Infection prevention and control measures for Ebola and Marburg Virus disease: A series of rapid reviews

KQ5 Head/Neck Covers- Initial Summary

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Nicole Shaver, nicole.shaver@uottawa.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada. ORCID 0000-0003-3210-8895

Ba' Pham, ba.pham@theta.utoronto.ca, Li Ka Shing Knowledge Institute, St. Michael's Hospital, Unity Health Toronto, Toronto, Ontario, Canada

Alexandria Bennett, d.bennett@uottawa.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada. ORCID 0000-0002-5977-2094

Andrew Beck, andrew.beck@uottawa.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada. ORCID 0000-0002-8308-2202

Becky Skidmore, bskidmore@rogers.com, Independent Information Specialist, Ottawa, Ontario, Canada.

Maura R. Grossman, maura.grossman@uwaterloo.ca, University of Waterloo, Waterloo, Ontario, Canada.

Gordon V. Cormack, gvcormac@uwaterloo.ca, University of Waterloo, Waterloo, Ontario, Canada.

Sharmistha Mishra, Sharmistha.Mishra@toronto.ca, Department of Medicine, St. Michael's Hospital, University of Toronto, Toronto, Ontario, Canada;

MAP Centre for Urban Health Solutions, Li Ka Shing Knowledge Institute, Unity Health Toronto, Toronto, Ontario, Canada;

Epidemiology Division and Institute of Health Policy, Management, and Evaluation, Dalla Lana School of Public Health, University of Toronto, Toronto, Ontario, Canada;

Institute of Medical Science, University of Toronto, Toronto, Ontario, Canada. ORCID: 0000-0001-8492-5470

Adrienne Chan, adrienne.chan@sunnybrook.ca, Sunnybrook Health Sciences Centre, Toronto; Dalla Lana School of Public Health, University of Toronto, Toronto, Ontario, Canada.

Lan Xu, lan.xu@sjtu.edu.cn, School of Medicine, Shanghai Jiao Tong University, China.

David Moher, dmoher@ohri.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada.

Melissa Brouwers, Melissa.Brouwers@uottawa.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada.

Andrea C. Tricco, Andrea.Tricco@unityhealth.to, Li Ka Shing Knowledge Institute, St. Michael's Hospital, Unity Health Toronto, Toronto, Ontario, Canada; Epidemiology Division and Institute of Health Policy, Management, and Evaluation, Dalla Lana School of Public Health, University of Toronto, Toronto, Ontario, Canada; Queen's Collaboration for Health Care Quality Joanna Briggs Institute Centre of Excellence, Queen's University, Kingston, Ontario, Canada.

Julian Little, jlittle@uottawa.ca, Knowledge Synthesis and Application Unit, School of Epidemiology and Public Health, Faculty of Medicine, University of Ottawa, Ottawa, Ontario, Canada.

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Competing interests: DM was involved in the 2015 rapid review by Hersi et al. [1] There are no other competing interests to acknowledge.

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Key Question

KQ5: Should Health workers in direct contact and/or indirect contact to patients with Ebola Virus Disease (EVD) or Marburg virus disease cover head and neck skin and mucous membranes or just cover mucous membranes?

Methods Summary

This is one of a series of rapid reviews answering 12 key questions related to three themes on infection prevention and control measures for filoviruses: (i) transmission/exposure (n=3 questions), (ii) personal protective equipment (PPE) (n=5), and (iii) decontamination and disinfection (n=4). Data sources include Medline, Embase, bio/medRxiv pre-print servers, Global Medicus Index, Epistemonikos, China National Knowledge Infrastructure (CNKI) and Wangfang database. We used an automation tool (CAL® tool) for titles/abstracts screening for relevant systematic reviews and primary comparative studies. Full-text screening, data extraction, risk of bias assessment, and GRADE (Grading of Recommendations Assessment, Development and Evaluation) for the certainty of evidence were completed independently by two reviewers with any disagreements resolved by consensus, with arbitration by a third reviewer, when needed.

Initial findings

We present study characteristics in Table 1 and a summary of findings in Tables 2-4.

Initially, 137 studies were screened in the CAL tool software and 42 studies were included for fulltext screening. Four studies met the eligibility criteria and were included (Appendix 2). A list of excluded studies with reasons for exclusion can be found in Appendix 1.

No studies provided direct information on the transmission or incidence of EVD or Marburg virus disease related to the use of personal protective equipment (PPE) for head and neck skin protection. We included two simulation studies that addressed outcomes related to heat stress for health care workers (HCW) donning extra head/neck covering PPE (hoods). Additionally, we included two crossover randomized controlled trials that simulated contamination events for HCWs while doffing PPE ensembles with and without neck covering.

Overall, for heat tolerance outcomes, we found very low certainty evidence that PPE ensembles with additional head/neck covering increased both physiological and subjective measures of heat exhaustion, compared to PPE with no cover of the head and neck. We found low to very low certainty of evidence that PPE ensembles with head/neck covering resulted in less contamination than PPE with no cover for the head and neck. We found low to very low certainty evidence that PPE ensembles that covered the head/neck resulted in more human errors during donning/doffing of equipment, compared to ensembles without head/neck cover.

Table 1. Characteristics of Included Studies

Citation	Study	Funding	Virus Species	Setting	# Total	# Health	Description	Study
Yearl	Design	Source	species		Workers	Facilities	Worker	[as reported]
rearj					Workers	1 actitics	Care/contact	by study
							with patients	authors]
Coca, 2017 ¹	Non- randomized simulation study	Not reported	N/A	Simulated ambient conditions for West African countries ^a	6 healthy individuals to simulate HCWs	N/A; one environmental chamber	Exercise intensity was set to the average for nursing care ^b	Evaluate the human physiological and subjective responses to continuous light exercise within environmental conditions similar to those in West Africa while wearing 3 different, commonly used PPE ensembles.
Coca, 2015 ²	Non- randomized simulation study	Not reported	N/A	Simulated ambient conditions for warmest months in West African countries ^c	N/A; sweating thermal manikins	N/A; one environmental chamber	Metabolic work rate (work intensity) was set to the average for nursing care ^d	The focus of the present study was to provide a baseline heat stress analysis of some of the PPE

Suen, 2018 ³	Crossover randomized	Public university	Fluorescent solution ^e	Air- conditioned	59 HCWs (all	N/A; one air- conditioned	Fluorescent solution	ensemble options used in West Africa in the fight against the spread of Ebola. Compare the efficacy of
	controlled trial	funded	on the PPE surface to simulate Ebola virus	room with an average temperature of 23 °C \pm 2 °C and a relative humidity of 60% \pm 3%	evaluated in each of PPE ensembles)	room	sprayed on PPE at the length of a stethoscope to simulate usual working distance between a patient and an HCW ^f ; contamination events monitored during doffing	three PPE ensembles for routine patient care and performing aerosol- generating procedures to prevent EVD transmission by measuring the degree of contamination of HCWs and the environment.
Zamora, 2006 ⁴	Crossover randomized controlled trial	Physicians' Services Incorporated Foundation and the Clinical Teachers' Association	Fluorescent solution ^g on the PPE surface to simulate HID	Not reported	50 HCWs	N/A	Participants' front face shield, torso, hands, forearms and elbows were contaminated with	Examine the difference in self- contamination rates and the level of contact and droplet

	of Queen's University			fluorescent solution/ paste	protection- associated with E-RCP and the PAPR
					system.

