Core–Shell Polydopamine/Cu Nanometer Rods Efficiently Deactivate Microbes by Mimicking Chloride-Activated Peroxidases

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ABSTRACT: Cu-modified nanoparticles have been designed to mimic peroxidase, and their potent antibacterial and anti-biofilm abilities have been widely investigated. In this study, novel core–shell polydopamine (PDA)/Cu(OH)$_2$SO$_4$ crystal (PDA/Cu) nanometer rods were prepared. The PDA/Cu nanometer rods show similar kinetic behaviors to chloride-activated peroxidases, exhibit excellent photothermal properties, and are sensitive to the concentrations of pH values and the substrate (i.e., H$_2$O$_2$). PDA/Cu nanometer rods could adhere to the bacteria and catalyze hydrogen peroxide (H$_2$O$_2$) to generate more reactive hydroxy radicals (*OH) against Staphylococcus aureus and Escherichia coli. Furthermore, PDA/Cu nanometer rods show enhanced catalytic and photothermal synergistic antibacterial activity. This work provides a simple, inexpensive, and effective strategy for antibacterial applications.

INTRODUCTION

Antibiotic-resistant bacterial infections are posing a serious threat to public health. Bacteriostatic antibiotics act on bacteria by targeting and disrupting their basic survival (bactericidal) and multiplication (bacteriostatic) processes. However, microorganisms rapidly evolve antibiotic resistance through gene mutation and horizontal gene transfer, changes in antibiotic targets, or modification of the antibiotic agents. Therefore, Cu-modified nanoparticles with efficient antibacterial properties were developed, and they use multiple mechanisms simultaneously to kill and/or inhibit the growth of microorganisms, overcoming microbial resistance mechanisms. Cu-modified nanoparticles were recently reported to exhibit peroxidase-like activity and consequently could conceivably be used as antibacterial materials. Nanoparticles working as artificial enzymes keep their catalytic activity in extreme pH and temperature conditions and have higher stability and lower price and are easier to be stored compared with natural enzymes. The peroxidase-like catalytic mechanism of Cu-based nanomaterials is generally summarized as a Fenton-like reaction. Briefly, Cu$^{2+}$ ions participate in catalyzing the conversion of hydrogen peroxide (H$_2$O$_2$) to hydroxyl radicals (*OH). *OH is much more effective against microorganisms than H$_2$O$_2$, since H$_2$O$_2$ is less reactive and can be detoxified by endogenous antioxidants. Accelerated rates of *OH production correlate remarkably with antibacterial activities.

In addition to Cu or Cu compound nanomaterials being used as nanozymes, it was also reported that the adhesion of Cu$^{2+}$ on organic nanomaterials significantly improved the enzyme-like catalytic activity of nanozymes and enhanced synergistic antibacterial activity. Polydopamine (PDA) has been recently widely used in antibacterial composite materials, benefiting from its advantages of easy synthesis, good biocompatibility, strong adhesion, excellent photothermal properties, and unique antibacterial ability. It was reported that PDA was used to modify the surface of nanomaterials to improve their antibacterial effect and ability to target the bacteria membrane. Moreover, the photothermal effect of PDA provided the composite nanozymes with additional photothermal therapy and near-infrared (NIR)-enhanced catalytic activity to augment their antibacterial activity.

In this study, we developed a simple method to prepare core–shell PDA/Cu$_4$(OH)$_6$SO$_4$ crystal (PDA/Cu) nanometer rods. PDA/Cu nanometer rods showed Cl$^-$-accelerated peroxidase-like activity. In phosphate-buffered saline (PBS) buffer containing sufficient Cl$^-$, the antibacterial activity of PDA/Cu and H$_2$O$_2$ against Staphylococcus aureus and Escherichia coli was much higher than that of H$_2$O$_2$ alone. Furthermore, PDA/Cu nanometer rods have an excellent photothermal property compared with non-PDA Cu$_4$(OH)$_6$SO$_4$ crystals, which endowed them with enhanced catalytic and photothermal synergistic antibacterial activity.
RESULTS AND DISCUSSION

