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Assembling a high-performance asymmetric supercapacitor based on pseudocapacitive S-doped VSe₂/CNT hybrid and 2D borocarbonitride nanosheets†

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The construction of high-performance asymmetric supercapacitors is vital for achieving a sustainable energy storage model. Layered 2D materials are considered pivotal contributing factors to modern-day energy storage solutions. Herein, the electrochemical performance of the S-doped VSe₂/CNT positrode and the BCN negatrode is explored for the construction of a compelling aqueous asymmetric supercapacitor. This study provides an understanding of the sulfur doping effects and contributions of CNT addition in the energy storage performance of VSe₂. The S doping and CNT addition played a significant role in boosting the pseudocapacitive energy storage performance of metallic VSe₂. Borocarbonitride (BCN), which is utilized as a conventional electric double layer material, shows outstanding performance as a negatrode owing to its inherent properties. An asymmetric supercapacitor (ASC) assembled using S-VSe₂/CNT and BCN yields a high energy density of 36.3 μW h cm⁻² with remarkable reversibility and initial capacitance retention of 87.2% even after 5000 cycles. The remarkable electrochemical performance of S-VSe₂/CNT//BCN ASC provides a significant reference for futuristic electronic and energy storage applications.

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1. Introduction

Supercapacitors are compelling energy storage devices to integrate with modern renewable energy harvesting systems.¹ They are widely acknowledged owing to their excellent power operation, rapid charge–discharge ability, extraordinary cycle life and safety.² Conventionally, supercapacitors are categorized into electric double layer capacitors (EDLCs) and pseudocapacitors based on electrochemical double layer formation and surface bound redox reactions at the electrode and electrolyte interface respectively.^{3,4} In the last few decades, numerous efforts have been made in the scientific community to enhance the performance of supercapacitors either by applying high energy density pseudocapacitor electrode materials or constructing asymmetric supercapacitors (ASCs) with a high operating window. Compared to the symmetric system, ASCs combine the working potential of both the positrode and negatrode to achieve a large operating window.⁵ The intrinsic limitation of bulk materials paved the way for utilizing various two-dimensional (2D) nanomaterials as electrode materials.

Recently, 2D transition metal dichalcogenides (TMDs) have gained attention in the field of supercapacitors owing to their excellent electronic conductivity, fast reversible redox reactions and numerous electrochemical active sites.^{6,7} TMDs are denoted by the general formula MX₂, where M represents transition metal Ti, V, Mo or W and X represents S or Se atom, in which a layer of transition metal atoms is sandwiched between layers of chalcogen atoms. Among TMDs, vanadium diselenide (VSe₂) is widely utilized in numerous energy storage and conversion applications owing to its exotic surface and electronic properties.^{8–10} It has been reported that strong electronic coupling interaction between V⁴⁺ atoms imports metallic properties to VSe₂.¹¹ The sandwiched Se–V–Se layers of VSe₂ are stacked by a weak van der Waals forces with an interlayer spacing of ~6.1 Å.⁸ The reasonably large interlayer spacing allows for the facile diffusion of large ions during the electrochemical reactions without any structural deformations. The potential multi-electron transfer per formula unit provides a large theoretical energy density for VSe₂.¹² Recently, our group explored VSe₂ and its nano carbon-based hybrids for supercapacitors.^{13,14} The nanocarbon additives, such as carbon nanotubes (CNTs) and graphene, enhance the energy storage performance of VSe₂ along with providing excellent structural and electrochemical stability.^{15–18} However, the energy storage performance of VSe₂-based electrodes is still far below the theoretically predicted values. Doping of TMDs with non-metal, such as

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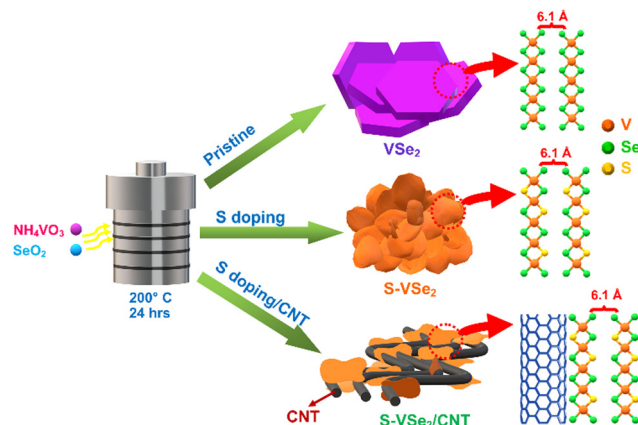
sulphur, is also a reported strategy to enhance energy storage performance.^{19–21} The anion substitution, such as sulphur, can alter the electronic and chemical atmosphere in the VSe_2 host system and generates more electrochemical active centres.^{22–24} The S doping affects surface properties and significantly affects reaction pathways and kinetic energy barriers for the electrochemical reactions.^{23,25,26} The strategy of synthesizing the S-doped VSe_2 /CNT hybrid can synergize the effects of both doping and hybrid formation. It is interesting when the S-doped VSe_2 /CNT positrode is combined with the borocarbonitride negatrode to achieve a high-energy density asymmetric supercapacitor.

Borocarbonitride (BCN), composed of graphene and BN domains and having only B–C, C–N, B–N and C–C bonds, is an emerging 2D material that exhibits excellent activity towards supercapacitor and hydrogen evolution reaction owing to high surface areas and abundant active sites, such as stone wale defects and sp^3 -carbon.^{27–29} The structural and compositional properties of the BCN significantly differ from those of B, N heteroatom-doped graphene because of the combined presence of BN and graphene domains.³⁰ BCN has excellent electrical conductivity, tunable surface area and enriched surface bound active centres for electrochemical adsorption contributed by both the BN and graphitic domains.^{28,30} Recent studies reveal the compelling supercapacitor performance of BCN in various electrolyte media. However, the true potential of BCN in supercapacitor applications must be tapped into further. In this study, the S-doped VSe_2 /CNT hybrid and BCN have been successfully employed as electrodes to construct a compelling asymmetric supercapacitor.^{30,31} The charge storage performance and mechanism of each electrode material are thoroughly studied using various electrochemical characterizations. The ASC constructed with S-doped VSe_2 /CNT as the positrode and BCN as the negatrode exhibits an outstanding energy storage performance with an energy density of $36.3 \mu\text{W h cm}^{-2}$ at a power density of 3.2 mW cm^{-2} . Remarkably, the S- VSe_2 /CNT//BCN ASC showed excellent cyclic stability by retaining 87.2% of its initial areal capacitance after 5000 charge/discharge cycles.

