

# Synergistic Effect of Redox Dual PdO<sub>x</sub>/MnO<sub>x</sub> Cocatalysts on the Enhanced H<sub>2</sub> Production Potential of a SnS/α-Fe<sub>2</sub>O<sub>3</sub> Heterojunction via Ethanol Photoreforming

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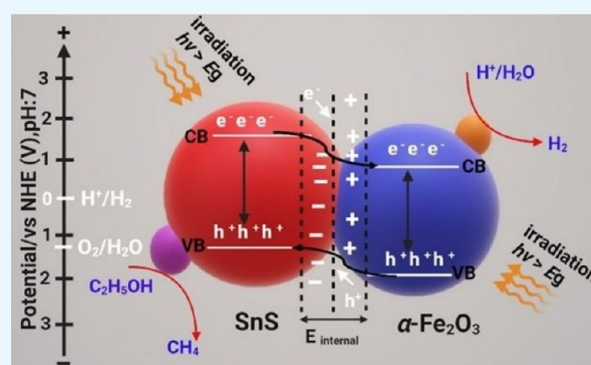
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**ABSTRACT:** In the quest for optimal H<sub>2</sub> evolution (HE) through ethanol photoreforming, a dual cocatalyst-modified heterocatalyst strategy is utilized. Tin(II) sulfide (SnS) was hybridized with α-Fe<sub>2</sub>O<sub>3</sub> to form the heterocatalyst FeOSnS with a p–n heterojunction structure as confirmed by X-ray diffraction (XRD), Fourier transform infrared spectroscopy (FT-IR), UV–vis diffusive reflectance spectroscopy (UV–vis DRS), and Brunauer–Emmett–Teller (BET) techniques. PdO<sub>x</sub> and PdO<sub>x</sub>/MnO<sub>x</sub> cocatalysts were loaded onto the FeOSnS heterocatalyst through the impregnation method, as verified by high-resolution transform electron microscopy (HRTEM), X-ray photoelectron spectroscopy (XPS), and elemental mapping. Photocatalytic ethanol photoreforming resulted in the production of H<sub>2</sub> as the main product with a selectivity of 99% and some trace amounts of CH<sub>4</sub>. The FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% photocatalyst achieved the highest HE rate of 1654 μmol/g, attributed to the synergistic redox contribution of the PdO<sub>x</sub> and MnO<sub>x</sub> species.



## 1. INTRODUCTION

Hydrogen (H<sub>2</sub>) is a crucial requirement for many industrial chemical processes such as ammonia synthesis (~50%), oil refining (~40%), methanol synthesis (~8%), and transportation. As of 2020, approximately 87 million tons of H<sub>2</sub> were generated worldwide, 95% of which is from fossil fuels by steam reforming of methane and other hydrocarbons as well as coal gasification. Nevertheless, the burning of fossil fuels releases large amounts of greenhouse gases such as CO<sub>2</sub> into the atmosphere, leading to global warming.<sup>1,2</sup> Solar energy is considered an ideal alternative to fossil fuels due to its high energy capacity (~1.2 × 10<sup>14</sup> kJ received at the Earth's surface every second), low cost, and abundance.<sup>3</sup> Solar-light-assisted splitting of H<sub>2</sub>O over a TiO<sub>2</sub> photocatalyst was proposed in 1972 by Honda and Fujishima as a sustainable way for H<sub>2</sub> production.<sup>4</sup> Currently, the best performance has been achieved by TiO<sub>2</sub>-based systems using UV light. The advantages of TiO<sub>2</sub> include availability, low cost, chemical stability, high chemical inertness, and nontoxicity.<sup>5</sup> Nevertheless, one disadvantage of TiO<sub>2</sub> is its limited activity in the visible spectrum due to its large band gap ( $E_g \approx 3.2$  eV) and fast recombination of photo-generated electron–hole pairs.<sup>6</sup>

In spite of the significant body of research conducted with TiO<sub>2</sub>, other UV-active photocatalysts such as ZnO<sup>7</sup> and BiPO<sub>4</sub><sup>8</sup> and even visible-active photocatalysts such as Bi<sub>2</sub>WO<sub>6</sub>,<sup>9</sup> WO<sub>3</sub>,<sup>10</sup> and BiVO<sub>4</sub><sup>11</sup> with wide band gaps ( $E_g < 3$  eV), the H<sub>2</sub>

production efficiency achieved in water photosplitting is still too low for industrial viability. This is due to the occurrence of energetically favorable backward H<sub>2</sub> and O<sub>2</sub> reactions to yield water, fast recombination of photogenerated electron and hole (e<sup>-</sup>/h<sup>+</sup>) pairs due to the strong Coulombic force, limited light absorption efficiency, low charge transport properties, photocorrosion, and instability in water solutions.<sup>12</sup> An alternative strategy is the photoreforming of aqueous solutions containing biomass-derived oxygenates such as ethanol, glycerol, and glucose. Additionally, photoreforming can lead to the production of benzaldehyde, formaldehyde, and cyclohexanone as industrially attractive products.<sup>13</sup>

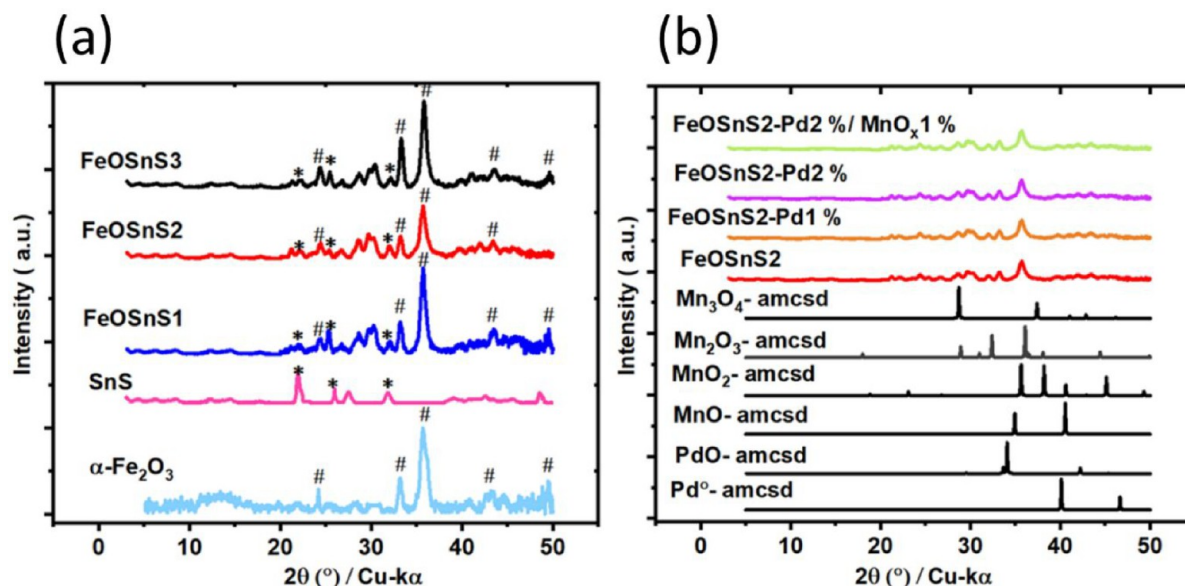
Hematite (α-Fe<sub>2</sub>O<sub>3</sub>) is an n-type photocatalyst and has been examined for photocatalytic H<sub>2</sub> production through H<sub>2</sub>O splitting due to its abundance, nontoxicity, good corrosion resistance, low cost, and high photo-/thermostability.<sup>14</sup> Importantly, the narrow band gap of 1.9–2.2 eV renders it with the ability to absorb about 40% of the incident visible solar

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**Figure 1.** (a) PXRD patterns of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and SnS, FeOSnS1–3. (b) PXRD patterns of FeOSnS2-PdO<sub>x</sub>/MnO<sub>x</sub> catalysts.

