Bactericidal effects of silver plus titanium dioxide-coated endotracheal tubes on Pseudomonas aeruginosa and Staphylococcus aureus

Keiko M Tarquinio1
Nikhil K Kothurkar2
Dharendra Y Goswami3
Ronald C Sanders Jr4
Arno L Zaritsky5
Ann Marie LeVine6
1Division of Pediatric Critical Care Medicine, Department of Pediatrics, Rhode Island Hospital, The Warren Alpert Medical School, Brown University, Providence, RI USA; 2Department of Chemical Engineering and Materials Science, Amrita School of Engineering, Ettimadai, Coimbatore, India; 3Clean Energy Research Center, University of South Florida, Tampa, FL, USA; 4Section of Pediatric Critical Care, Department of Pediatrics, College of Medicine, University of Arkansas for Medical Sciences, Arkansas Children’s Hospital, Little Rock, AR, USA; 5Executive Medical Director, Children’s Hospital of The King’s Daughters, Norfolk, VA, USA; 6Pediatric Critical Care Medicine, University of Michigan Medical School, C.S. Mott Children’s Hospital, Ann Arbor, MI, USA

Purpose: Ventilator-associated pneumonia (VAP) is a nosocomial infection resulting in significant morbidity and mortality. Pseudomonas aeruginosa (P. aeruginosa) and Staphylococcus aureus (S. aureus) are pathogens associated with VAP. Silver (Ag) coating of endotracheal tubes (ETTs) reduces bacterial colonization, however titanium dioxide (TiO2) coating has not been studied.

Methods: Five types of ETT coatings were applied over silica layer: Ag, solgel TiO2, solgel TiO2 with Ag, Degussa P25 TiO2 (Degussa TiO2), and Degussa TiO2 with Ag. After ETTs were incubated with P. aeruginosa or S. aureus; colonization was determined quantitatively.

Results: Pseudomonas aeruginosa and S. aureus grew for 5 days on standard ETTs. Compared to standard ETTs, P. aeruginosa growth was significantly inhibited by solgel TiO2 with Ag at 24 hours, and by Degussa TiO2 with Ag at 24 and 48 hours after inoculation. No significant difference in S. aureus growth was observed between the control and any of the five coatings for 5 days.

Conclusion: In vitro, solgel TiO2 with Ag and Degussa TiO2 with Ag both attenuated P. aeruginosa growth, but demonstrated no effect on S. aureus colonization. Further studies using alternative coating and incorporating UV light exposure are needed to identify their potential utility in reducing VAP.

Keywords: ventilator-associated pneumonia, Degussa titanium dioxide, solgel titanium dioxide, quantitative culture

Introduction
Ventilator-associated pneumonia (VAP) is one of the most common nosocomial infections among intensive care unit patients, which prolongs ventilation and hospitalization, resulting in significant morbidity, mortality and medical costs.1-3 Researchers have developed strategies to prevent VAP, including controlling gastric pH, intratracheal intubation with a high-pressure endotracheal tube (ETT) cuff, continuous subglottic suction to prevent microaspiration, promoting oral hygiene with chlorhexidine, and semi-recumbent positioning.4-9

One strategy to reduce VAP is to treat the ETT with an antiseptic agent to control ETT biofilm formation.10,11 A biofilm is an aggregate of a bacterial community attached to a solid surface and encased in an exopolysaccharide matrix, making it resistant to antibiotic penetration.12 Biofilm formation is most prominent in the distal third of the ETT and can occur as early as sixty hours following endotracheal intubation.13
Due to silver’s antimicrobial properties, Pacheco-Fowler and colleagues studied chlorhexidine and Ag carbonate impregnated ETTs in an in vitro model and observed that bacterial growth of *S. aureus*, *P. aeruginosa*, and other organisms was attenuated. Similarly, Rello et al found that bacterial colonization was delayed for seven days in adults intubated with Ag-coated ETTs compared with standard ETTs, and Kollef and colleagues concluded a statistically significant reduction in the VAP incidence in a randomized multicenter trial.