Synthesis and Characterization of PDA/Cu Nanometer Rods. PDA/Cu nanometer rods were synthesized via a simple method. The green suspension of Cu$_4$(OH)$_6$SO$_4$ crystals (brochantite) was prepared in the aqueous mixture of CuSO$_4$$\cdot$5H$_2$O and NaAc (pH = 5), in a molar proportion of Cu$^{2+}$/Ac$^- = 1/6$. The excessive addition of Ac$^-$ was to keep the aqueous solution in slightly acidic conditions to fit with the narrow stability field of brochantite. Subsequently, DA·HCl was added, and the green suspension turned brown immediately, reflecting the formation of PDA. The X-ray diffraction (XRD) pattern of the obtained Cu$_4$(OH)$_6$SO$_4$ crystals is shown in Figure 1a. The characteristic peaks of the obtained crystals are consistent with the reference brochantite, indicating that the main component of the green suspension is Cu$_4$(OH)$_6$SO$_4$ crystals. Moreover, it was concluded from the slight difference between the XRD patterns of the Cu$_4$(OH)$_6$SO$_4$ crystals and PDA/Cu nanometer rods that the coating with PDA did not disrupt the structure of the Cu$_4$(OH)$_6$SO$_4$ crystals greatly. As shown in the transmission electron microscopy (TEM) images and scanning electron microscopy (SEM) images (Figure 1b–d), the core–shell PDA/Cu nanometer rods had a length of 1–2 μm and a maximum width of 200 nm, with the thickness of the PDA...
coating of about 30 nm. The dynamic light scattering (DLS) study suggested that the PDA/Cu nanometer rods had a hydrodynamic diameter of 1237 ± 20 nm. The formation process of PDA/Cu nanometer rods was additionally monitored by SEM. According to the SEM images of PDA/Cu nanometer rods, the Cl⁻ promoted the etching of Cu nanometer rods and Cu⁺ alone was a weak oxidant under acidic conditions, the Cl⁻ from DA-HCl facilitated the reaction. This is because Cl⁻ increases the positive redox potential of Cu²⁺/Cu⁺, indicating that the complete decomposition was achieved and the decomposition product was CuO₂⁻, with a mass percentage of 43.2%.

**Figure 3.** (a) UV-vis absorption spectra of colorimetric substrates: (1) TMB + NH₄Cl + H₂O₂, (2) TMB + NH₄Cl + PDA/Cu, (3) TMB + H₂O₂ + PDA/Cu, (4) TMB + NH₄Cl + H₂O₂ + PDA/Cu, (5) OPD + NH₄Cl + PDA/Cu, (6) OPD + NH₄Cl + PDA/Cu, (7) OPD + H₂O₂ + PDA/Cu, (8) OPD + NH₄Cl + H₂O₂ + PDA/Cu (TMB: 0.5 mM, OPD: 0.5 mM, NH₄Cl: 100 mM, H₂O₂: 100 mM, PDA/Cu: 50 μg mL⁻¹, solution: 0.1 M 2-(N-morpholino)ethanesulfonic acid (MES) buffer, pH 5.5). (b) Kinetics of the TMB oxidation as a function of the H₂O₂ concentration. The lines show the linear data interpolation of the first 60 s.
requirement of oxidative power. Moreover, when Cl\textsuperscript{−} was added to the mixture containing TMB, PDA/Cu nanometer rods, and H\textsubscript{2}O\textsubscript{2}, more oxidized TMB was obtained, indicated by the higher absorption peaks at 370 and 652 nm. The same results were obtained from the absorption spectra of OPD. These results suggested that PDA/Cu nanometer rods had a peroxidase-like catalytic ability, and the addition of Cl\textsuperscript{−} accelerated this process. The Cl\textsuperscript{−}-accelerated mechanism could be explained by the fact that the coordination of Cl\textsuperscript{−} stabilized Cu\textsuperscript{2+} more than Cu\textsuperscript{2+}. This led to an enhanced formation of Cu\textsuperscript{3+} from Cu\textsuperscript{2+}, which was the rate-determining step of the Cu\textsuperscript{3+}-catalyzed decomposition of H\textsubscript{2}O\textsubscript{2}\textsuperscript{10,30}.