energy.<sup>15</sup> Nevertheless, the performance of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> is still limited by its low electrical conductivity ( $\sim 10$ – $14 \Omega^{-1} \text{ cm}^{-1}$ ), short hole-diffusion distance (2–4 nm), poor electron–hole pair lifetime (<10 ps), poor oxygen evolution reaction (OER) kinetics, and weak charge mobility ( $10^{-2}$  to  $10^{-1} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ).<sup>16,17</sup> Furthermore, its conduction band (CB) is not negative enough to reduce H<sup>+</sup> ions to H<sub>2</sub>.<sup>18</sup> Various strategies have been adopted to address these drawbacks associated with  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> for water splitting. These include the development of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> nanoarchitectures with sizes smaller than its hole diffusion length,<sup>19</sup> surface state passivating,<sup>20</sup> the creation of oxygen vacancies,<sup>21</sup> doping of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> with heteroatoms such as Sc, Ti, Cr, Mn, and Ni,<sup>22</sup> and construction of Z-scheme p–n junction heterostructures.<sup>23</sup> In contrast to water splitting, few studies have highlighted the H<sub>2</sub> production through ethanol photoreforming using  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>. For example, Carraro et al. reported that control of the crystal phase of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> can significantly improve the H<sub>2</sub> production potential. The results revealed H<sub>2</sub> production rates of 40, 225, and 125 mmol h<sup>-1</sup> m<sup>-2</sup> for  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>,  $\beta$ -Fe<sub>2</sub>O<sub>3</sub>, and  $\epsilon$ -Fe<sub>2</sub>O<sub>3</sub>, respectively, through photoreforming of ethanol/water solutions under irradiation from a 150 W Xe lamp.<sup>24</sup> Wender et al. reported an improved H<sub>2</sub> production rate of 546 mmol h<sup>-1</sup> g<sup>-1</sup> for an  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> nanoring loaded with a 7% Co(OH)<sub>2</sub> cocatalyst versus 350  $\mu\text{mol h}^{-1} \text{ g}^{-1}$  for pure  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>.<sup>25</sup>

Tin monosulfide (SnS) is a p-type semiconductor with excellent optoelectric characteristics.<sup>26</sup> SnS has an indirect band gap of 1–1.2 eV, a direct optical band gap of 1.2–1.5 eV, high absorption coefficient ( $10^4$ – $10^5 \text{ cm}^{-1}$  in the visible region), high conductivity, low cost, and high charge mobility.<sup>27</sup> Cocatalysts such as noble metals (Ni, Pt, Au, Pd, and Rh and Ag) or transition metal oxides (NiO, Co<sub>3</sub>O<sub>4</sub>, MnO<sub>x</sub>, and CuO<sub>x</sub>) have been extensively examined as important charge mediators to boost photocatalytic performance.<sup>28</sup>

In this contribution, we have optimized the conduction band energy of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> through hybridization with SnS to improve the reduction of H<sup>+</sup> ions to H<sub>2</sub> via ethanol photoreforming. Additionally, the FeOSnS heterocatalyst was decorated with PdO<sub>x</sub> and MnO<sub>x</sub> dual cocatalysts to boost the H<sub>2</sub> production efficiency. This is the first report on the tertiary mixture of  $\alpha$ -

Fe<sub>2</sub>O<sub>3</sub>/SnS with dual redox cocatalysts for photocatalytic H<sub>2</sub> production.

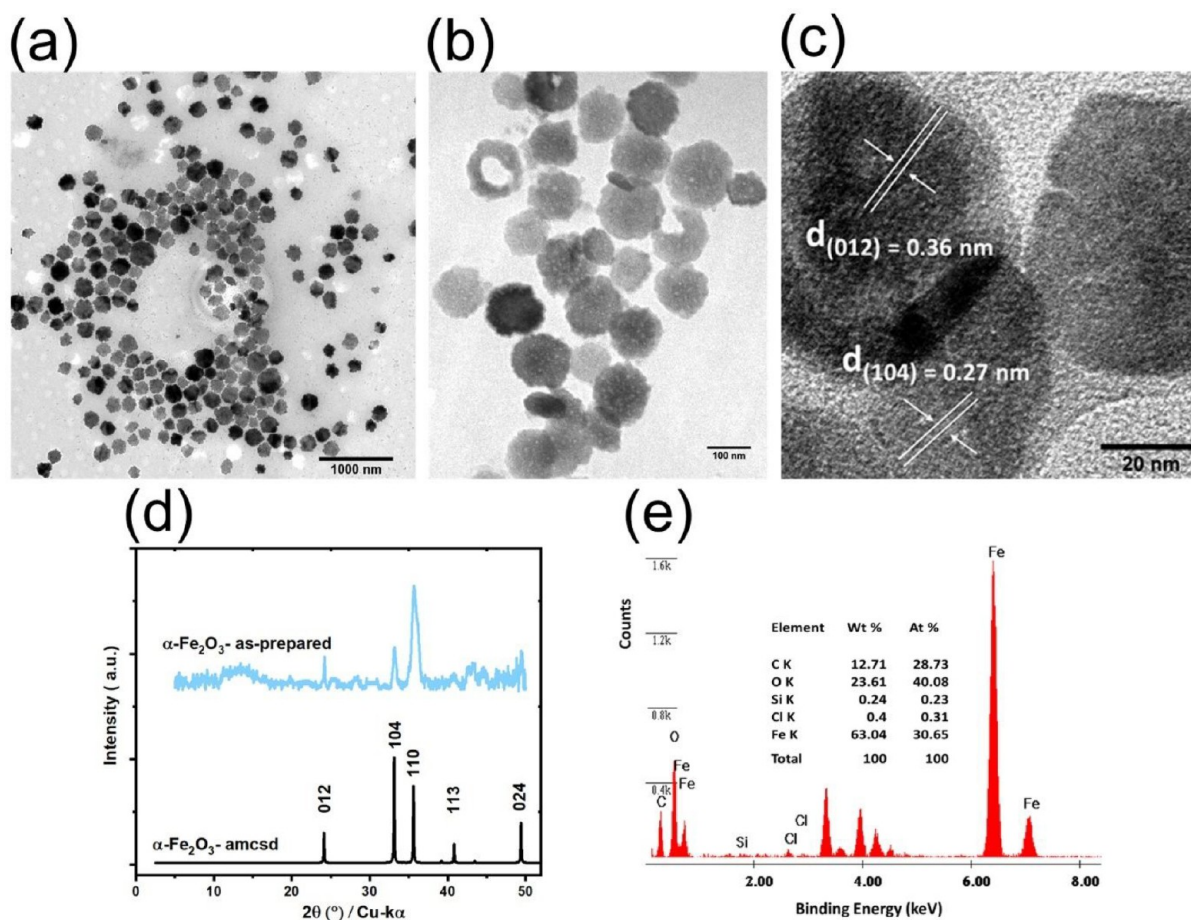
## 2. RESULTS AND DISCUSSION

### 2.1. Synthesis of the Heterocatalysts and Cocatalysts.

FeOSnS heterocatalysts were prepared using the *in situ* chemical precipitation method with SnS: $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> ratios of 1:1 w/w (FeOSnS1), 1.5:0.5 w/w (FeOSnS2), and 0.5:1.5 w/w (FeOSnS3). An impregnation method was then used to load PdO<sub>x</sub> and MnO<sub>x</sub> catalysts onto the FeOSnS2 catalyst, forming FeOSnS2-PdO<sub>x</sub> 1%, FeOSnS2-PdO<sub>x</sub> 2%, and FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalysts based on the mass percentages of the loaded cocatalysts. The synthesized catalysts were characterized by X-ray diffraction, transmission electron microscopy (TEM), scanning electron microscopy (SEM), X-ray photoelectron spectroscopy (XPS), atomic absorption spectroscopy (AAS), and UV–vis diffuse reflectance spectroscopy (UV–vis DRS).

**2.2. X-ray Diffraction.** The crystalline phase and purity of the as-prepared  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, SnS, FeOSnS1–3, and FeOSnS2-PdO<sub>x</sub>/MnO<sub>x</sub> catalysts were studied by powder X-ray diffraction (PXRD) (Figure 1). Pristine  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, which was synthesized from the decomposition of Fe(acac)<sub>3</sub> in DMF in the presence of PVP surfactant at 180 °C, presents distinct diffraction peaks at  $2\theta = 24.1^\circ$ ,  $33.1^\circ$  (main characteristic peak),  $35.6^\circ$ ,  $40.8^\circ$ , and  $49.4^\circ$ , assigned to the crystal planes of (012), (104), (110), (113), and (024), respectively (Figure 1(a)).<sup>29</sup> For pristine SnS, formed from SnCl<sub>2</sub> and sulfur in the presence of PVP in DMF, the characteristic diffraction peaks were observed at  $2\theta = 21.9^\circ$ ,  $25.9^\circ$ ,  $27.6^\circ$ ,  $31.9^\circ$ ,  $39.2^\circ$ ,  $42.6^\circ$ ,  $45.5^\circ$ , and  $48.5^\circ$  corresponding to the (011), (012), (102), (004), (113), (021), (015), and (023) diffraction planes, respectively (Figure 1(a)).<sup>30</sup>

In the PXRD pattern of FeOSnS1–3 heterocatalysts (Figure 1(a)), a series of characteristic diffractions peaks from  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> (indicated by the symbol #) and SnS (indicated by the symbol \*) are observed. In addition, the main diffraction peaks in the heterocatalysts have minor changes compared with pristine  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and SnS, suggesting that the hybridization had a negligible influence on the original crystal structure of the constituents. This also indicates that any improvement in the photocatalytic potential is not caused by alteration of the crystal



