



Green chemistry synthesis of biocompatible ZnS quantum dots (QDs): their application as potential thin films and antibacterial agent

Jejiron Maheswari Baruah¹ · Sanjeeb Kalita² · Jyoti Narayan¹

Received: 16 February 2018 / Accepted: 15 March 2019 / Published online: 27 March 2019
© The Author(s) 2019

Abstract

We are presenting here the synthesis of quantum dots (QDs) of direct band gap semiconductor, cubic ZnS through modified green chemistry-mediated chemical precipitation reaction. Green chemistry-synthesized (GCS) ZnS QDs were characterized using powder X-ray diffraction and high-resolution transmission electron microscope techniques. Analysis of results, revealed by both the techniques for the synthesized QDs, is complementary as far as the size range (2–6 nm) of ZnS QDs is concerned. UV–Vis spectrophotometric spectrum ($\lambda_{\text{max}} = 314 \text{ nm}$) showed a conspicuous blue shift than the bulk. The Fourier-transformed infrared spectra convincingly reported a Zn–S bond stretching frequency at 649 cm^{-1} . The characterized QDs were subjected to the preparation of thin films over SiO_2 template (57 nm thickness) using photoresist spin coating technique at the ambient condition. The surface topology of nanoscale-thick films was studied by atomic force microscope (roughness parameter—33.28 nm, rms; for a scan area of $3.48 \times 3.48 \mu\text{m}^2$). The symmetrical (skewness = 1.68) and random distribution (kurtosis = 2.93) of the peaks and valleys revealed the nanoscale-thick films of ZnS QDs. Zeta potential (−9.2 mV) fairly proved stable existence of ZnS QDs. The GCS QDs were found to be non-toxic toward L929 mouse fibroblastic cells and human erythrocytes. However, they demonstrated significant inhibitory effects against seven bacterial pathogens with an average zone of inhibition of 1.5 cm at 100 $\mu\text{g/ml}$ concentration. The minimum inhibitory concentrations determined were in the range of 75 to 125 $\mu\text{g/ml}$ for gram-positive and 100 to 150 $\mu\text{g/ml}$ for gram-negative bacterial pathogens.

Keywords Zinc sulfide quantum dots · Green synthesis · Thin films · Biocompatibility · Antibacterial effect

Introduction

Today's world is significantly pursuing the research that indulges in the design, development and implementation of chemical products and processes to reduce the use and

generation of products that are hazardous to human health and environment [1–3]. The sustainable and renewable green energy, as an alternative to carbon-emitting fossil fuels, has become the main target of green energy researchers [4–6]. Greenhouse gases are the main by-products of carbon-based traditional energy sources. To eliminate hazardous environmental issues, photovoltaic solar cells (PVSCs) apart from nuclear, thermal and wind energy have gained the momentum [7]. Presently, quantum dot-sensitized solar cell (QDSSC) is gaining the preference as the most effective PVSC [8–10], although it is still at the investigation level only.

The semiconducting quantum dots (QDs) have full access to the whole solar spectrum. They have the additional advantage of production of high quantum yield by the generation of multiple excitons [11–16]. Semiconducting QD (SQD) metamaterial has potential applications in the field of smart LEDs (light emitting diodes) [17–20], photocatalytic activity [21–24], space science [25, 26], biological fields such as biosensors, biomarkers and as bio-tagging agents [27–30].

Electronic supplementary material The online version of this article (<https://doi.org/10.1007/s40089-019-0270-x>) contains supplementary material, which is available to authorized users.

✉ Jyoti Narayan
jnarayan.nehu@gmail.com

- ¹ Synthetic Nanochemistry Laboratory, Department of Basic Sciences and Social Sciences (Chemistry Division), School of Technology, North Eastern Hill University, Shillong 793022, India
- ² Drug Discovery Lab, Biological and Chemical Sciences Section, Life Sciences Division, Institute of Advanced Study in Science and Technology (An Autonomous Institute Under Department of Science and Technology Government of India), Guwahati, Assam 781035, India



It has been observed that SQDs of IIB–VIA groups are efficient metamaterial for PVSC applications due to their direct band gap, large excitons binding energy [31, 32] and a high index of refraction [33]. The literature reveals that the nanometer range of the particles enhances the properties of semiconducting metamaterial due to its quantum confinement effect [34]. Among the nano-materials of group IIB–VIA, zinc sulfide (ZnS) QDs have wide and direct band gap $E_g = 3.58$ eV at 300 K (bulk cubic crystals), which might make it suitably promising optical material toward the design of solar cell. Keeping in view the above advantage and multi-faceted applications, the present work has been taken into consideration. The present work reports the green synthesis of SQDs of ZnS at the ambient conditions. The eco-friendly synthesis involves the use of water-soluble precursors and capping agents, resulting in the formation of non-hazardous by-products which can further easily be removed by repeated washing with water. The green chemistry synthesis (GCS) process was found to be cost-effective, reproducible, energy-efficient and sustainable. It has been observed that the mechanized thin film solar cells (TFSCs) are much thinner and light weighted as compared to the traditional first-generation solar cells [35, 36]. This has led to the development of thin films using nano-materials providing better efficiency as compared to the previously available one. The thin films are considered to be third-generation [37–42] engineered device that can hold the desired PVSC applications quite well. Keeping in view this, the colloidal solution of ZnS nanoparticles was used for the fabrication of variable nano-sized (< 100 nm) thin films in order to study their structural morphology and surface topology. In order to make the thin films eco-friendly, the QDs used for making thin films need to be analyzed for their biocompatibility and antibacterial activity. Thus, green-synthesized QDs of ZnS semiconducting metamaterial were further subjected to the studies on their biocompatibility and zone of inhibition (ZOI) over human erythrocytes, mouse fibroblast cells and seven (gram positive and gram negative) bacterial pathogens, respectively.

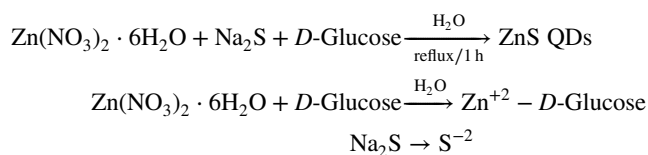
Experimental section

Materials and methods

All the chemicals used during the synthesis that were of analytical grade are reported under Supplementary information. Series of green synthetic reactions were carried out by either changing the concentration of precursors or capping reagent. At first, the $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (1 M, 1.487 g) was added to 5 ml of double distilled water (A). The solution was stirred well on a magnetic stirrer at room temperature (pH = 6.1) for about 5 min in order to get a homogeneous solution.

In the meantime, D-glucose (2 M, 1.9816 g) solution was prepared in 5 ml of double distilled water (B). Solution B was added dropwise to the solution A till a uniform homogeneous mixture was obtained (pH = 6.3). At this stage, the aqueous solution of Na_2S (2 M, 0.7808 g) in 5 ml double distilled water was added dropwise to the precursor solution till the entire solution became white in color (pH = 5.4). The resultant solution was refluxed for an hour. A yolk-colored solution was observed.

Reaction scheme



The resultant solution was subjected to centrifugation at 14,000 rpm, and the precipitate formed was washed several times with double distilled water (pH = 7.0) and acetone, respectively. The purified end product so formed was dried overnight by keeping in a vacuum desiccator. The dried sample was used for further characterization.

