

IPC Moments that Matter: Fundamentals for Impact

Professor Brett Mitchell & Amy Cartwright 16th October 2023





Moments that matter

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Acknowledgement of Country

I wish to acknowledge the Traditional Custodians of country throughout Australia and their connections to land, sea and community.

Pay my respect to their Elders past and present and extend that respect to all Aboriginal and Torres Strait Islander peoples today.

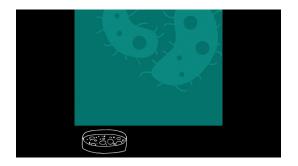


Declarations

Received no funding for this talk

 Recipient of grant funding from government and professional bodies through a competitive process

Start with story that some people may find upsetting



Moments that matter





#onesmallactofkindess

Kathy Koschel



Overview: Moments that matter

• Standard precautions

Air, skin and hands

Standard Precautions

Always follow these standard precautions



Perform hand hygiene before and after every patient contact



Clean and reprocess shared patient equipment



Use personal protective equipment when risk of body fluid exposure



Follow respiratory hygiene and cough etiquette



Use and dispose of sharps safely



Use aseptic technique

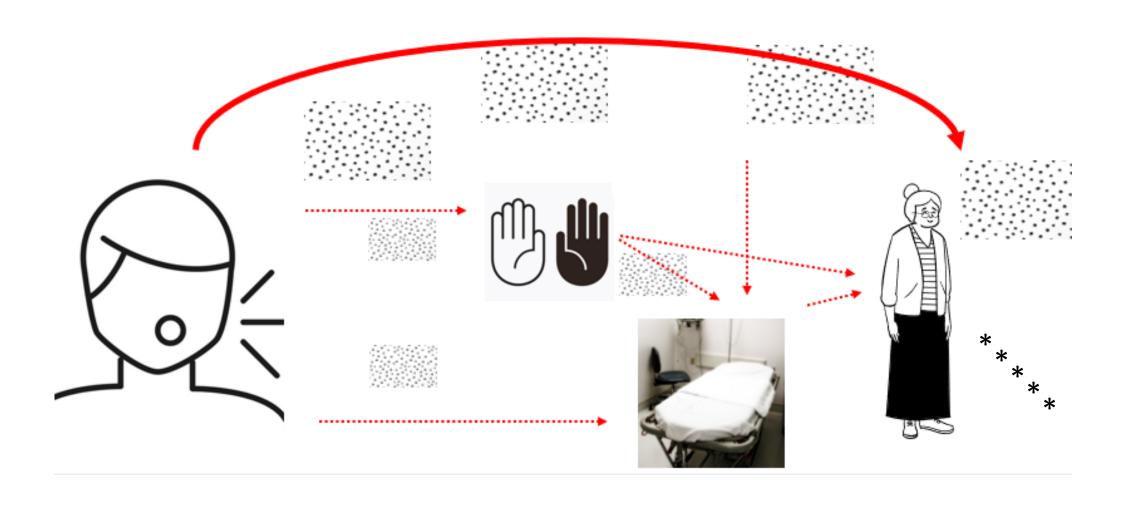


Perform routine environmental cleaning



Handle and dispose of waste and used linen safely

Why skin, surface and air?



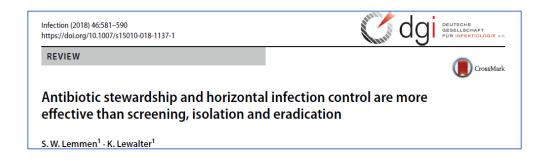
Horizontal approaches to IPC

Antibiotic stewardship

Antiseptic washing

Hand hygiene

Environment





Surface

 Pathogens can survive in the environment

 Increased risk of acquiring the sa pathogen from a prior room occi

 Improving routine and discharge cleaning has been shown to redu the risk of HAIs in RCTs.

