

Inhibited growth of *Pseudomonas aeruginosa* by dextran- and polyacrylic acid-coated ceria nanoparticles

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Abstract: Ceria (CeO₂) nanoparticles have been widely studied for numerous applications, but only a few recent studies have investigated their potential applications in medicine. Moreover, there have been almost no studies focusing on their possible antibacterial properties, despite the fact that such nanoparticles may reduce reactive oxygen species. In this study, we coated CeO₂ nanoparticles with dextran or polyacrylic acid (PAA) because of their enhanced biocompatibility properties, minimized toxicity, and reduced clearance by the immune system. For the first time, the coated CeO₂ nanoparticles were tested in bacterial assays involving *Pseudomonas aeruginosa*, one of the most significant bacteria responsible for infecting numerous medical devices. The results showed that CeO₂ nanoparticles with either coating significantly inhibited the growth of *Pseudomonas aeruginosa*, by up to 55.14%, after 24 hours compared with controls (no particles). The inhibition of bacterial growth was concentration dependent. In summary, this study revealed, for the first time, that the characterized dextran- and PAA-coated CeO₂ nanoparticles could be potential novel materials for numerous antibacterial applications.

Keywords: antibacterial, biomedical applications

Introduction

Among the rare earth elements (the 15 lanthanides, along with scandium and yttrium), cerium is the most abundant element, at 66.5 ppm in the Earth's crust compared with copper (60 ppm) or tin (2.3 ppm).^{1,2} As a highly abundant material, ceria (or cerium oxide [CeO₂]) is technologically important due to its wide range of applications, for example, in catalysts for the elimination of toxic auto exhaust gases,^{3,4} oxygen sensors,^{5,6} fuel cells,⁷⁻⁹ electrochromic thin films,^{10,11} and so on. Upon the transition from a macrosized particle to a nanocrystalline particle, CeO₂ significantly changes its physicochemical properties to, usually, possess a high density of nanocrystalline CeO₂ interfaces.¹² Thus, nanostructured CeO₂ has attracted extensive attention in the field of nanotechnology due to improvements in redox properties, transport properties, and high surface to volume ratios, which may be of value for various biological applications.

However, although CeO₂ nanoparticles have been studied for numerous traditional scientific and engineering applications, there have been almost no studies regarding the potential biomedical applications of CeO₂ until recently, when researchers demonstrated that CeO₂ nanoparticles possess antioxidant properties at physiological pH values. The ability of these nanoparticles to act as an antioxidant lies in their ability to reversibly switch from Ce³⁺ to Ce⁴⁺.¹³ In recent studies, this antioxidative ability enabled the application of CeO₂ nanoparticles to protect against radiation damage, oxidative stress, and inflammation.¹⁴⁻¹⁶ For example, Tsai et al reported

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that free radical scavenging could be achieved in murine insulinoma β TC-tet cells by the introduction of crystalline CeO₂ nanoparticles.¹⁷

Moreover, in previous studies, due to safety concerns for biomedical applications, the toxicity of zinc oxide (ZnO), CeO₂, and titanium oxide (TiO₂) was compared based on dissolution and oxidative stress properties.^{18,19} The results showed that the CeO₂ nanoparticles could suppress the generation of reactive oxygen species (ROS) and induce cellular resistance to an exogenous source of oxidative stress, to protect cells from oxidation injury. However, concerns over CeO₂ toxicity, biocompatibility, and avoidance of immune-system clearance remained. Thus, to further improve the clinical use of CeO₂ nanoparticles, previous efforts have coated CeO₂ with biocompatible dextran and PAA; enhanced stability in aqueous solutions has also been reported for these coated CeO₂ nanoparticles.^{15,31}

