Enhanced bioactivity of a GO–Fe$_3$O$_4$ nanocomposite against pathogenic bacterial strains

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Abstract: GO with –OH, –CHO, –CO, –COOH, and epoxide groups is considered more suitable than other commonly used materials for biomedical applications. The presence of –CO, –CHO, and –OH groups renders the easy functionalization of GO by biomolecules and drugs. Therefore, in this study, we designed a multifunctional GO–Fe$_3$O$_4$ nanocomposite and investigated its bactericidal activity against different bacterial strains with an evaluation of percentage inhibition. Our results revealed the potential of GO–Fe$_3$O$_4$ nanocomposites as an effective bactericidal, which can be used for dynamic applications in medical devices and food and other industries.

Keywords: graphene oxide, GO nanocomposites, GO–Fe$_3$O$_4$, pathogenic strains, antibacterial

Introduction

Nanotechnology enables engineering of the materials and tailoring of their physicochemical properties at a nanoscale level. The application of nanotechnology in biomedical fields is one of the major thrust areas that have attracted a great deal of interest. Being abundant in nature, carbon and its allotrope, graphene are considered more environmentally friendly. Graphene, a newly discovered allotrope of carbon is a single, tightly packed layer of pure carbon atoms bonded together in a hexagonal honeycomb lattice. GO, an oxidized form of graphene with –OH, –CHO, –CO, –COOH, and epoxide groups, is considered as more suitable for biomedical applications. The presence of –COOH, –CHO, and –OH groups renders the easy functionalization of GO by biomolecules and drugs. In recent years, multidrug resistance, shown by various microorganisms against existing antibiotics, necessitates the search for new antimicrobial agents, or modifications in existing agents, to improve their antimicrobial activity. Different material systems, such as Ag, ZnO, TiO$_2$, CuO, and Fe$_3$O$_4$, have been extensively used as antimicrobial agents.$^1$ Beside these, various investigations have been conducted on the antibacterial activity of graphene-based materials, namely graphene-based hybrids against Escherichia coli, Staphylococcus aureus, Salmonella typhimurium, Bacillus subtilis, and Enterococcus faecalis.$^2$$^4$ Tang et al.$^5$ reported that GO–Ag nanocomposite functions as a bactericide against the Gram-negative E. coli through disrupting the bacterial cell wall integrity, whereas it exhibits bacteriostatic effects on the Gram-positive S. aureus by dramatically inhibiting cell division.

In this study, we designed a multifunctional GO–Fe$_3$O$_4$ nanocomposite and investigated its bactericidal activity against different bacterial strains with an evaluation of percentage inhibition.
Materials and methods

In this study, GO was synthesized by the modified Hummers method.1 GO–Fe₃O₄ nanocomposites were synthesized by dispersing GO in deionized water (1:1 w/v) and sonicating for ~30 minutes in a conical flask to obtain a homogeneous suspension. FeCl₃ (65 mg) and FeCl₂·4H₂O (40 mg) (Fe²⁺/Fe³⁺=1:2) were dissolved in 50 mL of deionized water and were purged with N₂ for 30 minutes while undergoing constant stirring. Fifty milliliters of ammonia solution was then introduced dropwise with vigorous stirring. The pH was controlled between 11 and 12 throughout the reaction. The mixture was refluxed for ~3 hours at 75°C to form a stable suspension of GO–Fe₃O₄ nanocomposites. These nanocomposites were centrifuged at 10,000 rpm for 20 minutes with water to remove anionic and cationic impurities.

The GO and its nanocomposite were characterized by ultraviolet–visible (UV–Vis)–near-infrared spectroscopy, Fourier transform infrared spectra spectroscopy, and Tecnai G² transmission electron microscope.

Bacterial toxicities of GO and GO–Fe₃O₄ at different (0.01%-0.04%) concentrations were tested against four different pathogenic bacteria including two Gram-negative strains, Klebsiella pneumoniae (ATCC 13883) and Proteus hauseri (ATCC 13315), and two Gram-positive strains, S. aureus (ATCC 25923) and Streptococcus pyogenes (ATCC 19615). A turbidity assay was used to measure the bacterial growth at 660 nm using an UV–Vis spectrophotometer, whereas a cell-viability test was used to further measure the bacterial growth by the colony-forming unit method after treatment with GO and GO–Fe₃O₄. Colonies were counted and compared with control plates (graphene-based materials) to calculate percentage inhibition.

