

First UK trial of Tru-D: An automated UV-C room decontamination device

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BACKGROUND

The Tru-D™ rapid room disinfection device is a mobile, automated room decontamination technology utilizing ultraviolet-C (UV-C) irradiation to kill micro-organisms.

This device has been previously reported to significantly reduce nosocomial pathogens in the healthcare environment such as MRSA, VRE and Clostridium difficile. 1,2 making it an interesting potential alternative to systems such as vaporised hydrogen peroxide or dry-mist hydrogen peroxide for terminal disinfection of patient rooms.

However, until recently this technology has not been available in the UK.

PURPOSE

To perform a rapid trial of the first UK Tru-D unit in an NHS healthcare setting, looking principally at:

- · ease of use
- · time taken for room disinfection
- microbiological efficacy
 - total surface aerobic counts
 - specific pathogens (MRSA, VRE, MRA and Aspergillus)

MATERIALS AND METHODS

Deploying Tru-D

The unit was trialed in six side rooms within an Intensive Therapy Unit (ITU), an operating theatre and an ensuite ward sideroom. Measurements were taken to calculate room volume and time taken for disinfection was recorded at two settings (reflected UV-C dose of 12,000 µWs/cm² for vegetative bacteria & a sporicidal setting of 22,000 μ Ws/cm²).

Evaluating microbial efficacy

Assessment was performed using two methods:

Contact plates

Tryptone soya agar (TSA) contact plates were applied to surfaces both within the line of sight (LOS) of the Tru-D unit and areas not directly in the line of site (shadow). TSA contact plates were re-applied to surfaces directly adjacent to original sampling area at the end of the Tru-D disinfection cycle. The total number of colonies before and after disinfection were counted

Plastic Petri dishes seeded with multi-resistant clinical isolates

Suspensions containing MRSA, VRE, MRA or Aspergillus were produced (McFarland 0.5-1 [1.5-3.0 X108 cfu/ml]). A sterile cotton-wool swab was used to spread the inoculum evenly on Petri dishes to produce a confluent/semi-confluent growth of organisms on control plates (~104/105 colonies/plate). Seeded Petri dishes were then placed on surfaces (line of sight and shadow) and exposed to Tru-D. Control Petri dishes were kept outside. The total number of colonies recovered from TSA contact plates after disinfection were counted and the log₁₀ reduction compared to controls calculated (see Fig 2).

All TSA contact plates were incubated at 37°C for 48 hours aerobically.



Fig 1: Photograph showing inoculated Petri dishes from a suspension containing VRE



Fig 2: Photograph showing application of a TSA contact plate onto a Petri dish inoculated with MRA

BIBLIOGRAPHY

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RESULTS

Using the Tru-D Unit

The Tru-D unit was relatively easy to transport and operate. It was placed in the centre of the rooms and furniture moved away from the walls. The device was operated using a remote control with suitable warning signs to prevent inadvertent entry whilst UV-C disinfection was in process. It took approximately 30-40 minutes to decontaminate rooms at 12,000 μWs/cm² (for vegetative bacteria) and 60-90 minutes at the sporicidal setting (22,000 μWs/cm²) as shown in the table below.

	Room		Dose	Time taken	Dose	Time taken
	Surface	Volume				
	Area (m2)	(m3)	uWs/cm2	(mins)	uWs/cm2	(mins)
ITU Single Room A (1)	26	72	12,000	27		
ITU Single Room A (2)	26	72	12,000	39	22,000	73
ITU Single Room B	29	80	12,000	31		
ITU Single Room C	17	47	12,000	36		
ITU Single Room D	21	59	12,000	32		
ITU Single Room E	20	55	12,000	26		
ITU Single Room F	22	59			22,000	93
Operating theatre	42	126	12,000	49		
Stoke unit isolation room	16	39			22,000	60
Stroke unit ensuite bathroom	4	8			22,000	23

Contact plates

Between 0-40 (median 10) colony forming units (cfu) per contact plate could be recovered from surfaces in the cleaned, unoccupied operating theatre. Following the use the Tru-D unit (12,000 µWs/cm²) no organisms could be recovered from the environment (Figures 3 & 4).

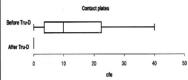


Fig 3: Box and whisker plot of the range of total aerobic cfu recovered from 15 replicate samples in an operating theatre before and after Tru-D

Before Tru-D After Tru-D Fig 4: Contact plates before and after the use of Tru-D

Seeded plastic Petri dishes

Tru-D demonstrated a high kill of MRSA both in line of sight (LOS) and shadow. Log₁₀ reductions of VRE, MRA and Aspergillus were slightly lower in shadowed areas and at 12,000 compared to 22,000 µWs/cm²

			reduction		
		Dose	LOS*	Shadow	
Operating theatre	MRSA	12,000	>4	>4	
	VRE	12,000	3.6	2.4	
	VRE	12,000	4.2	2.3	
	VRE	22,000	4.4	3.5	
ITU single room	MRA	12,000	>4	1.7	
	MRA	22,000	>4	3.0	
	Aspergillus	12,000	4.0	2.0	
	Aspergillus	22,000	4.3	1.0	
	*LO	B = line of	sight		



Fig 5: TSA contact plates from Petri dishes inoculated with MRSA before after the use of Tru-D at 12,000 µWs/cm2

Fig 6: TSA contact plates from Petri dishes inoculated with MRA (above) and Aspergillus before after the use of Tru-D at 12,000 µWs/cm2 and 22,000 µWs/cm2



CONCLUSIONS

- The Tru-D was easy to use and room disinfection times were relatively short. Without the need to inactivate room ventilation or smoke detectors, we were able to disinfect 3 ITU single rooms within 3 hours.
- 2. This device appears to achieve significant killing of key healthcare environmental pathogens including MRSA, VRE, MRA and Aspergillus.
- 3. Log₁₀ reductions were lower within shadowed areas compared to areas directly within line of sight, and we would recommend using the Tru-D at a reflected dose setting of 22,000 µWs/cm² for terminal room disinfection in most situations.

ACKNOWLDEGEMENTS

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