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Catalytic performance and antibacterial behaviour with molecular docking analysis of silver and polyacrylic acid doped graphene quantum dots†

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In this research, a fixed concentration (3 wt%) of Ag/PAA and PAA/Ag doped graphene quantum dots (GQDs) were synthesized using the co-precipitation technique. A variety of characterization techniques were employed to synthesize samples to investigate their optical, morphological, structural, and compositional analyses, antimicrobial efficacy, and dye degradation potential with molecular docking analysis. GQDs have high solubility, narrow band gaps, and are suitable for electron acceptors and donors but show less adsorption and catalytic behavior. Incorporating polyacrylic acid (PAA) into GQDs increases the catalytic and antibacterial activities due to the carboxylic group (–COOH). Furthermore, introducing silver (Ag) increased the degradation of dye and microbes as it had a high surface-to-volume ratio. In addition, molecular docking studies were used to decipher the mechanism underlying the bactericidal action of silver and polyacrylic acid-doped graphene quantum dots and revealed inhibition of β -lactamase and DNA gyrase.

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1. Introduction

Urban sprawl and rapid industry growth have exacerbated freshwater scarcity.¹ Industries like paper, textiles, food, leather, and plastic release harmful chemicals, synthetic dyes, heavy metals, and other organic pollutants into the water reservoir, leading to water contamination.² 7×10^5 tons of dyes originate from textile industries per year, such as safranin dye, methylene blue (MB), methyl orange, methyl red, and rhodamine B dye (RhB). RhB ($C_{28}H_{31}N_2O_3Cl$) belongs to the family of xanthene dyes, released directly or indirectly into water resources,

threatening human beings with various diseases like cancer, skin irritation, renal failure, respiratory disorders, eye burns, and hepatic dysfunction.^{3–5} Moreover, effluent water contains infective agents, including algae, viruses, bacteria, and other microorganisms.⁶ The most common multidrug resistance (MDR) bacterium, *Escherichia coli* (*E. coli*), is responsible for nosocomial infection⁷ and causes 2.5 million deaths of children every year from diarrhea.⁸ Degrading organic pollutants and removing metallic ions or pathogens from wastewater is imperative for water purification. Several techniques have been manifested for treating contaminated water, including ion exchange, membrane filtration, adsorption/precipitation, photo-catalysis, catalysis, electrochemical, and enzymatic decomposition.^{9–11} Among these, catalysis in the presence of a nano-material has gained much attention attributed to its environmentally friendly nature, cost-effectiveness, and excellent efficiency in water purification.¹² Nano-materials have proven to be a practical field for wastewater remediation due to their different physical and chemical properties, like shape, size, and surface area to volume ratio, which play a productive role in the purification of contaminated water.^{13–15} Nano-materials (ZnO, TiO₂, CeO₂, CdS, and La₂O₃) can remove heavy metal ions, toxic dyes, and other infectious bacteria.^{16,17} Among these, semiconductor nano-materials are successfully utilized for catalytic dye degradation even with a sufficient band gap.¹⁸ A two-dimensional (2D) gapless semiconductor graphene discovered in 2004 became of particular interest to researchers because of having excellent electrical properties, high surface

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area ($2630 \text{ m}^2 \text{ g}^{-1}$), less toxicity, and high thermal conductivity ($5000 \text{ W m}^{-1} \text{ K}^{-1}$).^{19,20} Graphene has been intensively used as a catalyst for degrading pollutants and has wide applications in sensors, solar cells, and batteries.^{21,22} Among graphene nanostructures, thin layers GQDs (100 nm) have gained a lot of attention^{23,24} for characteristics like photoluminescence, high stability, narrow bandgap energy, good electron acceptors, and donors.^{25,26} But its applications are limited due to low solubility, poor luminescence,²⁷ and inferior catalytic. To enhance the catalytic activity, doping with metals, nonmetals, and polymers has been suggested. Incorporating an organic polymer such as PAA increases the catalytic and antibacterial activities due to the presence of a carboxylic group ($-\text{COOH}$). It is a protective capping agent, capturing heavy metals from polluted water.²⁸ The doping of inorganic metals such as Ag increased the degradation of dye and microbes because of the high surface-to-volume ratio.²⁹ The combination of Ag and PAA dopants provides a large surface area for the adsorption of pollutants.³⁰ Two approaches (top-down and bottom-up) are applied to the synthesis GQDs in top-down method focuses on breaking the precursors such as carbon fibres, graphene sheets, and other carbonaceous compounds. In contrast, the bottom-up approach involves assembling basic units into nano-material.³¹ In the present study, a simple bottom-up method involving pyrolysis of glucose was used to prepare GQDs from the organic precursor to investigate the catalytic efficacy against RhB degradation and bactericidal potency for *E. coli*. This research contributes to discovering multifunctional, defensible, eco-friendly, and economic catalysts for maintaining water standards by reducing toxic products and blocking bacterial cell proliferation.

2. Experimental

2.1 Materials and reagents

Glucose ($\text{C}_6\text{H}_{12}\text{O}_6$, 99.5%), ammonia solution (33%), polyacrylic acid ($\text{C}_3\text{H}_4\text{O}_2$)_n, AgNO_3 , 99.8%, and HCl, 37% were acquired from Sigma Aldrich.

2.1.1 Synthesis of GQDs, PAA, and Ag-doped GQDs. To synthesize GQDs, pyrolysis of glucose was performed. Initially, 4 g of glucose was heated at 260°C to liquefy, and after 20 min, the yellow color shifted to orange. Subsequently, NH_3 solution (12.5%) was incorporated dropwise to liquefy glucose under continuous stirring at 70°C for 3 h to remove the ammonia odor. To obtain neutral pH, HCl was added, and the mixture was stirred magnetically at 150°C for 12 h and crushed to get a fine powder, as shown in Fig. 1.

2.1.2 Synthesis of Ag/PAA-doped GQDs. For doping, fixed amounts of (3 wt%) Ag and PAA were added into the prepared GQDs solution at pH around 12 under vigorous stirring (Fig. 1). The mixture was heated for two hours at 100°C , centrifuged twice at 7000 rpm for 7 min, then washed with DI water to obtain precipitates. Ultimately, the residues were dried overnight at 100°C to obtain a refined powder of Ag/PAA-GQDs.

2.1.3 Synthesis of PAA/Ag-doped GQDs. Fixed amounts of (3 wt%) PAA and Ag were introduced into the synthesized GQDs solution at pH ~ 12 under continuous stirring for the required nanocomposite (Fig. 1). To obtain precipitates; the mixture was

heated for 2 h at 100°C , centrifuged two times at 7000 rpm for 7 min and then cleaned with DI water. Finally, residues were dried overnight at 100°C to obtain a fine powder of PAA/Ag-GQDs.

