Antibacterial nanoparticles: enhanced antibacterial efficiency of coral-like crystalline rhodium nanoplates†

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This paper deals with the newly found antibacterial efficiency of coral-like crystalline Rh nanoplates. Rh nanoplates with rough surface morphology synthesized by inverse-directional galvanic replacement exhibited highly enhanced antibacterial efficiency compared to Rh\(^{3+}\) ion and Rh nanospheres. The observed antibacterial efficiency was comparable to Ag nanoplates, a well-known anticancer nanoagent. Results clearly demonstrate that the composition and morphology of a nanostructure play significant roles in antibacterial effects.

Bacteria live everywhere, covering most of where we live. Although some bacteria such as lactobacilli are beneficial to humans, the risks of chronic infection and sepsis caused by a bacterial infection also exist.\(^1\) Although the use of antibiotics is widely accepted as a general countermeasure against infections, drug overuse has brought about the emergence of super bacteria, multidrug-resistant bacterial strains that can be considered as a major crisis.\(^2\) As a new direction in developing antibiotics, approaches using antibacterial nanoparticles have attracted much attention.\(^3\) Typical examples include Gram-positive antibacterial metal oxides such as zinc oxide (ZnO) and titanium dioxide (TiO\(_2\)),\(^4,5\) and Gram-negative antibacterial metal nanoparticles such as silver (Ag) nanoparticles.\(^6\) Antibacterial effects of nanoparticles are known to rely on bacterial cell membrane disruption,\(^7\) reactive oxygen species generation,\(^8\) and intracellular antibacterial effects.\(^9\) For nanoparticles have such abilities, small dimensions and positive surface charge tend to be more favorable, but the exact features of inorganic nanostructures responsible for antibacterial effect remain to be elucidated. Therefore, the discovery of antibacterial nanoparticles based on new elements, morphology, composition, and physicochemical properties still merits efforts.

Synthesis via surface control or post-synthetic treatments are generally utilized to manufacture fascinating nanostructures with various shapes and elemental compositions.\(^10\) Among them, galvanic replacement based on spontaneous redox reactions between oxidation-prone nanoplates and reduction-prone replacing metal cations is generally used, since it is environment-friendly and convenient to perform.\(^11\) In typically such galvanic replacement setups, Ag or copper (Cu) for nanotemplates, and gold (Au), platinum (Pt), and palladium (Pd) for replacing cations, many of which are closely related to antibacterial properties.\(^12\) Therefore, we expected galvanic replacement can be a promising way to develop new antibacterial nanoparticles.

In this study, we identified the antibacterial property of coral-like crystalline rhodium nanoplates (RhNPs), which recently emerged for its new possibilities in biomedical applications.\(^13\) To verify the enhanced antibacterial activity of RhNPs based on the morphology and composition, a comparative experiment was performed using Ag nanoplates (AgNPs), conventionally small Rh nanospheres (RhNSs; 5 nm diameter), and Rh\(^{3+}\) ions (Fig. 1).\(^14\) Quantitative analysis showed that RhNPs exhibited a slightly improved effect when compared to AgNPs (typically used as an antibacterial nanomaterial),\(^15\) and free Rh ions and Rh nanospheres were not effective. The results not only demonstrate a new possibility of using metal elements beyond Ag for antibacterial nanostructures, but also show that structural properties play an important role.

RhNP preparation was accomplished using a two-step procedure: seed-mediated synthesis of sacrificial Ag nanoplate templates, followed by inverse-directional galvanic replacement with Rh\(^{3+}\). The high-temperature (190 °C) reaction along with the reducing agent and ethylene glycol (EG) co-solvent condition, which caused galvanic replacement depending on the shape of the AgNPs. A subsequent outward surface growth resulted in the formation of coral-like RhNPs. Similarly, control RhNS synthesis was accomplished via reductant-based synthesis under the same conditions without Ag

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†Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra00214f
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Coral-like nanoplates with high surface-to-volume ratio and many surface irregularities have a high potential for intimate interactions with bacterial or mammalian cell membranes.

