Quaternary Cu$_2$FeSnS$_4$/PVP/rGO Composite for Supercapacitor Applications

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ABSTRACT: Electrochemical energy storage is a current research area to address energy challenges of the modern world. The Cu$_2$FeSnS$_4$/PVP/rGO-decorated nanocomposite using PVP as the surface ligand was explored in a simple one-step solvothermal route, for studying their electrochemical behavior by designing asymmetric hybrid supercapacitor devices. The full cell three-electrode arrangements delivered 748 C/g (62.36 mA h/g) at 5 mV/s employing CV and 328 F/g (45.55 mA h/g) at 0.5 A/g employing GCD for the Cu$_2$FeSnS$_4$/PVP/rGO electrode. The half-cell two-electrode device can endow with 73 W h/kg and 749 W/kg at 1 A/g energy and power density. Furthermore, two Cu$_2$FeSnS$_4$/PVP/rGO//AC asymmetric devices connected in series for illuminating a commercial red LED more than 1 min were explored. This work focuses the potential use of transition-metal chalcogenide composite and introduces a new material for designing high-performance supercapacitor applications.

1. INTRODUCTION

In the developing modern era, the world is facing great hardships as fuel prices continue to rise, air pollution continues to rise, and fossil fuels become inadequate. In such a scenario, the so-called unconventional energies should be used to meet energy requirements. The one of the drawback is that there is no sustainable storage source to store such energies. Supercapacitors and batteries are a potential source of energies for practical use, while principle energy sources can be exhausted. Recently, supercapacitors have attracted a lot of interest due to their physicochemical characteristics because of double-layer capacitor (EDLC) or pseudocapacitor charge-storage mechanisms. Pseudocapacitors are having high storage and on the other hand having low cyclic rate, whereas EDLC materials as carbon-based electrode materials exhibit excellent cyclability. Nanomaterials are widely used to provide alternative energy sources to eco-friendly fossil fuels. For example, various alloys, metal nitrides, metal oxides, and sulfides have been used for supercapacitor use. Generally, NiO, SnO, MnO$_2$, RuO$_2$, V$_2$O$_5$, and Co$_3$O$_4$ and their composites are used as electrode materials. So far, RuO$_2$ has been reported as the best electrode material that can provide good performances due to its excellent theoretical specific capacity (1400–2000 F/g). Nevertheless, the high cost of production and the effects of agglomeration stand as high barriers for marketable use. Nowadays, the binary, ternary, and quaternary transition-metal composites are extensively investigated to improve the specific capacity and the performance of the supercapacitor. In recent days, research studies have been carried out on quaternary chalcogenides for its excellent redox activity, pseudocapacitance, and effective layering ion diffusion. It is a good idea to try and use the Cu$_2$FeSnS$_4$ material with the earth’s numerous replacement for the supercapacitor application which are tetrahedral integrated systems, wherein each sulfur anion is bound with 4 cations and vice versa. Numerous protocols are being accounted to synthesize Cu$_2$-II−IV−VI$_4$ group semiconductor materials in literature studies. According to the literature review, no similar efforts have been carried out for using such a composition in supercapacitor applications. Nanostructures greatly enhanced supercapacitor applications because of the suitable addition of carbon-based materials. Thus, improving ionic transport at the electrode/electrolyte interface accelerates redox reactions that occur in energy-harvesting devices. The earth-abundant quaternary chalcogenide materials are rarely reported in the supercapacitor applications. Furthermore, in stability point of view, the sulfide materials easily decayed in the aqueous electrolytes. Accordingly, retaining the stability is a curial factor in supercapacitor applications. The novel
combination of Cu$_2$FeSnS$_4$/PVP/rGO focuses the long-term stability in the two-electrode system up to 2000 cycles. Also, this work highlighted an alternative electrode with a novel composite to enhance the electrochemical performances. This work definitely grasps the attention of all researchers working in the areas of energy-related applications.

