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Synthesis and characterization of ZnS with controlled amount of S vacancies for photocatalytic H₂ production under visible light

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Controlling amount of intrinsic S vacancies was achieved in ZnS spheres which were synthesized by a hydrothermal method using Zn and S powders in concentrated NaOH solution with NaBH₄ added as reducing agent. These S vacancies efficiently extend absorption spectra of ZnS to visible region. Their photocatalytic activities for H₂ production under visible light were evaluated by gas chromatograph, and the midgap states of ZnS introduced by S vacancies were examined by density functional calculations. Our study reveals that the concentration of S vacancies in the ZnS samples can be controlled by varying the amount of the reducing agent NaBH₄ in the synthesis, and the prepared ZnS samples exhibit photocatalytic activity for H₂ production under visible-light irradiation without loading noble metal. This photocatalytic activity of ZnS increases steadily with increasing the concentration of S vacancies until the latter reaches an optimum value. Our density functional calculations show that S vacancies generate midgap defect states in ZnS, which lead to visible-light absorption and responded.

In solving the global energy need and environmental pollution, hydrogen has attracted much attention for its potential to replace fossil fuels. Currently, however, production of hydrogen is mostly based on fossil fuels and on a process requiring high energy consumption^{1,2}. Since the report of photocatalytic activity of TiO₂ for hydrogen production, photocatalysis has become a desirable approach toward producing hydrogen with clean, environmentally friendly and economical process. In the past few decades, numerous photocatalysts have been found to exhibit high activities for water splitting³⁻⁶. These photocatalysts are mostly active only under UV light, which accounts for only 4% of the total sunlight. For practical applications, therefore, photocatalysts for hydrogen production need to operate under visible light.

ZnS is one of the most widely investigated photocatalysts because it rapidly generates electron-hole pairs under photoexcitation, and exhibits a relatively high activity for H₂ production under UV light^{6,7}. ZnS has a hexagonal structure and forms nanosheets or nanorods with large specific surface area⁷⁻⁹. Nevertheless, ZnS still has a substantially negative potential for excited electrons, and it does not respond to visible light due to its large band gap (~3.6 eV)¹⁰. Attempts have been made to extend the optical absorption of ZnS into the visible region by doping it with transition-metal ions (Au, Ni, Cu)¹⁰⁻¹². However, these composite materials were differently synthesized and hard to evaluate their intrinsic property of materials. Surface defects can also enhance light harvesting in photocatalytic materials¹³, and can also serve as adsorption sites where a charge transfer to the adsorbed species can prevent the recombination of photogenerated electrons and holes (namely, photogenerated charge carriers). However, when present in excessive amount, defects can act as traps for charge carriers leading to the recombination of photogenerated electrons and holes and hence decreasing the photocatalytic activity¹⁴⁻¹⁷. Therefore, controlling the amount of defects is great important to photocatalytic reaction. Vacancies are another kind of intrinsic defects in crystals, which are easily formed in quasi-two-dimensional materials because the exposed atoms on their surface can escape from the lattice hence affecting their physical and chemical properties^{18,19}. Theoretical investigations have reported that S vacancies can decrease the band gap of ZnS, and the introduction of S vacancies is harder than Zn vacancies in ZnS crystals^{20,21}. McCloy et al. introduced different



vacancies into cubic ZnS crystals by chemical vapor deposition under various atmospheres and studied the change in their lattice parameters²². To our knowledge, there has been no study on S vacancies in ZnS with an aim for photocatalytic applications.

In the present work we first controlled amount of S vacancies in ZnS spheres, and characterized the ZnS samples with S vacancies by various experimental techniques and density functional calculations. The ZnS samples with controllable S vacancies were obtained via adding NaBH₄ as a reducing agent in NaOH solution, and found to possess good photocatalytic activities for H₂ production without loading noble metal under visible light irradiation. The probable cause for the visible-light photocatalytic activity of these ZnS materials was explored.

Results

XRD and SEM characterizations. The XRD profiles for the ZnS samples synthesized using NaBH₄ as reducing agent are given in Fig. 1, which is coordinated with that of ZnS^{ref} reported by Zhang et al.¹⁰. These patterns have a strong and sharp peak at $2\theta = 27.2^\circ$, which can be indexed as the (100) reflection of wurtzite ZnS, and the broader peak is the overlapping peaks of wurtzite (002), sphalerite (111), and wurtzite (101)^{10,23}. These suggest that the {001} facet of wurtzite crystals is exposed, which is confirmed by SEM images (see below). The peaks of ZnS decrease with increasing the amount of NaBH₄ (hence the amount of S vacancies), because S vacancies diminish the crystallinity of ZnS crystals.