Abbreviations: HCW, health care workers, HID, highly infectious diseases, NR, not reported, PPE, personal protective equipment

- a. For each testing protocol, three periods with different conditions were simulated: 15-minute pre-exercise stabilization period (22°C, 50% relative humidity) and a 60-minute exercise period (32°C, 92% relative humidity), followed by a 30-minute recovery period in ambient conditions (22°C, 50% relative humidity).
- b. The exercise protocol consisted of 60 minutes of continuous walking, within an environmental chamber, on a treadmill at an intensity of three METs (2.5 mph, 0% grade). This exercise intensity was chosen to represent the working intensity seen in hospital nurses during patient care, such as walking, standing, and carrying light objects.
- c. Two conditions were simulated. Condition A consisted of 32°C, 92% relative humidity, Condition B consisted of 26°C, 80% relative humidity
- d. Average work intensity for nursing corresponded to patient care that includes standing and walking slowly [2.5 mph] and carrying light objects [<11.3 kg]) of 3 METs (metabolic equivalent, or the measure of the intensity of aerobic exercise) over 80 min of continuous activity
- e. UV GERM Hygiene Spray, Glow Tec Ltd., London, England
- f. Three strokes of fluorescent solution were sprayed onto the face shield, two upper limb/ gloves and anterior surfaces of the gown at a distance of 60 cm from the participants (total 12 strokes per case). There was an average of 1.99 g fluorescent solution/per stroke.
- g. Fluorescein solution (1 mL of a 25% solution in 100 mL of sterile water). A Devilbiss atomizer (model DV15-RD, Sunrise Medical Products, Carlsbad, Calif.) was used to apply 5 mL of solution to each participant's front face shield and torso. "Invisible" Detection Paste (15 mL; Sirchie, Youngsville, NC) was applied from the forearms to the elbow and to the palmar aspects of participants' hands

Table 2. Summary of Findings: Heat Tolerance

Study	Intervention	Comparator	Mean (±	Mean (±	Pairwise	Quality	GRADE	Notes
details	(cover	(cover	SD) in	SD) in	comparison	Assessment ^a		
	head/neck and	mucous	intervention	comparator				
	mucus	membranes	group	group				
	membranes)	only)						
Time (min)	to reach critical co	re temperature	of 39°C under o	condition A ^b				
Coca, 2015 ²	E4 ^c	E2 ^d	62±6 min	78±7 min	P = 0.04	Moderate	$\oplus OOO$	None
	E3 ^e	$E2^d$	65±3 min	78±7 min	P = 0.04	risk of bias	Very low	
	E4 ^c	E1 ^f	62±6 min	+80 min	P < 0.05			
	E3 ^e	$\mathrm{E1^{f}}$	65±3 min	+80 min	P < 0.05			
Body surface	e skin temperature	• (°C) time to rea	ach critical cor	e temperature	of 39°C under	condition A ^b		
Coca, 2015 ²	E4 ^c	$E2^d$	38.4 ± 0.8	37.7 ± 0.2	NS	Moderate	$\oplus OOO$	None
	E3 ^e	$E2^d$	38.3 ± 0.2	37.7 ± 0.2	NS	risk of bias	Very low	
	E4 ^c	E1 ^f	38.4 ± 0.8	37.3 ± 0.3	P < 0.05			
	E3 ^e	E1 ^f	38.3 ± 0.2	37.3 ± 0.3	P < 0.05			
Heat sensati	ion ^g at time to reac	h critical core t	emperature of	39°C under co	ndition A ^b			
Coca, 2015 ²	E4 ^c	$E2^d$	3.8 ± 0.1	3.5 ± 0.2	NS	Moderate	$\oplus OOO$	None
	E3 ^e	$E2^d$	3.7 ± 0.1	3.5 ± 0.2	NS	risk of bias	Very low	
	E4 ^c	E1 ^f	3.8 ± 0.1	3.6 ± 0.2	NS			
	E3 ^e	E1 ^f	3.7 ± 0.1	3.6 ± 0.2	NS			
Discomfort ^h	at time to reach cr	ritical core temp	perature of 39°C	C under condit	tion A ^b			
Coca, 2015 ²	E4 ^c	$E2^d$	-3.4 ± 0.1	-3.2 ± 0.1	NS	Moderate	$\oplus OOO$	None
	E3 ^e	$E2^d$	-3.4 ± 0.1	-3.2 ± 0.1	NS	risk of bias	Very low	
	E4 ^c	E1 ^f	-3.4 ± 0.1	-3.2 ± 0.1	NS		-	
	E3 ^e	E1 ^f	-3.4 ± 0.1	-3.2 ± 0.1	NS			
Core temper	ature (°C) after 80	minutes of activ	vity under con	dition B ⁱ				
Coca, 2015 ²	E4 ^c	E2 ^d	38.9 ± 0.2	38.33 ± 0.1	P < 0.05	Moderate	$\oplus OOO$	None
	E3 ^e	$E2^d$	38.7 ± 0.1	38.33 ± 0.1	P < 0.05	risk of bias	Very low	
	E4 ^c	E1 ^f	38.9 ± 0.2	38.05 ± 0.1	P < 0.05		-	
	E3 ^e	E1 ^f	38.7 ± 0.1	38.05 ± 0.1	P < 0.05			
Body surface	e skin temperature	e (°C) after 80 m	inutes of activi	ty under cond	ition B ⁱ			