ype of bacterium	Duration of persistence (range)	Reference(s) [18, 25, 28, 29, 87, 88]	
Acinetobacter spp.	3 days to 5 months		
Bordetella pertussis	3 – 5 days	[89, 90]	
Campylobacter jejuni	up to 6 days	[91]	
Clostridium difficile (spores)	5 months	[92–94]	
Chlamydia pneumoniae, C. trachomatis	≤ 30 hours	[14, 95]	
Chlamydia psittaci	15 days	[90]	
Corynebacterium diphtheriae	7 days – 6 months	[90, 96]	
Corynebacterium pseudotuberculosis	I–8 days	[21]	
Escherichia coli	1.5 hours – 16 months	[12, 16, 17, 22, 28, 52, 90, 97–99]	
Enterococcus spp. including VRE and VSE	5 days – 4 months	[9, 26, 28, 100, 101]	
Haemophilus influenzae	12 days	[90]	
Helicobacter pylori	≤ 90 minutes	[23]	
Klebsiella spp.	2 hours to > 30 months	[12, 16, 28, 52, 90]	
Listeria spp.	I day – months	[15, 90, 102]	
Mycobacterium bovis	> 2 months	[13, 90]	
Mycobacterium tuberculosis	I day – 4 months	[30, 90]	
Neisseria gonorrhoeae	I – 3 days	[24, 27, 90]	
Proteus vulgaris	I – 2 days	[90]	
Pseudomonas aeruginosa	6 hours - 16 months; on dry floor: 5 weeks	[12, 16, 28, 52, 99, 103, 104]	
Salmonella typhi	6 hours – 4 weeks	[90]	
Salmonella typhimurium	10 days – 4.2 years	[15, 90, 105]	
Salmonella spp.	I day	[52]	
Serratia marcescens	3 days - 2 months; on dry floor: 5 weeks	[12, 90]	
Shigella spp.	2 days – 5 months	[90, 106, 107]	
Staphylococcus aureus, including MRSA	7 days – 7 months	[9, 10, 16, 52, 99, 108]	
Streptococcus pneumoniae	I – 20 days	[90]	
Streptococcus pyogenes	3 days - 6.5 months	[90]	
Vibrio cholerae	I – 7 days	[90, 109]	

- Kramer, K et al (2006). BMC Infectious Diseases, 130
- Mitchell, BG et al (2023). Infection Disease and Health
- Anderson, D et al (2017), 389(10071):805-814
- Mitchell BG et al (2019). Lancet Infectious Disease, 19 (4) 410-18

Surface

 Pathogens can survive in the environment

• Increased risk of acquiring the same pathogen from a prior room occupant

• Improving routine and discharge cleaning has been shown to reduce the risk of HAIs in RCTs.

