

What's new in HEH innovation: special focus on room disinfection

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Automated Whole Room Disinfection

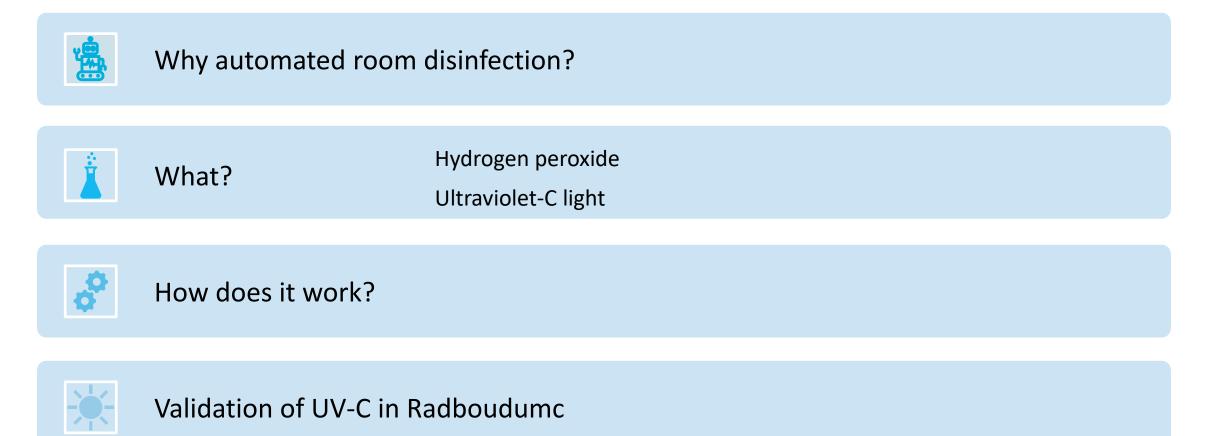
Why, What and How

Interclean Amsterdam 16-5-2024 Edmée Bowles, clinical microbiologist Radboudumc, Nijmegen

No disclosures



Content



Why automated room disinfection?

- Europe: 4.100.000 healthcare associated infections (HAI)/year*
- 90.000 deaths/year*
- Growing problem of antimicrobial resistance.



 Patients have an odds ratio of 2.45 to get infected with the microbes of the previous occupant of their room**

<u>https://www.ecdc.europa.eu/en/healthcare-associated-infections</u>
 Risk of organism acquisition from prior room occupant -Mitchell et al Inf dis Health 2023

Transmission through hospital environment

Mean length of stay 6 days **

One *Klebsiella pneumonia* in a room could infect up to 100 new patients

 Cleaning & disinfection is important weapon against AMR and HAI

Micro-organism **	Survival time
Staphylococcus aureus	<1min- 318 days
Klebsiella pneumoniae	0.57 - 600 days
Pseudomonas spp.	0.08 – 7 days

**Radboudumc 2022

**Porter et al- J. Hosp. Inf. 2024 https://doi.org/10.1016/j.jhin.2024.01.023

Cleaning & disinfection is our weapon against AMR



Manual mechanical cleaning and disinfection:

Labour intensive

Physically demanding

Error prone

Quality may vary during the day



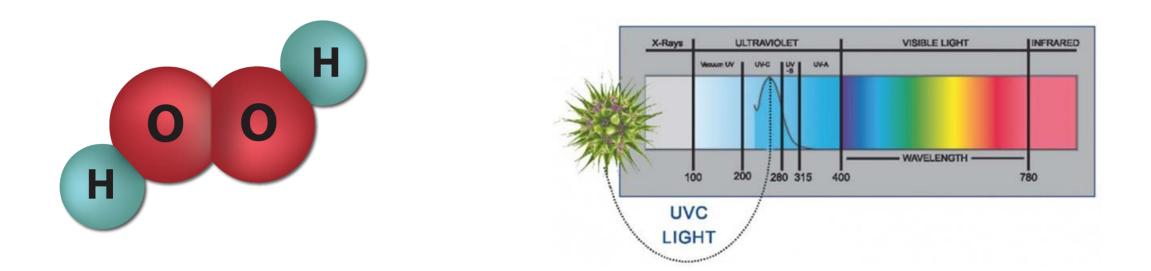
Automated whole room disinfection,

after manual cleaning, ensures a constant quality

What? Automated whole room disinfection

• Hydrogen peroxide (H_2O_2)