2. Results and discussion

2.1 Material characterization of S- VSe_2 /CNT

The synthesis route of VSe_2 , S-doped VSe_2 (S- VSe_2) and S- VSe_2 /CNT is depicted in Scheme 1, and the detailed synthesis procedure is provided in the ESI.† The carboxyl groups of functionalized CNT can act as nucleation sites that electrostatically absorb V^{4+} ion from the starting precursor and then culminate in the growth of S- VSe_2 sheets on CNT.^{32,33} Crystal-line phase analysis of the synthesised samples was carried out using XRD. XRD patterns of VSe_2 , S- VSe_2 and S- VSe_2 /CNT (Fig. 1a) show the highly crystalline diffraction pattern of the hexagonal VSe_2 phase (JCPDS: 89-1641). The XRD pattern of the S-doped sample agrees well with the undoped VSe_2 sample without the formation of any isolated VS_2 or VS phases. The intensity of the (00 l) c -axis peaks in the S- VSe_2 sample is substantially increased compared to that of pristine VSe_2 .



Scheme 1 Schematic representation of the synthesis of VSe_2 , S- VSe_2 and S- VSe_2 /CNT.

A similar intensity variation can also be observed in the S- VSe_2 /CNT sample, signifying that this relative intensity variation of the c -axis plane is caused by S doping in the VSe_2 lattice. This distinctly enhanced (00 l) reflections in the doped samples can be attributed to the formation of multilayer VSe_2 nanosheets compared to the bulk macro sheets of pristine VSe_2 .^{34,35} A broad reflection in the S- VSe_2 /CNT from the (002) plane of CNT is observed at $\sim 26^\circ$, confirming the incorporation of CNT. In the diffraction profile of S- VSe_2 /CNT, small reflections apart from VSe_2 and CNT are also observed owing to the presence of minute fractions of V_2O_5 .^{36,37} Further, the doping effects and hybrid formation in the samples are evaluated using Raman spectroscopy (Fig. 1b). Pristine VSe_2 shows a sharp characteristic A_{1g} vibration at 231 cm^{-1} and small E_g vibration at 137 cm^{-1} .³⁸ The presence of these vibrations can be observed in both S- VSe_2 and S- VSe_2 /CNT confirming the formation of VSe_2 . In S- VSe_2 , the E_g vibration is prominently visible probably because of the S-doping-induced VSe_2 multilayer nanosheet formation. In the case of S- VSe_2 /CNT, apart from the A_{1g} and E_g bands of VSe_2 , the characteristic D and G Raman bands of CNT are observed at 1345 cm^{-1} and 1595 cm^{-1} , respectively.³⁹ The presence of D and G vibrations explicitly confirms the presence of CNT in the hybrid.

XPS is implemented to study the chemical state and composition of the S- VSe_2 /CNT sample. The XPS survey provided in Fig. S1 (ESI†) shows the presence of V 2p, Se 3d, S 2p and C 1s signals. The high-resolution V 2p spectra show two peaks corresponding to the V 2p_{3/2} and V 2p_{1/2} states (Fig. 1c). The V 2p_{3/2} peak is deconvoluted into two peaks located at 516.5 eV and 517.6 eV, and the V 2p_{1/2} peak is deconvoluted into two peaks at 523.8 eV and 525.1 eV. In particular, peaks at 516.5 eV and 523.8 eV correspond to the V^{4+} state confirming the presence of VSe_2 formation.^{8,12,38} The peaks at 517.6 eV and 525.1 eV represent the V^{5+} state of V, showing the presence of V_2O_5 formation during the CNT hybrid synthesis.³⁶

The deconvoluted Se 3d spectrum shows two peaks at 54.7 eV (Se 3d_{5/2}) and 55.8 eV (Se 3d_{3/2}) attributed to the V–Se bonding in VSe_2 crystal (Fig. 1d).⁴⁰ Fig. 1e shows the high-resolution S 2p

