

Cite this: *Nanoscale Adv.*, 2021, 3, 486

Superior supercapacitive performance of $\text{Cu}_2\text{MnSnS}_4$ asymmetric devices

M. Isacfranklin,^a R. Yuvakkumar,^b *^a G. Ravi,*^a Dhayalan Velauthapillai,^b Mehboobali Pannipara^c and Abdullah G. Al-Sehemi^c

Renewable energy sources are considered the cornerstone of achieving a sustainable future for today's modern world. In the current research on energy storage devices, transition metal sulfides are being explored as an excellent electrode material. In this study, the electrochemical performance of $\text{Cu}_2\text{MnSnS}_4$ and its micro/nano-structural changes due to variations in the structure directing agents used in its fabrication were investigated. Synthesis of the sulfide sediments was confirmed *via* standard characterization techniques. The charge-storage electrochemical activity was explored *via* electrochemical characterization techniques. Furthermore, the electrochemical charge-storage behavior was enhanced by $\text{Cu}_2\text{MnSnS}_4$. The best cyclic stabilities obtained were 83%, 87%, and 89% of capacitive retention over 5000 cycles for AA, CA, and OA electrodes in three-electrode arrangements, respectively, while an 80.35% capacitance retention and 97.54% coulombic efficiency were achieved after 21 000 cycles in a fabricated device. The as-fabricated device exhibited an energy density of 27 W h kg^{-1} and power density of 759 W g^{-1} at a current density of 1 A g^{-1} in the two-electrode setup.

Received 16th September 2020

Accepted 9th November 2020

DOI: 10.1039/d0na00775g

rsc.li/nanoscale-advances

1. Introduction

Eco-friendly and efficient energy storage systems are emerging as an urgent need to meet the ever-increasing energy demands. Supercapacitors (SCs) in particular are attracting unparalleled attention because of their low weight, high power density, rapid charge discharge, extended cyclic ability, and robustness.^{1,2} However, the current carbon-based SCs are affected by a low energy density when compared to batteries. It is therefore attractive to have a wide range of applications for supercapacitors that can deliver high energy density, which has evolved into an essential requirement. It is still believed today for these supercapacitors that, it would be possible to improve the energy density by increasing the cell potential, which can be achieved by electrode combinations.^{3–5} Metal sulfides have many redox reactions to form good quality SCs with battery-type electrode materials and they are considered as positive electrode materials. Depending on the device, they may exhibit an extended potential window and higher energy density than conventional supercapacitors.^{6–10} Based on their electrochemical charge storage, generally pseudocapacitors (PCs) exhibit higher specific capacitance values than electric double layer capacitors (EDLCs).^{11,12} The reason is that in EDLCs, the

charges between the electrode and electrolyte interfaces are stored by processes such as the diffusion/charge accumulation process, whereas in pseudocapacitors, the charge is stored by redox reactions in the active substance (mass), such as the faradaic process.^{13,14} Carbon-aerogel, carbon nanotubes, carbon cloth, graphene, and activated carbon are used in EDLCs, whereas metal oxides/hydroxides/sulfides are extensively used as electrode materials in pseudocapacitors.^{15,16} Metal oxides/sulfides, such as MnCo_2O_4 , NiCo_2O_4 , FeCo_2O_4 , CoS , and Ni_3S_2 , have attracted more research attention because they provide significant beneficial characteristics, such as a high energy and power density, and more specific capacitance thanks to their rapid and reversible rich redox reactions on the electrode/electrolyte interface. Various transition metal sulfide electroactive materials have been explored for electrochemical capacitors (ECs) due to their rich valence and desirable chemical stability.^{17,18} Sulfides are considered to be excellent pseudocapacitive materials, especially thanks to their multiple redox reactions, improved electrical conductivity, and low band gap, which lead to their superior performance, better oxidation state, and longer cycling capability when compared to metal oxides/hydroxides.^{19,20} In general, the pseudocapacitance capabilities of these sulfides in alkaline solutions are exhibited in the faradaic reactions that are associated with the mutual transformation from ion diffusion into electrode materials with rich electroactive sites.

Cu_2XSnS_4 (X = Mn, Fe, Co, Ni, Zn, and Cd) (CXTS) materials have excellent structural and optical properties, absorption coefficients, and catalytic efficacy in redox reactions due to their

^aDepartment of Physics, Alagappa University, Karaikudi 630003, Tamil Nadu, India. E-mail: yuvakkumarr@alagappauniversity.ac.in; raviganesa@rediffmail.com

^bFaculty of Engineering and Science, Western Norway University of Applied Sciences, Bergen 5063, Norway

^cDepartment of Chemistry, King Khalid University, Abha, 61413, Saudi Arabia



multiple oxidation states.^{21,22} Among their most interesting benefits in practice are the production of materials with a high surface area, good porosity, and production of comparatively labor-intensive planar and conformal nanostructured layers using a simple, low cost, quick, and measurable process. The metal chalcogenides have attracted great research interest due to their enhanced electrochemical properties. Molecular metal chalcogenide species can be broken down from the bulk chalcogenide complex and formed as a result of the basic nucleophilic reactions involving the chalcogenide anions.^{23–25} The hydrothermal method is the best technique for designing a good quality micro/nano structure compared to all other methods. During the initial nucleation in the hydrothermal reaction, the acceleration of the endothermic reaction results in a disruption of the nuclei, which dissolves the urea ions, resulting in an insufficient activation surface energy and subsequent defects.^{26,27} Acidic acid, citric acid, and oxalic acid are all well-known structure directing agents that can be classified from dicarboxylic acid. They are also considered to be powerful reducing and chelating agent for metal cations. The acidic strength of oxalic acid is much greater than the other two, and therefore, it can lead to a better structure; thereby creating more ionic transport at the electrode/electrolyte interface, resulting in an improved electrochemical behavior.^{28,29} In this study, a solvothermal process was pursued utilizing a ethanol-mixed de-ionized water solution as the solvent. The structure directing agent's role in creating the Cu₂MnSnS₄ nanostructure could be visualized from the FE-SEM analysis. Characterization of the materials was performed by several physicochemical, chemical, and electrochemical techniques. Here, superior supercapacitive performances of Cu₂MnSnS₄ asymmetric devices could be reported. To the best of our knowledge, this was also the first study to investigate and report the use of Cu₂MnSnS₄ nanostructures fabricated based on the use of different directing agents for fabricating a positive electrode for supercapacitor applications.

2. Experimental

Copper(II) chloride dihydrate (CuCl₂·2H₂O, 99%), manganese(II) chloride tetrahydrate (MnCl₂·4H₂O, 99%), citric acid (C₆H₈O₇, 99%), thiourea (NH₂CSNH₂, 99%), and tin(IV) chloride pentahydrate (SnCl₄·5H₂O, 98%) were purchased. The solvothermal reaction was adapted to synthesize the Cu₂MnSnS₄ nanostructure. In a typical synthesis route, copper(II) chloride dihydrate (1 mole), manganese(II) chloride tetrahydrate (0.5 mole), tin(IV) chloride pentahydrate (0.5 mole), and structure directing acidic (AA), citric (CA), or oxalic (OA) acid (1 mole) were dissolved in 80 mL in a 1 : 1 ratio with ethanol and de-ionized water solvents under magnetic stirring until an homogeneous solution appeared. Thereafter, thiourea (0.4 mole) solution was added and the mixture was put in an autoclave at 180 °C for 12 h. The samples were collected after a thorough wash with de-ionized water, ethanol, and methanol with centrifugation at 4000 rpm. Finally, the collected samples were separately placed in a hot air oven for drying. The resultant products were used in the different characterizations.

The working electrode was formulated by mixing Cu₂MnSnS₄ (prepared with AA, CA, or OA) as an active cathode material, activated carbon as the anode material, and polyvinylidene fluoride (PVDF) as a binder in a 75 : 20 : 5, weight ratio. Then, all three materials were ground for at least 30 min. Thereafter, the mixed electrode material was dispersed in the solvent *N*-methyl-2-pyrrolidone (NMP) and ground for 1 h to prepare the electrode slurry. Well-cleaned nickel foam was employed as a substrate (1 × 2 cm²). A 1 × 1 cm² section of nickel foam was coated with the prepared slurry using a mercury premium art small size brush. The coated nickel foam was dried in a hot air oven at 80 °C for 12 h and the loading material was adjusted to 2 mg cm⁻² on the electrode surface. These Cu₂MnSnS₄ (AA, CA, OA) electrodes, platinum (Pt) wire, and Ag/AgCl were utilized as the working, counter, and reference electrodes. Electrochemical measurements in 2 M KOH electrolyte were performed using a BioLogic potentiostat. CVs at different scan rates ranging from 10 to 100 mV s⁻¹ were recorded at a 0.6 V constant potential, and 0.5 to 5A g⁻¹ current densities at 0.5 V potential were utilized for the galvanostatic charge discharge (GCD) readings. The electrochemical impedance spectroscopy (EIS) study results were analyzed by Nyquist plots and fitted by *Z* fit analysis. Eqn (1) was employed to estimate the specific capacitance from the GCD profile, and similarly the two-electrode (full-cell) system was set up with the two equal loadings mass balanced using the formula $\frac{Q^+}{Q^-} = \frac{m^+ \times C_{sp} \times \Delta V^+}{m^- \times C_{sp}^- \times \Delta V^-}$ and then coated on to the positive and negative electrodes. The full-cell device parameters such as energy (*E*) and power (*P*) density, were measured using eqn (2) and (3). Eqn (4) was used to calculate the coulombic efficiency of the materials from the GCD curves.

$$C_s = \frac{I \times \Delta t}{m \times \Delta V} \quad (\text{F g}^{-1}) \quad (1)$$

$$E = \frac{C \times \Delta V^2}{2 \times 3.6} \quad (\text{W h kg}^{-1}) \quad (2)$$

$$P = \frac{3600 \times E}{\Delta t} \quad (\text{W kg}^{-1}) \quad (3)$$

$$\eta\% = \frac{t_d}{t_c} \times 100\% \quad (4)$$

where, *t_d* and *t_c* are the discharging and charging time, respectively.^{30,31}

2.1 Trasatti method analysis

The Trasatti method was first used to identify the EDLC behavior and PC behavior by the eminent scientist S. Trasatti in 1990 using the formula $q_T = q_i + q_o$; $C_T = C_i + C_o$, where q_T is the total voltammetric charge, q_i is the charge stored at the inner surface, q_o is the charge stored at the outer surface, and C_T is the total capacity, which is the sum of the pseudocapacity and the electric double layer capacity. In general, in the outer surface $V \rightarrow \infty$; therefore it just allows the surface process to happen. Hence, the more accessible area is in the electrode electrolyte interface. In the same way, at the inner surface, $V \rightarrow 0$; therefore



it gives sufficient time for the diffusion and for more ions to react. Hence, it has less accessible area.

