Sacrificial-template-free synthesis of core-shell C@Bi$_2$S$_3$ heterostructures for efficient supercapacitor and H$_2$ production applications

S. V. Prabhakar Vattikuti$^1$, Anil Kumar Reddy Police$^2$, Jaesool Shim$^1$ & Chan Byon$^2$

Core-shell heterostructures have attracted considerable attention owing to their unique properties and broad range of applications in lithium ion batteries, supercapacitors, and catalysis. Conversely, the effective synthesis of Bi$_2$S$_3$ nanorod core@amorphous carbon shell heterostructure remains an important challenge. In this study, C@Bi$_2$S$_3$ core-shell heterostructures with enhanced supercapacitor performance were synthesized via sacrificial-template-free one-pot-synthesis method. The highest specific capacities of the C@Bi$_2$S$_3$ core shell was 333.43 F g$^{-1}$ at a current density of 1 A g$^{-1}$. Core-shell-structured C@Bi$_2$S$_3$ exhibits 1.86 times higher photocatalytic H$_2$ production than the pristine Bi$_2$S$_3$ under simulated solar light irradiation. This core-shell feature of C@Bi$_2$S$_3$ provides efficient charge separation and transfer owing to the formed heterojunction and a short radial transfer path, thus efficiently diminishing the charge recombination; it also facilitates plenty of active sites for the hydrogen evolution reaction owing to its mesoporous nature. These outcomes will open opportunities for developing low-cost and noble-metal-free efficient electrode materials for water splitting and supercapacitor applications.

Active supercapacitors are a class of energy storage devices that have fascinated the scientific community owing to their high power capacity with long cycle life, low cost, and low maintenance. Recently, supercapacitors are being widely used in memory backup, consumer electronics, and industrial power management$^{1-3}$. In addition, supercapacitors are safer and have a lesser environmental impact than batteries$^{2,3}$. Mainly, the specific capacitance of a supercapacitor depends on the electrode materials and thus, electrode materials of supercapacitors play an important role. There are several electrode materials available for supercapacitors; the most common are carbon materials owing to their high conductivity, large specific surface area, different forms, and abundance. However, their activity is not sufficient, owing to their low specific energy. In addition, the inset surface of carbon materials makes it difficult for electrolytes to further penetrate the internal layers of carbon$^4$. Different approaches have been employed to overcome these issues like introduction of metal oxides, hetero atoms, and coupling of two or more materials.

In recent years, heterostructure nanomaterials comprising two or three different functionalities showed enhanced or different physicochemical performances compared to single functionality materials$^5$. Among the various heterostructure features, core-shell structures exhibit superior performances owing to the combined effects of cores and shells$^{6,7}$. Meanwhile, metal sulfides including nickel sulfides, cobalt sulfides, and bismuth disulfides are very important semiconductor materials and have been employed to develop supercapacitor electrodes$^{8-11}$. Among these, bismuth sulfide (Bi$_2$S$_3$) has a lamellar structure with a 1.3–1.7 eV direct band gap and these features offers potential applications in the field of thermoelectric cooler devices$^{12}$, lithium-ion batteries$^{13}$, and optoelectronic devices$^{14}$ owing to the possibility of band gap tuning with different sizes of the subcomponent.

$^1$School of Mechanical Engineering, Yeungnam University, Gyeongsan, 712-749, South Korea. $^2$School of Mechanical and Nuclear Engineering, Ulsan National Institute of Science and Technology (UNIST), Ulsan, 44919, Republic of Korea. Correspondence and requests for materials should be addressed to S.V.P.V. (email: drprabu@ynu.ac.kr) or J.S. (email: jshim@ynu.ac.kr) or C.B. (email: cbyon@unist.ac.kr)
As the above-mentioned properties strongly rely on the size, shape and morphologies, diverse strategies are put forth to controlled synthesis of bismuth sulfide nanostructured materials. Several Bi\(_2\)S\(_3\) nanomaterials like nanorods, tubes, wires, ribbons are prepared by various synthesis procedures. However, Bi\(_2\)S\(_3\) as an electrode material possesses low conductivity as well as high volume expansion during the charge and discharge cycling, which are major challenges. In addition, the reversible specific capacity fades rapidly during cycling reactions in metal sulfides. This capacitance loss is mainly because of the volume expansion during the charge and discharge processes causing particle cracks and pulverization that breakdown the electrical contacts in the bismuth sulfide anode. Hence, improving the number of charge discharge cycles is a crucial task to address for the optimization of Bi\(_2\)S\(_3\) based anode materials for supercapacitor applications.

To overcome these drawbacks, different methods have been followed to enhance its electrochemical activity; one of the most efficient ways is to design nanohybrids with a core-shell structure using the allotropes of carbon materials, which can act as a projective/buffer layer to improve the volume expansion during charge and discharge processes\(^{15,24}\). Recently, Yang et al.\(^{17}\) fabricated strongly coupled Bi\(_2\)S\(_3\) nanocrystals anchored on carbon nanotube (CNT) backbones; the resulting Bi\(_2\)S\(_3@\)CNT enhanced the conductivity. Zhang et al.\(^{24}\) fabricated composites with CNTs twined with Bi\(_2\)S\(_3\) microspheres using a facile reflux synthetic route, and the CNTs@Bi\(_2\)S\(_3\) composite electrode showed a reversible specific capacity of 247.9 mAh g\(^{-1}\) after 50 cycles. Jung et al.\(^{18}\) and Jin et al.\(^{20}\) developed carbon-coated Bi\(_2\)S\(_3\) crystallites as electrode materials; the specific capacity and cyclability of C@Bi\(_2\)S\(_3\) hybrid were significantly enhanced compared to pristine Bi\(_2\)S\(_3\). However, many well-known electrode materials with unique structural integrity for supercapacitor applications remain unexplored to enhance the specific capacitance. Continuing with the development of carbon nanomaterials, graphene and CNTs are often referred to as supercapacitors owing to their high electron-storage capacities and good electrical conductivity\(^{21,22}\).

