

Cite this: *RSC Adv.*, 2019, 9, 1335

Received 24th November 2018

Accepted 28th December 2018

DOI: 10.1039/c8ra09660k

rsc.li/rsc-advances

Enhanced photoelectrochemical activity of Co-doped β - In_2S_3 nanoflakes as photoanodes for water splitting†

Supriya Pulipaka,^a A. K. S. Koushik,^a Melepurath Deepa ^b and Praveen Meduri ^{*a}

This work is primarily focused on indium sulfide (β - In_2S_3) and cobalt (Co)-doped β - In_2S_3 nanoflakes as photoanodes for water oxidation. The incorporation of cobalt introduces new dopant energy levels increasing visible light absorption and leading to improved photo-activity. In addition, cobalt ion centers in β - In_2S_3 act as potential catalytic sites to promote electro-activity. 5 mol% Co-doped β - In_2S_3 nanoflakes when tested for photoelectrochemical water splitting exhibited a photocurrent density of 0.69 mA cm^{-2} at 1.23 V, much higher than that of pure β - In_2S_3 .

Introduction

Photoelectrochemical (PEC) water splitting is attracting huge attention since solar energy is directly converted into hydrogen which can be used as fuel. A PEC semiconductor device in an aqueous electrolyte, irradiated with sunlight can directly split water into hydrogen and oxygen. The overall reaction in PEC is controlled by the oxygen evolution reaction (OER) at the anode due to its sluggish kinetics.¹ Fujishima and Honda first reported a PEC device by using titania (TiO_2) as a photoanode in 1972.² However, TiO_2 absorbs only 5% of the solar spectrum due to its large band gap of 3.2 eV and is primarily limited to the ultra-violet region. The requirement for visible light absorption drives the development of metal oxides with narrow band gap such as bismuth vanadate (BiVO_4),³ tungsten oxide (WO_3), and hematite (α - Fe_2O_3).^{4–6} Along with metal oxides, metal sulfides have garnered attention as promising materials due to their unique properties with good mechanical and thermal stability in electrochemical devices.⁷ Metal chalcogenides have received recent interest in photocatalysis owing to their narrow band gap and low-temperature synthesis.⁸ Among the metal chalcogenides, cadmium sulfide (CdS) with a band gap of 2.4 eV is a highly photoactive material under visible light. However, toxicity and photocorrosion make its use challenging for water oxidation.^{9–11} A recent study on multishelled hollow structures with CdS and zinc sulfide (ZnS) have shown enhanced PEC performance.¹² The other chalcogenide which has a favorable band gap of 2.0–2.3 eV is indium sulfide (β - In_2S_3). It is a n-type semiconductor

which is primarily used for solar cells with visible-light absorption.^{13–15} A variety of 2D β - In_2S_3 morphologies such as nanobelts and nanoflakes have been synthesized using metal ions such as calcium and aluminum to restrict the crystal growth in 2D wherein the nanoflakes have shown the best photoelectrochemical activity.¹⁶ A biomaterial template with amino acids has been used to synthesize high surface area β - In_2S_3 nanoflakes with increased number of surface active sites which are shown to promote the photocatalytic process of alcohol to aldehyde selective oxidation.¹⁷ However, the use of β - In_2S_3 nanoflakes for PEC water splitting to generate H_2 or O_2 has been sparsely reported despite good photocatalytic properties of β - In_2S_3 .^{18,19} Previous studies have strived for the improvement of β - In_2S_3 photocatalytic properties by the way of heterojunctions with TiO_2 , MoS_2 , CdS and CuS.^{20–22} Alongside, noble metal nanoparticles such as silver have also been used to improve efficiency but at the expense of increased cell costs.²³ The electro and photoactivity of β - In_2S_3 can be improved by doping with metals like zirconium,²⁴ cobalt,²⁵ tin^{26,27} and vanadium.²⁸ The present work explores the introduction of a transition metal, cobalt (Co) into β - In_2S_3 nanoflakes by chemical bath deposition (CBD) at relatively low temperatures. The doping of Co has been confirmed by optical studies. Co-doped β - In_2S_3 showed an improved PEC activity due to an increase in the photo and electroactivity. To the best of our knowledge, there is only one report on Co-doped β - In_2S_3 which is an atomic layer confined doping in three stacked layers and used as photoanode for PEC water oxidation. They reported a current density of 1.17 mA cm^{-2} at an applied potential of 1.5 V (vs. RHE) in the basic electrolyte.²⁵ Mesoporous TiO_2 offers a robust support for the deposition of sensitizers in solar energy conversion applications. In particular, metal sulfides can be easily deposited onto TiO_2 using a chemical bath because of good adhesion properties. The electronic communication between the sensitizer and TiO_2 is also high due to the strong electron affinity of TiO_2 .²⁹

^aDepartment of Chemical Engineering, Indian Institute of Technology Hyderabad, Kandi, Sangareddy-502285, Telangana, India. E-mail: meduripraveen@iith.ac.in

^bDepartment of Chemistry, Indian Institute of Technology Hyderabad, Kandi, Sangareddy-502285, Telangana, India

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ra09660k

The adhesion of β - In_2S_3 nanoparticles on to bare fluorine doped tin oxide (FTO) substrate is poor and does get peeled off during testing. TiO_2 coating on FTO leads to even deposition of β - In_2S_3 with improved adhesion. The photogenerated electrons of β - In_2S_3 can easily transfer to the conduction band of TiO_2 due to its proper band alignment, thus separating the photogenerated electron-hole pairs effectively. In the present work, all the doped and un-doped samples are deposited on TiO_2 layer.