Step 1. Collect cyclic voltammogram at various scan rates (10 to 100 mV s^{-1}).

Step 2. Calculate the specific capacity (C g^{-1}) at each scan rate using the following formula,

$$C_q = \frac{S}{2\nu \times \Delta U} \quad (5)$$

where C_q denotes the specific capacity (C g^{-1}), S is the area enclosed by the CV curve, ν is the scan rate (mV s^{-1}), and m is the active mass in the electrode surface.

Step 3. Estimation of the total capacity (C_T).

Let us assume semi-infinite linear ion diffusion (*i.e.*, ions randomly diffuse from the bulk electrolyte to the electrode/electrolyte interface, when $V \rightarrow 0$, $q \rightarrow q_T$). Based on the Cottrell equation, a linear correlation between the reciprocal of the calculated specific capacity ($1/C_q$) and the square root of scan rates ($\nu^{1/2}$) gives the maximum total capacity by extrapolation to the Y intercept.

$$\frac{1}{q(\nu)} = \text{const } \nu^{1/2} + \frac{1}{q_T} \quad (6)$$

Multiply both sides by dU :

$$\frac{dU}{q(\nu)} = \text{const } \nu^{1/2} + \frac{dU}{q_T} \quad (7)$$

$$\text{i.e., } \frac{1}{C(\nu)} = \text{const } \nu^{1/2} + \frac{1}{C_T} \quad (8)$$

Step 4. Estimation of the charge stored at the outer surface (C_o).

In the same way, assuming semi-infinite linear ion diffusion (*i.e.*, ions randomly diffuse from the bulk electrolyte to the electrode/electrolyte interface, when $V \rightarrow \infty$, $q \rightarrow q_o$), based on the Cottrell equation, a linear correlation between specific capacity (C_q) and the reciprocal of square root of scan rates ($\nu^{-1/2}$) gives the charge stored at outer surface (C_o) by extrapolation of the intercept to the Y axis.

$$q(\nu) = \text{const } \nu^{-1/2} + q_o \quad (9)$$

Divide both sides by dU

$$\frac{q(\nu)}{dU} = \text{const } \nu^{-1/2} + \frac{q_o}{dU} \quad (10)$$

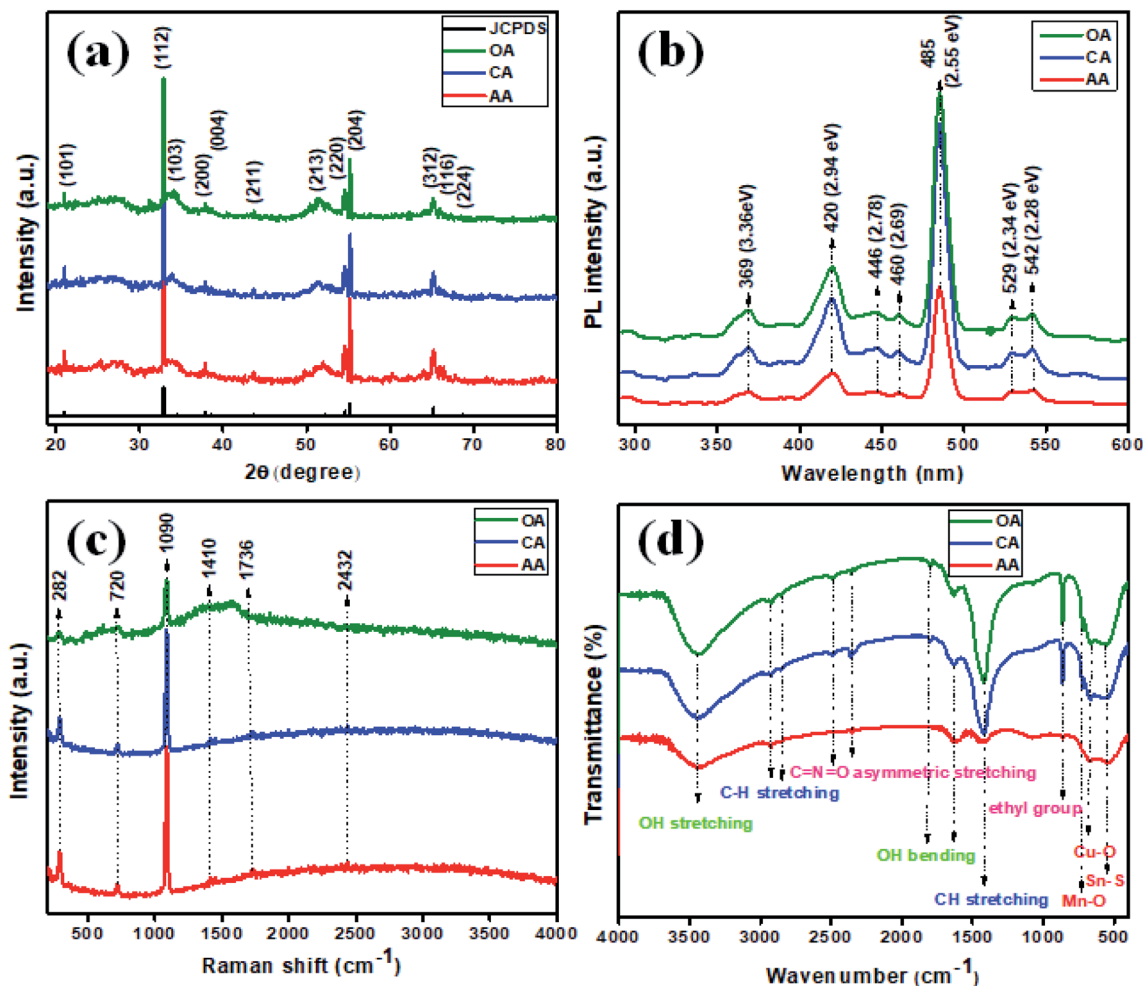


Fig. 1 (a) XRD, (b) photoluminescence, (c) Raman, (d) FTIR spectra of $\text{Cu}_2\text{MnSnS}_4$.



$$i.e., C(v) = \text{const } v^{-1/2} + C_o \quad (11)$$

Step 5. Estimation of the inner surface (C_i).

Based on the Cottrell equation, the maximum capacity (C_T) and charge stored at the outer surface (C_o) could be calculated. The maximum capacity (C_T) is the sum of the inner (C_i) and outer surface (C_o), i.e., $C_T = C_i + C_o$, where C_T is the total capacitance, C_i is responsible for the maximum pseudocapacitance (C_i), and C_o corresponds to the maximum electric double layer capacitance. Subtracting C_o from C_T gives the maximum pseudocapacitance (C_i).

$$C_o = C_T + C_i \quad (12)$$

Step 6. Estimation of the percentage capacitive contribution.

$$C_i \text{ (%) } = \frac{C_i}{C_T} \times 100\% \quad (13)$$

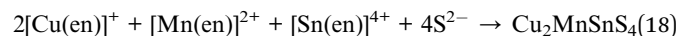
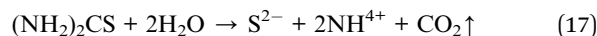
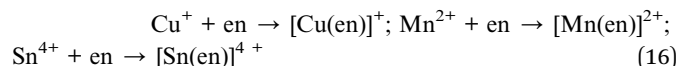
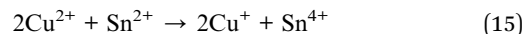
$$C_o \text{ (%) } = \frac{C_o}{C_T} \times 100\% \quad (14)$$

3. Results and discussion

The obtained material crystal structure was characterized using X-ray diffraction analysis (Fig. 1(a)), wherein the 2θ values of 20.99°, 32.89°, 34.17°, 37.83°, 38.62°, 43.66°, 51.57°, 54.54°, 55.23°, 65.21°, 66.21°, and 68.77° corresponded to the (hkl) planes of (101), (112)*, (103), (200), (004), (211), (213), (220), (204), (312), (116), and (224) respectively (JCPDS card: 51-0757). The good crystalline nature was evidenced in all the samples, as suggested by the peaks observed. The high intensity peak present at 32.89° with the plane orientation of (112)* inferred the as-fabricated $\text{Cu}_2\text{MnSnS}_4$ had a polycrystalline nature. The structural parameters of the $\text{Cu}_2\text{MnSnS}_4$ microstructure are tabulated in Table 1.