Visible-light-driven photocatalytic reforming of biomass-derived feedstock is a sustainable, renewable, and cost-effective method for H\(_2\) production. Recently, the enhanced photocatalytic activity of Bi\(_2\)S\(_3\)-based composites were explored for photocatalytic hydrogen production\(^{23–25}\). Kadam et al.\(^{26}\) reported a Bi\(_2\)S\(_3\) quantum dot/glass composite that demonstrated efficient hydrogen production (6418.8 \(\mu\)mol h\(^{-1}\) g\(^{-1}\)) under solar light irradiation. Feng et al.\(^{27}\) reported carbon-coated Co\(_9\)S\(_8\) nanoparticles as electrocatalysts for water splitting with trithiocyanuric acid as a surfactant for the formation of core-shell structures. However, surfactant- or sacrificial- template-free synthetic procedures remains a challenge, offering scope for the rational design of high-performance core-shell heterostructures.

In this work, we have attempted and reported a rational assembly of carbon on the surface of Bi\(_2\)S\(_3\) nanorods, formed as a core-shell heterostructure via a facile and simple one-step hydrothermal reaction. The resulting C@Bi\(_2\)S\(_3\) core-shell heterostructures contain well-defined cross linkages between the Bi\(_2\)S\(_3\) core and carbon shell. These highly porous structures allow the free permeation of electrolyte ensuring rapid movement of ions. Also, the carbon shell on the Bi\(_2\)S\(_3\) core acts as an additive effectually shielding the mechanical strain during the charge and discharge processes. Moreover, the crosslinked assemblies in the C@Bi\(_2\)S\(_3\) core-shell heterostructures facilitates charge transfer through multipath electron transfer, which could accelerate the electrochemical processes to enhance high energy storage and photocatalytic H\(_2\) evolution features.

### Experimental Synthesis of Bi\(_2\)S\(_3\) and C@Bi\(_2\)S\(_3\).

To develop the final C@Bi\(_2\)S\(_3\), the Bi\(_2\)S\(_3\) was initially synthesized as described below using a modified hydrothermal method from previously reported literature\(^{28}\). In detail, Bi(NO\(_3\)\(_3\))\(_3\)·5H\(_2\)O (0.2 M) is dissolved in 25 mL of deionized (DI) water in a glass beaker with vigorous stirring, and a mixture of sodium sulfide (Na\(_2\)S·9H\(_2\)O) (0.3 M) and 36 mL H\(_2\)O\(_2\) was added drop by drop into the above solution at 90 °C with continuous stirring for 1 h. Then, it is transferred to a Teflon-lined autoclave and maintained at 190 °C for 6 h. Once it is cooled to room temperature naturally, the product was collected by centrifuging at 6000 rpm, and then was washed three times with ethanol. Then, the product was dried in a vacuum oven at 100 °C for 3 h.

In a typical synthetic procedure for C@Bi\(_2\)S\(_3\), 5 mg of the as-prepared Bi\(_2\)S\(_3\) was added in 30 mL ethanol and stirred to ensure homogenous mixing. Then, 15 mL of D-glucose solution (0.02 M) was mixed with the above solution, followed by continuous stirring for 60 min, and then transferred into a Teflon-lined stainless-steel autoclave and maintained at 190 °C for 6 h, and then cooled to room temperature naturally. The effect of hold time was explored by interrupting the shell growth at different times: 4, 6, 8, and 12 h, with the other parameters kept constant. Finally, the products were washed thoroughly with ethanol and water thrice and dried at 100 °C for 3 h. The schematic of the preparation procedures for the C@Bi\(_2\)S\(_3\) is presented in Fig. 1.
Characterization. The as-synthesized samples were characterized by X-ray powder diffraction (XRD) using a Shimadzu Labx X-ray diffractometer, XRD 6100 model. The XRD patterns of the samples were obtained using Cu Kα radiation (λ = 0.154 nm) at 40 kV and 30 mA. Transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM) observations were performed with a Hitachi H-7000 at 110 kV and a Tecnai G2 F 20 s-twin TEM at an accelerating voltage of 300 kV, respectively. Energy-dispersive X-ray spectroscopy (EDS) was performed to analyze the elemental composition. The scanning electron microscope (SEM) images were obtained using a Hitachi S-4800. X-ray photoelectron spectroscopy (XPS) analysis was conducted using a Thermo Scientific instrument and K-alpha surface analysis. The nitrogen adsorption–desorption isotherms were recorded using a Micromeritics ASAP 2420 surface area analyzer at liquid nitrogen temperature. Prior to gas adsorption, all photocatalysts were degassed for 3 h at 200°C. The infrared (IR) spectra were recorded using an Avatar 370 Fourier transform infrared (FTIR) spectroscope. The spectra were acquired at a resolution of 4 cm⁻¹ in the spectral range of 4000–400 cm⁻¹ using 32 scans. Raman spectra were verified using a Jobin-Yvon Horiba (Lab Ram HR-800) single monochromator attached with a Peltier-cooled charged coupled device (CCD) and used 532 nm Argon (Ar⁺) laser excitation source at 25°C under back scattering geometry.

2.3 Electrochemical Performance Measurements. All the electrochemical measurements were performed using the electrochemical workstation (Biologic SP-200) using a three-electrode configuration in a 0.5 M Na₂SO₄ aqueous electrolyte. The as-synthesized samples supported on ITO substrate directly served as the working electrode, while a saturated calomel electrode (SCE) and a platinum wire were used as the reference and the counter electrodes, respectively. Cyclic voltammetry (CV) and galvanostatic charge–discharge (GCD) measurements were performed to examine the electrochemical performance of the working electrodes. Electrochemical impedance spectroscopy (EIS) was conducted by applying an alternating current voltage with a 0-mV amplitude in the frequency range of 0.01 Hz to 1 MHz. The specific capacities (Cs, C g⁻¹) of the Bi₂S₃ samples were estimated from the GCD curves based on the following equation:

\[ C_s = \frac{I \times \Delta t}{m}, \]

where I (mA) indicates the discharge current, \( \Delta t \) (s) refers to the discharge time, and m (mg) corresponds to the mass of the active material. The corresponding specific capacitance can be estimated according to the following equation:

\[ C = \frac{2I}{m \Delta V \int_{t_1}^{t_2} V(t) \, dt}, \]

where \( \Delta V \) (V) is the potential range, \( V(t) \) is the operating of potential, and \( t_1 \) and \( t_2 \) denote to the initial and terminal discharge time of the GCD curves, respectively.