Study	Intervention	Comparator	Mean (±	Mean (±	Pairwise	Quality	GRADE	Notes
details	(cover	(cover	SD) in	SD) in	comparison	Assessment ^a		
	head/neck and	mucous	intervention	comparator				
	mucus	membranes	group	group				
	membranes)	only)						
Coca, 2015^2	E4 ^c	E2 ^d	37.6 ± 0.4	36.4 ± 0.4	P < 0.05	Moderate	$\oplus \bigcirc \bigcirc \bigcirc \bigcirc$	None
	E3 ^e	E2 ^d	36.9 ± 0.2	36.4 ± 0.4	NS	risk of bias	Very low	
	E4 ^c	E1 ^f	37.6 ± 0.4	35.8 ± 0.6	P < 0.05			
	E3 ^e	E1 ^f	36.9 ± 0.2	35.8 ± 0.6	NS			
Heat sensati	ion ^g after 80 minut	es of activity un	der condition	B ⁱ				
Coca, 2015 ²	E4 ^c	$E2^d$	3.2 ± 0.6	2.5 ± 0.6	NS	Moderate	$\oplus \bigcirc \bigcirc \bigcirc$	None
	E3 ^e	$E2^d$	2.5 ± 0.4	2.5 ± 0.6	NS	risk of bias	Very low	
	E4 ^c	$\mathrm{E1^{f}}$	3.2 ± 0.6	2.4 ± 0.5	P < 0.05			
	E3 ^e	E1 ^f	2.5 ± 0.4	2.4 ± 0.5	NS			
Discomfort ^h	after 80 minutes o	f activity under	condition B ⁱ					
Coca, 2015 ²	E4 ^c	$E2^d$	-3.2 ± 0.2	-2.6 ± 0.4	P < 0.05	Moderate	$\oplus \bigcirc \bigcirc \bigcirc$	None
	E3 ^e	$E2^d$	-3 ± 0.2	-2.6 ± 0.4	NS	risk of bias	Very low	
	E4 ^c	E1 ^f	-3.2 ± 0.2	-2.3 ± 0.3	P < 0.05			
	E3 ^e	E1 ^f	-3 ± 0.2	-2.3 ± 0.3	NS			
Core Tempe	rature (°C) at end	of exercise						
Coca, 2017 ¹	E3 ^j	$E1^k$	38.91 ± 0.29	38.18 ± 0.46	P < 0.05	High risk of	$\oplus \bigcirc \bigcirc \bigcirc$	None
	$E2^{l}$	$E1^k$	38.78 ± 0.36	38.18 ± 0.46	P < 0.05	bias	Very low	
Skin Tempe	rature (°C) at end	of exercise						
Coca, 2017 ¹	E3 ^j	$E1^k$	37.94 ± 0.15	36.12 ± 0.65	NS	High risk of	$\oplus \bigcirc \bigcirc \bigcirc$	None
	$E2^{l}$	$E1^k$	37.21 ± 0.21	36.12 ± 0.65	NS	bias	Very low	
Heart Rate (beats per minute)	at end of exerci	ise					
Coca, 2017 ¹	E3 ^j	$E1^k$	163 ± 17.52	135.57 ±	P < 0.05	High risk of	$\oplus \bigcirc \bigcirc \bigcirc$	None
				15.05		bias	Very low	
	$E2^{l}$	E1 ^k	156 ± 16.71	135.57 ±	P < 0.05			
				15.05				
Average swe	at weight loss (kg)) per hour						
Coca, 2017 ¹	E3 ^j	$E1^k$	1.48 ± 0.47	0.94 ± 0.40	P = 0.000	High risk of	$\oplus \bigcirc \bigcirc \bigcirc$	None
			kg	kg		bias		

Study	Intervention	Comparator	Mean (±	Mean (±	Pairwise	Quality	GRADE	Notes
details	(cover	(cover	SD) in	SD) in	comparison	Assessment ^a		
	head/neck and	mucous	intervention	comparator				
	mucus	membranes	group	group				
	membranes)	only)						
	$E2^{l}$	$E1^k$	1.26 ± 0.53	0.94 ± 0.40	P = 0.032		Very low	
			kg	kg				
Heat Sensat	ion ^g at end of exer	cise						
Coca, 2017 ¹	E3 ^j	$E1^k$	3.86 ± 0.38	3.29 ± 0.49	P < 0.05	High risk of	$\oplus O O O$	None
	$E2^{l}$	E1 ^k	3.86 ± 0.38	3.29 ± 0.49	P < 0.05	bias	Very low	
Thermal Co	mfort ^m at end of ex	kercise						
Coca, 2017 ¹	E3 ^j	$E1^k$	2.71 ± 2.56	2.71 ± 0.76	NS	High risk of	$\oplus OOO$	None
	$E2^{l}$	$E1^k$	3.57 ± 0.79	2.71 ± 0.76	P < 0.05	bias	Very low	
Rated perce	ived exertion ⁿ at er	nd of exercise					•	
Coca, 2017 ¹	E3 ^j	$E1^k$	15.29 ± 2.50	11.86 ± 2.12	P < 0.05	High risk of	$\oplus OOO$	None
	$E2^{l}$	$E1^k$	14.43 ± 3.10	11.86 ± 2.12	P < 0.05	bias	Very low	
Breathing co	omfort° at end of e	xercise					•	
Coca, 2017 ¹	E3 ^j	$E1^k$	5.14 ± 0.69	3.57 ± 1.27	P < 0.05	High risk of	$\oplus OOO$	None
	$E2^{l}$	E1 ^k	5.29 ± 1.11	3.57 ± 1.27	P < 0.05	bias	Very low	
Wetness ^p at	end of exercise							
Coca, 2017 ¹	E3 ^j	E1 ^k	2.86 ± 0.38	2.86 ± 0.38	NS	High risk of	000	None
	$E2^{l}$	$E1^k$	2.86 ± 0.38	2.86 ± 0.38	NS	bias	Very low	

Abbreviations: NS, non-significant (P>0.05); SD, standard deviation

a. Quality assessment of studies was completed using the ROBINS-I scale for observational/non-randomized studies. For the mannequin simulation study (Coca et al. 2015), quality assessment was performed under assumption that mannequin could be treated as a volunteer and humanized.

b. Condition A consisted of 32°C, 92% relative humidity

c. Ensemble 4 (E4): medical scrubs, socks, and rubber boots, impermeable coverall, Tyvek hood with an integrated splash-resistant surgical mask; rubber surgical apron, splash-resistant goggles, surgical nitrile inner gloves, heavy-duty nitrile outer gloves, N95 mask, a fluid-resistant surgical cap. The Tyvek hood provided the head and neck cover.

d. Ensemble 2 (E2): medical scrubs, socks, rubber boots with a mid-calf-length, disposable, fluid-resistant surgical gown, a polyethylene surgical apron, a face shield, disposable nitrile examination inner gloves, N95 mask, a fluid-resistant surgical cap. The cap provided some head covering, but the majority of head and neck skin remained exposed.