The growth mechanism of the $\text{Cu}_2\text{MnSnS}_4$ product in the presence of the chloride precursor in the solvothermal reaction mixture in the well-closed Teflon tube produced Cu^{2+} , Mn^{2+} , Sn^{4+} , and S^{2-} . Most importantly, when the compounds were combined with thiourea, sulfur ions were released in the autoclave during the reaction mixture. Here, the S^{2-} ions reacted with the metal ions Cu, Mn, and Sn during the nucleation

formation of $\text{Cu}_2\text{MnSnS}_4$, as suggested by the following equations.^{32,33}



The small crystalline size and poor crystallinity of the material were explored for improving the electrochemical performance. This is because of the fact that poor crystalline materials create more ion transport channels than crystalline materials, which is an important factor in electrochemical supercapacitors. Photoluminescence was utilized as an eminent tool to investigate the emission behavior of the synthesized $\text{Cu}_2\text{MnSnS}_4$ (AA, CA, OA). In Fig. 1(b), PL emission bands could be observed at the wavelengths: 369 (3.36), 420 (2.94), 446 (2.78), 460 (2.69), 485 (2.55), 529 (2.34), and 542 nm (2.28 eV). Here, a higher intensity suggests a faster recombination rate. Seven photoluminescence peaks could be detected and with almost the same PL emission bands, expect for the intensity. The strong and broad blue emission band located at 485 nm was in good agreement with the Cu-S hierarchical structures, which absorb more photons to produce electron-hole pairs, which may start from the excitonic band edge or free-charge carrier transition at 2.55 eV. Weak green emission bands were observed at 529 and 542 nm. The blue emission band at 420 nm had a slightly broad band, which represented defects present in the prepared samples. The emission peaks assigned at 446 and 460 nm indicated different defects, which largely depended on the high density of sulfur and tin vacancies.^{34,35} The lattice vibrations of a material can be measured using Raman spectroscopy and so was used here, as shown in Fig. 1(c). The characteristic Raman band occurred at 282 cm^{-1} , which was attributed to the Sn-S phases in $\text{Cu}_2\text{MnSnS}_4$. The Cu-S stretching mode vibration was detected at 449 cm^{-1} . The small peak located at 720 cm^{-1} revealed a second-order Raman scattering mode of the prepared samples. In the synthesized product, the main broad peak existed in the Vis-NIR region at 1090 cm^{-1} . C-C stretching vibrations were observed for the remaining bands present at 1736 and 2432 cm^{-1} .³⁶ FTIR spectrometry is a well-known technique used to characterize the molecular fingerprint region of synthesized products, and so was used here (Fig. 1(d)). Initially, a small quantity of the sample was mixed with potassium bromide (KBr) and finely grounded and then pressed well into a hydraulic press to be converted into a pellet shape. The acquired spectrum was within the range of 4000 to 400 cm^{-1} . The characteristic spectrum 3448 cm^{-1} corresponded to the OH stretching vibration of water molecules observed on the sample. The vibrations at 1639 and 1820 cm^{-1} were because of the bending vibrations of OH water molecules. The absorption bands located at 2853 and 2929 cm^{-1} were due to the CH stretching vibration mode. The

Table 1 Structural parameters of the $\text{Cu}_2\text{MnSnS}_4$ microstructure

Physical Quantity (Symbol) (Units)	Value
Lattice parameter (a) (Å)	$a = 5.513, c = 10.826$
Crystalline size (nm)	56
Microstrain (ϵ) $\times 10^{-3}$	0.618
Dislocation density (δ) $\times 10^{14}$	3.18
Cell volume (V) (nm^3)	329.11



asymmetric stretching vibration of C=N=O was indicated by the vibration band at 2491 cm^{-1} . The absorption vibration bands that existed in the fingerprint region indicated the metal oxide and metal sulfide vibrations. The bands at 729 , 664 , and 554 cm^{-1} signified the Mn–O, Cu–O, and Sn–S vibrations.^{37,38}

Fig. 2 shows the morphological images of the prepared $\text{Cu}_2\text{MnSnS}_4$ with using different structure directing agents, *i.e.*, acidic acid (AA), citric acid (CA), and oxalic acid (OA), as studied using FE-SEM analysis. Irregular shapes with non-uniform size

morphologies were found in the AA and CA samples, whereas well-formed flower shapes were found with the addition of oxalic acid, which exhibited a strong role in the flower-shape formation. The thickness of a few micrometer of the 2D thick plate-like petals was loosely covered with rose-like thrones. The structure of the flower shape facilitates high ionic transport and increases the accessible specific surface area. However, the petals in the flower were tightly stacked and appeared as thin pieces.³⁹ It was clearly evidenced that when using the structure

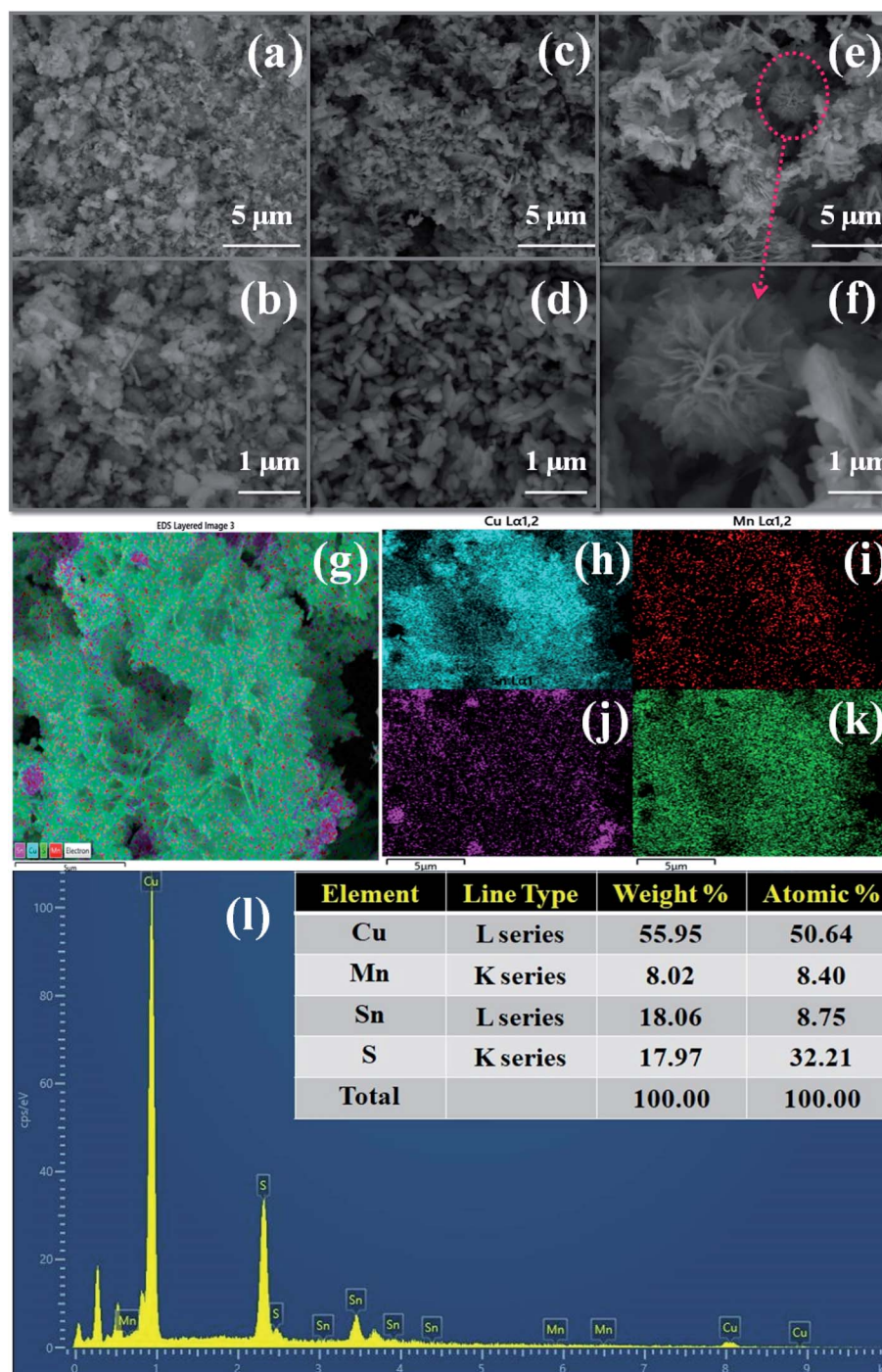


Fig. 2 $\text{Cu}_2\text{MnSnS}_4$ morphology (a and b) AA, (c and d) CA, (e and f) OA, (g–k) EDS mapping, (l) EDAX spectrum of the $\text{Cu}_2\text{MnSnS}_4$ –OA sample.



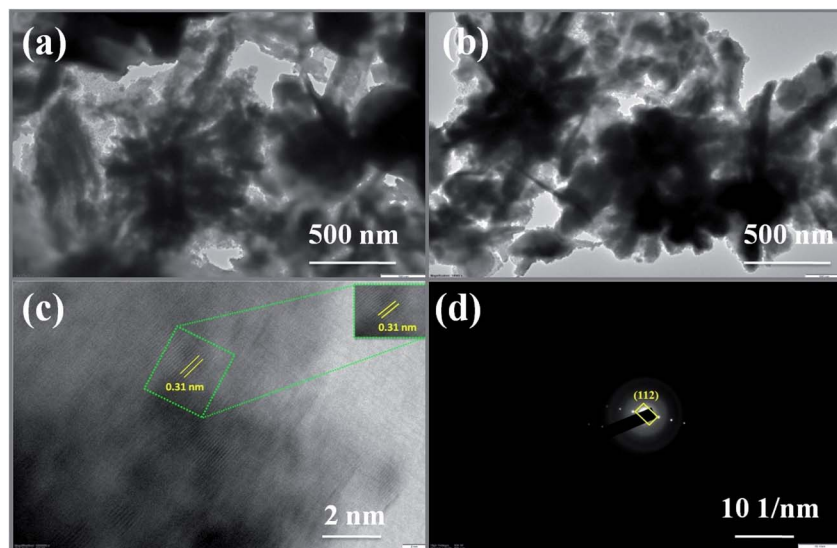


Fig. 3 (a and b) TEM images, (c) lattice fringes, (d) SAED pattern of $\text{Cu}_2\text{MnSnS}_4\text{-OA}$.