2. MATERIALS AND METHODS

Preparation of GO and rGO was followed by the modified Hummer’s method. First, 0.6 g of graphite flakes was blended with 23 mL of H$_2$SO$_4$ and stirred well for an hour. Then, in an ice bath, the mixture was stirred for 30 min and 3 g of potassium permanganate was added. After that, 46 mL of double-distilled water was diluted with the resulting solution. Then, temperature was increased rapidly to 100 °C and the appearance of the color was changed to brown. Finally, the yellow color appeared after the addition of 10 mL of hydrogen peroxide (H$_2$O$_2$). The sediment was desiccated at 100 °C for 12 h to obtain the GO product, and then, 100 mg of GO was mixed with 100 mL of water. After that, 500 mg of NaBH$_4$ was appended and stirred at room temperature for 12 h. The reduced GO was gradually obtained. In the synthesis process of Cu$_2$FeSnS$_4$, Cu$_2$FeSnS$_4$/PVP, and Cu$_2$FeSnS$_4$/PVP/rGO, initially, 0.1 M copper (II) chloride dehydrate, 0.05 M ferric chloride anhydrous, and 0.05 M tin (IV) chloride pentahydrate were suspended in 80 mL of water and stirred continuously for 30 min and found to be greenish brown. Then, a 0.4 M thiourea solution was stirred for another 1 h. The abovementioned mixture was then poured into a 100 mL autoclave at 160 °C for 12 h. Finally, the Cu$_2$FeSnS$_4$ product was obtained. Subsequently, a similar procedure was followed for the addition of polyvinyl pyrrolidone (PVP) as a structure-directing reagent in the chemical process to regulate crystal growth and nucleation, reducing surface tension and stabilizing metal nanoparticles. Third, the rGO was dispersed into the Cu$_2$FeSnS$_4$/PVP compound to form another compound called Cu$_2$FeSnS$_4$/PVP/rGO. Subsequently, the obtained Cu$_2$FeSnS$_4$, Cu$_2$FeSnS$_4$/PVP, and Cu$_2$FeSnS$_4$/PVP/rGO powders were given for characterization, and electrochemical performance was carried out for all samples. The Cu$_2$FeSnS$_4$ compound was formed by dissolving water in metal chlorides and precursors of thiourea, and necessary chemical reactions during the synthesis of materials are described.

\[
\begin{align*}
\text{CuCl}_2 & \rightarrow \text{Cu}^{2+} + 2\text{Cl}^-; \\
\text{SnCl}_4 & \rightarrow \text{Sn}^{4+} + 4\text{Cl}^-; \\
\text{FeCl}_2 & \rightarrow \text{Fe}^{2+} + 2\text{Cl}^-; \\
\text{CS(NH}_2\text{)}_2 & \rightarrow S^{2-} + \text{H}_2 + \text{NH}_2 - \text{CaNH} \\
\text{CuCl}_2 + \text{FeCl}_2 + \text{CS(NH}_2\text{)}_2 & \rightarrow [\text{CuFeSn(CS(NH}_2\text{)}_2^m]^{n+} \\
\text{The chemical composition of Cu/Fe/Sn/S is} & \text{2:1:1:4 M.} \\
2\text{Cu}^{2+} + \text{Sn}^{2+} & \rightarrow 2\text{Cu}^+ + \text{Sn}^{4+} \\
\text{Cu}^+ + \text{en} & \rightarrow [\text{Cu(en)}]^+; \\
\text{Fe}^{2+} + \text{en} & \rightarrow [\text{Fe(en)}]^2+; \\
\text{Sn}^{4+} + \text{en} & \rightarrow [\text{Sn(en)}]^4+; \\
(\text{NH}_2\text{)}_2\text{CS} + 2\text{H}_2\text{O} & \rightarrow S^{2-} + 2\text{NH}_4^+ + \text{CO}_2^+ \\
\end{align*}
\]
2[Cu(en)]^+ + [Fe(en)]^2+ + [Sn(en)]^4+ + 4S^{2-} → Cu_2FeSnS_4