Instruments used for characterization

The X-ray diffraction patterns of synthesized ZnS QDs are recorded with Bruker D8 Advance X-ray diffractometer. The primary X-ray wavelength Cu-K α 1 is at 1.5405 Å with a 2.2-kW power source. The sample was recorded in a wide range of Bragg's angle from 5° to 90° to get the best profile for the sample. The high-resolution transmission electron micrographs are obtained from JEOL JEM-2100 Transmission Electron Microscope with a resolution of 1.9 Å to 1.4 Å. The images are captured with a 2.672 × 2.672 k high-resolution CCD camera at 200 kV acceleration voltage. The absorption spectrum (λ_{max}) is recorded with Ocean Optics DH-2000-BAL UV–visible spectrophotometer. The FTIR spectrum is recorded with PerkinElmer FTIR spectrophotometer (Spectrum 400) in the range of 400–4500 cm^{-1} . The zeta potential was measured with Anton Paar Litesizer 500.

Preparation of thin films

The variable concentration of purified and characterized nano-crystals of ZnS was redispersed in double distilled water followed by sonication for about 10 min. Variable volumes of sonicated ZnS colloidal solution were used for making nano-sized thin films using photoresist spin coater (Ducom, PR-6-M2) at different rpm over well-cleaned SiO_2 templates. The thin films were dried at 50 °C for 5 h. Dried thin films were subjected to analysis regarding their thickness and surface structure using film thickness measurement

instrument (Filmetrics F20, thickness range 15 nm–70 μm ; visible wavelength range from 380 to 1050 nm) and atomic force microscope (AFM) (Nanosurf Easyscan 2 in dynamic mode), respectively. To analyze the surface roughness, 70- μm scan head was used in AFM to record the inference in non-contact tapping mode using Tap190Al-G probe.

Scheme for the analysis of biocompatibility and antibacterial activity

Biocompatibility of synthesized ZnS QDs was evaluated through hemolytic activity [43] against human erythrocytes and MTT assay [44] [(3-[4, 5-assay dimethylthiazole-2-yl]-2, 5-diphenyl tetrazolium bromide), a yellow tetrazole against L929 mouse fibroblast cell lines. The antibacterial activity of ZnS QDs was performed in terms of zone of inhibition, minimum inhibitory concentration, minimum bactericidal concentration and killing kinetic assay [45, 46]. The details of materials used and the description of all the assays are reported in Supplementary information.

Results and discussion

Mechanism involved in the synthesis of ZnS QDs

It has been observed that most of the green chemistry-synthesized nanoparticle reactions are carried out in the aqueous medium. In the present study, the suitability of the reaction in the water is probably because of the reaction between Zn^{+2} (borderline soft acid) and S^{-2} (soft base). According to the *hard and soft acid and base (HSAB)* theory, the reactions carried between a soft acid and a soft base give rise to a product having low-solubility product in the aqueous medium [2]. This helps in the separation of the required pure compounds easily from the solvent. The by-products formed during the reaction can be easily removed from the medium. Addition of D-glucose (capping agent) was done to restrain the size of the end product. The density functional theory (DFT) calculation (Supplementary section) showed that the attachment of the ZnS molecule is with the oxygen atom bonded to the fourth carbon of D-glucose molecule. The conformation manages to have the least energy associated with it ($-722,197.6382$ kcal/mol). The attachment of the ZnS molecule to the oxygen atom bonded to the third carbon may also be possible due to comparable bond energy associated with it (less by 0.2181 kcal/mol). The plausible reaction mechanism is reported in Supplementary section. Prepared nano-material was subjected to various microscopic and spectroscopic characterization techniques.

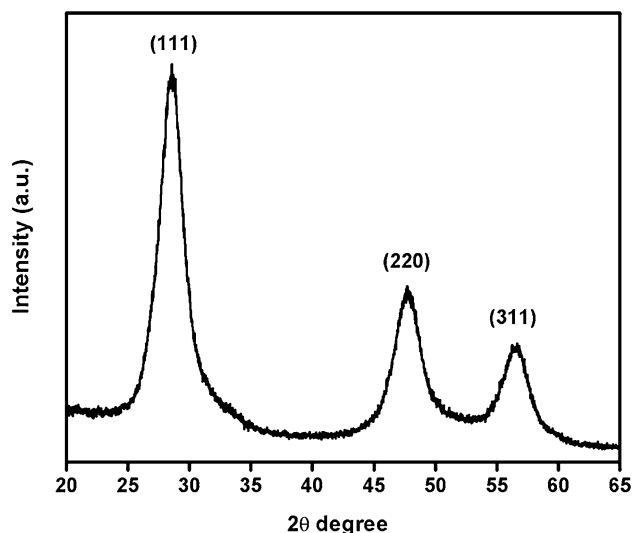


Fig. 1 Powder XRD pattern of ZnS nano-crystals revealing the corresponding Miller indices of diffraction planes

Powder X-ray diffraction analysis

This powerful technique gives accurate information about the zinc blende crystal structure of ZnS nanoparticles (Fig. 1). The broader FWHM of the peaks informs us about the generation of nanoparticles. The most intense peak was found along the plane (111), with the d-spacing 3.1 Å, indicating the formation of the majority of the nano-crystallites along this axis. Other prominent peaks were found for the planes (220) and (311) at the d-spacing 1.9 Å and 1.6 Å, respectively.

The X-ray analysis reveals the identity of the cubic crystallites of ZnS quantum dots having FCC structure. The unit-cell edge parameter ‘ a ’ and the volume of each unit cell were calculated to be 539.96 pm and 1.57×10^{-22} cm³, respectively. Broadening of the XRD peaks indicates the smaller particle size of ZnS nanoparticles. The Scherrer formula (Eq. 1) reveals that β (FWHM) is inversely proportional to τ (crystallite size). The actual broadening due to crystallite size, β_a , and strain are calculated after instrumental broadening, β_i , is deducted from the overall, observed broadening, β_o , considering Cauchy (Lorentzian) [47] (Eq. A) profile as follows:

$$\beta_a = \beta_o - \beta_i \quad (\text{A})$$

$$\tau = \kappa \lambda / \beta \cos \theta \quad (1)$$

The average grain (τ) size, 4.09 nm, was calculated using Eq. (1). The lattice strain of the nanoparticles was calculated using Williamson and Hall model (Eq. 2).

$$\beta_{hkl} \cos \theta = \kappa \lambda / L + 4\eta \sin \theta \quad (2)$$



where β_{hkl} denotes line broadening at the half of the intensity, θ Bragg angle (in degree), κ shape factor, λ X-ray wavelength, L mean size of the particle, and η lattice strain

The average value of lattice strain (η) along all the 3 prominent planes calculated is 0.0258. The highest strain of 0.0381 was found along the most intense peak in (111) plane. The increase in lattice strain is also attributed to the increased surface energy when the size of the particle is very small (QDs).

The specific surface area (S) was also calculated with the help of Brunauer, Emmett and Teller (BET) method [48] (Eq. 3) from the powder XRD spectrum. Since the surface-to-volume ratio of the nanoparticle is higher, its surface activity will enhance immensely.

$$S_{\text{BET}} = 6000/D_p * \rho \quad (3)$$

where S_{BET} denotes specific surface area, D_p size of the particles, and ρ density of the material

To analyze the surface activity, the density of ZnS QDs (ρ) is calculated as 4.11 g/cm³. Calculation reveals the value of S_{BET} as 356.93 m²/g, and the higher value indicates its expected efficiency in terms of the extent of adsorption.

XRD data were further used for calculating the dislocation density (δ) of ZnS QDs per unit volume:

$$\delta = 15\beta \cos \theta / 4aD \quad (4)$$

where δ denotes dislocation density, β line broadening at the half of the intensity (FWHM—full width half maximum), θ Bragg angle (in degree), a lattice edge parameter, and D particle size.

The dislocation density for ZnS QDs along the highest intensity peak along (111) plane was calculated to be $6.40 \times 10^{14} \text{ m}^{-2}$. The existence of dislocation density may be attributed to lattice strain which is observed more along (111) plane. The smaller the particle size, the higher will be the possibility of dislocations as they have more tendencies to stabilize their higher surface energy. Dislocations are basically the topological defects, which may be screw and edge, and have been visualized through HRTEM for ZnS QDs.

