by the National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

Conflict of Interest

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This work was supported by the Centers for Disease Control and Prevention, National Institute of Occupational Safety and Health, cooperative agreement 75D30118C02645.

Human Subjects

This study received human subjects' approval from the Institutional Review Board at The University of Texas-Health Science Center in Houston, Houston Texas; Baylor College of Medicine, Houston, Texas; and Emory University, Atlanta, Georgia.

Attribution Statement

N95 is a certification mark of the U.S. Department of Health and Human Services (HHS) registered in the United States and several international jurisdictions.

ORCID iD

Lisa Pompeii (D) https://orcid.org/0000-0001-5017-6483

References

Ahmed, J., Malik, F., Bin Arif, T., Majid, Z., Chaudhary, M. A., Ahmad, J., Malik, M., Khan, T. M., & Khalid, M. (2020). Availability of Personal Protective Equipment (PPE) among US and Pakistani doctors in COVID-19 pandemic. *Cureus*, 12(6), e8550. https://doi.org/10.7759/cureus.8550

- Auerbach, M. A., Abulebda, K., Bona, A. M., Falvo, L., Hughes, P. G., Wagner, M., Barach, P. R., & Ahmed, R. A. (2021). A national US survey of pediatric emergency department coronavirus pandemic preparedness. *Pediatric Emergency Care*, 37(1), 48–53. https://doi. org/10.1097/PEC.00000000000002307
- Baracco, G., Eisert, S., Eagan, A., & Radonovich, L. (2015). Comparative cost of stockpiling various types of respiratory protective devices to protect the health care workforce during an influenza pandemic. *Disaster Medicine and Public Health Preparedness*, 9(3), 313–318. https://doi.org/10.17226/12748
- Centers for Disease Control and Prevention. (2020). Elastomeric respirators: Strategies during conventional and surge demand situations: Conventional, contingency, and crisis strategies. https://stacks.cdc.gov/pdfjs/web/viewer.html?file=https://stacks.cdc.gov/view/cdc/95188/cdc_95188_DS1.pdf
- Ciconte, R., & Danyluk, Q. (2013). Assessment and determination of practical considerations for wide-scale utilization of elastometric balf-facepiece respirators during a pandemic or outbreak situation. Workers' Compensation Board of Nova Scotia.
- de Perio, M. A., Dowell, C. H., Delaney, L. J., Radonovich, L. J., Kuhar, D. T., Gupta, N., Patel, A., Pillai, S. K., & D'Alessandro, M. (2020). Strategies for optimizing the supply of N95 filtering facepiece respirators during the coronavirus disease 2019 (COVID-19) pandemic. Disaster Medicine and Public Health Preparedness, 14(5), 658–669. https://doi.org/10.1017/dmp.2020.160
- Harris, P. A., Taylor, R., Minor, B. L., Elliott, V., Fernandez, M., O'Neal, L., McLeod, L., Delacqua, G., Delacqua, F., Kirby, J., Duda, S. N., & Consortium, R. E. (2019). The REDCap consortium: Building an international community of software platform partners. *Journal of Biomedical Informatics*, 95, 103208. https://doi.org/10.1016/j.jbi.2019.103208
- Hines, S. E., Brown, C., Oliver, M., Gucer, P., Frisch, M., Hogan, R., Roth, T., Chang, J., & McDiarmid, M. (2019a). Storage and availability of elastomeric respirators in health care. *Health Security*, 17(5), 384–392. https://doi.org/10.1089/hs.2019.0039
- Hines, S. E., Brown, C., Oliver, M., Gucer, P., Frisch, M., Hogan, R., Roth, T., Chang, J., & McDiarmid, M. (2019b). User acceptance of reusable respirators in health care. *American Journal of Infection Control*, 47(6), 648–655. https://doi.org/10.17226/25275
- Hines, S. E., Mueller, N., Oliver, M., Gucer, P., & McDiarmid, M. (2017). Qualitative analysis of origins and evolution of an elastomeric respirator-based hospital respiratory protection program. *Journal of the International Society for Respiratory Protection*, 34(2), 95–110.
- Hines, S. E., Thurman, P., Zhuang, E., Chen, H., McDiarmid, M., Chalikonda, S., Angelilli, S., Waltenbaugh, H., Napoli, M., Haas, E., McClain, C., Sietsema, M., & Fernando, R. (2023). Elastomeric halfmask respirator disinfection practices among healthcare personnel. *American Journal of Industrial Medicine*, 66(12), 1056–1068. https:// doi.org/10.1002/ajim.23538
- Lane, M., Pompeii, L., Rios, J., Benavides, E., Kasbaum, M., Patlovich, S., Ostrosky-Zeichner, L., Hornbeck, A., McClain, C., Fernando, R., Sietsema, M., & Kraft, C. (2024, 2024/01/24/). Provider experiences with daily use of elastomeric half-mask respirators in health care. *American*

