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Computationally validated magnesium and graphene oxide anchored SnO₂ quantum dots for RhB reduction and antibacterial activity

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Developing multi-functional catalysts for wastewater purification and bactericidal inactivation is quite interesting but remains a significant challenge. Herein, varying concentrations (2 and 4 wt%) of magnesium (Mg) doped with a fixed amount of graphene oxide–stannic oxide quantum dots (GO–SnO₂ QDs) were prepared by a low temperature co-precipitation approach. This research provides dual approaches, experimental as well as theoretical, to study the properties of doping-dependent SnO₂ for rhodamine B degradation and MDR *S. aureus* inactivation. The addition of two-dimensional GO exposed more active sites, and Mg metal facilitates the transportation of charge carriers for catalytic and antibacterial activity. Advanced characterization confirmed the multiple phases (tetragonal, orthorhombic, and anorthic), polycrystalline behavior, high light absorption intensity, and the QD-like morphology of ternary catalysts. The combination of metal, 2D materials, and metal oxides was proven to be beneficial as they synergistically increased the catalytic activities. Among all synthesized samples, optimized 4% Mg/GO–SnO₂ revealed considerable (94.7%) rhodamine B (RhB) degradation in acidic medium. The optimized sample exhibited a 3.65 ± 0.03 mm inhibition zone towards MDR *S. aureus* in comparison to ciprofloxacin. Mg/GO-doped SnO₂ exhibited significant inhibitory effects on DNA gyrase and tyrosyl-tRNA synthetase (TyrRS), as corroborated by the *in silico* approach, indicating its potential as a therapeutic agent for bactericidal action.

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1 Introduction

Organic dyes and other potentially hazardous substances are released by industries into freshwater reservoirs, which exhibit complex detrimental impacts on aquatic ecosystems.¹ The effluent of hazardous chemicals by textile manufacturers contributes significantly towards the detrimental impacts on

the ecosystem and human wellness.² RhB is a cationic, extremely water-soluble dye that induces thyroid damage, allergic dermatitis, and adverse respiratory effects.^{3–5} Additionally, bovine mastitis is a particularly prevalent disease affecting dairy cows worldwide. The illness is multifactorial, caused by various infectious agents, leading to significant economic losses and negative impacts on the dairy industry, characterized by decreased milk yield and heightened culling and treatment costs.⁶ *Staphylococcus aureus* (*S. aureus*) is often regarded as a causative agent of infectious mastitis. The excessive use of antibiotics in the treatment of mastitis is the main factor contributing to increased resistance in dairy cows.^{7,8} The extensive use of antibiotics poses a considerable risk to human health, primarily through the development of bacterial resistance and the contamination of milk with antibiotic residues. An urgent need exists to develop a novel and safe approach for mastitis treatment that prioritizes public health.⁹

Adsorption, reverse osmosis, and ion exchange are the primary techniques for removing dyes from contaminated water, each offering unique benefits and limitations.¹⁰ In particular, catalytic decomposition has attracted interest as a pragmatic, energy-efficient, and environmentally sustainable approach.¹¹ In recent decades, semiconductor nanomaterials have attracted considerable attention from researchers in environmental development

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due to their notable physicochemical properties, especially their reduced toxicity and environmental sustainability.¹² Various methods have been employed to prepare metal oxides, including ZnO (hydrothermal method and nanoflake-like morphology),¹³ Al₂O₃ (sol-gel method and nanoparticles),¹⁴ V₂O₅ (ultrasound-assisted synthesis and nanoparticles (NPs)),¹⁵ and SnO₂ (co-precipitation method and NPs).¹⁶ Among these, co-precipitation-synthesized SnO₂ has now been suggested as an efficient catalyst owing to its chemical stability, cost-effectiveness and low toxicity.¹⁷ This method provides economic viability and regulates NP synthesis, surface area, and stoichiometry without needing high-temperature systems.^{18–20} Furthermore, SnO₂ is an n-type semiconductor with a wide bandgap energy (E_g), high electron mobility, and low resistivity.^{21–24} However, SnO₂ has a rapid e⁻/h⁺ recombination rate, which limits its application for RhB degradation.²⁵

To address this issue, carbon-based nanomaterials such as g-C₃N₄, graphene oxide (GO), and carbon nanotubes were employed due to their chemical stability and high conductivity.²⁶ GO, a 2D aromatic compound, is considered the best option for improving the catalytic reduction of RhB as well as antibacterial activity due to its biocompatibility, extensive surface area, low E_g and high adsorption capacity.²⁷ So, we selected GO as a doping agent. The dye reduction efficacy of SnO₂ was further enhanced by doping with metals (Be, Mg, Ca and Sr), as they produce lattice imperfections, including oxygen vacancies and enhanced the formation of reactive oxygen species (ROS).¹²

To our knowledge, no reports have investigated the co-precipitation synthesis of Mg/GO-SnO₂ QDs for catalytic and antibacterial activity.

2 Experimental section

2.1 Materials

SnCl₂·2H₂O (purity~99%), graphite powder (99%), MgCl₂·6H₂O, 99%, sodium nitrate (NaNO₃, 99.9%), hydrogen

peroxide (H₂O₂), potassium permanganate (KMnO₄, 99.5%) and NaOH (99%) were acquired from Sigma Aldrich (Germany) while, H₂SO₄, HCl, and H₃PO₄ were obtained from Analar.