The absorbance at 652 nm was considered as a measure of the progress of TMB oxidation. The initial stage was linearly approximated and the slope was considered as the initial reaction rate (see Figure 3b). The plots of the initial reaction rates as a function of the concentrations of TMB and H\textsubscript{2}O\textsubscript{2} clearly followed a Michaelis–Menten kinetic (see Figure S4 in the SI). The kinetic parameters of K\textsubscript{m} and V\textsubscript{max} were obtained from the nonlinear parameter fitting of the Michaelis–Menten equation to the experimental data. Table 1 compared the kinetic parameters of PDA/Cu nanometer rods with horseradish peroxidase (HRP) and other peroxidase mimics. These results show that PDA nanometer rods have higher V\textsubscript{max} for both H\textsubscript{2}O\textsubscript{2} and TMB than other nanoenzymes.

### Temperature, pH, and Solution-Dependent Peroxidase-Like Activity

Selected factors that may affect the peroxidase-like catalytic activity of PDA/Cu nanometer rods were explored. For that purpose, the initial rate versus Cl\textsuperscript{−} concentrations were analyzed (Figure 4a). A linear Cl\textsuperscript{−}-accelerating effect on the peroxidase-like activity of PDA/Cu nanometer rods in the Cl\textsuperscript{−} concentrations in the range from 10\textsuperscript{−3} to 125 mM was observed. Figure 4b shows the catalytic activity as a function of pH in the range of 3.0–8.0, with an optimum between 4.5 and 5.0.

To study the relationship between catalytic activity and temperature, we chose OPD as the chromogenic substrate instead of TMB, which had two oxidation products at 60 °C with different UV–vis absorption spectra (see Figure S5 and Scheme S1a in the SI). In contrast, OPD provided a color response that maintained its maximum absorption peak without unacceptable changes at elevated temperatures (Figure 5a and Scheme S1b). PDA/Cu nanometer rods showed a higher initial rate at higher temperatures between 25 and 65 °C, with a good linear relation between the natural logarithm of the kinetic constant and the reciprocal temperature (Figure 5b). The data fitted well with the Arrhenius equation, and the calculated activation energy of the reaction was 78.48 ± 2.86 kJ mol\textsuperscript{−1} (Figure 5c). The final absorbance of H\textsubscript{2}O\textsubscript{2} + OPD was much lower than that of the H\textsubscript{2}O\textsubscript{2} + OPD + PDA/Cu system or the H\textsubscript{2}O\textsubscript{2} + OPD + PDA/Cu + NH\textsubscript{3}Cl system (see Figure S6 in the SI), indicating the lower catalytic activity of the H\textsubscript{2}O\textsubscript{2} + OPD system. Compared with the H\textsubscript{2}O\textsubscript{2} + OPD + PDA/Cu system (E\textsubscript{a} = 84.49 ± 4.24 kJ mol\textsuperscript{−1}), the OPD + NH\textsubscript{3}Cl + H\textsubscript{2}O\textsubscript{2} + PDA/Cu system (E\textsubscript{a} = 78.48 ± 2.86 kJ mol\textsuperscript{−1}) had faster kinetics at the same temperature, and the reaction mechanism required lower activation energy.

The above-mentioned results indicated that PDA/Cu nanometer rods had higher stability at extreme temperature conditions and were easier to be stored compared with natural enzymes.\textsuperscript{35} Additionally, PDA/Cu nanometer rods showed higher V\textsubscript{max} than other nanoenzymes and had higher catalytic activity at higher temperatures, while some other nanoenzymes had optimum activity at a temperature of around 40 °C\textsuperscript{31,36} or showed stable catalytic activity in a wide temperature range.\textsuperscript{7}

The peroxidase-like activity of PDA/Cu nanometer rods depended on the applied buffer solution (Figure 5d and Table S1 in the SI). The catalytic activity of PDA/Cu nanometer rods in the MES buffer was higher than that in the BR buffer at the same pH. This was because Ac\textsuperscript{−} was tightly adsorbed on Cu\textsuperscript{2+} and resulted in a negatively charged surface. In contrast, 2-(N-morpholino)ethanesulfonic acid (MES) hardly was bound to Cu\textsuperscript{2+} at pH < 7.\textsuperscript{37} The catalytic activity of the PDA/Cu nanometer rods in the PBS buffer (pH 5.0) reached a particularly high level due to the high concentration of Cl\textsuperscript{−} in the PBS buffer.

