Toxic effects of graphene and related materials on bacteria

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Abstract. Graphene and related materials (GRMs) have been reported to have extensive applications in many areas. The widespread use of GRMs makes them inevitably enter the environment through various links causing adverse effects on organisms. Bacteria were representatively used to review the toxicological effects of GRMs on biological organisms in this paper. We comprehensively summarize the recent researches about negative effects of GRMs on bacteria. This is conductive to the evaluation of the ecological risk assessment of GRMs.

1 Introduction

Graphene is a two-dimensional material composed of carbon atoms arranged in a net of fused hexagons[1], including single-layer graphene, few-layer graphene and multi-layer graphene[2]. Graphene, graphene oxide (GO), reduced graphene oxide (rGO) and graphene-based materials formed by combining with various functional groups, organic compounds, metal ions, metal particles, etc. are collectively referred to as graphene and related materials(GRMs)[3-5]. GRMs have a wide range of potential applications in the fields of medicine, electronics, chemicals, water treatment, etc. because of its exceptional optical, mechanical and electrical properties[6-8]. The widespread use of GRMs makes them inevitably enter the water environment through various links resulting in exposure of organisms to the GRMs.

There have been many studies reported the negative effects of GRMs on different types of living organisms and a great possibility of bioaccumulation[9, 10]. Studies using the body burden factor (BBF) to evaluate the accumulation of 14C-labeled graphene in E. coli, T. thermophila, D. magna, and D. rerio have showed that organisms exposed to graphene-containing culture medium have high potential of graphene accumulating[11]. Researches on the ecological safety of GRMs have increased rapidly and received extensive attention. However, these studies are chaotic and lacking in sorting and induction. Therefore, we summarized the characteristics and general patterns of toxic effects of different kinds of GRMs on bacteria, which were selected as representatives. The review of this article is worthy of reference for the future exploration of the environmental safety of GRMs.

2 Toxicity characteristics of GRMs on bacteria

2.1 Common toxic effects of GRMs on bacteria

Table 1 summarized the effects of GRMs on bacteria. On the whole, the antibacterial properties of GRMs has been recognized. They can reduce the cell viability, destroy cell wall integrity, inhibit the formation of bacterial biofilm, alter energy metabolism and cause oxidative damage on bacteria[12-15]. What’s more, there are studies reported that its antibacterial effect is comparable to certain antibiotics[16]. In any case, more researches are needed about the antibacterial properties of different types GRMs to take advantage of this excellent feature. A disk diffusion test conducted by Kadiyala et al. showed that the diameter of inhibition zone (DIZ) of GO on gram-negative (E. coli) and gram-positive bacterial strain (S. aureus) are 19 mm and 21 mm, respectively, which is bigger than gentamicin (9.3 mm and 12.17 mm respectively)[17]. But it is unsure whether GRMs exert greater effects on all the gram-negative bacteria than gram-positive ones. GRMs could be the promising materials for inhibiting resistant bacteria, because one of its antibacterial mechanisms is based on the physical damage. GO with concentrations higher than 10 mg/L could damage the resistant plasmids of E. coli HB101 consequently reduce its resistance to antibiotics[18]. Once GRMs were successfully used to inactivate various resistant bacteria, there will be a leap forward in the treatment of resistant bacteria. Among the influencing factors, the shape, size and surface coating etc. are factors that are relatively easy to regulate.

Table 1 summarized the effect of GRMs on bacteria.

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2.2 Effect of surface modification on the toxicities of GRMs

Surface modification of GRMs can both increase and decrease the toxicities of GRMs. Understanding the changes of toxic effects of surface modification is useful.

Table 1. Toxic effects of different GRMs on different bacteria.