• Ultraviolet-C light



How does it work? Hydrogen peroxide

- $H_2O_2 \rightarrow H_2O + O_-$.
- O⁻ (oxigen radicals) kill bacteria
- Degrades into water and oxygen

 \rightarrow environementally friendly

- H_2O_2 is **toxic**: No people in the room!
- Room needs to be prepared for adequate concentration
- Two methods: Aerosolised H_2O_2 and H_2O_2Vapor



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Aerosolised HP (aHP) 5-7% H2O2

HP vapor (HPV) 30-35% H2O2

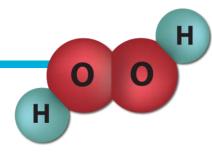
How does it work? aerosolised HP (aHP)

- 5-7% H_2O_2 is fogged into the room \rightarrow "dry mist"
- Room must be cleaned manually
- Literature suggests that the airvents and doors be sealed to prevent leakage of H_2O_2 .
- Turn off the smoke alarm.
- Open the cupboard doors and drawers, make sure the "mist" can reach the back side of the matrass etc.
- Measure the concentration (ppm with datalogger)
- Room can only be re-entered when [H₂O₂] is < 1ppm. (average 2-3 hours)

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Aerosolised HP (aHP) 5-7% H2O2



In vitro efficacy - aerosolised HP (aHP) review

Table 2 In vitro efficacy of aHP for the preselected set of microorganisms, expressed in log-reduction

	Micro-organism	Effect in log ₁₀ reduction, median (range)	N (ref)
Viruses	Norovirus (Surrogate)	2.5 (0.5–2.7) 4.5 (>4–5.3)	3 [12, 18, 19] 3 [18–20]
Bacteria	Acinetobacter (CPE	2 (1->4)	2 [12, 16]
	VRE	1-1.7	1 [21]
	ESBL	>6	1 [14]
	MRSA	>4 (2->6)	4 [12, 14, 16, 17]
Spores Yeast	C. difficile C. auris	4.9 (0.13->5)	4 [12, 14, 22, 23]

Disinfection is: > 5log reduction in bacterial load



Aerosolised HP (aHP) 5-7% H2O2

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van der Starre et al. Antimicrobial Resistance & Infection Control (2022)

aHP) H

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Pittfalls, Pro's and Con's – aerosolised HP (aHP)

Pitfalls:

- PPM needs to be monitored during the process
- H_2O_2 particles are affected by gravity.

Pro's

- User friendly, easy to transport
- Curtains can stay in the room, they are disinfected.

Con's

- Room needs to prepared.
- Room cannot be entered for 2-3 hours
- Doesn't reach log 5 reduction



Aerosolised HP (aHP) 5-7% H2O2

How does it work? HP Vapor

HOOH

- **30-35% H₂O₂ evaporation** (heat and multiple nozzles)
- Room must be cleaned and prepared
- Time and labour intensive preparation.
- Vents and doors *must* be sealed to prevent leaking of H_2O_2 .
- Must be operated by well trained person
- Thorough validation process



In vitro efficacy - HP Vapour review

Table 4 In vitro efficacy of H₂O₂ vapour for the preselected set of micro-organisms, expressed in log-reduction

	Micro-organism	Effect in log ₁₀ reduction, median (range)	N (ref)
Viruses	NoV (Surrogaat)	>4 4.4 (3-≥6)	1 [18] 4 [33, 34, 42, 43]
Bacteria	Acinetobacter	>5 (>4->6)	5 [12, 14, 35, 44, 45]
	CPE	>6	2 [44, 45]
	VRE	>6 (>4->6)	3 [35, 44, 45]
	ESBL		
	MRSA	>6 (3->6)	7 [12, 14, 35, 50 44, 46, 47]
Spores	C. difficile	>6 (>5,7->6)	6 [12, 23, 31, 37, 44, 48]
Yeasts	C. auris		

• HP vapour effectively reduces norovirus and the preselected set of bacteria.