High-resolution transmission electron microscope analyses

HRTEM micrographs allow us to achieve the shape and size of each and every individual nano-crystallite in the most precise way. The lattice alignments, as well as their directions and defects, can also be detected with the help of this tool. Nanoparticles of ZnS ranging from 3 to 6 nm in diameter with spherical shape (larger surface energy) were observed in the scale of 5 nm (Fig. 2a inset). A large number of clusters of ZnS QDs (homogeneously distributed) were also observed in the range of 50 nm (Fig. 3). The average

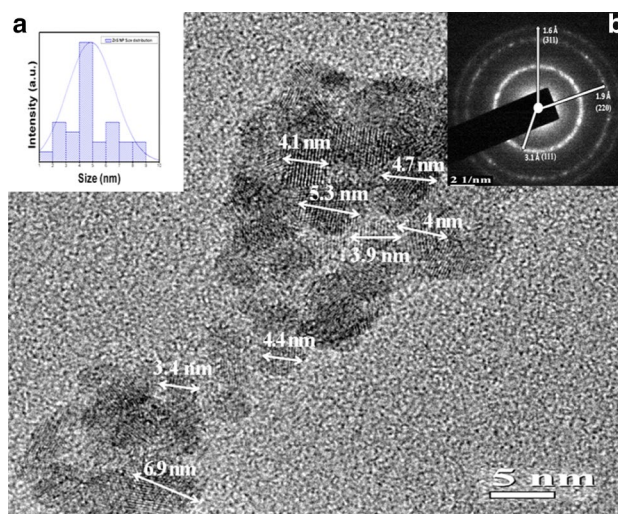


Fig. 2 HRTEM image of colloidal distribution of spherical-shaped ZnS QDs with **a** size distribution and **b** with SAED pattern

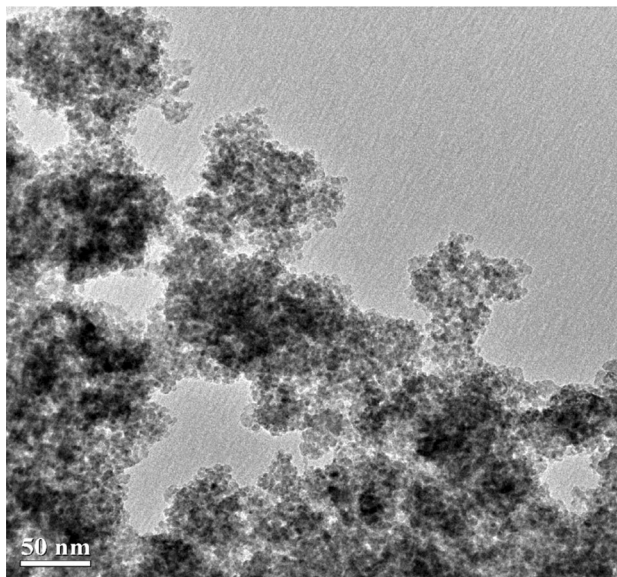


Fig. 3 TEM image exhibiting clustering of ZnS QDs on copper grid

particle size for around 20 nano-crystallites was recorded to be 4 nm, which is in good agreement with the results revealed by the XRD analysis also. SAED (selected area electron diffraction) pattern clearly reveals the polycrystallinity of the ZnS QDs with the d-spacing (Fig. 2b inset) of 3.1 Å, 1.9 Å and 1.6 Å, which is in confirmation with the XRD results. The imprinting polycrystallinity with its every bright spot, observed in the SAED pattern, confirms the presence of nano-crystallites feature of ZnS QDs. The zigzag alignment of atoms in the FCC ZnS nanoparticle reveals the presence of planar stacking fault along the (200) plane, which is evident from (Fig. 4). The FFT pattern of



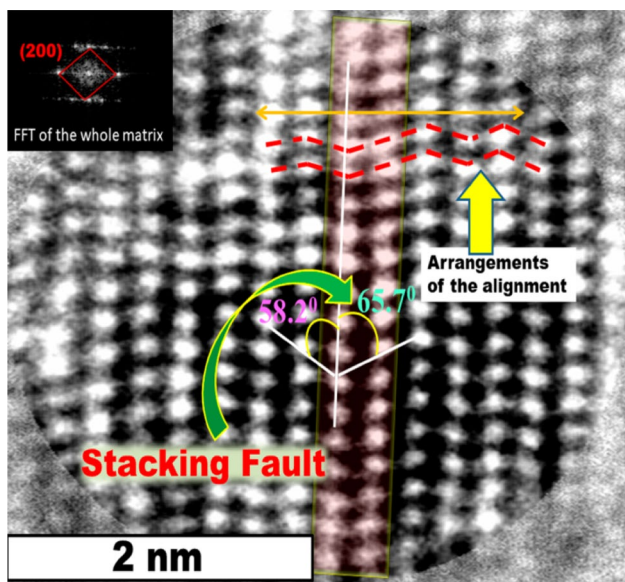


Fig. 4 Stacking fault (shaded area)

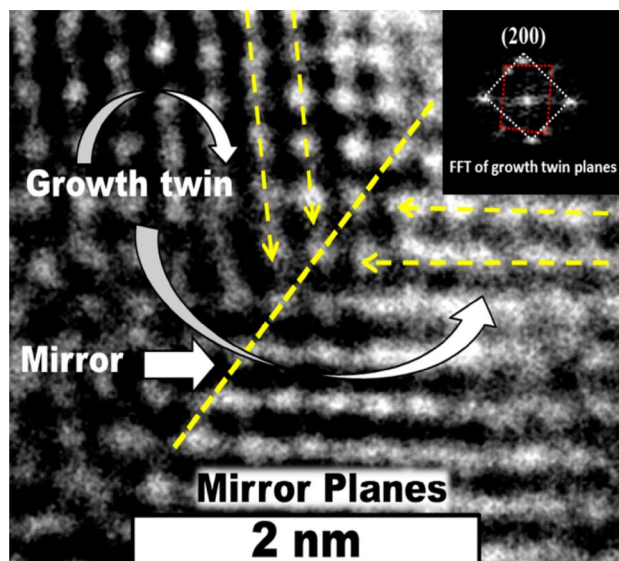


Fig. 6 Growth mirror twin planes

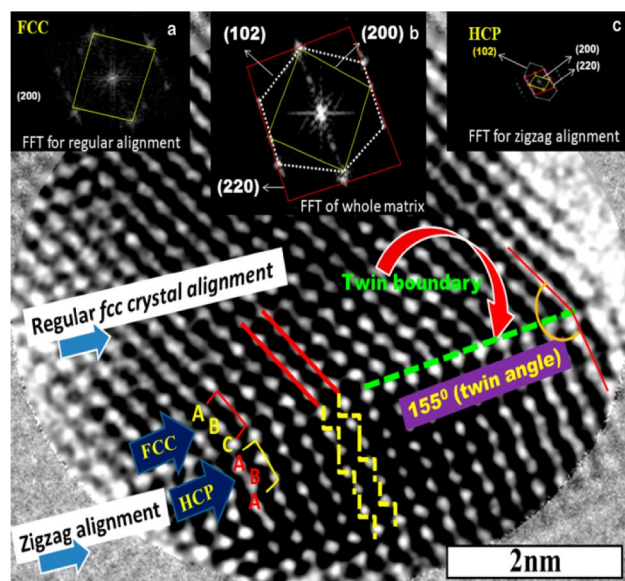


Fig. 5 Two differently aligned (*ccp* and *hcp*) planar conformations within the same grain