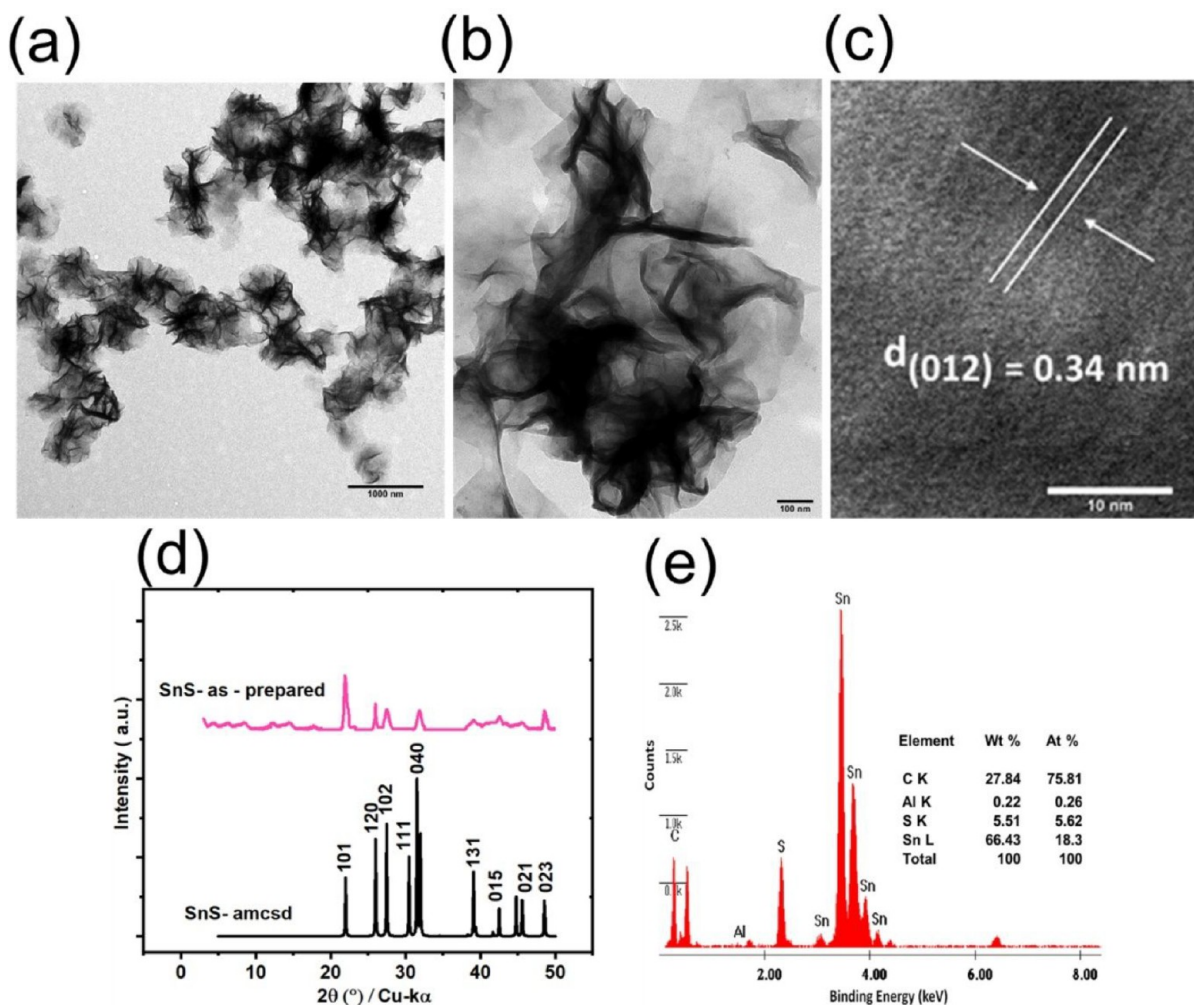
**Figure 2.** (a)–(c) TEM and HRTEM image, (d) PXRD pattern (pattern for  $\alpha\text{-Fe}_2\text{O}_3$  (AMCSD card no. 0000143) is shown for comparison), and (e) EDS spectra of the  $\alpha\text{-Fe}_2\text{O}_3$  catalyst.

structure of the constituents.<sup>31</sup> It is worth noting that we used an *in situ* chemical precipitation method to synthesize FeOSnS heterocatalysts rather than direct physical mixing. The reason was to improve the charge transfer between the two photocatalysts through the strong chemical bonding formed at the interface.<sup>32</sup> In the case of the FeOSnS2-PdO<sub>x</sub>/MnO<sub>x</sub> catalyst, no obvious characteristic peaks of PdO<sub>x</sub> species (metallic Pd or PdO) or MnO<sub>x</sub> species (MnO, Mn<sub>2</sub>O, Mn<sub>2</sub>O<sub>3</sub>, and Mn<sub>3</sub>O<sub>4</sub>) were detected (Figure 1(b)). This was due to the low loading amount (1 and 2%) of the PdO<sub>x</sub> and MnO<sub>x</sub> cocatalysts and the high dispersion of catalyst on the heterostructure, in accordance with other results in the literature.<sup>33–35</sup>

**2.3. Transmission Electron Microscopy.** Transmission electron microscopy images of synthesized  $\alpha\text{-Fe}_2\text{O}_3$  display nanometer sized particles with high monodispersity and an average diameter of  $120 \pm 24$  nm (Figure 2(a) and 2(b)). The high-resolution TEM (HRTEM) image reveals lattice spacing of 0.36 and 0.27 nm, referring to (012) and (104) crystal planes, respectively (Figure 2(c)).<sup>36</sup> The observed values match with the PXRD pattern of the standard American Mineralogist Crystal Structure Database (AMCSD card no. 0000143) for rhombohedral (hexagonal)  $\alpha\text{-Fe}_2\text{O}_3$  with lattice constants of  $a = b = 0.503$  nm and  $c = 1.377$  nm (Figure 2(d)). The characteristic peaks for iron (Fe), oxygen (O), and carbon (C) were detected in the EDS spectrum of  $\alpha\text{-Fe}_2\text{O}_3$  (Figure 2(e)).<sup>37</sup> We have attributed the trace amounts of chloride to the KCl used for the shape-controlled synthesis.

The TEM image of SnS reveals the characteristic formation of interconnected flower-like structures (Figure 3(a)).<sup>38</sup> Each nanoflower consists of thin 2D nanosheets with sharp edges growing anisotropically in all directions (Figure 3(b)).<sup>39</sup> The HRTEM image reveals lattice spacing of 0.34 nm ascribed to the (012) crystal plane (Figure 3(c)).<sup>40</sup> The observed values match with the PXRD pattern of the standard American Mineralogist Crystal Structure Database assigned to the orthorhombic phase of SnS with lattice parameters  $a = 0.433$ ,  $b = 1.11$ , and  $c = 0.398$  nm (AMCSD card no. 0018115) (Figure 3(d)). The EDS spectra of SnS exhibit the signals for C, Sn, and S atoms (Figure 3(e)). The presented data reveal an Sn/S atomic ratio of 70.3, indicating the synthesized SnS catalyst deviated from stoichiometry. This high sulfur deficiency is due to postcalcination of SnS which results in significant evaporation of sulfur due to its high volatility.<sup>41</sup> Banu et al. reported the atomic ratio of Sn/S = 21.97 after annealing the prepared SnS thin films at 500 °C for 30 min.<sup>42</sup>

The TEM image of the  $\alpha\text{-Fe}_2\text{O}_3$  and SnS heterostructure (FeOSnS) shows that  $\alpha\text{-Fe}_2\text{O}_3$  is dispersed onto the surface of SnS nanoflowers with close contact between the two components (Figure 4(a)). This is favored for forming a heterostructural interface through interactions between the two photocatalysts, which can promote effective charge separation.<sup>43,44</sup> Furthermore,  $\alpha\text{-Fe}_2\text{O}_3$  and SnS are shown in the TEM images to retain their original shape without morphological changes after hybridization.<sup>45</sup> The TEM image of FeOSnS2-PdO<sub>x</sub> 2% shows the morphology changes after loading PdO<sub>x</sub>.