For electrochemical measurements and electrode preparations, Cu_2FeSnS_4 (80%) was mixed with activated charcoal (10%), acetylene black (5%), polyvinylidene fluoride (PVDF) (5%), and solution of binder in N-methyl-2-pyrrolidone (NMP) on nickel foam. Filter paper soaked in 2 M potassium hydroxide was used as a separator. An asymmetric cell was constructed with 2 M KOH with Cu_2FeSnS_4 and AC as positive and negative electrodes which were separated by KOH-soaked filter paper configuration to form a supercapacitor. The two-electrode cell system was configured as follows: Cu_2FeSnS_4/PVP/rGO//AC using biological SP-150 instruments. The following mathematical relationship explored to calculate

specific capacity from CV: \( C_v = \int \frac{I \times dv}{2 \times s \times \Delta V \times m} \) C/g

specific capacitance from GCD: \( C_g = \frac{I \times \Delta t}{m \times \Delta V} \) F/g

It is necessary to specify capacitance in F/g to the specific capacity in C/g or mA h/g to show battery-like behavior.

specific capacity from CV: C
\[
C = \frac{\int I \times dv}{2 \times 3.6 \times \Delta V \times m} \text{(mA h/g)}
\]

specific capacity from GCD: \( C_g = \frac{I \times \Delta t}{3.6 \times m} \) (mA h/g)

To fabricate the hybrid ASC supercapacitor, Cu_2FeSnS_4/rGO as a cathode and AC as an anode were used to explore Cu_2FeSnS_4/rGO/AC. The cathode electrode was designed by blending 80:10:5:5 wt% of active material, activated carbon, acetylene black, and polyvinylidene fluoride (PVDF), respectively. The anode electrode was the combination of activated carbon (80 wt%), acetylene black (10 wt%), and polyvinylidene fluoride (PVDF) (10 wt%) using N-methyl pyrrolidone as a solvent. All the material mingled well to make slurry. Thereafter, the mixture was coated uniformly on Ni foam (1 \times 1 \text{ cm}^2) and vacuum air-dried at 80 °C. The charge balancing was optimized by proper mass proportion of the cathode and anode; \( m_c/m_a = Q_c/Q_a \times V_a/V_c \). To calculate the specific capacity, energy, and power density, one can employ the following formula.\(^{21,22}\)

\[
E = \frac{C \times \Delta V^2}{7.2} \text{(W h/g)}
\]

\[
P = \frac{3600 \times E}{\Delta t} \text{(W/kg)}
\]

3. RESULTS AND DISCUSSION

The XRD diffraction pattern of the synthesized pure Cu_2FeSnS_4 and the addition of PVP and rGO are explored in Figure 1a (JCPDS 44-1476). The formation of (112), (200), (220), (204), (312), and (224) planes in the corresponding diffraction peaks in all the samples suggested the formation of the well-crystallized Cu_2FeSnS_4 orthogonal structure. No other secondary diffraction peaks were observed. The Scherrer formula \( D = 0.89 \lambda / \beta \cos \theta \) was used and the crystallite size of 27 nm was calculated for the high intensity diffraction pattern of the (112) plane. The dislocation density \( \delta = 1.37 \times 10^{15} \text{ lines/meter}^2 \) and microstrain \( \epsilon = 1.28 \times 10^{-3} \) were calculated for the high intensity 2θ = 28.40° peak using the respective formulas \( \delta = 1/D^2 \) and \( \epsilon = \beta \cos \theta/4. \)