2.2 Synthesis of SnO₂ and Mg/GO-SnO₂

SnO₂ was prepared based on our previous studies with a few characterization results reused (<https://doi.org/10.1039/D3RA00698K>).²⁸ The prepared material is named SnO₂ (sample 1) in the subsequent text. Graphene oxide (GO) was prepared by the modified Hummer approach, as reported earlier.²⁹ GO-SnO₂ was prepared by adding 2 wt% GO in SnCl₂·2H₂O solution under the same conditions and this is named GO-SnO₂ (sample 2) in the remaining part of the manuscript. Similarly, ternary catalysts (Mg/GO-SnO₂) were prepared using 2 and 4 wt% Mg in GO/SnCl₂·2H₂O solution, keeping the same conditions (Fig. 1) and are referred to as 2 and 4% Mg/GO-SnO₂ (sample no 3 and 4) in the subsequent text.

2.3 Catalytic dye degradation activity

The catalytic RhB reduction efficacy of SnO₂, GO-SnO₂ and Mg/GO-SnO₂ was evaluated in the dark in the presence of a reducing agent (NaBH₄). Firstly, 0.1 M NaBH₄ was injected into 5 mL RhB solution (5 mg/500 mL) under constant agitation. Subsequently, 400 μL of SnO₂, GO-SnO₂ and Mg/GO-SnO₂ solutions (5 mg/10 mL) were integrated into the above solution, and RhB reduction was recorded using a UV-vis spectrophotometer after specific intervals.

2.4 Segregation and characterization of *S. aureus*

Samples of raw milk were collected from mastitis-positive cows milked at different locations in Punjab, Pakistan, including marketplaces and veterinary facilities. The milk specimens were promptly delivered to the laboratory at 4 °C following collection. The growth potential of *Staphylococcus aureus* (*S. aureus*) in

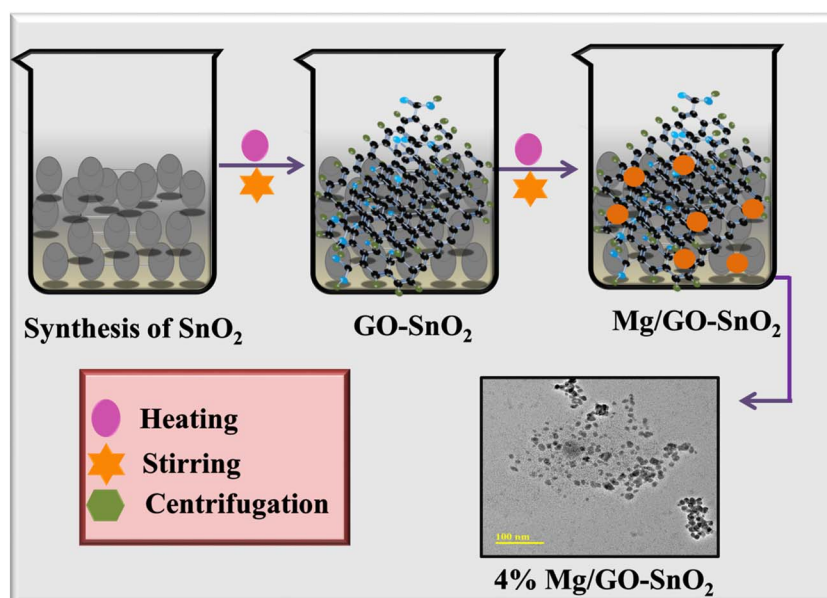


Fig. 1 Schematic preparation of Mg/GO-SnO₂.



unpasteurized milk was evaluated by cultivating specimens thrice on mannitol salt agar (MSA) following aerobic incubation at a specific temperature (37 °C) for 48 hours. The purified culture attained was characterized morphologically and biochemically using Gram staining and different biochemical tests. Furthermore, the susceptibility of isolated *S. aureus* for different antibiotics was analyzed and the resulting bacteria found resistant to at-least three drugs were classified as multiple drug resistant (MDR).⁷

2.4.1 Antibacterial effectiveness. Ten distinct MDR *S. aureus* strains obtained from mastitis milk were assessed for the antibacterial effectiveness of SnO₂, GO-SnO₂ and (2 and 4 wt%) Mg/GO-SnO₂ using the well diffusion approach. Petri dishes containing MSA were swabbed with 0.5 McFarland suspension of MDR *S. aureus*. Control SnO₂ and doped SnO₂ were loaded into 6 mm wells on MSA plates at concentrations of 500 and 1000 µg/50 µL. The positive control treatment (5 µg/50 µL) included ciprofloxacin, while the substitute control was composed of (50 µL) DI water. The extent of inhibition domains was assessed quantitatively utilizing a digital vernier scale after incubation at 37 °C for 24 hours. The statistical validation of quantified inhibition domains (mm) considering a significance level <5% was performed using one way analysis of variance (ANOVA) with SPSS 24.0.

2.5 Molecular docking analysis

A plethora of nanocomposites have shown efficiency in the elimination of bacteria. QDs have been shown to have biological potential, but further study is needed to pinpoint how exactly they exert their therapeutic effects. Researchers from all over the world are now able to undertake in-depth assessments of a wide range of interactions and developments owing to the widespread use of computational approaches, in particular molecular docking studies. In light of the notable antibacterial

properties demonstrated by the synthesized nanocomposites, our principal aim in this study was to assess their viability as molecular docking-based inhibitors of specific enzymes. We tested the inhibitory effect of Mg/GO-doped SnO₂ on DNA gyrase and tyrosyl-tRNA synthetase (TyrRS) to establish its potential against *S. aureus*. Crystal structures of DNA gyrase_{*S. aureus*} and TyrRS_{*S. aureus*} (PDB codes 5CTU and 1JJJ; <https://www.rcsb.org/>) were downloaded from the RCSB PDB. For the docking study, we used the SYBYL-X 2.0 program. To construct 3D models of the compounds of interest and investigate interactions between QDs and active site residues of the proteins of interest, we used the SYBYL-X 2.0 program in the same way as described in our previous paper.^{30,31}