2.2 Instrumental measurements

2.2.1 Catalytic activity (CA). The CA of pure and doped GQDs was examined in the presence of a reducing agent (NaBH_4) and an oxidizing agent, rhodamine B (RhB), for the degradation of RhB. Freshly prepared 1.5 mL of RhB solution was added to the quartz cuvette containing 400 μL of prepared 0.1 M NaBH_4 . Moreover, 400 μL of the prepared sample was dissolved into the solution. In the presence of NaBH_4 , RhB was reduced to leuco RhB (LRhB), confirming the dye degradation. The following equation calculated degradation efficiency: degradation (%) = $(C_0 - C_t)/C_0 \times 100\%$, where C_0 and C_t are the initial and final concentrations of RhB.

2.2.2 Catalysis mechanism. The mechanism for the catalytic degradation of RhB involves the redox reaction between NaBH_4 and RhB. Initially, reactants adsorbed onto the surface of nano-material, NaBH_4 dissociates into BH_4^- and H^+ ions which RhB accepts, favoring the breakdown of organic dye. However, in the presence of NaBH_4 reaction was slow; to accelerate the degradation rate, synthesized specimens were added as catalysts. The enhanced catalytic rate was related to the significant interaction among the nucleophilic reducing agent and electrophilic dye on the large surface area of quantum dots. The catalyst functions as an electron relay, permitting the transfer of electrons and detached H atoms from BH_4^- to the cationic dye that results in the breakage of the double bond by π conjugation. Finally, pink RhB was reduced to colorless LRhB,^{32,33} as shown in Fig. 2.

2.3 Biological activity

2.3.1 Sample collection. Direct milking into sterile glassware collected raw milk samples from lactating cows marketed at various markets, veterinary facilities, and farms in Punjab, Pakistan. Milk samples were brought to the lab after being acquired at 4°C . Gram-negative (G –ve) *E. coli* bacteria found in raw milk were counted on MacConkey agar (MA). Cultured specimens were incubated for 48 h at 37°C .

2.3.2 Identification and characterization of bacterial isolates. Using Bergey's Manual of Determinative Bacteriology as a reference, different biochemical and morphological strategies as gram staining based on colony morphology were applied to identify *E. coli* bacteria.³⁴

2.3.2.1 Antibiotic susceptibility. On Mueller Hinton agar (MHA), the Bauer *et al.*³⁵ disk diffusion method was employed to conduct the antibiotic susceptibility test. The test was conducted to analyze whether the *E. coli* was resistant to the following antibiotics (classes); ceftriaxone (Cro) $30 \times 10^{-6} \text{ g}$ (cephalosporins), gentamicin (Gm) $10 \mu\text{g}$ (aminoglycosides), ciprofloxacin (Cip) $5 \times 10^{-6} \text{ g}$ (Quinolones), tetracycline (Te) $30 \times 10^{-6} \text{ g}$ (tetracyclines), imipenem (Imi) $10 \times 10^{-6} \text{ g}$ (carbapenem), amoxycillin (A) $30 \times 10^{-6} \text{ g}$ (penicillins), and azithromycin (Azm) $15 \times 10^{-6} \text{ g}$ (macrolides).³⁶ *E. coli* purified



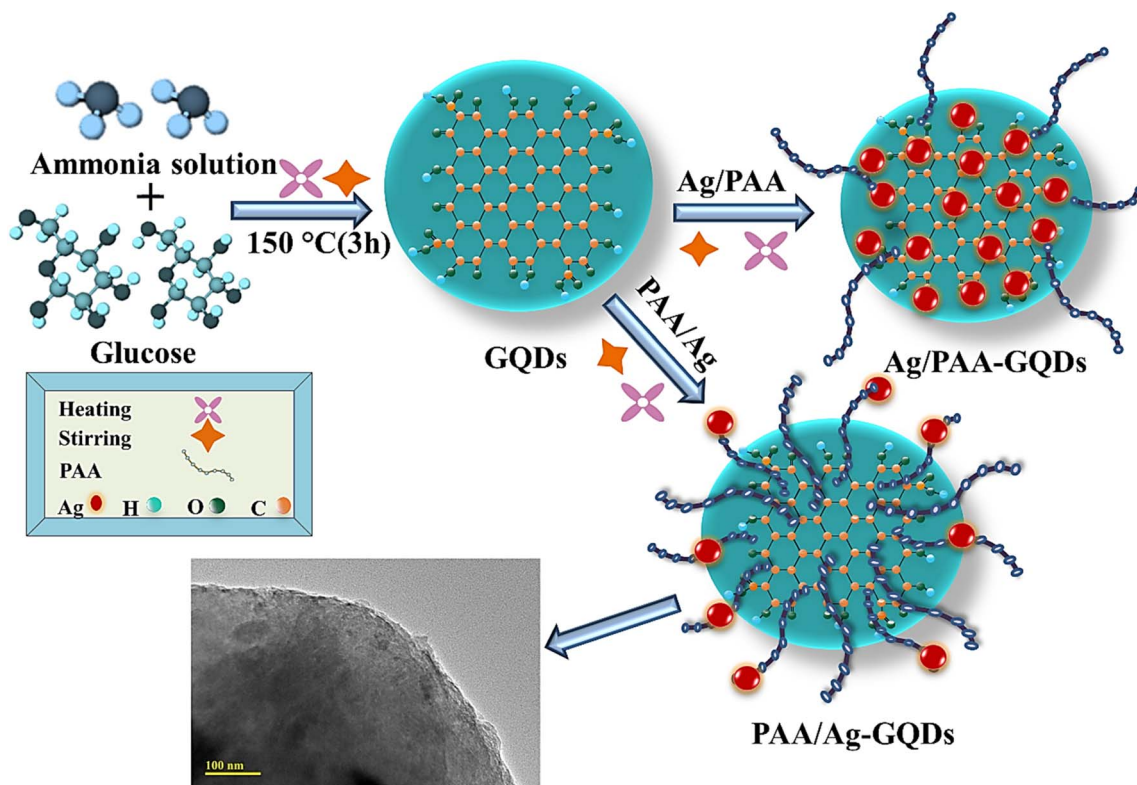


Fig. 1 Schematic synthesis of GQDs, Ag/PAA-GQDs and PAA/Ag-GQDs.