The morphologies of the prepared ZnS samples are shown in Fig. 2. The nanosheets of microspheres become crimped gradually with increasing the amount of NaBH₄, because S vacancies enhance the stress in the nanosheets of ZnS. The morphology of the microsphere turns into a blossom as the amount of NaBH₄ increases to 0.03 mol, in which nanosheets are already unfolded and broken. The morphologies of the microspheres are gradually ruptured, crisp and unfolded as the amount of NaBH₄ is increased, which may decrease the specific surface areas of the samples reducing the photocatalytic activity.

Optical absorption. The UV-Vis diffuse reflectance spectra of the ZnS samples obtained under different conditions are presented in Fig. 3, with their colour changes presented in the inset. It is clearly seen that ZnS^{ref} is a typical direct gap semiconductor with no

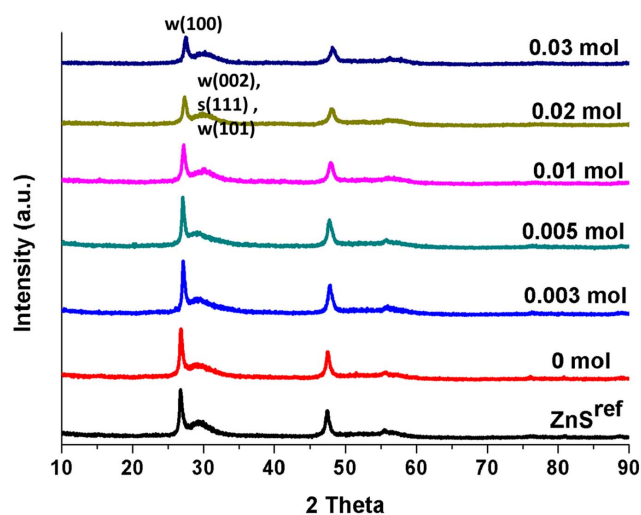


Figure 1 | XRD patterns obtained for ZnS^{ref} and the ZnS samples prepared using different amounts of NaBH₄ (0–0.03 mol), w(100) = wurtzite(100), w(002) = wurtzite (002), s(111) = sphalerite (111), and w(101) = wurtzite (101).

absorption in the visible light region. The ZnS samples, which obtained by the reaction between Zn and S powder without adding NaBH₄ (0 mol), exhibited visible light absorption, but one cannot control their visible-light absorption and their photocatalytic activities are low. NaBH₄ is a strongly reducing agent, and can reduce Zn²⁺ ions thereby creating S vacancies due to the charge balance requirement. Fig. 3 shows the absorption spectra measured for ZnS^{ref} and ZnS prepared without using NaBH₄ as well as those prepared using NaBH₄. The amount of NaBH₄ used for synthesis clearly affects the visible-light absorption; the larger the amount of NaBH₄, the stronger the absorption in the visible light.

Being a source of H⁻, NaBH₄ can reduce H₂O to generate hydrogen. However, under the high-concentration NaOH solution used in our synthesis, the concentration of H⁺ in the solution is negligible so that reduction of H₂O by H⁻ is negligible. Thus, H⁻ reduces Zn²⁺ ions of the ZnS crystal lattice thereby forming Zn⁰. To keep the charge balance, the amount of S atoms in the ZnS crystal lattice should decrease leading to S vacancies. With increasing the amount of NaBH₄, more S vacancies are formed resulting in a darker colour of the samples (see the inset of Fig. 3). Therefore, visible-light absorption of ZnS crystals can be tuned by the concentration of S vacancies, which in turn can be controlled by varying the amount of NaBH₄ in the synthesis process.

XPS. The chemical states of Zn²⁺ ions in ZnS samples were investigated by XPS (Full XPS survey scan display there was no impurity elements in the sample (Fig. S5, ESI[†])). As shown in Fig. 4, the binding energies of the Zn 2p_{3/2} and Zn 2p_{1/2} peaks in ZnS^{ref} are about 1021.5 eV and 1044.5 eV, respectively²⁴. When Zn and S powders were used for the synthesis without adding NaBH₄, the resulting ZnS sample weakly absorbs visible light and a lower binding energy for both Zn 2p_{3/2} and Zn 2p_{1/2}. This is attributed to the presence of some S vacancies²⁰. With increasing the amount of NaBH₄ during the synthesis, the binding energy for the Zn 2p_{3/2} and Zn 2p_{1/2} peaks of the ZnS samples shift gradually to a lower energy. This indicates that, to a certain degree, the concentration of S vacancies can be controlled by varying the amount of NaBH₄ used. As depicted in Fig. 5, the Zn 2p_{3/2} peaks can be decomposed into two Gaussian peaks. With increasing the amount of NaBH₄ from 0 to 0.03 mol, the major Zn 2p_{3/2} peaks change from 1020.8 eV to 1018.9 eV, and the minor Zn 2p_{3/2} peaks from 1019.1 eV to 1017.4 eV. At the same time, the major peaks of Zn 2p_{1/2} change from 1043.8 eV to 1041.8 eV, and the minor Zn 2p_{1/2} from 1042.1 eV to 1039.9 eV. All these peaks are lower in energy than that of ZnS^{ref}. It is reasonable to assign the major peaks to the Zn²⁺ ions away from the S vacancy sites, and the minor peaks to those Zn²⁺ ions adjacent to the S vacancy sites, and that the presence of Zn⁰ atoms around the S vacancy sites (see below) lowers the binding energies of the Zn²⁺ ions. EDS results also proved that absence of S element in ZnS samples (Fig S6, ESI[†]). The XPS results of S element in different samples were also investigated, and peaks of S 2p were all shifted to higher energy (Fig S7, ESI[†]).

