

Aerosol generation during surgical tracheostomy in a patient with COVID-19

Forbes McGain, Ruhi S Humphries, Juan Carlos Mora, Patrick Timms, Fiona Hill and Craig French

Significant concern exists regarding the risk of transmission of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) to health care workers during aerosol generating procedures.^{1,2} The risk of viral transmission to health care workers during tracheostomy insertion is unknown.

To address this uncertainty, we continuously measured aerosol particle concentration during a planned, semi-elective surgical tracheostomy insertion on a SARS-CoV-2-positive patient. The procedure was performed in April 2020 at Western Health, Melbourne, Australia. Written informed consent for tracheostomy was obtained from the patient's legally approved representative. The Western Health Human Research Ethics Committee advised that ethics approval was not required to conduct this environmental study.

We used two spectrometers to measure aerosol particles: the portable Mini Wide Range Aerosol Sizer 1371 (MiniWRAS) (GRIMM Aerosol Technik, Ainring, Germany) and the Aerodynamic Particle Sizer (APS) (TSI, Shoreview, MN, USA). The APS spectrometer detected larger aerosols (> 0.37 μm) and the MiniWRAS spectrometer measured smaller particles (0.01–0.35 μm). Human-generated respiratory air particles range mostly from 0.5 μm to 20 μm .³ We measured operating theatre aerosol counts for 24 hours before the procedure to obtain baseline background aerosol concentration. During the procedure, the aerosol detector inlet was positioned 30 cm directly above the patient's neck, representing the surgeon's breathing air space.

Total intravenous anaesthesia was maintained with propofol and fentanyl, and neuromuscular blockade with cisatracurium. Synchronised intermittent mandatory volume controlled ventilation was delivered throughout. Surgeons used bipolar diathermy for electrocautery during dissection of the neck tissues. Several minutes before tracheal incision, the endotracheal tube (ETT) was advanced under apnoeic conditions towards the carina to avoid accidental cuff damage when tracheal incision occurred. Ventilation was recommenced to allow for several minutes of pre-oxygenation before tracheal incision. At this point, the high pressure alarm occurred. This was clinically suggestive of inadvertent endobronchial intubation; however, due to the

risk of aerosolisation this was not verified via bronchoscopy. The high pressure alarm was rectified by withdrawing the ETT by 2 cm. Tracheal incision and tracheostomy tube insertion were conducted under apnoeic conditions. A 5-second period of accidental circuit disconnection occurred before completion of surgery.

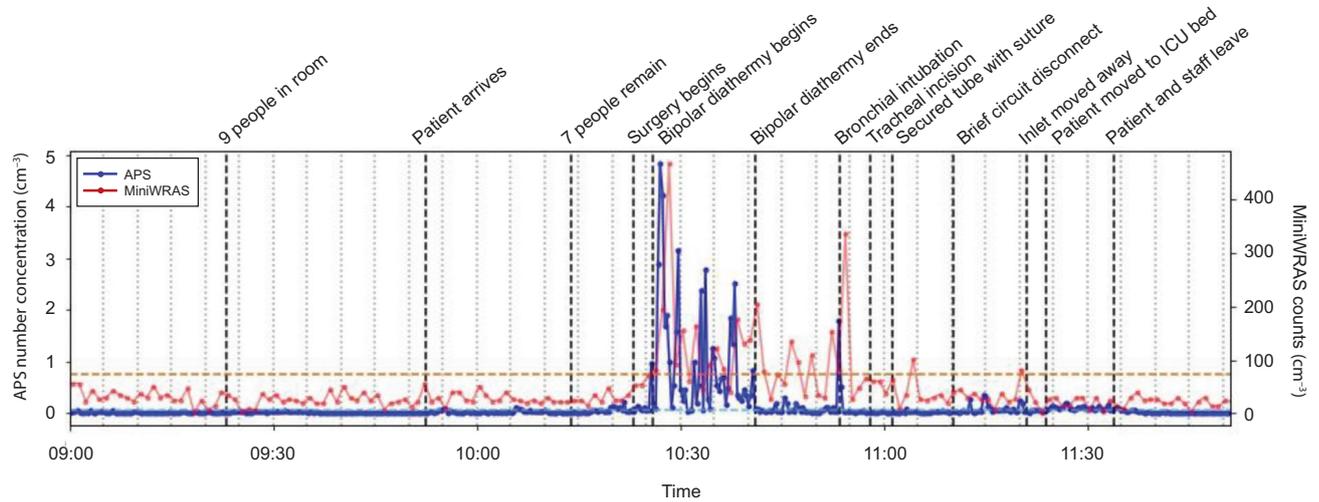
Figure 1 provides a time series of particle concentrations detected by the APS (larger size) and MiniWRAS (smaller size) spectrometers. Table 1 details the total particle concentrations measured by the spectrometers at different times during the procedure. Particle counts for the majority of the tracheostomy procedure were low; similar to delivering oxygen at 15 L/min via a simple face mask.⁴ Tracheal incision and brief circuit disconnection did not cause a significant rise in aerosol counts. The most significant increase in particle counts was detected during diathermy (50-fold increase) and at the time of inadvertent bronchial intubation (30-fold increase). The sudden and sharp rise in aerosol count at this time was unexpected and could not be attributed to another clinical incident. While the high peak airway pressure alarm was being investigated, the peak pressure limit was increased temporarily. We suspect that this led to airway pressures surpassing the ETT cuff pressure and thus allowing turbulent gas flow to bypass the ETT and generate aerosols via the patient's upper airway. These high aerosol counts are similar to quantities detected during non-invasive ventilation⁴ and are greater than the chosen safe threshold for low aerosol generation.