Photocatalytic H₂ production. Using the as-prepared photocatalysts, the photocatalytic hydrogen production was carried out by water splitting under simulated solar light irradiation. The reaction was carried out in a 150-mL tube-like quartz reactor with a round bottom. 5 mg of the photocatalyst was added to 50 mL of 5% aqueous lactic acid solution in a quartz reactor, which was then sealed with an air-tight rubber septum. The reactor was evacuated for 20 min, and the solution was purged with N₂ gas for 20 min. The reaction was carried out using a 300 W Xe lamp (Max 303) with a light intensity of 50 mW cm⁻². The amount of H₂ gas generated during the reaction was monitored at one-hour intervals by collecting the gas samples in an airtight syringe. The analysis of the sample was carried out using a gas chromatograph (6500GC system-YL Instrument) equipped with a thermal conductivity (TC) detector and a molecular sieve 5Å column with He as the carrier gas.

Results and Discussion

On the basis of the attempted experimental analysis, the formation mechanism of the core-shell structure and accompanying morphology evolution process is illustrated in Fig. 1. In this work, D-glucose was selected as a carbon source owing to its (i) environmental biocompatibility, (ii) unique and stable polymerized structure, and (iii) abundance. Moreover, the in-situ carbon coating also considerably simplifies the process and reduces the cost. Generally, D-glucose serving as a surfactant can generate carbon via a solid thermal decomposition reaction. In the present synthesis methodology, we used ethanol as a solvent to disperse the Bi₂S₃ nanorods and D-glucose was introduced as the carbon source. When the D-glucose is subjected to the hydrothermal treatment at 190°C, more than the glycosidation temperature, it tends to form C-C bonding that leads to the aromatization and carbonization. As the solution reach critical supersaturation, crosslinking between these macromolecules tends to the carbonization that results in a short single burst of nucleation. Here, the highly dispersed Bi₂S₃ nanorods, possessing highly reactive surface exposed outside, offers its surface to catalyze the carbonization of glucose (nucleation) that leads to in-situ deposition of carbon products around the Bi₂S₃ nanorods to produce carbonaceous shell. The growth of nuclei is subject to the reaction temperature, time and concentration. Herein, we studied the effect of reaction time (4, 6, 8, and 12 h holding times), keeping the other two conditions constant, on the thickness of the carbon shell and discussed in the HRTEM section. Under the complex polymerization of D-glucose at hydrothermal conditions (4, 6, 8, and 12 h holding times), Bi₂S₃ nanorod arrays employed as the templates form scalable core-shell C@ Bi₂S₃, where Bi₂S₃ nanorods are initially coated by carbonaceous nucleation on their outer surface. Under hydrothermal treatment, the D-glucose was degraded to carbon and anchored onto the surface of Bi₂S₃ nanorods. As reported in the literature, the thickness of the carbonaceous layer typically depends upon the synthesis conditions. For example, a glucose-derived carbon precursor can be readily integrated onto nanostructures.

\[ \frac{m}{V} \frac{dV}{dt} \]

\[ V(t) \]

\[ t_1 \]
in solution, which is carbothermally reduced to metallic ions by carbon with respect to hydrothermal temperature. Saravanakumar et al. demonstrated dextrose as a carbon source for the synthesis of carbon-coated V_2O_5 nanorods. They demonstrated that the controlled hydrothermal treatment led the formation of rod-like carbon structures. In this work, the overall carbonization process is controlled based on the holding time to avoid the destruction of Bi_2S_3 nanorod arrays. It is well known that above 220 °C, sulfur is decomposed and oxides were formed. Therefore, we fixed the hydrothermal temperature at 190 °C and studied the effect of holding time on the carbonaceous layer formation.

The morphological studies were performed using HRTEM and SEM. A representative HRTEM image of pristine Bi_2S_3 is observed in Fig. 2. As can be seen, the nanorods are a few nanometers in length (Fig. 2a). The high-magnification HRTEM image (Fig. 2b) indicates high crystallinity of the Bi_2S_3 nanorods with a lattice spacing of 0.36 nm that can be assigned to the (130) plane of orthorhombic Bi_2S_3, indicating that Bi_2S_3 nanorods favorably grew parallel direction to the (130) plane. The inset in Fig. 2b presents the relative SAED image reveals some bright spot patterns of pristine Bi_2S_3, which is also consistent with the single crystalline Bi_2S_3 indexed as the (130) zone c-axis.

The decoration of the carbon layer on the surface of individual Bi_2S_3 nanorods with respect to the holding time is examined via HRTEM. Figure 3 shows the HRTEM images of C@Bi_2S_3 samples. The nanorods are fully enclosed by the carbon material, thus forming a core-shell structure. In the case of the C@Bi_2S_3 sample, a typical core-shell shape of organic/inorganic hybrid nanostructure is formed, with well-organized atomic structures of carbon layers on the surface of Bi_2S_3 nanorod. It can be inferred that the ultra-thin shell-like carbon layer is formed by the nucleation of carbonaceous molecules. Figure 4 shows the high-magnification HRTEM images of pristine Bi_2S_3 and C@Bi_2S_3 at different holding times: 4, 6, 8, and 12 h. It is noteworthy that as the holding time is increased, the thickness of the carbon layer increases. Moreover, on the Bi_2S_3 nanorod surface, there is a ~2 nm thick coating layer, which is probably amorphous carbon resulting from the polymerization of glucose for all the C@Bi_2S_3 samples. It was observed that the thickness of the carbon layer slightly increased with an increase in the holding time. However, the carbon layer becomes dense with increasing holding times which is due to the excess carbon particles deposited on the Bi_2S_3 nanorod. The HRTEM images depict a dense carbon layer on the Bi_2S_3 nanorods surface as highlighted by the yellow arrows. HR-TEM performed on the grown carbon layer on the Bi_2S_3 nanorods surface reveals its structural uniformity and amorphous nature. At different holding times, the Bi_2S_3 rods were assembled by the amorphous carbon particles with variable thickness and densities. In addition, the mesoporous features of the carbon layer attributed roughness to the surfaces of the Bi_2S_3 nanorods, leading to enhanced electrochemical properties. In other words, the homogeneous carbon layer structure could efficiently enhance the conductivity of the electrodes, limit the volume change of Bi_2S_3 materials, and keep the Bi_2S_3 electrode stable during oxidation–reduction reactions. Based on the HRTEM analysis, the C@Bi_2S_3 sample at an 8 h holding time is considered for further characterization and applications in this work.