Titanium dioxide is widely used as an antimicrobial agent, but it is not commonly employed in the medical setting. In a process known as photocatalysis, TiO$_2$ is activated into potent oxidative species when exposed to UV light, with antiviral, antibacterial as well as fungicidal actions. Recent investigations have suggested that the photocatalytic activity is important in destroying a range of bacteria including *S. aureus* and *P. aeruginosa* in aqueous solutions. Photocatalytic activity also is implicated in the inhibition of biofilm formation on TiO$_2$ based dental implants. In view of these antimicrobial actions it is plausible that TiO$_2$ coating of ETTs, potentially in combination with UV light, may sterilize ETTs in patients’ airways, thereby reducing the risk of VAP.

We conducted a pilot study of bacterial growth behavior on TiO$_2$-coated ETTs without UV exposure to evaluate the antimicrobial effect of TiO$_2$ on polyvinyl chloride (PVC) which is used to manufacture ETTs. Initial experiments examined TiO$_2$ without UV light exposure for two reasons. First, the Ag and TiO$_2$ combination has not been studied on PVC material. Second, if the Ag and TiO$_2$ coating has antimicrobial activity without photocatalysis, this would reduce potential toxicity from UV exposure. The objective of this study was to determine if TiO$_2$-coated ETTs with or without Ag would reduce bacterial colonization by two common VAP pathogens (ie, *P. aeruginosa* or *S. aureus*) in an in vitro model compared with standard ETTs.

### Material and methods

#### Solgel TiO$_2$ synthesis

Sols of TiO$_2$ were made using a method adapted from Mills and colleagues. A solution of 10 mL titanium isopropoxide (AC19470 98+%; Acros Organics, Geel, Belgium) in 2.32 g glacial acetic acid (A465-250; Fisher Chemical, Pittsburgh, PA), was stirred into 60 mL deionized water acidified with 0.59 g concentrated nitric acid (S719721; Fisher Chemical). The reaction solution was then heated rapidly and held at 80°C for 8 hours. The solution was concentrated by heating at 150°C to 11 wt% TiO$_2$, followed by dilution with ethanol (AC61510; Acros Organics) to obtain a translucent white sol containing 6 wt% TiO$_2$ sol.

#### Endotracheal tubes

The inner lumens of the ETTs were first coated with silica (SiO$_2$) which served as a passivating layer to prevent the TiO$_2$ from degrading the polyvinyl chloride (PVC) and to improve surface wettability by the subsequently applied TiO$_2$ coatings. The SiO$_2$ layer was applied using a nebulizer spray of 15 wt% colloidal SiO$_2$ in deionized water through the ETT, through a custom-designed nozzle, for 1 minute with an air flow rate of 10 liters per minute. The ETT was then dried with 85°C air for 1 minute.

The respective TiO$_2$ and Ag treatments are summarized in Table 1. Five types of coatings were applied over the SiO$_2$ layer (Figure 1): 1) solgel TiO$_2$, 2) solgel TiO$_2$ with Ag, 3) Degussa TiO$_2$ (Evonik Degussa, Parsippany, NJ, USA), 4) Degussa TiO$_2$ with Ag, and 5) Ag (not shown in Figure 1). Each of the coating solutions A–D had 0.19 wt% TiO$_2$ on a dry weight basis. Water and ethanol were used as solvents in variable ratios for each solution. Solgel TiO$_2$ was used for solutions A and B, while C and D solutions consisted of solgel and Degussa TiO$_2$ in the ratio 20:80 wt/wt. Silver nitrate (AgNO$_3$) was added to solutions B and D. For solution E, AgNO$_3$ 0.1 wt% was mixed in ethanol-water (50:50 wt/wt). Two types of TiO$_2$ coatings were applied; solgel