Study or Subgroup	Experimental (4		Control (-v		Moint	Odds Ratio	Odds Ratio
	Events	Total	Events	rotal	vveignt	M-H, Random, 95% CI	M-H, Random, 95% CI
I.1.1 MRSA		4405-	20-	00005	246	0.04 10 40	
Anderson	103	11005	725	293386	7.1%	3.81 [3.10, 4.69]	-
luang	57	1454	248	8697	7.0%	1.39 [1.04, 1.86]	
fitchell	74	884	163	5344	7.0%	2.90 [2.18, 3.86]	
Subtotal (95% CI)		13343		307427	21.1%	2.50 [1.38, 4.54]	_
otal events	234		1136				
Heterogeneity: Tau² = 1 Test for overall effect: 2			< 0.00001)	; P= 94%			
I.1.2 VRE							
Anderson	89	4083	423	307241	7.1%	16.16 [12.83, 20.36]	-
orees	19	138	31	500	6.4%	2.42 [1.32, 4.43]	
ord	47	149	89	300	6.8%	1.09 [0.71, 1.67]	
luang	58	1291	256	9058	7.0%	1.62 [1.21, 2.16]	-
hou	69	3556	92	4929	7.0%	1.04 [0.76, 1.43]	
Subtotal (95% CI)	0.5	9217	52	322028	34.3%	2.36 [0.61, 9.15]	
otal events	282	0211	891	OLLULO	0.110.10	Lieu [eie ij ei ie]	
Heterogeneity: Tau ² = 1 est for overall effect: 2	2.35; Chi ² = 329.); l²= 99%			
.1.3 ESBL							
Iseir	8	50 50	50	461 461	5.9%	1.57 [0.70, 3.52]	
Subtotal (95% CI)		50		461	5.9%	1.57 [0.70, 3.52]	
otal events	8		50				
leterogeneity: Not app est for overall effect: 2		3)					
.1.4 Klebsiella sp. or							100,200
jao	32	648	235	8723	6.9%	1.88 [1.29, 2.74]	
Subtotal (95% CI)	227	648	00000	8723	6.9%	1.88 [1.29, 2.74]	-
otal events leterogeneity: Not app			235				
Test for overall effect: 2	Z = 3.26 (P = 0.00	01)					
.1.5 Clostridioides di							
Inderson	43	3797	1278	307890	7.0%	2.75 [2.02, 3.73]	
haughnessy	10	91	77	1679	6.2%	2.57 [1.28, 5.15]	
Subtotal (95% CI)		3888		309569	13.2%	2.72 [2.05, 3.60]	•
fotal events Heterogeneity: Tau² = 1	53 0.00; Chi² = 0.03	df=1 (P:	1355 = 0.86); F=	0%			
est for overall effect: 2	Z = 7.01 (P < 0.00	0001)					
.1.6 Acinetobacter							
Iseir	16	52	41	459	6.3%	4.53 [2.32, 8.86]	
Subtotal (95% CI)	856	52	- 55	459	6.3%	4.53 [2.32, 8.86]	•
otal events	16		41				
leterogeneity: Not app	olicable						
est for overall effect.	Z = 4.42 (P < 0.00	001)					
.1.7 Pseudomonas				426	6.5%	1.96 [1.12, 3.45]	
Vseir	21	85	61				_
.1.7 Pseudomonas Jseir Subtotal (95% CI)	21	85 85		426	6.5%	1.96 [1.12, 3.45]	-
Vseir	21 21		61				•
lseir Subtotal (95% CI) Total events Heterogeneity: Not app	21 olicable	85					•
lseir Subtotal (95% CI) Otal events Heterogeneity: Not app	21 olicable	85					•
lseir Subtotal (95% CI)	21 blicable Z = 2.35 (P = 0.02	85	61	426	6.5%	1.96 [1.12, 3.45]	
lseir Subtotal (95% CI) Total events Heterogeneity: Not app Test for overall effect: 2 L.1.8 Norovirus Traenkel	21 olicable	85 2) 1016		426 32772	6.5% 5.7%	1.96 [1.12, 3.45]	-
dseir Subtotal (95% CI) Fotal events Heterogeneity: Not app est for overall effect: 2 .1.8 Norovirus Fraenkel Subtotal (95% CI)	21 Dilicable Z = 2.35 (P = 0.02	85	61	426	6.5%	1.96 [1.12, 3.45]	
Iseir Subtotal (95% CI) Total events Heterogeneity: Not applest for overall effect: 2 .1.8 Nor ovirus Traenkel Subtotal (95% CI) Total events	21 Discable Z = 2.35 (P = 0.02 5	85 2) 1016	61	426 32772	6.5% 5.7%	1.96 [1.12, 3.45]	
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Iseir unbtotal (95% CI) otal events eleterogeneity. Not appest for overall effect: 2.1.8 Norovirus raenkel unbtotal (95% CI) otal events eleterogeneity. Not appest for overall effect: 2 otal (95% CI)	21 olicable Z = 2.35 (P = 0.00 5 5 olicable Z = 2.54 (P = 0.01	85 1016 101 6	61 49 49	426 32772	6.5% 5.7% 5.7%	1.96 [1.12, 3.45]	-
dseir Subtotal (95% CI) Fotal events Heterogeneity: Not app est for overall effect: 2 .1.8 Norovirus Fraenkel Subtotal (95% CI)	21 olicable Z = 2.35 (P = 0.00 5 olicable Z = 2.54 (P = 0.01 651 0.81; Chi² = 357.	85 2) 1016 1016 1) 28299 84, df = 14	61 49 49 3818	32772 32772 32772	6.5% 5.7% 5.7%	1.96 [1.12, 3.45] 3.30 [1.31, 8.31] 3.30 [1.31, 8.31]	005 02 1 5 2