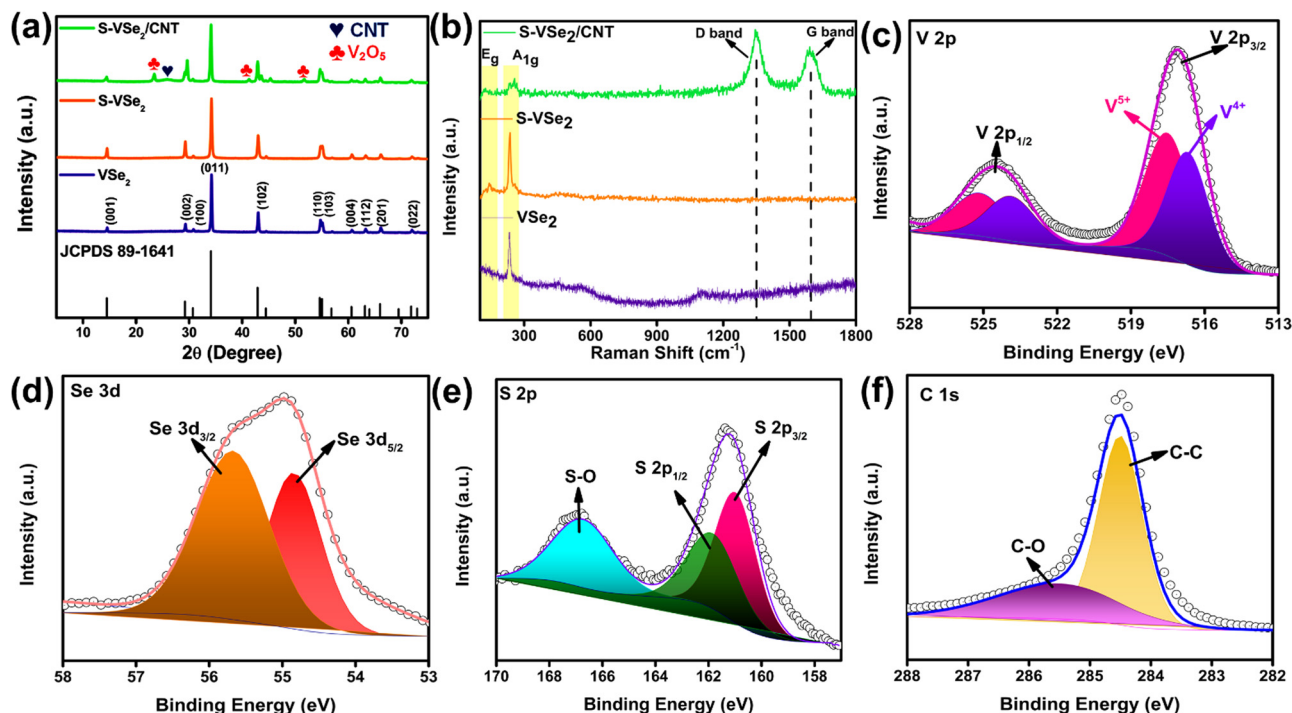


Fig. 1 (a) XRD pattern and (b) Raman spectra of VSe₂, S-VSe₂ and S-VSe₂/CNT; high-resolution XPS spectra of (c) V 2p, (d) Se 3d, (e) S 2p and (f) C 1s in S-VSe₂/CNT sample.

spectrum deconvoluted into two peaks at 161.1 eV and 162.08 eV corresponding to S 2p_{3/2} and S 2p_{1/2} states of S²⁻ ions.^{24,41,42} These results imply the substitutional S doping by replacing Se atoms in the VSe₂ lattice. A broad peak observed at 167.28 eV corresponds to the formation of SO₂ after the surface oxidation of the S-VSe₂/CNT sample.^{24,41} The C 1s core level spectrum shows an intense peak at 284.4 eV due to the sp² hybridized graphitic carbon atoms, which double down the presence of CNT; peak 285.5 eV can be attributed to C-O bond formation due to the presence of phenols or ether (Fig. 1f).³³ The oxygenated species of vanadium and carbon can cause the origin of the interfacial binding of S-VSe₂ and CNT during the hybrid formation.⁴³

FESEM analysis reveals the morphologies of the synthesized samples. Pristine VSe₂ exhibits a large hexagonal sheet morphology similar to previous reports (Fig. 2a and b). After S doping, a significant change is observed in the morphology of VSe₂. The S-VSe₂ samples show a nanoflower type of morphology with each flower comprising thin S-VSe₂ nanosheets (Fig. 2c and d). Compared to the bulk VSe₂, this nanoflower in S-VSe₂ exposes many electrochemical active centres that eventually enhance the adsorption rate and energy storage performance. In the case of S-VSe₂/CNT, it can be observed that the S-VSe₂ nanosheets were anchored by the multiwalled CNT (Fig. 2e and f). This interconnected network of CNT and S-VSe₂ nanosheets provides more active centres and short diffusion pathways for ions that enhance the electrochemical performance of the S-VSe₂/CNT sample. Fig. 2g depicts the EDS elementary mapping of S-VSe₂/CNT showing the uniform distribution of V, Se, and C in the sample, confirming homogeneity.

2.2 Electrochemical Assessment of S-VSe₂/CNT

Electrochemical analysis of VSe₂, S-VSe₂ and S-VSe₂/CNT is studied in a conventional three-electrode setup with a glassy carbon working electrode and a 0.5 M K₂SO₄ aqueous electrolyte. All the electrodes are assessed in a potential window ranging from -0.25 to 0.65 V. Pristine VSe₂ shows a quasi-rectangular CV shape, which indicates the pseudocapacitor behaviour of the electrode. The broad redox peaks in the cyclic voltammogram suggest a possible intercalation of the K⁺ ions during the electrochemical reactions (Fig. S2a, ESI[†]).^{5,12} Fig. 3a illustrates S-VSe₂ with a significant enhancement in current response and enclosed CV area compared to pristine VSe₂. This clearly suggests that S doping triggered a substantial enhancement in energy storage performance in VSe₂. The electrochemical performance of pristine VSe₂ and S-VSe₂ is illustrated in Fig. S2 (ESI[†]). Similarly, the addition of CNT into S-VSe₂ has further exceeded the energy storage performance (Fig. 3a). The CV performance of S-VSe₂/CNT at various scan rates is shown in Fig. 3b. The CV shape of the S-VSe₂/CNT electrode reflects the signatures of an EDLC behavior, which are obviously contributed by the presence of CNT in the sample. During all scan rates, S-VSe₂/CNT electrode maintained a similar CV shape that indicates the reversibility of the electrode during the electrochemical activity. To confirm the cation contribution, we tested S-VSe₂/CNT electrode in KCl; in both cases, the electrode retained a similar CV shape. This confirms the dominance of cation contribution in pseudocapacitive activity (Fig. S3, ESI[†]).