<table>
<thead>
<tr>
<th>Tube</th>
<th>Solution</th>
<th>TiO$_2$ source</th>
<th>Solvent</th>
<th>AgNO$_3$; TiO$_2$ (mol%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>A</td>
<td>Solgel</td>
<td>Water: Ethanol (60:40)</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>B</td>
<td>Solgel</td>
<td>Water: Ethanol (60:40)</td>
<td>5.6</td>
</tr>
<tr>
<td>3</td>
<td>C</td>
<td>Solgel: Degussa (20:80)</td>
<td>Water: Ethanol (3:97)</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
<td>D</td>
<td>Solgel: Degussa (20:80)</td>
<td>Water: Ethanol (5:95)</td>
<td>5.6</td>
</tr>
<tr>
<td>5</td>
<td>E</td>
<td>No TiO$_2$</td>
<td>Water: Ethanol (50:50)</td>
<td>0.1 wt% AgNO$_3$</td>
</tr>
</tbody>
</table>

Abbreviations: Ag, silver; AgNO$_3$, silver nitrate; Degussa TiO$_2$, Degussa P25 titanium dioxide; TiO$_2$, titanium dioxide.
Bactericidal effects on titanium dioxide-coated endotracheal tubes

Preparation of S. aureus

Frozen S. aureus (ATCC #25923) was rehydrated with TSB in the same fashion as P. aeruginosa, and stored at −80°C. Frozen S. aureus was inoculated into 50 mL TSB in a flask, and incubated overnight at 37°C. Turbid S. aureus bacterial broth was transferred to a 50 mL sterile culture tube, centrifuged for 15 minutes, and the supernatant was discarded. A concentrated broth was made by the addition of 1 mL of TSB to the pellet of S. aureus. Optical density was measured to obtain 10^8 CFU/mL for ETT inoculation.

Bacterial inoculation of ETTs

Each ETT was marked at 4 cm intervals for later division into six equal pieces, then immersed in 75% alcohol for 15 minutes to sterilize, rinsed with 20 mL 0.9% sterile saline, and air dried. After sterilization, each ETT was cut into six pieces, sealed on one end with Tegaderm® (3M, St. Paul, MN), filled with either 8 μl of P. aeruginosa broth plus 800 μl TSB or 100 μl S. aureus broth plus 700 μl TSB, then sealed with Tegaderm. All ETT pieces were placed in a 2 liter dry flask and agitated at 37°C overnight. On the following day, the bacterial fluid was drained from each ETT piece.

Peroxidase assay after coating ETTs

Colorimetric measurement was used to detect whether TiO_2 on ETTs generated peroxidase as part its antimicrobial property. EM Quant Peroxide Test Strips (EMD Chemicals Inc. Gibbstown, NJ, USA) which measure peroxidase between 1–100 parts-per million were used. Tryptic soy broth 800 μL was placed on the TiO_2-coated ETT for 24 hours at 37°C in the dark. Following incubation, the colorimetric strip was dipped into the TSB and the value of peroxidase production was determined.

Observation of bacterial growth

Bacterial inoculated ETT pieces were placed in an incubator to simulate the in vitro human trachea with no ambient light, a relative humidity of >90%, temperature of 37°C and at standard atmospheric pressure. Bacterial growth was observed for 24, 48, 72, 96, 120, and 144 hours after inoculation on the standard and five test ETTs. The inner surface of bacteria was collected by washing with 0.5 mL of sterile PBS and swabbing every 24 hours after inoculation. The collected fluid was serially diluted and plated on tryptic soy agar for incubation overnight at 37°C. Colony counts were expressed as CFU/mL, and each experiment was repeated four times.

Preparation of P. aeruginosa

Freeze-dried P. aeruginosa (ATCC #25668; American Type Culture Collection, Manassas, VA, USA) was rehydrated with nutrient broth in aliquots to be stored at −80°C. After a loopful of frozen bacteria was plated and incubated, a single colony was inoculated into 4 mL tryptic soy broth (TSB) and incubated overnight at 37°C. The following day, the turbid bacterial broth was vortexed and inoculated. To determine the exact inoculums, 100μL of bacterial broth was serially diluted with phosphate-buffered saline (PBS), then optical density was measured to obtain 10^6 colony forming units/milliliter (CFU/mL). to increase adherence to PVC for substrate depositition and Degussa TiO_2 for potentially high photocatalytic performance. Since Degussa TiO_2 suspensions had very poor adhesion to the ETTs, they were mixed with a small amount of sols of solgel TiO_2. After TiO_2 coating, the ETTs were dried with 85°C air for 1 minute, followed by further drying in an air-circulating oven at 100°C for 15 minutes, then packed while warm.
Statistical analysis
Data were reported as mean ± standard error of mean (SEM). Bacterial colony counts were analyzed using Mann–Whitney U test comparing coated ETTs with standard ETTs. A P-value of < 0.05 was considered statistically significant.