- Kramer, K et al (2006). BMC Infectious Diseases, 130
- Mitchell, BG et al (2023). Infection Disease and Health
- Anderson, D et al (2017), 389(10071):805-814
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Riskiest room





Nurses' and midwives' cleaning knowledge, attitudes and practices: An Australian study

Brett G. Mitchell ^{a,b,*}, Philip L. Russo ^{c,d}, Martin Kiernan ^{a,e}, Cassie Curryer ^a







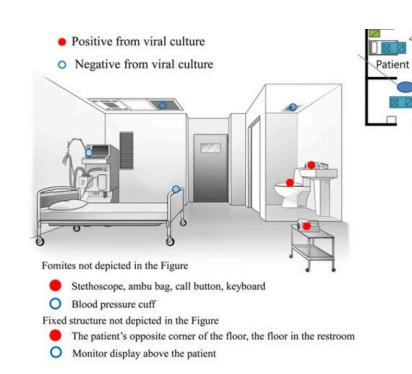
В

96%: Room A

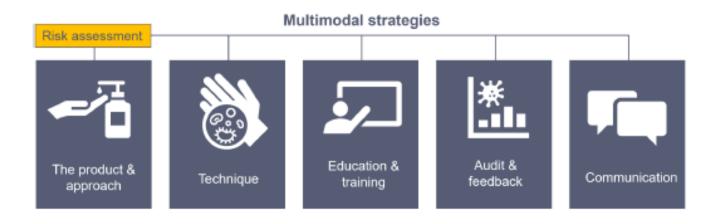
3%: Don't know

Surface

- Several modes of transmission dynamics, most commonly direct contact or aerosolization were identified
 - Even fast walking
 - Rashid, T., et al (2017) Epidemiol Infect 145(2): 347-357.
- Bhalla, A. (2004). "Acquisition of nosocomial pathogens on hands after contact with environmental surfaces near hospitalized patients." <u>Infect Control Hosp Epidemiol</u> 25(2): 164-167.
- Extensive viable MERS-CoV contamination of the air and surrounding materials in MERS outbreak units



Cleaning: considerations

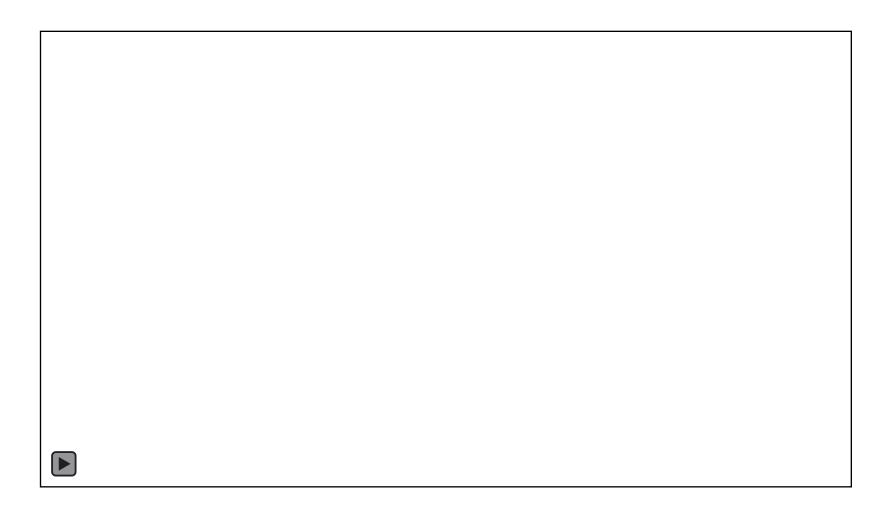


Product and approach

- Health and safety
- Preparation
- Contact time
- Reprocessing

- Storage
- Compatibility
- Efficacy
- Transferability of pathogens
- Practical considerations