To understand the charge storage mechanism in S-VSe₂/CNT electrode, we have the induced power law $i = av^b$ in the CV curve, where i is the current response at a specific potential,

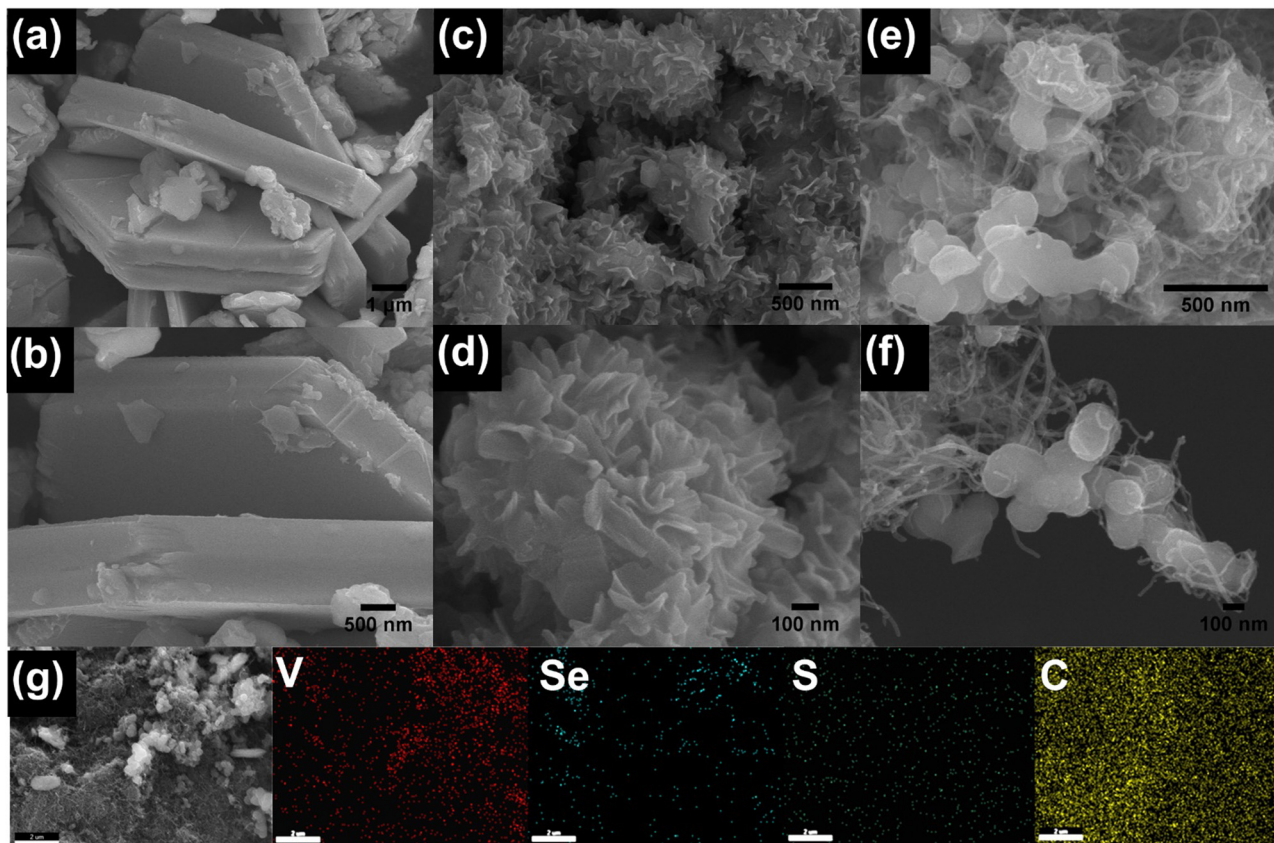


Fig. 2 Low and high magnified FESEM images of (a and b) VSe_2 , (c and d) $S-VSe_2$, (e and f) $S-VSe_2/CNT$ and (g) EDS elemental mapping of $S-VSe_2/CNT$ sample showing the uniform distribution of V, Se, S, and C.

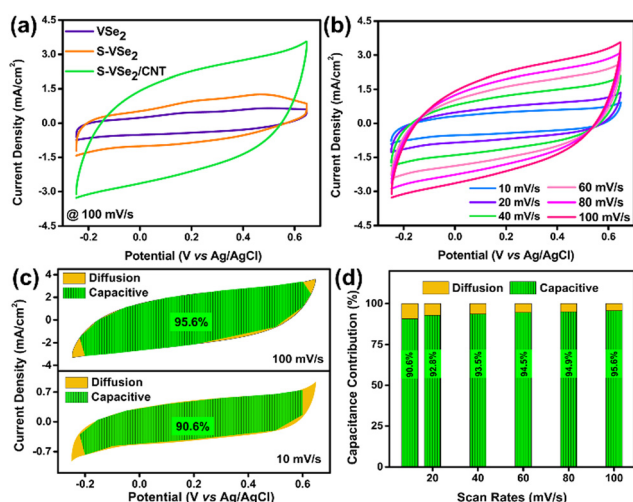


Fig. 3 Electrochemical measurements of VSe_2 , $S-VSe_2$ and $S-VSe_2/CNT$ electrodes in 0.5 M K_2SO_4 . (a) CV profile of VSe_2 , $S-VSe_2$ and $S-VSe_2/CNT$ at a scan rate of 100 mV s^{-1} , (b) CV profile of $S-VSe_2/CNT$ in scan rates ranging from 10 mV s^{-1} to 100 mV s^{-1} , (c) segregated CV curves of $S-VSe_2/CNT$ into capacitive and diffusive regions at selected scan rates and (d) normalized capacitive and diffusive contribution in all the scan rates.

a and b are the adjustable parameters, and ν is the scan rate.⁴⁴ Generally, if $b = 1$, then the mechanism is dominated by

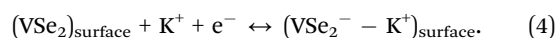
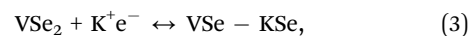
capacitive contribution, and if $b = \frac{1}{2}$, then the mechanism is dominated by diffusion reactions. The power law can be rewritten as follows:⁴⁴

$$i(V) = k_1\nu + k_2\nu^{1/2}, \quad (1)$$

OR

$$i(V)/\nu^{1/2} = k_1\nu^{1/2} + k_2. \quad (2)$$

By calculating k_1 and k_2 , the CV profiles can be deconvoluted into capacitive and diffusive contributions. The deconvoluted CV profile of $S-VSe_2/CNT$ shows complete domination of capacitive behaviour over diffusion. Fig. 3c shows the segregation of capacitive and diffusive contributions at lower and higher scan rates. At a lower scan rate of 10 mV s^{-1} , the capacitive contribution is 90.6%, which clearly suggests that the electrochemical mechanism is dominated by capacitive contributions obtained from the intercalation pseudocapacitance of $S-VSe_2$ and the EDLC of CNT. The power law-induced deconvolution is performed at all scan rates, and the results are shown in Fig. 3d. The electrochemical reactions in VSe_2 involve fast K^+ ion accessibility, which can be represented as follows:⁴⁵