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van der Starre et al. Antimicrobial Resistance & Infection Control (2022)

Pittfalls, Pro's and Con's – aerosolised HP

Pitfalls:

- Needs thorough validation process .
- Check compatibility of equipment with H_2O_2 .

Pro's

Good in vitro efficacy

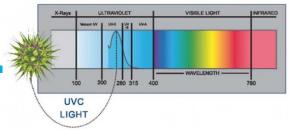
Con's

- Needs intensive training
- Preparation and vaporing is time consuming



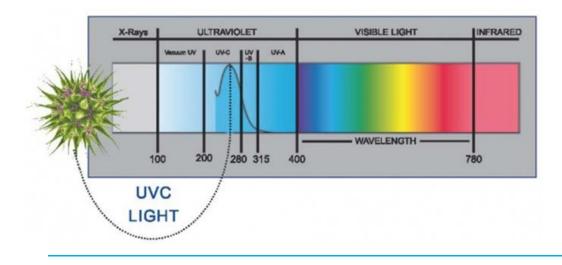
HP vapor (HPV)

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How does it work? Ultraviolet C light (UV-C)

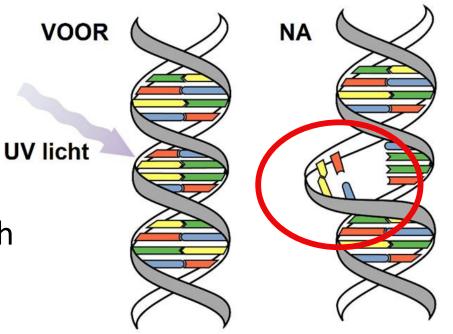
- Ultraviolet C light
 - stationary or mobile





How does it work? UV-C

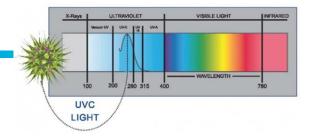
- UV-C has a wavelength of 254 nm
- UV-C light damages DNA/RNA → disrupts celldivision of micro-organisms
- Two thymines next to each other are linked to each other by UV-C.