To build on these exciting medical applications of CeO₂ nanoparticles, the present study investigated, for the first time, the influence of dextran- and PAA-coated CeO₂ nanoparticles on bacteria growth. *Pseudomonas aeruginosa* is an aerobic gram-negative bacterium that is an important cause of various infections, especially in hospitals. Hospitalized patients may be colonized with *P. aeruginosa* on admission or during their hospital stay, and *P. aeruginosa* can be isolated from nearly any conceivable source within hospitals.^{20,21} It accounts for 11%–13.8% of all nosocomial infections when a microbiological isolate is identifiable.²² Infections caused by *P. aeruginosa* are not only common,^{23,24} but are also associated with high morbidity and mortality when compared with other bacterial pathogens.^{25,26} An additional concern involves the increased antimicrobial resistance of nosocomial *P. aeruginosa* isolates.²⁷ Even more troublesome, *P. aeruginosa* attaches to both biotic and abiotic surfaces, forming a biofilm. Once formed, the *P. aeruginosa* infections are much more difficult to treat, due to their formation of exopolysaccharides. Biofilms of *P. aeruginosa* have been found on numerous medical devices, including central venous catheters, contact lenses, mechanical heart valves, intrauterine devices, and indwelling urinary catheters.^{28,29} Thus, it is significant to develop active and biocompatible molecules that do not rely on antibiotics, to which bacteria may develop a resistance, and that kill bacteria at early stages of infections, before biofilm formation.

Therefore in this study, for the first time, due to concerns over the biocompatibility, toxicity, and immune system clearance of pristine CeO₂ nanoparticles, we

investigated the effectiveness of dextran- and PAA-coated CeO₂ nanoparticles in inhibiting bacterial growth, specifically, *P. aeruginosa*. The results showed that such coated CeO₂ nanoparticles significantly inhibited the growth of *P. aeruginosa*, revealing a promising application of CeO₂ nanoparticles to prevent bacterial infections that should be further explored.

Materials and methods

Synthesis, coating, and characterization of the cerium oxide nanoparticles

The dextran- or PAA-coated CeO₂ nanoparticles were synthesized according to previously described procedures.^{15,30,31} Briefly, a solution of cerium (III) nitrate (1.0 M, 99%) (Sigma-Aldrich Corp, St Louis, MO, USA) in water (5.0 mL) was mixed with an aqueous solution of either PAA (0.5 M) (Sigma-Aldrich Corp) or dextran (1.0 M) (Sigma-Aldrich Corp). The resulting mixtures were then added to an ammonium hydroxide solution (30.0 mL, 30%) (Sigma-Aldrich Corp) under continuous stirring for 24 hours, at room temperature. The preparation was then centrifuged at 4000 rpm for two 30-minute cycles to settle down any debris and large agglomerates. The supernatant solution was then purified from free polymers and other reagents and then concentrated using a 30K Amicon® Ultra-4 Centrifugal Filter Units (EMD Millipore Corp, Billerica, MA, USA). The final dextran- and PAA-coated CeO₂ nanoparticle preparations were stored at a 1.5 mg CeO₂/mL concentration in deionized water until further used; such concentrations have previously been shown to be nontoxic to mammalian cells.^{15,30,31}

The various polymer-coated CeO₂ nanoparticle preparations were characterized by dynamic light scattering (DLS) to determine the size, zeta potential, and surface charge of the nanoparticles (Zetasizer; Malvern Instruments Ltd, Malvern, UK). Fourier transform infrared (FTIR) spectrometry analysis was performed on vacuum dried samples to verify the surface functionalities of the nanoparticles (Spectrum™ 100 Optica; PerkinElmer Inc, Waltham, MA, USA).

Bacterial assays

A bacteria cell line of *P. aeruginosa* was obtained in freeze-dried form from the American Type Culture Collection (catalog number 27853; Manassas, VA, USA). The bacterial cells were propagated in 30 mg/mL tryptic soy broth (TSB) (catalog number 22092; Sigma-Aldrich Corp) for 24 hours in an incubator (37°C, humidified, 5% carbon dioxide). When the second passage of bacteria reached its stationary phase,

the second passage was frozen in one part TSB and one part 40% sterile glycerol (catalog number G5516; Sigma-Aldrich Corp). Before bacterial seeding, a sterile 10 μ L loop was used to withdraw bacteria from the prepared frozen stock, the bacteria were streaked onto a TSB Agar plate (catalog number A1296; Sigma-Aldrich Corp), and the TSB agar plate was incubated for 20 hours, to form single bacterial colonies. Then, a single colony was collected using a sterile loop and was incubated in a 15 mL sterile test tube containing 3 mL of TSB, for 20 hours. The bacterial solution was diluted to a concentration of 10^7 bacteria/mL, which was assessed by measuring the optical density of the bacterial solution, using a standard curve correlating optical densities and bacterial concentrations. The optical densities were measured at 562 nm, using a SpectraMax M5 Microplate Reader (Molecular Devices LLC, Sunnyvale, CA, USA).

