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## Urea-driven hydrothermal synthesis of $\text{Mn}_2\text{O}_3$ : electrochemical performance across various electrolytes for supercapacitor applications

Alisha Dhakal,<sup>ID \*a</sup> Felio Perez<sup>b</sup> and Sanjay R Mishra<sup>\*a</sup>

In this study, cubic  $\text{Mn}_2\text{O}_3$  was synthesized using different urea concentrations (3, 6, 9, and 12 mM) via a hydrothermal method. During synthesis, an increase in urea content resulted in decreased particle and crystallite sizes and increased lattice parameters, with a concomitant increase in the surface area and number of  $\text{Mn}^{3+}$  ions in  $\text{Mn}_2\text{O}_3$  particles. The electrochemical performance of the  $\text{Mn}_2\text{O}_3$ -9 mM urea sample outperformed samples prepared with other urea contents. The  $\text{Mn}_2\text{O}_3$ -9 mM urea sample exhibited high specific capacitance ( $C_{\text{sp}}$ ) values in 1 M and 3 M KOH electrolytes, achieving  $881.3 \text{ F g}^{-1}$  and  $1043.2 \text{ F g}^{-1}$ , respectively, at a scan rate of  $1 \text{ mV s}^{-1}$ . Furthermore, at a current density of  $1 \text{ A g}^{-1}$ , the  $C_{\text{sp}}$  of  $\text{Mn}_2\text{O}_3$  in 1 M KOH was  $758.5 \text{ F g}^{-1}$ . The values increased to  $891.4 \text{ F g}^{-1}$  with energy density and power density of  $44.7 \text{ W h kg}^{-1}$  and  $398.1 \text{ W kg}^{-1}$ , respectively, in 3 M KOH. Owing to the superior electrochemical performance of the  $\text{Mn}_2\text{O}_3$ -9 mM urea sample, its electrochemical performance was assessed in basic KOH and NaOH and neutral  $\text{Na}_2\text{SO}_4$  and  $\text{NaNO}_3$  aqueous electrolytes. Moreover, the  $\text{Mn}_2\text{O}_3$ -9 mM urea sample demonstrated a  $C_{\text{sp}}$  of 721.0 and  $446.3 \text{ F g}^{-1}$  in 3 M concentrations of NaOH and  $\text{NaNO}_3$  electrolytes, respectively. The  $\text{Mn}_2\text{O}_3$ -9 mM urea sample with the highest content of  $\text{Mn}^{3+}$  ions displayed the highest  $C_{\text{sp}}$  in KOH electrolytes compared with the others owing to the smaller hydration radii of  $\text{K}^+$  and high ionic diffusivity and conductivity of  $\text{OH}^-$  compared with other basic and neutral salts. These results highlight that the synthesis process, electrolyte choice, and concentration of electrolytes significantly influence the electrochemical properties of  $\text{Mn}_2\text{O}_3$  battery-type, emphasizing their critical role in optimizing material performance for supercapacitor applications.

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## 1. Introduction

Considering the increasing concerns about global warming and depletion of fossil fuels, more sophisticated energy storage technologies and the immediate development of clean alternative energies are required.<sup>1</sup> In recent years, supercapacitors, often called as electrochemical capacitors, have gained popularity as one of the most appealing energy storage technologies.<sup>2</sup> Supercapacitors are the most promising options for next-generation energy storage devices because they feature faster charge-discharge rates, better power densities, and longer life spans than existing energy storage devices like batteries.<sup>3–5</sup> Supercapacitors store energy either *via* ion adsorption (electrochemical double layer capacitors, and EDLCs) or fast redox reaction (pseudocapacitors). Sometimes, both types of charge storage mechanisms

occur (ion adsorption and fast redox reaction), and they are called hybrid capacitors.<sup>6,7</sup>

EDLCs, which are based on carbon-based compounds (*e.g.*, carbon-based materials and graphenes), usually display very high surface areas, and their electrochemical capacitance is directly proportional to the surface area accessible to the electrolytes.<sup>4,8–11</sup> These supercapacitors can store significantly more energy per unit mass or volume than conventional capacitors by several orders of magnitude. This is because charge separation occurs over a minimal distance within the electrical double-layer at the interface between the electrode and the adjacent electrolyte,<sup>4,12</sup> and a larger amount of charge can be stored on the highly extended electrode surface area created by numerous pores within the high surface-area electrode material.<sup>13</sup> Pseudocapacitors have better energy density than EDLCs, drawing more attention to the energy storage field.<sup>14,15</sup> Among the transition metal oxides (oxides of Mn, Mo, W, Ru, Ni, and Co) used as pseudocapacitors,  $\text{RuO}_2 \cdot \text{xH}_2\text{O}$  shows excellent electrochemical performance;<sup>16</sup> however, its toxicity, less abundance of raw materials, and high cost limit its commercial application.