Experimental

Chemicals

Indium chloride from Alfa Aesar; thioacetamide, ethyl cellulose and terpineol from Sigma Aldrich; titanium tetrachloride (TiCl_4) and cobalt acetate tetrahydrate from Merck are used without further purification. TiO_2 powder (P25) is obtained from Evonik. Deionized water with a resistivity of $18.2 \Omega \text{ cm}$ is obtained from a Millipore Direct-Q3 UV system. FTO substrates obtained from Pilkington are cleaned in soap solution, 30% HCl, twice distilled water, ethanol and acetone in that particular sequence prior to use.

Preparation of electrodes

Synthesis of TiO_2 layers. TiO_2 paste is prepared by mixing 100 mg of TiO_2 powder with 50 mg of ethyl cellulose and 1 mL of terpineol which is then applied over a FTO plate by doctor blading. TiO_2 film is further dried at 60°C for 30 min followed by annealing at 500°C for 30 min. An extra layer of TiO_2 is deposited using the same procedure as above for reducing the surface roughness of the deposited TiO_2 film. These samples are then immersed in an aqueous solution of 0.4 M TiCl_4 at 70°C for 20 min. The coated substrate is rinsed in ultrapure water to remove unadsorbed excess TiO_2 and further annealed at 500°C for 30 min.

Synthesis of $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ electrodes. $\text{InCl}_3 \cdot 4\text{H}_2\text{O}$ (0.22 g) and thioacetamide (0.11 g) are dissolved in distilled water (12.5 mL). Different mol% (2, 5 and 8 mol%) of cobalt precursor is added to synthesize Co-doped $\beta\text{-In}_2\text{S}_3$ electrodes. The TiO_2 coated FTOs are dipped in the precursor solution and kept at 80°C for 2 h under stirring conditions. The samples are washed with water to remove excess material and precursor solution on the surface; dried at 60°C for 12 h and annealed at 120°C for 1 h in vacuum. $\text{TiO}_2/\beta\text{-In}_2\text{S}_3$ electrodes are synthesized by following the same procedure without addition of cobalt precursor.

Fig. 1a depicts the fabrication of photoelectrodes. Fig. 1b shows the as-synthesized powders of $\beta\text{-In}_2\text{S}_3$ and Co-doped $\beta\text{-In}_2\text{S}_3$ nanoflakes with a difference in color indicating a change in the optical emission properties. The mol% that is referred to in the entire document represents added mol% in the solution. The actual wt% is calculated from energy dispersive spectroscopy (EDS) which is given in the Fig. S1 in ESI.†

Characterization

XRD graphs of the samples are obtained on a PANalytical X'PertPRO instrument with $\text{Cu-K}\alpha$ ($\lambda = 1.5406 \text{ \AA}$) radiation.

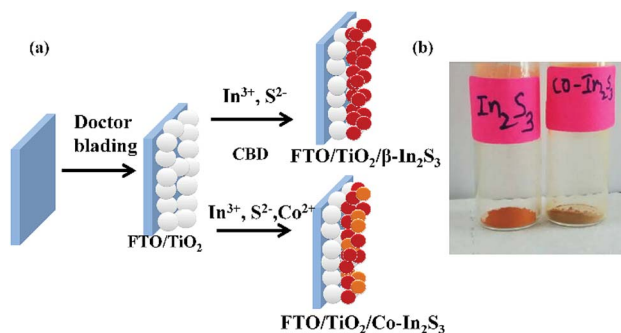


Fig. 1 (a) Schematic representation of the synthesis process using CBD and (b) color difference between as-synthesized $\beta\text{-In}_2\text{S}_3$ and Co-doped $\beta\text{-In}_2\text{S}_3$.

HRTEM (High Resolution Transmission Electron Microscopy) images of $\beta\text{-In}_2\text{S}_3$ and Co-doped $\beta\text{-In}_2\text{S}_3$ samples are obtained on a HRTEM JEOL JEM 2100 with a LaB_6 filament operating at 200 kV. The samples are prepared by dispersing the powder in ethanol followed by ultrasonication for 1 hour before drop casting onto a carbon-coated copper grid. The solvent is evaporated and further used as a specimen for TEM. The optical absorption spectra of the films are measured in the diffuse reflectance mode and converted to absorbance using Kubelka-Munk function on a UV-Vis-NIR spectrophotometer (Shimadzu UV-3600). Raman spectra of the samples are recorded on a Bruker Senterra Dispersive Raman Microscope spectrometer. The laser excitation wavelength is fixed at 532 nm. Photoluminescence (PL) spectra of films are measured on a Horiba Fluoromax-4 fluorescence spectrometer; a suitable filter is utilized during the measurement and background correction is applied. Surface morphology analysis is performed using a field emission scanning electron microscope (Carl Zeiss Supra 40 FE-SEM). The specific surface area is calculated from N_2 adsorption-desorption isotherms measured using a Micromeritics ASAP 2020 instrument using the Brunauer-Emmett-Teller (BET) method. Pore volume and pore-size distribution are calculated by using the Barrett-Joyner-Halenda (BJH) method.