Figure 5a shows the SEM images of C@Bi_2S_3 sample. The SEM images indicate that the C@Bi_2S_3 structure comprises a large amount of homogeneous nanorods, which are covered by the amorphous carbon material. SEM observation indicates that the carbon was densely packed and uniformly covered the entire surface of Bi_2S_3 nanorods, resulting in the formation of core-shell structures. SEM images show that the nanorods have a length of approximately a few nanometers to submicron. Figure 5(b,c) presents the EDX elemental mapping of pristine Bi_2S_3 and C@Bi_2S_3 samples. This result also confirmed the deposition of carbon materials throughout the Bi_2S_3 nanorods and provides evidence for the existence of carbon materials in the C@Bi_2S_3 sample. Furthermore, the HRTEM elemental mapping of the C@Bi_2S_3 sample is shown in Fig. 5(d–g). It can be seen that the Bi_2S_3 nanorod is also decorated with the carbon layer.

The phase purity and structural features of the as-synthesized samples were verified via XRD, and the results are shown in Fig. 6. All the diffraction peaks in the both XRD patterns can be indexed to the orthorhombic phase.
of Bi$_2$S$_3$, which was well matched with JCPDS No. 89–8964. Interestingly, the weak peaks also match well with the standard pattern. In case of the C@Bi$_2$S$_3$ sample, similar diffraction peaks were observed owing to the amorphous nature of the carbon materials. As observed in the XRD result, the intensities of the diffraction peaks at 2θ = 11.08, 15.72, 17.56, 23.71, 24.92, 28.64, 31.11, 33.92, 35.91, 39.89, 42.69, 46.5, 48.44, 52.76, 62.71, 69.44, and 78.46 degrees corresponding to the (110), (020), (120), (101), (130), (230), (221), (410), (240), (141), (421), (431), (022), (312), (370), (561), and (670) planes of orthorhombic Bi$_2$S$_3$ respectively, agree well with reported values in literature. Both the pristine Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples exhibit good crystallinity and all the peaks can be clearly attributed to the orthorhombic phase of Bi$_2$S$_3$. Carbon is not recognized in the pattern of the C@Bi$_2$S$_3$ sample.

Figure 3. HRTEM image of the growth of C@Bi$_2$S$_3$ at different holding times (a) 4, (b) 6, (c) 8, and (d) 12 h, (e) high magnification image and (f) SAED pattern of C@Bi$_2$S$_3$ at a holding time of 8 h.
demonstrating the disordered nature of the carbon layer\textsuperscript{31}. In fact, the carbon component in the as-prepared C@Bi\textsubscript{2}S\textsubscript{3} sample is confirmed via SEM and HRTEM elemental mapping.

To confirm the nature of carbon shell existing on the Bi\textsubscript{2}S\textsubscript{3} nanorods, Raman spectroscopy of pristine Bi\textsubscript{2}S\textsubscript{3} and C@Bi\textsubscript{2}S\textsubscript{3} samples was recorded. As shown in Fig. 7, both samples demonstrate scattering bands at 258, 347, 426, 632, and 963 cm\textsuperscript{-1}, which are well consistent with the reported literatures for the crystalline Bi\textsubscript{2}S\textsubscript{3}\textsuperscript{32}. From the partial enlarged drawing of C@Bi\textsubscript{2}S\textsubscript{3} Raman spectra (inset of Fig. 7), two weak peaks appeared at 1354 and 1575 cm\textsuperscript{-1} are observed. These two new peaks of C@Bi\textsubscript{2}S\textsubscript{3} sample are ascribed to the D and G bands of carbon, respectively\textsuperscript{13}. Thus the Raman confirms the presence of carbon in the synthesized C@Bi\textsubscript{2}S\textsubscript{3} sample.

The FTIR spectrum of the C@Bi\textsubscript{2}S\textsubscript{3} core-shell structure within the range 400–4000 cm\textsuperscript{-1} is shown in Fig. 8. The peaks at 435, 573, 788, and 857 cm\textsuperscript{-1} can be assigned to Bi-S bonding\textsuperscript{28}. The vibrational stretching peak at 1045–1250 cm\textsuperscript{-1} can be attributed to the asymmetric S-H bond of Bi\textsubscript{2}S\textsubscript{3}. As reported in the previous study\textsuperscript{33}, the

Figure 4. High magnification HRTEM image of the (a,b) pristine Bi\textsubscript{2}S\textsubscript{3} and C@Bi\textsubscript{2}S\textsubscript{3} at holding times of (c) 4 h, (d) 6 h, (e) 8 h and (f) 12 h.
The existence of polymerized carbon layer could be identified through the FTIR analysis (Fig. 8). The carbon layer is identified by the appearance of C=O, C=C, O-H, and C-OH modes of vibration peaks at 1689, 1914, 3110–3670, and 1410 cm⁻¹, respectively. This indicates the presence of a large number of residual hydroxyl groups and intermolecular hydrogen bonds, which reflects the dispersion of water molecules.

The textural features of pristine Bi₂S₃ and C@Bi₂S₃ were investigated using N₂ adsorption-desorption isotherms present in type-H₃ hysteresis loops (Fig. 9a), which indicates the typical mesoporous characteristics. The Brunauer–Emmett–Teller (BET) surface areas of the pristine Bi₂S₃ and C@Bi₂S₃ samples estimated from N₂ isotherms were determined to be 22.58 and 43.63 m² g⁻¹, respectively. Compared to the previously reported Bi₂S₃@C (surface area is 11.2 m² g⁻¹), the as-prepared C@Bi₂S₃ possess higher BET surface area. The Barrett–Joyner–Halenda (BJH) pore-size distribution curves (Fig. 9b) of the pristine Bi₂S₃ and C@Bi₂S₃ samples display an average pore size of 15.1 and 16.2 nm, respectively, which further reveals their mesoporous features. The enhancement of the specific surface area and pore size by the introduction of carbon material characteristics clearly indicates that the C@Bi₂S₃ sample can store more charge for enhanced specific capacity owing to the enlarged active sites and the good approachability of electrolyte ions.