The GCD curves of S-VSe₂/CNT show a quasi-triangular shape without any potential independent region in current densities ranging from 1 to 8 mA cm⁻² (Fig. 4a). Further, the GCD curve limits the IR drop and exhibits excellent coulombic efficiency, corroborating the reversible electrochemical reactions involved with the S-VSe₂/CNT electrode. Areal capacitance as a function of current density for VSe₂, S-VSe₂ and S-VSe₂/CNT is illustrated in Fig. 4b. S doping increased the areal capacitance of S-VSe₂ more than twice that of pristine VSe₂. The addition of CNT further enhanced the areal capacitance to 83 mF cm⁻², which is a significant enhancement from S-VSe₂. The enhancement in the capacitance value of the S-VSe₂/CNT electrode can be attributed to the synergistic effect between S-VSe₂ and CNT. Cyclic stability is one of the most important factors in analysing the performance of supercapacitor electrodes. S-VSe₂/CNT exhibited a capacitance retention of 87.5% after 5000 charge/discharge cycles with a coulombic efficiency of more than 97% (Fig. 4c). High capacitance retention and coulombic efficiency indicate excellent electrochemical stability and reversibility of the S-VSe₂/CNT electrode. The kinetics of the electrochemical reactions of all the samples are analysed using EIS (Fig. 4d). The intersection at the high-frequency region is usually considered the intrinsic resistance of the electrode (R_s), and the diameter of the semi-circular region at

the mid-high-frequency region represents the charge transfer resistance (R_{ct}). The Nyquist plot reveals that S doping reduced charge transfer resistance of VSe₂, which emphasises the enhanced electrochemical performance in S-VSe₂.⁴⁶ After the addition of CNT, the R_{ct} value further decreased significantly. This correlates with the excellent charge storage performance of the electrode.¹⁴ Pristine VSe₂ shows a low R_s value owing to its inherent metallic nature. An increase in the R_s value of the S-VSe₂ sample is due to an altered electronic atmosphere after the doping of a non-metal. The addition of highly conducting CNT reduces the R_s value, which clearly affects the charge storage performance of the S-VSe₂/CNT electrode. R_s and R_{ct} values of all the samples are provided in Table S1 (ESI†). The diffusion-controlled high-frequency region of the S-VSe₂/CNT electrode exhibits capacitor-type behaviour correlating with results obtained after the deconvolution of CV.

S doping alters the electronic atmosphere inside the VSe₂ lattice and provides better electronic conductivity.^{23,42} The formation of the nanoflower morphology of VSe₂ after S doping improves the electrode/electrolyte contact area, and the increased number of active sites can enhance the rate of electrochemical reactions.⁴⁷ CNT addition provides further improvement in electrochemical performance owing to the synergistic effect between S-VSe₂ and CNT.⁴⁸ The presence of CNT facilitates the accommodation of more electrolyte ions, and this enhances the electrochemical reactions. The highly conducting CNT network decreases ionic and electronic pathways between S-VSe₂ nano sheets and fastens the rate of electrochemical reactions. CNT acts as a conductive skeleton and provides excellent electrochemical stability to S-VSe₂.⁴⁹ This hybrid architecture of S-VSe₂/CNT prevents the volume expansion and restacking of S-VSe₂ layers during K⁺ intercalation and deintercalation. The plausible charge storage mechanism of the S-VSe₂/CNT electrode is schematically depicted in Fig. 4e.

2.3 Material Characterization of the BCN

The synthesis procedure of the BCN nanosheets is schematically illustrated in Fig. 5a. The structural characteristics of the prepared BCN are analysed using XRD. The XRD pattern of the BCN shows characteristic peaks of the (002) and (100) planes located at 25.8° and 42.7°, respectively (Fig. S3a, ESI†). The broadening of the (002) peak indicates the formation of few-layer thick BCN sheets.³⁰ Raman spectra show two distinct peaks at 1349 and 1596 cm⁻¹, respectively (Fig. 5b), corresponding to in-plane B-N vibrations and graphitic D bands.^{27,30} Moreover, the peak observed at 1596 cm⁻¹ (G band) is attributed to the sp²-bonded carbon atoms of the graphene domains. FESEM images of the prepared BCN are provided in Fig. 5c and d. The BCN shows a sheet-like morphology with porous structures on the surface. This porous sheet-like morphology enhances electrochemical adsorption and allows for the use of almost the entire available electrode surface for electrochemical reactions. EDS elemental mapping is provided in Fig. S4e (ESI†), showing a uniform distribution of B, C and N in the BCN sample. XPS analysis was implemented to understand the elemental composition of the BCN sample.

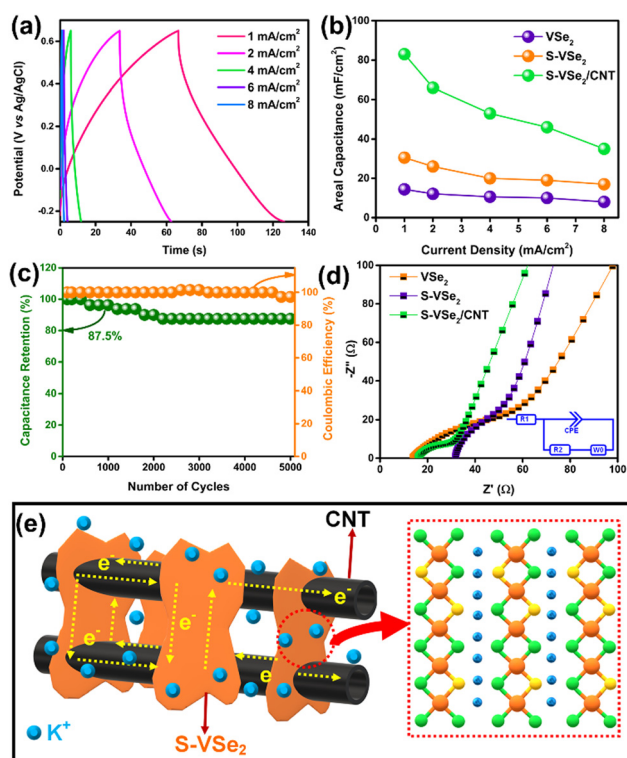


Fig. 4 (a) GCD curves of S-VSe₂/CNT electrode in various current densities, (b) areal capacitances of VSe₂, S-VSe₂ and S-VSe₂/CNT plotted against corresponding current densities, (c) cyclic performance of S-VSe₂/CNT during 5000 cycles at a current density of 10 mA cm⁻², (d) Nyquist plots of VSe₂, S-VSe₂ and S-VSe₂/CNT and (e) schematic illustration of the possible charge storage mechanism in S-VSe₂/CNT electrode involving intercalation of K⁺ ions in S-VSe₂ sheets.