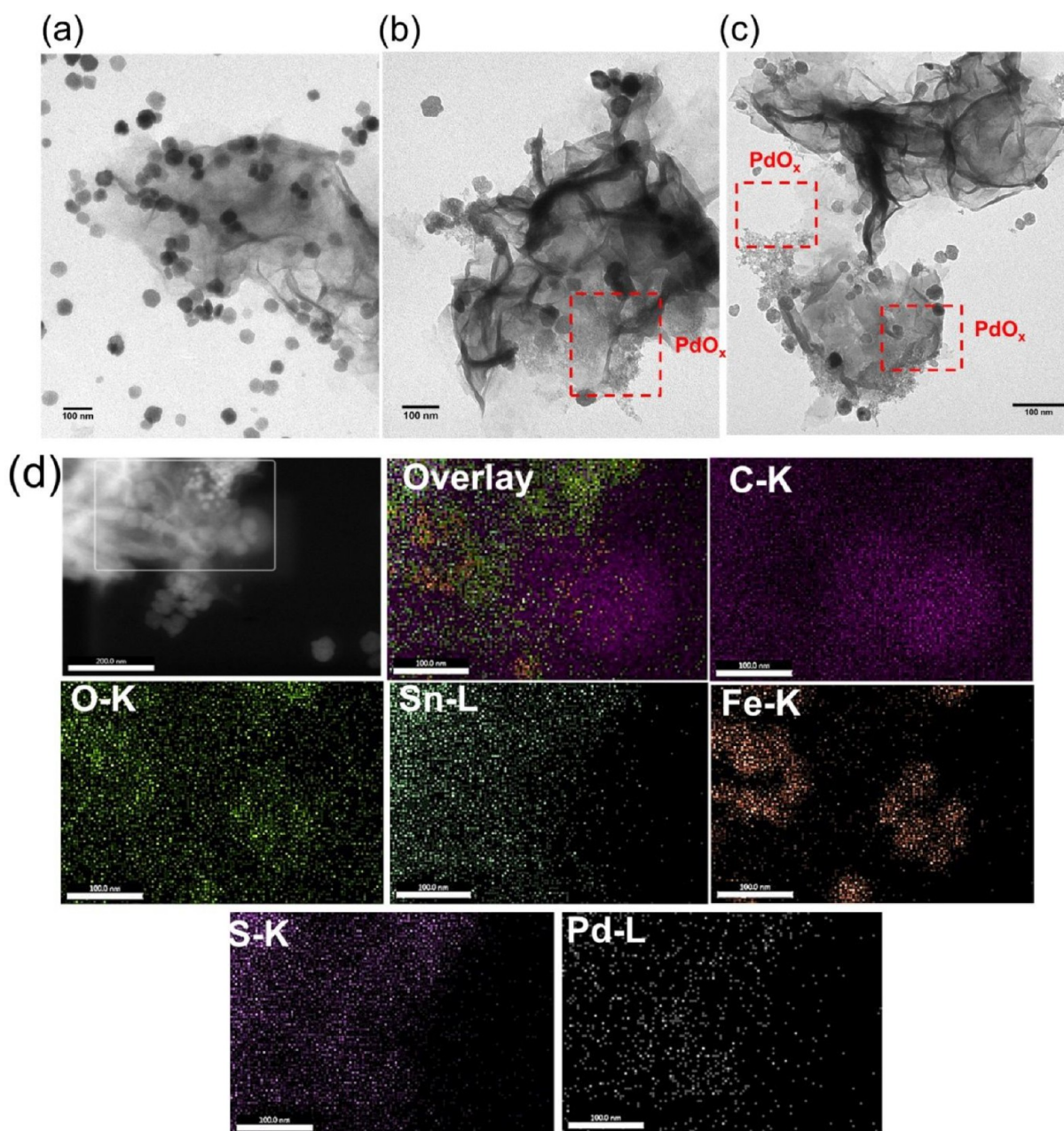
**Figure 3.** (a)–(c) TEM and HRTEM images, (d) PXRD pattern (a pattern for SnS (AMCSD 0018115) is shown for comparison), and (e) EDS spectra of the SnS catalyst.

nanoparticles, in which the PdO<sub>x</sub> nanoparticles are observed as black dots, dispersed on the surface of the heterocatalyst (Figure 4(b)). The size distribution profile shows an average size of 3.7 ± 1 nm (Figure S1). The TEM image of FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% exhibits the same morphology as FeOSnS2-PdO<sub>x</sub> 2% with PdO<sub>x</sub> nanoparticles seen as black dots, with an average size of 2.8 ± 0.87 nm (Figures 4(c) and S1). Notably, the MnO<sub>x</sub> cocatalyst is not localized due to the low concentration. Scanning TEM (STEM) imaging using a High-Angle Annular Dark Field (HAADF) detector and an Energy-Dispersive Spectroscopy (EDS) elemental mapping evidenced the presence of C, O, Fe, Sn, S, and Pd elements without any signal for Mn in the FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalyst (Figure 4(d)).

**2.4. Heterostructure Formation.** The heterostructure formation was further studied using the Brunauer–Emmett–Teller (BET) method. We calculated the specific surface area ( $S_{\text{BET}}$ ) and pore volume of the synthesized catalysts from adsorption–desorption isotherms. SnS presented  $S_{\text{BET}}$  and pore volume values of 60.9 m<sup>2</sup> g<sup>-1</sup> and 0.063 cc/g which were higher than that of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> with  $S_{\text{BET}}$  and pore volume values of 24.6 m<sup>2</sup> g<sup>-1</sup> and 0.031 cc/g, respectively (Figure 5(a)). The higher surface area of the SnS catalyst is due to its hierarchical porous architecture, whereas in the case of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, a small surface area and low pore volume indicate that the  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> primary crystals are densely packed as evidenced by TEM results.<sup>46</sup> Integration

of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> with SnS resulted in FeOSnS heterocatalysts (FeOSnS1–3) with reduced specific surface areas and pore volumes (Figure 5(a) and Table S1). This is due to the decrease in the pore volume of SnS or the blocking of the porous channels of SnS when coupling with  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, as reported by Yousatit et al.<sup>47</sup> The pore volume and diameter values for FeOSnS1–3 catalysts were not consistent with the initial ratios (Table S1), indicating that  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> nanoparticles are not uniformly distributed on SnS which agrees with the TEM results. Furthermore, loading of the cocatalyst reduced the surface area and pore volume of the FeOSnS2 heterocatalyst from 24.9 m<sup>2</sup>/g and 0.008 cc/g down to 15.6 m<sup>2</sup>/g and 0.005 cc/g for FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% (Figure 5(b) and Table S1). This phenomenon is attributed to the partial surface coverage of the heterocatalyst by the deposited cocatalysts.<sup>48</sup>

The heterostructure formation was further confirmed by attenuated total reflection-Fourier transform infrared (ATR-FTIR) and UV–vis diffuse reflectance spectroscopy (UV–vis DRS) results (see Figures S2 and S3 in the Supporting Information). XRD and FT-IR studies could not confirm the presence of the cocatalysts in the FeOSnS heterostructures, but EDS analysis was able to. Unfortunately, as the detection limit of EDS used in this study was <1 wt %, the technique was not able to give accurate quantitative values (see Figure S4 in the Supporting Information). Atomic absorption spectroscopy

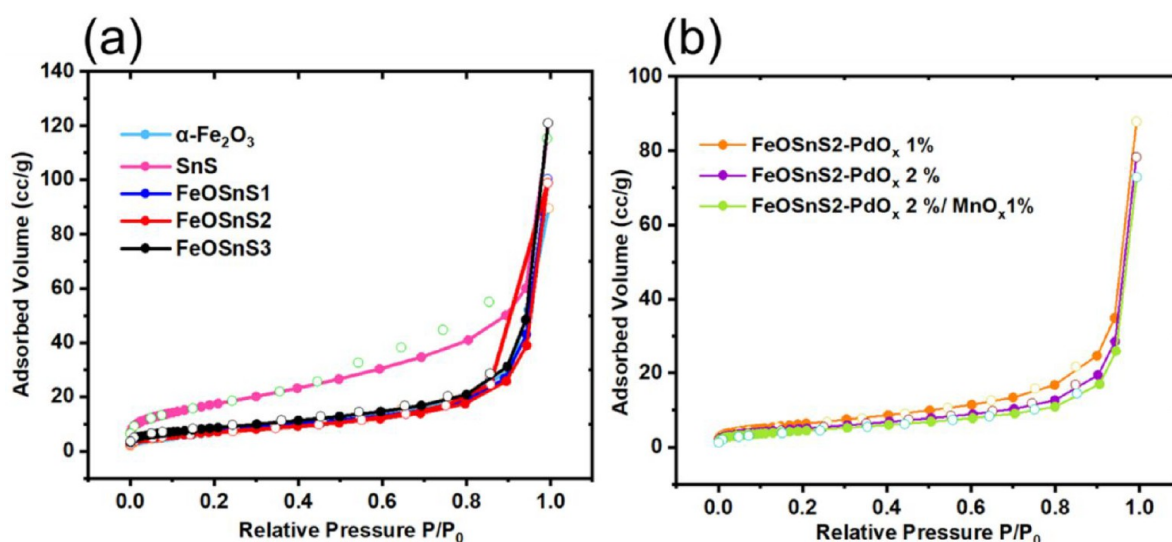


**Figure 4.** (a)–(c) TEM images of FeOSnS<sub>2</sub>, FeOSnS<sub>2</sub>-PdO<sub>x</sub> 2%, and FeOSnS<sub>2</sub>-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalysts. (d) HAADF-STEM and EDX elemental mapping images of FeOSnS<sub>2</sub>-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalyst.