The chemical composition and chemical status of the elements in synthesized C@Bi₂S₃ core-shell structure were measured using x-ray photoelectron spectroscopy, as shown in Fig. 10. The Fig. 10a displays the XPS survey spectra of the pristine Bi₂S₃ and C@Bi₂S₃ samples. The prominent peaks at 158.56 and 163.81 eV in Fig. 10b are typical for Bi 4f 7/2 and Bi 4f 5/2, respectively, while the peaks at 161.34 and 162.52 eV are ascribed to the divalent anionic state of sulfur in Bi₂S₃. Figure 10c presents the peak at 163.53 eV corresponding to the binding energies of S 2p. The C 1s scan for carbon displayed in Fig. 10d has peaks at 284.53, 286.08, and 288.54 eV. The C 1s peak at 284.53 eV is dominated by elemental carbon, while two shoulders located at 286.08 and 288.54 eV are ascribed to
Figure 6. XRD pattern of pristine Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples at 8 h.

Figure 7. Raman spectra of pristine Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples at 8 h.
Figure 8. FTIR spectrum of the C@Bi$_2$S$_3$ sample at 8 h.

Figure 9. N$_2$ adsorption/desorption isotherm and, (b) Barrett–Joyner–Halenda (BJH) pore size distribution data of the pristine Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples at 8 h.
the existence of C–OH and C–O–C bonds, respectively. The presence of these small peaks suggests the existence of some residual groups due to the insufficient reduction of D-glucose.

In order to estimate the electrochemical properties of the pristine Bi₂S₃ and C@Bi₂S₃ samples, cyclic voltammetry (CV) and galvanostatic charge/discharge tests, electron impedance, and photocurrent tests were performed. Figure 11a presents the CV curves for the pristine Bi₂S₃ and C@Bi₂S₃ core-shell structures at a scanning rate of 10 mV s⁻¹ for a potential ranging from −0.6 to 0 V (vs SCE) in a 0.5 M Na₂SO₄ electrolyte. As shown in Fig. 11a, the area of the C@Bi₂S₃ core-shell structure is larger than that of pristine Bi₂S₃, representing a greater specific capacitance. Figure 11(b,c) displays the CV curves of the pristine Bi₂S₃ and C@Bi₂S₃ core-shell structures at various scanning rates. The voltammograms were recorded between −0.6 and 0 V at potential scan rates ranging from 1 to 250 mV s⁻¹ in a 0.5 M Na₂SO₄ solution. No redox peak is observed at different scan rates in the CV curves, suggesting that the electrode is charged and discharged at a pseudo-constant rate over the complete voltammetric cycle. The CV curve at faster scan rates has a larger area with a greater charge capacitance compared with that at a lower scan rate.

Figure 11d presents the constant current charge/discharge curves of the pristine Bi₂S₃ and C@Bi₂S₃ samples at a current density of 1 A g⁻¹. During the charging and discharging steps, the charge curve of the samples were not symmetric to its corresponding discharge counterpart with a negligible internal resistance (IR) drop, indicating the supercapacitive phenomenon. The GCD curve of the C@Bi₂S₃ sample exhibits a longer discharging time compared with the charging time. This phenomenon is repeatedly observed in Bi₂S₃-based electrode systems; however, the investigation of the exact mechanism of this phenomenon is still under progress. In addition, the GCD curves of the C@Bi₂S₃ sample at different current densities (1, 2, 3, 4, 5, 6 and 7 A g⁻¹) is presented in Fig. 11e.

The EIS analysis has been recognized as one of the principal methods to investigate the fundamental behavior of electrode materials, especially for supercapacitors. The impedance of both the samples are tested at frequencies varying between 0.1 Hz and 1 MHz at an open circuit potential with an AC perturbation of 0 mV. Figure 11f shows the Nyquist plots of the pristine Bi₂S₃ and C@Bi₂S₃ electrodes. Both the samples demonstrated a semicircular arc at higher frequencies and a straight line at lower frequencies, indicating that the electrode processes are organized by the charge transfer (i.e., electrochemical reaction) in the former region and by the diffusion of charges (i.e., mass transfer) in the absolute region. The intercept on the Z' axis refers to Rₛ, including the solution resistance and the contact resistance, whereas the dimension of the semicircle reflects the electrode film resistance (Rₚ), which is consistent with reported literature. EIS parameters, including the solution resistance (Rₛ), charge transfer resistance (Rₚ), and time constant (τ), were obtained as 1.2 Ω, 5.02 Ω, and 0.32 ms, respectively, for the C@Bi₂S₃ electrode, indicating fast charge/discharge processes, small solution resistance, and superior electrical conductivity at the electrode/electrolyte interface. Therefore, the carbon layer can efficiently bind with Bi₂S₃ nanorods, and thus improve electrical contact with the current collector.
confirms that more active carbon participates in the charge/discharge reaction process, leading to a higher charge carrier mobility.

Figure 12a presents the relationship between the capacitance and current density. The specific capacitance is 143.24, 129.87, 124.69, 110.45, 104.83, 96.21, and 90.61 F g\(^{-1}\) for pristine Bi\(_2\)S\(_3\) electrode, while the specific capacitance of the C@Bi\(_2\)S\(_3\) electrode is 333.43, 302.21, 280.16, 257.03, 243.97, 223.89, and 210.84 F g\(^{-1}\) at 1, 2, 3, 4, 5, 6, and 7 A g\(^{-1}\), respectively. The obtained value of the specific capacitance of the C@Bi\(_2\)S\(_3\) electrode is higher than the previously reported values for carbon-decorated MoS\(_2\) (210 F g\(^{-1}\) at 1 A g\(^{-1}\))\(^{37}\). As the current density increased from 1 to 7 A g\(^{-1}\), the specific capacitance linearly decreased to 90.6 and 210.84 F g\(^{-1}\) with a 63.2% retention of its original value. These findings specify that both the electrodes have a remarkable rate capability.