3 Results and discussion

The structural properties of SnO₂ and Mg/GO-SnO₂ were characterized by XRD analysis, over a 2θ range of 20 to 70° (Fig. 2a). Peaks at 26.4, 33.7, 37.8, 51.5, 54.5, 61.6, and 65.5 correspond to the tetragonal phase (*P4₂/mnm*, no. 136) of SnO₂, as confirmed by JCPDS card no. 01-077-0452. Other diffraction peaks at 24.9° (111), 29.8° (101), 31.2° (020), 45.4° (024), and 47.0° (115) are ascribed to orthorhombic (*Cmc2₁*, no. 36) and tetragonal phases (*P4/nmm*, no. 129) of SnO (JCPDS card no. 01-077-2296/00-006-0395). Additional peaks at 21.7° (020), 56.3° ($\bar{3}\bar{1}1$) and 58.6° (0 $\bar{3}$ 2) confirmed the anorthic structure (*P**, no. 2) of Sn₂O₃ (JCPDS card no. 00-025-1259). Compared to the pure sample, the relative peak intensity or crystallinity of GO-SnO₂ decreased owing to various functional groups, including carboxylic acid and hydroxyl.^{32,33} With Mg, the crystallinity was further reduced as Mg²⁺ (0.072 nm) has a larger ionic radius compared to Sn⁴⁺ (0.071 nm).¹² SAED patterns of GO-SnO₂ and 4% Mg/GO-SnO₂, containing identifiable bright rings showing polycrystalline behavior (Fig. 2b and c).

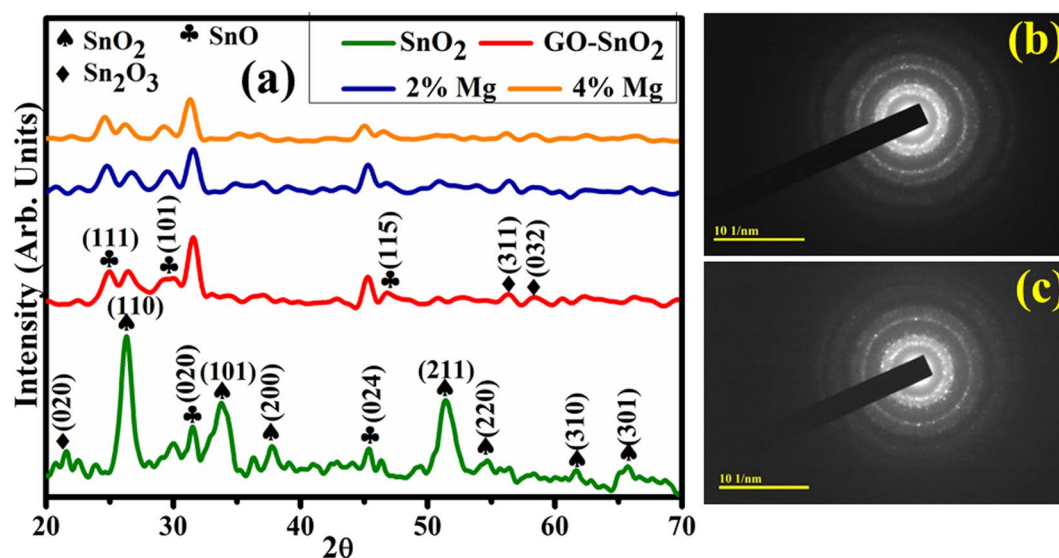


Fig. 2 (a) Diffraction patterns of SnO₂,³⁴ GO-SnO₂, 2% Mg/GO-SnO₂ and 4% Mg/GO-SnO₂ and (b and c) SAED analysis of GO-SnO₂ and 4% Mg/GO-SnO₂.



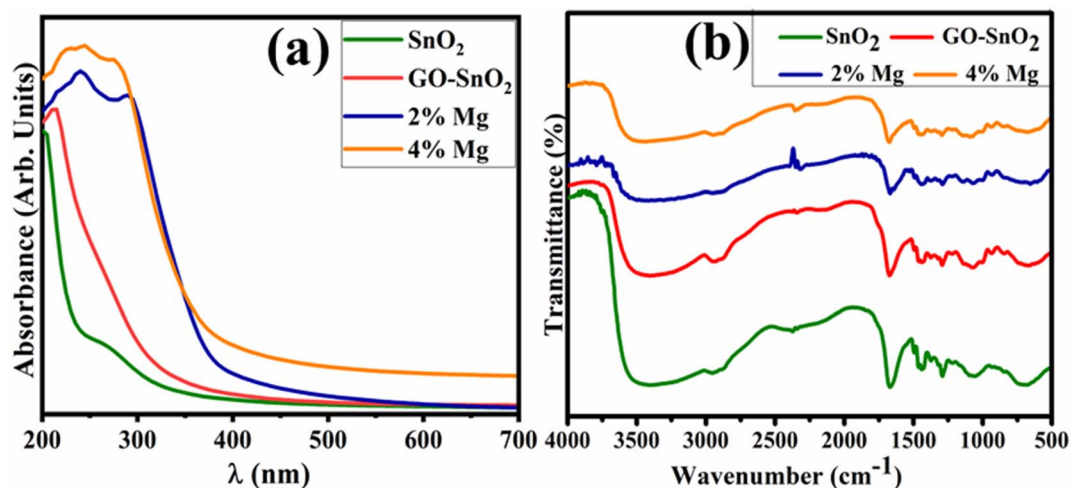


Fig. 3 (a) UV-vis spectra, (b) FTIR analysis of SnO₂, GO-SnO₂, 2% Mg/GO-SnO₂ and 4% Mg/GO-SnO₂.