Photoelectrochemical performances

Photoelectrochemical data is studied using Autolab PGSTAT 302N equipped with NOVA 1.11 software. The light source is a white LED (40 W) and the intensity of the light is adjusted to 1 sun by keeping the optimum distance between the light and the sample. Light intensities are measured with a LOT-Oriel radiometer-photometer (ILT: 1400). A three electrode cell with the prepared $\beta\text{-In}_2\text{S}_3$ (or) Co-doped $\beta\text{-In}_2\text{S}_3$ samples as working electrode, a platinum wire as the counter electrode and Ag/AgCl as the reference electrode is used. 0.5 M Na_2SO_4 (buffered to $\text{pH} = 7$) is used as an electrolyte for all the measurements. The measured Ag/AgCl reference electrode potentials are converted to reversible hydrogen electrode (RHE) by using Nernst equation $E_{\text{RHE}} = E_{\text{Ag}/\text{AgCl}} + 0.0591 \text{ pH} + 0.199 \text{ V}$. The photocurrent is measured by linear sweep voltammetry in the range of 0 to 1.6 V vs. RHE at a scan rate of 10 mV s^{-1} . The time dependent



currents are measured at a bias potential of 1.23 V vs. RHE. Prior to all the measurements argon gas is purged for 30 min to remove the dissolved oxygen in the solution.

Results and discussion

XRD patterns of as-synthesized samples are shown in Fig. 2a. Peaks for TiO_2 correspond to the body-centered tetragonal structure of TiO_2 (PDF: 894921). The diffraction patterns in the range of $10\text{--}90^\circ$ can be indexed as cubic phase $\beta\text{-In}_2\text{S}_3$ structure (PDF: 650459). The diffraction peak of $\beta\text{-In}_2\text{S}_3$ at $2\theta = 33.45^\circ$ can be prominently seen which is indexed to (400) plane of $\beta\text{-In}_2\text{S}_3$ and the other peaks overlap with TiO_2 . Fig. 2b shows the XRD peaks without TiO_2 . No distinct characteristic peaks are visible in XRD of the doped samples which rules out the formation of a new compound, potentially cobalt sulfide (CoS_2). The broadened peaks do indicate that the as-synthesized samples consist of nanomaterials *i.e.* nanoflakes in this case. The $\beta\text{-In}_2\text{S}_3$ crystal has a defect spinel lattice structure; there are twelve tetrahedral sites in $\beta\text{-In}_2\text{S}_3$ of which eight are occupied by In^{3+} ion whereas four are empty. These empty sites can be potentially occupied by other metals,³⁰ paving ways for doping of metals in $\beta\text{-In}_2\text{S}_3$ at synthesis temperatures of $\beta\text{-In}_2\text{S}_3$ *i.e.*, low temperatures. Cobalt-doped in our study is believed to occupy one or more empty tetrahedral sites of $\beta\text{-In}_2\text{S}_3$ which is confirmed by optical studies and discussed a little later. We observe a shift of 0.1° for the (400) plane in the doped sample and the crystallite size decreases with increasing Co^{2+} ion concentration in doped samples. The smaller size of Co^{2+} ion (70 pm) in comparison to the In^{3+} ion (80 pm) occupying the empty tetrahedral sites and binding to the other elements in the crystal causes a reduction in the overall crystallite size of the material. The average crystallite sizes of $\beta\text{-In}_2\text{S}_3$, Co-doped $\beta\text{-In}_2\text{S}_3$ with 2, 5 and 8 mol% are 9.22 nm, 9.2 nm, 8.08 nm, and 7.77 nm respectively. Fig. 2c

shows the UV-Vis-NIR diffuse reflectance spectra for both doped and undoped samples. The undoped $\beta\text{-In}_2\text{S}_3$ nanoflakes have an absorption peak at 650 nm which correlates to a band gap of 1.9 eV. The doped samples have three sub-bandgap absorption peaks in the range of 650–900 nm corresponding to new energy levels in addition to the absorption peak at 650 nm. The position of the $\beta\text{-In}_2\text{S}_3$ absorption peak in the visible region did not alter with the doping level. The three peaks seem to overlay against each other and appear as one broad peak.¹⁶ The broad peak is indicative of the tetrahedrally coordinated Co^{2+} ion transitions, ${}^2\text{E}(\text{G}) \rightarrow {}^4\text{A}_2(\text{F})$, ${}^4\text{T}_1(\text{P}) \rightarrow {}^4\text{A}_2(\text{F})$, ${}^2\text{A}_1(\text{G}) \rightarrow {}^4\text{A}_2(\text{F})$ respectively.²⁵ The twelve In^{3+} sites in the spinel $\beta\text{-In}_2\text{S}_3$ structure have tetrahedral coordination while there is no tetrahedral Co^{2+} co-ordination in CoS_2 (cubic pyrite structure). Hence, the broad peak above indicates that Co^{2+} ions are located at tetrahedral sites in $\beta\text{-In}_2\text{S}_3$ which is indicative of doping.^{31,32} The incorporation of Co^{2+} ion in the vacant sites does improve photoabsorption. Cobalt sites in $\beta\text{-In}_2\text{S}_3$ can act as trapping sites for the photogenerated electrons/holes which will prevent their recombination.