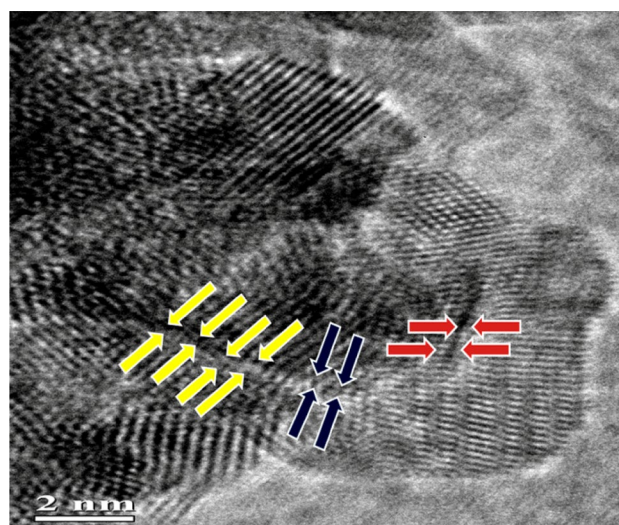


Fig. 7 Edge dislocation leading to indistinct grain boundary

(200) plane is reported in Fig. 4 inset. The formation of zigzag alignment may be attributed to the occurrence of two partial dislocations of atoms in the respective planes. Using the *ImageJ* software, the angles of atoms in each alignment in both sides of the stacking fault were found to be 58.2° and 65.7°. The presence of (102) plane (FFT of Fig. 5 inset) also speaks about the occurrence of the hexagonal plane ($d_{102} = 2.3 \text{ \AA}$). TEM micrographs also reveal the existence of twin boundary growth (Fig. 5) in ZnS nanoparticles. An angle of 155° was calculated between the two differently

aligned planes observed in the twin boundary area. The regular FCC (*ccp*) and zigzag (*hcp*) crystal alignments were observed in the area under investigation. The FFT patterns of the crystal alignments are reported in Fig. 5a, b, c inset.

The loss in contrast through (200) plane is probably due to the presence of stacking fault, which usually has been observed to occur during the synthesis of QDs [49]. From Fig. 6, it is observed that the twin growths tend to merge each other resulting in the formation of mirror images of each other. Such types of micro-twin growths can be

pointed out in quite a number of QD alignments of ZnS, where these localized defects try to form parallel planes at the junction of contact twin. Since the edge dislocation (Fig. 7) can be seen prominently in the alignments of ZnS nanoparticles, the grain boundary cannot be distinctly visualized.

Absorption spectra

The λ_{\max} for the ZnS QDs was found to be at 314 nm (blue shift). Unlike conspicuous absorption peaks which can be observed for bulk materials, the formation of QDs of ZnS is signified relatively by broader peaks (Fig. 8).

This is attributed to the generation of variation in the band gap for each nano-sized particle of ZnS. The optical band gap of the particles was found to be 3.95 eV. The higher value of the band gap of ZnS QDs by 0.37 eV than the bulk cubic ZnS confirms the formation of nanoparticles in the solution. The particle size was calculated with the Brus Eq. (5) and found to be 5.8 nm, which is in agreement with the absorption spectrum:

$$E_{g(\text{QD})} = E_{\text{bulk}} + h^2/8R^2(1/m_e^* + 1/m_h^*) - 1.786e^2/4\pi\epsilon_0\epsilon_r R \quad (5)$$

where E_{gap} denotes band gap energy of the bulk ZnS, m_e^* effective mass of the excited electron [50], m_h^* effective mass of the electron-hole, r radius of the particle, h Planck's constant, e charge of electron, ϵ_0 vacuum permittivity, and ϵ_r dielectric constant of ZnS.

The value of the third quantity in Eq. (5) is often neglected for semiconductors due to the high value of dielectric constant.

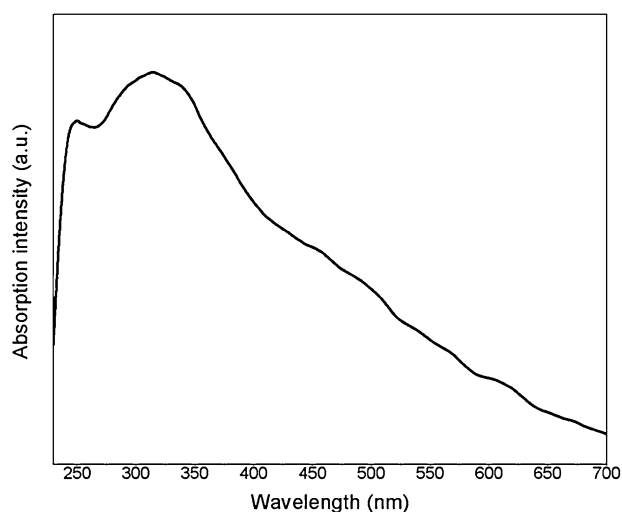


Fig. 8 UV-visible absorption spectra (314 nm) of colloidal ZnS QDs synthesized

Table 1 Comparative size of ZnS QDs calculated using different characterization techniques

Average size of ZnS nanoparticle			Optical band gap $E = hc/\lambda$	Blue shift ΔE
XRD	TEM	UV-visible		
4.09 nm	5 nm	5.76 nm	3.95 eV	0.37 eV

The comparison of the particle size of ZnS nanoparticles calculated by different techniques is reported in Table 1.

Zeta potential

The surface charge (zeta potential) of ZnS nanoparticles was found to be negative (-9.2 mV), attributed to the fairly stable configuration of synthesized QDs.

FTIR spectra

The FTIR spectrum (Fig. 9) shows a strong peak at 649 cm^{-1} , which is significant for Zn-S (metal sulfide) bond. The other strong peaks at 1635 cm^{-1} and 3411 cm^{-1} reveal the presence of H_2O bending and -OH stretching. The presence of a later peak confirms the presence of capping of D-glucose molecule. Another peak at 1010 cm^{-1} stands for C-O stretching.

Atomic force microscopy

From the AFM topologies, large aggregated nanoparticles are observed. The aggregation may be due to the particle coalescing (Fig. 10). The darker and brighter portions in Figs. 10, 11 correspond to the valleys and peaks

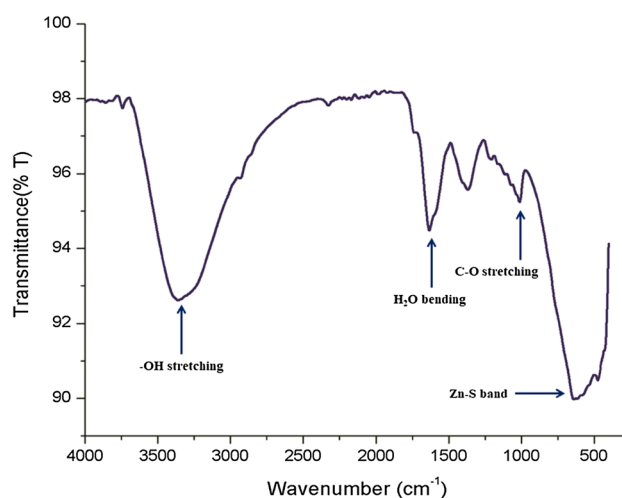


Fig. 9 FTIR spectra showing characteristic metal-sulfur bond as well as the interaction of ZnS QDs with D-glucose molecule as a capping agent