directing agents AA and CA in constant hydrothermal reactions, the particles agglomerated to form irregular flakes (Fig. 2(a–d)), while OA exhibited a fine flower-shaped morphology, as shown in Fig. 2(e and f). When compared to the AA and CA samples, the OA samples in the ethanol and distilled water mixture in the solvothermal route co-ordinate metal ions to urea and so are stronger combined. However, the petals in the flowers are stacked tightly, resembling thin slices. Initially, nanocrystals of $\text{Cu}_2\text{MnSnS}_4$ are formed as the seeds for microcrystal formation. The aggregated nanocrystals become larger and they coalesce to

grow into flakes due to the CA addition in the hydrothermal reactions. The hydrogen bond interacted with urea to form self-assembled flower-like microstructures by the addition of oxalic acid at a constant temperature. By the proper selection of structure directing agents, the sheet-like structures self-assembled and coalesce into a flower-shaped morphology. During this formation, the surface energy decreased.⁴⁰

Elemental analysis or chemical characterization of an area can be studied using EDS analysis. Fig. 2(g–k) shows the EDS SEM mapping images of the $\text{Cu}_2\text{MnSnS}_4\text{-OA}$ sample. The

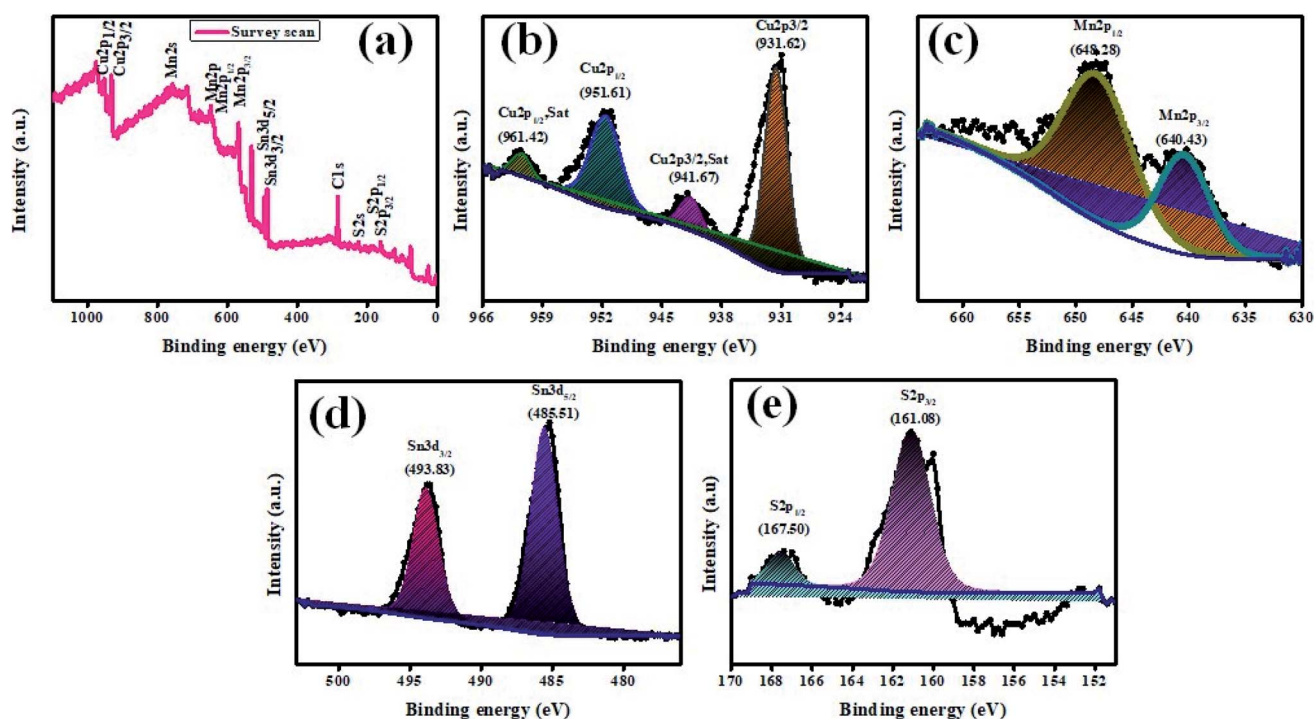


Fig. 4 XPS images of $\text{Cu}_2\text{MnSnS}_4$.



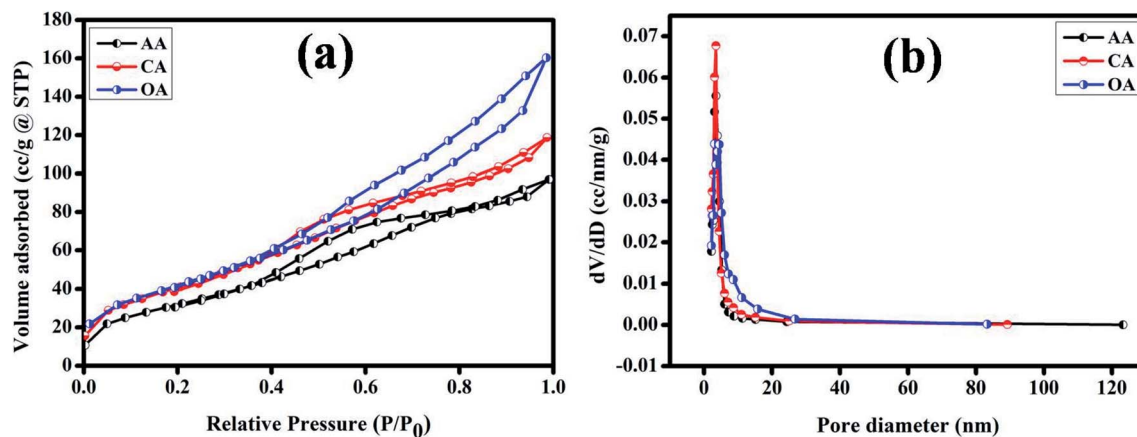


Fig. 5 (a) N_2 adsorption–desorption (b) pore-size distribution of the Cu_2MnSnS_4 –(AA, CA, OA) electrodes.

higher peak in a spectrum revealed a higher concentration in the elemental composition. Fig. 2(l) shows the EDAX spectrum of the Cu_2MnSnS_4 –OA electrode, which was composed of Cu, Mn, Sn, and S elements with good atomic and weight percentages as shown in the inset of Fig. 2(l). The elemental purity of the samples was confirmed by the absence of impurity peaks.⁴¹ Fig. 3 presents the TEM images of the synthesized Cu_2MnSnS_4 nanostructure with the addition of oxalic acid. Non-uniform flower-like structures were located in many places with sizes in the 1–3 μm range. The well-resolved lattice fringe distances were approximately 0.31 nm, which well matched with the (112) crystal plane for the Cu_2MnSnS_4 –OA electrode, as shown in Fig. 3(c). The selected area electron diffraction (SAED) pattern of the Cu_2MnSnS_4 –OA electrodes showed the good crystallinity of the sample (Fig. 3(c)), which correlated with the X-ray results.

The elemental constituents present in the product Cu_2MnSnS_4 were studied using XPS (Fig. 4). Fig. 4(a) presents the survey spectrum of the as-obtained Cu_2MnSnS_4 compound. The Cu binding energy species are shown by major peaks at 931.62 eV ($Cu2p_{3/2}$) and 951.61 eV ($Cu2p_{1/2}$) and the corresponding satellite peaks at 941.67 eV ($Cu2p_{3/2}$, Sat) and 961.42 eV ($Cu2p_{1/2}$, Sat) (Fig. 4(b)). Here, the peak splitting of 19.99 eV represents the Cu(I) configurations. Two 2p peaks appeared at 648.28 eV for $Mn2p_{1/2}$ and at 640.43 eV for $Mn2p_{3/2}$ (Fig. 4(c)). The peak splitting of 7.85 eV indicated the Mn(II) configuration. The binding energies located at 485.51 and 493.83 eV were due to $Sn3d_{5/2}$ and $Sn3d_{3/2}$, respectively (Fig. 4(d)). The peak splitting of 8.52 eV indicates the Sn(IV) configuration. The binding energies from 160 to 164 eV were related to the sulfide phases (Fig. 4(e)). $S2p_{3/2}$ and $S2p_{1/2}$ showed peaks at 161.08 and 160.01 eV, respectively, with the peak splitting of 1.07 eV.⁴²

The surface area ($m^2 g^{-1}$) and pore-size ($cm^3 g^{-1}$) distribution of the synthesized Cu_2MnSnS_4 electrodes were measured using Brunauer–Emmett–Teller (BET) analysis, as displayed in Fig. 5(a). All the fabricated electrodes exhibited type IV isotherms, which are a general distinctive isotherm of mesoporous specimens (Table 2). The Cu_2MnSnS_4 –OA electrodes provide high electrolyte contact at the electrode and electrolyte

interface because of its large surface area. This leads to better ionic transport for faradaic reactions than the other two electrodes. The pore-size distribution was observed employing the Barrett–Joyner–Halenda (BJH) approach, which suggested that the entire prepared electrode had pore sizes in the 3 to 50 nm range, which clearly signified that the three samples were in the mesoporous range (Fig. 5(b)). Furthermore, the high specific surface area and mesoporous nature of the Cu_2MnSnS_4 –OA electrode allows more specific parts of the electrolyte to diffuse effectively at the electrode with minimum internal resistance.⁴³

Electrochemical studies, such as CV, GCD, and EIS, were used to investigate the charge-storage performances of Cu_2MnSnS_4 (AA, CA, OA) in three-electrode setups. Fig. 6(a–c) show typical CV plots from 10 to 100 $mV s^{-1}$. The pair of redox reactions that are invisible in electrical double layer capacitors are evident in the CV curves and confirmed its battery-type behavior, which is the key characteristic of the pseudocapacitive mechanism.⁴⁴ The good redox peaks indicated charge storage derived from the Faraday process in battery-type electrodes. The apparent changes at high scan rates showed the improved rate performances of the electrode.⁴⁵ A pair of oxidation and reduction reactions of the anodic and cathodic voltage peaks at 10 $mV s^{-1}$ scan rate was observed at $E_a = 0.50, 0.45, 0.41$ V and $E_c = 0.38, 0.33, 0.28$ V for AA, CA, and OA, respectively. When the scan rate was increased from 10 to 100 $mV s^{-1}$, the redox peaks moved toward higher and lower energies and the corresponding peak current (i_p) also increased.