The electronic structure of SnO₂ and Mg/GO-SnO₂ was characterized through UV-vis spectroscopy (Fig. 3a). SnO₂ showed absorption from 200 to 700 nm with a pronounced hump around 285 nm assigned to π - π^* transition.³⁴ With Mg and GO, the absorption intensity of SnO₂ enhanced, introducing a redshift attributed to increased defect sites.³⁵ From the equation $E_g = 1240/\lambda_{\text{onset}}$,³⁶ the calculated E_g of SnO₂ decreased from 4.30 to 4.19 eV with the addition of GO and Mg.³³ The

functional group identification of SnO₂ and Mg/GO-SnO₂ was analyzed utilizing FTIR analysis (Fig. 3b). The strong and broad bands at 3300–3500 and 1600–1640 cm⁻¹ are attributed to -OH stretching and bending modes, respectively.³⁷ The bands ~1120–1381 cm⁻¹ are attributed to Sn-OH bending vibration. The bands ~500–650 cm⁻¹ correspond to O-Sn-O bending and Sn-O stretching modes, respectively.³⁸ With the addition of GO, variations in intensity and position were observed, caused by

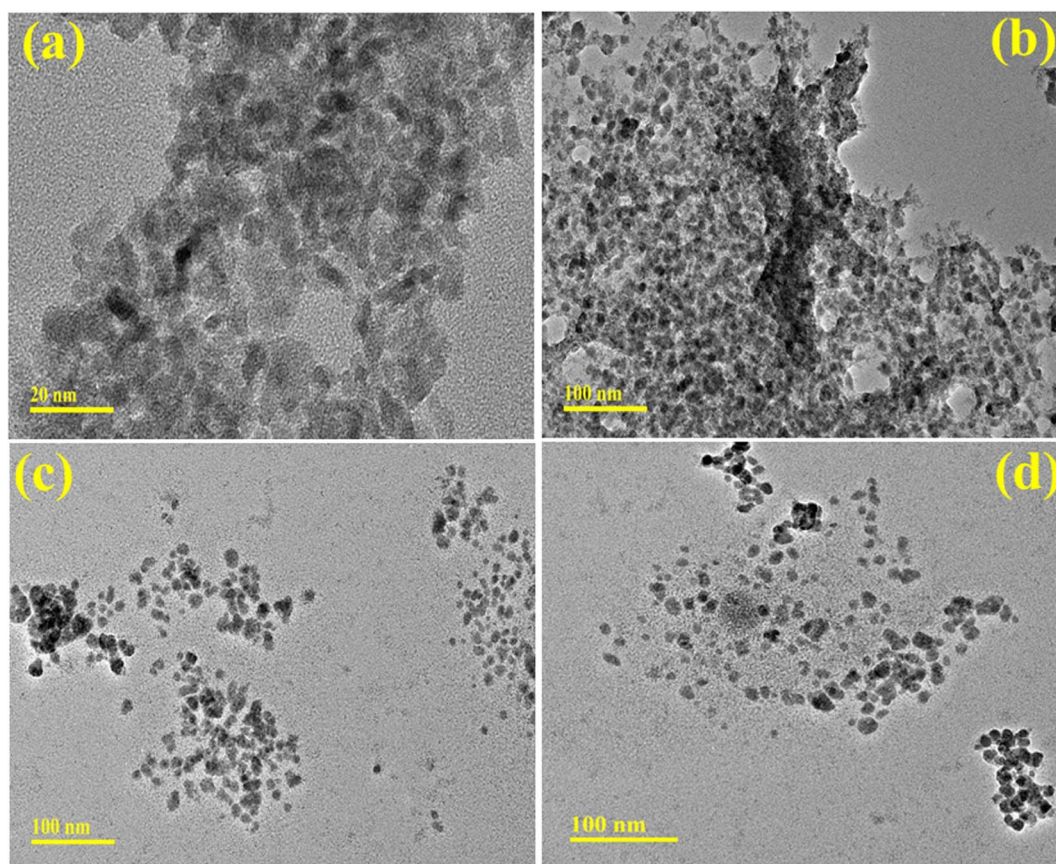


Fig. 4 (a–d) TEM analysis of (a) SnO₂,³⁴ (b) GO-SnO₂, (c) 2% Mg/GO-SnO₂, and (d) 4% Mg/GO-SnO₂.