cultures were cultivated, and turbidity was brought to 0.5 MacFarland. After that, it was spread-plated on Muller Hinton Agar (MHA) (Oxoid Limited, Basingstoke, UK), and antibiotic disks were positioned apart from the inoculated infected plate's

surface to preclude the overlap of inhibitory zones. Clinical and Laboratory Standard Institute was utilized to explain the results after the incubation of plates for 24 h at 37 °C.³⁷ Bacterium was proclaimed MDR if shown to resist at least three drugs.³⁸

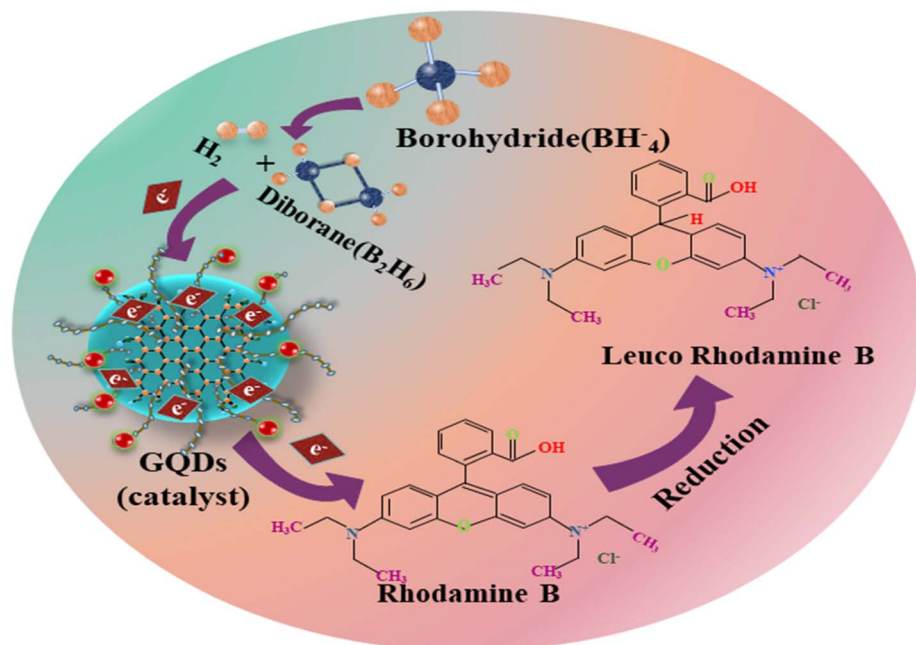


Fig. 2 Schematic diagram of the catalysis mechanism of GQDs, Ag/PAA-GQDs, and PAA/Ag-GQDs.



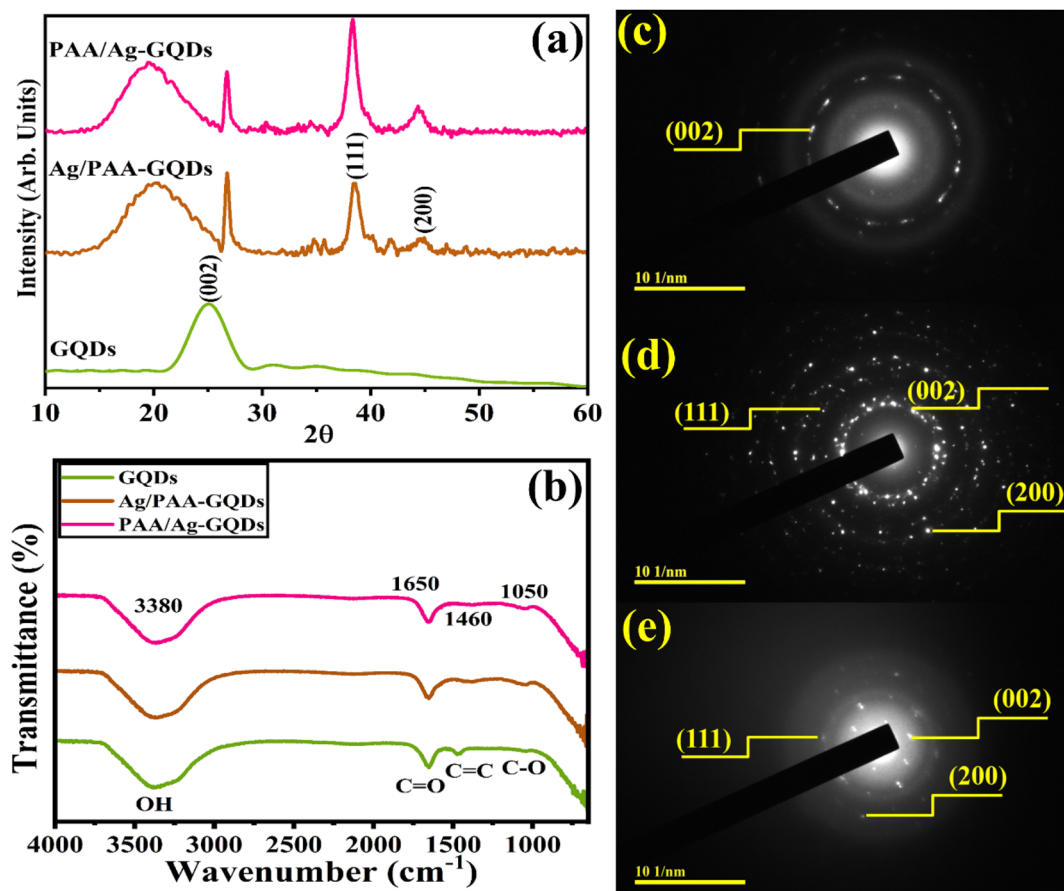


Fig. 3 (a) XRD patterns, (b) FTIR spectra, (c–e) SAED images of Ag/PAA-GQDs and PAA/Ag-GQDs.