The extinction spectrum of RhNP identified through UV-visible-near infrared spectroscopy showed a weak UV-localized plasmon resonance peak at 273 nm and uniformly high optical density over the whole wavelength region with weak and broad bands at around 320–1100 nm, which probably originated from the coral-like two-dimensional interconnected complex structure (Fig. 2b). The hydrodynamic diameter of RhNP was observed to be 68.1 nm for maximum number distribution, which is smaller than that measured using the TEM images, because calculations carried out using the Stocks–Einstein equation assume a spherical shape. Although the polydispersity index (PDI) was excellent (0.146 ± 0.016), it did not satisfy the recommended parameters for antibacterial nanoparticle efficacy due to the highly negative surface charge (ζ-potential: −27.1 ± 0.3) and relatively large sizes (120 nm; Fig. 2c).

High-resolution (HR)-TEM observations and fast Fourier transforms of the images provided highly crystalline face-centered cubic (fcc) structures (Fig. S2†). Moreover, a slight shift from the X-ray diffraction pattern was observed possibly due to the partially-presenting Rh–Ag alloy composition (Fig. 2d).

To investigate the antibacterial effect of RhNP further, three additional control materials were prepared and their antibacterial properties against Escherichia coli were tested. AgNPs, a well-known antibacterial material and a template for galvanic replacement, were prepared using the same particle count as RhNPs to identify constituent element-based differences. In addition, Rh³⁺ ions and RhNS were prepared with the same Rh content via inductively-coupled plasma mass spectroscopy. All nanoparticles were prepared at a similar degree of surface negative charge, which excluded the antibacterial effect from the charge property (Fig. S3 and S4†).

Compared to untreated control (negative), RhNPs showed remarkable inhibition of E. coli growth; the inhibition was even slightly better than AgNPs (positive) when treated at the same concentration. Contrastingly, free Rh³⁺ ions and RhNSs did not show noticeable effects even though they had the same Rh contents. From these results, both the constituent element and morphological characteristics were confirmed to be crucial factors (Fig. 3a). Similar results were also obtained for the dose-dependent antibacterial effects of RhNPs (Fig. 3b).

SEM images of RhNP-treated E. coli provided more detailed information regarding the observed antibacterial effects. Surface adsorption of a large number of nanoplates was observed in E. coli treated with RhNPs when compared to control (Fig. 4a and b). Magnified SEM images (Fig. 4c and d) show that RhNPs were adsorbed in a form of clusters on the surfaces of E. coli. The results also indicated that RhNPs bigger than a certain size were required for interaction with E. coli surfaces, and that relatively small RhNPs were ineffective due to difficulties in accessing and interacting with the E. coli surface. Negatively-charged particles may seem hard to adhere to cells due to the negatively-charged cell membrane, so the adhesion

**Fig. 1** Schematic illustration of the antibacterial effect of Rh nanoparticles in spherical and coral-like plate morphologies.

**Fig. 2** Characterization of prepared RhNPs. (a) TEM, (b) UV-Vis spectrometer, (c) DLS and zeta-potential, and (d) XRD pattern exhibited the successful synthesis of crystalline monodisperse RhNPs. The scale bar is 200 nm.
presumably occurred between RhNPs and protein corona clusters. Such adsorption of RhNPs resulted in cell membrane weakening (Fig. 4c and d; red arrows) or disruption (Fig. 4d; yellow arrow), which account for the observed antibacterial property.

Conclusion

In this study, we reported the antibacterial effect of RhNPs that varies depending on their chemical composition and morphology. Recent studies based on RhNPs with contact lengths of 100 nm or more showed interesting results. Notably, previous studies reported difficulties in synthesizing Rh nanoparticles above 5 nm. Although the actual antibacterial effects and partial mechanisms of RhNP were identified in the present study, specific factors for the RhNP antibacterial properties were not elucidated. However, it is important to continue exploring the biological applications of Rh, because the antibacterial effects of nanomaterials still remain unclear.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Grant No. (NRF-2016R1C1B1008090) through the National Research Foundation of Korea (NRF) and a grant from Kyung Hee University in 2016 (KHU-20160610).

Notes and references