To explore the electrochemical stability and durability of the C@Bi\(_2\)S\(_3\) core-shell structure, consecutive GCD cycles were conducted at a constant current density of 4 A g\(^{-1}\). Figure 12b presents the relationship between

![Graphs showing electrochemical properties](image-url)
the specific capacitance and coulombic efficiency with respect to the number of cycles. The specific capacitance remained 97.36% of initial value even after 1000 cycles of operation. In addition, the morphology of C@Bi_2S_3 core-shell structure was examined using HRTEM, as shown in Fig. 12c. It can be seen that the core-shell feature of C@Bi_2S_3 still maintained even after suffering intense cycling test, which contributes to the remarkable cyclic stability. Moreover, the EIS of the 1st and 1000th cycle samples was conducted and compared. Figure 12d shows the Nyquist plots for the C@Bi_2S_3 electrode before and after 1000 cycles. It can be seen that both the curves contained a vertical line parallel to the imaginary axis at a low frequency, signifying a pseudocapacitor behavior. In addition, the R_{ct} value was consistent before and after 1000 cycles (5.02 and 5.08 Ω), respectively, which is reliable with the remarkable stability, as confirmed by the cycling tests, suggesting the interconnected structure of core-shell features of C@Bi_2S_3.

Figure 13a presents the photocatalytic H_2 production over the pristine Bi_2S_3 and C@Bi_2S_3 samples via water splitting using a 5 vol% lactic acid solution as a sacrificial agent. Due to the low bandgap, the recombination of electrons and holes in the Bi_2S_3 semiconductor is rapid leading to lower quantum yields. Here the carbon shell on the Bi_2S_3 prevents the recombination process by accepting an electron from the conduction band of Bi_2S_3. Also the high conductivity of carbon shell affluences the transportation of electrons to the reaction center that is the surface of carbon shell where the proton reduction process occurs. It is obvious that the photocatalytic H_2 generation increases with an increase in the irradiation time. Interestingly, the amount of H_2 generation is higher over the core-shell structure of C@Bi_2S_3 (754.34 μmol h^{-1} g^{-1}) compared to the pristine Bi_2S_3 (408.54 μmol h^{-1} g^{-1}), which is a 1.84-fold improvement in H_2 production over the core-shell structure of C@Bi_2S_3. It is believed that the improved photocatalytic activity may be associated with the core-shell feature of C@Bi_2S_3, which leads to close contact between the organic/inorganic frame networks. This core-shell interconnected assembled structure could also transport photo-induced electrons and holes to the binding sites; thus, the efficient oxidation of water occurred in the presence of lactic acid as a sacrificial agent, which is also consistent with previously reported literatures. In addition, the core-shell heterostructure may facilitate remarkable electron transfer compared with the pristine one, which can efficiently improve hydrogen production. As it already discussed in Fig. 10f in which the EIS of the C@Bi_2S_3 sample showed lower semicircle as compared to the pristine Bi_2S_3. The lower arc for the C@Bi_2S_3 sample is resulted from the less resistance of the material for the charge transfer process which is crucial for the electron transfer to the protons during the water splitting reaction to produce hydrogen. Thus, the presence of carbon shell improves the charge transfer characteristics of Bi_2S_3 that might be the reason for enhanced activity.

The effect of carbon shell thickness on the photocatalytic activity for the hydrogen production has been studied over C@Bi_2S_3 catalysts prepared at various reaction times (Fig. S1). All the C@Bi_2S_3 photocatalysts showed
superior H₂ production than the pristine Bi₂S₃. This infers that the presence of carbon shell has strong influence on the enhanced photocatalytic activity of Bi₂S₃. Maximum H₂ production was observed over the C@Bi₂S₃−8h catalyst which was prepared at 8 h of reaction time. However, the C@Bi₂S₃−12h sample prepared at 12 h reaction time showed slightly lower photocatalytic activity than the C@Bi₂S₃−8h sample. The excess reaction time might have resulted in the formation of overgrown thick carbon shell over the Bi₂S₃ in which the photoexcited electron in the Bi₂S₃ core has to travel longer path to reach the surface of the carbon shell where the proton reduction takes place. This results in the recombination of excited electrons with the holes and subsequent reduction in the photocatalytic activity. Also the excess reaction time might have formed individual carbon particles which may not be in direct contact with the C@Bi₂S₃ core shell structure. In the present study, the C@Bi₂S₃ core shell material prepared at 8 h reaction time has shown the optimum photocatalytic activity and hence we believe that 8 h reaction time is ideal for the growth of carbon shell on the surface of Bi₂S₃ core. Thus the C@Bi₂S₃−8h catalyst was selected for further studies.

Figure 13. (a) Photocatalytic H₂ production and, (b) Time on stream photocatalytic H₂ production activity over the pristine Bi₂S₃ and C@Bi₂S₃ samples under simulated solar light irradiation in 5 vol% lactic acid: water solution, (c) effect of lactic acid concentration on the photocatalytic H₂ production over 5 mg of pristine Bi₂S₃ and C@Bi₂S₃ samples and (d) recycling studies over the pristine Bi₂S₃ and C@Bi₂S₃ samples under simulated solar light irradiation in 5 vol% lactic acid.
rate, which reflects that the as-prepared samples are more stable, including those used in cyclic operations. Therefore, it can be concluded that the as-prepared samples, especially, C@Bi$_2$S$_3$ core-shell structures are more stable and reusable visible light–driven photocatalysts, which represent a fruitful methodology for harvesting solar energy for H$_2$ evolution and supercapacitor applications.

For a deeper insight into the behavior of photogenerated charge carriers, the profiles of transient photocurrent vs. time were obtained for the as-prepared Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples. The charge carrier separation phenomena of these samples were tested via on/off photocurrent at a bias voltage of 0.4 mV under simulated solar light irradiation, as shown in Fig. 14a. The results indicate that the C@Bi$_2$S$_3$ core-shell structure exhibits a remarkably higher photocurrent compared to the pristine Bi$_2$S$_3$, which is consistent with the above electrochemical performance (Fig. 11). The higher photocurrent over the core-shell structure of C@Bi$_2$S$_3$ specified that there was effective transfer of interfacial photogenerated charge carriers from Bi$_2$S$_3$ and improvement in electron-hole separation$^{18}$. This time-dependent response of the C@Bi$_2$S$_3$ sample is preferable for optoelectronic device applications. The photoexcited electron and holes can facilitate free radicals such as superoxide radicals and holes, which have been previously confirmed by electrochemical performance tests (Fig. 11). All these results confirm that more electrons may participate for photocatalytic water splitting and H$_2$ generation.