2.3.2.2 Antimicrobial activity. Agar well diffusion procedure was employed to assay the *in vitro* antimicrobial potential of pure and doped GQDs upon ten isolates of MDR *E. coli* collected from mastitic milk by swabbing 1.5×10^8 CFU mL⁻¹ (0.5 McFarland standard) MDR *E. coli* on MA. A sterile cork bore was used to bore well on MA plates having a diameter of 6 mm. Each well was filled with distinct concentrations as (0.5 mg/50 μL)

and (1.0 mg/50 μL) of prepared samples by micropipette in contrast to ciprofloxacin (0.005 mg μL⁻¹) and deionized water (50 μL) referred as a positive and negative control, respectively.³⁹

2.3.2.3 Statistical analysis. The inhibition zone in millimeters (mm) and inhibition zone diameters were determined by one-way analysis of variance (ANOVA) utilizing SPSS 20 for estimating the antimicrobial efficacy.⁴⁰

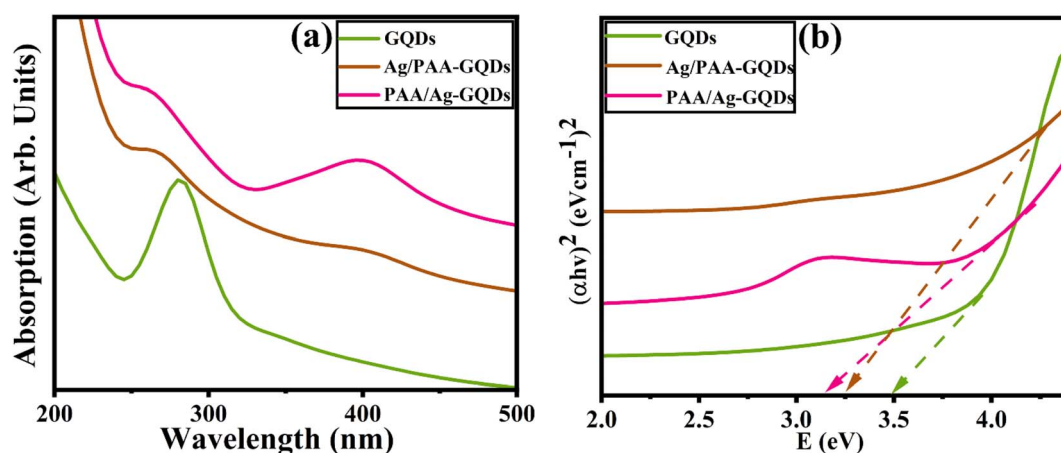


Fig. 4 (a) UV-visible spectra, (b) band gap energies of GQDs, Ag/PAA-GQDs, and PAA/Ag-GQDs.

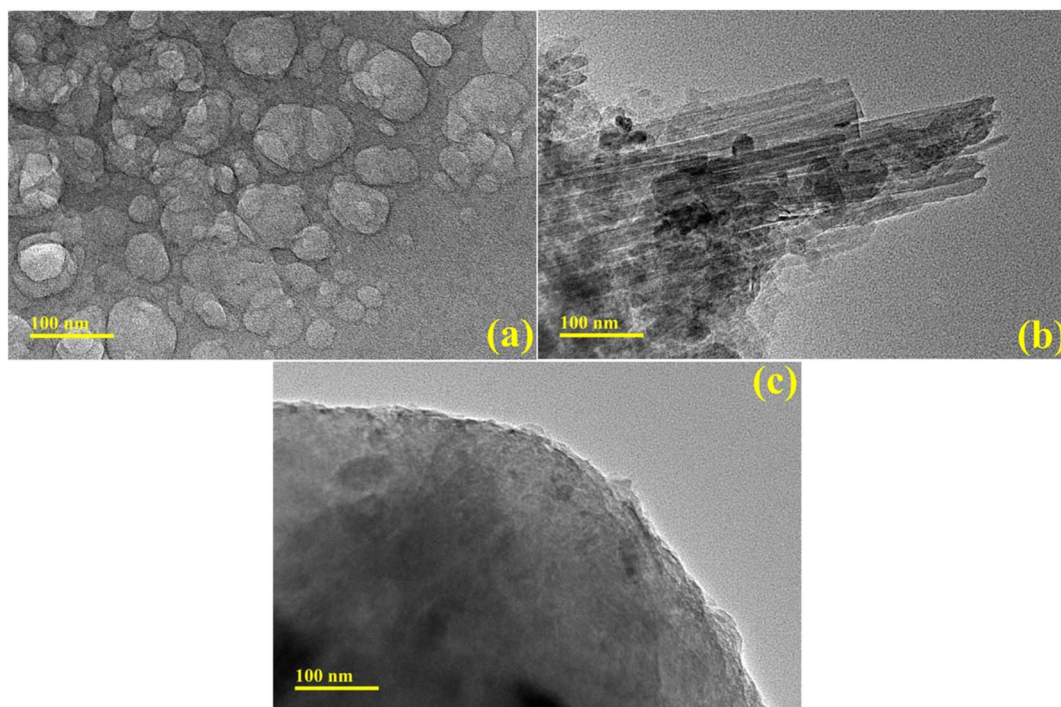


Fig. 5 TEM analysis of (a) GQDs, (b) Ag/PAA-GQDs, and (c) PAA/Ag-GQDs samples.

2.4 Molecular docking analysis

The cell wall production process disruption has been suggested as an attractive target for antibiotic research and as a possible mechanism behind the bactericidal action of

different nanostructures. Enzymes pertaining to peptidoglycan production have enormous significance for the identification of novel antibacterial drugs since their inhibition results in the destruction of the cell wall and, eventually, the death of bacteria.^{41,42} Similarly, enzymes pertaining to nucleic

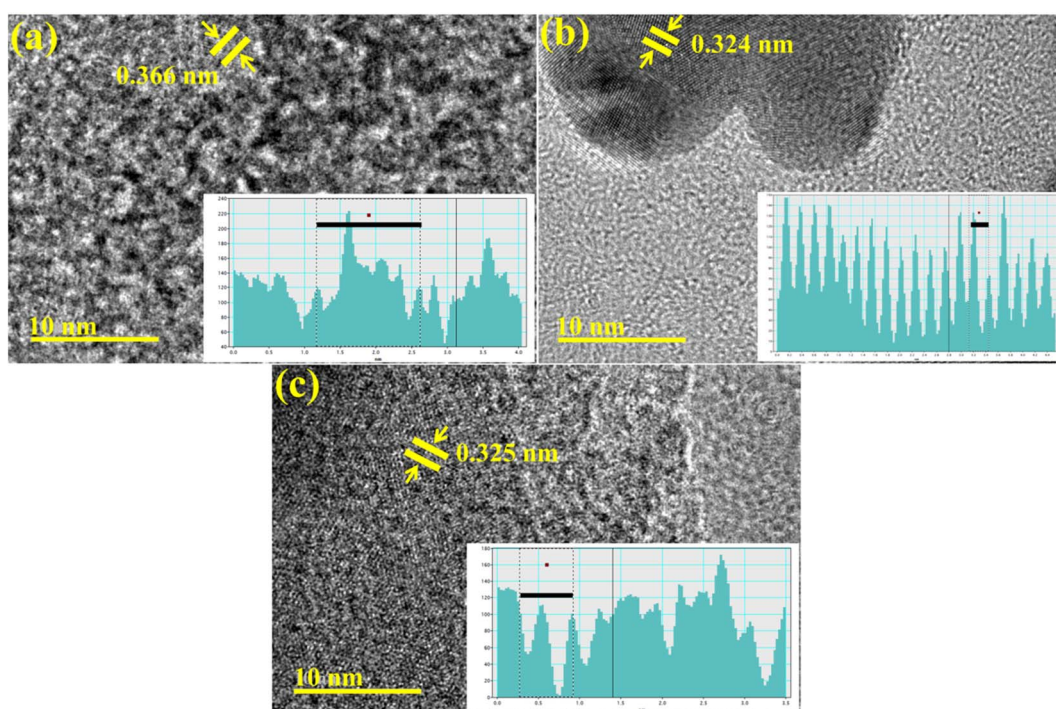


Fig. 6 HR-TEM micrographs of synthesized (a) GQDs, (b) Ag/PAA-GQDs, and (c) PAA/Ag-GQDs.