Raman spectra in Fig. 2d shows a characteristic peak at 305 cm^{-1} which pertains to $\text{A}_{1\text{g}}$ mode confirming the structure and phase of $\beta\text{-In}_2\text{S}_3$ nanoflakes.³³ This peak is ascribed to symmetric stretching of the vibrating mode in InS_4 tetrahedra. Fig. 2d shows a considerable shift from 305 cm^{-1} to 297.7 cm^{-1} when $\beta\text{-In}_2\text{S}_3$ is doped with cobalt. The incorporation of Co^{2+} ions in the tetrahedral sites and binding to other elements (sulfur) can lead to bond stretching of In–S which is seen as a blueshift in the observed spectra. Furthermore, only a slight shift from 297.7 cm^{-1} to 296.3 cm^{-1} is observed when the Co mol% is increased in the sample. It shows that once bond stretching happens, further addition of Co^{2+} does not alter the binding to sulfur which is indicative of Co^{2+} doping rather than formation of CoS_2 . In addition, it also indicates the saturation of Co^{2+} ions in the $\beta\text{-In}_2\text{S}_3$ structure beyond a certain concentration which is reflected in the electrochemical data.

SEM images in Fig. 3a and b show the agglomerated particles of $\beta\text{-In}_2\text{S}_3$ and Co-doped $\beta\text{-In}_2\text{S}_3$ deposited on TiO_2 respectively. These uniform microspheres have an average size of $1\text{--}1.5\text{ }\mu\text{m}$ in both the doped and undoped samples. These microspheres consist of bundles of nanoflakes as shown in TEM images. Fig. 3c and d show nanoflakes of 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ which are of $20\text{--}30\text{ nm}$ width and $40\text{--}50\text{ nm}$ length. These small nanoflakes tend to collate during stirring in the CBD process which leads to the formation of microspheres or nanoflowers.^{22,33}

In order to confirm the presence of Co doping in $\beta\text{-In}_2\text{S}_3$, Energy Dispersive Spectroscopy (EDS) analysis and elemental mapping studies have been carried out. Fig. 4a shows the SEM image of $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$. The cross-sectional view of the $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ film in Fig. 4b shows homogenous and well-integrated layers which lead to low recombination losses.³⁴ Owing to the porosity of TiO_2 layer, Co-doped $\beta\text{-In}_2\text{S}_3$ nanoparticles easily percolate through and coat the TiO_2 particles. Thus, the resulting $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ film appears as a single homogeneous entity and the thickness of the film is inferred to be nearly $8.10\text{ }\mu\text{m}$. Fig. 4c–e show the elemental

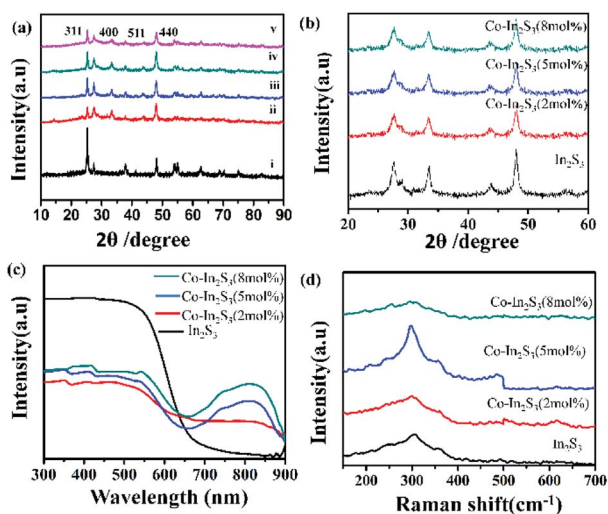


Fig. 2 (a) X-ray diffraction pattern of (i) TiO_2 (ii) $\text{TiO}_2/\beta\text{-In}_2\text{S}_3$ (iii) (iv) and (v) $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ with 2, 5, 8 mol% respectively (b) X-ray diffraction pattern without TiO_2 (c) UV-Vis-NIR diffuse reflectance spectra (d) Raman spectra.