Surface areas of ZnS. An important factor affecting the photocatalytic activity of a ZnS sample is its surface area (Table 1). Fig S2 (ESI[†]) shows the typical isotherms of N₂ adsorption onto the ZnS sample obtained using 0.01 mol of NaBH₄, with the pore size distributions in the inset. These isotherms exhibit type IV isotherms with hysteresis loops according to the IUPAC classification²⁵, which implies the presence of mesopores. The pore size distribution curves (the inset in Fig S2) indicate that the sample possesses mesopores and macropores in a wide size range of 10 to over 90 nm. Use of NaBH₄ can increase the BET of the sample to a certain degree. The ZnS sample obtained with 0.01 mol of NaBH₄ possesses a relatively large specific surface area, while use of a greater amount of NaBH₄ decreases by destroying the microspheres.

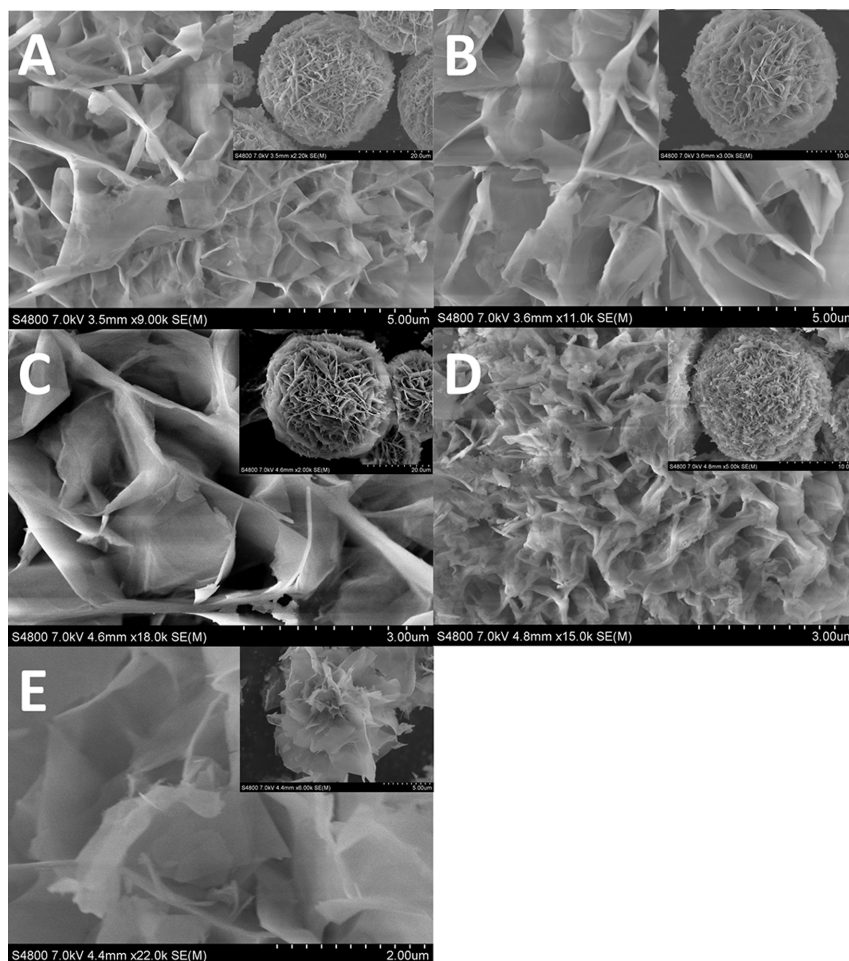


Figure 2 | SEM images of the ZnS samples synthesized using different amounts of NaBH_4 . (A) 0.003, (B) 0.005, (C) 0.01, (D) 0.02, and (E) 0.03 mol NaBH_4 .