Figure 1b reveals the crystalline nature of the Cu_2FeSnS_4 using the Vesta software. Figure 1c reveals the photoluminescence spectra of pure Cu_2FeSnS_4 and the addition of PVP and rGO samples under an excitation wavelength of 270 nm. The PL spectra displayed strong peaks centered at 545 and 368 nm and the weak peaks centered around 485 and 298 nm; 368 nm may be because of band-to-band transition. The peak assigned to 545 nm in the visible region is the reason for the surface defects during the growth process. Also, the emission occurred in the visible region may be the cause of oxygen vacancies and the recombination of electron trapping in the crystal interstitials. The green emission at 485 nm is because of surface vacancies which may be because of electron transition. The absorption peak at 298 nm is attributed to the quantum effect related to the copper sulfide.\(^{24,25}\) The infrared spectrum explored the ion state in a crystal due to crystal vibrations, as displayed in Figure 1d. Vibrations of the infrared absorption group in solids typically occur at 100–1000; 668 cm\(^{-1}\) indicates stretching vibration of the metal–thiourea complex-C=S of the synthesized product. The band located at 1020 cm\(^{-1}\) also denotes stretching vibration of the C=S band. The 1638 cm\(^{-1}\) band is responsible for the metal–thiourea complex-N=C–N stretching and NH\(_2\) bending vibrational mode. The carbon dioxide peak is revealed at 2368 cm\(^{-1}\); the peak at 3467 cm\(^{-1}\) is attributed to O–H stretching vibration of H\(_2\)O.\(^{26}\)

The identification of the various compositional phases of the synthesized material has been revealed through the Raman studies (Figure 2). The characteristic Raman vibrational modes are observed at 144, 214, 476, and 2116 cm\(^{-1}\). In the Cu_2FeSnS_4/PVP/rGO product, the formation of the D and G band and 2D bands is located at 1353, 1597, and 2400–2500 cm\(^{-1}\). The peak at 476 cm\(^{-1}\) revealed the sulphur S=S mode. The peak assigned to 214 cm\(^{-1}\) is corresponding to the pure sulfur anion in the region of the copper metal.\(^{27}\) The disorder-induced D band and first-order graphite G band \( (I_D/I_G) \) ratio is used to measure disorder. Intensity is proportional to disorder. In our case, the \( I_D/I_G = 0.99 \) explored nanocrystalline graphitic nature. The peak at 1353 cm\(^{-1}\) attributed to the D
band refers to defective sites with vacancies and grain boundaries and the G band at 1597 cm$^{-1}$ responsible for the $E_{2g}$ phonon first-order scattering of the sp$^2$ carbon–carbon bond.\textsuperscript{28}

Figure 3a–c reveals SEM micrographs of synthesized Cu$_2$FeSnS$_4$ nanostructures among the addition of PVP and rGO. Almost all the morphological images exhibit agglomerated nanoparticles. It is clearly evidenced with the average 200 nm size. The surface ligand PVP had a major impact on the nucleation and growth.\textsuperscript{29} Figure 3d shows the selected area of the EDS spectrum which clearly evidenced the formation of sphere-like morphological structures. The EDAX mapping with the atomic and weight percentages of Cu$_2$FeSnS$_4$/PVP/rGO is shown in Figure 3e. Figure 3f–h reveals TEM analysis of the Cu$_2$FeSnS$_4$/PVP/rGO sample. The clear-cut spherical ball-structured morphology along with the nanosheet is due to rGO content confirmed through the TEM analysis with the SAED pattern of a series of diffraction rings. The crystal plane of (112) was marked in the SAED pattern, and the gap between the fringes is around 0.38 nm.\textsuperscript{30} The carbon-based rGO products were combined with these quarterly products formed between surface and graphene layers. rGO dispersed with Cu$_2$FeSnS$_4$/PVP facilitates electrolyte diffusion through their electrodes and reduces product internal resistance; as a result, the conductivity has been greatly improved.