### Peroxidase-Like Catalytic Antimicrobial Activity

Two samples of potentially pathogenic bacteria (the Gram-positive bacteria S. aureus and the Gram-negative bacteria E. coli) were chosen to evaluate the antibacterial activity of the PDA/Cu + H\textsubscript{2}O\textsubscript{2} + Cl\textsuperscript{−} system. The experiments were performed in the PBS buffer at pH 5.0 because PDA/Cu showed the highest catalytic activity at pH 5.0 (Figure 4b), and the high...
concentration of Cl\(^-\) in the PBS buffer greatly increased the Cl\(^-\)-accelerated peroxidase-like activity (Figure 5d). The PBS buffers of pH 5.0 and 7.4 had no significant influence on the survival of both sample bacterial strains, as demonstrated by colony-forming units (CFUs) after incubation for 1 h at 37 °C (see Figure S7 in the SI).

Agar plates were used to determine the survival of bacteria after incubation with PDA/Cu nanometer rods and H\(_2\)O\(_2\) for 1 h. As shown in the photographs of agar plates (Figure 6a), both the growth of *S. aureus* and *E. coli* were suppressed by PDA/Cu nanometer rods in combination with H\(_2\)O\(_2\), whereas both agents alone had no such significant effect.

SEM images were taken to visualize the changes in bacterial membrane and structure after exposure to different treatments (Figure 6b). After incubation with PDA/Cu nanometer rods, both *S. aureus* and *E. coli* had a smooth membrane and an intact cell structure consistent with the control, indicating that PDA/Cu nanometer rods did not cause fatal damage to the bacteria. In contrast, partial deformation and shrinkage of bacteria appeared after incubation with H\(_2\)O\(_2\). Furthermore, in the group of PDA/Cu nanometer rods with H\(_2\)O\(_2\), the bacteria showed major damage and lost their intact spherical-like or rod-like shape. The adhesion of bacteria to PDA/Cu nanometer rods was also observed, which would facilitate the rapid and precise attack of *OH* from the surface of PDA/Cu nanometer rods to the cell membrane.

A plate-counting method was applied for quantifying the influence of the different agents on the survival of the bacteria. PDA/Cu nanometer rods in the concentration range of 0–20 μg mL\(^{-1}\) alone did not show high antibacterial activity (Figure 6c,d). After the addition of H\(_2\)O\(_2\), the number of viable bacteria decreased significantly with the increase in the PDA/Cu nanometer rod concentrations. The survival rates of both *S. aureus* and *E. coli* were less than 0.01% when the concentration of PDA/Cu nanometer rods was 10 μg mL\(^{-1}\). H\(_2\)O\(_2\) was able to kill 99.9% of *S. aureus* and *E. coli* at concentrations of 100 and 10 mM, respectively (Figure 6e,f).

Compared with *E. coli*, *S. aureus* required higher concentrations of H\(_2\)O\(_2\) to effectively inactivate bacteria. The remarkable H\(_2\)O\(_2\) resistance of *S. aureus* was known to be caused by antioxidant enzymes (superoxide dismutases, catalases, and peroxiredoxins), which participated in the detoxification of O\(_2^-\) and H\(_2\)O\(_2\) but not of *OH*.\(^{15,38}\) The PDA/Cu nanometer rods catalyzed the conversion of H\(_2\)O\(_2\) to the potent antibacterial agent *OH*. Approximately 20 mM H\(_2\)O\(_2\) was required for *S. aureus* and 5 mM for *E. coli* to achieve a similar antibacterial activity.

To clarify whether the PDA/Cu nanometer rods or the released Cu\(^{2+}\) played a major role in the antibacterial process, the release of Cu\(^{2+}\) in the PBS buffer (pH 5.0) was quantified by photometric analysis with the chromogenic probe dicyclohexanone oxaly dihydrazone using the absorbance at 540 nm (Figure 7a).\(^{53,39}\) According to TGA and DTG results, the molar concentration of Cu in 1 mg mL\(^{-1}\) PDA/Cu nanometer rods was 5.4 mM (Figure 2d). In contrast, the absorbance of filtrates of PDA/Cu nanometer rods after different incubation times was much lower than that of 0.6 mM Cu\(^{2+}\), indicating that very little Cu\(^{2+}\) was released into the solution.