UVC

VISIBLE LIGHT

Source: INFO UV | Safety Science



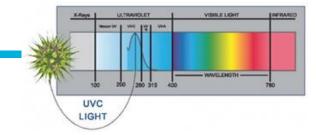
How does it work? UV-C

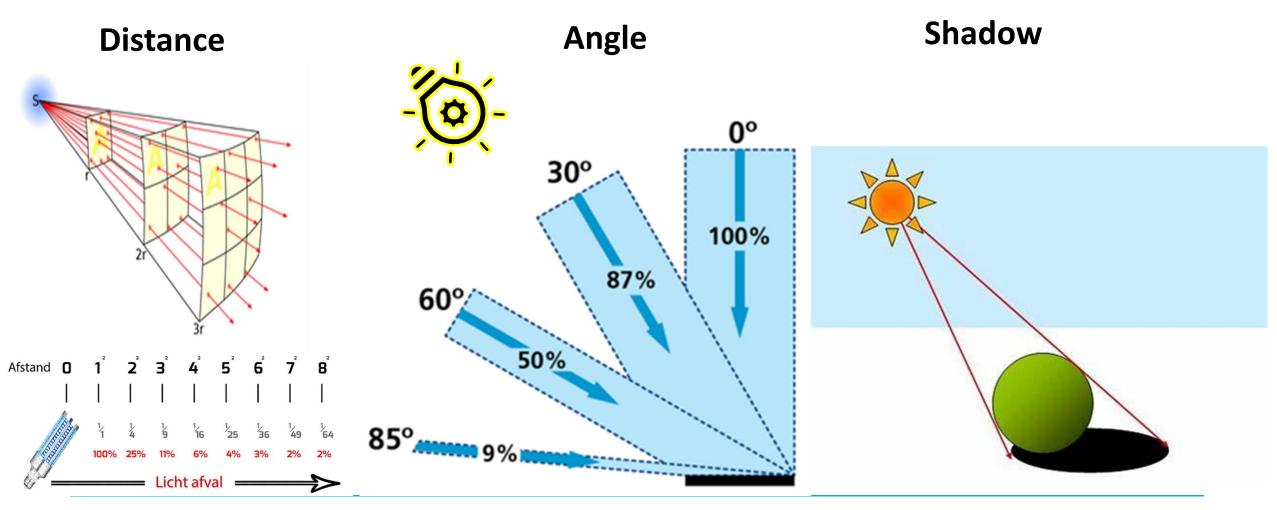
- Stationary and mobile systems.
- Clean before disinfection with UV-C
- Disinfection time: stationary +/- 40-50 min, mobile +/-15-20 min
- Mobile: needs space to drive around
- Room is immediately accessible after disinfection.
- Needs validation:

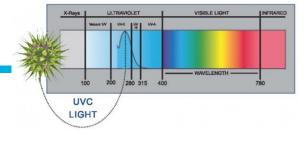
Shadow places must be disinfected manually



Intensity of UV-c depends on:







In vitro efficacy of UV-C review

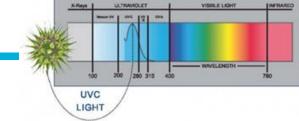
Table 6 In vitro efficacy of UV-C for a preselected set of micro-organisms under optimal and suboptimal circumstances

	Micro-organism	Optimal circumstances; effect in log ₁₀ reduction, median (range)	Suboptimal circumstances; effect in log ₁₀ reduction, median (range)	N (ref)
Viruses	Norovirus			
Bacteria	Acinetobacter	≥4 (≥4->8)	3 (< 1–4)	6 [58, 61–65]
	CPE	4–5	1-5	1 [66]
	ESBL	>8	>3	2 [62, 65]
	MRSA	4 (2-9)	<3 (<1->6)	13 [58, 60–63, 65–72]
	VRE	3.9 (2->8)	<3 (<1->4)	10 [58, 60–63, 67–71]
Spores	C. difficile	2.5 (1->5)	<2 (0->3)	11 [31, 57, 60, 61, 66–72]
Yeasts	C. auris	>5 (3.99->6)	3.3 (<2->4)	4 [72–75]

Big difference btwn studies and btwn optimal and suboptimal conditions

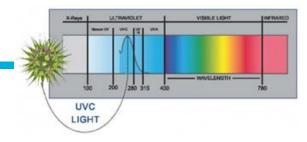
van der Starre et al. Antimicrobial Resistance & Infection Control (2022)

Microbiological evaluation of efficacy mobile UV-c robot in Radboudumc



- Regular patient room
- Demonstrate a log 5 reduction
- Make logarhitmic reference series for 5 relevant micro-organisms (Escherichia coli, Staphylococcus aureus, Enterococcus faecium, Pseudomonas aeruginosa, Acinetobacter baumanni complex)
- Inoculate Rodac-plates with bacterial suspension (0,5McFarland=log 0) and place them on various places in the patient room.
- Irradiate the plates by UV-c robot.
- Of each micro-organism a control log 0 plate was not irradiated