(AAS) was therefore utilized to confirm the content of the cocatalyst in the synthesized FeOSnS<sub>2</sub> heterocatalyst (see Table S2 in the Supporting Information).

**2.5. X-ray Photoelectron Spectroscopy.** To further explore the presence of the cocatalysts and identify the related oxidation states, XPS was performed. Wide-scan survey spectra revealed photoelectron lines at binding energies (BEs) of 23.6 eV (Sn 4d), 153–175 eV (S 2p), 278–300 eV (C 1s), 328–348 eV (Pd 3d), 480–505 eV (Sn 3d), 520–545 eV (O 1s), 556–630 eV (Mn 2p), and 697–745 eV (Fe 2p), at the surface ( $\leq 3$  nm), as illustrated in Figure 6a. This is consistent with the EDS results and revealed the successful impregnation of PdO<sub>x</sub> and MnO<sub>x</sub> cocatalysts on FeOSnS<sub>2</sub>. The high-resolution XPS window of the C 1s core level (Figure 6(b)) is deconvoluted into three peaks at BEs of 284.8, 285.7, and 288.3 eV, respectively, corresponding to C–C sp<sup>2</sup> from PVP molecules

and adventitious carbon, a C–OH bond, and a carbonyl (–OC=O) bond, respectively.<sup>49</sup> The O 1s core-level spectra (Figure 6(c)) exhibit a high intensity peak at  $\sim 530.3$  eV corresponding to the lattice oxygen of iron–oxygen bonds (Fe–O) of the  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> component in the heterocatalyst. The high intensity of the peak is due to the high number of iron ions (Fe<sup>3+</sup>) strongly interacting with the lattice oxygen (O<sup>2-</sup>) in the crystal lattice. The peak centered at 532.2 eV is associated with the surface –OH groups (adsorbed water) and oxygen bonded with the PVP C atoms (Fe–O–C bond).<sup>50</sup> As can be seen in the wide-scan survey spectra and high-resolution S 2p photoelectron spectra (Figure 6(d)), the peaks at 161–162 and 162–163 eV corresponding to S 2p<sub>3/2</sub> and S 2p<sub>1/2</sub> are absent. This is due to sulfur being present in amounts lower than the detection limit for XPS,<sup>51</sup> which is in accordance with EDS and elemental mapping results.<sup>52,53</sup> Sn manifests itself with strong peaks



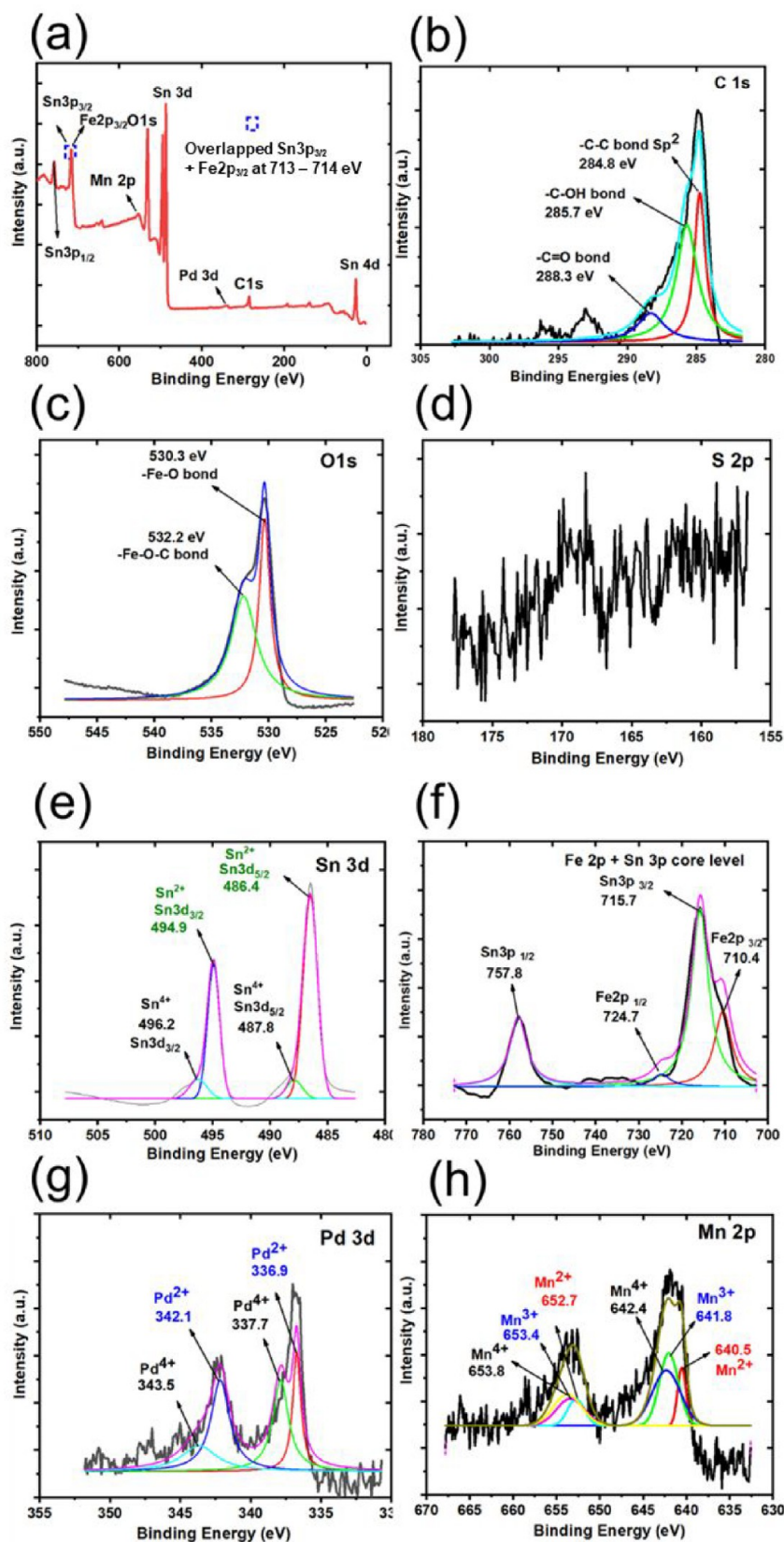
**Figure 5.** (a,b) N<sub>2</sub> adsorption–desorption isotherm of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, SnS, FeOSnS1–3, and FeOSnS2-PdO<sub>x</sub>/MnO<sub>x</sub> catalysts measured at 77 K. Filled circles = adsorption points; empty circles = desorption points.

centered at 24 eV (Sn 4d), 486.4 eV (Sn 3d<sub>5/2</sub>), 494.9 eV (Sn 3d<sub>3/2</sub>), 713.6 eV (Sn 3p<sub>3/2</sub>), and 755.2 eV (Sn 3p<sub>1/2</sub>) (Figure 6(a)). The deconvolution of the Sn 3d energy state (Figure 6(e)) revealed spin–orbit doublet peaks at 486.4 eV (Sn 3d<sub>5/2</sub>) and 494.9 eV (Sn 3d<sub>3/2</sub>) with a separation of 8.5 eV which confirms the oxidation state of +2 and the formation of single-phase SnS. In addition, two small peaks at slightly higher BEs of 487.8 (Sn 3d<sub>5/2</sub>) and 496.2 eV (Sn 3d<sub>3/2</sub>) are assigned to Sn<sup>4+</sup>, indicating the partial oxidation of the edges of SnS due to the unstable state of Sn<sup>2+</sup> at the annealing temperature (500 °C).<sup>54</sup> Overall, XPS analysis indicates SnS as the major phase and SnS<sub>2</sub> as an impurity in the FeOSnS2/PdO<sub>x</sub>/MnO<sub>x</sub> catalyst. The high-resolution Fe 2p spectrum in Figure 6(f) exhibits the BEs of Fe 2p<sub>3/2</sub> and Fe 2p<sub>1/2</sub> at 710.9 and 724.5 eV with additional satellite peaks at 734.7 eV. Taken together, the data verify the valence state of Fe ions as +3 in  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>. The result indicates that calcination at 500 °C does not change the valence state of Fe<sup>3+</sup> to reduced or oxidized forms (e.g., Fe<sup>2+</sup> or Fe<sup>4+</sup>). Notably, as marked in the dotted square in the low-resolution survey spectrum (Figure 6(a)), the core-level lines assigned to Sn 3p located at BEs of 715.6 eV for Sn 3p<sub>3/2</sub> and 757.7 eV for Sn 3p<sub>1/2</sub> overlap with the Fe 2p core levels.<sup>55</sup> The core-level high-resolution orbital scan of the Pd 3d spectra were deconvoluted into four peaks centered at BEs of 336.9, 335.4, 342.4, and 343.5 eV, respectively (Figure 6(g)). The high intensity peaks at BEs of 336.9 (3d<sub>5/2</sub>) and 342.4 eV (3d<sub>3/2</sub>) are assigned to Pd<sup>2+</sup> from the palladium precursor (PdCl<sub>2</sub>) or PdO, which is in agreement with the reported value of the BE at 336.8–337.4 eV.<sup>56</sup> The very low intensity peaks at BEs of 337.7 and 343.5 eV are attributed to Pd<sup>4+</sup>, suggesting that a low percentage of Pd<sup>2+</sup> has been oxidized during the impregnation/calcination process.<sup>57</sup> There was no metallic Pd (Pd<sup>0</sup>) detected on the surface of the catalyst with typical BEs of 335.1 and 341.1 eV.<sup>58</sup> The high-resolution spectrum of Mn 2p exhibited two main peaks at BEs of 641.8 and 653.3 eV assigned to Mn 2p<sub>3/2</sub> and Mn 2p<sub>1/2</sub>, respectively. The spin–orbit splitting energy value of Mn 2p was 11.5 eV, close to 11.6 eV as reported by Yu et al.<sup>59</sup> The peaks were deconvoluted to identify the chemical valence of the MnO<sub>x</sub> cocatalyst. As can be seen from Figure 6(h), the Mn 2p<sub>1/2</sub> and Mn 2p<sub>3/2</sub> peaks can be resolved into two pairs of triplet peaks, respectively. The Mn 2p<sub>3/2</sub> peaks at BEs of 640.6, 641.8, and 642.4 eV correspond to