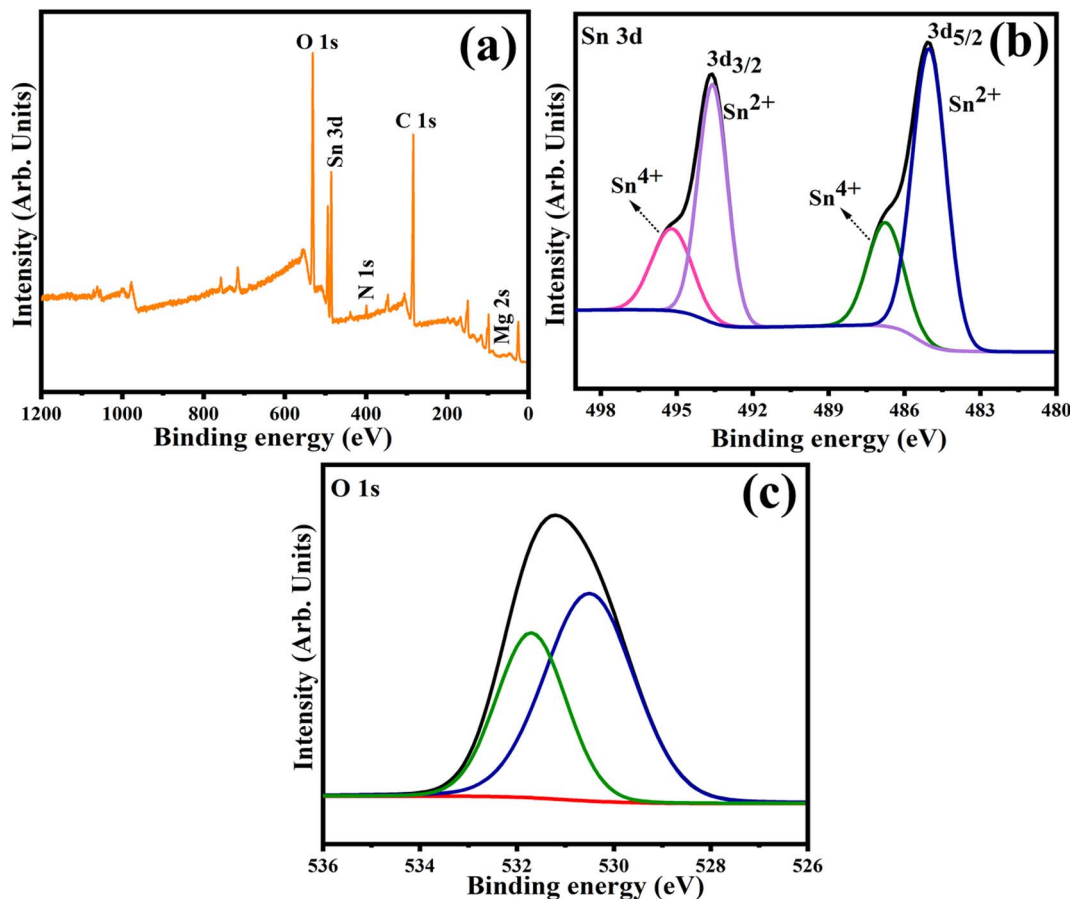


Fig. 5 (a) XPS survey spectrum of 4% Mg/GO-SnO₂ and the high-resolution spectra of (b) Sn 3d, and (c) O 1s.

the contribution of functional groups in GO-SnO₂.³⁹ The same trend was observed with Mg, possibly due to the introduction of Mg²⁺ into the SnO₂ crystal structure.⁴⁰

EDS analysis characterized the elemental configuration of SnO₂ and Mg/GO-SnO₂ (Fig. S1a-d). Sn, O, C, and Mg peaks clarified the formation of Mg/GO-SnO₂. The Cl peak emerged as a result of using the precursor (SnCl₂·2HO) in SnO₂ preparation. A Na peak appeared as NaOH was utilized during the synthesis process. Various concentrations of elements (Sn, O, C, Mg, and Na) *via* specific colors were analyzed through EDS mapping images (Fig. S1a'-d').

The morphological properties of SnO₂ and Mg/GO-SnO₂ were characterized through TEM analysis (Fig. 4a-d). Fig. 4a demonstrates the non-uniform, randomly oriented, interconnected network of SnO₂ quantum dots with an average size of 4.55 nm measured through ImageJ software. With GO, overlapping and covering of nanosheets with interconnected QDs were observed (Fig. 4b). With 2% Mg, agglomeration decreased, and the same behavior was observed with a higher concentration (4%) of Mg (Fig. 4c and d).

To characterize the oxidation states of 4% of Mg doped GO-SnO₂, XPS analysis was utilized. Fig. 5a clarifies the presence of O 1s, Sn 3d, N 1s, C 1s, and Mg 2s. In the Sn 3d spectrum, peaks at 486.75 eV (Sn⁴⁺) and 485.02 eV (Sn²⁺) are ascribed to Sn 3d_{5/2} and peaks at 495.2 eV (Sn⁴⁺) and 493.58 eV (Sn⁴⁺) are attributed to Sn

3d_{3/2} (Fig. 5b).⁴¹ In the high resolution O 1s spectrum, peaks at 530.5 eV (lattice oxygen bound to Sn) and 531.7 eV (oxygen deficient region in SnO₂) are observed, as shown in Fig. 5c.⁴²

The degradation of RhB dye by prepared SnO₂ and Mg/GO-SnO₂ QDs was analyzed through a UV-vis spectrophotometer (absorption mode). The mechanism for the reduction of RhB

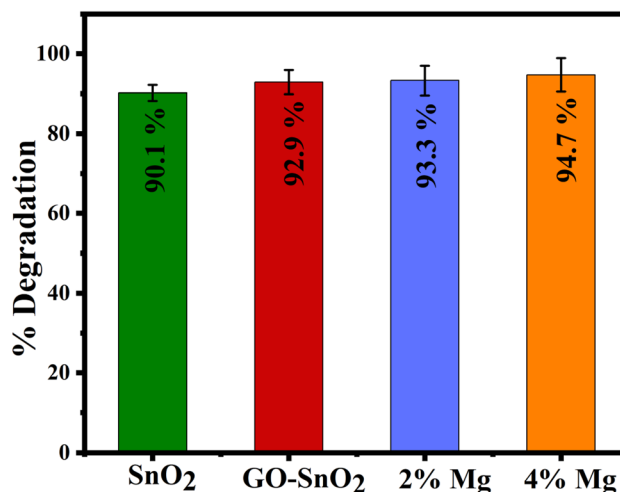


Fig. 6 Catalytic efficacy of SnO₂, GO-SnO₂, 2% Mg/GO-SnO₂, and 4% Mg/GO-SnO₂ in acidic medium.