At low scan rates, the electrolyte ions and active species have sufficient time to interact with each other. This increases the probability of redox reactions occurring; nevertheless, at high

Table 2 BET analysis parameters

Product	BET surface area ($m^2 g^{-1}$)	Cumulative pore volume ($cm^3 g^{-1}$)	Pore diameter (nm)
Cu_2MnSnS_4 –AA	143.369	0.157	3.479
Cu_2MnSnS_4 –CA	166.025	0.186	3.492
Cu_2MnSnS_4 –OA	179.276	0.254	3.091



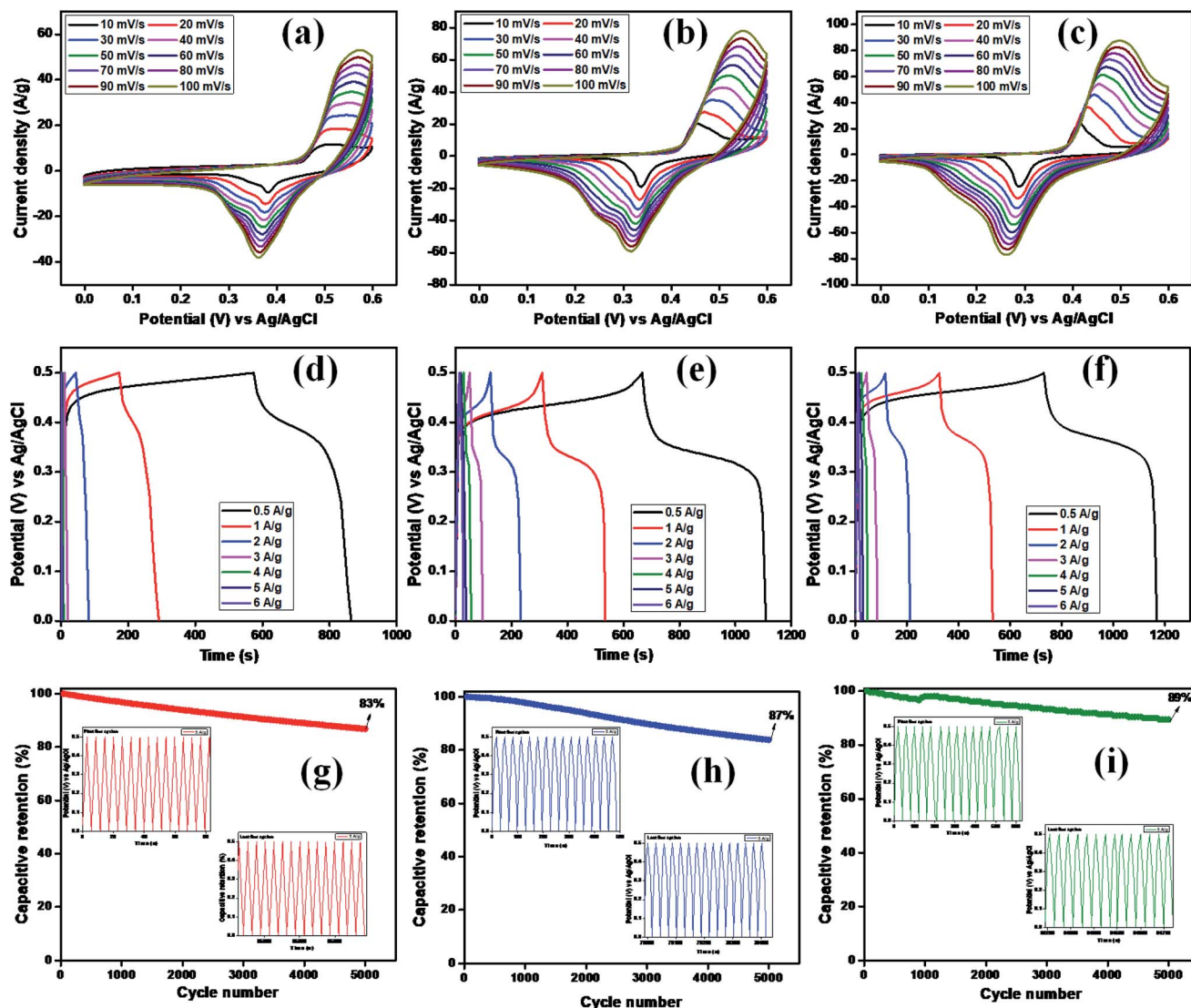


Fig. 6 (a–c) CV, (d–f) GCD, (g–i) capacitive retention of $\text{Cu}_2\text{MnSnS}_4$ (AA, CA, OA).

scan rates, the interaction of ions is not perfect due to their short residence time, which limits the different reactions between them. Cyclic voltammograms are affected by diffusion-controlled and kinetic-controlled processes. If a redox process is only affected by diffusion, the peak potential should generally be independent of the scan rate. Nevertheless, in contrast, if the electrode kinetics is dominant, the peak potential is affected by the voltammetric changes due to the increasing scan rates. Furthermore, the nature of the redox process can be easily identified by the cathodic and anodic

shifts. In view of the fact that this electrode material exhibited a battery-like behavior, there should be an apparent change in the peak shift.⁴⁴

Furthermore, this reduces the reaction kinetics because of the material polarization and ohmic resistance during the redox process, resulting in the formation of reversible peaks. The anodic peak current increased when the scan rate increased, which also led to an increase in the CV curves area directly proportional to the specific capacity. In all three samples, the lower scan rate (10 mV s^{-1}) exhibited a higher capacity, which

Table 3 Specific capacity of the different scan rates

Scan rate (mV s^{-1})	10	20	30	40	50	60	70	80	90	100
Specific capacitance (C g^{-1})–AA	268	198	165	143	127	114	103	95	88	83
Specific capacitance (C g^{-1})–CA	294	237	225	217	207	194	182	170	159	149
Specific capacitance (C g^{-1})–OA	302	285	279	272	262	249	237	224	213	204



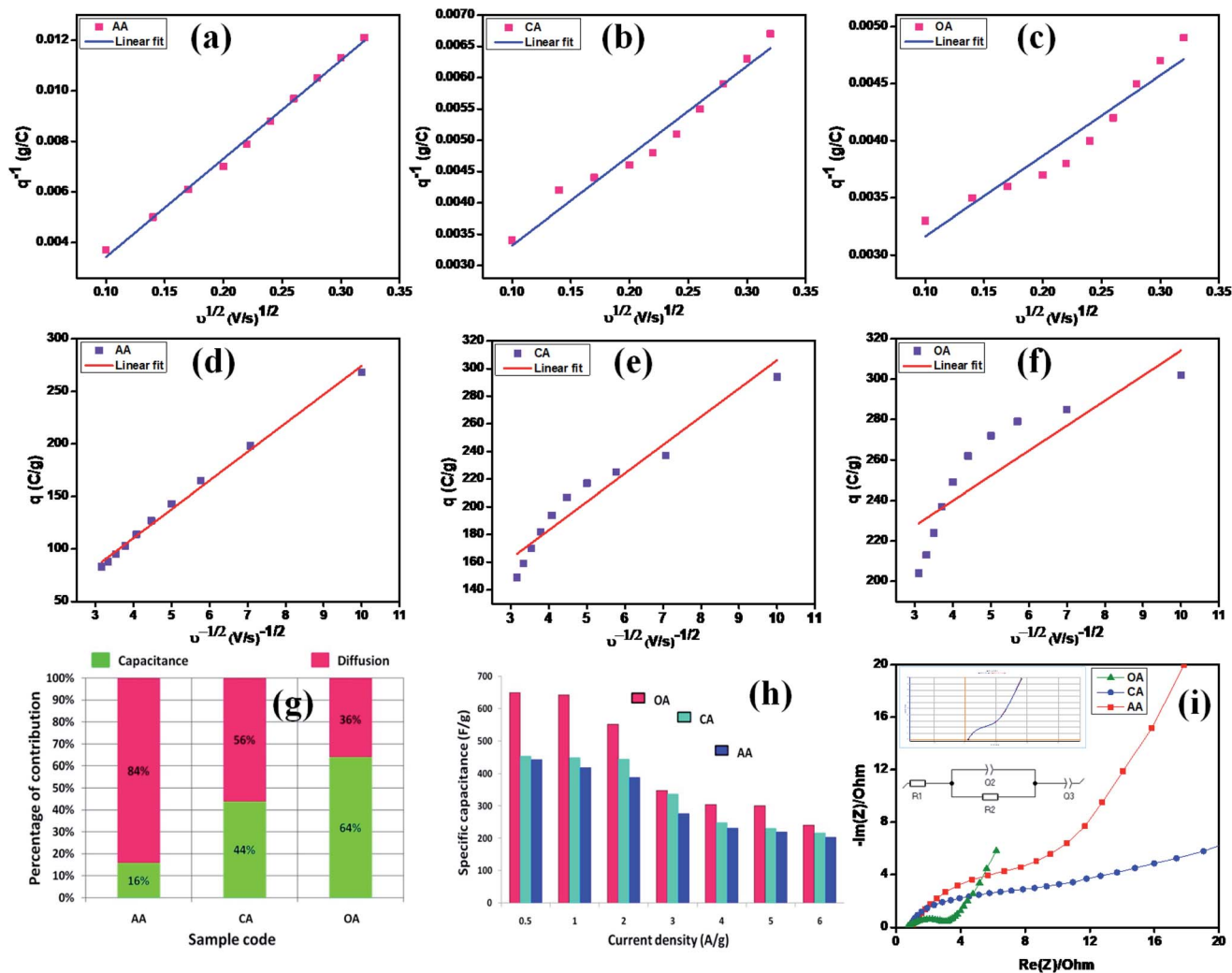


Fig. 7 (a and d), (b and e), and (c and f) linear fits of C vs. $v^{-1/2}$ and C^{-1} vs. $v^{1/2}$ for AA, CA, and OA, (g) capacitive (EDLC) and diffusive (PC) charging derived from Trasatti's method, (h) specific capacitance from GCD, and (i) Nyquist plot.

gives sufficient time for the ions to diffuse and react in the electrode/electrolyte interface. As a result, there is less accessible area occupied at the lower surface of the electrode for the electrochemical reaction process to occur.⁴⁶ The calculated specific capacity values are summarized in Table 3. The OA electrode displayed the highest specific capacity, which may be due to the well-arranged structure of the samples as proven in the FE-SEM analysis. This is because of the obtained nanostructures, which enhance the electrode specific surface area and rational pore-size distribution and greatly enhances the specific capacity, as clearly evidenced from the BET surface analysis.