Results and discussion

The synthesis of GO involves the oxidation of graphite by strong oxidizing agents leading to the formation of –COOH, C=O, and epoxide functionalities in the edges and surfaces below and above the plane of the GO nanosheets. Furthermore, high energy sonication provided the effective exfoliation of graphite into few layer GO nanosheets. Fourier transform infrared spectra bands observed at 3,400, 1,720, 1,620, 1,380, and 1,220 cm⁻¹ were attributed to –OH, –CO, aromatic C=C, –COOH, and epoxy groups present on the GO nanosheets. Further, the decrease in the intensities of the infrared band and the absence of band at 1,720 cm⁻¹ for the GO–Fe₃O₄ nanocomposite are attributed to the chemical deposition of iron ions onto the GO nanosheets, and the presence of IR band at 560 cm⁻¹ related to the Fe–O bond confirms the attachment of Fe₃O₄ onto the surface of GO (Figure 1A).

Figure 1B shows the optical absorption spectra (UV–Vis) of the GO nanosheets exhibiting an absorption peak at 230 nm, corresponding to the π→* transition of aromatic C–C bonds, and a shoulder at ~300 nm, which is attributed to the n→* transitions of C=O bonds. The UV–Vis absorption of the GO–Fe₃O₄ nanocomposite has no significant visible peak, indicating possible surface interactions.

Figure 2 shows the structural and size characterization of GO–Fe₃O₄ using transmission electron microscopy (TEM). TEM micrograph revealed that Fe₃O₄ nanoparticles of ~3–10 nm in size are uniformly distributed onto GO nanosheets.

The bacterial toxicity experiment revealed a dramatic decrease in the number of bacteria in response to an increase in the concentration of GO–Fe₃O₄ nanocomposites. Significantly, we found that GO–Fe₃O₄ nanocomposites almost completely suppressed the growth of P. hauseri, leading to a viability loss of up to 98%, whereas the metabolic activity of K. pneumoniae and S. aureus cells was reduced by up to 87% and 83%, respectively. In a similar manner, the cell viability of S. pyogenes was reduced by up to 90% and 83% with a GO–Fe₃O₄ nanocomposite of 0.03% and 0.04%, respectively (Figures 3A and B).
Compared to GO–Fe$_3$O$_4$ nanocomposites, the GO nanosheets displayed weaker antimicrobial activity toward Gram-positive and Gram-negative bacteria. This can be explained on the basis of the negative charge found on the surface of the bacterial cells. The bacterial cells and the GO nanosheets are negatively charged and would repel each other under experimental conditions. However, in the case of Gram-negative bacteria, lipopolysaccharide subunits in the outer membrane contain sugars, phosphates, and lipids, whereas in Gram-positive bacteria, there is the presence of phosphate groups in teichoic acids of the lipid bilayer. These groups facilitate the Hydrogen bonding with the oxygenate group, such as –OH, –COOH, –CO, and epoxide of GO, and assist in wrapping GO nanosheets around the bacterial cells, blocking the cells from taking up nutrition from the growth medium, and eventually leading to cell death.\(^3\)

In comparison with GO, a significant decrease in the negative charge was observed on the GO–Fe$_3$O$_4$ nanocomposites, which results in a better contact between bacterial cells and the GO–Fe$_3$O$_4$ enhancing its antibacterial activity by exposing the bacterial cell to the Fe$_3$O$_4$ nanoparticles. The close interaction inflicts oxidative stress inside the bacterial cell and consequent damage to proteins, membranes, and DNA, leading to cell death.

**Conclusion**

Overall, our results demonstrate the potential of GO–Fe$_3$O$_4$ nanocomposites as an effective bactericidal which can be used for dynamic applications in the field of medical devices and food and other industries.

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**Disclosure**

The authors report no conflicts of interest in this work.

**References**