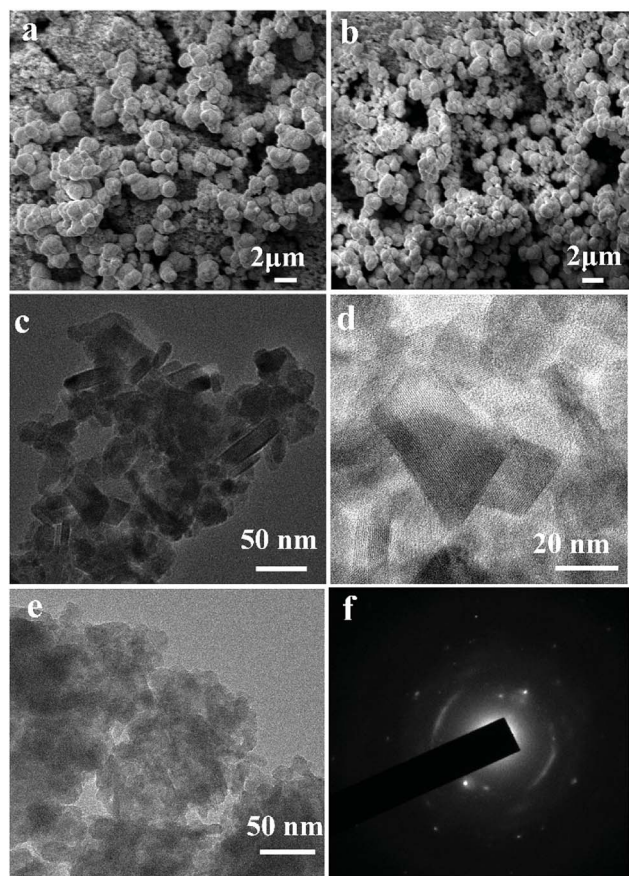


Fig. 3 SEM images of (a) $\text{TiO}_2/\beta\text{-In}_2\text{S}_3$ and (b) $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ (5 mol%). TEM images of (c) 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ with clearly visible nanoflakes (d) individual nanoflake of 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ (e) nanoflakes of 8 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ and (f) SAED pattern of 8 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ indicating crystallinity.

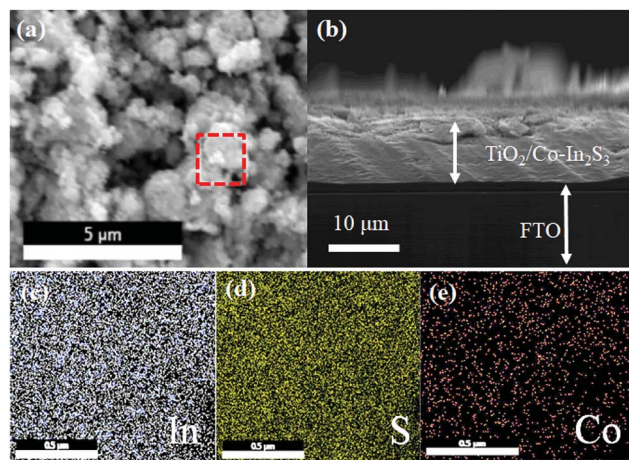


Fig. 4 SEM images of (a) $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ (b) cross sectional view of $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ film and (c–e) EDS elemental mapping of selected area of inset in (a).

mapping of 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$. Fig. 4e clearly demonstrates the uniform distribution of cobalt in the $\beta\text{-In}_2\text{S}_3$ crystal structure. Line scan of an individual Co-doped $\beta\text{-In}_2\text{S}_3$ particle

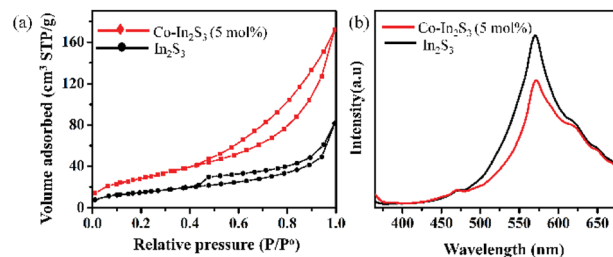


Fig. 5 Comparison of $\text{TiO}_2/\beta\text{-In}_2\text{S}_3$ and $\text{TiO}_2/\text{Co-doped } \beta\text{-In}_2\text{S}_3$ (5 mol%) properties using (a) nitrogen adsorption-desorption isotherms and (b) photoluminescence (PL) spectra.

also confirms the uniform distribution of cobalt in the particle which is shown in ESI (Fig. S1).†

The specific surface area and porosity are calculated by N_2 adsorption-desorption isotherms (Fig. 5b). Both the isotherms are identified as type IV with a hysteresis loop which is a characteristic of mesoporous materials. Mesopores primarily arise due to the spaces between the nanoflakes in the microspheres.³⁵ The BET surface area of the $\beta\text{-In}_2\text{S}_3$ and 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ are estimated to be $52.17 \text{ m}^2 \text{ g}^{-1}$ and $101.15 \text{ m}^2 \text{ g}^{-1}$, respectively. The doping of Co does increase specific surface area significantly. The values are reported in ESI (Table 1).† Generally, doping is believed to inhibit the particle growth and lead to an increase in the specific surface area. The as-prepared Co-doped $\beta\text{-In}_2\text{S}_3$ showed a higher specific surface area than most of the reported values.³⁶ High surface area can provide more active sites for the adsorption of reactants on the surface which improves interfacial charge transfer thus improving the photocatalytic properties.