The specific capacity, stability, and time reversibility of the $\text{Cu}_2\text{MnSnS}_4$ electrode material were studied by GCD characterization study. Fig. 6(d–f) display the GCD curves comparison of the prepared as-fabricated $\text{Cu}_2\text{MnSnS}_4$ electrode materials. The highest reversible time was observed for the OA sample than AA and CA. The calculated specific capacitance values for OA, CA, and AA were 649, 453, and 442 F g^{-1} at 0.5 A g^{-1} . The cone

diagram displays the specific capacitance values at different current density ranges. The long-term cyclic stability up to 5000 cycles was assessed and capacitive retentions of 83%, 87%, and 89% were achieved for the AA, CA, and OA electrodes, respectively (Fig. 6g–i). Due to the electrode properties, such as internal resistance, polarization, and electrolytic ions insertion, and removal due to mechanical stress, its specific capacitance

Table 4 EIS plot Z fit analysis

Parameter	Equivalent circuit: $R_1 + Q_2/R_2 + Q_3$			Unit
	OA	CA	AA	
$R_1 = R_s$	0.623	0.740	0.940	Ohm
Q_2	0.009	0.002	0.006	$\text{F s}^{(a^{-1})}$
a_2	0.467	0.706	0.739	—
$R_2 = R_{ct}$	3.106	6.850	8.231	Ohm
Q_3	0.209	0.079	0.079	$\text{F s}^{(a^{-1})}$
a_3	0.749	0.401	0.718	—



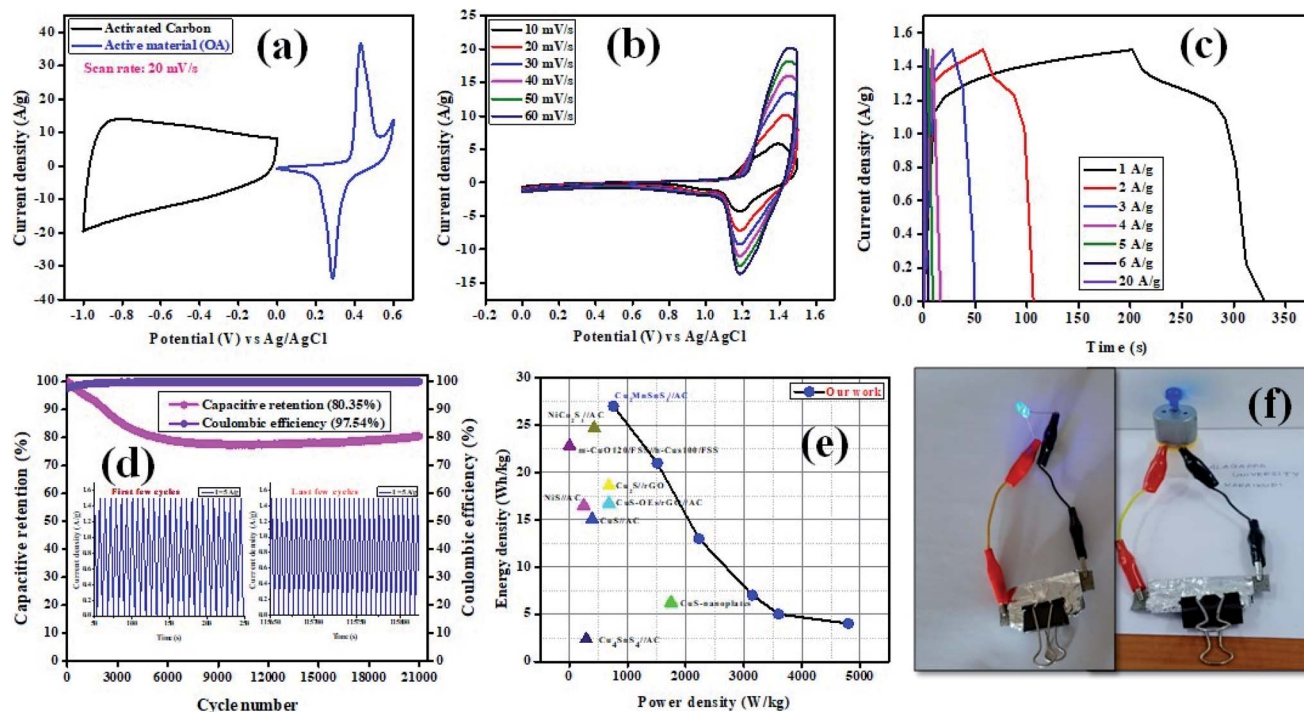


Fig. 8 (a) CV curves of $\text{Cu}_2\text{MnSnS}_4\text{-OA}$ and AC electrodes at 20 mV s^{-1} in a three-electrode system, (b) CV curves, (c) GCD curves, (d) capacitive retention and coulombic efficiency of the ASC device, (e) Ragone plot with other literature values, (f) practical demonstration of an ASC device.

value decreases due to a high current density.⁴⁷ According to the CV and GCD studies, the $\text{Cu}_2\text{MnSnS}_4\text{-OA}$ electrode may be preferred as a suitable electrode material.

The CV studies further indicated that the redox reaction is a diffusion-controlled process, which may be the main cause of the pseudocapacitive behavior with the increasing scan rates. The two differentiation methods, namely (i) Trasatti method and (ii) Dunn method, were then used to analyze the diffusion-controlled faradaic and non-diffusion-controlled battery-type processes of the synthesized electrodes.⁴⁸ In the Dunn method, the peak current (i_p) vs., the scan rate gives the power law $i_p = av^b$. The linear fit slope values give the 'b' value. A value of $b = 0.5$ means diffusion-controlled faradaic process, while $b = 1$ means a non-diffusion-controlled process due to surface redox reactions. Among the two studied methods, the Trasatti method is the simplest one to describe the mechanism of the electrode. Fig. 7(a, d, b, e, c and f) show the linear fit of C vs. $v^{-1/2}$ and C^{-1} vs. $v^{1/2}$ for the AA, CA, and OA samples, respectively. The non-linear behavior of the graphs explored was due to the ohmic drop of the intrinsic resistance.⁴⁹ The specific capacities of the AA, CA, and OA samples with their EDLC (capacitive) and pseudocapacitive (diffusive) behavior are shown in Fig. 7(g). The AA sample exhibited 16% and 84% capacity derived from the EDLC and PC contributions, while CA offered 44% and 56% capacity derived from the EDLC and PC contributions, and OA delivered 64% and 36% capacity for the EDLC and PC.

The interfacial structure and its kinetic response due to electrical variation were studied by electrochemical impedance spectroscopy. The opposition force to an electrical current is known as impedance and it is denoted by the unit of resistance

(Ω). There are three factors that determine the concepts of resistance used in an AC circuit: (i) the alternating current (AC) and voltage in phase, (ii) frequency independent, and (iii) all current and voltages. Nevertheless, the majority of these failed because the phase angle was not zero and almost in all cases capacitive and/or inductive effects were present at almost all the frequencies. Hence, it becomes an issue of frequency dependence. In these cases, the concept of impedance was used to denote the electrical current opposition in AC circuits.⁵⁰ Here, EIS spectra were plotted by the real vs. the imaginary part of the impedance, which determines the inter phase resistance between the working electrode and the electrolyte. Fig. 7(i) shows Nyquist plots of the OA, CA, and AA samples with the consequent equal circuit model. The overall impedance contribution was simplified to transform the EIS spectrum into

Table 5 Capacitive parameters from the GCD plot in a two-electrode configuration

Potential window (V)	Current (A)	Discharge time (s)	Specific capacitance (F g^{-1})	Energy density (W h kg^{-1})	Power density (W kg^{-1})
1.5	1	128	85	27	759
	2	50	67	21	1512
	3	21	42	13	2229
	4	8	21	7	3150
	5	5	17	5	3600
	6	3	12	4	4800
	20	0.6	8	2.5	15 000



Table 6 Comparison of the electrochemical performances between the assembled Cu₂MnSnS₄//AC ASC device with other literature values