was observed in the dark (absence of sunlight) and in the presence of synthesized catalysts (SnO_2 and $\text{Mg}/\text{GO}-\text{SnO}_2$) and a reducing agent (NaBH_4). During this process, prepared samples serve as an electron transfer medium, transferring e^- from NaBH_4 to the colored RhB dye (Fig. S2). The transfer of electrons to RhB causes the breakdown of the colored dye into less harmful compounds. SnO_2 , $\text{GO}-\text{SnO}_2$, 2% $\text{Mg}/\text{GO}-\text{SnO}_2$, and 4% $\text{Mg}/\text{GO}-\text{SnO}_2$ showed 90.1, 92.9, 93.3, and 94.7% degradation efficacy in acidic medium, respectively (Fig. 6). The catalytic efficacy is a multidimensional process controlled by a variety of parameters such as size, dimension and surface area of the synthesized catalysts and pH of the catalyst medium. At pH 4, the immediate availability of H^+ ions protonated the surface, significantly enhancing the ionic interaction of RhB on

the catalyst surface and demonstrating a high reduction of RhB.²⁸ Compared to SnO_2 , $\text{GO}-\text{SnO}_2$ revealed improved catalytic efficiency, ascribed to $\pi-\pi$ conjugation with the RhB chromophore group and high charge transfer efficacy.⁴³ With the dopants (Mg and GO), the catalytic efficacy is enhanced as the reduced size of SnO_2 leads to more active sites for dye reduction.

The antibacterial potential of SnO_2 , $\text{GO}-\text{SnO}_2$, 2% $\text{Mg}/\text{GO}-\text{SnO}_2$, and 4% $\text{Mg}/\text{GO}-\text{SnO}_2$ was analyzed *in vitro* (Fig. S3), and the obtained outcomes are tabulated in Table 1. The inhibition zone diameters were found to be 0 ± 0.0 mm at 0.5 mg/50 μL (lowest concentration) and ranged from 1.85 ± 0.05 – 3.65 ± 0.03 mm at 1.0 mg/50 μL (highest concentration). The findings were contrasted with those of DI water (0 ± 0.0 mm) and ciprofloxacin, which exhibited a 6.85 ± 0.01 mm inhibitory zone. The optimized $\text{Mg}/\text{GO}-\text{SnO}_2$ demonstrated the highest antibacterial efficacy due to various synergistic processes. The catalyst likely facilitated the generation of reactive oxygen species (ROS), including hydroperoxyl radicals, hydroxyl radicals, superoxide anions, and hydrogen peroxide, which produced oxidative stress, damaging bacterial proteins and DNA ultimately leading to cell death. The 4% $\text{Mg}/\text{GO}-\text{SnO}_2$ QDs immobilized bacterial cells, inhibiting their multiplication, while the disruption of glycolysis impeded energy generation. Moreover, the liberation of metal ions interferes with bacterial enzymes and proteins. The combined effects lead to improved antibacterial effectiveness of 4% $\text{Mg}/\text{GO}-\text{SnO}_2$, the catalyst,

Table 1 Antibacterial efficacy of pure and $\text{Mg}/\text{GO}-\text{SnO}_2$ QDs

Samples	Zone of inhibition (mm)	
	500 $\mu\text{g}/50 \mu\text{L}$	1000 $\mu\text{g}/50 \mu\text{L}$
SnO_2	0 ± 0.0	1.85 ± 0.05
GO/SnO_2	0 ± 0.0	2.15 ± 0.04
2% Mg	0 ± 0.0	2.95 ± 0.04
4% Mg	0 ± 0.0	3.65 ± 0.03
Ciprofloxacin	6.85 ± 0.01	6.85 ± 0.01
DIW	0 ± 0.0	0 ± 0.0