- *journal of infection control.* Advance online publication. https://doi.org/https://doi.org/10.1016/j.ajic.2024.01.015
- LaVela, S. L., Kostovich, C., Locatelli, S., Gosch, M., Eagan, A., & Radonovich, L. (2017). Development and initial validation of the Respirator Comfort, Wearing Experience, and Function Instrument [R-COMFI]. *Journal of Occupational and Environmental Hygiene*, 14(2), 135–147. https://doi.org/10.1080/15459624.2016.1237025
- Locatelli, S. M., LaVela, S. L., & Gosch, M. (2014). Health care workers' reported discomfort while wearing filtering face-piece respirators. Workplace Health & Safety, 62(9), 362–368. https://doi. org/10.3928/21650799-20140804-03
- National Institute for Occupational Safety and Health. (2022). Evaluation of exhalation resistance and inspired carbon dioxide concentration in elastomeric half-mask respirators with modified or covered exhalation valves [Technical Report] (By Strickland, K. T., Fernando, R., Schall, J., Walbert, G., Brannen, J., U.S. Department of Health and Human Services, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health, DHHS (NIOSH) Publication No. 2022-109). https://doi.org/10.26616/NIOSHPUB2022109
- Occupational Safety and Health Administration. (2011). Occupational Safety & Health Standard: Personal Protective Equipment (29 Standard 1910.134). https://www.osha.gov/laws-regs/regulations/standardnumber/1910/1910.134
- Pompeii, L. A., Kraft, C. S., Brownsword, E. A., Lane, M. A., Benavides, E., Rios, J., & Radonovich, L. J. Jr (2020). Training and fit testing of health care personnel for reusable elastomeric half-mask respirators compared with disposable N95 respirators. *JAMA*, 323(18), 1849–1852. https://doi. org/10.1001/jama.2020.4806
- Radonovich, L. J. Jr., Cheng, J., Shenal, B. V., Hodgson, M., & Bender, B. S. (2009). Respirator tolerance in health care workers. *JAMA*, 301(1), 36–38. https://doi.org/10.1001/jama.2008.894
- Radonovich, L. J. Jr., Wizner, K., LaVela, S. L., Lee, M. L., Findley, K., & Yorio, P. (2019). A tolerability assessment of new respiratory protective devices developed for health care personnel: A randomized simulated clinical study. *PLOS ONE*, 14(1), e0209559. https://doi.org/10.1371/journal.pone.0209559
- Radonovich, L. J. Jr., Yanke, R., Cheng, J., & Bender, B. (2010). Diminished speech intelligibility associated with certain types of respirators worn by healthcare workers. *Journal of Occupational and Environmental Hygiene*, 7(1), 63–70. https://doi. org/10.1080/15459620903404803
- U.S. Department of Veterans Affairs, Veterans Health Administration, Office of Public Health and Environmental Hazards, National Institute for Occupational Safety and Health, National Personal Protective Technology Laboratory, National Aeronautics and Space Administration, Occupational Safety and Health Administration. (2009). Better respiratory equipment using advanced technologies for bealthcare employees (Project B.R.E.A.T.H.E.): An interagency working group of the U.S. federal government. https://stacks.cdc.gov/view/cdc/112432
- Wizner, K., Stradtman, L., Novak, D., & Shaffer, R. (2016). Prevalence of respiratory protective devices in U.S. Health care facilities: Implications for emergency preparedness. Workplace Health & Safety, 64(8), 359–368. https://doi.org/10.1177/2165079916657108