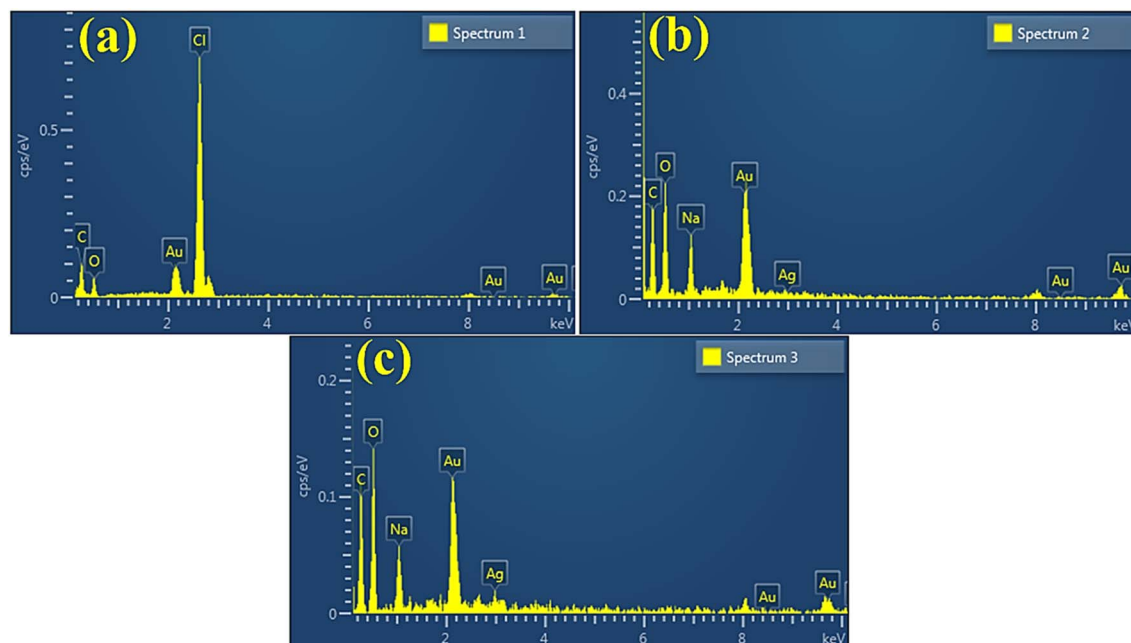


Fig. 7 EDS analysis of (a) GQDs (b) Ag/PAA-GQDs (c) PAA/Ag-GQDs.

acid biosynthesis, particularly DNA gyrase, have a significant impact on the identification of antibiotics.⁴³ Here, we evaluated the inhibitory ability of silver and polyacrylic acid-doped graphene quantum dots against β -lactamase and DNA gyrase enzymes from *E. coli*. Crystal structures of β -lactamase and DNA gyrase from *E. coli* were acquired from RCSB PDB (<https://www.rcsb.org>) with PDB codes 4KZ9 (resolution: 1.72)⁴⁴ and 5MMN (resolution: 1.90).⁴⁵ The docking investigation was executed using the SYBYL-X 2.0 program, as described in our previous studies.^{46,47} The 3D structures of the chosen compounds were generated using the Sybyl-X2.0/SKETCH module (Fig. S1†). Subsequently, energy reduction was performed using the Tripos force field, using Gasteiger Hückel atomic charges. The Surflex-Dock module, a component of the SYBYL-X 2.0 molecular modelling software program, was used to conduct flexible molecular docking simulations. These simulations aimed to investigate the binding interactions between nanoparticles and the active site residues of certain proteins. Hydrogen atoms were inadvertently omitted. The allocation of atomic types and application of atomic charges were carried out in accordance with the AMBER 7 FF99 force field. Ultimately, by using the Powell method with a convergence gradient of 0.5 kcal (mol⁻¹ Å⁻¹) over a span of 1000 cycles, the energy was successfully reduced in order to mitigate steric conflicts. In order to ensure accuracy and reliability, a minimum of 20 optimal docked conformations were definitively preserved for each ligand–receptor complex system. The Hammerhead scoring system was used for the evaluation of the most favorable putative ligand conformations. The Surflex dock module employs an empirically derived consensus scoring function called cScore to generate and prioritize hypothetical orientations of ligand fragments.

3. Results and discussion

3.1 Characterization of catalyst

The crystalline phase purity and structural information of synthesized samples were assessed through XRD ranging from 10° to 60° (Fig. 3a). XRD peak of GQDs at 25.2° corresponds to the (002) plane.⁴⁸ Upon Ag/PAA doping, two distinct peaks were observed at 38.2° and 44.4° for (111) and (200) planes of cubic crystal structure (JCPDS No. 01-087-0718). Peaks shift towards a higher 2θ value upon incorporating Ag, confirmed interstitial site of Ag.⁴⁹ Furthermore, a broad diffraction peak observed at 18.7° corresponds to PAA and shows its amorphous nature.⁵⁰ FTIR Spectroscopy was utilized to determine the nature of surface functional groups. The transmittance band at 3380 cm⁻¹ was ascribed to the stretching vibration of the hydroxyl (–OH) group.⁵¹ The vibration band at 1460 cm⁻¹ is assigned to the vibration of –C=C bonds of the aromatic system.^{52,53} The band of the –C=O group (stretching mode) was centered at around 1650 cm⁻¹, evidenced by the edges functionality of the –C=O group.^{52,54,55} The weak –C–O stretching peak was also observed at ~1050 cm⁻¹.⁵⁶ Incorporating Ag and PAA into prepared GQDs indicated that one of the bands was shifted with decreased intensity due to metal–polymer interactions⁵⁷ (Fig. 3b). The selected area diffraction (SAED) pattern of GQDs, Ag/PAA-GQDs, and PAA/Ag-GQDs NSs showed bright rings associated with distinct XRD planes (002), (111), and (200) (Fig. 3c–e).