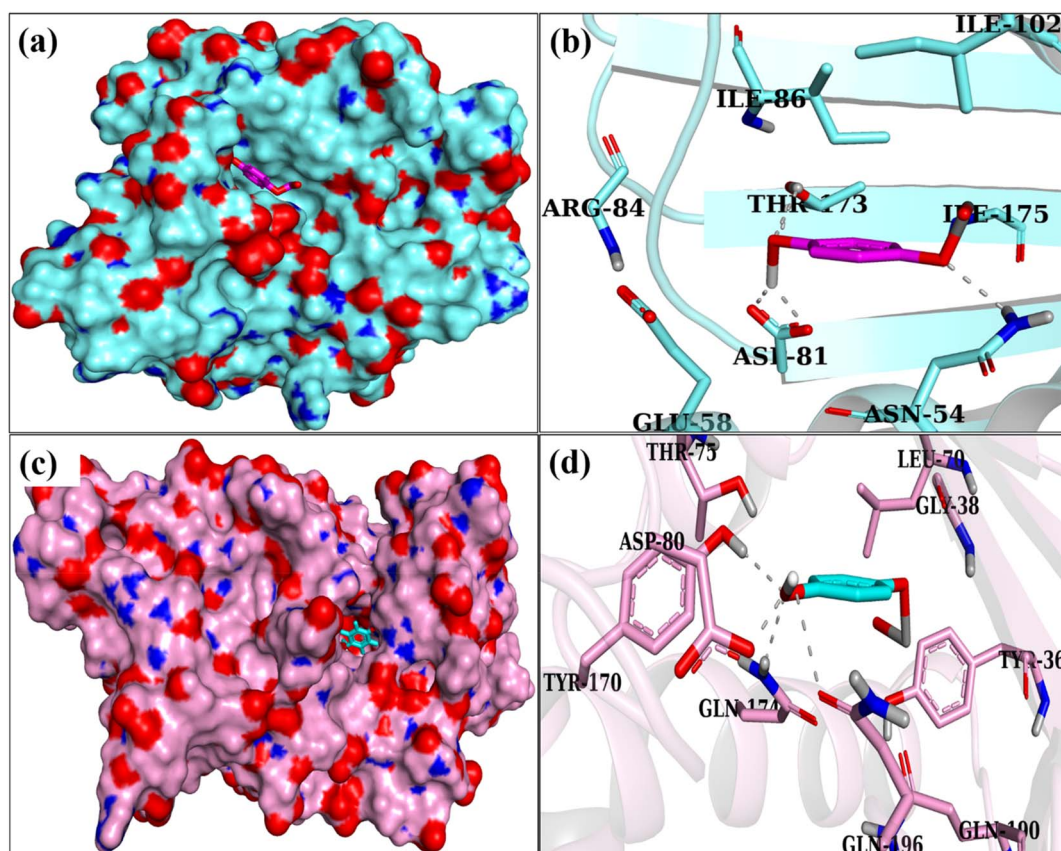


Fig. 7 The corresponding binding pockets (a and c) and binding interaction patterns (b and d) of Mg/GO -doped SnO_2 with active site residues of DNA gyrase and tyrosyl tRNA synthetase enzymes.



highlighting its promise for advantageous antimicrobial applications.^{44–48}

The assessment of antioxidant activity of Mg/GO–SnO₂ was performed using DPPH free radical scavenging at 517 nm wavelength, with concentrations ranging from 25–125 μg mL⁻¹ (Fig. S4). An improvement in scavenging efficacy is dependent on concentration, suggesting effective electron transfer. The 4% Mg-doped sample exhibited noteworthy antioxidant performance, attributed to increased availability of active sites and improved charge-transfer characteristics stemming from Mg integration, thus facilitating more efficient reduction of DPPH. The findings underscore the potential applications of QDs in the fields of biomedicine and therapeutic strategies.

In recent decades, molecular docking predictions have gained a lot of attention as a powerful tool for understanding different types of biological processes. There has been a lot of progress in recent years in identifying nanostructures with antibacterial capabilities, but there is still room for improvement in understanding the mechanisms that underlie these effects. The docking investigation of the DNA gyrase enzyme against *S. aureus* demonstrated that Mg/GO doped SnO₂ exhibited the most favorable configuration. Hydrogen bonding interactions were seen between Asn54, Gly85, and Thr173 in this conformation, leading to a binding score of 4.58 (Fig. 7a and b). Similar behavior was observed for the binding of Mg/GO doped SnO₂ into the active pocket of TyrRS_{*S. aureus*}, leading to an

overall binding score of 4.51. Fig. 7c and d highlight the significant hydrogen bonding interactions between Asp80, Thr170, Gln174, and Gln196.

3.1 DFT and MESP studies

The electronic structures and reactivity of Mg/GO-doped SnO₂ in gaseous and solvent phases were clarified by DFT and MESP calculations using the B3LYP functional with the SVP basis set in Gaussian 09 (Rev. E.01) (Fig. 8a). The greatest electronegativity areas are indicated as deep red hue on MESP maps, highlighting regions for electrophilic and nucleophilic attacks on molecules. The oxygen atoms in Mg/GO-doped SnO₂ rings have Mulliken charges of –0.888036 and –0.836170, indicating a preponderance of negatively charged areas in respective aqueous and gas phases. The docking results indicate that proximity of oxygen atoms amplifies the capacity for hydrogen bonding in hinge and solvent-exposed areas of Mg/GO-doped SnO₂. The dominant green coloration on the Mg/GO-doped SnO₂ surface suggests an increased presence of neutral regions, potentially involved in van der Waals or hydrophobic interactions. The molecular surface depicted in Fig. 8b illustrates frontier orbitals, specifically the HOMO and LUMO for Mg/GO-doped SnO₂, which signify the molecule's capacity for effective electronic transitions, irrespective of increased chemical reactivity. The recorded energy gaps of Mg/GO-doped SnO₂ for gaseous and aqueous phases are 2.773 and 0.525 eV. The