| GRMs            | Species               | Exposure condition                                                                 | Effects                                                                                           | Reference |
|-----------------|-----------------------|-----------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------|-----------|
| Ag-GO           | Natural aquatic microorganisms | Exposed to Ag-GO concentration of 10, 50, or 100 mg/L at mimic natural aquatic environment | Reduced enzymes activities; Decreased nitrification rate; ROS was generated continuously          | [19]      |
| Cu2O-Graphene   | V. cholerae           | The bacteria culture medium was incubated with 100 or 300 μg contained Cu2O-Graphene disc loaded at 4°C for 24 h | 300 μg-DIZ: 20 mm; 200 μg-DIZ: 14 mm; 300 μg-DIZ: 20 mm                                        | [20]      |
| Cu2O-Graphene   | S. pyogenes           | Incubated with 100 or 300 μg contained Cu2O-Graphene disc loaded at 4°C for 24 h | 100 μg-DIZ: 0 mm; 200 μg-DIZ: 14 mm; 300 μg-DIZ: 22 mm                                           | [20]      |
| AuNPs-rGO-NC    | E. coli               | Incubated with 100 ± 15 μg AuNPs-rGO-NC contained disk placed on at 32°C for 24 h and incubated with 10 to 100 μg/L AuNPs-rGO-NC in an orbital shaker maintained at 180 rpm at 30°C | DIZ: 33 mm; AI: 3.548; MIC: 30 μg/mL; MBC: 40 μg/mL                                             | [17]      |
| AuNPs-rGO-NC    | B. subtilis           | Incubated with 100 ± 15 μg AuNPs-rGO-NC contained disk placed on at 32°C for 24 h and incubated with 10 to 100 μg/L AuNPs-rGO-NC in an orbital shaker maintained at 180 rpm at 30°C | DIZ: 30 mm; AI: 2.323; MIC: 40 μg/mL; MBC: 50 μg/mL                                             | [17]      |
| AuNPs-rGO-NC    | S. aureus             | Incubated with 100 ± 15 μg AuNPs-rGO-NC contained disk placed on at 32°C for 24 h and incubated with 10 to 100 μg/L AuNPs-rGO-NC in an orbital shaker maintained at 180 rpm at 30°C | DIZ: 29 mm; AI: 3.382; MIC: 50 μg/mL; MBC: 60 μg/mL                                             | [17]      |
| GO              | B. subtilis           | Incubated with 100 ± 15 μg GO contained disk placed on at 32°C for 24 h and incubated with 10 to 100 μg/L GO in an orbital shaker maintained at 180 rpm at 30°C | DIZ: 21 mm; AI: 1.626; MIC: 60 μg/mL; MBC: 80 μg/mL                                             | [17]      |
| rGO             | E. coli               | 50 μL rGO solution were placed into a well with diameter of 1 cm                   | DIZ: 18 ± 0.5 mm; Cell wall integrity reduced with increasing time                               | [15]      |
| ZnO-rGO         | E. coli               | 50 μL ZnO-rGO solution were placed into a well with diameter of 1 cm               | DIZ: 28 ± 0.7 mm; A sudden decrease of integrity after 20 min                                    | [15]      |
| GO              | E. coli               | Dispersed GO was inoculated with bacteria of 70 μl 108 CFU/ml, incubated at 37°C; Bacteria of 108 CFU/ml were cultured on nutrient agar plates supplemented with GO of 250 ppm | MIC: 125 ppm; Removal efficiency: about 70%                                                     | [21]      |
| GO              | S. aureus             | Dispersed GO NPs was inoculated with 70 μl 108 CFU/ml of the bacteria, incubated at 37°C | MIC: 225 ppm                                                                                     | [21]      |
| G-Fe3O4         | E. coli               | Dispersed G-Fe3O4 NPs was inoculated with 70 μl 108 CFU/ml of the bacteria, incubated at 37°C; Bacteria of 108 CFU/ml were cultured on nutrient agar plates supplemented with G-Fe3O4 of 250 ppm | MIC: 100 ppm; Removal efficiency: about 92%                                                      | [21]      |
| G-Fe3O4         | S. aureus             | Dispersed G-Fe3O4 NPs was inoculated with 70 μl 108 CFU/ml of the bacteria, incubated at 37°C | MIC: 200 ppm                                                                                     | [21]      |
| rGO-ZnO         | K. pneumonia          | RGO-ZnO nanocomposite was loaded (20, 30, 50 μL) in 4 mm diameter of agar wells, incubated at 38°C for 24 h | DIZ: 14 ± 0.44 mm                                                                                | [22]      |
| rGO-ZnO         | B. thuringiensis      | rGO-ZnO nanocomposite was loaded (20, 30, 50 μL) in 4 mm diameter of agar wells, incubated at 38°C for 24 h | DIZ: 14 ± 0.73 mm                                                                                | [22]      |
for choosing appropriate types of modifier to achieve the specific purpose of toxicity regulation. The factors determining the modification effect include the kinds of surface coating and the ratio of GRMs to modifiers. Common types of modifying include metal or metal oxide nanoparticles, metal ion, inorganic salts and other organics. Generally speaking, the compound is more toxic when it is combined with a substance that itself is antibacterial[25, 26]. For instance, GRMs decorated with Au-NPs, Cu-NPs, Cu2O, curcumin or polyindole have stronger inhibitory effects than the bare GRMs[17, 20-29]. This may be the synergy effect of GO and modifiers. But there are also exceptions. Unlike this enhanced toxicity, Gao et al. reported the decreased joint toxicity while investigating the combined toxicity of GO and heavy metal cation (Me(II)) (Cd2+, Co2+ and Zn2+) [30]. The authors gave some reasons for this phenomenon like the adsorption of GO reduced the content of free Me(II), Me(II) weakened the sharp edges of GO and bacteria are difficult to assimilate the complexes. In other word, it is because the combination of graphene and Me(II) reduces the probability of Me(II) entering cells. At the same time, the deeper reason may be that Me(II) need to enter the bacteria firstly and then bind to the deeper reason may be that Me(II) need to enter the bacteria firstly and then bind to the deeper reason may be that Me(II) need to enter the bacteria firstly and then bind to the deeper reason may be that Me(II) need to enter the bacteria firstly and then bind to the bacteria contained agar at 37°C for 12 h; Put disks on the bacteria contained agar at 37°C for 12 h