Evaluation in the patient room

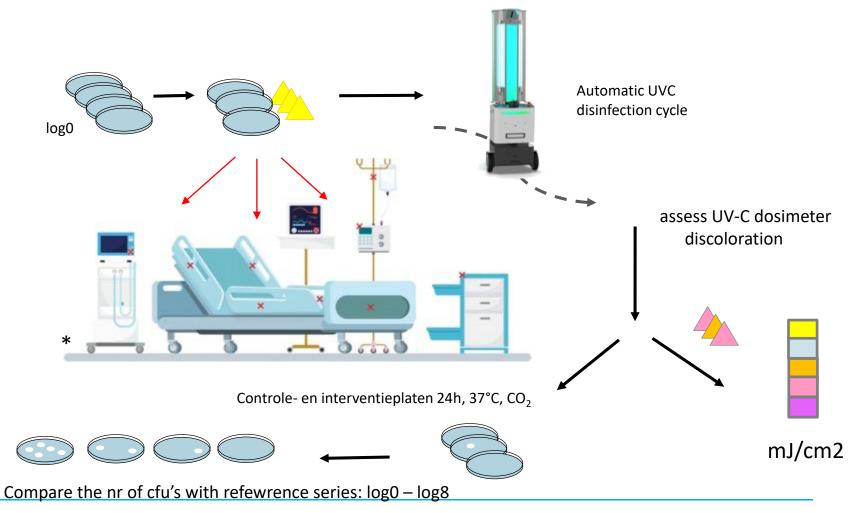


49 relevant positions in the patientroom, sluice and bathroom.

Inoculated Rodacplates

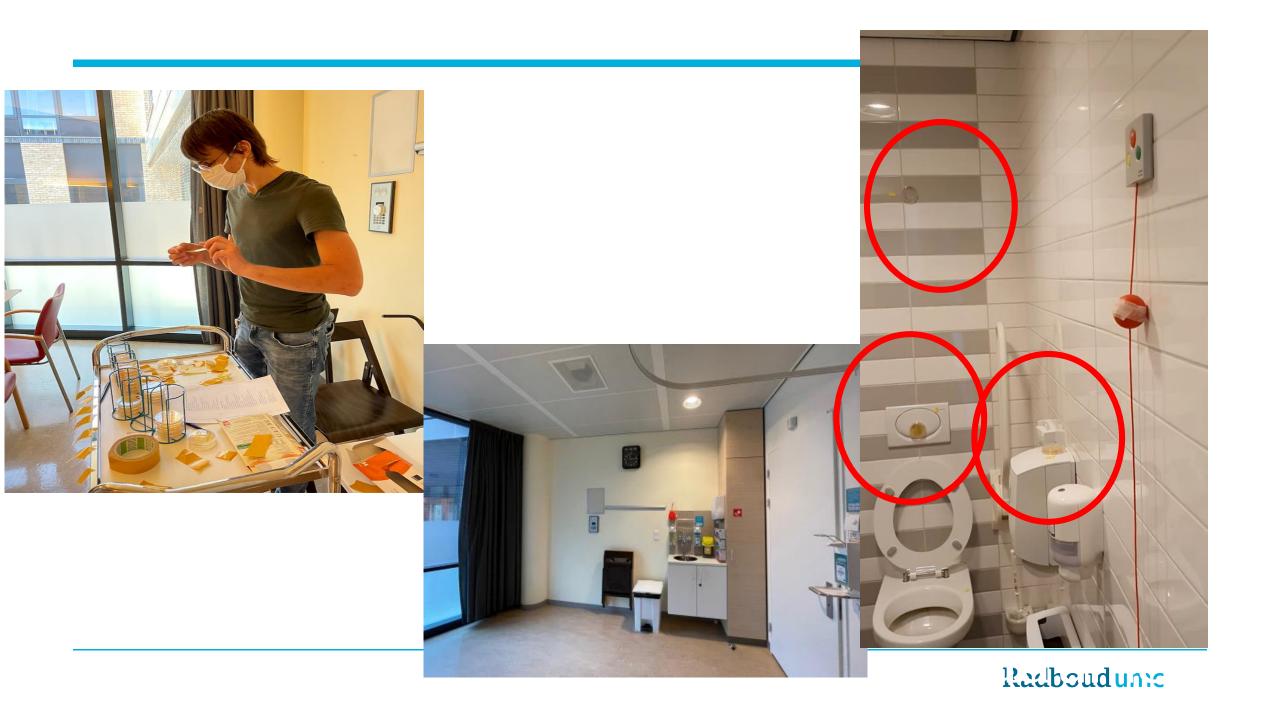
(log0) (

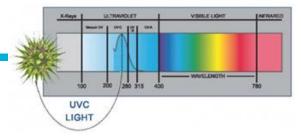
- UV-C dosimeter sticker
- Horizontal &
 vertical placed rodac plates and dosimeters