The surface composition of synthesized Cu$_2$FeSnS$_4$/PVP/rGO nanostructures has been explored. Figure 4a reveals the survey spectrum of the prepared product. Peaks located at 931.30 and 951.16 eV are attributed to the Cu$_2$P$_{3/2}$ and Cu$_2$P$_{1/2}$ species. The peak splitting of 19.31 eV denotes the Cu(I) configuration (Figure 4b). The most popular peaks at 709.60 eV (Fe$_2$p$_{3/2}$) and 724.42 eV (Fe$_2$p$_{1/2}$) signify the Fe(II) configuration, and the peak centered at 715.60 is responsible for Sn$_2$p$_{3/2}$ (Figure 4c). 485.60 and 494 eV correspond to Sn$_3$d$_{5/2}$ and Sn$_3$d$_{3/2}$ as displayed in Figure 4d, which specifies the Sn(IV) configuration. The peak expected at 160–164 eV is corresponding to the sulfide phases. The single peak in Figure 4e at 161.32 eV is the reason for the Sn(−II) oxidation state. Finally, the carbon peaks are positioned at 283.66 eV for C$_1$s which could be the reason for the presence of rGO in the synthesized material, as displayed in Figure 4f.
The major states of the Cu, Fe, Sn, and S elements are +1, +2, +4, and −2, for Cu₂FeSnS₄ chemical formula.³¹

Electrochemical processes were studied at 2 M KOH to test electrochemical behavior. Cyclic voltammetry (CV) is an electrochemical tool commonly used to reveal oxidation and reduction processes of molecular species, electrochemical reactions caused by the electron transfer process, and the performances of the supercapacitor. In the CV technique, current is recorded when the electric potential varies with the corresponding scan rates. Figure 5a–c reveals the CV graphs of the prepared Cu₂FeSnS₄ and its composite at different sweep currents (5–50 mV/s) in 0–0.6 V. Pseudocapacitors store charges by the active substance surface redox reaction and OH⁻ ions in the KOH solution. Charge is stored for the electric double layer at the electrode–electrolyte interface. As a result of the current responses, a pair of redox peaks clearly express battery-type properties, resulting in electrode materials showing reversible Faradaic reactions. Faradaic redox reactions of charge-storage mechanisms are as follows: Cu₂FeSnS₄ + 9OH⁻ → 2Cu(OH)₂ + M(SOH) + Sn(OH)₄ + 3S, where M signifies Fe. The CV profiles of Cu₂FeSnS₄ explored the active redox pairs of Cu⁺/Cu²⁺, Fe²⁺/Fe³⁺, and Sn⁴⁺/Sn⁵⁺ operating in a constant-potential window of OH anions in an alkaline electrolyte.³² The electrochemical redox reaction occurs only when the electrons are transmitted from a Ni foam substrate that is oxidized to one that is being reduced. Almost all CV curves revealed similar patterns of current responses (redox peaks), while increasing the scan rate except for systematic anodic and cathodic shifts in the positive and negative directions, respectively, because of the fast Faradaic reaction. Electrolyte ion diffusion is indicated by opposition of peak shift at low potential. These CV results should be consistent with the electrochemical reactions at a high scan rate. The behavior of these redox reactions ensures that the appropriate chosen electrode can provide better reversibility.³³ Hence, it shows the battery-type behavior, and the specific capacity was calculated for the synthesized product using eq 1. The calculated specific capacity values are 398 C/g (33.19 mA h/g), 602 C/g (50.13 mA h/g), and 748 C/g (62.36 mA h/g) at a 5 mV/s scan rate for Cu₂FeSnS₄, Cu₂FeSnS₄/PVP, and Cu₂FeSnS₄/PVP/rGO, respectively, and the remaining specific capacity values of all other scan rates are pictorially illustrated (Figure 5g).