Figure 14. Transient photocurrent studies of the pristine Bi$_2$S$_3$ and C@Bi$_2$S$_3$ samples.

Figure 15. Schematic illustration of the plausible mechanism of photocatalytic H$_2$ production over the C@Bi$_2$S$_3$ core-shell structure under simulated solar light irradiation.
Based on the above results and discussion, a plausible mechanism of photocatalytic performance can be described as shown in Fig. 15. When C@Bi₂S₃ is exposed to visible light irradiation, Bi₂S₃ is excited and generates electron–hole pairs. The holes generated on the Bi₂S₃ nanorods can easily transfer to carbon because of the intimate contact between carbon and Bi₂S₃. Meanwhile, carbon can serve as a catalytic site for H₂ evolution. The following reasons can clarify the improved photocatalytic hydrogen evolution rate for the C@Bi₂S₃: (i) improved visible light absorption property helps in producing more electrons and holes, which advances photocatalytic activity, (ii) the remarkable specific surface area of the C@Bi₂S₃ core–shell structure provide more reactive sites for promoting hydrogen evolution, (iii) skillfully designed heterostructures prominently endorse the separation of electrons and holes, and (iv) the excellent properties of carbon materials such as high conductivity and more active sites benefit charge carrier separation and charge transfer to diminish the electron–hole pair recombination and enhance the photocatalytic hydrogen evolution rate. Finally, the introduction of carbon can reduce the H² reduction overpotential and act as a catalytic active site.

Conclusions
We attempted and demonstrated a promising core-shell design of C@Bi₂S₃ nanorods. The as-fabricated core-shell heterostructure can be directly employed as an electrode for supercapacitors, demonstrating a remarkable specific capacity of 333.43 F g⁻¹ at a current density of 1 A g⁻¹ and a considerable cycling stability with a capacity retention of 97.36% after 1000 cycles at a current density of 4 A g⁻¹. The remarkable cycling performance, combined with robust rate capability, exhibit the potential of C@Bi₂S₃ to be used as a novel electrode material for a supercapacitor. In addition, the core-shell structure of the C@Bi₂S₃ electrode demonstrated a H₂ evolution rate of 754.34 µmol h⁻¹ g⁻¹, a 1.84-fold improvement over the pristine Bi₂S₃. This outstanding improvement was due to enhanced specific surface area with site edges of carbon that helps in the effective separation and transportation of photoinduced carriers. It is expected that the proposed synthetic strategy has potential for the exploration of advanced electrochemical cell materials.