The antibacterial activity of recollected PDA/Cu nanometer rods from the PBS buffer (pH 5.0) was tested. The nearly identical CFU counts results between PDA/Cu nanometer rods and the recollected ones indicated that the antimicrobial activity mainly remained. In conclusion, the recollected PDA/Cu nanometer rods, after being dispersed in the PBS buffer (pH 5.0), had equal enzyme mimicry-assisted antimicrobial activity to kill 99.9% of *S. aureus* and *E. coli* (Figure 7b). These results demonstrated that PDA/Cu nanometer rods adhered to the bacteria and generated efficiently *OH* in the presence of H\(_2\)O\(_2\), causing rapid damage to the bacteria.

**Photothermal and Catalytic Synergistic Antibacterial Therapy.** Numerous studies have reported PDA and PDA composites as photothermal materials with a strong ability to
absorb near-infrared light and convert light energy to heat.\textsuperscript{40} It was coordinated with the properties of PDA/Cu nanometer rods that exhibited higher catalytic activity at higher temperatures (Figure 5c). Therefore, we studied the photothermal properties of PDA/Cu nanometer rods. The PDA/Cu nanometer rods displayed a monotonously changing UV–vis absorption spectrum (Figure 8a), with a peak at 270 nm and smaller shoulders at 370, 460, and 660 nm, the shoulders could be related to the partially oxidated state of the metal-oxidized polydopamine.\textsuperscript{23,41} Moreover, compared to the Cu\textsubscript{2}SO\textsubscript{4}·(OH)\textsubscript{6} crystals (Figure S8a in the SI), PDA/Cu nanometer rods displayed a remarkably higher NIR absorption, which was contributed by the PDA layer and was favorable for photothermal therapy. With the NIR irradiation at 808 nm (0.5 W cm\textsuperscript{-2}), the temperature of PDA/Cu nanometer rods increased over time (Figure 8b). The changes in temperatures

Figure 6. (a) Typical photographs of the agar plates of \textit{S. aureus} and \textit{E. coli} (both about 5 \times 10\textsuperscript{7} CFU mL\textsuperscript{-1}) after incubation with different agents with the following concentrations (PDA/Cu nanometer rods: 20 μg mL\textsuperscript{-1}, \textit{H\textsubscript{2}O\textsubscript{2}}: 50 mM (\textit{S. aureus}) or 5 mM (\textit{E. coli})). In the control experiments, nothing was added. (b) SEM images of \textit{S. aureus} and \textit{E. coli} (both about 1 \times 10\textsuperscript{8} CFU mL\textsuperscript{-1}) after incubation with different agents with the following concentrations (PDA/Cu nanometer rods: 40 μg mL\textsuperscript{-1}, \textit{H\textsubscript{2}O\textsubscript{2}}: 200 mM (\textit{S. aureus}) or 50 mM (\textit{E. coli})), PDA/Cu nanometer rods were marked with blue arrows and cell membrane damage was marked with red arrows. Colony-forming units (CFUs) of \textit{S. aureus} (c) and \textit{E. coli} (d) after incubation with different concentrations of PDA/Cu nanometer rods with and without \textit{H\textsubscript{2}O\textsubscript{2}} (50 and 5 mM, respectively). CFUs of \textit{S. aureus} (e) and \textit{E. coli} (f) after incubation with different concentrations of \textit{H\textsubscript{2}O\textsubscript{2}} with or without PDA/Cu nanometer rods (20 μg mL\textsuperscript{-1}).
Figure 7. (a) Release of Cu$^{2+}$ from PDA/Cu nanometer rods. (b) CFU of S. aureus and E. coli after incubation with H$_2$O$_2$ (50 and 5 mM, respectively), PDA/Cu nanometer rods (20 $\mu$g mL$^{-1}$) + H$_2$O$_2$, and the recollected PDA/Cu nanometer rods (PDA/Cu (r), 20 $\mu$g mL$^{-1}$) + H$_2$O$_2$.

Figure 8. (a) UV–vis absorption spectra of PDA/Cu nanometer rods with different concentrations. (b) Photothermal effects of PDA/Cu nanometer rods at different concentrations at 2.0 W cm$^{-2}$ ($\lambda$ = 808 nm) and (c) under different power densities (200 $\mu$g mL$^{-1}$). Photothermal and catalytic synergistic antibacterial properties: (d) typical photographs of S. aureus and E. coli agar plates. (e, f) CFU of S. aureus and E. coli (PDA/Cu nanometer rods: 80 $\mu$g mL$^{-1}$, H$_2$O$_2$: 20 mM (S. aureus) or 5 mM (E. coli), irradiation laser: 808 nm, 40 mW cm$^{-2}$; all groups were incubated for 20 min).
were dependent on the concentrations of PDA/Cu nanometer rods and the power densities (Figure S8b,c). As shown in Figure S8b,c in the SI, the photothermal conversion efficiency (η) of PDA/Cu nanometer rods was 22.39%, which was greatly higher than that of pure Cu(OH)₂SO₄ crystals (0.14%).