Results
The coating was slightly opaque using solgel TiO₂ compared with Degussa TiO₂. Samples with Ag-containing coatings appeared dark due to the reduction of AgNO₃ to Ag. The coating was particulate in appearance to the naked eye and peeled off while collecting the bacteria by mechanical force. Peroxidase production was not detected on TiO₂-coated ETTs.

Pseudomonas aeruginosa
Growth peaked at 72 hours and continued for 144 hours on standard ETTs. Pseudomonas aeruginosa growth was more robust on Ag-coated ETTs compared to standard ETTs at 120 hours (P = 0.04) and 144 hours (P = 0.06) despite initial slow growth in the first 24 hours (Figure 2). There was no difference in P. aeruginosa growth over 144 hours among standard, solgel TiO₂ and Degussa TiO₂-coated ETTs (Figure 3). Compared to standard ETTs, solgel TiO₂ with Ag inhibited P. aeruginosa growth at 24 hours (P = 0.02), and Degussa TiO₂ with Ag inhibited P. aeruginosa growth at 24 and 48 hours after inoculation (P = 0.02 and P = 0.02, respectively) (Figure 4).

Staphylococcus aureus
Growth was observed up to 144 hours. Staphylococcus aureus growth peaked at 48 hours after inoculation on both standard and Ag-coated ETTs without a difference over 144 hours (Figure 5). Furthermore, there was also no difference in S. aureus growth over 144 hours between standard, solgel TiO₂ and Degussa TiO₂-coated ETTs (Figure 6). Compared to standard ETTs, neither solgel TiO₂ with Ag nor Degussa TiO₂ with Ag showed a difference in S. aureus growth over a 144 hour period except solgel TiO₂ with Ag at the 144 hour point after inoculation in which there was a significant attenuation of S. aureus growth compared to the standard ETT (Figure 7).

Discussion
Clinically, VAP is diagnosed after patients have been mechanically ventilated for more than 48 hours similar to the duration of bacterial growth in our simulated model. The growth of P. aeruginosa was consistent with the findings of other investigators under comparable conditions when monitoring growth in the first 24 hours. An initial suppression of P. aeruginosa in Ag-coated ETT was also
observed in sections of intraventricular tubing impregnated with Ag.23 Moreover, our experience was similar to results of others who observed an undulating pattern of bacterial growth over a 50 hour period in Ag-coated ETTs.24 The second \textit{P. aeruginosa} growth peak observed in Figure 2 was thought to be an experimental variation due to the small sample size.

A similar growth pattern on Ag-coated ETT was also observed in a study evaluating the growth of \textit{P. aeruginosa} and \textit{S. aureus} on PVC even in the presence of antimicrobials. Both \textit{P. aeruginosa} and \textit{S. aureus} produce a biofilm; its robustness is strain dependent.25–27 Biofilm–generating species of \textit{S. aureus} exhibit accelerating growth kinetics in the first 6 days in animal models. Furthermore, these strains form microcolonies on catheter surfaces that act as sites of biofilm formation.28 Thus, it is likely that \textit{S. aureus} growth was probably blunted in our model until a glycocalyx developed, then growth proceeded in the presence of nutrients and favorable conditions.

The antimicrobial properties of Ag result from Ag ions binding to bacterial sulphhydryl- or histidyl-containing proteins, which disrupt transmembranous energy metabolism and electrolyte transport systems.29 Because the cell wall composition and thickness vary between gram-positive and gram-negative bacteria, silver’s antimicrobial effect may be greater in \textit{P. aeruginosa} than in \textit{S. aureus} during the initial phase of bacterial multiplication.