<sup>a</sup> Department of Physics and Materials Science, The University of Memphis, Memphis, TN 38152, USA. E-mail: adhakal@memphis.edu, srmishra@memphis.edu

<sup>b</sup> Department of Integrated Microscopy Center, The University of Memphis, Memphis, TN 38152, USA



Many researchers have focused on developing electrode materials that are abundant in nature, low-cost, and environmentally friendly. Among them, Mn-oxides like  $\text{MnO}_2$ ,  $\text{Mn}_2\text{O}_3$ , and  $\text{Mn}_3\text{O}_4$  received great attention due to their excellent applications in many fields, including catalysis, magnetic materials, electronics, and supercapacitors.<sup>17</sup> Moreover, they can provide various oxidation states for redox charge transfer, enabling high energy and theoretical specific capacitance.<sup>18–20</sup> The specific capacitance of the materials depends upon various factors like structure, porosity, and electrolytes. Wenya Li *et al.* synthesized porous  $\text{Mn}_2\text{O}_3$  by a facile hydrothermal process and exhibited a specific capacitance of  $191.1 \text{ F g}^{-1}$  at a current density of  $100 \text{ mA g}^{-1}$  in  $0.5 \text{ M Na}_2\text{SO}_4$  electrolyte.<sup>21</sup> Zhen-Yu Li *et al.* synthesized  $\text{Mn}_2\text{O}_3$  nanowalls on nickel foam through the facial hydrothermal route and obtained a specific capacitance of  $480 \text{ F g}^{-1}$  at  $10 \text{ mV s}^{-1}$  scan rate in  $6 \text{ M KOH}$  electrolyte.<sup>22</sup> Ravinder N. Reddy *et al.* prepared amorphous  $\text{Na}_{0.25}\text{MnO}_2 \cdot 0.5\text{H}_2\text{O}$  using the sol-gel method and reported the specific capacitance of  $110 \text{ F g}^{-1}$  at a scan rate of  $5 \text{ mV s}^{-1}$  in  $2 \text{ M KCl}$  electrolyte.<sup>23</sup> Recently, A. Dhakal *et al.* synthesized  $\text{Mn}_2\text{O}_3\text{-}6$  using urea and piperazine by a hydrothermal process and obtained the specific capacitance of  $952 \text{ F g}^{-1}$  at a scan rate of  $1 \text{ mV s}^{-1}$  in  $6 \text{ M KOH}$  electrolyte.<sup>24</sup>

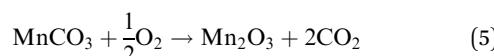
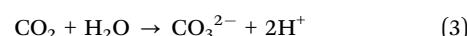
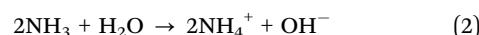
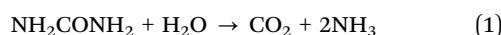
In this work, we synthesized  $\text{Mn}_2\text{O}_3$  using the hydrothermal process and observed the variation of structural and morphological changes with varying concentrations of urea during the synthesis of  $\text{Mn}_2\text{O}_3$ . The electrochemical performance of the as-synthesized  $\text{Mn}_2\text{O}_3$  electrode was assessed in different electrolytes. The choice of electrolyte affects the ionic conductivity, potential window, and overall electrochemical stability, directly impacting the supercapacitor's energy storage capability and efficiency.<sup>25</sup> The suitability of  $\text{Mn}_2\text{O}_3$  as an electrode material for supercapacitors was evaluated in four different electrolytes: potassium hydroxide (KOH), sodium hydroxide (NaOH), sodium sulfate ( $\text{Na}_2\text{SO}_4$ ), and sodium nitrate ( $\text{NaNO}_3$ ) in  $1$  and  $3 \text{ M}$  aqueous solution. Meanwhile, hydroxide salts are strong bases providing a high ionic strength due to the  $\text{OH}^-$  concentration, which can influence the conductivity and the diffusion of reactants. Sulfate and nitrate salts are natural salts having low ionic strength, but provide a more stable environment for electrode materials, reducing the risk of corrosion or degradation compared with alkaline solutions. Thus, selecting an appropriate electrolyte is crucial as it influences the electrode's electrokinetic behavior regarding reaction kinetics, stability, and conductivity, affecting energy and power density performance. By exploring various electrolytes, this study aims to optimize the electrochemical performance of  $\text{Mn}_2\text{O}_3$ , highlighting the profound impact of electrolyte selection on supercapacitor efficiency and stability.