With temperature-dependent peroxidase-like activity and photothermal performance, the photothermal and catalytic synergistic antibacterial activity of PDA/Cu nanometer rods were investigated. The synergistic antibacterial effect of PDA/Cu nanometer rods against bacteria was clearly observed through the typical photographs of the agar plate incubated with S. aureus and E. coli (Figure 8d). The results indicated that Figure 6 PDA/Cu nanometer rods had enhanced nanzyme-photothermal antibacterial activity, efficiently killing bacteria in 20 min (Figure 8e,f).

**CONCLUSIONS**

In summary, core–shell PDA/Cu nanometer rods were obtained with the Cu₄(OH)₂SO₄ crystal core and PDA coating. The formation of Cu₄(OH)₂SO₄ crystals and polymerization of DA was carried out in a weakly acidic aqueous solution, accompanied by the reduction of Cu²⁺ ions. The obtained PDA/Cu nanometer rods acted similar to the Cl⁻-activated peroxidase with the activation energy of 78.48 ± 2.86 kJ mol⁻¹. PDA/Cu nanometer rods showed tunable peroxidase-like catalytic activity, which could be easily enhanced by increasing the temperature as well as increasing the concentration of Cl⁻. Further studies demonstrated that PDA/Cu nanometer rods showed enhanced catalytic and photothermal synergistic antibacterial activity. In addition, PDA/Cu nanometer rods potentially functionalize material surfaces for efficient synergistic antibacterial properties due to the unique ability of the PDA coating to deposit on almost all inorganic and organic substrates.

**EXPERIMENTAL SECTION**

**Materials.** CuSO₄·5H₂O (99.0%), o-phenylenediamine (OPD, 98.5%), H₂O₂ (30.0%), ammonium chloride (NH₄Cl, 99.5%), sodium phosphate dibasic dodecahydrate (Na₂HPO₄·12H₂O, 99.0%), potassium phosphate monobasic (KH₂PO₄, 99.5%), sodium chloride (NaCl, 99.5%), potassium chloride (KCl, 99.5%), acetic acid (HAc, 99.5%), phosphoric acid (H₃PO₄, 85%), boric acid (H₃BO₃, 99.5%), sodium hydroxide (NaOH, 96%), and hydrochloric acid (HCl, 36.0–38.0%) were purchased from Sinopharm Chemical Reagent Co., Ltd., Shanghai. Sodium acetate (NaAc, 99%) was bought from Acrros Organics, Belgium. DA-HCl (98%) and 3,3′,5,5′-tetramethylbenzidine (TMB) (98%) were purchased from Aláddin Industrial Inc, Shanghai. 2-(N-Morpholino)-ethanesulfonic acid (MES, 99%) was purchased from Macklin Biochemical Co. Ltd. Shanghai. All these chemicals were used directly without any further purification. All aqueous solutions were prepared in ultrapure water with a resistivity of 18.2 MΩ cm⁻¹ (Millipore Simplicity). The composition and the pH range of MES, Britton–Robinson (BR), and phosphate-buffered saline (PBS) buffer solutions are listed in Table S1 in the SI.

**Characterization.** The morphologies of Cu₄(OH)₂SO₄ crystals and PDA/Cu nanometer rods were characterized by transmission electron microscopy (TEM) on JEM-2100 Plus (JOEL, Japan) and scanning electron microscopy (SEM) on Zeiss Merlin Compact (Oxford, U.K.). The dynamic light scattering (DLS) measurements were performed on Malvern Instruments Zetasizer Nano (Malvern Instruments, U.K.) at 25 °C. TGA and DTG analysis of PDA/Cu nanometer rods were performed on a Mettler-Toledo TGA2 (Mettler-Toledo, Switzerland) instrument by heating 10 mg of PDA/Cu nanometer rods at a rate of 10 °C min⁻¹ from 30 to 1000 °C in a flow of air. The X-ray diffraction (XRD) measurements were recorded on an Xpert Pro X-ray diffractometer (Panaco, the Netherlands). The Fourier transform infrared (FT-IR) was conducted on a Nicolet IS10 Fourier transform infrared spectroscope (Thermo Fisher Scientific). The X-ray photoelectron spectra (XPS) were carried out on an Escalab 250Xi photoelectron spectrometer (Thermo Fisher Scientific). The photothermal properties were measured with a NIR laser (Hi-Tech Optoelectronics Co. Ltd., China).