Photocatalytic production of H_2 . The ability of the ZnS samples (0.5 g) for photocatalytic hydrogen generations under visible light ($\lambda > 420$ nm) irradiation was evaluated by using 100 mL aqueous solution containing Na_2S (0.25 M) and Na_2SO_3 (0.5 M) as sacrificial agents. The H_2 production of the ZnS sample (obtained with 0.01 mol of NaBH_4) as a function of the irradiation time is presented in Fig. 6A, which shows that the activity of the sample

has no decrease after 7 h irradiation (The XPS results of this sample, before and after photocatalytic reaction, were also show in Fig S4, ESI[†]). The hydrogen production rates of the ZnS samples synthesized using different amounts of NaBH_4 (0–0.03 mol), displayed in Fig. 6B, show that the production rate increases until the amount of NaBH_4 reaches 0.01 mol but drops abruptly when it is beyond 0.01 mol. This means that the S vacancies enhance the visible light absorption resulting in an enhanced high photocatalytic activity. However, when present in excess amount, vacancies act as

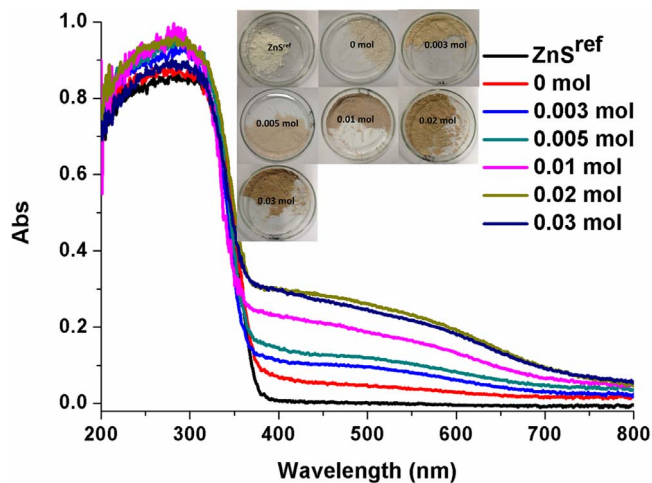


Figure 3 | UV-Vis diffuse reflectance spectra of ZnS synthesized using 0–0.03 mol of NaBH_4 .

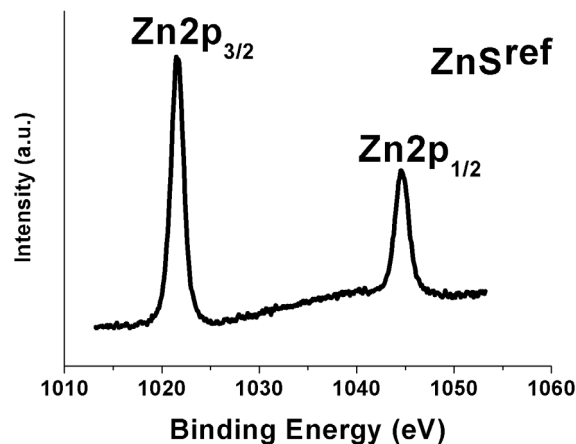


Figure 4 | XPS results of ZnS^{ref} sample.