Photoluminescence data presented here indicates the recombination intensity. Lower PL intensity indicates less recombination of photo-induced electron-hole pairs, and higher photocatalytic activity.^{27,37} Fig. 5b shows the PL spectra of both pure $\beta\text{-In}_2\text{S}_3$ and 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$. PL quenching is shown at an excitation wavelength of 350 nm. 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ displays significantly diminished intensity as compared to pure $\beta\text{-In}_2\text{S}_3$ which is indicative of the efficient charge separation of photogenerated electron-hole pairs and lower recombination. Improved efficiency is due to Co doping which delayed the recombination rate of photogenerated charge carriers of $\beta\text{-In}_2\text{S}_3$.

Fig. 6a shows the schematic of 3-electrode setup where the electron transfer mechanism is explained. The photogenerated electrons under irradiation move from $\beta\text{-In}_2\text{S}_3$ to TiO_2 due to their proper band alignment. The holes in $\beta\text{-In}_2\text{S}_3$ help in water oxidation. Photoanode studies are generally carried out in basic medium as oxygen evolution is higher with high hydroxide ion concentration. Even though the currents are high, extreme pH conditions will damage other parts of the cell with prolonged use leading to maintenance and safety issues in practical applications.³⁸ Therefore, we use neutral pH solution for all our testing purposes unless indicated otherwise. Fig. 6b shows linear sweep voltammograms (LSV) of the $\beta\text{-In}_2\text{S}_3$, Co-doped $\beta\text{-In}_2\text{S}_3$ electrodes in Argon gas purged 0.5 M Na_2SO_4 (pH = 7) under light and dark conditions. All the electrodes showed



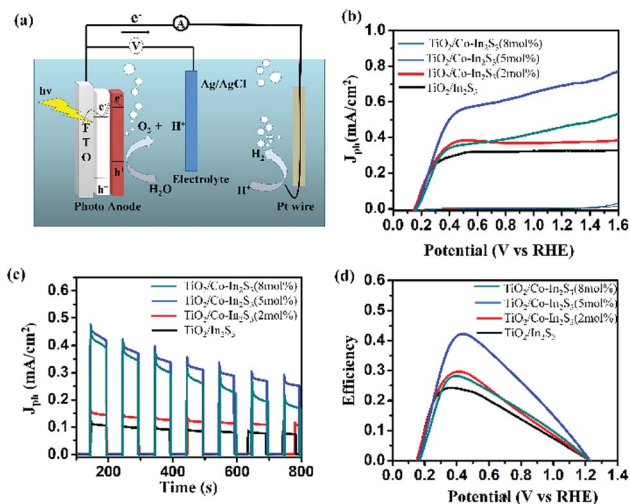


Fig. 6 Photoelectrochemical properties of $\text{TiO}_2/\beta\text{-In}_2\text{S}_3$, TiO_2/Co -doped $\beta\text{-In}_2\text{S}_3$ with 2, 5 and 8 mol%. (a) Three electrode measurement setup (b) linear sweep voltammograms in light and dark conditions (c) chrono-amperometry curves under chopped light at 1.23 V vs. RHE and (d) photoconversion efficiency of all samples.

negligible current in the potential range of 0 to 1.6 V vs. RHE under dark conditions. Current densities are reported at a potential of 1.23 V vs. RHE. The photocurrent density of pure $\beta\text{-In}_2\text{S}_3$ is 0.32 mA cm^{-2} . The photocurrent densities of 2, 5 and 8 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ are 0.37, 0.69 and 0.46 mA cm^{-2} respectively. The photoanodes exhibited an onset potential of 0.14 V vs. RHE which is quite low in comparison to the literature reported values.²⁴ The low onset potential indicates good charge carrier separation and transport in the photoanode. Among all the doped samples, 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ shows the best performance with double the current density of pure $\beta\text{-In}_2\text{S}_3$. Particularly, Co^{2+} ions in $\beta\text{-In}_2\text{S}_3$ nanoflakes act as trapping sites for the photogenerated electrons which will prevent the recombination of electron-hole pairs and lead to enhancement of photoactivity. It leads to higher density of charge carriers. As cobalt is a transition metal with well-known electroactivity,^{25,39,40} it also acts as an active site for the promotion of water oxidation reaction. Hence doped $\beta\text{-In}_2\text{S}_3$ leads to the enhancement of both photo and electro activity. However, the increase in cobalt concentration beyond 5 mol% does not increase the current density. The probable cause for this is that the low temperatures limit the diffusion of cobalt ion inside the material but rather deposit on the surface beyond a certain concentration. Hence, the activity goes down after 5 mol% doping of cobalt.