Energy devices, such as diathermy, generate aerosolised particles,⁵ but the risk of infection transmission via these bioaerosols is unknown. The only evidence that viral transmission is possible is in a bovine model.⁶ We are not aware of any evidence in humans that diathermy-generated aerosols can carry active virus.⁵

Our study has limitations. We only used a single sampling point, which could potentially underestimate particle concentrations; however, it was positioned in close proximity to the surgical field and likely provides a reliable estimate of aerosol burden related to the surgical procedure. We did not measure or confirm viral aerosol presence. We

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Figure 1. Timeline of aerosol concentration detected by Aerodynamic Particle Sizer (APS)* and Mini Wide Range Aerosol Sizer (MiniWRAS)† spectrometers‡



* GRIMM Aerosol Technik, Ainring, Germany. † TSI, Shoreview, MN, USA. ‡ The **x-axis** has lighter grey, vertical dotted bars indicating 5-minute intervals. Important events are labelled along the timeline as darker, black dashed bars. Pre-oxygenation occurred between bipolar diathermy ending and bronchial intubation. A useful threshold for a significant rise in particle counts is up to 3 standard deviations (99.5%) above the baseline mean of the theatre preparation period with all staff present (Table 1, Activity 2). The horizontal dashed orange line indicates this threshold for the APS measurements, while the light blue dashed line (mixed into the background blue line) represents this for the MiniWRAS measurements. Clearly, only diathermy, bronchial intubation and inadvertent circuit disconnection post-tracheal incision lead to aerosolisation above this background aerosol count. The **y-axis** gives the number of aerosols per cm⁻³ (mL), with smaller particles measured by the MiniWRAS spectrometer and larger particles by the APS spectrometer.

Table 1. Total aerosol concentration recorded by the Aerodynamic Particle Sizer (APS)* and Mini Wide Range Aerosol Sizer (MiniWRAS)† spectrometers during the tracheostomy‡

Activity	Particles per mL								Number of samples taken	
	Mean		10th %		90th %		Maximum		Mini-WRAS	APS
	Mini-WRAS	APS	Mini-WRAS	APS	Mini-WRAS	APS	Mini-WRAS	APS	Mini-WRAS	APS
1. Background empty theatre (overnight)	29	0.001	11	0.000	46	0.003	188	0.019	360	1080
2. Theatre preparation (all staff present)	32	0.020	15	0.009	50	0.035	57	0.057	23	69
3. Bipolar diathermy	136	0.997	51	0.092	187	2.671	468	4.828	15	45
4. Bronchial intubation	209	0.594	-	0.035	-	1.407	-	1.790	2	4
5. Tracheal incision	48	0.029	-	0.028	-	0.031	-	0.032	1	3
6. Brief circuit disconnect	60	0.018	-	0.013	-	0.023	-	0.025	1	3
7. Patient moved to ICU bed	41	0.089	-	0.020	-	0.035	-	0.038	1	3
8. Patient and staff leave	26	0.100	-	0.064	-	0.117	-	0.126	1	3
9. Background stabilised theatre 30 min after surgery	15	0.001	-	0.087	-	0.115	-	0.120	1	3

ICU = intensive care unit. * GRIMM Aerosol Technik, Ainring, Germany. † TSI, Shoreview, MN, USA. ‡ APS particle size measurement from > 0.37 µm. The MiniWRAS 1371 spectrometer was used to determine the concentration of smaller particles (0.01–0.37 µm). Statistics were not shown when the number of samples was low.¹⁻²

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cannot extrapolate our findings to an emergent surgical tracheostomy or percutaneous tracheostomy.

This case demonstrates it is possible to achieve low aerosol particle generation during the majority of the conduct of a surgical tracheostomy. While diathermy generates aerosols, it is unlikely any virus survives the high energy discharge related to it. However, subclinical or clinical leakage of exhaled airway gas around the endotracheal tube also caused increased aerosol particle generation. This reinforces the need for full personal protective equipment with contact and aerosol precautions. Our data suggest that particle generation dangers during a surgical tracheostomy are generally low and can be mitigated by careful health care worker collaboration. Surgical tracheostomy can be performed safely in patients with coronavirus disease 2019 (COVID-19).

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Competing interests

None declared.

Author details

Forbes McGain^{1,2,3}
Ruhi S Humphries⁴
Juan Carlos Mora¹
Patrick Timms¹
Fiona Hill¹
Craig French^{1,3,5}

- 1 Western Health, Melbourne, VIC, Australia.
- 2 School of Public Health, University of Sydney, Sydney, NSW, Australia.
- 3 Centre for Integrated Critical Care, University of Melbourne, Melbourne, VIC, Australia.
- 4 Climate Science Centre, CSIRO Oceans and Atmosphere, Melbourne, VIC, Australia.
- 5 Department of Epidemiology and Preventive Medicine, Monash University, Melbourne, VIC, Australia.

Correspondence: forbes.mcgain@wh.org.au

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