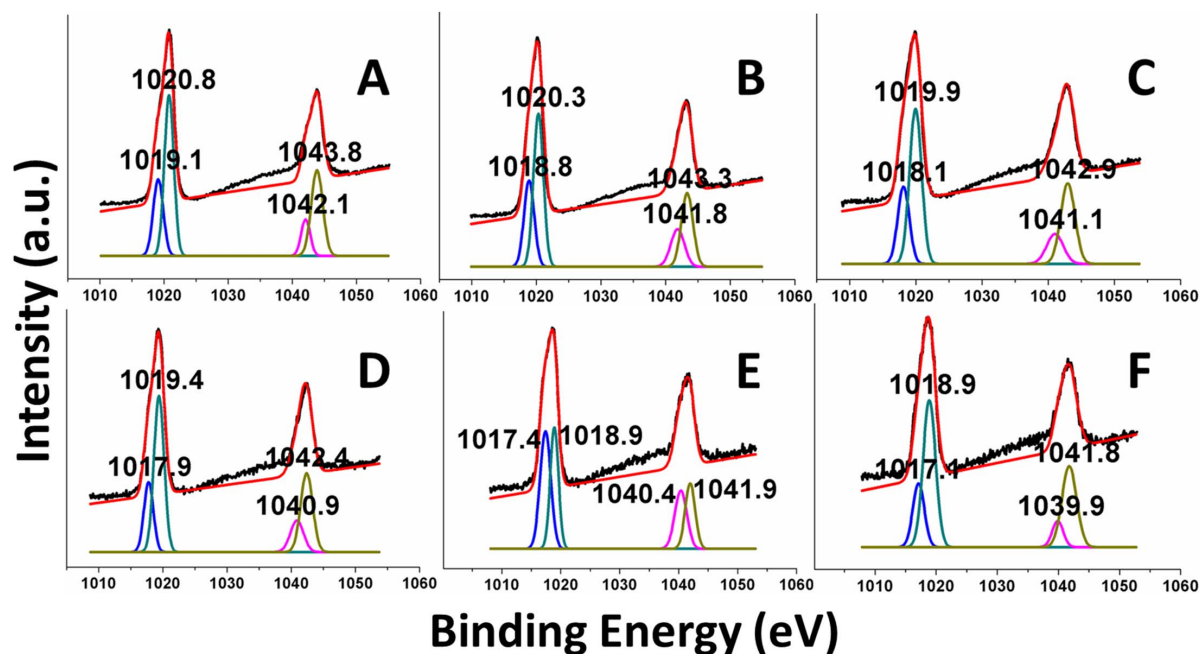


Figure 5 | Zn $2p_{3/2}$ spectra of the ZnS samples synthesized using different amounts of NaBH_4 . (A) 0, (B) 0.003, (C) 0.005, (D) 0.01, (E) 0.02, and (F) 0.03 mol of NaBH_4 .

recombination centers for photogenerated carriers decreasing the photocatalytic activity. With increasing the amount of NaBH_4 to 0.02 and 0.03 mol, lots of hierarchical microspheres are broken, which can also contribute to the precipitous decline of the H_2 production rate. Cycling photocatalytic experiments of ZnS were also done for evaluating the stability of the ZnS samples (Fig S8, ESI[†]), which exhibited good stability after 3 times photocatalytic reactions.

PL spectra. The photocatalytic activity is enhanced when photogenerated electron-hole pairs are efficiently separated. PL emission is an effective way of estimating the ability of a sample to separate these photogenerated carriers. The PL spectra of different ZnS samples are presented in Fig. 7, which were carried out on a Hitachi F-4500 fluorescence spectrophotometer at room temperature and obtained with excitation wavelength at 300 nm. It is observed that ZnS^{ref} has the highest intensity of photoluminescence emission, indicating that the photogenerated carriers are quickly recombined. The PL intensity of the ZnS samples decreases with increasing the amount of NaBH_4 reaches 0.01 mol, but decreases when it is beyond 0.01 mol. It clearly indicates that the S vacancies can become centers for the recombination of photogenerated electrons and holes when present in excessive amount. This finding is consistent with the results of the photocatalytic activities of these ZnS samples.

Density functional characterization of the defect states. The density of states (DOS) plot calculated for ZnS with no S vacancy is presented in Fig. 8A, which shows a band gap of ~ 3.0 eV. The

DOS plots calculated for the model ZnS systems with one S vacancy in the $2 \times 2 \times 2$, $4 \times 2 \times 2$ and $4 \times 4 \times 2$ supercells are presented in Figs. 9B–D, respectively, where the insets show the projected DOS (PDOS) plots calculated for the four Zn atoms surrounding the S vacancy. These plots reveal that the S vacancy generates new states within the band gap of defect-free ZnS. The presence of these midgap defect states causes a visible-light absorption for the ZnS samples with S vacancies. It is of interest to discuss the formation of these defect states from the viewpoint of the local electronic structure around an S vacancy.

The examination of the relaxed structures around the S vacancies show that one of the four ZnS_3 pyramids surrounding the S vacancy becomes strongly pyramidal (with $\angle \text{S-Zn-S} \approx 95^\circ$) while the remaining three ZnS_3 pyramids do not undergo a strong change in shape (with $\angle \text{S-Zn-S} \approx 105^\circ$). In ZnS each S^{2-} ion is surrounded by four Zn^{2+} ions to form a SZn_4 tetrahedron (Fig. 9A), and each Zn^{2+} ion surrounded by four S^{2-} ions to form a ZnS_4 tetrahedron. To a first approximation, the bonding in ZnS can be described in terms of the sp^3 hybridization for the Zn and S atoms so that each Zn–S bond results from a bonding combination of a filled sp^3 orbital of S^{2-} (i.e., a sp^3 lone pair) with an empty sp^3 orbital of Zn^{2+} (Fig. 9B). When an S atom is removed from a Zn_4 tetrahedron, each of the four Zn atoms surrounding the S vacancy becomes a ZnS_3 pyramid with one sp^3 dangling bond. If the structure around the vacancy site is not relaxed, then the interactions between the four dangling bonds lead to the midgap states 1a and 1t ($1a < 1t$) of the Zn_4 tetrahedron lying approximately in the middle of the gap between the valence band maximum (VBM) and conduction band minimum (CBM) (left of