Fig. 6c shows chronoamperometry (CA) curves of all the synthesized electrodes under chopped light at 1.23 V vs. RHE. High and stable photocurrent density is obtained for 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ when compared to other electrodes. Fig. 6d shows the solar to hydrogen (STH) conversion efficiency curves as a function of the applied voltage, which is calculated using the equation

$$\eta = \frac{J_{\text{ph}}[1.23 - V]}{P_{\text{in}}}$$

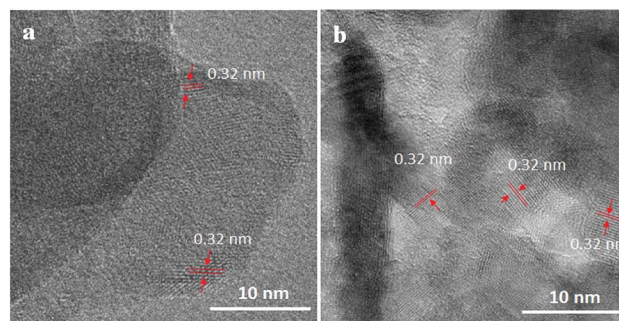


Fig. 7 TEM images of (a) 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ before testing and (b) TiO_2/Co -doped $\beta\text{-In}_2\text{S}_3$ (5 mol%) after testing.

where J_{ph} = photocurrent density (mA cm^{-2}) at the measured bias, V = applied bias potential versus reversible hydrogen electrode (RHE). P_{in} = power density of light illumination, 100 mW cm^{-2} .

5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ achieves the highest efficiency of 0.42% at 0.42 V, which is double than pure $\beta\text{-In}_2\text{S}_3$ (0.24% at 0.37 V). However, the onset potentials of all the electrodes are the same suggesting no occurrence of compositional and structural modification by introduction of cobalt. We also analyzed the samples as well as electrolyte after water oxidation for a span of 1 h. The electrolyte after prolonged illumination is mixed with thioacetamide and heated to 80°C for 2 h under stirring to check for indium dissolution. There is no change in the color of the electrolyte, which is indicative of no indium dissolution. In addition, we analyzed the TEM images before and after testing which are shown in Fig. 7. The d -spacing of (311) plane in Co-doped $\beta\text{-In}_2\text{S}_3$ is 0.32 nm. Co-doped $\beta\text{-In}_2\text{S}_3$ samples analyzed using TEM show no change in the d -spacing even after testing indicating no structural transformation to indium oxide. This shows that indeed, the stability of the Co-doped $\beta\text{-In}_2\text{S}_3$ is good and have a strong potential to be used practically.

Conclusions

In summary, Co-doped $\beta\text{-In}_2\text{S}_3$ nanoflakes are synthesized using chemical bath deposition at relatively low temperatures. 5 mol% Co-doped $\beta\text{-In}_2\text{S}_3$ nanoflakes with a current density of 0.69 mA cm^{-2} at 1.23 V vs. RHE and onset potential of 0.14 V vs. RHE showed superior performance in comparison to pure $\beta\text{-In}_2\text{S}_3$. In addition, it can be seen that the Co-doped $\beta\text{-In}_2\text{S}_3$ samples are stable in the neutral electrolyte.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We would like to thank Indian Institute of Technology, Hyderabad for the financial support (seed grant no: SG/IITH/F148/2016-17/SG-6) in providing the necessary research facilities. We



would also like to thank the TEM and XRD facilities in the institute for their help.