UV-VIS spectroscopy was utilized to analyze the optical properties of synthesized samples. GQDs showed absorption in the range of ~260–320 nm,⁵⁸ and a strong absorption peak was observed at ~280 nm, attributed to π – π^* electronic transition of graphitic C=C domains in sp² cluster^{59,60} as elaborated in (Fig. 4a). Tauc plot was used to calculate the band gap energy



(E_g) of QGDs as 3.5 eV, consistent with previously published data.^{61,62} E_g of QGDs in the presence of capping was decreased from 3.5 to 3.10 and 3.2 eV. Upon doping, E_g values are the consequence of the enhanced quantum confinement effect with a decrease in domain size in QGDs⁶³ (Fig. 4b).

TEM analysis was performed to examine synthesized products' morphology and structural properties. TEM images of the control sample revealed the formation of quantum dots and, upon doping of Ag-PAA nanorods of Ag, occupied the surface of QGDs. Incorporating PAA-Ag in QGDs demonstrated an aggregation (Fig. 5a–c).

Additionally, the interlayer distance of QGDs, Ag/PAA-QGDs, and PAA/Ag-QGDs was calculated from HR-TEM micrographs using Gatan software as (0.366 nm, 0.324 nm, 0.325 nm) (Fig. 6a–c).

EDS spectra revealed carbon and oxygen peaks which confirmed the presence of QGDs. The prominent oxygen peak in doped samples generated from PAA, chemical formula $(C_3H_4O_2)_n$.⁶⁴ The chloride (Cl) and sodium (Na) peaks ascribed to HCl and NaOH were used in the synthesis to sustain the pH. The Ag peak confirmed the existence of Ag in doped samples, while small Au peaks attributed to the coating on the samples minimized charging effects^{52,65} (Fig. 7a–c). Moreover, EDS mapping of the synthesized doped specimen was utilized to analyze its elemental constituent distribution pattern to check additional interfacial contact (Fig. 8a–c). Two components (Cu

and O) were found to spread in doped samples. As mentioned, O, Cu, and Zn were assigned to contamination, the sample holder used for EDS analysis.

3.2 Catalytic properties of QGDs, Ag/PAA-QGDs, and PAA/Ag-QGDs

A UV-Vis spectrophotometer (200–800 nm) was used to assess the catalytic efficiency of Ag/PAA-QGDs and PAA/Ag-QGDs with $NaBH_4$ in acidic, basic, and neutral conditions. The spectroscopic results revealed the RhB degradation with some approximate errors (72.4 ± 0.8 , 66.4 ± 0.9 , $66.1 \pm 1\%$) in neutral medium (pH = 7), ($54. \pm 1.3$, 50.2 ± 2 , $57.4 \pm 2.3\%$) in basic medium (pH = 12) and (62.1 ± 1.1 , 63.5 ± 1.8 , $44.7 \pm 2.2\%$) in acidic medium (pH = 4) illustrated in (Fig. 9a–c). The reduction of RhB occurred slowly (40 min) without a catalyst, which was calculated with some errors in acidic, basic, and neutral media, 25.2 ± 2.28 , 21.8 ± 2.3 , and $17.0 \pm 1.2\%$, respectively (Fig. 9d). At pH = 7, the catalyst surface (QGDs) typically acquires a negative charge, which promotes the adsorption of positively charged RhB, thus accelerating the degradation rate. In an acidic medium, catalytic activity was increased, ascribed to the production of H^+ ions that adhere to the nano-material. In a basic medium, the concentration of hydroxyl ions increases, resulting in product oxidation and a significant reduction in catalytic performance. Upon doping of Ag/PAA, the degradation potential decreased as the Ag occupied active sites on a nano-

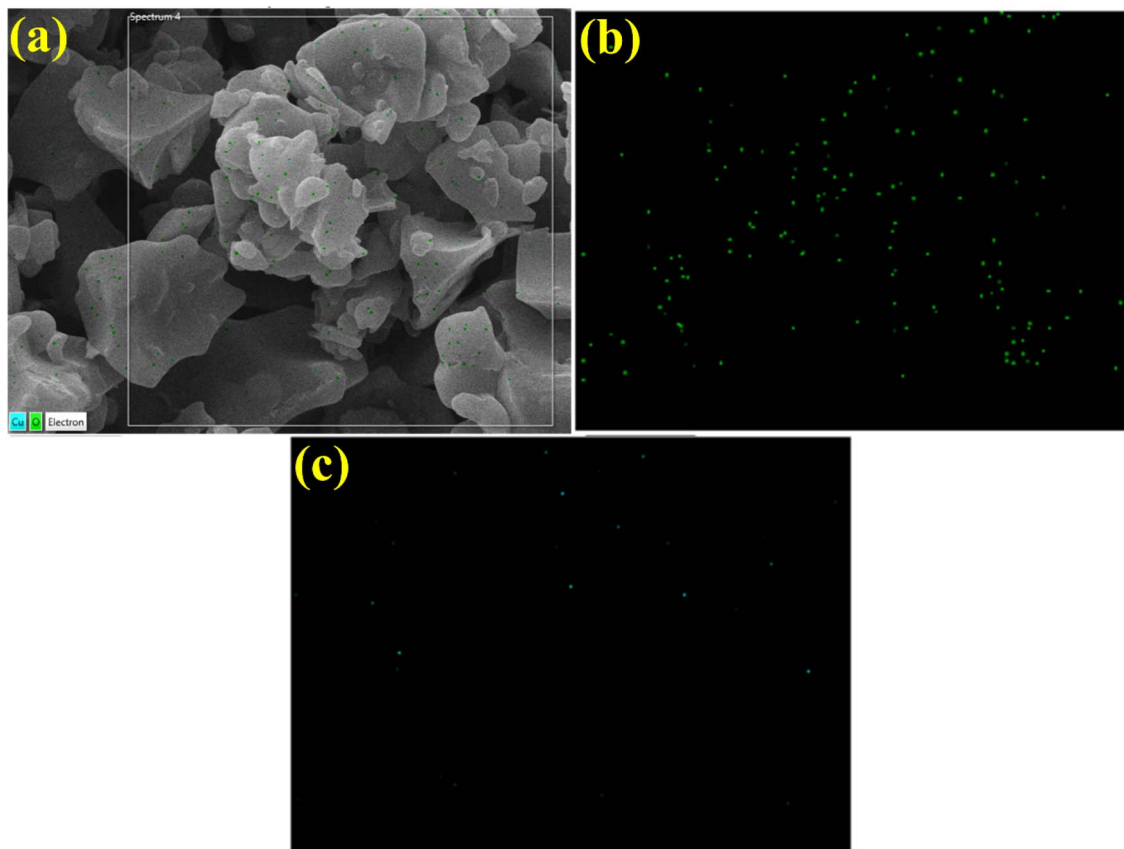


Fig. 8 EDS mapping of (a) QGDs (b) Ag/PAA-QGDs (c) PAA/Ag-QGDs.