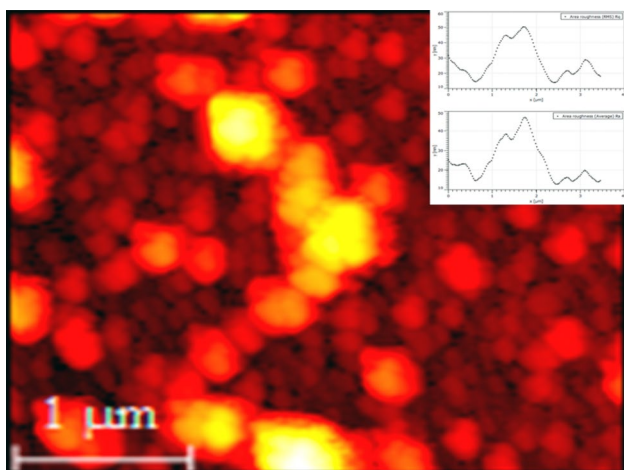


Fig. 10 Top view morphology of ZnS QDs thin film observed under AFM in dynamic mode

on the surface of the nano-sized thin film of ZnS QDs. The ratio of average roughness, root mean square (R_q) and average roughness (R_a) is found to be 1.33, which is in accordance with the height distribution predicted by Gaussian for most of the engineering surface (1.31) [51]. The height symmetry of the surface of thin films can be understood by skewness ($R_{sk} = 1.68$) (Table 2). The positive R_{sk} value indicates the presence of a bumpy surface. Since the value is not much higher than zero, it relates to a nearly symmetrical distribution of peaks and valleys about the mean line position. The peakedness parameter, kurtosis (R_{ku}) (Table 2), shows a value of 2.93 for the nano-sized thin film of ZnS QDs, which corresponds to the random distribution of peaks and valleys, known as mesokurtic behavior. The positive value of R_{sk} and R_{ku} reveals that the nano-sized ZnS thin films are suitable for tribological applications [52]. The maximum peak to valley height difference ($R_t = 210.78$ nm) has an important impact on the surface properties of thin films. It gives us the idea about the maximum roughness of the topology of the thin film of ZnS nanoparticles. *Gwyddion* free software was used to calculate the average thickness (50 ± 20 nm) using AFM data of thin films, which is in good agreement with the thickness (57 nm) measured with the help of *Filmetrics* (Supplementary information). Since QD thin films (QDTFs) can absorb and convert harmful UV lights to use visible light, they can be used

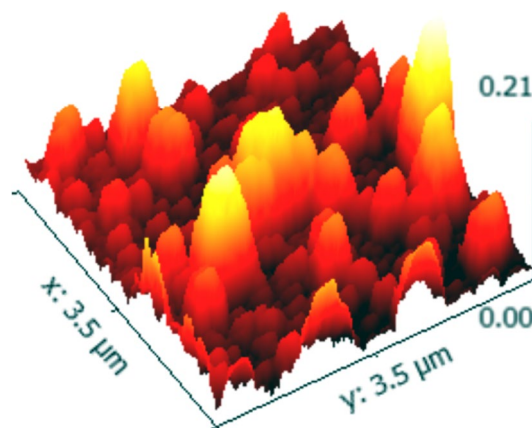


Fig. 11 3D view of Fig. 10 ZnS QDs thin film observed under AFM

for the future fabrication of QDTF solar cells. These cells can expand the photoresponse range and therefore can enhance the photoelectric conversion efficiently [53].

Biocompatibility analysis of green chemistry-synthesized ZnS nanoparticles (NPs)

Cytotoxicity against mammalian cell

The synthesized ZnS NPs demonstrated noticeable biocompatibility as is evident from cytotoxicity (MTT) assay results. The tested NPs (up to 200 μg/ml concentration) did not induce any significant cytotoxicity toward L929 mouse fibroblastic cells even after 72 h of post-treatment. The synthesized ZnS NPs exhibited time- and concentration-dependent decrease in cell viability which is in the range between 99.60 and 78.23%. This is also in agreement with the previous reports (Fig. 12) [54]. Even, with 200 μg/ml concentration and 72 h of incubation, the synthesized ZnS NPs did not induce any significant cytotoxicity to the tested mammalian cell. The higher percentage of mammalian cell viability demonstrated by the tested ZnS NPs may be attributed to repulsive interaction that mediated inefficient uptake of negatively charged ZnS NPs through negatively charged cell membrane [55]. Before further investigation of the synthesized ZnS NPs in preclinical and clinical settings, various biomedical applications and the high dose tolerance exhibited by the mammalian cell line for a prolonged duration (72 h or more hours) are very important.

Table 2 Characteristic roughness parameters of ZnS QDs thin film evaluated using AFM at $3.48 \times 3.48 \mu\text{m}^2$ of the scan area

Average particle size	Average roughness (R_a)	Average roughness (R_q) (rms)	R_q/R_a	Skewness R_{sk}	Kurtosis R_{ku}	The maximum peak to valley height difference (R_t)
56 ± 10 nm	24.97 nm	33.28 nm	1.33	1.68	2.93	210.78 nm

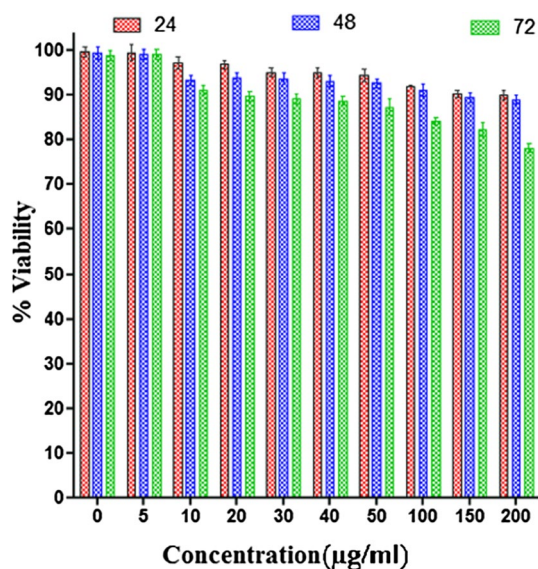


Fig. 12 Cytotoxicity evaluation (MTT) of ZnS NPs (The data are presented as mean \pm S.E. of five replicates)

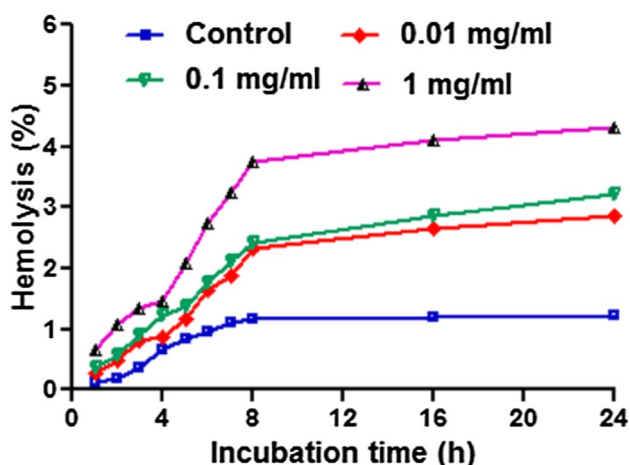


Fig. 13 Hemolysis of the RBCs incubated with different concentrations of the ZnS NPs at 37 °C temperature

Hemolytic activity evaluation

RBC constitutes 40–50% (v/v) of human whole blood and is responsible for the transport of carbon dioxide and oxygen. Hemolysis refers to the release of hemoglobin due to the disintegration of the three-layered RBC membrane, which is considered one of the most crucial parameters for evaluation of biocompatibility of biomaterials. The tested ZnS NPs in a concentration ranging from 0.01 to 1 mg/ml demonstrated mild to moderate hemolysis up to incubation period of 24 h in a dose- and time-dependent manner. For all the tested concentration of ZnS NPs, as compared to control,

the RBCs maintained its membrane integrity and did not lyse significantly up to even 4 h of time point, whereas from 8 h onward, all the tested concentrations of ZnS NPs started inducing, a mild to moderate level of lysis of erythrocytes, reaching 2.86%, 3.21% and 4.32% for 0.01 mg/ml, 0.1 mg/ml and 1 mg/ml concentrations, respectively (Fig. 13). The negative surface charge of the RBC repulsed the ZnS NPs and obstructed interaction, which may be attributed to the higher level of hemocompatibility of the tested ZnS NPs. The mild or moderate hemolytic activity exhibited by the ZnS NPs may be due to the hydrophobic interaction that mediated damage of RBC membrane and subsequent release of hemoglobin [56]. These preliminary data on hemocompatibility of the synthesized ZnS NPs will help to standardize, in future, the bio-safe concentration for its prospective biomedical applications.