Two concentrations of both dextran- or PAA-coated CeO₂ nanoparticles were separately tested against bacterial growth. The solutions of CeO₂ nanoparticles were mixed with bacterial solutions (10^7 bacteria/mL) at a 1:10 or 1:5 volume ratio and cultured for either 1, 6, or 24 hours in the incubator. Solutions of sterile TSB were mixed with bacteria solutions as blank controls. After the treatment, the solutions with bacteria were diluted with a phosphate-buffered saline (PBS) (catalog number P4417; Sigma-Aldrich Corp) solution and were spread on agar plates, and the bacteria colonies were counted after 18 hours of incubation.

Bacterial tests were conducted in triplicate and repeated three times. Data were collected, and the differences were assessed with the probability associated with a one-tailed Student's *t*-test. Statistical analyses were performed using Microsoft® Excel® 2010 (Microsoft Corporation, Redmond, WA, USA).

Results and discussion

Cerium oxide nanoparticle characterization

The CeO₂ nanoparticle preparations used in this study were aqueous suspensions of nanoparticles coated with either PAA or dextran. The synthetic procedure used to make these nanoparticles is highly reproducible and yielded nanoparticles with a CeO₂ core of 3–4 nm, by transmission electron microscopy (TEM).^{15,30,31} Specifically, the hydrodynamic diameter of the PAA-coated CeO₂ nanoparticles was 5 nm, while the size for the dextran-coated CeO₂ nanoparticles was 14 nm.^{15,30,31} Confirmation of the polymer coating was accessed by FTIR, as previously described,^{15,30,31} and by zeta potential, confirming the presence of a highly negative charged polymer coating for the PAA-coated CeO₂ nanoparticles ($\zeta = -45$ mV) and a lower negative charge (close to neutral) for the dextran-coated CeO₂ nanoparticles ($\zeta = -2$ mV).

Bacterial assays

As shown in Figures 1 and 2, the CeO₂ nanoparticles did not show significant inhibition of *P. aeruginosa* growth compared with the TSB controls after 1 hour of treatment. However, after 6 hours of treatment, both concentrations of the dextran-coated CeO₂ nanoparticles were found to have significantly inhibited the growth of *P. aeruginosa*, by about 20% compared with the TSB controls ($P < 0.05$). The PAA-coated CeO₂ nanoparticles showed significant inhibition of *P. aeruginosa* growth only for the higher concentration (bacterial solution:CeO₂ nanoparticles = 5:1).

Most importantly, the growth of *P. aeruginosa* was strongly inhibited in the presence of either the dextran- or PAA-coated CeO₂ nanoparticles after 24 hours compared

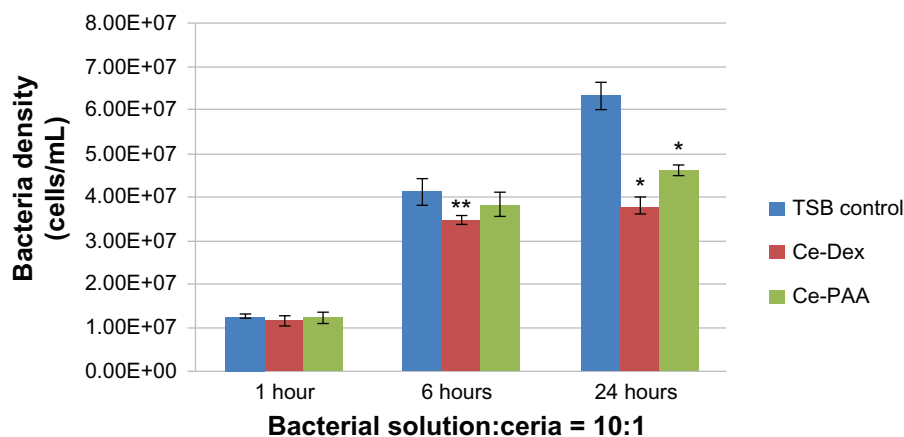


Figure 1 Inhibited *Pseudomonas aeruginosa* growth in the presence of ceria nanoparticles at three time points tested (1, 6, and 24 hours).