The characteristics of discolored Ag-coated particles were not analyzed in this experiment. However, AgNO$_3$ is known to undergo photochemical reduction to metallic Ag upon exposure to light as described by the reaction:

\[ 2\text{AgNO}_3 \rightarrow 2\text{Ag} + 2\text{NO}_2 + \text{O}_2 \]

This leads to the tiny particles precipitation of Ag that appears black or brown. Future studies are needed to define the specific mechanism involved in the antimicrobial properties of Ag as well as the physiological impact of discolored Ag. Silver nitrate was not expected to react with SiO$_2$ under ambient conditions, since SiO$_2$ is quite nonreactive.

Application and treatment of ETTs with these chemicals may also change the surface charge and hydrophobicity, as in this study, thus changing the PVC characteristics from hydrophobic to hydrophilic.

This study demonstrated that the addition of TiO$_2$ significantly augmented the antibacterial effects of Ag on \textit{P. aeruginosa} during the first 72 hours, but had no effect on \textit{S. aureus}. In our limited study, TiO$_2$ photocatalysis was not observed as determined by the release of hydrogen peroxide.
as a part of reactive oxygen species. Silver and TiO$_2$ might have reacted with each other releasing Ag ion, and this reactant might have a synergistic antimicrobial effect even in a dark environment without photocatalysis.

In contrast, Yao showed that application of thin films of Ag and TiO$_2$ for sterilization purposes on silicone catheters promptly eliminates *S. aureus* within 90 minutes without TiO$_2$-photocatalysis. They also showed inhibition of growth of *Escherichia coli* (*E. coli*) and *P. aeruginosa*, although they only observed growth for 24 hours.\textsuperscript{30} Li and colleagues demonstrated that AgNO$_3$ and TiO$_2$ pulverized to surgical masks reduced *E. coli* and *S. aureus* by 100%. In Li’s study, Ag plus TiO$_2$ pulverized masks were exposed to UV-C irradiation prior to bacterial inoculation.\textsuperscript{31} *Pseudomonas aeruginosa* behaves very differently in the initial phase of bacterial attachment to the surface compared to *S. aureus*. Rogers and colleagues demonstrated that *S. aureus* multiplies and produces a biofilm on the surface that thickens monotonically, becoming hard during growth and softening during starvation. In comparison, *P. aeruginosa* showed much less reproducible behavior; the organism was loosely adhered and detached from the surface after several hours *in vitro*.\textsuperscript{32} Until *P. aeruginosa* attachment is established, this initial motile motion phase may be inhibited or delayed by synergistic bactericidal activity of Ag and either form of TiO$_2$ as we demonstrated.

Our study has several limitations. First, it was conducted in an *in vitro* model that was designed to replicate the human tracheal environment. However, the results may not be applicable to the *in vivo* setting where other factors may impact the ETT surface such as frequent suctioning, nebulizer therapy or host factors. Second, the concomitant use of Ag and TiO$_2$ would seem to be marginally effective against VAP if it does not inhibit a common cause of VAP (ie, *S. aureus*). Third, minimizing the particulate surface that facilitates bacterial binding could be effective by reducing the surface area for bacterial adherence and growth. Fourth, the coating, peeled off under mechanical stress, could be dislodged by a suction catheter and produce potentially harmful Ag particles in the human airway and lung parenchyma. Nevertheless, titanium dioxide is known to be safe in humans and is used in such products as cosmetics, toothpaste, and sunscreen. Hence, the combination of Ag and TiO$_2$ needs to be evaluated in a clinical setting. Lastly, this study did not directly evaluate the attractive feature of TiO$_2$ photocatalysis. The potency of TiO$_2$ either individually or in combination with Ag needs to...
be examined in conjunction with photocatalysis. However, exposing tracheal mucosa to UV light involves potential toxicity. This potential risk and, more importantly, its clinical feasibility, need to be ascertained in future studies.

Conclusion

One strategy to reduce the incidence of VAP is to pretreat ETT with an antiseptic agent to reduce or delay bacterial biofilm formation. The use of TiO$_2$ with Ag-coated ETTs effectively inhibited \textit{P. aeruginosa} growth, but not \textit{S. aureus} growth, in an in \textit{vitro} model of a human trachea environment.

Disclosures

The authors have no conflicts of interest to disclose in this work.

References