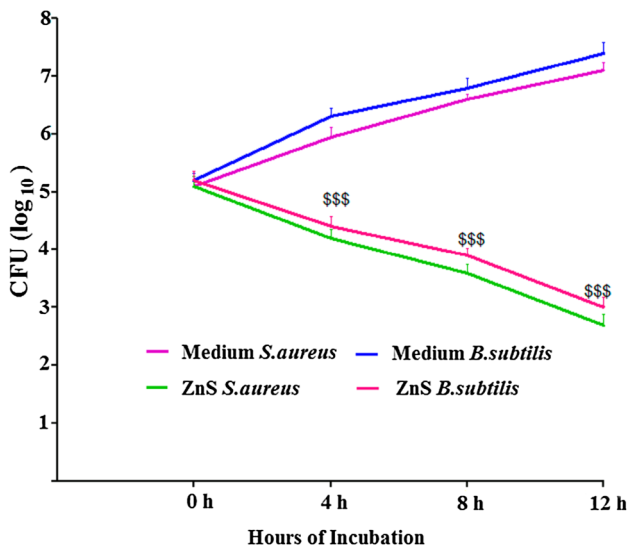
In vitro Antibacterial activity of ZnS nanoparticles (NPs)

To explore antibacterial effectiveness of the synthesized ZnS NPs, zone of inhibition (ZOI), minimum inhibitory concentration (MIC), minimum bactericidal concentration (MBC) and time-kill assay studies were performed against seven pathogenic bacterial strains. The tested ZnS NPs demonstrated noticeable zone of inhibition with a concentration-dependent increase in antibacterial activity against all the tested bacterial pathogens (Table 2). It was observed that in spite of possessing thicker peptidoglycan layer, gram-positive bacteria exhibited more susceptibility toward the tested ZnS NPs as compared to gram-negative bacteria; this may be due to the absence of an outer membrane. We conclude that the outer lipopolysaccharide membrane present in gram-negative bacteria offers a significant obstruction toward nanoparticle penetration or diffusion and consequently protects the cellular ultra-structures from ZnS NPs-mediated disturbances and subsequent dysfunction of the cell. *S. aureus* and *P. aeruginosa* were found to be most and least susceptible to the tested ZnS NPs. The pathogenic bacterial strains demonstrated variable MIC/MBC values ranging from 75 to 125 μ g/ml for gram-positive bacteria and 100 to 150 μ g/ml for gram-negative bacterial pathogens, respectively (Table 3). Since *S. aureus* and *B. subtilis* were found to be most susceptible bacterial strains, these were further evaluated for killing kinetics assay. The tested ZnS NPs demonstrated significant enhanced killing in terms of reduction in CFU count of *S. aureus* and *B. subtilis* in 4-, 8- and 12-h time points as compared to the non-treated strains (Fig. 14). All experiments were done in triplicate and results are expressed in mean, and standard deviations were negligible.

In order to exhibit antibacterial functions, NPs need to undergo interaction with the bacterial cell. The prominent

Table 3 Antibacterial activity profile of ZnS NPs in terms of zone of inhibition (ZOI), minimum inhibitory concentration (MIC) and minimum bactericidal concentration (MBC)

Sl. no	Bacterial strains	100 µg/ml	ZOI (cm) 200 µg/ml	300 µg/ml	MIC/MBC (µg/ml)
1.	<i>Bacillus cereus</i>	1.3	2.5	3.1	100/125
2.	<i>Bacillus subtilis</i>	1.6	2.3	2.8	75/125
3.	<i>Klebsiella pneumonia</i>	1.1	2.1	2.7	125/150
4.	<i>Escherichia coli</i>	1.9	2.6	3	100/150
5.	<i>Staphylococcus aureus</i>	2.1	2.9	3.4	75/100
6.	<i>Pseudomonas aeruginosa</i>	0.7	1	1.4	125/150
7.	<i>Staphylococcus epidermidis</i>	1.8	2.7	2.9	100/125

**Fig. 14** Time-kill assay of *S. aureus* (MTCC 3160) and *B. subtilis* (MTCC 441) were incubated with ZnS NPs. Surviving CFU at selected time points is shown. All the results were expressed in mean \pm S.D. ($n=3$). \$\$\$ $p < 0.001$, comparison of medium *S. aureus* and medium *B. subtilis* with ZnS QDs *S. aureus* and ZnS *B. subtilis* at different time intervals

accepted forms of interactions can be established by van der Waals forces [57], electrostatic forces [58], hydrophobic interactions [59] and ligand–receptor interactions [60]. While the NPs, capped with biocompatible cappers, are capable of forming such interactions with the bacterial cell membrane, they are able to cross the cell membrane and get into the metabolic pathways, thereby influencing the function and the shape of the cell membrane. The high antibacterial performance of the synthesized ZnS QDs is due to their very small size (~ 5 nm), i.e., within 10 nm range as observed in the present study. It is an established fact that the lesser the size (< 10 nm) of the nanoparticles, the better is their diffusion efficacy to penetrate through the pores of the cell membrane, and hence, they can easily go inside a microbial cell [61]. The synthesized ZnS QDs are expected to readily access the interior of the bacterial cell and can interact with the basic components of the cell (viz. DNA,

ribosome, lysosomes, etc.). These may lead to the generation of oxidative stress, protein deactivation, electrolyte balance disorder, enzyme inhibition and changes in cell membrane permeability inside the cell and finally cause death to the cell [62–64]. Conclusively, it can easily be understood that it is the advantage of the size (~ 5 nm) of the synthesized ZnS QDs (< 10 nm) that makes these nanoparticles (QDs) very promising as antibacterial agent. The rapid killing efficiency of the tested ZnS NPs is thus helpful in impeding biofilm formation at a faster pace. Findings of these experiments unveil the positive prospective attribute of the synthesized ZnS NPs utility in biomedical applications as anti-infective agents and hold tremendous potential in the application in superficial antibacterial therapeutics, implant coating, as well as in antibacterial textile material development with various advantageous functionalities.

Conclusion

We have presented a simple, time economic, cost-effective, reproducible and therefore successful method of synthesizing ZnS QDs. The method is an environmentally benign process, fulfilling the principles of *green chemistry*. The quantum dot thin films (QDTF) with its specified surface morphologies can absorb and transform harmful UV light to usable visible light efficiently, making them suitable for solar cell applications. The synthesized 2–6-nm nanoparticles were specially found to be biocompatible and also proficient antibacterial agents over a wide range of antibacterial studies. The synthesized QDs did not exert any adverse effect on mammalian cells.