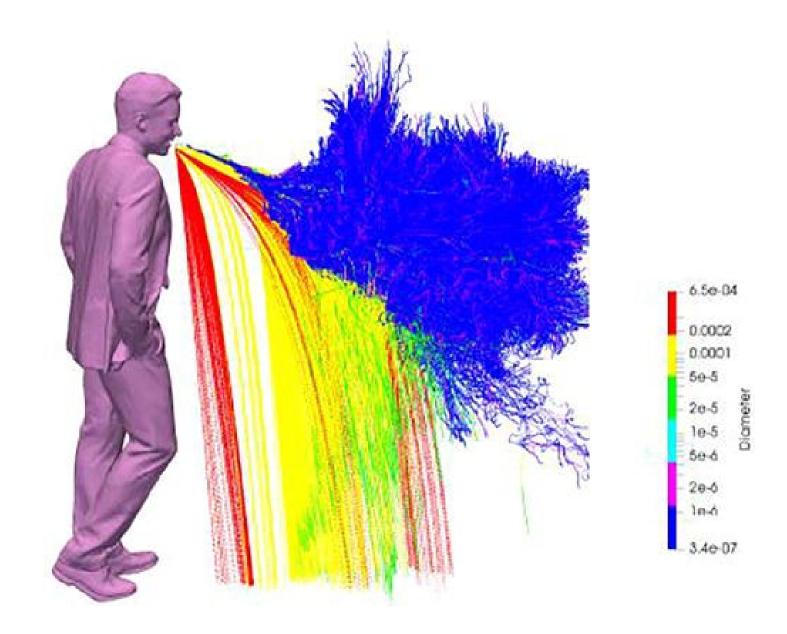
Air



Phase 1 Phase 2 Phase 3 Generation and exhalation Transport Inhalation, deposition and infection · Generation mechanisms · Size distribution of inhalable aerosols Settling velocity and residence time in air · Size change during transport Viral load at generation sites Deposition mechanisms · Size distribution of exhaled aerosols · Persistence of viruses in aerosols Size-dependent deposition sites Number of virions in aerosol · Environmental factors: temperature, humidity, · Deposition site susceptibility airflow and ventilation, UV radiation · <5 µm •100-5 µm Oral Laryngeal Bronchial Bronchiolar Alveolar

Fig. 1. Airborne transmission of respiratory viruses.

Phases involved in the airborne transmission of virus-laden aerosols include (i) generation and exhalation; (ii) transport; and (iii) inhalation, deposition, and infection. Each phase is influenced by a combination of aerodynamic, anatomical, and environmental factors. (The sizes of virus-containing aerosols are not to scale.)



Trivedi, S et al (2021). *Physics of Fluids*, 33(11), 115130.