References
1. Bin, X. et al. Facile synthesis of nitrogen-doped porous carbon for supercapacitors. J. Mater. Chem. A 1, 4565–4570 (2013).
2. Zhang, L. L. et al. Graphene-based materials as supercapacitor electrodes. J. Mater. Chem. 20, 5983–5992 (2010).
3. Zhang, L. L. et al. Carbon-based materials as supercapacitor electrodes. Soc. Rev. 38, 2520–2531 (2009).
4. Tahir, M. et al. Multifunctional g–C₃N₄ Nanolayers: A Template-Free Fabrication and Enhanced Optical, Electrochemical, and Photocatalyst Properties. ACS Appl. Mater. Interfaces. 6. 1258–1265 (2014).
5. Wang, H. et al. Strongly coupled inorganic–nano-carbon hybrid materials for energy storage. Chem. Soc. Rev. 42, 3088–3113 (2013).
6. Zhou, W. et al. One-step synthesis of Ni₅S₉ nanorods@Ni(OH)₂ nanosheet core–shell nanostructures on a three dimensional graphene network for high-performance supercapacitors. Energy Environ. Sci. 6, 2216–2221 (2013).
7. Wang, H. et al. Exploiting Core–Shell Synergy for Nanosynthesis and Mechanistic Investigation. Acc. Chem. Res. 46, 1636–1646 (2013).
8. Raj, C. J. et al. Electrochemical capacitor behavior of copper sulfide (CuS) nanoplatelets. J. Alloy. Compd. 586, 191–196 (2014).
9. Fang, Z. et al. Epitaxial Growth of CdS Nanoparticle on Bi₂S₃ Nanowire and Photocatalytic Application of the Heterostructure. J. Phys. Chem. C. 115(29), 13968–13976 (2011).
10. Meng, X. et al. Cobalt Sulfide/Graphene Composite Hydrogel as Electrode for High-Performance Pseudocapacitors. Sci. Rep. 6, 21717 (2016).
11. Zhu, T. et al. Hierarchical nickel sulfide hollow spheres for high performance supercapacitors. RSC Adv. 1, 397–400 (2011).
12. Pawar, S. H. et al. Growth of Bi₂S₃ film using a solution-gas interface technique. Thin Solid Films 110, 165–170 (1983).
13. Zhao, Y. et al. One-pot facile fabrication of carbon-coated Bi₂S₃ nanomeshes with efficient Li-storage capability. Nano Res. 7(5), 765–773 (2014).
14. Nayak, B. B. Structural characterization of Biₓₐ₋ₓShₓS₃ films prepared by the dip–dry method. Thin Solid Films. 105, 17–24 (1983).
15. Zhou, X. Facile synthesis of MoSₓ@CMK-3 nanocomposites as an improved anode material for lithium-ion batteries. Nanoscale 4, 5868–5873 (2012).
16. Xu, C. et al. Controlled soft-template synthesis of ultrathin C@Fe₃O₄ nanosheets with high Li–storage performance. ACS Nano 6, 4713–4721 (2012).
17. Yang, W. et al. A Bi₂S₃@CNT nanocomposite as anode material for sodium batteries. Materials Letters 167, 102–105 (2016).
18. Zhang, Z. et al. Facile synthesis of dandelion-like Bi₂S₃ microspheres and their electrochemical properties for lithium-ion batteries. Mater. Lett. 91, 100–102 (2013).
19. Jung, H. et al. Bismuth sulfide and its carbon nanocompositesfor rechargeable lithium-ion batteries, Electrochim. Acta. 56, 2135–2141 (2011).
20. Jin, R. et al. Carbon coated flower like Bi₂S₃ grown on nickel foam as binder-free electrodes for electrochemical hydrogen and Li-ion storage capacities. Electrochim. Acta. 173, 458–464 (2015).
21. Robel, I. et al. Single-Walled Carbon Nanotube-CdS Nanocomposites as Light-Harvesting Assemblies: Photoinduced Charge-Transfer Interactions. Adv. Mater. 17, 2458–2463 (2005).
22. Qu, F. L. et al. Colorimetric Platform for Visual Detection of Cancer Biomarker Based on Intrinsic Peroxidase Activity of Graphene Oxide. Biosens. Bioelectron. 26, 3927–3931 (2011).
23. Brahim, R. et al. Visible light induced hydrogen evolution over the heterosystem Bi₂S₃/TiO₂. Catalysis Today 122, 62–65 (2007).
24. Zeng, Q. et al. Combined nanostructured Bi₂S₃/TNA photoanode and Pt/Si/PVC photocathode for efficient self-biasing photoelectrochemical hydrogen and electricity generation. Nano Energy. 9, 152–160 (2014).
25. Han, M. et al. 3D Bi₂S₃/TiO₂ cross-linked heterostructure: An efficient strategy to improve charge transport and separation for high photoelectrochemical performance. J Power Sources. 329, 23–30 (2016).
26. Kadam, S. B. et al. A stable Bi₂S₃ quantum dot–glass nanosystem: size tuneable photocatalytic hydrogen production under solar light. RSC Adv. 5, 58485–58490 (2015).
27. Feng, L. L. et al. Carbon-Armored Co₃S₄ Nanoparticles as All-pH Efficient and Durable H₂-Evolving Electrocatalysts. ACS Appl. Mater. Interfaces 7, 980–988 (2015).
28. Vattikuti, S. V. et al. Synthesis, characterization, and optical properties of visible light driven Bi₂S₃ nanorod photocatalysts. J Mater Sci: Mater Electron 28, 14282–14292 (2017).
29. Saravanakumar, B. et al. High performance supercapacitor based on carbon coated V₂O₅ nanorods. J. Electroanalyt. Chem. 758, 111–116 (2015).
30. Sun, X. M. et al. Colloidal carbon spheres and their core/shell structures with noble-metal nanoparticles. Angew. Chem. Int. Ed. 43, 597–601 (2004).
31. Zhang, C. et al. Facile synthesis of carbon-coated MoS2 nanorods with enhanced lithium storage properties. *Electrochim. Commun.* **20**, 7–10 (2012).
32. Zuo, X. et al. Facile synthesis of Bi2S3-C composite microspheres as low-cost counter electrodes for dye-sensitized solar cells. *RSC Adv.* **4**, 57412–57418 (2014).
33. Kaneti, Y. L. et al. Carbon-Coated Gold Nanorods: A Facile Route to Biocompatible Materials for Photothermal Applications. *ACS Appl. Mater. Interfaces* **7**, 25658–25668 (2015).
34. Hu, S. et al. Preparation of carbon coated MoS2 flower-like nanostructure with self-assembled nanosheets as high-performance lithium-ion battery anodes. *J. Mater. Chem. A.* **2**, 7862–7872 (2014).
35. Sun, Y. Ultrafine MoO2 nanoparticles embedded in a carbon matrix as a high-capacity and long-life anode for lithium-ion batteries. *J. Mater. Chem.* **22**, 425–431 (2012).
36. Ma, L. et al. Controlled assembly of Bi2S3 architectures as Schottky diode, supercapacitor electrodes and highly efficient photocatalysts. *RSC Adv.* **4**, 41636–41641 (2014).
37. Hu, B. et al. Synthesis of porous tubular C/MoS2 nanocomposites and their application as a novel electrode material for supercapacitors with excellent cycling stability. *Electrochem. Acta* **100**, 24–28 (2013).
38. Pandey, K. et al. Influence of current collector electrode on the capacitive performance of electrodeposited PANI: insight gained from frequency and time domain analysis. *RSC Adv.* **4**, 53740–53751 (2014).
39. Li, Q. et al. Design and Synthesis of MnO2/Mn/MnO2 Sandwich-Structured Nanotube Arrays with High Supercapacitive Performance for Electrochemical Energy Storage. *Nano Lett.* **12**(7), 3803–3807 (2012).
40. Ni, M. et al. A review and recent developments in photocatalytic water-splitting using TiO2 for hydrogen production. *Renew Sust. Energ. Rev.* **11**, 401–425 (2007).
41. Feng, J. et al. Co(OH)2@PANI hybrid nanosheets with 3D networks as high-performance electrocatalysts for hydrogen evolution reaction. *Adv. Mater.* **27**, 7051–7057 (2015).
42. Feng, J. et al. Efficient hydrogen evolution on Cu nanodots-decorated Ni3S2 nanotubes by optimizing atomic hydrogen adsorption and desorption. *J. Am. Chem. Soc.* **140**(2), 610–617 (2018).
43. Bera, S. et al. Fabrication of Bi2S3/ZnO heterostructures: an excellent photocatalyst for visible-light-driven hydrogen generation and Photoelectrochemical properties. *New J. Chem.* **42**, 541–554 (2018).

**Acknowledgements**

This research was supported by the National Research Foundation of Korea (NRF) and funded by the Ministry of Science, ICT, and Future Planning (2017R1A2B1004860). This work was supported by the National Research Foundation of Korea (NRF) grant funded by the Korea government (2017R1A4A101558).

**Author Contributions**

S.V.P.V. designed, conducted the synthesis, and supervised the project. P.A.K.R. performed the test of the electrochemical surface area. J.S. and C.B. partly designed and supervised the project. All authors contributed to data analysis and approved the final version of the manuscript.

**Additional Information**

*Supplementary information* accompanies this paper at https://doi.org/10.1038/s41598-018-22622-0.

**Competing Interests:** The authors declare no competing interests.

**Publisher’s note:** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

---

**Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit [http://creativecommons.org/licenses/by/4.0/](http://creativecommons.org/licenses/by/4.0/).

© The Author(s) 2018