The electrode materials are also determined by charge–discharge capabilities by varying different current densities (Figure 4d–f). The calculated values are 139 F/g (19.30 mA h/g), 259 F/g (34.72 mA h/g), and 328 F/g (45.55 mA h/g) at 0.5 A/g for Cu₂FeSnS₄, Cu₂FeSnS₄/PVP, and Cu₂FeSnS₄/PVP/rGO, respectively, and all other specific capacitance values are illustrated (Figure 5h). The capacitance decrease is proportional to current density increase as a result of scan rate and redox reaction in the CV. At high current densities, only outer active electrode surface is used for redox reactions which may be due to time limitations. Nevertheless, low current density is used for the entire inner and outer electrode surface to promote efficient redox activity. Due to the lack of sufficient electrolyte ions in the higher current densities, it does not have sufficient reactive kinetics to migrate and diffuse the electrolyte ions to the electrodes. Further IR losses and concentration polarization may be in addition to the decrease in capacitance of the high current density. Galvanostatic charge–discharge (GCD) curves are similar to curves at EDLC electrodes, with a slight different linear nature than the triangular nature.³⁴ The retained stability is a crucial factor in supercapacitor applications. The suitable incorporation of the carbon-based rGO materials solve this problem effectively along with PVP, forming the well-shaped structure as clearly evidenced in the selected area of EDAX and TEM images. Figure 5i indicates the cyclic stability graph of the best performed Cu₂FeSnS₄/
PVP/rGO electrode. The 86.40% of capacity was retained over the 5000 charge−discharge cycles.

EIS analysis was used to evaluate the charge-transfer characteristic of three electrodes (Figure 5j). Furthermore, the addition of rGO in the Cu2FeSnS4/PVP/rGO electrode enhances the low charge-transfer resistance which improved electronic conductivity of the prepared electrodes. Equivalent series resistance (ESR) consisting of electrolyte resistance, substrate/active material contact resistance, and internal resistance has been located at the joint in the real part at high frequency expanse in the Nyquist plot. Equivalent charge-transfer resistance ($R_{ct}$) and solution resistance ($R_s$) and other phase elements ($Q_3$) values are summarized in Table 1. The incorporation of rGO increased the adhesion and facilitates charge transport between rGO and Ni foam current collector. It has been fitted using Z fit analysis in biologic SP-150 instrument according to the equivalent circuit, as shown in Figure 5k. The lowest solution resistance shows that
the synthesized Cu$_2$FeSnS$_4$/PVP/rGO electrode provides the superior conductivity.

3.1. Trasatti Method. The Trasatti method is used to evaluate capacitive contribution from electrical double-layer and pseudocapacitive reactions. The detailed procedure and the method of calculations are clearly mentioned in our previous work.\textsuperscript{36} The capacitive and diffusion contributions in terms of the specific capacity in percentages of all the electrodes are shown in Figure 5g. Figure 5h displays the capacitive- and diffusion-controlled contributions calculated from CV scans at 5 mV/s for the best-performing Cu$_2$FeSnS$_4$/PVP/rGO electrode.\textsuperscript{37}