Notes: The volume ratio of bacterial solution (10^7 bacteria/mL) and ceria nanoparticles was 10:1. Data is represented as mean \pm standard deviation, $n = 3$; * $P < 0.01$ compared with the TSB control group, after 24 hours of treatment. ** $P < 0.05$ compared with the TSB control group, after 6 hours of treatment.

Abbreviations: Ce, ceria; Dex, dextran; PAA, polyacrylic acid; TSB, tryptic soy broth.

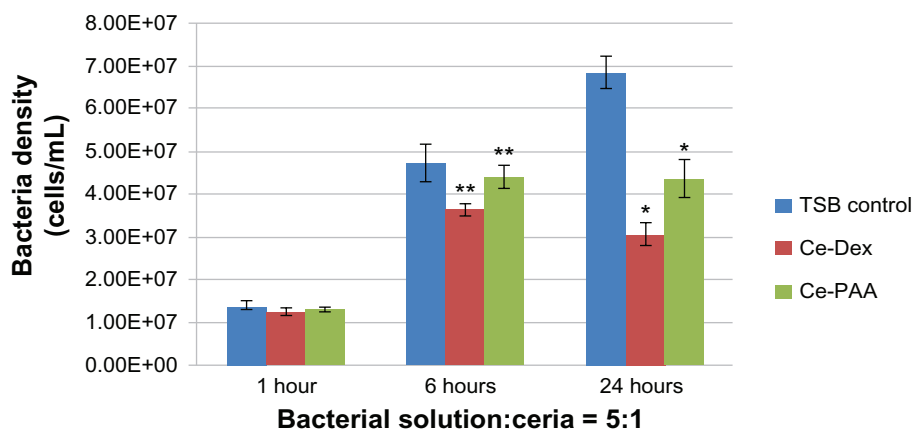


Figure 2 *Pseudomonas aeruginosa* growth in the presence of ceria nanoparticles at three time points tested (1, 6, and 24 hours).

Notes: The volume ratio of bacterial solution (10^7 bacteria/mL) and ceria nanoparticles was 5:1. Data is represented as mean \pm standard deviation, $n = 3$; * $P < 0.01$ compared with the TSB control group, after 24 hours of treatment. ** $P < 0.05$ compared with the TSB control group after, 6 hours of treatment.

Abbreviations: Ce, ceria; Dex, dextran; PAA, polyacrylic acid; TSB, tryptic soy broth.

with the TSB controls ($P < 0.01$). When the volume ratio of bacterial solution to CeO_2 nanoparticles was 10:1, *P. aeruginosa* growth was inhibited by 40.12% and 26.88% in the presence of dextran- and PAA-coated CeO_2 nanoparticles, respectively, compared with the TSB controls. When the concentration of CeO_2 nanoparticles doubled, the inhibition of *P. aeruginosa* growth reached 55.41% and 36.44% for the dextran- and PAA-coated CeO_2 nanoparticles, respectively. Critically, all of this inhibition occurred without the use of antibiotics, with only the use of the CeO_2 nanoparticles.

Based on the results of bacterial assays, the inhibition to *P. aeruginosa* growth by dextran- and PAA-coated CeO_2 nanoparticles was more significant as the treatment time increased to 24 hours. In addition, the dextran-coated CeO_2 nanoparticles showed a stronger inhibition of bacterial growth than did the PAA-coated CeO_2 nanoparticles, especially after treatment for 6 and 24 hours. One possible reason the dextran coating and PAA coating could affect the bacterial growth differently is that the size of the dextran-coated CeO_2 nanoparticles was 14 nm and the size of PAA-coated CeO_2 nanoparticles was 5 nm, which translates to an altered ability to interact with bacteria membranes. In addition, as mentioned, the surface energy of the dextran- and PAA-coated CeO_2 nanoparticles was different, which could also have strongly influenced their interactions with bacteria cell membranes. Moreover, as expected, from 6 hours to 24 hours, the bacteria propagated quickly in the TSB control, but the bacteria numbers decreased even more during this time period with the treatment with both kinds of coated CeO_2 nanoparticles.