- e. Ensemble 3 (E3): medical scrubs, socks, rubber boots with a Tyvek coverall, Tyvek hood with an integrated splash-resistant surgical mask; a rubber surgical apron, splash-resistant goggles, surgical nitrile inner gloves; heavy-duty nitrile outer gloves, a duckbill N95 filtering face piece respirator, and a fluid-resistant surgical cap. The Tyvek hood provided the head and neck cover.
- f. Ensemble 1 (E1): medical scrubs, socks, rubber boots with a mid-calf-length, disposable, fluid-resistant surgical gown, a fluid-resistant 3-ply surgical mask, a disposable polyester lens face shield, disposable nitrile examination gloves. Head and neck skin was exposed.
- g. Heat sensation (rated from -4 [very cold] to 4 [very hot]
- h. Thermal comfort rated from -4 [very uncomfortable] to 4 [very comfortable])
- i. Condition B consisted of 26°C, 80% relative humidity
- j. Ensemble 3 (E3): medical scrubs, socks and rubber boots, Tychem QC highly impermeable coverall (DuPont), Médecins Sans Frontières (MSF) custom-made Tyvek (DuPont) hood with integrated splash-resistant surgical mask, rubber surgical apron, splashresistant goggles, surgical nitrile inner gloves, heavy-duty nitrile outer gloves, duckbill N95 filtering face piece respirator and fluidresistant surgical cap
- k. Ensemble 2 (E2): medical scrubs, socks and rubber boots, Microgard coverall, Tyvek hood with integrated splash-resistant surgical mask, rubber surgical apron, splash-resistant goggles, surgical nitrile inner gloves, heavy-duty nitrile outer gloves, duckbill N95 filtering face piece respirator, fluid-resistant surgical cap
- Ensemble 1 (E1): medical scrubs; socks and rubber boots; a midcalf-length disposable, fluid-resistant surgical gown, Performance Surgical Gown 7696C; polyethylene surgical apron, face shield, disposable nitrile examination inner gloves, duckbill N95 surgical filtering face piece respirator, and fluid-resistant surgical cap
- m. Subjective thermal comfort was measured on a scale of 1 to 4 (where 1 = not uncomfortable and 4 = very uncomfortable)
- n. Rate of perceived exertion was measured by using the OMNI 6-20 exertional scale
- o. Breathing comfort was measured by using a scale of 1 to 7 (where 1 = no discomfort and 7 = intolerable discomfort)
- p. Subjective wetness was measured by using a scale of 1 to 5 (where 1 = dry and 5 = soaked)

Table 3. Summar	v of Findings:	Contamination	during	doffing	of PPE

Study	Intervention	Comparator	Outcome in	Outcome	Statistical	Quality	GRADE	Notes
details	(cover	(cover mucous	intervention	in control	test	Assessment		
	head/neck and	membranes	group	group				
	mucus	only)						
0 11	membranes)				1 (1 1 2)			
Overall c	ontamination durin	g doffing of PPE: S	mall sized conta	minated pate	ches (< 1 cm ²), n	nedian		
Suen,	PPE1 [°] - Hospital	PPE3 ^c - HA	5.00	7.00	ANOVA:	Low risk of	$\Theta \Theta \bigcirc \bigcirc \bigcirc$	None
20183	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value $= 0.05$			
	PPE2 ^d -	PPE3 ^c - HA	7.00	7.00		Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
	DuPont TM	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Hair and	head contaminatio	on during doffing of	PPE: Small size	d contaminat	ed patches (< 1	cm²), median		
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	1.00	2.50	ANOVA:	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value $= 0.68$			
	PPE2 ^d -	PPE3 ^c - HA	2.00	2.50		Low risk of	$\Theta \Theta \bigcirc \bigcirc$	None
	DuPont TM	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Neck (an	terior) contaminati	ion during doffing o	f PPE: Small siz	ed contamina	ted patches (<	1 cm ²), median		
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	2.50	11.00	ANOVA:	Low risk of	$\Theta \Theta \bigcirc \bigcirc$	None
2018 ³	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.		2011	
	PPE set	1						

Study	Intervention	Comparator	Outcome in	Outcome	Statistical	Quality	GRADE	Notes
details	(cover	(cover mucous	intervention	in control	test	Assessment ^a		
	head/neck and	membranes	group	group				
	mucus	only)						
	membranes)							
		care and			PPE3 : p-			
		performing AGPs			value $= 0.095$			
	PPE2 ^d -	PPE3 ^c - HA	5.00	11.00		Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
	DuPont™	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Neck (po	osterior) contamina	tion during doffing	of PPE: Small si	zed contamin	nated patches (<	< 1 cm²), mediar	1	
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	2.00	18.50	ANOVA:	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value = 0.824			
	$PPE2^{d}$ -	PPE3 ^c - HA	1.00	18.50		Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
	DuPont™	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Overall c	ontamination durin	ng doffing of PPE: E	Extra large sized	contaminated	d patches (≥ 5ci	m²), median		
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	39.00	47.00	ANOVA:	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value =			
	PPE2 ^d -	PPE3 ^c - HA	43.00	47.00	< 0.001*	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
	DuPont™	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						

Study	Intervention	Comparator	Outcome in	Outcome	Statistical	Quality	GRADE	Notes
details	(cover	(cover mucous	intervention	in control	test	Assessment		
	head/neck and	membranes	group	group				
	mucus	only)						
	membranes)							
Hair and	head contaminatio	on during doffing of	'PPE: Extra larg	e sized conta	minated patche	s (≥ 5cm²), mec	lian	
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	0.00	0.00	ANOVA:	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value = N/A			
	PPE2 ^d -	PPE3 ^c - HA	17.00	0.00		Low risk of	$\Theta \Theta \bigcirc \bigcirc$	None
	DuPont™	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Neck (an	terior) contaminati	ion during doffing o	f PPE: Extra lar	ge sized cont	aminated patch	es (≥ 5cm²), me	edian	
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	0.00	24.00	ANOVA:	Low risk of	$\Theta \Theta \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value = N/A			
	PPE2 ^d -	PPE3 ^c - HA	0.00	24.00		Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
	DuPont™	isolation gown for				bias	Low	
	Tyvek®, Model	routine patient						
	1422A	care and						
		performing AGPs						
Neck (po	osterior) contamina	tion during doffing	of PPE: Extra la	rge sized con	taminated patch	hes (≥ 5 cm ²), m	edian	
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA	0.00	0.00	ANOVA:	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority	isolation gown for			PPE1 vs.	bias	Low	
	Standard Ebola	routine patient			PPE2 vs.			
	PPE set	care and			PPE3 : p-			
		performing AGPs			value = \dot{N}/A			