Positive material//negative material	Cell configuration	Electrolyte	Specific capacitance (F g ⁻¹) @ GCD	Energy density (W h kg ⁻¹)	Power density (W kg ⁻¹)	References
CuS-nanoplatelets	Thin film (three-electrode)	1 M LiClO ₄	55.55 @ 3 A g ⁻¹	6.23	1750	53
CuS//AC	Two-electrode (asymmetric)	2 M KOH	48.2 @ 0.5 A g ⁻¹	15.06	392.9	54
CuS-OEs/rGO//AC	Two-electrode (asymmetric)	3 M KOH	37.2 @ 10 A g ⁻¹	16.7	681	55
NiS//AC	Two-electrode (asymmetric)	2 M KOH	119 @ 1 A g ⁻¹	16.5	250	56
Cu ₂ S//rGO	Two-electrode (asymmetric)	6 M KOH	93.8 @ 1 A g ⁻¹	18.6	681.2	57
NiCo ₂ S ₄ //AC	Two-electrode (asymmetric)	2 M KOH	—	24.7	428	58
Cu ₄ SnS ₄ //AC	Two-electrode (solid state)	1 M PVA + NaOH	34.9	2.4	291	59
m-CuO120/FSS//h-CuS100/FSS	Two-electrode (solid state)	2 M PVA + KOH	123	22.8	2.5	60
Cu ₂ MnSnS ₄ //AC	Two-electrode (asymmetric)	2 M KOH	85	27	759	This work

its corresponding equivalent circuit and the physical process involved in the samples was determined using equivalent circuit parameters. The following fitted equivalent circuit ($R_1 + Q_2/R_2 + Q_3$) was used to assess the different components of the cell resistance, where R_1 corresponds to the ohmic drop in the electrolyte, R_2 refers to the charge ion displacement through the electrode mass interface, and Q_2 and Q_3 represent the constant phase components.⁵¹ All the electrical components values are summarized in Table 4 and clearly show that the value of resistance decreases while the capacitance increases gradually, which significantly improve the bias potential. This behavior enhances the electron-transfer dynamics and causes significant changes, leading to improvements in the charge-transfer process. Also, from Table 4, the OA sample exhibited lower R_{ct} values than the other two, which further supported the CV and GCD results.

3.1 Two-electrode fabrication of an ASC device

The electrochemical performance of Cu₂MnSnS₄//AC was evaluated under a two-electrode configuration *via* a specifically assembled asymmetric supercapacitor (ASC), with Cu₂MnSnS₄ as the working positive electrode and activated carbon (AC) as the negative electrode in 2 M KOH as the electrolyte. The operation voltage of the full cell was set to 0–1.5 V for all the studies. The loading mass of an electrode was estimated employing the formula: $m^+/m^- = c^- \times 1V^-/c^+ \times 1V^+$. Fig. 8(a) displays a comparative graph of the positive and negative electrodes. In order to analyze a proper voltage window for the ASC device, the potential windows were varied in the CV study, as shown in Fig. 8(b), which revealed both the battery-type and electric double layer effects.⁵² Very quick charge-transfer performance was exhibited by Cu₂MnSnS₄//AC ASC when the current density was increased from 10 to 60 mV s⁻¹ and there was no distortion of the CV shape found. Fig. 8(c) presents GCD curves of Cu₂MnSnS₄//AC ASC, from which the electrochemical performance, such as energy and power density, was calculated (Table 5). For the highest current density of 20 A g⁻¹, 2.5 W h kg⁻¹ and 15 000 W kg⁻¹ energy and power density were obtained, respectively. Here in order to remove the stability barriers, we used a switch lock-type device to test the stability behavior. Initially the stability decreased due to the fast reaction kinetics of the electrode/electrolyte interactions. Then after

a few cycles more (1000 cycles), constant stability was reached, which may be due to the stabilized reaction kinetics. The best cyclic stability test was carried out for 21 000 cycles and 80.35% capacity as well as 97.54% coulombic efficiency were retained, thus indicating the quality of the chosen electrode, as shown in Fig. 8(d). Fig. 8(e) displays a Ragone plot of the energy *vs.* the power density with other comparative literature values, and these are summarized in Table 6. Further, for the practical demonstration of real-time application, two Cu₂MnSnS₄//AC ASCs were cascaded and were able to light up a blue LED and run a small-sized motor (Fig. 8(f)). Hence, Cu₂MnSnS₄//AC ASC devices may be suitable for electrochemical storage applications.

In this study, a Cu₂MnSnS₄ electrode with a high working potential for aqueous asymmetric supercapacitors was proposed. The structure directing agent's role in fabrication of the Cu₂MnSnS₄ nanostructure was investigated. Here, OA electrodes showed the highest capacity and best cyclic stability. Moreover, the fabricated device exhibited a high energy density and power density.

4. Conclusions

In summary, Cu₂MnSnS₄ electrode materials fabricated with the use of varying structure directing agents (AA, CA, OA) in a three-electrode configuration were successfully developed. Two electrode devices and their electrochemical measurements were tested to determine the best electrode. The calculated specific capacitance values for electrodes fabricates with the use of OA, CA, and AA were 649, 453, and 442 F g⁻¹ at 0.5 A g⁻¹. The electrodes demonstrated 83%, 87%, and 89% capacity retention and cycling stability after 5000 cycles for the AA, CA, and OA electrodes. An asymmetric device was designed and exhibited 27 W h kg⁻¹ energy and 759 W kg⁻¹ power density at 1 A g⁻¹. Moreover, the device provided 80.35% capacity retention and 97.54% Coulombic efficiency after 21 000 cycles. Thus excellent electrochemical effects were obtained for the Cu₂MnSnS₄-OA//AC electrode.

Conflicts of interest

We hereby declare that there are no conflicts of interest.



Acknowledgements

This work was supported by RUSA, UGC-SAP, DST-FIST, DST-PURSE grants. The authors extend their appreciation to the Deanship of Scientific Research at King Khalid University for funding this work through General Research Project under grant number GRP-180-41.

References

- Z. Liu, S. Zhang, L. Wang, T. Wei, Z. Qiu and Z. Fan, *Nano Select*, 2020, **1**, 244–262, DOI: 10.1002/nano.202000011.
- E. Hossain, H. M. R. Faruque, M. Sunny, S. Haque, N. Mohammad and N. Nawar, *Energies*, 2020, **13**, 3651, DOI: 10.3390/en13143651.
- S. Kumar, G. Saeed, L. Zhu, K. N. Hui, N. H. Kim and J. H. Lee, *Chem. Eng. Trans.*, 2020, **403**, 126352, DOI: 10.1016/j.cej.2020.126352.
- Z. Yang, J. Tian, Z. Ye, Y. Jin, C. Cui, Q. Xie, J. Wang, G. Zhang, Z. Dong, Y. Miao and X. Yu, *Energy Storage Materials*, 2020, **33**, 18–25, DOI: 10.1016/j.ensm.2020.07.020.
- W. Liu, M. Li, G. Jiang, G. Li, J. Zhu, M. Xiao, Y. Zhu, R. Gao, A. Yu, M. Feng and Z. Chen, *Adv. Energy Mater.*, 2020, **10**, 2001275, DOI: 10.1002/aenm.202001275.
- S. Biswas, V. Sharma, D. Mandal, A. Chowdhury, M. Chakravarty, S. Priya, C. C. Gowda, P. De, I. Singh and A. Chandra, *CrystEngComm*, 2020, **22**, 1633–1644, DOI: 10.1039/C9CE01547G.
- T. Wang, F. Yu, X. Wang, S. Xi, K. J. Chen and H. Wang, *Electrochim. Acta*, 2020, **334**, 135586, DOI: 10.1016/j.electacta.2019.135586.
- A. M. Zardkhouhou and S. S. H. Davarani, *Chem. Eng. J.*, 2020, **402**, 126241, DOI: 10.1016/j.cej.2020.126241.
- C. Xiong, Q. Yang, W. Dang, M. Li, B. Li, J. Su, Y. Liu, W. Zhao, C. Duan, L. Dai and Y. Xu, *J. Power Sources*, 2020, **447**, 227387, DOI: 10.1016/j.jpowsour.2019.227387.
- Y. Wang, X. Wang, X. Li, Y. Bai, H. Xiao, Y. Liu and G. Yuan, *Chem. Eng. J.*, 2021, **405**, 126664, DOI: 10.1016/j.cej.2020.126664.
- K. Robert, D. Stievenard, D. Deresmes, C. Douard, A. Iadecola, D. Troadec, P. Simon, N. Nuns, M. Marinova, M. Huvé and P. Roussel, *Energy Environ. Sci.*, 2020, **13**, 949–957, DOI: 10.1039/C9EE03787J.
- M. Isacfranklin, R. Yuvakkumar, G. Ravi, S. I. Hong, D. Velauthapillai, M. Thambidurai, C. Dang, T. S. Algarni and A. M. Al-Mohaimed, *Carbon*, 2020, **172**, 613–623, DOI: 10.1016/j.carbon.2020.10.081.
- K. Thiagarajan, T. Bavani, P. Arunachalam, S. J. Lee, J. Theerthagiri, J. Madhavan, B. G. Pollet and M. Y. Choi, *J. Nanomater.*, 2020, **10**, 392, DOI: 10.3390/nano10020392.
- R. Wang and Y. Li, *Energy Storage Materials*, 2020, **31**, 230–251, DOI: 10.1016/j.ensm.2020.06.012.
- T. Okamura and K. Nakagawa, *ECSarXiv*, 2020, DOI: 10.1149/osf.io/9gjym.
- Y. Liu, X. Xu and Z. Shao, *Energy Storage Materials*, 2020, **26**, 1–22, DOI: 10.1016/j.ensm.2019.12.019.
- K. N. Hui, K. San Hui, Z. Tang, V. V. Jadhav and Q. X. Xia, *J. Power Sources*, 2016, **330**, 195–203, DOI: 10.1016/j.jpowsour.2016.08.116.
- S. Liu, D. Ni, H. F. Li, K. N. Hui, C. Y. Ouyang and S. C. Jun, *J. Mater. Chem. A*, 2018, **6**, 10674–10685, DOI: 10.1039/C8TA00540K.
- Q. Zhang, W. B. Zhang, P. Hei, Z. Hou, T. Yang and J. Long, *Appl. Surf. Sci.*, 2020, **527**, 146682, DOI: 10.1016/j.apsusc.2020.146682.
- R. Pothu, R. Bolagam, Q. H. Wang, W. Ni, J. F. Cai, X. X. Peng, Y. Z. Feng and J. M. Ma, *Rare Met.*, 2020, **258**, 1–21, DOI: 10.1007/s12598-020-01470-w.
- H. Hua, S. Liu, Z. Chen, R. Bao, Y. Shi, L. Hou, G. Pang, K. N. Hui, X. Zhang and C. Yuan, *Sci. Rep.*, 2016, **6**, 20973, DOI: 10.1038/srep20973.
- X. Wang, R. Cao, S. Zhang, P. Hou, R. Han, M. Shao and X. Xu, *J. Mater. Chem. A*, 2017, **5**, 23999–24010, DOI: 10.1039/c7ta06809c.
- J. S. Wei, T. B. Song, P. Zhang, Z. Y. Zhu, X. Y. Dong, X. Q. Niu and H. M. Xiong, *ACS Appl. Energy Mater.*, 2020, **3**, 6907–6914, DOI: 10.1021/acsaem.0c00990.
- R. S. Gohar, I. Ahmad, A. Shah, S. Majeed, M. Najam-Ul-Haq and M. N. Ashiq, *J. Electrochem. Energy Convers. Storage*, 2020, **31**, 101621, DOI: 10.1016/j.est.2020.101621.
- P. Wonner, T. Steinke, L. Vogel and S. M. Huber, *Chem.–Eur. J.*, 2020, **26**, 1258–1262, DOI: 10.1002/chem.201905057.
- R. Henríquez, C. Vásquez, E. Muñoz, P. Grez, F. Martín, J. R. Ramos-Barrado and E. A. Dalchiele, *Phys. E Low-dimens. Syst. Nanostruct.*, 2020, **118**, 113881, DOI: 10.1016/j.physe.2019.113881.
- J. H. M. van Gils, E. van Dijk, A. Peduzzo, A. Hofmann, N. Vettore and M. P. Schützmann, *PLoS Comput. Biol.*, 2020, **16**(5), e1007767, DOI: 10.1371/journal.pcbi.1007767.
- T. T. Trinh, K. Q. Tran, X. Q. Zhang, R. A. van Santen and E. J. Meijer, *Phys. Chem. Chem. Phys.*, 2015, **17**, 21810–21818, DOI: 10.1039/c5cp02068a.
- J. Qin, Q. Li, Y. Liu, A. Niu and C. Lin, *Ecotoxicol. Environ. Saf.*, 2020, **201**, 110873, DOI: 10.1016/j.ecoenv.2020.110873.
- S. Asaithambi, P. Sakthivel, M. Karuppaiah, G. U. Sankar, K. Balamurugan, R. Yuvakkumar, M. Thambidurai and G. Ravi, *J. Electrochem. Energy Convers. Storage*, 2020, **31**, 101530, DOI: 10.1016/j.est.2020.101530.
- M. Isacfranklin, G. Ravi, R. Yuvakkumar, P. Kumar, D. Velauthapillai, B. Saravanakumar, M. Thambidurai and C. Dang, *Ceram. Int.*, 2020, **46**, 16291–16297, DOI: 10.1016/j.ceramint.2020.03.186.
- L. Chen, H. Deng, J. Tao, H. Cao, L. Huang, L. Sun, P. Yang and J. Chu, *RSC Adv*, 2015, **5**, 84295–84302, DOI: 10.1039/c5ra14595c.
- C. Sripan, D. Alagarasan, S. Varadharajaperumal, R. Ganesan and R. Naik, *Curr. Appl. Phys.*, 2020, **20**, 925–930, DOI: 10.1016/j.cap.2020.05.003.
- M. Devika, N. Koteeswara Reddy, M. Prashantha, K. Ramesh, S. Venkatramana Reddy, Y. B. Hahn and K. R. Gunasekhar, *Phys. Status Solidi*, 2010, **207**, 1864–1869, DOI: 10.1002/pssa.200925379.