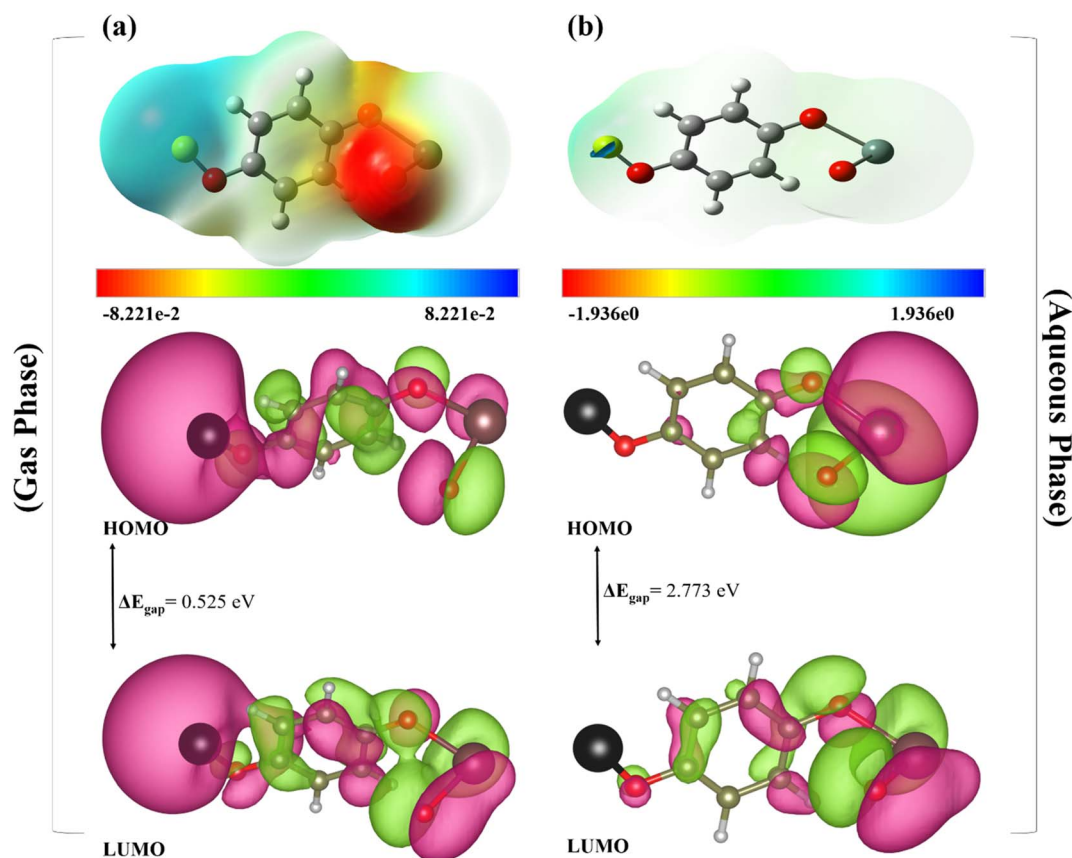


Fig. 8 MESP and HOMO–LUMO analysis of the selected ligand Mg/GO–doped SnO₂: (a) gas phase and (b) aqueous phase.



Table 2 DFT calculations (quantum chemical descriptors) of the selected ligands

Ligand	Dipole moment (Debye)	HOMO (a.u.)	LUMO (a.u.)	Energy gap (ΔE_{gap})	Ionization potential (eV)	Electron affinity (eV)	Electronegativity χ (eV)	Electrochemical potential μ (eV)	Hardness η (eV)	Softness S (eV^{-1})	Electrophilicity ω (eV)
Mg/GO-doped SnO_2 (gas)	6.4896	-0.17464	-0.1553	0.52 eV	4.753	4.227	4.490	-4.490	0.263	3.802	38.34
Mg/GO-doped SnO_2 (aqueous)	32.5162	-0.15290	-0.0507	2.77 eV	4.162	1.387	2.775	-2.775	1.388	0.720	2.775

difference in energy between the two phases can be attributed to the influence of solvent. Within the aqueous environment, the polar solvent enhances the stabilization of molecular orbitals through dielectric screening, consequently augmenting the energy separation between these orbitals.

The solvent phase of Mg/GO-doped SnO_2 exhibited a dipole moment of 32.5162 debye, implying stable interactions, as outlined in Table 2. In conclusion, the aqueous phase exerts a stabilizing effect on the electronic structure, leading to wider energy gaps and moderated reactivity characteristics. This thorough electronic structure analysis explores structural attributes and functional performance, fostering exploration of nanostructures with enhanced therapeutic effectiveness.

4 Conclusion

SnO_2 and Mg/GO- SnO_2 QDs were effectively prepared for catalytic RhB reduction and bactericidal inactivation. XRD depicted the reduction in crystallinity with GO and Mg. UV-vis spectroscopy indicated increased absorption upon doping and a decreased E_g of SnO_2 from 4.30 to 4.19 eV. TEM analysis revealed multiple-shaped SnO_2 QDs and a decrease in QD size was observed with GO and Mg. The optimized Mg/GO- SnO_2 showed 94.7% RhB reduction in an acidic medium and similarly, a 3.65 ± 0.03 mm inhibition zone against MDR *S. aureus*. The results of molecular docking simulations exhibited concordance with the observed *in vitro* bactericidal effects against MDR *S. aureus*, indicating that the inhibition of DNA gyrase and TyrRS enzyme could potentially serve as a mechanism underlying the antibacterial properties of the unique Mg/GO doped SnO_2 nanocomposites. In summary, the optimized Mg/GO- SnO_2 synthesized through a co-precipitation approach exhibits greater potential as a catalyst for the reduction of RhB and the bactericidal inactivation of MDR *S. aureus*. The present research was conceived as a preliminary screening investigation; moreover, thorough evaluations based on MIC, alongside *in vivo* efficacy and cytotoxicity of the synthesized QDs are required in future.

Author contributions

Eman Rai – investigation, writing original draft, data curation, Muhammad Imran – conceptualization, supervision, methodology, Iram Shahzadi – software, writing original draft, formal analysis (computational study), Ali Haider – investigation, methodology (antibacterial part), Sawaira Moeen – writing review and editing, visualization, data curation, Anwar Ul-Hamid – resources, TEM investigation, I Boukhris – resources, writing review and editing, validation, Muhammad Ikram – conceptualization, funding acquisition, project administration, writing review and editing. The 1st author completed her research work under the supervision of 2nd author (currently in KFUPM).

Conflicts of interest

The authors declare no known conflict of interest.



Data availability

The data used in this manuscript can be made available upon reasonable request.

Supplementary information (SI) is available. See DOI: <https://doi.org/10.1039/d5na00915d>.

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