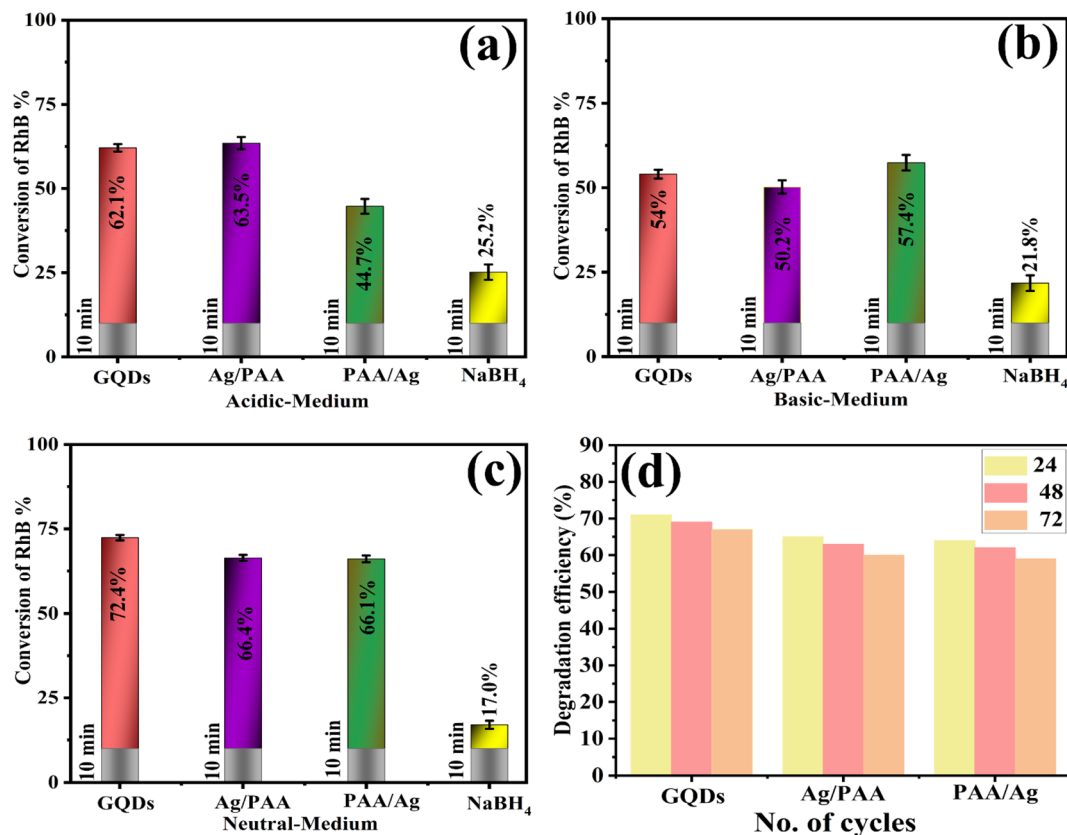


Fig. 9 Catalytic potential of GQDs, Ag/PAA-GQDs, and PAA/Ag-GQDs in (a) acidic, (b) basic, and (c) neutral media (d) catalysis recyclability studies.

material, reducing PAA adsorption. The incorporation of PAA/Ag in pure GQDs led to an increase in catalytic activity due to the presence of COOH, which exhibited electrostatic interactions with the catalyst.⁶⁶ The catalytic activity of pristine and doped GQDs is compared with the literature, as illustrated in Table 1.

3.3 Efficiency of GQDs, Ag/PAA-GQDs and PAA/Ag-GQDs

Moreover, the reusability of pure and doped GQDs catalysts was assessed by performing degradation using previously used samples. During each cycle, the catalyst was separated from the solution using centrifugation, washed with DI water, and heated overnight at 60 °C. The reusability of GQDs, Ag/PAA-GQDs, and PAA/Ag-GQDs were investigated over three

consecutive cycles (24 h, 48 h, and 72 h) for the RhB degradation, as depicted in Fig. 9d.

X-ray photoelectron spectroscopy (XPS) was employed to characterize the spectrum of PAA/Ag-GQDs. The intent was to ascertain the elemental makeup and chemical nature of constituent elements. Fig S2a–d† exhibits the narrow range XPS spectra of generated samples, specifically highlighting the Ag O 1s and Ag 3d peaks. The measured binding strength of Ag O 1s in PAA/Ag-GQDs Fig. S2a† was determined to be 530.8 eV, therefore consistent with previous investigations.^{72,73} The determination of the precise position of the binding strength spike was accomplished by considering the ionic and electro-negativity characteristics of constituent atoms within the molecule. The findings are comparable with previous characterization techniques⁷⁴ and confirm that silver nanoparticles are

Table 1 Oxidation of RhB with H₂O₂ in the presence of different catalytic systems

Nano-materials	Catalyst concentration	RhB concentration	Conversion of RhB (%)	Ref.
Graphene quantum dots decorated titania nanosheets	2 mL	10.20 mg L ⁻¹	56	67
Graphene quantum dots by P doping	100 mg	10 mg L ⁻¹	58	68
Graphene quantum dot-based hydrogels	100 µg mL ⁻¹	10 mg L ⁻¹	62	69
Graphene quantum dots from corn powder	1 mg mL ⁻¹	10 mg L ⁻¹	45	70
Ag and polyacrylic acid (PAA) doped SrO	400 µL	3 mL	60.7	71
Silver and polyacrylic acid doped graphene quantum dots	400 µL	1.5 mL	66.4	Present work

Table 2 Antimicrobial efficacy of GQDs, Ag/PAA-GQDs and PAA/Ag-GQDs

Samples	Inhibition areas (mm)	Inhibition areas (mm)
	0.5 mg/50 μ L	1.0 mg/50 μ L
GQDs	2.55	4.15
Ag/PAA-GQDs	3.15	5.25
PAA/Ag-GQDs	3.65	5.85
Ciprofloxacin	5.75	5.75
DI water	0	0

efficiently acquired following NaBH_4 reduction. On the contrary, the binding energies of 365.9 eV and 371.9 eV, accordingly, correspond to Ag $3d_{5/2}$ and Ag $3d_{3/2}$ Fig. S2b†. ^{75–78}