Importantly, the mechanism of how dextran- and PAA-coated CeO_2 nanoparticles prevented bacterial growth

remains unexplored. Since CeO_2 nanoparticles have been shown to suppress the generation of ROS,¹⁹ their antibacterial properties demonstrated here, for the first time, could be different from those of many other nanoparticles (such as ZnO nanoparticles, silver [Ag] nanoparticles, and so on) that are believed to kill bacteria mainly through the generation of ROS. Although requiring additional investigation, the possible mechanism for the antibacterial action of the coated CeO_2 nanoparticles formulated here could be that the dextran and PAA coatings interacted with specific proteins on the bacterial cell membranes to alter the permeability of the cell membranes, thus, killing the bacterial cells. Moreover, it is possible that the dextran or PAA coated CeO_2 nanoparticles studied here may have promoted the adsorption, from the TSB, of proteins that are known to inhibit bacteria growth (such as fibronectin, mucin, vitronectin, etc).³² Future studies should also investigate how universal such bacterial inhibition is, by exposing such coated CeO_2 nanoparticles to other bacteria (such as *Staphylococcus aureus* and *Escherichia coli*).

Conclusion

In conclusion, the growth of *P. aeruginosa* was significantly inhibited in the presence of either dextran- or PAA-coated CeO_2 nanoparticles after 24 hours. The effectiveness of bacterial-growth inhibition reached 55.41% for the dextran-coated CeO_2 nanoparticles compared with the TSB control. This study revealed, for the first time, that the characterized coated CeO_2 nanoparticles could potentially serve as a novel antibacterial agent in biomedical applications. Further studies, such as investigation of longer-term treatment and elucidating the mechanism of action, are important

to achieve a better understanding and the application of such coated CeO₂ nanoparticles in numerous antibacterial applications.

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Disclosure

The authors report no conflicts of interest in this work.