Table 1 | BET surface areas and pore structures of the ZnS samples obtained using different amounts of NaBH_4

NaBH_4 (mol)	Surface area (m^2/g)	Pore volume (cc/g)	Pore size (Å)
0	113.1	0.55	97.7
0.003	87.4	0.65	148.3
0.005	93.9	0.36	94.2
0.01	129.3	0.61	87
0.02	112.9	0.5	89.1
0.03	90.1	0.53	117.6

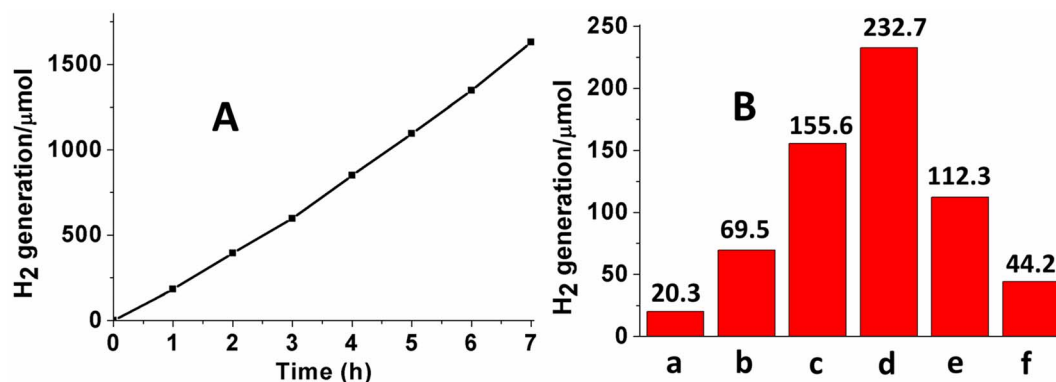


Figure 6 | Photocatalytic reactions results. (A) Photocatalytic production of H_2 by the ZnS sample (0.5 g) obtained using 0.01 mol $NaBH_4$ as a function of the visible-light irradiation time. (B) Hydrogen production rate by ZnS samples obtained using different amounts of $NaBH_4$ (a = 0 mol, b = 0.003 mol, c = 0.005 mol, d = 0.01 mol, e = 0.02 mol, and f = 0.03 mol).

Fig. 9C) (see Fig S3, ESI[†] for the DOS plot in the case of no geometry relaxation around the S vacancy). When the structure around the vacancy site is relaxed, the energy gap between the 1a and 1t levels becomes large with the 1t level splitting (right of Fig. 9C) due to the symmetry lowering. As a consequence, the 1a level comes close to the VBM and becomes doubly filled, while the three empty levels come close to the CBM. In the relaxed structure around an S vacancy, one ZnS_3 pyramid that becomes strongly pyramidal ($\angle S-Zn-S \approx 95^\circ$) has its dangling bond doubly filled becoming a lone pair (Fig. 9D). Then the optical excitations associated with the filled defect level close the VBM and the empty defect levels below the CBM are responsible for the visible-light absorption of the ZnS samples with S vacancies. Furthermore, trapping of photogenerated electrons and holes by these defect states helps slow down their recombination, thereby enhancing the photocatalytic activities of ZnS samples with S vacancies.

In summary, aimed at finding photocatalytic ZnS systems for H_2 production operating under visible-light irradiation, we synthesized a new ZnS samples with controlled amount of S vacancies by the hydrothermal reaction of zinc and sulfur powders with $NaBH_4$ added as a reducing agent. The concentration of S vacancies in the ZnS samples can be easily controlled by varying the amount of $NaBH_4$. The as-prepared ZnS samples exhibit enhanced visible-light absorption with increasing the concentration of S vacancies. The photocatalytic activities of these samples for H_2 production, evaluated under visible light irradiation, show that the photocatalytic activity increases with increasing the concentration of S vacancies until it reaches the optimum value, but decreases sharply when it goes

beyond the optimum value because, when present in excessive amount, S vacancies destroy both crystal structure and morphologies of the samples and also play as recombination sites of photogenerated electrons and holes. Our density functional analysis reveals that S vacancies generate midgap defect states in ZnS, which are responsible for visible-light absorption and also act as trap centers for electrons and holes improving the separation efficiency of photo-generated charge carriers. This work provides a novel method to induce and control S vacancies in crystals for enhancing their photocatalytic H_2 production under visible light.

Methods

Materials. All chemicals in this work (i.e., Zn powder, sublimed sulfur, NaOH, $NaBH_4$, $Na_2S \cdot 9H_2O$, Na_2SO_3) were AR reagents, and were used without any further purification.