## 2. Experimental section

### 2.1. Synthesis of $\text{Mn}_2\text{O}_3$

The  $\text{Mn}_2\text{O}_3$  compound was synthesized using a hydrothermal method. Manganese(II) nitrate tetrahydrate (2 mmol,  $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ , Sigma-Aldrich Chemicals) and urea ( $x = 3, 6, 9$ , and

$12 \text{ mmol, NH}_2\text{CONH}_2$ , Sigma-Aldrich Chemicals) were mixed in a  $2:x$  ratio in  $40 \text{ mL}$  of deionized water (DI). The solution was magnetically stirred for 1 hour. Consequently, the  $40 \text{ mL}$  of precursor solution was transferred into a stainless-steel autoclave with a Teflon-lined  $50 \text{ mL}$  beaker. The mixture was autoclaved and maintained at a vacuum furnace temperature of  $180 \text{ }^\circ\text{C}$  for 24 h. After cooling to room temperature, the sample was centrifuged and washed multiple times with DI water to achieve a pH of 7. The residue was dried on a hot plate at  $60 \text{ }^\circ\text{C}$  and finally heated in a furnace at  $900 \text{ }^\circ\text{C}$  for three hours. The resulting black powder was  $\text{Mn}_2\text{O}_3\text{-}x$  ( $x = 3, 6, 9$ , and  $12 \text{ mM}$  urea) particles, which are named  $\text{Mn}_2\text{O}_3\text{-}3$ ,  $\text{Mn}_2\text{O}_3\text{-}6$ ,  $\text{Mn}_2\text{O}_3\text{-}9$ , and  $\text{Mn}_2\text{O}_3\text{-}12$ , respectively. The main reactions were related to the following process: (1) urea decomposes at high temperatures and pressure and generates  $\text{CO}_3^{2-}$  and  $\text{OH}^-$  (2)  $\text{Mn}^{2+}$  reacts with  $\text{CO}_3^{2-}$  to form a precipitate of  $\text{MnCO}_3$ . Finally,  $\text{Mn}_2\text{O}_3$  can be obtained after thermal treatment (eqn (1)–(5)).<sup>26,27</sup> Fig. 1 shows the schematic diagram of the  $\text{Mn}_2\text{O}_3$  hydrothermal synthesis process.



### 2.2. Preparation of electrode

Nickel foam with an area of  $1 \text{ cm} \times 1 \text{ cm}$  was cleaned by ultrasonication in  $20 \text{ mL}$  of  $37\%$  HCl solution for 10 minutes. Afterward, the nickel foam was rinsed several times with deionized water and ultrasonicated for 10 minutes in deionized water and ethanol. The cleaned nickel foam was dried at  $100 \text{ }^\circ\text{C}$  for 2 hours in a furnace. For electrode preparation,  $1 \text{ mL}$  of *N*-methylpyrrolidone (NMP) was added to a mixture of  $6 \text{ mg}$   $\text{Mn}_2\text{O}_3$  and  $1 \text{ mg}$  polyvinylidene fluoride (PVDF) and magnetically stirred for 6 hours to ensure uniform distribution. Subsequently,  $0.5 \text{ mL}$  of the solution was applied to the surface of the nickel foam and heated in a furnace at  $100 \text{ }^\circ\text{C}$  for 20 hours. The weight of the sample loaded on the nickel foam was determined by subtracting the clean nickel foam's weight from the nickel foam's weight with the sample applied.

### 2.3. Characterization

$\text{Mn}_2\text{O}_3$  particles were characterized using an X-ray diffractometer (Bruker D8 Advance, Germany) with a  $\text{Cu K}\alpha$  radiation source ( $\lambda = 0.15406 \text{ \AA}$ ,  $40 \text{ kV}$ ,  $40 \text{ mA}$ , step size of  $0.04$ , scan rate of  $0.2 \text{ min per step}$ ,  $20^\circ \leq 2\theta \leq 70^\circ$ ). The morphology of  $\text{Mn}_2\text{O}_3$  particles was evaluated using a field emission scanning electron microscope (FESEM, Hitachi S-4700) with an acceleration voltage of  $20 \text{ kV}$  and an emission current of  $10 \text{ mA}$ . The surface composition and chemical state of  $\text{Mn}_2\text{O}_3$  particles were analyzed using X-ray photoelectron spectroscopy (XPS) with a Thermo Scientific K-Alpha system, employing a  $0.1 \text{ eV}$  step size