**Preparation of PDA/Cu Nanometer Rods.** PDA/Cu nanometer rods were prepared as previously described with modifications. In a typical procedure, an aqueous solution of CuSO₄·SH₂O (4 mM) and NaAc (24 mM) was kept at 55 °C with gentle stirring for 1 h to prepare the bladed Cu₄(OH)₂SO₄ crystals. During this time the mixture changed from a blue solution to a green suspension (at pH 5.0). Then, DA-HCl (2 g L⁻¹) was added, and the mixture was kept at 55 °C for 9 h with gentle stirring to prepare PDA/Cu nanometer rods. To analyze the kinetics of the growth of the PDA coating, the samples were collected at 0.5, 1, and 3 h after the addition of DA-HCl for further characterization. All these samples were purified by repeated centrifugation (12 000 rpm, 8 min) and dried via the freeze-drying process (~60 °C, 1 Pa, 48 h) on an LGI-10 vacuum freeze dryer (Beijing Songyuan Huaxing Technology Develop Co. Ltd., China). Two grams per liter DA-HCl was added to the Tris–HCl buffer (0.1 M, pH 8.5) and stirred for 12 h to prepare self-polymerized PDA.

**Peroxidase-Like Activity of PDA/Cu Nanometer Rods and Kinetic Assay.** The peroxidase-like activity measurements were carried out in an MES aqueous buffer (pH 5.5) at 25 °C, monitoring the absorbance changes at 652 nm every 10 s over 3 min on a Hitachi U-2900 UV–vis spectrophotometer (Hitachi High-Tech, Japan) with a circulating bath. To calculate the kinetic constants, experiments were performed by determining the concentrations of H₂O₂ (100 mM), NH₄Cl (100 mM), and PDA/Cu nanometer rods (50 μg mL⁻¹) and changing the concentration of TMB (0.05–0.6 mM) or by determining the concentrations of TMB (0.5 mM), NH₄Cl (100 mM), and PDA/Cu nanometer rods (50 μg mL⁻¹) and changing the concentration of H₂O₂ (10–120 mM).

The kinetic parameters V max and K m were calculated by fitting to the Michaelis–Menten equation (eq 1)

\[ v = \frac{V_{\text{max}}[S]}{K_m + [S]} \]  

\[ [S] \] is the substrate concentration, V max is the maximum reaction rate, and K m is the Michaelis constant, v is the initial reaction rate. Briefly, v = k / e, k is the slope of the linear change in absorbance and e is the molar absorption coefficient of a colorimetric substrate.

**Temperature, pH, and Solution-Dependent Peroxidase-Like Activity.** The effect of Cl⁻ concentration on catalytic activity was performed with fixed concentrations of TMB (0.5 mM), H₂O₂ (100 mM), and PDA/Cu nanometer rods (50 μg mL⁻¹).

The pH-dependent peroxidase-like catalytic activity was carried out at 25 °C in BR buffers with different pH values, and
the concentrations of TMB, H₂O₂, PDA/Cu nanometer rods, and NH₄Cl were 0.5 mM, 50 mM, 20 μg mL⁻¹, and 50 mM, respectively. The temperature-dependent peroxidase-like catalytic activity of PDA/Cu nanometer rods toward OPD was measured in the BR buffer (pH 5.0), with the fixed concentrations of OPD (0.5 mM), H₂O₂ (20 mM), PDA/Cu nanometer rods (20 μg mL⁻¹), and NH₄Cl (20 mM). The parameters Eₐ was calculated by fitting to the Arrhenius equation (eq 2).

\[
\ln k_T = -\frac{E_a}{RT} + \ln A
\]  
(2)

R, T, Eₐ, A, and \( k_T \) are the molar gas constant, the temperature, the activation energy, the frequency factor, and the rate constant, respectively.