References

- Ahrens TJ, editor. *Global Earth Physics: A Handbook of Physical Constants*. Washington, DC: American Geophysical Union; 1995.
- Lide D, editor. *CRC Handbook of Chemistry and Physics*. 88th ed. Boca Raton, FL: CRC Publishing Co; 2007.
- Trovarelli A. Catalytic properties of ceria and CeO₂-containing materials. *Catalysis Reviews*. 1996;38(4):439–520.
- Kašpar J, Fornasiero P, Graziani M. Use of CeO₂-based oxides in the three-way catalysis. *Catalysis Today*. 1999;50(2):285–298.
- Beie HJ, Gnörich A. Oxygen gas sensors based on CeO₂ thick and thin-films. *Sens Actuators B Chem*. 1991;4(3–4):393–399.
- Jasinski P, Suzuki T, Anderson HU. Nanocrystalline undoped ceria oxygen sensor. *Sens Actuators B Chem*. 2003;95(1–3):73–77.
- Park S, Vohs JM, Gorte RJ. Direct oxidation of hydrocarbons in a solid-oxide fuel cell. *Nature*. 2000;404(6775):265–267.
- Sun C, Hui R, Roller J. Cathode materials for solid oxide fuel cells: a review. *J Solid State Electrochem*. 2010;14(7):1125–1144.
- Sun C, Stimming U. Recent anode advances in solid oxide fuel cells. *J Power Sources*. September 27, 2007;171(2):247–260.
- Porqueras I, Person C, Corbella C, Vives M, Pinyol A, Bertan E. Characteristics of e-beam deposited electrochromic CeO₂ thin films. *Solid State Ionics*. 2003;165(1–4):131–137.
- Özer N. Optical properties and electrochromic characterization of sol-gel deposited ceria films. *Solar Energy Materials and Solar Cells*. 2001;68(3–4):391–400.
- Ivanov VK, Usatenko AV, Shcherbakov AB. Antioxidant activity of nanocrystalline ceria to anthocyanins. *Russian Journal of Inorganic Chemistry*. 2009;54(10):1522–1527.
- Tarnuzzer RW, Colon J, Patil S, Seal S. Vacancy engineered ceria nanostructures for protection from radiation-induced cellular damage. *Nano Lett*. 2005;5(12):2573–2577.
- Das M, Patil S, Bhargava N, et al. Auto-catalytic ceria nanoparticles offer neuroprotection to adult rat spinal cord neurons. *Biomaterials*. 2007;28(10):1918–1925.
- Perez JM, Asati A, Nath S, Kaittanis C. Synthesis of biocompatible dextran-coated nanoceria with pH-dependent antioxidant properties. *Small*. 2008;4(5):552–556.
- Lee SS, Zhu H, Contreras EQ, Prakash A, Puppala HL, Colvin VL. High temperature decomposition of cerium precursors to form ceria nanocrystal libraries for biological applications. *Chem Mater*. 2012;24(3):424–432.
- Tsai YY, Oca-Cossio J, Agering K, et al. Novel synthesis of cerium oxide nanoparticles for free radical scavenging. *Nanomedicine (Lond)*. 2007;2(3):325–332.
- Nel A, Xia T, Mädler L, Li N. Toxic potential of materials at the nanolevel. *Science*. 2006;311(5761):622–627.
- Xia T, Kovochich M, Liong M, et al. Comparison of the mechanism of toxicity of zinc oxide and cerium oxide nanoparticles based on dissolution and oxidative stress properties. *ACS Nano*. 2008;2(10):2121–2134.
- Bonten MJ, Bergmans DC, Speijer H, Stobberingh EE. Characteristics of polyclonal endemicity of *Pseudomonas aeruginosa* colonization in intensive care units. Implications for infection control. *Am J Respir Crit Care Med*. 1999;160(4):1212–1219.
- Pirnay JP, De Vos D, Cochez C, et al. Molecular epidemiology of *Pseudomonas aeruginosa* colonization in a burn unit: persistence of a multidrug-resistant clone and a silver sulfadiazine-resistant clone. *J Clin Microbiol*. 2003;41(3):1192–1202.
- Lizioli A, Privitera G, Alliata E, et al. Prevalence of nosocomial infections in Italy: result from the Lombardy survey in 2000. *J Hosp Infect*. 2003;54(2):141–148.
- Rello J, Ollendorf DA, Oster G, et al; VAP Outcomes Scientific Advisory Group. Epidemiology and outcomes of ventilator-associated pneumonia in a large US database. *Chest*. 2002;122(6):2115–2121.
- Kollef MH, Shorr A, Tabak YP, Gupta V, Liu LZ, Johannes RS. Epidemiology and outcomes of health-care-associated pneumonia: results from a large US database of culture-positive pneumonia. *Chest*. 2005;128(6):3854–3862.
- Osmon S, Ward S, Fraser VJ, Kollef MH. Hospital mortality for patients with bacteremia due to *Staphylococcus aureus* or *Pseudomonas aeruginosa*. *Chest*. 2004;125(2):607–616.
- Harbarth S, Ferrière K, Hugonnet S, Ricou B, Suter P, Pittet D. Epidemiology and prognostic determinants of bloodstream infections in surgical intensive care. *Arch Surg*. 2002;137(12):1353–1359; discussion 1359.
- Gaynes R, Edwards JR; National Nosocomial Infections Surveillance System. Overview of nosocomial infections caused by gram-negative bacilli. *Clin Infect Dis*. 2005;41(6):848–854.
- Donlan RM, Costerton JW. Biofilms: survival mechanisms of clinically relevant microorganisms. *Clin Microbiol Rev*. 2002;15(2):167–193.
- Costerton JW, Stewart PS, Greenberg EP. Bacterial biofilms: a common cause of persistent infections. *Science*. 1999;284(5418):1318–1322.
- Asati A, Santra S, Kaittanis C, Nath S, Perez JM. Oxidase-like activity of polymer-coated cerium oxide nanoparticles. *Angew Chem Int Ed Engl*. 2009;48(13):2308–2312.
- Asati A, Santra S, Kaittanis C, Perez JM. Surface-charge-dependent cell localization and cytotoxicity of cerium oxide nanoparticles. *ACS Nano*. 2010;4(9):5321–5331.
- Kawakubo M, Ito Y, Okimura Y, et al. Natural antibiotic function of a human gastric mucin against *Helicobacter pylori* infection. *Science*. Aug 13 2004;305(5686):1003–1006.

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