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https://doi.org/10.1016/j.infpip.2023.100322

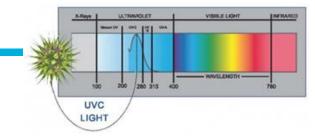




After UV-C

Before

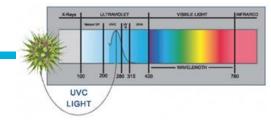




After UV-C

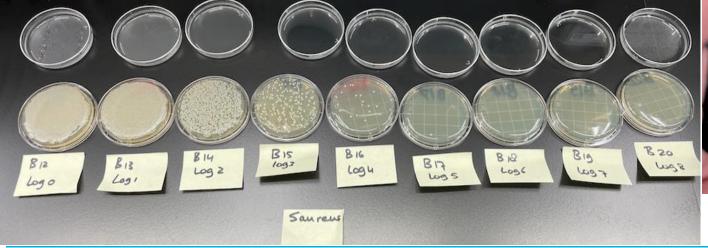
Before





Logaritmic reference series S. aureus

Position	Micro organism	Log	Cfu's	minimal log reduction
B12	S.aureus	log 0	full	0
B13		log1	full	1
B14		log2	full	2
B15		log3	>100	3
B16		log4	23	4
B17		log5	1	5
B18		log6	1	6
B19		log7	0	7
B20		log8	0	8





https://doi.org/10.1016/j.infpip.2023.100322

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Results

KRays LLTRADOLET VISILE LIGHT HATRA

Table II

Overview of UV-C dosage received per plate, and reduction of CFU's per plate. The results are presented separately for the plates showing \geq 5-log and <5-log reduction respectively. Results are split between plates placed on UV-accessible surfaces (unshaded surfaces below elbow height), and plates placed on surfaces with limited UV-accessibility ([partly] shaded and/or above elbow height)

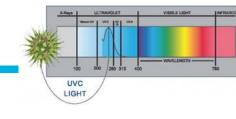
	UV-C dosage ≥50 mJ/cm², n	UV-C dosage 25-50 mJ/cm ² , n	UV-C dosage 25 mJ/cm ² , n	UV-C dosage <25 mJ/cm ² , n	UV-C dosage variable or not measured, n	Total, n
UV- accessible surfaces, n	29	1	0	1	1	32
≥5-log reduction, n	28 (1 plate contaminated)	1	0	1	1	31
<5-log reduction, n		0	0	0	0	0
surfaces with limited UV- accessibility, n	5	1	5	6	1	18
≥5-log reduction, n	4	1	4	3	1	13
<5-log reduction, n	1	0	1	3	0	5

Pro's

- Adequate log reduction
- Standardised disinfection, constant predictable quality
- For the cleaning staff: less repetitive movements, less physically strenuous
- Less/no chemicals -> Environmentally friendly
- Less sickleave

Con's

- Investment
- Needs validation in a new setting.
- Training of staff
- Beware of the shadow!!
- Beds create shadow
- Check compatibility with (medical) equipment



UV-c robot does not:

- Clean the room
- Disinfect objects in the shadow
- Disinfect inside cupboards, wet surfaces



Take home message:

- Manual cleaning is essential prior to disinfection, even with automated disinfection systems.
- Every new system needs to be validated and monitored.
- Adequately executed UV-C disinfection ensures consistent disinfection quality.
- Beware of the shadow!



Questions?