Acknowledgements The research scholar is grateful to University Grants Commission, New Delhi, India, for providing the financial support. The authors are thankful to Centre for Advanced Studies in Chemistry, NEHU, Shillong.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

- Li, H., Shih, W.Y., Shih, W.-H.: Synthesis and characterization of aqueous carboxyl-capped CdS quantum dots for bioapplications. *Ind. Eng. Chem. Res.* **46**, 2013–2019 (2007). <https://doi.org/10.1021/ie060963s>
- Jing, L., Kershaw, S.V., Li, Y., et al.: Aqueous based semiconductor nanocrystals. *Chem. Rev.* **116**, 10623–10730 (2016). <https://doi.org/10.1021/acs.chemrev.6b00041>
- Dahl, J.A., Maddux, B.L.S., Hutchison, J.E.: Toward greener nanosynthesis. *Chem. Rev.* **107**, 2228–2269 (2007). <https://doi.org/10.1021/cr050943k>
- Lu, Y.B., Li, L., Su, S.C., et al.: A novel TiO₂ nanostructure as photoanode for highly efficient CdSe quantum dot-sensitized solar cells. *RSC Adv.* **7**, 9795–9802 (2017). <https://doi.org/10.1039/c6ra26029b>
- Wang, J., Li, Y., Shen, Q., et al.: Mn doped quantum dot sensitized solar cells with power conversion efficiency exceeding 9%. *J. Mater. Chem. A* **4**, 877–886 (2016). <https://doi.org/10.1039/c5ta09306f>
- Nozik, A.J.: Exciton multiplication and relaxation dynamics in quantum dots: applications to ultrahigh-efficiency solar photon conversion. *Inorg. Chem.* **44**, 6893–6899 (2005). <https://doi.org/10.1021/ic0508425>
- Rühle, S., Shalom, M., Zaban, A.: Quantum-dot-sensitized solar cells. *Chem Phys Chem* **11**, 2290–2304 (2010). <https://doi.org/10.1002/cphc.201000069>
- Salant, A., Shalom, M., Hod, I., et al.: quantum dot sensitized solar cells with improved efficiency prepared using electrophoretic deposition. *ACS Nano* **4**, 5962–5968 (2010). <https://doi.org/10.1021/nn1018208>
- Beard, M.C.: Multiple exciton generation in semiconductor quantum dots. *J. Phys. Chem. Lett.* **2**, 1282–1288 (2011). <https://doi.org/10.1021/jz200166y>
- Hanna, M.C., Nozik, A.J.: Solar conversion efficiency of photovoltaic and photoelectrolysis cells with carrier multiplication absorbers. *J. Appl. Phys.* **100**, 074510 (2006). <https://doi.org/10.1063/1.2356795>
- Semonin, O.E., Luther, J.M., Choi, S., et al.: Peak external photocurrent quantum efficiency exceeding 100% via MEG in a quantum dot solar cell. *Science* **334**, 1530–1533 (2011). <https://doi.org/10.1126/science.1209845>
- Kamat, P.V., Tvrđy, K., Baker, D.R., Radich, J.G.: Beyond photovoltaics: semiconductor nanoarchitectures for liquid-junction solar cells. *Chem. Rev.* **110**, 6664–6688 (2010). <https://doi.org/10.1021/cr100243p>
- Nozik, A.: Quantum dot solar cells. *Physica E* **14**, 115–120 (2002). [https://doi.org/10.1016/s1386-9477\(02\)00374-0](https://doi.org/10.1016/s1386-9477(02)00374-0)
- Kamat, P.V.: Quantum dot solar cells. the next big thing in photovoltaics. *J. Phys. Chem. Lett.* **4**, 908–918 (2013). <https://doi.org/10.1021/jz400052e>
- Yu, W.W., Qu, L., Guo, W., Peng, X.: Experimental determination of the extinction coefficient of CdTe, CdSe, and CdS nanocrystals. *Chem. Mater.* **15**, 2854–2860 (2003). <https://doi.org/10.1021/cm034081k>
- Wang, J., Li, Y., Shen, Q., et al.: Mn doped quantum dot sensitized solar cells with power conversion efficiency exceeding 9%. *J. Mater. Chem. A* **4**, 877–886 (2016). <https://doi.org/10.1039/c5ta09306f>
- Kershaw, S.V., Jing, L., Huang, X., et al.: Materials aspects of semiconductor nanocrystals for optoelectronic applications. *Mater Horizons* **4**, 155–205 (2017). <https://doi.org/10.1039/c6mh00469e>
- Kamat, P.V.: Meeting the clean energy demand: nanostructure architectures for solar energy conversion. *J. Phys. Chem. C* **111**, 2834–2860 (2007). <https://doi.org/10.1021/jp066952u>
- Michalet, X.: Quantum dots for live cells, in vivo imaging, and diagnostics. *Science* **307**, 538–544 (2005). <https://doi.org/10.1126/science.1104274>
- Talapin, D.V., Lee, J.-S., Kovalenko, M.V., Shevchenko, E.V.: Prospects of colloidal nanocrystals for electronic and optoelectronic applications. *Chem. Rev.* **110**, 389–458 (2010). <https://doi.org/10.1021/cr900137k>
- Kamat, P.V.: Photochemistry on nonreactive and reactive (semiconductor) surfaces. *Chem. Rev.* **93**, 267–300 (1993). <https://doi.org/10.1021/cr00017a013>
- Thompson, T.L., Yates, J.T.: TiO₂-based photocatalysis: surface defects, oxygen and charge transfer. *Top. Catal.* **35**, 197–210 (2005). <https://doi.org/10.1007/s11244-005-3825-1>
- Harris, C., Kamat, P.V.: Photocatalysis with cdse nanoparticles in confined media: mapping charge transfer events in the subpicosecond to second timescales. *ACS Nano* **3**, 682–690 (2009). <https://doi.org/10.1021/nn800848y>
- Tachikawa, T., Fujitsuka, M., Majima, T.: Mechanistic insight into the TiO₂ photocatalytic reactions: design of new photocatalysts. *J. Phys. Chem. C* **111**, 5259–5275 (2007). <https://doi.org/10.1021/jp069005u>
- <http://www.google.com/patents/US7916065>. Accessed 29 Mar 2011
- NASA <https://www.nasa.gov/feature/goddard/2017/nasa-and-mit-collaborate-to-develop-space-based-quantum-dot-spectrometer>. Accessed 14 Feb 2017
- Wegner, K.D., Hildebrandt, N.: Quantum dots: bright and versatile in vitro and in vivo fluorescence imaging biosensors. *Chem. Soc. Rev.* **44**, 4792–4834 (2015). <https://doi.org/10.1039/c4cs00532e>
- Xing, Y., Chaudry, Q., Shen, C., et al.: Bioconjugated quantum dots for multiplexed and quantitative immunohistochemistry. *Nat. Protoc.* **2**, 1152–1165 (2007). <https://doi.org/10.1038/nprot.2007.107>
- Yezhelyev, M.E.V., Al-Hajj, A., Morris, C., et al.: In situ molecular profiling of breast cancer biomarkers with multicolor quantum dots. *Adv. Mater.* **19**, 3146–3151 (2007). <https://doi.org/10.1002/adma.200701983>
- Smith, A.M., Dave, S., Nie, S., et al.: Multicolor quantum dots for molecular diagnostics of cancer. *Exp. Rev. Mol. Diagn.* **6**, 231–244 (2006). <https://doi.org/10.1586/14737159.6.2.231>
- Elward, J.M., Chakraborty, A.: Effect of dot size on exciton binding energy and electron-hole recombination probability in cdse quantum dots. *J. Chem. Theory Comput.* **9**, 4351–4359 (2013). <https://doi.org/10.1021/ct400485s>
- Franceschetti, A., Zunger, A.: Direct pseudopotential calculation of exciton coulomb and exchange energies in semiconductor quantum dots. *Phys. Rev. Lett.* **78**, 915–918 (1997). <https://doi.org/10.1103/physrevlett.78.915>
- Kasap, S.O., Capper, P.: Springer Handbook of Electronic and Photonic Materials. Springer, Boston (2007)
- Alivisatos, A.P.: Semiconductor clusters, nanocrystals, and quantum dots. *Science* **271**, 933–937 (1996). <https://doi.org/10.1126/science.271.5251.933>
- Nozik, A.J., Beard, M.C., Luther, J.M., et al.: Semiconductor quantum dots and quantum dot arrays and applications of multiple exciton generation to third-generation photovoltaic solar cells.