Study details	Intervention (cover head/neck and mucus membranes)	Comparator (cover mucous membranes only)	Outcome in intervention group	Outcome in control group	Statistical test	Quality Assessment ^a	GRADE	Notes
	PPE2 ^d - DuPont [™] Tyvek®, Model 1422A	PPE3 ^c - HA isolation gown for routine patient care and performing AGPs	0.00	0.00		Low risk of bias	⊕⊕⊖⊖ Low	None
Overall c	ontamination durin	ng doffing of PPE, a	ny size, n (%)					
Zamora, 2006 ⁴	PAPR ^e	E-RCP ^f	13 (26%)	48 (96%)	Mainland– Gart: p <0.001	Some concerns	⊕○○○ Very low	None
Face con	tamination during	doffing of PPE, any	size, n (%)		•			
Zamora, 2006 ⁴	PAPR ^e	E-RCP ^f	0	2 (4%)	Mainland– Gart: p=1	Some concerns	⊕○○○ Very low	None
Back of the	he head contamina	tion during doffing	of PPE, any size	e, n (%)	•			
Zamora, 2006 ⁴	PAPR ^e	E-RCP ^f	0	0	Mainland– Gart: undefined	Some concerns	⊕○○○ Very low	None
Neck (an	terior) contaminati	ion during doffing o	f PPE, any size,	n (%)				
Zamora, 2006 ⁴	PAPR ^e	E-RCP ^f	3 (6%)	48 (96%)	Mainland– Gart: p<0.001	Some concerns	⊕○○○ Very low	None
Zamora, 2006 ⁴	PAPR ^e	E-RCP ^f	1 (2%)	9 (18%)	Mainland– Gart: p=0.012	Some concerns	⊕OOO Very low	None

a. Quality assessment of studies was completed using the Cochrane RoB 2 for randomized trials.

b. Hospital Authority Standard Ebola PPE set (PPE 1): a neck-to-ankle overall with an overlying water-resistant gown double and long nitrate gloves, boots, hood, disposable face shield and N95 respirator. Order of doffing: gloves, gown, boots, hood, N95.

c. HA isolation gown for routine patient care and performing AGPs (PPE3): pure cotton surgical scrub suit, appropriate size gowns and gloves and the known best-fitted respirator model (3 M 1860, 1860s and 1870). Order of doffing: gloves, gown, full face shield, cap, N95 respirator.

- **d.** DuPont[™] Tyvek®, Model 1422A (PPE2): head-to-ankle overall with a zipper on the front. The whole outfit includes double gloves, boots, disposable face shield and an N95 respirator. A plastic apron was used to cover up the front zipper before use. Order of doffing: apron, hood, coverall/outer gloves, face shield, N95 respirator, boots, inner gloves.
- e. PAPR (powered air-purifying respirator): Tyvek hood, Bouffant hair cover, Economy impact goggle, Air-mate breathing tube, face-shield, HEPA filter unit, N95 mask any of several modes (8210, 1860s, PFR95, 7210, 695), Gloves (Non-latex, latex, latex surgical), Tyvek coveralls with hood, Tyvek boot covers, Astound impervious surgical gown
- f. E-RCP (Enhanced respiratory and contact precautions) contains a bouffant hair cover, economy impact goggle, face-shield, N95 mask any of several modes (8210, 1860s, PFR95, 7210, 695), gloves (Non-latex, latex), astound impervious surgical gown

Table 4. Summary of Findings: Human factors: Deviation rate (%) during donning and doffing of personal protective equipment

Study	Intervention (cover	Comparator (cover	Outcome in	Outcome	Quality	GRADE	Notes
details	head/neck and	mucous membranes	intervention	in control	Assessment ^a		
	mucus membranes)	only)	group	group			
Overall de	eviation rate (%) during	donning of PPE					
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA isolation	6.06	3.70	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	PPE2 ^d - DuPont TM	PPE3 ^c - HA isolation	6.00	3.70			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Deviation	rate (%) during donnin	ng of hood					
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA isolation	20.00	N/A	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	PPE2 ^d - DuPont TM	PPE3 ^c - HA isolation	3.33	N/A			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Deviation	rate (%) during donnin	ng of faceshield					
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA isolation	11.67	6.67	Low risk of	$\oplus \oplus \bigcirc \bigcirc$	None
2018^{3}	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	PPE2 ^d - DuPont TM	PPE3 ^c - HA isolation	15.00	6.67			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Overall de	eviation rate (%) during	doffing of PPE					

Study	Intervention (cover	Comparator (cover	Outcome in	Outcome	Quality	GRADE	Notes
details	head/neck and	mucous membranes	intervention	in control	Assessment ^a		
	mucus membranes)	only)	group	group			
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA isolation	2.95	3.52	Low risk of	$\Theta \Theta O O$	None
2018^{3}	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	$PPE2^{d} - DuPont^{TM}$	PPE3 ^c - HA isolation	9.48	3.52			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Deviation	rate (%) during doffing	g of hood	[]				
Suen,	PPE1 ^b - Hospital	PPE3 ^c - HA isolation	5.00	N/A	Low risk of	$\Theta \Theta O O$	None
20183	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	$PPE2^{a} - DuPont^{TM}$	PPE3 ^c - HA isolation	8.33	N/A			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Deviation	rate (%) during doffing	g of faceshield					
Suen,	PPE1 [°] - Hospital	PPE3 ^c - HA isolation	6.67	10.00	Low risk of	$\Theta \Theta \bigcirc \bigcirc \bigcirc$	None
20183	Authority Standard	gown for routine patient			bias	Low	
	Ebola PPE set	care and performing					
		AGPs					
	PPE2 ^d - DuPont TM	PPE3 ^c - HA isolation	11.67	10.00			
	Tyvek®, Model	gown for routine patient					
	1422A	care and performing					
		AGPs					
Total don	ning errors, n (%)				2		
Zamora,	PAPR ^e	E-RCP ^r	19 (38%)	2 (4%)	Some	$\oplus \bigcirc \bigcirc \bigcirc \bigcirc$	None
20064					concerns	Very low	
Total doff	ing errors, n (%)						