Improving air to reduce the risk of infection

Aerosol transmission risk

Pathogens attached to aerosols – transmission route

Ventilation – changing air – bringing in new area

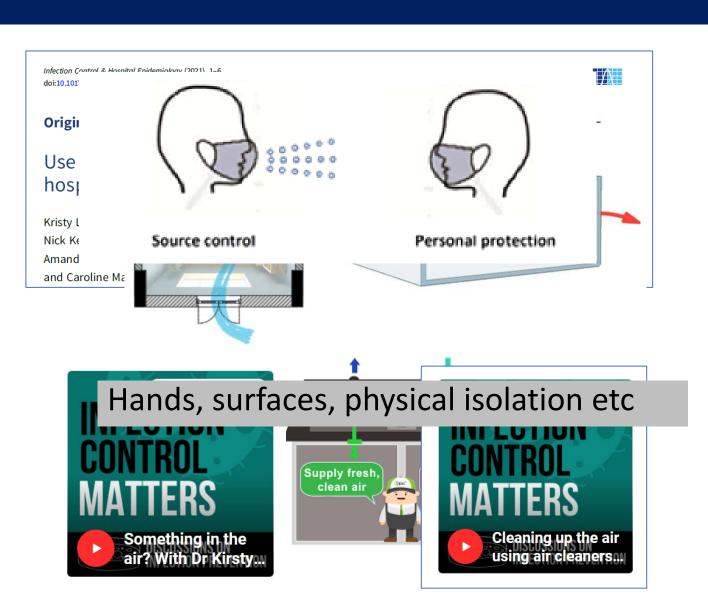
Air cleaning – remove particles from (human) from the air

Improving air to reduce the risk of infection

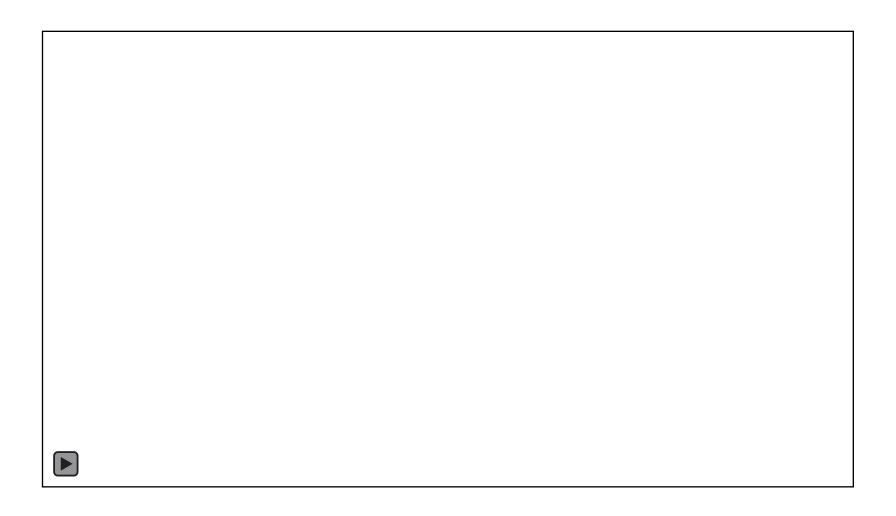
- Improving clean air exchange
 - Natural
 - Mechanical