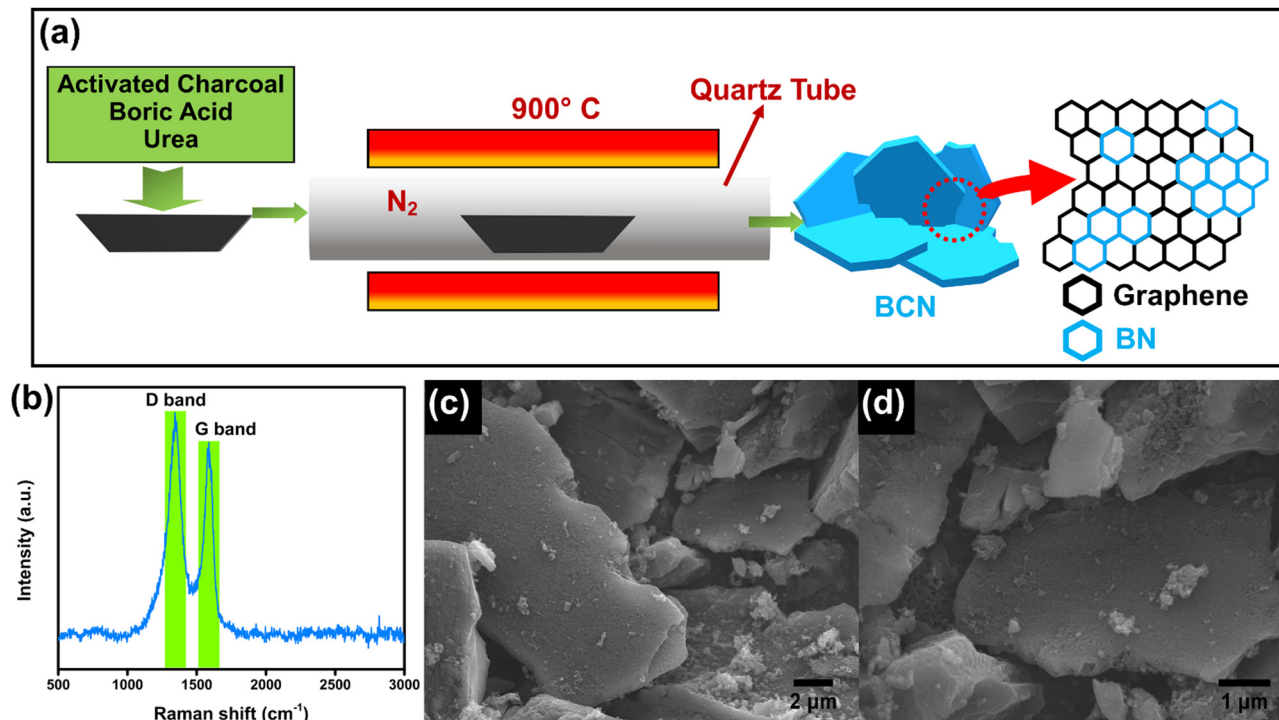


Fig. 5 (a) Schematic illustration of the synthesis of BCN sheets using the urea route, (b) Raman spectrum of BCN, (c) and (d) FESEM images showing the porous sheet like morphology of BCN.

The high-resolution B 1s spectrum provided in Fig. S4b (ESI[†]) depicts two distinct peaks located at 190.9 and 192.0 eV corresponding to B–C and B–N bonds. The C 1s spectrum of BCN was deconvoluted into three peaks at 283.8, 284.5 and 285.4 eV, representing C–B, C–C and C–N bonds, respectively (Fig. S4c, ESI[†]). The deconvoluted N 1s spectrum shows N–B and N–C bonding at binding energies of 398.4 and 400.1 eV, respectively (Fig. S4d, ESI[†]).³¹

2.4 Electrochemical assessment of BCN

Similar to S-VSe₂/CNT, the electrochemical performance of BCN is also analyzed in a 0.5 M K₂SO₄ electrolyte by employing a three-electrode cell. A cyclic voltammogram of BCN shows neat EDLC behaviour in a potential window of –1.0 to 0.0 V (Fig. 6a). From a lower scan rate to a higher scan rate, the BCN electrode maintained a rectangular shape without any deformation, showing excellent reversibility during electrochemical reactions. To further understand the charge storage mechanism of BCN, we segregated CV curves into capacitive and diffusive contributions using power law. The segregated CV profile of the BCN shows a complete dominance of capacitive contribution over diffusion due to the EDLC mechanism of the electrode. The capacitive contribution of the BCN electrode increases from 94.3% at 10 mV s^{–1} to 97% at 100 mV s^{–1} (Fig. 6b and c). Dominant capacitive performance is beneficial for the BCN electrode in terms of excellent cyclic stability and high-rate performance.