Study	Intervention (cover	Comparator (cover	Outcome in	Outcome	Quality	GRADE	Notes
details	head/neck and	mucous membranes	intervention	in control	Assessment ^a		
	mucus membranes)	only)	group	group			
Zamora,	$PAPR^{e}$	E-RCP ^f	6 (12%)	12 (24%)	Some	$\oplus \bigcirc \bigcirc \bigcirc$	None
2006^{4}					concerns	Very low	
Error in a	pplication of goggles d	uring donning, n (%)					
Zamora,	PAPR ^e	E-RCP ^f	2 (4%)	0	Some	$\oplus \bigcirc \bigcirc \bigcirc$	None
2006^{4}					concerns	Very low	
Failure to	zip up coveralls or put	hood over head during d	onning, n (%)				
Zamora,	PAPR ^e	E-RCP ^f	1 (2%)	N/A	Some	$\oplus \bigcirc \bigcirc \bigcirc$	None
2006^{4}					concerns	Very low	
Error in a	pplication of bouffant h	air-cover during donning	z, n (%)				
Zamora,	PAPR ^e	E-RCP ^f	N/A	1 (2%)	Some	$\oplus \bigcirc \bigcirc \bigcirc$	None
2006^{4}					concerns	Very low	
Error in re	emoval of face shield du	uring doffing, n (%)					
Zamora,	PAPR ^e	E-RCP ^f	N/A	1 (2%)	Some	$\oplus O O O$	None
20064					concerns	Very low	
Error in re	emoval of hair-cover du	ring doffing, n (%)				• • • • • •	
Zamora,	PAPR ^e	E-RCP ^f	N/A	2 (4%)	Some	$\oplus \bigcirc \bigcirc \bigcirc$	None
20064					concerns	Very low	

a. Quality assessment of studies was completed using the Cochrane RoB 2 for randomized trials.

b. Hospital Authority Standard Ebola PPE set (PPE 1): a neck-to-ankle overall with an overlying water-resistant gown double and long nitrate gloves, boots, hood, disposable face shield and N95 respirator. Order of doffing: gloves, gown, boots, hood, N95.

- c. HA isolation gown for routine patient care and performing AGPs (PPE3): pure cotton surgical scrub suit, appropriate size gowns and gloves and the known best-fitted respirator model (3 M 1860, 1860s and 1870). Order of doffing: gloves, gown, full face shield, cap, N95 respirator.
- **d.** DuPont[™] Tyvek®, Model 1422A (PPE2): head-to-ankle overall with a zipper on the front. The whole outfit includes double gloves, boots, disposable face shield and an N95 respirator. A plastic apron was used to cover up the front zipper before use. Order of doffing: apron, hood, coverall/outer gloves, face shield, N95 respirator, boots, inner gloves.
- e. PAPR (powered air-purifying respirator): Tyvek hood, Bouffant hair cover, Economy impact goggle, Air-mate breathing tube, faceshield, HEPA filter unit, N95 mask - any of several modes (8210, 1860s, PFR95, 7210, 695), Gloves (Non-latex, latex, latex surgical), Tyvek coveralls with hood, Tyvek boot covers, Astound impervious surgical gown

f. E-RCP (Enhanced respiratory and contact precautions) contains a bouffant hair cover, economy impact goggle, face-shield, N95 mask - any of several modes (8210, 1860s, PFR95, 7210, 695), gloves (Non-latex, latex), astound impervious surgical gown

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Appendix 1. Excluded Studies List – By Reason for Exclusion:

Abstract Only

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Commentary

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Narrative review

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Martin-Moreno JM, Llinás G, Martínez-Hernández J. Response to "MacIntyre et al., 2014: Respiratory protection for healthcare workers treating Ebola virus disease (EVD): are facemasks sufficient to meet occupational health and safety obligations?" *International Journal of Nursing Studies*. 2014;51(12):1693. doi:10.1016/j.ijnurstu.2014.10.005

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No relevant comparisons

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Doshi RH, Hoff NA, Mukadi P, et al. Seroprevalence of ebola virus among health care workers in the Tshuapa district democratic republic of congo. *Am J Tropic Med Hygiene*. Published online 2016.

Dunn AC, Walker TA, Redd J, et al. Nosocomial transmission of Ebola virus disease on pediatric and maternity wards: Bombali and Tonkolili, Sierra Leone, 2014. *American Journal of Infection Control.* 2016;44(3):269-272. doi:10.1016/j.ajic.2015.09.016

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Raj D, Hornsey E, Perl TM. Personal protective equipment for viral hemorrhagic fevers: *Current Opinion in Infectious Diseases.* 2019;32(4):337-347. doi:10.1097/QCO.000000000000562

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No relevant outcome data

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<mark>No PDF</mark>

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With strengthened guidelines for health care workers, the CDC ups its game against the deadly Ebola virus. *ED management : the monthly update on emergency department management.* 2014;26(12):133-136.

Appendix 2. Eligibility Criteria

Question (5): Should Health workers in direct contact and/or indirect contact to patients with Ebola Virus Disease (EVD) or Marburg virus disease cover head and neck skin and mucous membranes or just cover mucous membranes?

)	
Setting	Health care facilities, ETU, <u>community (e.g. burial teams)</u>
	*Contexts to consider: ETU use vs. healthcare facility;
	outbreak vs readiness vs. high alert scenario.
Population	Staff in HCF, ETU, community (e.g. burial teams)
Background interventions	The mucous membranes of eyes, mouth and nose are
(Standard of care)	covered by PPE. Use of a head cover that covers head and
[L] SEP	neck.
Intervention	Use a cover for the head and neck.
Comparator(s)	Not use a cover for head and neck.
	Direct contact, indirect contact
Outcome	Infection with Ebola or Marburg, PPE breaches, compliance
	related to heat and comfort, <u>dehydration, heat tolerance, human</u>
	factors, health worker confidence
	Indirect evidence: Lassa fever
Potential effect modifiers	Frequency and type of exposure, vaccination

Appendix 3. GRADE Assessment: Heat Tolerance

	_		Certainty a	ssessment			№ of p	atients	Effec	:t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% CI)	Absolute (95% Cl)	Certainty	Importance
Time (min)	to reach critical	core temperature	of 39°C under cor	dition A								
1	observational studies	not serious ^a	not serious	very serious ^b	serious	none	3	3	-	MD 16 min fewer (30.78 fewer to 1.22 fewer)		
									-	MD 13 min fewer (25.2 fewer to 0.79 fewer)		
									-	MD 18 min fewer (27.7 fewer to 8.2 fewer)		
									-	MD 15 min fewer (20.06 fewer to 9.9 fewer)		
Body surfa	ce skin temperat	ture (°C) at time to	reach critical cor	e temperature of 3	9°C under conditi	on A					-	
1	observational studies	not serious ^a	not serious	very serious ^b	serious⁰	none	3	3	-	MD 0.7 C higher (0.62 lower to 2.02 higher)		
									-	MD 0.6 C higher (0.14 higher to 1.05 higher)		
									-	MD 1.1 C higher (0.26 lower to 2.46 higher)		
									-	MD 1 C higher (0.42 higher to 1.57 higher)		
Heat sensa	tion at time to re	each critical core t	emperature of 39°	C under condition	Α							