- 35 S. Varadharajaperumal, C. Sripan, R. Ganesan, G. Hegde and M. N. Satyanarayana, *Cryst. Growth Des.*, 2017, **17**, 5154–5162, DOI: 10.1021/acs.cgd.7b00632.
- 36 L. Chen, H. Deng, J. Tao, W. Zhou, L. Sun, F. Yue, P. Yang and J. Chu, *J. Alloys Compd.*, 2015, **640**, 23–28, DOI: 10.1016/j.jallcom.2015.03.225.
- 37 S. Riyaz, A. Parveen and A. Azam, *Perspectives in Science*, 2016, **8**, 632–635, DOI: 10.1016/j.pisc.2016.06.041.
- 38 Y. Liu, X. Mi, J. Wang, M. Li, D. Fan, H. Lu and X. Chen, *Inorg. Chem. Front.*, 2019, **6**, 948–954, DOI: 10.1039/C9qi00020h.
- 39 Y. Zuo, J. Li, X. Yu, R. Du, T. Zhang, X. Wang, J. Arbiol, J. Llorca and A. Cabot, *Chem. Mater.*, 2020, **32**, 2097–2106, DOI: 10.1021/acs.chemmater.9b05241.
- 40 D. Sharma, A. V. Menon and S. Bose, *Nanoscale Adv*, 2020, **2**, 3292–3303, DOI: .
- 41 X. Liang, P. Guo, G. Wang, R. Deng, D. Pan and X. Wei, *RSC Adv*, 2012, **2**, 5044–5046, DOI: 10.1039/c2ra20198d.
- 42 Y. Xie, C. Zhang, G. Yang, J. Yang, X. Zhou and J. Ma, *J. Alloys Compd.*, 2017, **696**, 938–946, DOI: 10.1016/j.jallcom.2016.12.043.
- 43 Y. A. Kumar, K. D. Kumar and H. J. Kim, *Electrochim. Acta*, 2020, **330**, 135261, DOI: 10.1016/j.electacta.2019.135261.
- 44 Y. Wang, Y. Song and Y. Xia, *Chem. Soc. Rev.*, 2016, **45**, 5925–5950, DOI: 10.1039/c5cs00580a.
- 45 T. S. Mathis, N. Kurra, X. Wang, D. Pinto, P. Simon and Y. Gogotsi, *Adv. Energy Mater.*, 2019, **9**, 1902007, DOI: 10.1002/aenm.201902007.
- 46 P. Himasree, I. K. Durga, T. N. V. Krishna, S. S. Rao, C. V. M. Gopi, S. Revathi, K. Prabakar and H. J. Kim, *Electrochim. Acta*, 2019, **305**, 467–473, DOI: 10.1016/j.electacta.2019.03.041.
- 47 J. Theerthagiri, K. Karuppasamy, G. Durai, A. U. H. S. Rana, P. Arunachalam, K. Sangeetha, P. Kuppasami and H. S. Kim, *J. Nanomater.*, 2018, **8**, 256, DOI: 10.3390/nano8040256.
- 48 D. Guragain, C. Zequine, R. K. Gupta and S. R. Mishra, *Processes*, 2020, **8**, 343, DOI: 10.3390/pr8030343.
- 49 S. Kaipannan, K. Govindarajan, S. Sundaramoorthy and S. Marappan, *ACS Omega*, 2019, **4**, 15798–15805.
- 50 W. Chen, X. Yu, Z. Zhao, S. Ji and L. Feng, *Electrochim. Acta*, 2019, **298**, 313–320, DOI: 10.1016/j.electacta.2018.12.096.
- 51 G. M. Kumar, F. Xiao, P. Ilanchezhian, S. Yuldashev and T. W. Kang, *RSC Adv*, 2016, **6**, 99631–99637.
- 52 X. He, X. Mao, C. Zhang, W. Yang, Y. Zhou, Y. Yang and J. Xu, *J. Mater. Sci.: Mater. Electron.*, 2020, **31**, 2145–2152, DOI: 10.1007/s10854-019-02737-4.
- 53 C. J. Raj, B. C. Kim, W. J. Cho, W. G. Lee, Y. Seo and K. H. Yu, *J. Alloys Compd.*, 2014, **586**, 191–196, DOI: 10.1016/j.jallcom.2013.10.056.
- 54 J. Zhang, H. Feng, J. Yang, Q. Qin, H. Fan, C. Wei and W. Zheng, *ACS Appl. Mater. Interfaces*, 2015, **7**, 21735–21744, DOI: 10.1021/acsami.5b04452.
- 55 Y. Cui, J. Zhang, G. Li, Y. Sun, G. Zhang and W. Zheng, *Chem. Eng. J.*, 2017, **325**, 424–432, DOI: 10.1016/j.cej.2017.05.069.
- 56 S. Nandhini and G. Muralidharan, *Appl. Surf. Sci.*, 2018, **449**, 485–491, DOI: 10.1016/j.apsusc.2018.01.024.
- 57 T. Zhao, X. Peng, X. Zhao, J. Hu, W. Yang, T. Li and I. Ahmad, *Compos. B Eng.*, 2019, **163**, 26–35, DOI: 10.1016/j.compositesb.2018.11.023.
- 58 Y. Zhu, X. Ji, Z. Wu and Y. Liu, *Electrochim. Acta*, 2015, **186**, 562–571, DOI: 10.1016/j.electacta.2015.10.176.
- 59 A. C. Lokhande, A. Patil, A. Shelke, P. T. Babar, M. G. Gang, V. C. Lokhande, D. S. Dhawale, C. D. Lokhande and J. H. Kim, *Electrochim. Acta*, 2018, **284**, 80–88, DOI: 10.1016/j.electacta.2018.07.170.
- 60 A. M. Patil, V. C. Lokhande, T. Ji and C. D. Lokhande, *Electrochim. Acta*, 2019, **307**, 30–42, DOI: 10.1016/j.electacta.2019.03.108.