Mn<sup>2+</sup> (MnO), Mn<sup>3+</sup> (Mn<sub>2</sub>O<sub>3</sub>), and Mn<sup>4+</sup> cations (MnO<sub>2</sub>), respectively. Similarly, the Mn 2p<sub>1/2</sub> peaks appeared at BEs of 652.4, 653.3, and 653.8 eV which correspond to Mn<sup>2+</sup>, Mn<sup>3+</sup>, and Mn<sup>4+</sup> cations, respectively. These results indicate the coexistence of MnO, Mn<sub>2</sub>O<sub>3</sub>, and MnO<sub>2</sub> species. The loaded manganese oxide is labeled as MnO<sub>x</sub> (1 < x < 2).<sup>60</sup>

**2.6. Photocatalytic Reforming of Ethanol.** Our initial investigations of the photocatalytic performance of as-synthesized  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>, SnS, and FeOSnS1–3 heterocatalysts focused on their ability to act as water-splitting catalysts. While FeOSnS2 was found to have the optimum photocatalytic activity, H<sub>2</sub> generation was low over all samples (Figure S7 in the Supporting Information). Therefore, our efforts turned to the photocatalytic reforming of ethanol using the cocatalysts. To explore the effect of the cocatalysts on the photocatalytic activity, single PdO<sub>x</sub> and dual PdO<sub>x</sub>/MnO<sub>x</sub> cocatalysts were loaded on FeOSnS2, and studies of the photocatalytic reforming of ethanol in ethanol/water solutions compared to bare FeOSnS2 were performed. The results are shown in Figure 7 and Table 1. A blank test was carried out without photoirradiation in the presence of both ethanol and photocatalyst (entry 1), resulting in negligible photocatalytic production rates. This confirms that this reaction does proceed photocatalytically with photoirradiation in the presence of the photocatalyst. The photocatalytic reforming of ethanol using ethanol/water solutions with single and dual cocatalysts loaded on FeOSnS2 resulted in the production of H<sub>2</sub>, with only trace amounts of CH<sub>4</sub> produced which is consistent with the results presented in the literature for both single and dual cocatalyst systems.<sup>61</sup> Product selectivity of H<sub>2</sub> among other possible gases reached 99% (Table 1), with the formation of other oxygenated organic compounds unable to be detected; therefore, this method produces very high levels of pure H<sub>2</sub> gas.

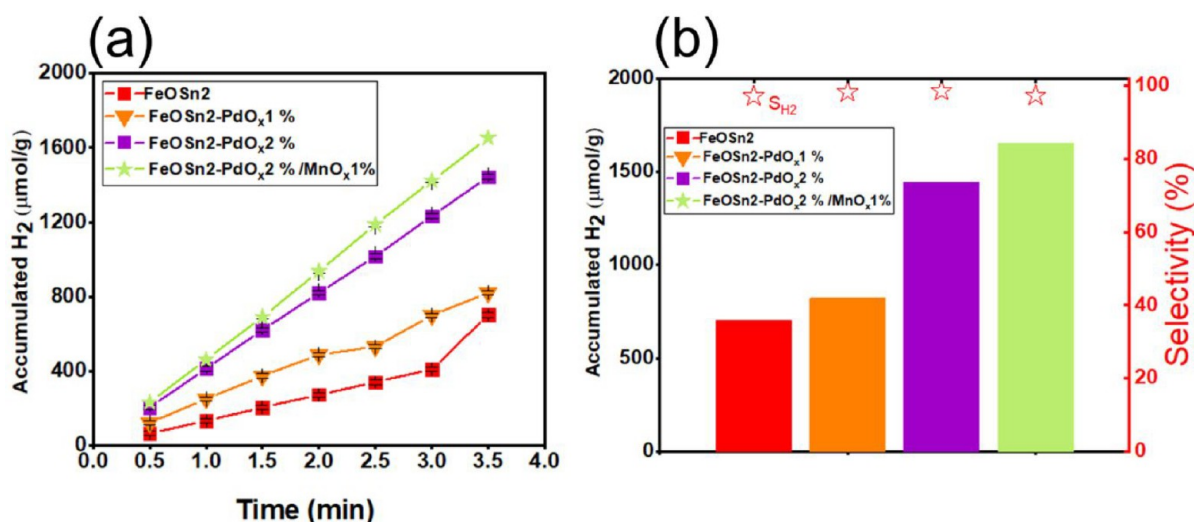
Time course data for H<sub>2</sub> formation in the photocatalytic reforming of ethanol over the FeOSnS2 and FeOSnS2/PdO<sub>x</sub>/MnO<sub>x</sub> catalysts are shown in Figure 7(a). An overall increase in H<sub>2</sub> formation yield with time of illumination was observed for all samples, but a low amount of H<sub>2</sub> (42  $\mu$ mol/g) was produced from the noncatalyzed irradiation of ethanol after 3.5 h of illumination (Table 1). The FeOSnS2 sample produced a high amount of H<sub>2</sub> (702  $\mu$ mol/g) after 3.5 h of illumination,



**Figure 6.** (a) XPS survey spectra and high-resolution XPS spectra of (b) C 1s, (c) O 1s, (d) S 2p, (e) Sn 3d, (f) Fe 2p, (g) Pd 3d, and (h) Mn 2p for FeOSnS<sub>2</sub>-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalyst.

indicating the successful trapping of  $h^+$  by ethanol during the reaction (Figure 7b).<sup>62</sup> The consumption of the photogenerated holes by ethanol results in the accumulation of photogenerated

electrons on the photocatalyst surface, leading to a higher H<sub>2</sub> formation.<sup>34</sup> Our prepared FeOSnS<sub>2</sub> heterocatalyst presents better performance than previously reported photocatalysts



**Figure 7.** (a) Time course and (b) accumulated  $\text{H}_2$  evolution amount from ethanol photoreforming over 1 g of  $\text{FeOSnS}_2\text{-PdO}_x/\text{MnO}_x$  catalysts after a 3.5 h illumination (source: mercury lamp ( $4.4 \text{ mW cm}^{-2}$  measured at  $\lambda = 254 \pm 10 \text{ nm}$ )).