1	observational studies	not serious ^a	not serious	very serious ^b	serious⁰	none	3	3	-	MD 0.3 higher (0.05 lower to 0.65 higher)		
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	_		Certainty a	assessment	_		Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
									-	MD 0.2 higher (0.15 lower to 0.55 higher)		
									-	MD 0.2 higher (0.15 lower to 0.55 higher)		
									-	MD 0.1 higher (0.25 lower to 0.45 higher)		

Discomfort at time to reach critical core temperature of 39°C under condition A

1	observational studies	not serious ^a	not serious	very serious ^b	serious℃	none	3	3	-	MD 0.2 lower (0.42 lower to 0.02 higher)	
									-	MD 0.2 lower (0.42 lower to 0.02 higher)	
									-	MD 0.2 lower (0.42 lower to 0.02 higher)	
									-	MD 0.2 lower (0.42 lower to 0.02 higher)	

Core temperature (°C) after 80 minutes of activity under condition B

1	observational studies	not serious ^a	not serious	very serious ^b	serious	none	3	3	-	MD 0.57 C higher (0.21 higher to 0.92 higher)	
									-	MD 0.37 C higher (0.14 higher to 0.59 higher)	
									-	MD 0.85 C higher (0.49 higher to 1.2 higher)	
									-	MD 0.65 C higher (0.42 higher to 0.87 higher)	

Body surface skin temperature (°C) after 80 minutes of activity under condition B

			Certainty a	ssessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
1	observational studies	not serious ^a	not serious	very serious ^b	serious	none	3	3	-	MD 1.2 C higher (0.29 higher to 2.1 higher)		
									-	MD 0.5 C higher (0.21 lower to 1.21 higher)		
									-	MD 1.8 C higher (0.64 higher to 2.95 higher)		
									-	MD 1.1 C higher (0.08 higher to 2.11 higher)		

Heat sensation after 80 minutes of activity under condition B

1	observational studies	not serious ^a	not serious	very serious ^b	serious∘	none	3	3	-	MD 0.7 higher (0.66 lower to 2.06 higher)	
									-	MD 0 higher (1.15 lower to 1.15 higher)	
									-	MD 0.8 higher (0.45 lower to 2.05 higher)	
									-	MD 0.1 higher (0.92 lower to 1.12 higher)	

Discomfort after 80 minutes of activity under condition B

1	observational studies	not serious ^a	not serious	very serious ^b	serious℃	none	3	3	-	MD 0.6 lower (1.31 lower to 0.11 higher)	
									-	MD 0.4 lower (1.11 lower to 0.31 higher)	
									-	MD 0.9 lower (1.47 lower to 0.32 lower)	

			Certainty a	assessment			Nº of p	patients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
									-	MD 0.7 lower (1.27 lower to 0.12 lower)		

Core Temperature (°C) at end of exercise

1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 0.73 C higher (0.14 lower to 1.6 higher)	
									-	MD 0.6 C higher (0.33 lower to 1.53 higher)	

Skin Temperature (°C) at end of exercise

1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 1.8 C higher (0.75 higher to 2.88 higher)	
									-	MD 1.09 C higher (0 higher to 2.18 higher)	

Heart Rate (beats per minute) at end of exercise

1	observational studies	serious₫	not serious	very serious ^b	serious	none	3	3	-	MD 27.43 BPM higher (9.59 lower to 64.45 higher)	
									-	MD 20.43 BPM higher (15.61 lower to 56.47 higher)	

Average sweat weight loss (kg) per hour

1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 0.54 kg higher (0.44 lower to 1.52 higher)	
									-	MD 0.32 kg higher (0.74 lower to 1.38 higher)	

Heat Sensation at end of exercise

	Certainty assessment							atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 0.57 higher (0.42 lower to 1.56 higher)		
									-	MD 0.57 higher (0.42 lower to 1.56 higher)		

Thermal Comfort at end of exercise

1	observational studies	serious ^d	not serious	very serious ^b	serious⁰	none	3	3	-	MD 0 (4.28 lower to 4.28 higher)	
									-	MD 0.86 higher (0.89 lower to 2.61 higher)	

Rated perceived exertion at end of exercise

1	observational studies	serious₫	not serious	very serious ^b	serious℃	none	3	3	-	MD 3.43 higher (1.82 lower to 8.68 higher)	
									-	MD 2.57 higher (3.45 lower to 8.59 higher)	

Breathing comfort at end of exercise

1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 1.57 higher (0.74 lower to 3.88 higher)	
									-	MD 1.72 higher (0.98 lower to 4.42 higher)	

Wetness at end of exercise

1	observational studies	serious ^d	not serious	very serious ^b	serious	none	3	3	-	MD 0 (0.86 lower to 0.86 higher)	
									-	MD 0 (0.86 lower to 0.86 higher)	

Cl: confidence interval; MD: mean difference

Explanations

a. Coca et al., 2015 was judged to be at moderate risk of bias. The mannequins were treated in the quality assessment, as if the mannequin were a volunteer. There was a lack of information reported for several ROBINS-I domains, including outcome measurement. There was no outcome assessor blinding, though outcomes were less vulnerable to bias, due to simulated nature of the study.

b. Downrated due to simulation study and non-human participants, as well as other differences in evaluated PPE equipment other than just head/neck cover vs. no cover.

c. Few participants and optimal information size (OIS) threshold not met.

d. We rated Coca et al., 2017, at a high risk of bias because of no demonstration of data availability for all the study participants and lack of blinding of the outcome assessor. Outcomes like thermal comfort, heat sensation, rating of perceived exertion, breathing comfort, and wetness were subjective measures which could potentially be more vulnerable to bias.