Preparation of ZnS. Samples of ZnS with S vacancies were synthesized by mixing Zn powder and sublimed sulfur in 50 ml of 21 M NaOH solution. After the suspension was cooled to room temperature, a various amount of $NaBH_4$ (0–0.03 mol) was added to control the amount of S vacancies in ZnS crystals. After constant stirring for 2 h, the mixed solution was transferred into a 120 ml sealed Teflon-lined autoclave and was heated at $230^\circ C$ for at least 12 h. The autoclave was slowly cooled down to room temperature, and the obtained sample was washed with distilled water, and was finally dried at $40^\circ C$ overnight. For the purpose of comparison, we prepared a reference ZnS sample following the method proposed by Zhang et al.¹⁰ with slight modification¹⁰, which will be hereafter referred to as ZnS^{ref} ; a certain amount of ZnO was dissolved in 50 ml of 21 M NaOH solution, and $Na_2S \cdot 9H_2O$ was added after the solution cooled down. The suspension was transferred into a 120 mL sealed Teflon-lined autoclave and followed by hydrothermal treatment at $230^\circ C$ for at least 12 h. The sample was also collected after washing and drying.

Sample characterizations. XRD patterns were recorded on a Bruker AXS D8 Advance powder diffractometer (Cu $K\alpha$ X-ray tube, $\lambda = 0.154056$ nm). The morphologies of the samples were obtained using SEM (Hitachi S-4800). The chemical energy-dispersive X-ray spectrum (EDS) were examined by an energy dispersive X-ray spectrometer equipped in the SEM machine. The surface areas of the as-prepared samples were measured by the BET method using nitrogen adsorption-desorption isotherms on a Micromeritics ASAP 2020 apparatus at liquid nitrogen temperature. UV-Vis diffuse reflectance spectra were obtained for dry-pressed disk samples by using a Shimadzu UV 2550 recording spectrophotometer equipped with an integrating sphere, and $BaSO_4$ was used as a reference. PL spectra were carried out on a Hitachi F-4500 fluorescence spectrophotometer at room temperature and the excitation wavelength was 300 nm.

Details of calculations. Our density functional calculations employed the projector augmented wave method as implemented in the Vienna ab initio simulation package^{26–29}, the generalized gradient approximation of PBE for exchange and correlation corrections^{30,31} with plane wave cutoff energies of 400 eV, and a threshold of self-consistent-field energy convergence of 10^{-4} eV. The atomic positions were fully optimized until all the residual forces are smaller than 0.01 eV/Å. The structure of hexagonal ZnS was constructed on the basis of $2 \times 2 \times 2$, $4 \times 2 \times 2$ and $4 \times 4 \times 2$ supercells containing 32, 64 and 128 atoms, respectively. Various model structures of sulfur-deficient ZnS were constructed by removing one sulfur atom from the 32-atom $2 \times 2 \times 2$ supercell (see Fig. S1 ESI[†]), 64-atom $4 \times 2 \times 2$ supercell, and 128-atom $4 \times 4 \times 2$ supercell.

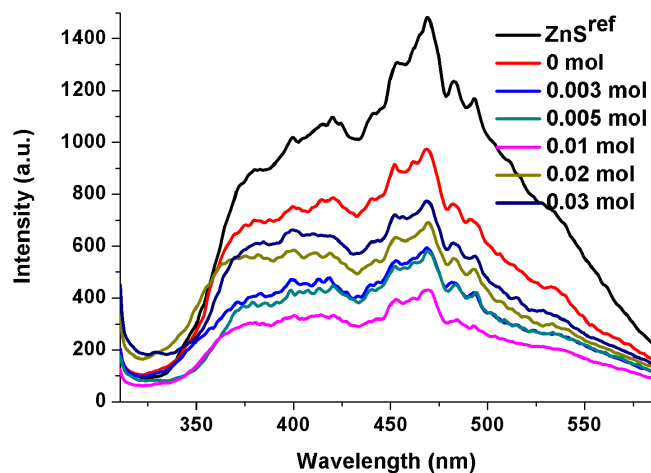


Figure 7 | PL spectra of different ZnS samples ($\lambda_{exc} = 300$ nm).