The GCD curves of the BCN electrode exhibited a nearly triangular shape based on the EDLC behaviour (Fig. 6d). The linear charge/discharge curves in current densities ranging

from 1 to 8 mA cm^{–2} suggest the excellent capacitive behaviour of the BCN electrode. The areal capacitance of the BCN electrode calculated from the GCD is shown in Fig. 6e. The BCN attained an areal capacitance of 115 mF cm^{–2} at a current density of 1 mA cm^{–2}, and the electrode showed a great rate capability by retaining almost 69% of its initial capacitance after an eight-fold increment in current density. The cyclic stability of BCN is measured, and the electrode exhibited a 93.7% capacitance retention after 5000 GCD cycles (Fig. 6f). During the cyclic study, we observed that the capacitance value exceeded 100% and decreased after reaching the maximum value. The increase in capacitance is quite obvious in porous materials owing to the entrapment of ions inside the pores and the enhancement of pore size during long-cycle operation.^{50,51} BCN showed near 100% coulombic efficiency during the cyclic study, which emphasises the high reversibility and exceptional electrochemical stability of the BCN electrode. The Nyquist plot of BCN reveals a capacitor-type behaviour with an *R*_{ct} value of 6.2 Ω (Fig. S5, ESI[†]).

2.5 Performance of S-VSe₂/CNT//BCN asymmetric supercapacitor

To evaluate the practical application of S-VSe₂/CNT and BCN, an asymmetric supercapacitor (ASC) was constructed with S-VSe₂/CNT as the positive electrode and BCN as the negative electrode (Fig. 7a). Before assembling, mass balance theory was implemented to attain the charge balance in both electrodes, *i.e.*, *m*_–/*m*₊ = 0.64. The potential stability of S-VSe₂/CNT//BCN ASC is studied at various potential windows, and the optimized

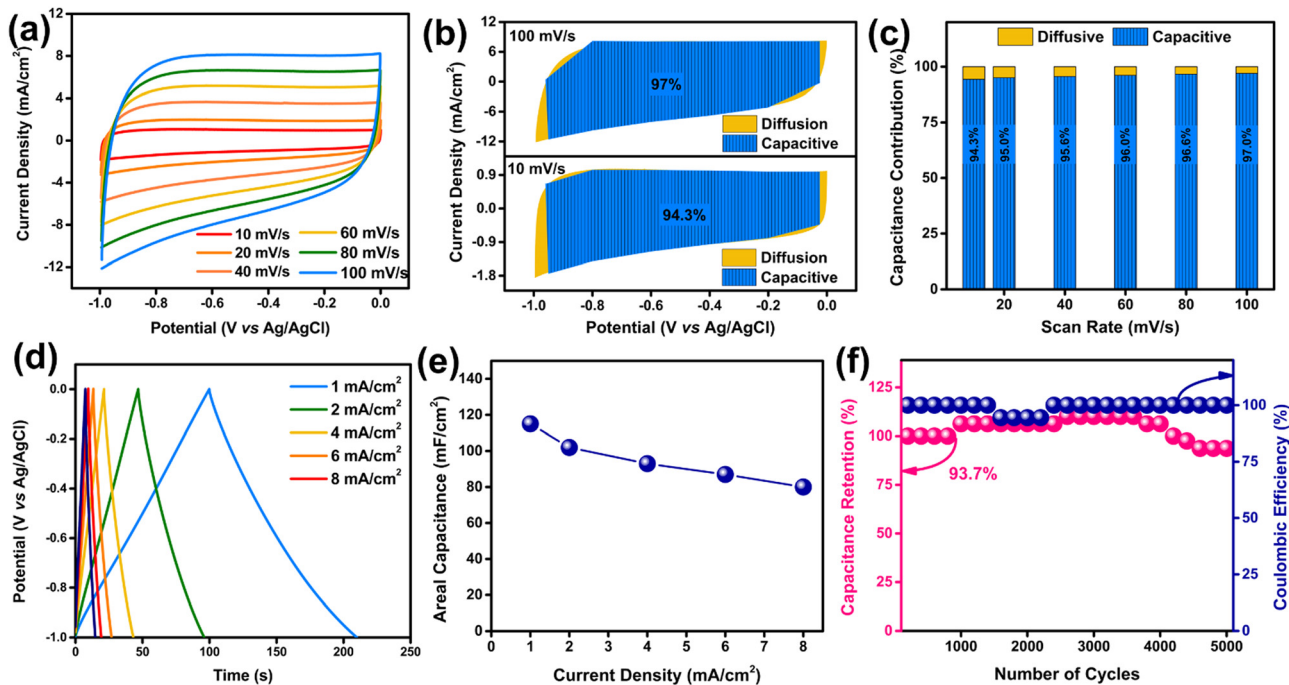


Fig. 6 Electrochemical measurements of BCN sheets in 0.5 M K_2SO_4 . (a) CV profile of BCN in scan rates ranging from 10 mV s^{-1} to 100 mV s^{-1} , (b) segregated CV curves of BCN into capacitive and diffusive regions at selected scan rates, (c) normalized capacitive and diffusive contributions in all the scan rates, (d) GCD curves of BCN in various current densities, (e) areal capacitance of BCN plotted against corresponding current densities and (f) cyclic performance of BCN electrode during 5000 cycles at a current density of 10 mA cm^{-2} .

working window of the device is set at 1.65 V (Fig. S6a, ESI[†]). The CV curve of ASC is analogous to that of S-VSe₂/CNT and BCN, showing that the working mechanism of ASC is an

admixture of pseudocapacitance and EDLC (Fig. 7b). During all the scan rates of the ASC, no considerable distortions are observed in the CV shape, signifying the excellent capacitive