3.4 Biological activity

Antibacterial activity of doped and pure GQDs against *E. coli* was evaluated by agar well diffusion strategy. The inhibition zones were recorded as (4.15–5.85 mm) and (2.55–3.65 mm) at maximum and minimum doses, summarized in (Table 2). An inhibition region of 5.75 mm ciprofloxacin (positive control) and 0 mm of DI water (negative control) was calculated as illustrated in Fig S3a and b.† The inhibition diameter of pure GQDs was increased as the dopant was incorporated. Nano-material generated oxidative stress related to the crystallinity, surface area, and diffusion ability. Ag/PAA-GQDs exhibited superior antimicrobial performance because Ag provides a large surface area, producing more reactive oxygen species (ROS) that lead to cell necrosis. Carboxyl and the hydroxyl group of PAA increased the production of ROS, leading to the extrusion of cytoplasmic components that eventually caused bacterial death.

Doped GQDs destroy the bacterial cell by membrane distortion, enzymes inactivation, proteins denaturation, leakage of cytoplasmic components and DNA deterioration, *etc.*, ^{66–68,79–81} as displayed (Fig. S4†).

In the past few decades, there has been significant interest in molecular docking predictions for deciphering the enigma behind many biological functions. The significance of cell wall synthesis (*i.e.*, peptidoglycan production) and the nucleic acid biosynthetic route for identifying antibiotics is well established. ^{82,83} In spite of the fact that the antibacterial activity of several nanostructures has been described in recent years, ^{84,85} the specific mechanism of their actions requires additional investigation. The silver and polyacrylic acid-doped graphene quantum dots had a high binding score (5.13) inside the binding pocket of the β -lactamase enzyme in *E. coli*. The binding interaction pattern with important amino acid residues is shown in (Fig. 10a–c) *via* H-bonding with Gln120 and Asn152. The molecular docking predictions of silver and polyacrylic acid doped graphene quantum dots against DNA gyrase of *E. coli* revealed H-bonds with key amino acid residues such as Arg76, Thr165, and Gly77 (shown in Fig. 10d–f).

The binding tendency of silver and polyacrylic acid doped graphene quantum dots revealed through molecular docking predicted these NPs as potential inhibitors of β -lactamase and DNA gyrase enzyme that is suggested to be further confirmed by *in vitro* enzyme inhibition techniques.

4. Summary

In this research work, GQDs, Ag/PAA-doped GQDs and PAA/Ag-doped GQDs were successfully synthesized by the cost-effective carbonization method to remove various organic and inorganic hazardous pollutants. XRD diffraction peak of the GQDs is centered at 25.2° corresponding to the (002) plane with a *d*-

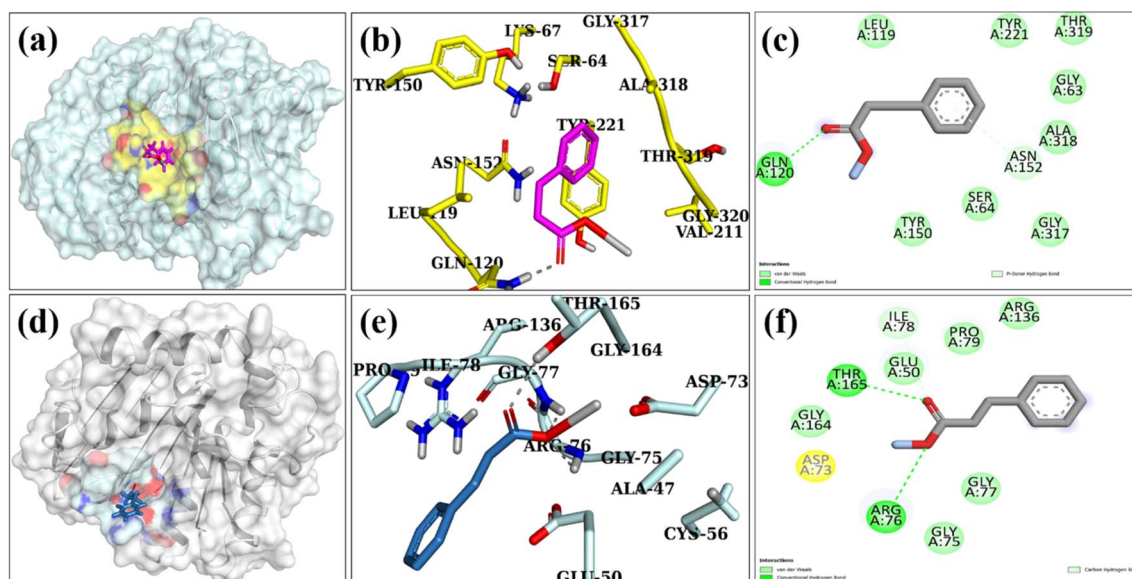


Fig. 10 Binding interaction pattern with active site residues of β -lactamase (a–c) and DNA gyrase (d–f) enzyme from *E. coli* where (a and d) represents binding pocket, (b and e) 3D structure and (c and f) 2D structure of NPs in the active site of selected proteins.



spacing of 0.366 nm. The increase in E_g from 3.03 to 3.14 eV accompanied by blue shift was exposed by UV-Vis spectroscopy. FTIR confirmed the presence of GQDs by displaying the vibration band of the sp^2 carbon plane. TEM results confirmed that Ag particles attached to the surface of GQDs and PAA formed a layer on the surface of GQDs additionally, HRTEM micrographs revealed interlayer d -spacing (0.366 nm, 0.324 nm, 0.325 nm). EDS spectra revealed the presence of C, O, Au, and Ag, confirming the elemental composition of pristine and doped GQDs. The maximum RhB deterioration rate of 57.42% and 66.41% in basic and neutral media was observed for pure and PAA/Ag-GQDs, respectively. *In silico* docking studies identified inhibition of β -lactamase and DNA gyrase as potential mechanisms underlying silver and polyacrylic acid doped graphene quantum dots bactericidal behavior. Furthermore, the significant inhabitation zone (5.85 mm) of PAA/Ag-GQDs against *E. coli* was recorded. In conclusion, these findings imply that synthesized pure and doped GQDs effectively eliminate toxic effluents from industrial wastewater (dye degradation) and are effective against pathogens, low cost, environment-friendly, and can be used in the future.

Conflicts of interest

The authors declare “no conflict of interest”.

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