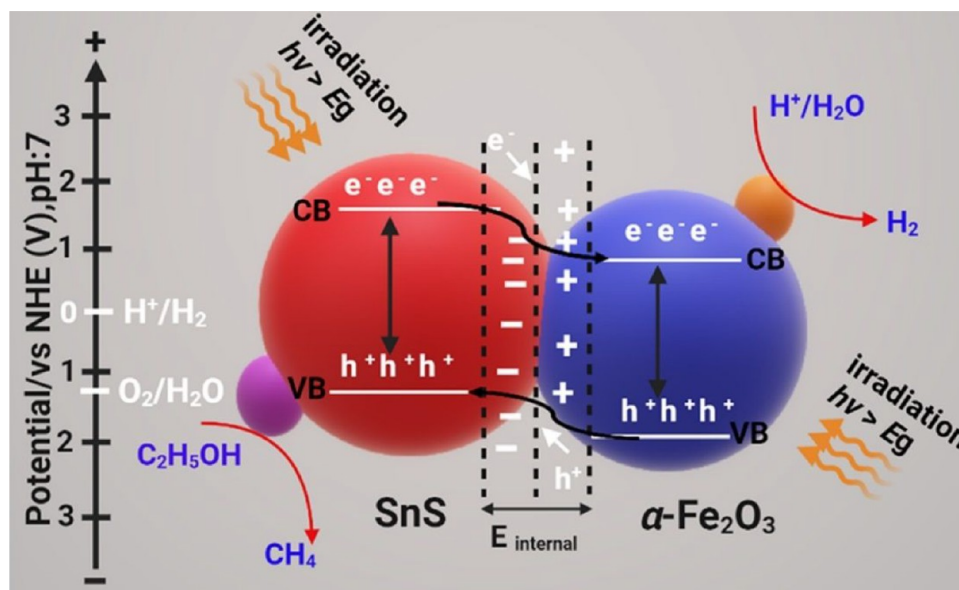
**Table 1. Product Yields for the Photocatalytic Reforming by Photocatalysts**

Entry	Gas	Photocatalyst	Production rate ( $\mu\text{mol/g}$ )				
			$\text{H}_2$	$\text{CH}_4$	$\text{CO}$	$\text{CO}_2$	$\text{S}_{\text{H}_2}$
	Ar	No light	-	-	-	-	-
1	Ar	$\text{FeOSnS}_2$	702	1.75	-	-	99.75
2	Ar	$\text{FeOSnS}_2\text{-PdO}_x$ 1%	821	4.75	-	-	99.42
3	Ar	$\text{FeOSnS}_2\text{-PdO}_x$ 2%	1444	5.5	-	-	99.62
4	Ar	$\text{FeOSnS}_2\text{-PdO}_x$ 2%/ $\text{MnO}_x$ 1%	1654	11	-	-	99.33

consisting of  $\alpha\text{-Fe}_2\text{O}_3$  modified with other metal sulfides. For example, Lu et al. reported an HE rate of  $6.9 \mu\text{mol/g/h}$  over Zn-doped  $\alpha\text{-Fe}_2\text{O}_3$ -modified  $\text{WS}_2$  after 2 h solar light irradiation.<sup>63</sup> Kadam et al. reported the highest HE rate of  $136 \mu\text{mol/g/h}$  for a 10% Mo-doped SnS photocatalyst at 400 nm.<sup>64</sup> When  $\text{PdO}_x$  species were loaded as cocatalysts on  $\text{FeOSnS}_2$ ,  $\text{H}_2$  formation

yield was increased from  $702 \mu\text{mol/g}$  for  $\text{FeOSnS}_2$  to  $821 \mu\text{mol/g}$  for  $\text{FeOSnS}_2\text{-PdO}_x$  1% and  $1444 \mu\text{mol/g}$  for  $\text{FeOSnS}_2\text{-PdO}_x$  2% after 3.5 h of illumination. The high  $\text{H}_2$  formation yield for  $\text{FeOSnS}_2\text{-PdO}_x$  2% with a higher loading amount of  $\text{PdO}_x$  demonstrates that the photogenerated electrons and holes could be efficiently separated once  $\text{PdO}_x$  was loaded on the surface of  $\text{FeOSnS}_2$  by the impregnation method. The loading of the  $\text{MnO}_x$  species as an additional cocatalyst on  $\text{FeOSnS}_2\text{-PdO}_x$  2% can increase the  $\text{H}_2$  formation yield from  $1444 \mu\text{mol/g}$  for  $\text{FeOSnS}_2\text{-PdO}_x$  2% to  $1654 \mu\text{mol/g}$  for the  $\text{FeOSnS}_2\text{-PdO}_x$  2%/  $\text{MnO}_x$  1% photocatalyst under the same conditions. The formation of  $\text{PdO}_x$  and  $\text{MnO}_x$  cocatalysts was evidenced by XPS and UV-vis DRS analysis, confirming the role of dual cocatalysts of  $\text{PdO}_x/\text{MnO}_x$  in photocatalytic reforming of ethanol, where the  $\text{PdO}_x$  serves as an electron trap (catalytic sites for  $\text{H}_2$  reduction) and  $\text{MnO}_x$  serves as a hole trap (catalytic sites for oxidation reaction). This vicinal charge separation by dual cocatalysts of  $\text{PdO}_x/\text{MnO}_x$  can also lead to efficient

**Scheme 1. Band Bending and Carrier Transport at the Surface or Interface for  $\text{FeOSnS}_2/\text{PdO}_x/\text{MnO}_x$  Catalysts**





photocatalytic reforming of ethanol. Since reduction and oxidation reactions take place in a pair, any improvements in the oxidation would result in improved reduction and subsequent higher H<sub>2</sub> formation rates.<sup>65</sup> This indicates that the coexistence of reductive and oxidative cocatalysts could synergistically improve the photocatalytic potential. López-Martínez et al. observed the same phenomenon where the maximum HE rate of 728 μmol/m<sup>2</sup> was achieved over dual AuPd cocatalysts loaded onto SnS after 3 h irradiation (Xe lamp of 450 W and 100 mW/cm<sup>2</sup>).<sup>66</sup>

**2.7. Proposed Mechanism for H<sub>2</sub> Production.** The possible charge separation at the interface of the PdO<sub>x</sub>/MnO<sub>x</sub> dual cocatalyst loaded on FeOSnS2 for the photocatalytic reforming of ethanol is depicted in Scheme 1. Before contact, the conduction band and valence band edge positions ( $E_{VB}$  and  $E_{CB}$ ) of SnS and  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> catalysts are independent of each other and were calculated to be 2.89 and 0.96 eV for  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and 1.04 and -0.40 eV for SnS (Supporting Information). The coupling of p-type SnS and n-type Fe<sub>2</sub>O<sub>3</sub> forms a p-n junction at the heterostructure interfaces, which leads to the formation of an internal electric field ( $E_{internal}$ ) between the p-type side and the n-type side and band bending.<sup>67</sup> At the p-n junction the electrons and holes are separated, and enhancement of the photo-generated electrons transferring from the CB of p-type SnS to the CB of n-type  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> (Scheme 1) occurs. Loading PdO<sub>x</sub> and MnO<sub>x</sub> on the FeOSnS2 surface improves the interface reaction speed because the cocatalysts can reduce the activation energy of the reduction and oxidation reaction for the photocatalytic reforming of ethanol. The TEM images revealed the PdO<sub>x</sub> NPs were deposited on the surface of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and SnS. However, we could not determine the exact position of MnO<sub>x</sub> on the surface of FeOSnS2, and it seems that MnO<sub>x</sub> is randomly distributed on the surface of the FeOSnS2 catalyst. Considering the improved HE rate for the dual cocatalyst, one possible explanation may be that the loaded cocatalysts are sufficiently spatially separated that the recombination of electron/hole pairs is minimized. Therefore, PdO<sub>x</sub> functions as the reduction site and receives the photoexcited electrons from the CB of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> to produce H<sub>2</sub>, while MnO<sub>x</sub> functions as the oxidation site and receives the photoexcited holes from the CB of SnS to produce CH<sub>4</sub>.

### 3. CONCLUSION

We have successfully synthesized FeOSnS heterocatalysts using a simple *in situ* chemical precipitation method. The prepared FeOSnS2 heterocatalyst exhibits excellent HE performance relative to pristine  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and SnS, due to the new heterostructural interface provided by this technique. Loading of redox PdO<sub>x</sub> and MnO<sub>x</sub> cocatalysts did not influence the crystal structure of the FeOSnS2 heterocatalyst, but they were partially oxidized during loading. The loaded cocatalysts masked the pores of the FeOSnS2 heterocatalyst, resulting in the reduction of the active surface area ( $S_{BET}$ ). HE results revealed the success of the undertaken strategies, namely, the integration of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> and SnS and coloading of redox PdO<sub>x</sub> and MnO<sub>x</sub> cocatalysts. The highest HE rate was achieved for FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1%, ascribed to the synergistic redox contribution of the PdO<sub>x</sub> and MnO<sub>x</sub> species. In order to increase the HE rate further, two strategies will be considered in future work including (i) the loading of cocatalysts using different methods onto the optimal heterocatalyst and (ii) altering the ratios of the loaded PdO<sub>x</sub> and MnO<sub>x</sub> cocatalysts.

## 4. EXPERIMENTAL SECTION

**4.1. Materials.** Iron(III) acetylacetonate (Fe(acac)<sub>3</sub> ≥ 99.9% trace metals basis), poly(vinylpyrrolidone) (PVP, average molecular weight 40,000 g mol<sup>-1</sup>), *N,N*-dimethylformamide (DMF), sulfur powder (99.99%), and tin chloride dihydrate (SnCl<sub>2</sub>·2H<sub>2</sub>O, 99%), were purchased from Sigma-Aldrich. All other chemicals were of analytical grade and used as received from commercial sources without further purification.