| rGO-ZnO | *B. cereus* | rGO-ZnO nanocomposite was loaded (20, 30, 50 µL) in 4 mm diameter of agar wells, incubated at 38°C for 24 h | DIZ: 15 ± 0.55 mm [22] |
|---------|-------------|--------------------------------------------------------------------------------------------------|------------------------|
| rGO-ZnO | *P. aeruginosa* | rGO-ZnO nanocomposite was loaded (20, 30, 50 µL) in 4 mm diameter of agar wells, incubated at 38°C for 24 h | DIZ: 12 ± 0.65 mm [22] |
| GO-Ag   | *E. coli*   | Bacterial suspensions with different concentrations of GO-Ag (from 1.0 to 100.0 µg/mL) added were incubated at 37°C for 18 h | MIC: 0.01 mg/mL [23] |
| GO-Ag   | *S. aureus* | Bacterial suspensions with different concentrations of GO-Ag (from 1.0 to 100.0 µg/mL) added were incubated at 37°C for 18 h | MIC: 0.17 mg/mL [23] |
| GO-Ag   | *E. coli*   | Bacteria were cultured in connect with foils coated with GO-Ag at 37°C | Antibacterial activity: 88.6%; Membrane Integrity: 66.3%; Enhanced ROS [14] |
| GO-Ag   | *S. aureus* | Bacteria were cultured in connect with foils coated with GO-Ag at 37°C | Antibacterial activity: 79.6%; Membrane Integrity: 59.4%; Enhanced ROS [14] |
| Spindle-shaped GO | *E. coli* | Cell solutions added spindle-shaped GO inoculated at 37°C for 12 h; Put disks on the bacteria contained agar at 37°C for 12 h | MIC: 125 µg/mL; R: 15 ± 0.2% [24] |
| Spindle-shaped GO | *S. typhimurium* | Cell solutions added spindle-shaped GO inoculated at 37°C for 12 h; Put disks on the bacteria contained agar at 37°C for 12 h | MIC: 125 µg/mL; R: 15 ± 0.2% [24] |

Caption: Diameter of inhibition zone (DIZ); Activity index (AI); Minimum inhibition concentration (MIC); Minimum bactericidal concentration (MBC); The normalized viability ratio (R)

Graphene oxide (GO); Graphene (G); AuNPs-rGO nanocomposite (AuNPs-rGO-NC); Reduced graphene oxide (rGO); Zinc oxide decorated rGO (ZnO-rGO)

2.3 Effect of inherent properties of GRMs on its toxicities.

Some inherent properties of GRMs such as size and morphology will affect its antibacterial characteristics. Generally speaking, materials with smaller size exert greater effects on organisms. Smaller-sized GO sheet showed stronger antibacterial property due to larger defect density, while measuring the antibacterial activity on GO-coated surface[31]. This could also be because the smaller-sized has a larger specific surface area with a stronger adsorption effect, and is easier to enter the organism accordingly. In term of morphology, wrinkled GO nanosheets with different roughness grade have different antibacterial capabilities, which is related to the relative dimension between the surface roughness grade and the bacterial size[32]. But, there is a lack of comparison of antibacterial properties between GRMs with the same size but wrinkled and smooth surfaces, respectively.

3 Conclusion

In general, there are many factors that contribute to the toxicities of GRMs on bacteria. Toxicities of GRMs with different purity, morphology, thickness, lateral dimension, surface charge and coating may vary greatly.
So, we can select appropriate types of GRMs according to our needs in practical. GRMs with higher toxicity can be chosen in the process of sterilization and elimination of harmful algae, but for materials that are frequently or directly in contact with human beings, the GRMs with lower toxicity should be selected through optimization. The researches on the toxic effects of GRMs make great contributions to the utilization and environmental risk assessment of GRMs.

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