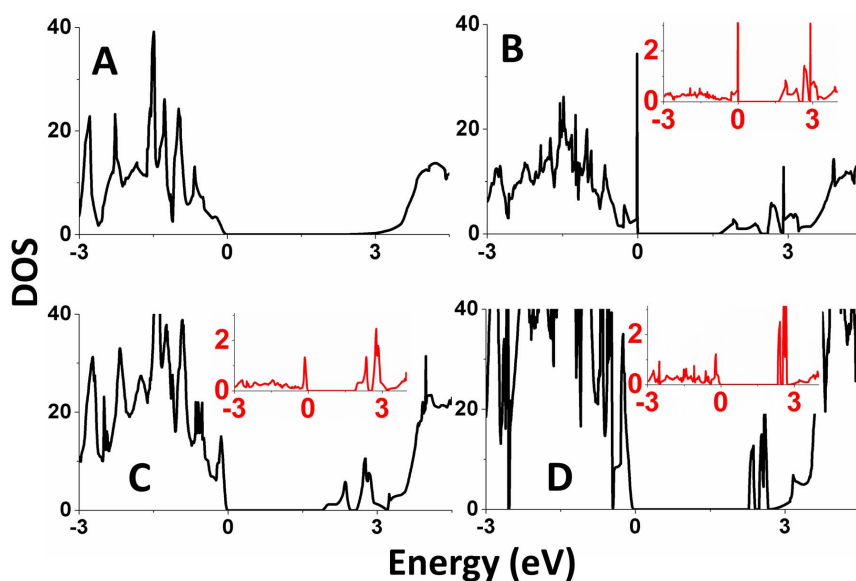


Figure 8 | Total DOS and PDOS plots calculated for various ZnS with and without S vacancy. The PDOS plots, shown in the inset, are calculated for the four Zn atoms surrounding the S vacancy. (A) Hexagonal ZnS without S vacancy in which there are 32 atoms per $2 \times 2 \times 2$ supercell. (B) ZnS in which there is one S vacancy per $2 \times 2 \times 2$ supercell. (C) ZnS in which there is one S vacancy per $4 \times 2 \times 2$ supercell. (D) ZnS in which there is one S vacancy per $4 \times 4 \times 2$ supercell.

Photocatalytic activity for H₂ production. A top irradiation vessel connected to a glass-enclosed gas-circulation system was used to evaluate the photocatalytic hydrogen evolution of the ZnS samples. In a typical photocatalytic experiment, 0.5 g sample was suspended in 100 mL aqueous solution containing Na₂S·9H₂O (0.35 M) and Na₂SO₃ (0.25 M) as sacrificial agents. The temperature was maintained at 5°C. The visible light source was obtained from a 300 W Xe arc lamp (PLS-SXE 300,

Beijing Trusttech Co. Ltd.) equipped with an ultraviolet cutoff filter ($\lambda > 420$ nm). The amount of H₂ produced was determined with a gas chromatograph (Varian GC-3800) equipped with thermal conductivity detector.

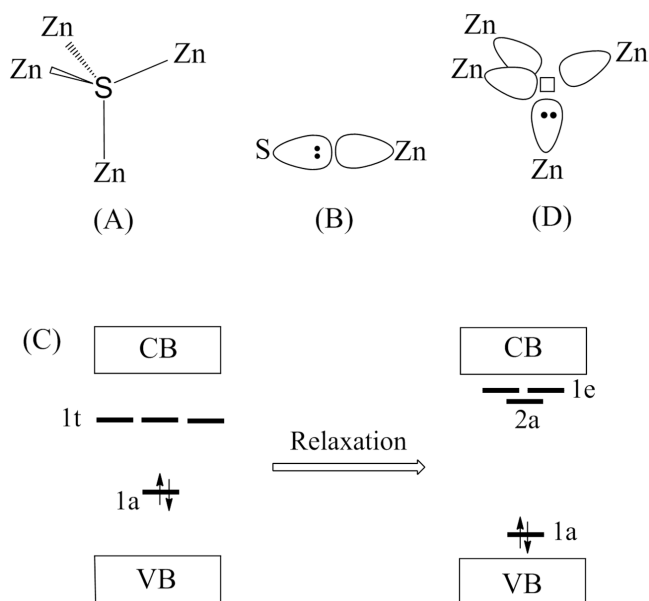


Figure 9 | The examination of the relaxed structures. (A) SZn₄ tetrahedron of ZnS. (B) Zn-S bonding as a combination of a filled sp³ hybrid orbital of S²⁻ with an empty sp³ orbital of Zn²⁺. (C) Schematic diagram showing the midgap defect states of ZnS created by an S vacancy before (left) and after (right) relaxing the structures around the vacancy site. (D) Overall effect of an S vacancy showing the formation of one of the four ZnS₃ pyramids surrounding the S vacancy getting two electrons. The VB and CB refer to valence and conduction bands, respectively. Here, for simplicity, it was assumed that the unrelaxed Zn₄ tetrahedron has a Td symmetry, and the relaxed Zn₄ tetrahedron a C₃ symmetry.

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Author contributions

G.W., B.B.H., Z.Z.L. and Z.Y.W. conceived the project, carried out the experiments and analysed the experimental data. Z.J.L., Y.D. and M.-H.W. performed the density functional theory calculations. All authors contributed to writing the manuscript.

Additional information

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