**4.2. Synthesis of the  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> Catalyst.** The  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> catalyst was synthesized through our previously published hydrothermal route.<sup>68</sup>

**4.3. Synthesis of the SnS Catalyst.** The flower-like SnS catalyst was synthesized by the above hydrothermal method, except SnCl<sub>2</sub> (0.2 mmol), PVP (0.4 mmol), and sulfur powder (0.1 mmol) precursors were mixed in DMF (20 mL) under stirring (500 rpm) at 70 °C for 120 min followed by hydrothermal reaction at 180 °C for 6 h and an eventual washing process as above.

**4.4. Synthesis of  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>-SnS (FeOSnS) Heterocatalysts.** The *in situ* chemical precipitation method was utilized to synthesize FeOSnS heterocatalysts using different mass ratios of as-prepared SnS and  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> constituents to find the optimal formulation for coating with cocatalysts in the next step. In a typical experiment, SnS: $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> (1:1 w/w) was dispersed by ultrasonication in 50 mL of EtOH/H<sub>2</sub>O solution (20:80 v/v) for 30 min and stirred in a fume hood at 80 °C for 24 h. The resultant product was collected by centrifugation, rinsed with distilled water and absolute ethanol three times, and dried at 80 °C for 6 h in an electric oven. The obtained heterocatalyst was denoted as FeOSnS1. Similarly, FeOSnS2 (SnS: $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> = 1.5:0.5 w/w) and FeOSnS3 (SnS: $\alpha$ -Fe<sub>2</sub>O<sub>3</sub> = 0.5:1.5 w/w) were also prepared following the above procedure. The prepared FeOSnS1-3 heterocatalysts were then placed inside ceramic crucibles and calcined under Ar(g) at 500 °C for 5 h.

**4.5. Loading of PdO<sub>x</sub>/MnO<sub>x</sub> Cocatalysts onto FeOSnS2.** An impregnation method was utilized to load cocatalysts on the FeOSnS heterocatalyst. To synthesize FeOSnS2-PdO<sub>x</sub> 1%, 0.5 g of FeOSnS2 was soaked in a solution (100 mL) containing the Pd precursor (0.008 g PdCl<sub>2</sub>) and continuously stirred in a water bath at 120 rpm at 80 °C until the evaporation of the water from the suspension. The resulting solid was washed with absolute ethanol and distilled water three times and dried at 70 °C in an electric oven overnight followed by a calcination on a ceramic crucible in air at 350 °C for 2 h. FeOSnS2-PdO<sub>x</sub> 2% and FeOSnS2-PdO<sub>x</sub> 2%/MnO<sub>x</sub> 1% catalysts were prepared using the same procedure with stoichiometric amounts of Pd (0.016 g of PdCl<sub>2</sub>) and MnO<sub>x</sub> precursors (0.016 g of PdCl<sub>2</sub> and 0.027 g of Mn(NO<sub>3</sub>)<sub>2</sub>·4H<sub>2</sub>O).

**4.6. Ethanol Reforming for Hydrogen Production.** The photocatalytic H<sub>2</sub> production was commenced by irradiation of the solution using a 100 W high-pressure mercury lamp (4.4 mW cm<sup>-2</sup> measured at  $\lambda = 254 \pm 10$  nm). The amount of H<sub>2</sub> in the outlet gas was quantified by an online gas chromatograph (Shimadzu, GC-8A, TCD, Shincarbon ST column, argon carrier). The experiment was carried out using a 20:80 ethanol-water solution where ethanol was used as the sacrificial reagent to consume the photogenerated holes during the reaction. The selectivity toward H<sub>2</sub> evolution compared with CH<sub>4</sub>,  $S_{H_2}$ , was calculated using the formula:

$$S_{H_2} (\%) = R_{H_2} / (R_{H_2} + R_{CH_4}) \times 100$$

where  $R_{\text{H}_2}$  and  $R_{\text{CH}_4}$  describe the production rate of  $\text{H}_2$  and  $\text{CH}_4$ , respectively. The amount of CO and  $\text{CO}_2$  produced was negligible and not considered in this equation.

## 5. CHARACTERIZATION

Powder X-ray diffraction (PXRD) patterns were recorded on a Rigaku Spider X-ray diffractometer with  $\text{Cu K}\alpha$  radiation ( $\lambda = 1.5406 \text{ \AA}$ ), at 40 kV and 50 mA from  $10^\circ$  to  $80^\circ$  in the Bragg configuration. The morphology of the catalysts was examined using transmission electron microscopy (TEM, Tecnai G2 Spirit Bio-TWIN, acceleration voltage of 200 Kv) and scanning electron microscopy (SEM, FE-SEM FEI Quanta 200). ImageJ software was acquired for postprocessing and particle size analysis. The elemental mapping images and energy-dispersive X-ray spectroscopy (EDX) spectra of catalysts were probed by an energy-dispersive spectrometer connected to field-emission scanning electron microscopy (FE-SEM FEI Quanta 200) to verify the chemical composition. The spectral data were collected with a silicon EDAX unit (NJ, USA) running Genesis Spectrum software (version 5.21). The EDS spectra were generated for each catalyst by a line scan of several randomly selected areas under different magnifications.

The chemical state of catalysts was probed with X-ray photoelectron spectroscopy (XPS) (Kratos Axis Ultra<sup>DLD</sup> X-ray Photoelectron Spectrometer) at the pressure of  $10^{-9}$  Torr with Al  $\text{K}\alpha$  X-rays (1486.69 eV) as the X-ray source. All spectra were calibrated with respect to the C 1s signal from adventitious hydrocarbon set at 285 eV. The high-resolution narrow-scan XPS spectra included nonlinear (Shirley) background subtraction and peak deconvolution by using mixed Gaussian–Lorentzian functions.

The mass percentages of  $\text{PdO}_x$  and  $\text{MnO}_x$  cocatalysts were analyzed by atomic absorption spectroscopy (AAS; GBC Scientific PerkinElmer Instrument) with a slit width of 0.2 nm, a lamp current of 5 mA, and an air-acetylene flame, with a Pd hollow cathode lamp (340.5 nm) and Mn hollow cathode lamp (403.1 nm) for  $\text{PdO}_x$  and  $\text{MnO}_x$ , respectively. For AAS analysis, approximately 5 mg of catalyst was digested in a mixture of concentrated HCl (37%, 0.5 mL),  $\text{HNO}_3$  (70%, 1.0 mL), and  $\text{H}_2\text{O}_2$  (32%, 1.0 mL) at  $100^\circ\text{C}$  overnight followed by dilution to 50 mL with Milli-Q water. Standard 1000 ppm Pd in HCl and Mn in  $\text{HNO}_3$  (Sigma-Aldrich) solutions were used to prepare metal calibration standard solutions (10–100 ppm) from which the intensity vs concentration plot was acquired for quantification.<sup>69</sup> The  $\text{N}_2$  adsorption–desorption isotherms were analyzed on a Quantachrome BELSORP Mini Autosorb nitrogen-adsorption apparatus. Samples were evacuated at  $120^\circ\text{C}$  for 20 h prior to Brunauer–Emmett–Teller (BET) measurements with autosorb at liquid nitrogen temperature (77 K). The BET specific surface area ( $S_{\text{BET}}$ ) of the catalysts was investigated by a multipoint BET method utilizing the adsorption data in the relative pressure ( $P/P_0$ ) range of 0–1.

The optical characteristics and band gap values of the catalysts were acquired via a UV–vis diffuse reflectance spectrometer (UV–vis DRS; V-640 (JASCO)) with an integrating sphere attachment in the wavelength range of 200–800 nm at RT. High-purity barium sulfate ( $\text{BaSO}_4$ ) powder was used as a reflectance standard. Photoluminescence (PL) emission spectra were collected at RT using a Horiba Scientific Fluoromax-4 Spectrofluorometer. All samples with concentrations of 0.06 mg/mL in EtOH were excited at 330 nm with a 150 W ozone-free xenon arc lamp as the excitation source, and the emission spectra were recorded from 330 to 620 nm.

## ■ ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acsomega.2c05410>.

TEM images and corresponding  $\text{PdO}_x$  particle size distribution for  $\text{FeOSnS}_2\text{-PdO}_x$  2% and  $\text{FeOSnS}_2\text{-PdO}_x$  2%/MnO<sub>x</sub> 1%, ATR-FTIR results, table of physicochemical properties, UV–vis DRS and EDS spectra, table of wt % of the heterocatalysts, PL spectra, hysteresis loops, photocatalytic water splitting, and accompanying explanations (PDF)

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### Author Contributions

H.E. designed and performed the synthetic experiments, collected characterization data, and wrote the draft manuscript. T.S. performed the  $\text{H}_2$  production experiments and contributed to the writing of the  $\text{H}_2$  production results. Y.Z. performed the BET and XRD measurements. S.G.T., H.Y., J.K.B., and P.G.P. provided an overall review of the manuscript. All authors read and approved the final manuscript.

### Notes

The authors declare no competing financial interest.

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