Air scrubber / purifiers

Source control

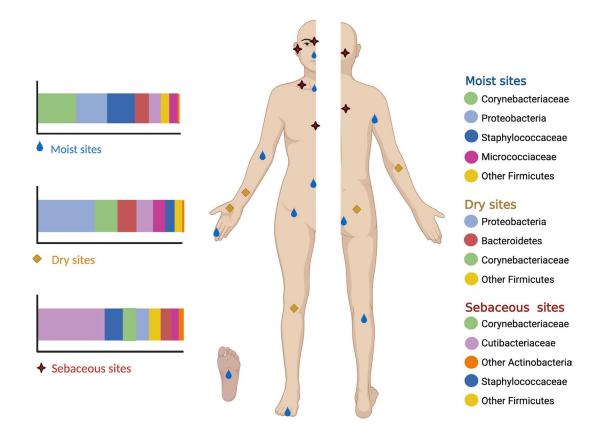


Skin



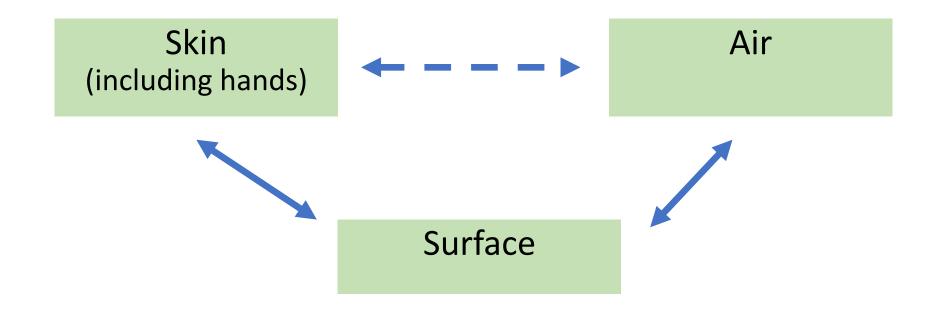
Skin

- Protective interface between internal organs and the environment
- The skin encounters a host of toxins, pathogenic organisms, and physical stresses.
- Skin functions as more than a physical barrier: it is an active immune organ.



Skin's role in transmission

- Skin role in both
 - Exogenous (from others)
 - Endogenous (from yourself)



Skin



Skin antisepsis















Skin: evidence

Hand hygiene

• Pittet, D., et al. (2000). Effectiveness of a hospital-wide programme to improve compliance with hand hygiene. *The Lancet*, *356*(9238), 1307-1312.

Peripheral vascular device insertion and management

- Webster, J et al (2019). Clinically-indicated replacement versus routine replacement of peripheral venous catheters. *Cochrane Database of Systematic Reviews*, (1).
- Rickard, C. et al. (2018). Dressings and securements for the prevention of peripheral intravenous catheter failure in adults (SAVE): a pragmatic, randomised controlled, superiority trial. *The Lancet*, 392(10145), 419-430.



- Assess the need
- Inform and partner with patients
- Ensure competency
- Right insertion site
- Maximise success of first insertion
- Insert and secure
- Document decisions and care
- Routine use: inspect, access
- Review ongoing need
- Remove safely and replace if needed

(Australian Commission on Safety and Quality in Healthcare, 2021)

Surgical site infection

- Seidelman, et al (2023). Surgical site infection prevention: a review. *JAMA*, *329*(3), 244-252.
- Tanner, J. et al. (2021). Preoperative hair removal to reduce surgical site infection. *Cochrane database of systematic reviews*, (8).
- Chen, S., et al (2020). Preoperative antisepsis with chlorhexidine versus povidone-iodine for the prevention of surgical site infection: a systematic review and meta-analysis. World Journal of Surgery, 44, 1412-1424.

Chlorhexidine bathing

- Climo et al (2013). Effect of daily chlorhexidine bathing on hospital-acquired infection. New England Journal of Medicine, 368(6), 533-542.
- Musuuza, J. S., et al (2019). The impact of chlorhexidine bathing on hospital-acquired bloodstream infections: a systematic review and metaanalysis. BMC infectious diseases, 19(1), 1-10.
- Lewis, S. et al (2019). Chlorhexidine bathing of the critically ill for the prevention of hospital-acquired infection. *Cochrane Database of Systematic Reviews*, (8).

- Decolonisation
- Hair removal
- Antibiotic prophylaxis
- Temperature and glucose management
- Asepsis
- Skin preparation
- Oxygenation

Central line associated infections

- Buetti, N., et al (2022). Strategies to prevent central line-associated bloodstream infections in acute-care hospitals: 2022 Update. *Infection Control & Hospital Epidemiology*, 43(5), 553-569.
- O'Grady, N. P. (2023). Prevention of Central Line—Associated Bloodstream Infections. *New England Journal of Medicine*, 389(12), 1121-1131.
- Ullman et al (2015). Dressings and securement devices for central venous catheters (CVC). *Cochrane Database of Systematic Reviews*, (9).