Appendix 4. GRADE Assessment: Contamination during doffing of PPE

		_	Certainty a	ssessment			№ of p	atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
Overall cor	tamination duri	ng doffing of PPE	: Small sized cont	aminated patches	(< 1 cm2), median							
1	randomised trials	not serious	not serious	serious⁵	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Hair and he	ad contaminati	on during doffing	of PPE: Small size	d contaminated p	atches (< 1 cm2), ı	nedian	L	I				
1	randomised trials	not serious	not serious	serious ^b	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Neck (anter	ior) contaminat	tion during doffing	of PPE: Small siz	ed contaminated	oatches (< 1 cm2),	median	•	•		•		
1	randomised trials	not serious	not serious	serious⁵	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Neck (posterior) contamination during doffing of PPE: Small sized contaminated patches (< 1 cm2), median												
1	randomised trials	not serious	not serious	serious⁵	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Overall cor	tamination duri	ing doffing of PPE	: Extra large sized	contaminated pat	ches (≥ 5cm2), me	edian	L	I				
1	randomised trials	not serious	not serious	serious ^b	serious ^a	none	59	59	not estimable	not estimable		
Hair and he	ad contaminati	on during doffing	of PPE: Extra larg	e sized contamina	ted patches (≥ 5cr	n2), median						
1	randomised trials	not serious	not serious	serious ^b	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Neck (anter	ior) contaminat	tion during doffing	of PPE: Extra larg	ge sized contamin	ated patches (≥ 5c	cm2), median						
1	randomised trials	not serious	not serious	serious⁵	serious ^a	none	59	59	not estimable	not estimable	$\bigoplus_{Low} \bigcirc \bigcirc$	
Neck (post	erior) contamina	ation during doffin	ng of PPE: Extra la	rge sized contami	nated patches (≥ 5	ocm2), median						
1	randomised trials	not serious	not serious	serious ^b	serious ^a	none	59	59	not estimable	not estimable		

Overall contamination during doffing of PPE, any size, n (%)

	Certainty assessment							atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
1	randomised trials	serious	not serious	serious₫	serious ^a	none	13/50 (26.0%)	48/50 (96.0%)	RR 0.27 (0.17 to 0.43)	701 fewer per 1,000 (from 797 fewer to 547 fewer)		

Face contamination during doffing of PPE, any size, n (%)

Back of the head contamination during doffing of PPE, any size, n (%)

1	randomised trials	serious	not serious	serious ^d	serious ^a	none	0/50 (0.0%)	0/50 (0.0%)	not estimable	not estimable		
											VCIYIOW	1

Neck (anterior) contamination during doffing of PPE, any size, n (%)

1	randomised trials	serious	not serious	serious ^d	serious ^a	none	3/50 (6.0%)	48/50 (96.0%)	RR 0.1200 (0.0378 to 0.3533)	845 fewer per 1,000 (from 924 fewer to 621	
										fewer)	l

Neck (posterior) contamination during doffing of PPE, any size, n (%)

CI: confidence interval; RR: risk ratio

Explanations

a. Few participants and optimal information size (OIS) threshold not met.

b. Downrated due to simulation study: Fluorescent contamination as a surrogate outcome for EVD/Marburg Virus Disease, other differences in evaluated PPE equipment other than just head/neck cover vs. no cover.

c. Downrated due to concerns with risk of bias. Unclear risk of bias for several domains, including allocation bias, blinding of participants, and unclear if all outcomes were reported.

d. Downsted due to simulation study: Fluorescent contamination as a surrogate outcome, other differences in evaluated PPE equipment other than just head/neck cover vs. no cover.

			Certainty a	ssessment			№ of patients		Effect			
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
Overall deviation rate (%) during donning of PPE												
1	randomised trials	not serious	not serious	serious ^a	serious	none	59	59	-	-	$\bigoplus_{Low} \bigcirc \bigcirc$	
Deviation ra	te (%) during de	onning of hood										
1	randomised trials	not serious	not serious	serious ^a	serious⁵	none	59	59	-	-	$\bigoplus_{Low} \bigcirc \bigcirc$	
Deviation ra	te (%) during de	onning of faceshie	eld									
1	randomised trials	not serious	not serious	seriousa	serious⁵	none	59	59	-	-	$\bigoplus_{Low} \bigcirc \bigcirc$	
Overall devi	ation rate (%) d	uring donning of	PPE									
1	randomised trials	not serious	not serious	seriousa	serious⁵	none	59	59	-	-	$\bigoplus_{Low} \bigcirc \bigcirc$	
Deviation ra	te (%) during de	offing of hood										
1	randomised trials	not serious	not serious	seriousa	serious⁵	none	59	59	-	-	$\bigoplus_{Low} \bigcirc \bigcirc$	
Deviation ra	te (%) during de	onning of faceshie	əld				•	•		••		
1	randomised trials	not serious	not serious	serious ^a	serious⁵	none	59	59	-	-		
Total donni	ng errors, n (%)											
1	randomised trials	serious	not serious	Serious₫	serious ^b	none	19/50 (38.0%)	2/50 (4.0%)	RR 9.50 (2.33 to 38.70)	340 more per 1,000 (from 53 more to 1,000 more)		
Total doffing errors, n (%)												
1	randomised trials	serious	not serious	serious₫	serious ^b	none	6/50 (12.0%)	12/50 (24.0%)	RR 0.42 (0.17 to 1.03)	139 fewer per 1,000 (from 199 fewer to 7 more)		
Error in app	lication of gogg	les during donnir	ng, n (%)									
1	randomised trials	serious	not serious	serious ^d	serious⁵	none	2/50 (4.0%)	0/50 (0.0%)	RR 5.00 (0.25 to 101.60)	0 fewer per 1,000 (from 0 fewer to 0 fewer)		

Appendix 5. GRADE Assessment: Deviation rate (%) during donning and doffing of personal protective equipment

			Certainty a	issessment			№ of p	№ of patients		t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	A cover for the head and neck	No cover for the head and neck	Relative (95% Cl)	Absolute (95% Cl)	Certainty	Importance
Failure to zi	Failure to zip up coveralls or put hood over head during donning, n (%)											
1	randomised trials	serious∝	not serious	seriousd	serious ^b	none	1/50 (2.0%)	N/A	not estimable	not estimable		
Error in app	Error in application of bouffant hair-cover during donning, n (%)											
1	randomised trials	serious∝	not serious	serious ^d	serious	none	N/A	1/50 (2.0%)	not estimable	not estimable		
Error in rem	Error in removal of face shield during doffing, n (%)											
1	randomised trials	serious∝	not serious	seriousd	serious	none	N/A	1/50 (2.0%)	not estimable	not estimable		
Error in rem	Error in removal of hair-cover during doffing, n (%)											
1	randomised trials	serious	not serious	seriousd	serious ^b	none	N/A	2/50 (4.0%)	not estimable	not estimable		

Cl: confidence interval; RR: risk ratio

Explanations

a. Downrated due to simulation study: Fluorescent contamination as a surrogate outcome for EVD/Marburg Virus Disease, other differences in evaluated PPE equipment other than just head/neck cover vs. no cover.
b. Few participants and optimal information size (OIS) threshold not met.
c. Downrated due to concerns with risk of bias. Unclear risk of bias for several domains, including allocation bias, blinding of participants, and unclear if all outcomes were reported.
d. Downrated due to simulation study: Fluorescent contamination as a surrogate outcome, other differences in evaluated PPE equipment other than just head/neck cover vs. no cover.