Notes and references

- 1 E. Fabbri, A. Haberer, E. Waltar, R. Kotz and R. J. Schmidt, *Catal. Sci. Technol.*, 2014, **4**, 3800–3821.
- 2 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- 3 R. A. He, S. W. Cao and J. G. Yu, *Acta Phys.-Chim. Sin.*, 2016, **32**, 2841–2870.
- 4 K. Sivula, F. Le Formal and M. Gratzel, *ChemSusChem*, 2011, **4**, 432–449.
- 5 S. V. Kershaw, A. S. Susa and A. L. Rogach, *Chem. Soc. Rev.*, 2013, **42**, 3033–3087.
- 6 P. Zhang, L. Yu and X. W. Lou, *Angew. Chem., Int. Ed.*, 2018, **57**, 15076–15080.
- 7 P. Kulkarni, S. K. Nataraj, R. G. Balakrishna, D. H. Nagaraju and M. V. Reddy, *J. Mater. Chem. A*, 2017, **5**, 22040–22094.
- 8 W. Sheng, Y. Song, M. Dou, J. Ji and F. Wang, *Appl. Surf. Sci.*, 2018, **436**, 613–623.
- 9 Z. Sun, Q. Yue, J. Li, J. Xu, H. Zheng and P. Du, *J. Mater. Chem. A*, 2015, **3**, 10243–10247.
- 10 M. Nguyen, P. D. Tran, S. S. Pramana, R. L. Lee, S. K. Batabyal, N. Mathews and M. Graetzel, *Nanoscale*, 2013, **5**, 1479–1482.
- 11 J. Zhang, Z. Zhu, Y. Tang, K. Mullen and X. Feng, *Adv. Mater.*, 2014, **26**, 734–738.
- 12 P. Zhang, B. Y. Guan, L. Yu and X. W. Lou, *Chem*, 2018, **4**, 162–173.
- 13 H. Jia, W. He, X. Chen, Y. Lei and Z. Zheng, *J. Mater. Chem. A*, 2011, **21**, 12824–12828.
- 14 J. Kim, H. Hiroi, T. K. Todorov, O. Gunawan, M. Kuwahara, T. Gokmen, D. Nair, M. Hopstaken, B. Shin, Y. S. Lee and W. Wang, *Adv. Mater.*, 2014, **26**, 7427–7431.
- 15 S. K. Sarkar, J. Y. Kim, D. N. Goldstein, N. R. Neale, K. Zhu, C. M. Elliott, A. J. Frank and S. M. George, *J. Phys. Chem. C*, 2010, **114**, 8032–8039.
- 16 Y. Tian, L. Wang, H. Tang and W. Zhou, *J. Mater. Chem. A*, 2015, **3**, 11294–11301.
- 17 T. Li, S. Zhang, S. Meng, X. Ye, X. Fu and S. Chen, *RSC Adv.*, 2017, **7**, 6457–6466.
- 18 F. Jiang, T. Harada, Y. Kuang, T. Minegishi, K. Domen and S. Ikeda, *J. Am. Chem. Soc.*, 2015, **137**, 13691–13697.
- 19 M. Q. Yang, B. Weng and Y. J. Xu, *Nano Res.*, 2013, **29**, 10549–10558.
- 20 J. Y. Do, R. K. Chava, S. K. Kim, K. Nahm, N. K. Park, J. P. Hong, S. J. Lee and M. Kang, *Appl. Surf. Sci.*, 2018, **451**, 86–98.
- 21 B. Sun, F. Shan, X. Jiang, J. Ji and F. Wang, *Appl. Surf. Sci.*, 2018, **435**, 822–831.
- 22 C. Gao, J. Li, Z. Shan, F. Huang and H. Shen, *Mater. Chem. Phys.*, 2010, **122**, 183–187.
- 23 R. Xu, H. Li, W. Zhang, Z. Yang, G. Liu, Z. Xu, H. Shao and G. Qiao, *Phys. Chem. Chem. Phys.*, 2016, **18**, 2710–2717.
- 24 L. Wang, L. Xia, Y. Wu and Y. Tian, *ACS Sustainable Chem. Eng.*, 2016, **4**, 2606–2614.
- 25 F. Lei, L. Zhang, Y. Sun, L. Liang, K. Liu, J. Xu, Q. Zhang, B. Pan, Y. Luo and Y. Xie, *Angew. Chem., Int. Ed.*, 2015, **54**, 9266–9270.
- 26 M. Kilani, C. Guasch, M. Castagne and N. Kamoun-Turki, *J. Mater. Sci.*, 2012, **47**, 3198–3203.
- 27 J. Feng, Z. Yang, S. He, X. Niu, T. Zhang, A. Ding, H. Liang and X. Feng, *Chemosphere*, 2018, **212**, 114–123.
- 28 C. Tapia, S. P. Berglund, D. Friedrich, T. Dittrich, P. Bogdanoff, Y. Liu, S. Levchenko, T. Unold, J. C. Conesa, A. L. De Lacey and M. Pita, *J. Phys. Chem. C*, 2016, **120**, 28753–28761.
- 29 J. Piris, A. J. Ferguson, J. L. Blackburn, A. G. Norman, G. Rumbles, D. C. Selmarten and N. Kopidakis, *J. Phys. Chem. C*, 2008, **112**, 7742–7749.
- 30 Z. Zheng, J. Yu, S. Cheng, Y. Lai, Q. Zheng and D. Pan, *J. Mater. Sci.: Mater. Electron.*, 2016, **27**, 5810–5817.
- 31 Y. Lu, Y. Lin, D. Wang, L. Wang, T. Xie and T. Jiang, *Nano Res.*, 2011, **11**, 1144–1152.
- 32 R. He, B. Tang, C. Ton-That, M. Phillips and T. Tsuzuki, *J. Nanopart. Res.*, 2013, **15**, 2030.
- 33 A. K. Nayak, S. Lee, Y. Sohn and D. Pradhan, *CrystEngComm*, 2014, **16**, 8064–8072.
- 34 O. Niitsoo, S. K. Sarkar, C. Pejoux, S. Rühle, D. Cahen and G. Hodes, *J. Photochem. Photobiol. A*, 2006, **181**, 306–313.
- 35 C. Wei, W. Guo, J. Yang, H. Fan, J. Zhang and W. Zheng, *RSC Adv.*, 2014, **4**, 50456–50463.
- 36 Y. Cheng, H. Niu, J. Chen, J. Song, C. Mao, S. Zhang and Y. Gao, *J. Nanopart. Res.*, 2017, **19**(5), 166.
- 37 J. Liqiang, Q. Yichun, W. Baiqi, L. Shudan, J. Baojiang, Y. Libin, F. Wei, F. Honggang and S. Jiazhong, *Sol. Energy Mater. Sol. Cells*, 2006, **90**, 1773–1787.
- 38 S. Y. Chae, P. Sudhagar, A. Fujishima, Y. J. Hwang and O. S. Joo, *Phys. Chem. Chem. Phys.*, 2015, **17**, 7714–7719.
- 39 A. Kaushik, B. Dalela, S. Kumar, P. A. Alvi and S. Dalela, *J. Alloys Compd.*, 2013, **552**, 274–278.
- 40 C. Bi, L. Pan, M. Xu, J. Yin, Z. Guo, L. Qin, H. Zhu and J. Q. Xiao, *Chem. Phys. Lett.*, 2009, **481**, 220–223.