- Chem. Rev. **110**, 6873–6890 (2010). <https://doi.org/10.1021/cr900289f>
36. Form EIA-63B, Annual Photovoltaic Module/Cell Manufacturers Survey, Energy Information Administration, USA, 2006
 37. Ginley, D., Green, M.A., Collins, R.: Solar Energy conversion toward 1 terawatt. MRS Bull. **33**, 355–364 (2008). <https://doi.org/10.1557/mrs2008.71>
 38. King, R.R., Law, D.C., Edmondson, K.M., et al.: 40% efficient metamorphic GaInP/GaInAs/Ge multijunction solar cells. Appl. Phys. Lett. **90**, 183516 (2007). <https://doi.org/10.1063/1.2734507>
 39. Dimroth, F., Kurtz, S.: High-efficiency multijunction solar cells. MRS Bull. **32**, 230–235 (2007). <https://doi.org/10.1557/mrs2007.27>
 40. Luque, A., Martí, A., Nozik, A.J.: Solar cells based on quantum dots: multiple exciton generation and intermediate bands. MRS Bull. **32**, 236–241 (2007). <https://doi.org/10.1557/mrs2007.28>
 41. Green, M.A.: Third Generation Photovoltaics: Advanced Solar Energy Conversion. Springer, Berlin (2006)
 42. Martí, A.: Next Generation Photovoltaics: High Efficiency Through Full Spectrum Utilization. Institute of Physics Publishing, Bristol (2004)
 43. Kalita, S., Kandimalla, R., Devi, B., et al.: Dual delivery of chloramphenicol and essential oil by poly- ϵ -caprolactone–Pluronic nanocapsules to treat MRSA-Candida co-infected chronic burn wounds. RSC Adv. **7**, 1749–1758 (2017). <https://doi.org/10.1039/c6ra26561h>
 44. Kandimalla, R., Kalita, S., Choudhury, B., et al.: Fiber from ramie plant (*Boehmeria nivea*): a novel suture biomaterial. Mater. Sci. Eng. C **62**, 816–822 (2016). <https://doi.org/10.1016/j.msec.2016.02.040>
 45. Kotoky, J., Kandimalla, R., Kalita, S., et al.: Chloramphenicol encapsulated in poly- ϵ -caprolactone–pluronic composite: nanoparticles for treatment of MRSA-infected burn wounds. Int. J. Nanomed. (2015). <https://doi.org/10.2147/ijn.s75023>
 46. Kalita, S., Kandimalla, R., Sharma, K.K., et al.: Amoxicillin functionalized gold nanoparticles reverts MRSA resistance. Mater. Sci. Eng. C **61**, 720–727 (2016). <https://doi.org/10.1016/j.msec.2015.12.078>
 47. General Area Detector Diffraction System (GADDS) User Manual (2005). Bruker Advanced X-Ray Solutions. https://depts.washington.edu/moleng/wordpress/wp-content/uploads/2015/03/GADDS_Manual.pdf. Accessed Jan 2005
 48. Brunauer, S., Emmett, P.H., Teller, E.: Adsorption of gases in multimolecular layers. J. Am. Chem. Soc. **60**, 309–319 (1938). <https://doi.org/10.1021/ja01269a023>
 49. Murray, C.B., Norris, D.J., Bawendi, M.G.: Synthesis and characterization of nearly monodisperse CdE (E = sulfur, selenium, tellurium) semiconductor nanocrystallites. J. Am. Chem. Soc. **115**, 8706–8715 (1993). <https://doi.org/10.1021/ja00072a025>
 50. Manzoor, K., Johny, S., Thomas, D., et al.: Bio-conjugated luminescent quantum dots of doped ZnS: a cyto-friendly system for targeted cancer imaging. Nanotechnology **20**, 065102 (2009). <https://doi.org/10.1088/0957-4484/20/6/065102>
 51. Thomas, T.R.: Rough Surfaces. Imperial College Press, London (1999)
 52. Kumar, B.R., Rao, T.S.: AFM Studies on surface morphology, topography and texture of nanostructured zinc aluminum oxide thin films. Dig. J. Nanomater. Biostruct. **7**, 1881–1889 (2012)
 53. Chen, Y., Li, S., Huang, L., Pan, D.: Single-step direct fabrication of luminescent Cu-doped Zn_xCd_{1-x}S quantum dot thin films via a molecular precursor solution approach and their application in luminescent, transparent, and conductive thin films. Nanoscale **6**, 9640–9645 (2014). <https://doi.org/10.1039/c4nr02237h>
 54. Jayasree, A., Sasidharan, S., Koyakutty, M., et al.: Mannosylated chitosan-zinc sulphide nanocrystals as fluorescent bioprobes for targeted cancer imaging. Carbohydr. Polym. **85**, 37–43 (2011). <https://doi.org/10.1016/j.carbpol.2011.01.034>
 55. Chen, L., Mccrate, J.M., Lee, J.C.-M., Li, H.: The role of surface charge on the uptake and biocompatibility of hydroxyapatite nanoparticles with osteoblast cells. Nanotechnology **22**, 105708 (2011). <https://doi.org/10.1088/0957-4484/22/10/105708>
 56. Li, S., Guo, Z., Zhang, Y., et al.: Blood compatibility evaluations of fluorescent carbon dots. ACS Appl. Mater. Interfaces. **7**, 19153–19162 (2015). <https://doi.org/10.1021/acsami.5b04866>
 57. Armentano, I., Arciola, C.R., Fortunati, E., Ferrari, D., Mattioli, S., Amoroso, C.F., Rizzo, J., Kenny, J.M., Imbriani, M., Visai, L.: The interaction of bacteria with engineered nanostructured polymeric materials: a review. Sci. World J. **2014**, 1–18 (2014). <https://doi.org/10.1155/2014/410423>
 58. Li, H., Chen, Q., Zhao, J., Urmila, K.: Enhancing the antimicrobial activity of natural extraction using the synthetic ultrasmall metal nanoparticles. Sci. Rep. (2015). <https://doi.org/10.1038/srep11033>
 59. Luan, B., Huynh, T., Zhou, R.: Complete wetting of graphene by biological lipids. Nanoscale. **8**, 5750–5754 (2016). <https://doi.org/10.1039/C6NR00202A>
 60. Gao, W., Thamphiwatana, S., Angsantikul, P., Zhang, L.: Nanoparticle approaches against bacterial infections. Wiley Interdiscip. Rev.: Nanomed. Nanobiotechnol. **6**, 532–547 (2014). <https://doi.org/10.1002/wnan.1282>
 61. Mukha, I.P., Eremenko, A.M., Smirnova, N.P., Mikhienkova, A.I., Korzhak, G.I., Gorchev, V.F., Chunikhin, A.Y.: Antimicrobial activity of stable silver nanoparticles of a certain size. Appl. Biochem. Microbiol. **49**, 199–206 (2013). <https://doi.org/10.1134/S0003683813020117>
 62. Xu, Y., Wei, M.-T., Ou-Yang, H.D., Walker, S.G., Wang, H.Z., Gordon, C.R., Guterman, S., Zawacki, E., Applebaum, E., Brink, P.R., Rafailovich, M., Mironava, T.: Exposure to TiO₂ nanoparticles increases Staphylococcus aureus infection of HeLa cells. J. Nanobiotechnol. (2016). <https://doi.org/10.1186/s12951-016-0184-y>
 63. Shrivastava, S., Bera, T., Roy, A., Singh, G., Ramachandrarao, P., Dash, D.: Characterization of enhanced antibacterial effects of novel silver nanoparticles. Nanotechnology. **18**, 225103 (2007). <https://doi.org/10.1088/0957-4484/18/22/225103>
 64. Yang, W., Shen, C., Ji, Q., An, H., Wang, J., Liu, Q., Zhang, Z.: Food storage material silver nanoparticles interfere with DNA replication fidelity and bind with DNA. Nanotechnology. **20**, 085102 (2009). <https://doi.org/10.1088/0957-4484/20/8/085102>

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