**Peroxidase-Like Catalytic Antimicrobial Activity.** The antibacterial activity was measured against the Gram-positive bacteria *S. aureus* (*S. aureus* ATCC 25923) and the Gram-negative bacteria *E. coli* (*E. coli* DSM 4230) in the PBS buffer (pH 5.0, adjusted with 2 mol L⁻¹ HCl). Briefly, a single colony was picked and cultured in 10 mL of LB medium at 37 °C. After incubating overnight in a shaking bath (shaking rate = 120 rpm), the bacterial suspension was diluted and had the final optical density of 0.5 (OD₆₀₀nm = 0.5), which was measured on a VICTOR X5 microplate reader (PerkinElmer, Waltham).

To find out the relationship between the antibacterial performance and the concentrations of H₂O₂ or PDA/Cu nanometer rods, the prepared bacteria suspension (50 μL) was incubated with a range of different concentrations of H₂O₂ and PDA/Cu nanometer rods for 1 h (final volume 1 mL) and studied by the plate-counting method. Furthermore, the samples after incubation were diluted tenfold and inoculated on a LB agar medium (50 μL per plate), incubated at 37 °C for 20 h, and then photographed.

The morphology of bacteria exposed to different treatments was observed by SEM. *S. aureus* and *E. coli* in the groups of the PBS buffer (pH 5.0), PDA/Cu nanometer rods, H₂O₂, and PDA/Cu nanometer rods + H₂O₂ were washed with PBS (pH 7.4), centrifuged (6000 rpm, 5 min), and resuspended in 2.5% glutaraldehyde (12 h, 4 °C). The samples were then dehydrated with a series of ethanol solutions (50–100%) and dried in air.

To identify the role of PDA/Cu nanometer rods and the released Cu²⁺ in the antibacterial process, the release of Cu²⁺ was quantified. PDA/Cu nanometer rods were suspended in PBS with a concentration of 1 μg mL⁻¹, incubated at 37 °C, and the filtrate at different times (5 min, 1 h, 3 h, and 18 h) was collected. The samples of the PBS buffer (pH 5.0), CuSO₄ solutions (0.6 and 2 mM), PDA/Cu nanometer rods (1 mg mL⁻¹), and the filtrates were added into the probe solution (0.05 wt % dicyclohexanoneoxalyl dihydrazone). All of the samples were incubated at 25 °C for 10 min and measured on a VICTOR X5 microplate reader. PDA/Cu nanometer rods incubated in the PBS buffer for 1 h were centrifuged to remove the supernatant, redispersed in water, and subjected to antibacterial experiments.

**Photothermal Performance of PDA/Cu Nanometer Rods.** PDA/Cu nanometer rods aqueous solutions (2 mL) at different concentrations (20–400 μg mL⁻¹) were exposed to NIR laser for 20 min (808 nm, 0.5 W cm⁻²). The water and Cu₄(OH)₃SO₄ crystals (200 μg mL⁻¹) were also performed under the same conditions. In addition, the temperature changes of PDA/Cu nanometer rods (200 μg mL⁻¹) under different power densities (0.5, 1.0, and 2.0 W cm⁻²) were recorded. The heating and cooling curve data of Cu₄(OH)₃SO₄ crystals (200 μg mL⁻¹) and PDA/Cu nanometer rods (200 μg mL⁻¹) were then used to calculate the photothermal conversion efficiency (η), as described by Shu et al.²⁵

**Photothermal and Catalytic Synergistic Antibacterial Therapy.** A suspension of *S. aureus* and *E. coli* (50 μL, OPD₀₀₀nm = 0.5) was incubated with H₂O₂, DA/Cu nanometer rods, and H₂O₂ + PDA/Cu nanometer rods. The bacteria survival of different groups was immediately studied by the plate-counting method after being irradiated for 20 min using a NIR laser (808 nm, 40 mW cm⁻²). The samples without laser were incubated at room temperature (25 °C) for 20 min as control groups.

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsomega.2c00986.

Characterization results, peroxidase-like activity measurements, bacteria in the PBS buffer of pH 5.0, photothermal performance, composition of the applied buffers, and the mechanism of OPD and TMB color reaction (PDF)

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