Urinary catheters

- Meddings, et al (2010). Systematic review and metaanalysis: reminder systems to reduce catheterassociated urinary tract infections and urinary catheter use in hospitalized patients. Clinical Infectious Diseases, 51(5), 550-560.
- Fasugba, O., et al (2019). Chlorhexidine for meatal cleaning in reducing catheter-associated urinary tract infections: a multicentre stepped-wedge randomised controlled trial. *The Lancet Infectious Diseases*, 19(6), 611-619.

Some examples

Some examples

- Insert only if required
- Meatal cleaning prior to insertion
- Aseptic technique
- Prompt removal
- Manual decontamination of hubs

Pneumonia prevention

- Mitchell, et al (2019). Strategies to reduce nonventilator-associated hospital-acquired pneumonia: a systematic review. *Infection, disease & health,* 24(4), 229-239.
- Wolfensberger, et al (2023). Prevention of nonventilator-associated hospital-acquired pneumonia in Switzerland: a type 2 hybrid effectiveness implementation trial. The Lancet Infectious Diseases.

 Improving quality and frequency of oral care

Dysphagia identification and management

Mobilisation

"We do not remember days, we remember moments." - Cesare Pavese, 1908-1950



Clean hospitals day – 20th October





Moments that matter

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Educational tools to build the fundamentals of IPC in daily practice.



Amy Cartwright, Clinical Educator





Clinical education

- Clinical education team of clinical staff
- Examples of the types of education and training we facilitate
- Full product roll out training
- Refresher training on current products
- Targeted training in areas needing extra support
- Study day presentations
- Production of educational materials



Hospital simulation

- As a clinical education and training team we regularly make use of our hospital simulation suite for training events both internally and externally.
- This is an opportunity to keep our skills up to date as well as help educate our internal teams.







Training

What can our training look like:

- Instructions for use
- Principles of cleaning
- 5 Moments for environmental decontamination
- Specific equipment training
- Product storage and dispensers



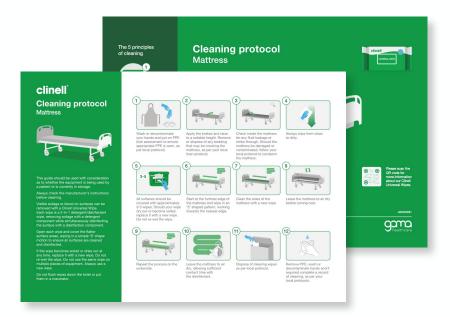




Training case study

Face-to-face training session on environmental decontamination:

- Sessions can be initiated either by the facility (usually the IPC team) or proactively by the clinical education team.
- Dates for the training, products to be covered, and any literature to be left with areas and key messages will all be discussed and planned beforehand.
- On day 1 of training a meeting with the organiser/IPC is always ideal to reiterate the key messages we will be sharing, be made aware of the areas we are to visit and if there are any PPE considerations in any departments (such as masks)





Case study - tips and ideas

- The main objectives of the sessions will include:
- Educating users on the appropriate use of our products for surface disinfection
- Discussing the appropriate applications and any limitations of the products,
- Demonstrating the proper techniques for use
- Educating users on the 5 principles of cleaning and how to apply them with using our products
- Helping users to understand the importance of surface decontamination
- Providing users with the knowledge and skills to use the products safely and effectively
- Supports ongoing Infection Prevention and Control training



Case study - tips and ideas

- Training usually takes place in the clinical department and can be done multiple times to capture as many staff as possible with little disruption to their jobs.
- All staff are asked to complete a register which are given to the organisations contact (usually IPC) who we have been liaising with at the end of the training along with a summary of the training.
- This kind of training is usually carried out over a few days to enable us to capture as many areas as possible.



Benefits and challenges of face-to-face training

Benefits:

- Ward/department based
- Quick/not too time-consuming
- Key essential points covered
- Good attendance
- Department-specific queries responded to
- Takes some pressure off IPC teams to carry out training

Challenges:

- Can still be difficult for staff to be released
- Only have time to deliver quick messages

Thank you.