3.2. Two-Electrode Device Performances. The two-electrode hybrid device has been fabricated, and the ASC device was assembled with Cu$_2$FeSnS$_4$/PVP/rGO as the cathode and AC as the anode, with positive and negative electrodes marked as Cu$_2$FeSnS$_4$/PVP/rGO//AC ASC.\textsuperscript{38} In fact, the asymmetric and hybrid supercapacitors are the same;
nevertheless, they differ in the appropriate electrode configuration based on their mechanisms. In both cases, EDLC and pseudocapacitive or battery-type properties are explored. The role of the prepared electrode is to increase the energy density by increasing the working voltage without significantly reducing the power density. In our case, the CV curves exhibit the pair of oxidation and reduction which shows the battery-type behavior. The battery-type electrode material combined with the EDLC-based activated carbon material will be termed as the hybrid device, whereas in the case of asymmetric, two different capacitive mechanisms such as EDLC and pseudocapitance are combined. The electrolyte is the crucial factor to determine the potential window. In general, the aqueous electrolytes have a maximum potential window up to 1 V. The polymer electrolyte may have the potential up to 4 V. In addition to that, the three-electrode and two-electrode system measurement varies in different ways such as $E_{\text{work}} - E_{\text{ref}}$ for the three-electrode system and $E_{\text{work}} - E_{\text{aux}}$ for the two-electrode system. Hence, for both two-electrode and three-electrode system, working voltage was dependent on the electrode configuration. In our case, we have fixed the potential 1.5 V to evaluate the device performances by varying the scan rate and current density in CV and GCD, respectively. To make an asymmetric device, we used Cu₂FeSnS₄ material as the active cathode and activated carbon as an anode. In between, filter paper was used as a separator. When we optimize device working voltage, we need to take the sum of the voltage differences. The difference of the negative electrode is 1 V and the positive electrode is 0.5 V in GCD. Hence, the sum of the 1.5 V is estimated to evaluate the device performances. We also tested with higher potentials. Nevertheless, the CV graph revealed the oxidation hump in the higher voltages greater than 1.5 V. Hence, we optimize the 1.5 V to evaluate the device performances. Figure 6a explores CV of the positive electrode (0–0.6 V) and negative electrode (−1.0–0 V) at 10 mV/s. Figure 6b displays CV of the ASC hybrid device with different scan rates over a wide potential window up to 1.5 V. Figure 6c shows the GCD profile of the ASC hybrid device with various current densities between 1 and 20 A/g. The fabricated ASC hybrid device revealed the 73 W h/kg and 749 W/kg energy and power density, respectively. Table 2 explores electrochemical device performances of the

| Potential window (V) | Current (A) | Specific capacitance (F g⁻¹) | Discharge time (s) | Energy density (W h kg⁻¹) | Power density (W kg⁻¹) |
|----------------------|------------|-------------------------------|-------------------|--------------------------|-----------------------|
| 1.5                  | 1          | 234                           | 351               | 73                       | 749                   |
| 2                    | 195        | 146                           | 61                | 1504                     |
| 3                    | 136        | 54                            | 43                | 2276                     |
| 5                    | 113        | 34                            | 35                | 3705                     |
| 10                   | 106        | 16                            | 33                | 7425                     |
| 15                   | 90         | 9                             | 28                | 11,200                   |
| 20                   | 66         | 5                             | 21                | 15,120                   |

Figure 7. (a) Comparative graph, (b) CV, (c) GCD, and (d) stability graph of the hybrid device.
The capacitive retention and Coulombic efficiency of the fabricated hybrid device are shown in Figure 6d. The small red LED was illuminated with the help of the hybrid device which is inserted in Figure 7d. For the comparison of the energy and power density with our fabricated ASC hybrid device, various reported literature studies are compared, as shown in Table 3. Figure 8 shows the Ragone plot of energy versus power density in comparison with other reported results. In addition, cyclic stability was explored for the Cu2FeSnS4/PVP/rGO//AC ASC hybrid device over 20,000 charge–discharge cycles at 5 A/g current density in GCD curves. The 63% of capacitive retention and the 99.99% of Coulombic efficiency were retained over 20,000 cycles. The fabricated hybrid device exhibited excellent performance and the cyclic stability which proves the importance of the Cu2FeSnS4/PVP/rGO electrode with the energy-storage applications.

4. CONCLUSIONS

In summary, the quaternary Cu2FeSnS4/PVP/rGO electrode has been fabricated and used as a potential candidate in supercapacitor application. The remarkable electrochemical performances have been exhibited. The designed Cu2FeSnS4/PVP/rGO electrode offers the enhanced conductivity. The active materials of Cu2FeSnS4/PVP/rGO revealed an excellent specific capacity of 748 C/g at 5 mV/s in CV and retained 86.40% capacity after 5000 cycles. It exhibits 73 W h/kg energy and 749 W/kg power densities. It has a 63% capacity retention and the Coulombic efficiency of 99.99% over 20,000 cycles and explored excellent cyclic stability of the ASC hybrid device. Therefore, this work will serve as a good research electrode material among researchers in the practical need for greater energy use.

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Notes
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