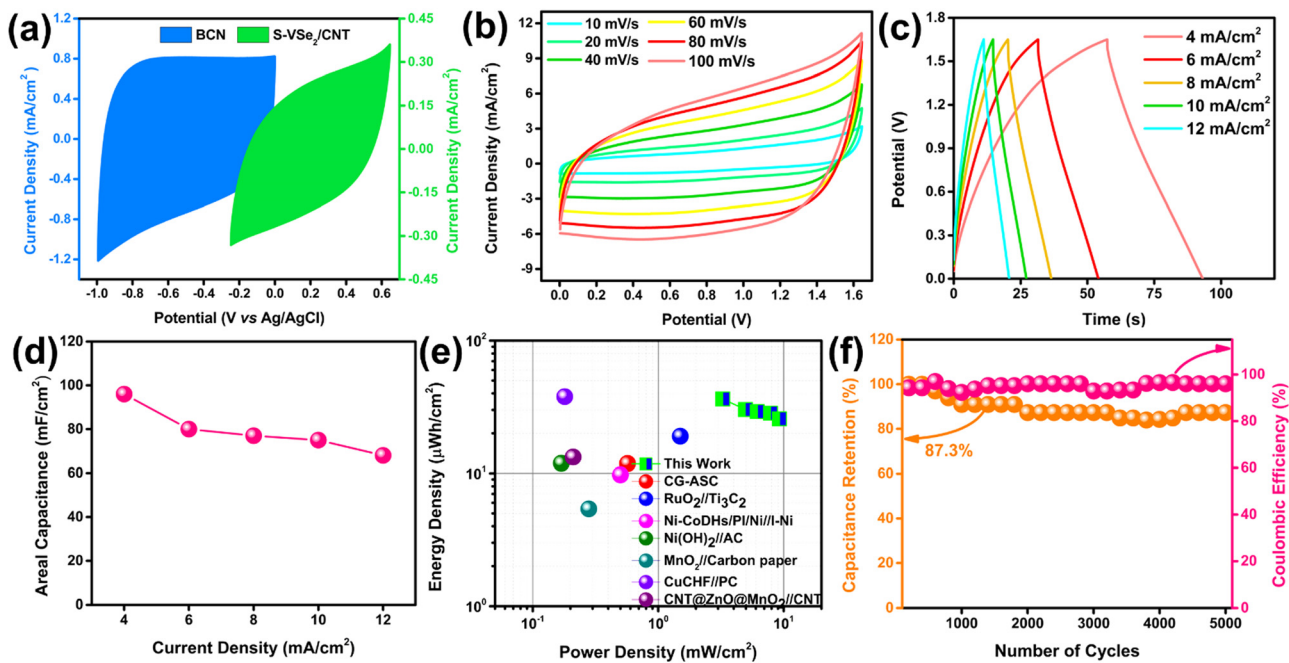
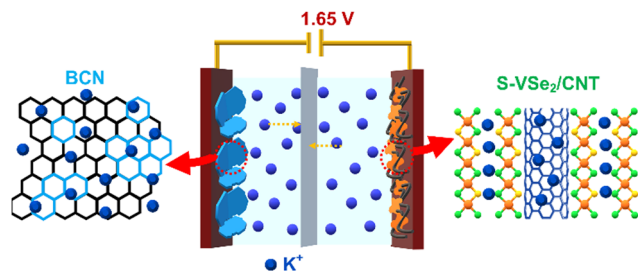


Fig. 7 Electrochemical performance of S-VSe₂/CNT//BCN asymmetric supercapacitor. (a) CV curves of S-VSe₂/CNT and BCN at 100 mV s^{-1} , (b) CV profile of S-VSe₂/CNT//BCN ASC in scan rates ranging from 10 mV s^{-1} to 100 mV s^{-1} , (c) GCD curves of S-VSe₂/CNT//BCN ASC in various current densities, (d) areal capacitance of the ASC plotted against current densities, (e) the Ragone plot of S-VSe₂/CNT//BCN ASC compared with reported ASCs and (f) cyclic performance of S-VSe₂/CNT//BCN ASC during 5000 cycles at 15 mA cm^{-2} .



Scheme 2 Schematic representation of the working of S-VSe₂/CNT//BCN ASC.

behaviour of the device. The GCD profiles of the S-VSe₂/CNT//BCN ASC show a near-symmetric shape without any plateaus, signalling the absence of diffusive contributions (Fig. 7c). The areal capacitance obtained from the GCD profiles of ASC is provided in Fig. 7d. The ASC delivered a high areal capacitance of 96 mF cm⁻² at a current density of 4 mA cm⁻² and retained 68 mF cm⁻² of its initial capacitance at a high current density of 12 mA cm⁻², exhibiting the high-rate capability of the device. The Ragone plot is important for evaluating the energy storage capability of a full cell. S-VSe₂/CNT//BCN ASC achieved a high energy density of 36.3 μW h cm⁻² at a power density of 3.2 mW cm⁻², and the ASC retained 70% of the initial energy density at a large power density of 9.2 mW cm⁻² (Fig. 7e). The energy storage performance of S-VSe₂/CNT//BCN ASC is comparable to many recently reported values^{52–59} (Table S2, ESI[†]). Fig. 7f illustrates the cyclic performance of ASC, exhibiting an 87.2% capacitance retention after 5000 charge/discharge cycles performed at a current density of 15 mA cm⁻² with a coulombic efficiency of more than 96%. Comparing the Nyquist plots of the ASC before and after stability, no considerable deviations in their shape were observed (Fig. S6b, ESI[†]). This corresponds to the electrochemical stability of the full cell. The working mechanism of S-VSe₂/CNT//BCN ASC is illustrated in Scheme 2.

3. Conclusions

In summary, S-doped VSe₂/CNT is successfully synthesized using a facile hydrothermal method. By applying sophisticated characterization techniques, the effects of S doping and CNT hybrid formation in VSe₂ are analysed. The substitutional doping of S atoms in the VSe₂ lattice can provide a considerable improvement in the charge storage capability of S-VSe₂, but this improvement is limited to a certain extent. The S-VSe₂/CNT hybrid as a positrode conferred excellent charge storage capabilities with remarkable cyclic stability compared to S-VSe₂ and pristine VSe₂. The enhanced electrochemical activity of S-VSe₂/CNT can be attributed to the synergistic effect between S-VSe₂ and CNT. BCN, an intriguing layered 2D material with EDLC behaviour prepared *via* a solid-state urea route, was electrochemically analysed as a negatrode. The material and electrochemical characterization provide insight into the chemical structure of BCN and its prodigious charge storage capability. An asymmetric supercapacitor was constructed with

S-VSe₂/CNT and BCN, showing exceptional energy density and power density along with long cycle life. The VSe₂/CNT//BCN ASC inherits all the individual advantages of each electrode material to deliver a promising ASC for next-generation energy storage applications.

Conflicts of interest

The authors declare no conflict of interest.

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