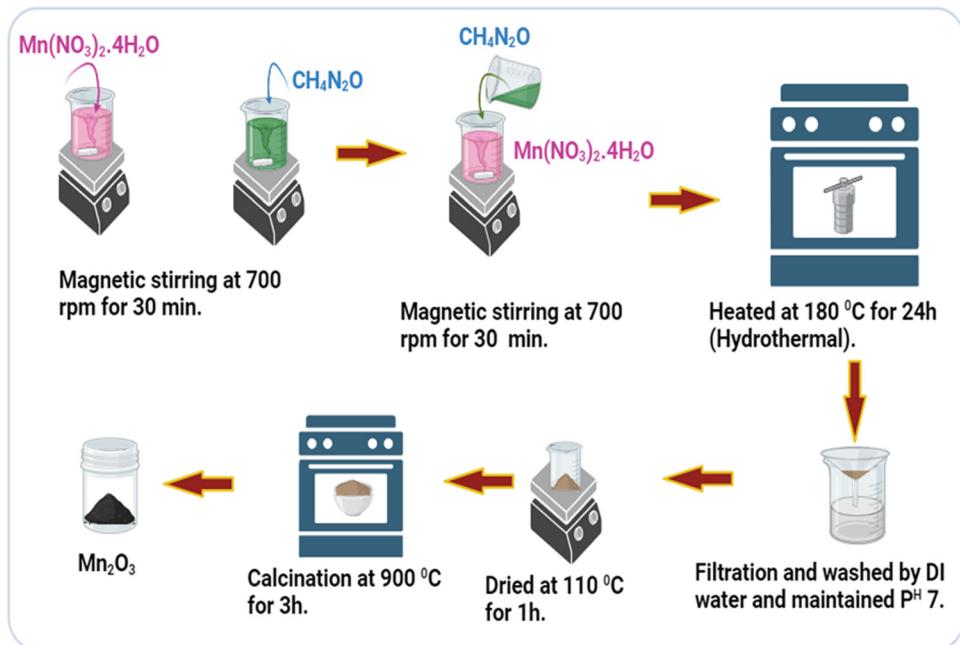


Fig. 1 Schematic of the hydrothermal synthesis process of  $\text{Mn}_2\text{O}_3$ . Created with BioRender.com.

and a standard Al X-ray source (1486.6 eV photons). A surface area analyzer measured the BET surface area and pore size distribution (Quantachrome Nova Touch NT2LX-2) at 77 K using nitrogen as an adsorbate gas. The electrochemical performance of  $\text{Mn}_2\text{O}_3$  particles was evaluated using an electrochemical analyzer (Gamry IFC 1000-07045, USA). The electrochemical properties were studied in a three-electrode configuration within the voltage range of 0 to +0.6 V in 1 and 3 M base electrolytes, *viz.*, KOH, NaOH, and neutral electrolytes, *viz.*,  $\text{NaNO}_3$  and  $\text{Na}_2\text{SO}_4$ , with a platinum plate as the counter electrode, an Ag/AgCl reference electrode, and the nickel foam ( $1 \times 1 \text{ cm}^2$ ) with the loaded sample as the working electrode. In this study, we did not use a 3 M  $\text{Na}_2\text{SO}_4$  electrolyte due to issues with the solubility limit of salt in aqueous solution at room temperature.<sup>28</sup>

### 3. Results and discussion

#### 3.1. Structure and morphology

**3.1.1. X-ray diffraction study.** The phase formation and crystallographic information of  $\text{Mn}_2\text{O}_3\text{-}x$  determined by analyzing the XRD patterns (Fig. 2(a)) and using Rietveld refinement with the GSAS-II program (Fig. 2(b)) present the Rietveld refined XRD patterns, featuring sharp diffraction peaks, which demonstrate the crystalline nature of the sample. The diffraction peaks are attributed to cubic phase  $\text{Mn}_2\text{O}_3$  (PDF 00-041-1442) with a space group of  $Ia\bar{3}$ .

The Williamson–Hall (W–H) equation is used to calculate the crystallite size ( $D$ ) and microstrain ( $\varepsilon$ ) of  $\text{Mn}_2\text{O}_3$ :<sup>29</sup>

$$\beta \times \cos \theta = \frac{k\lambda}{D} + 4\varepsilon \times \sin \theta \quad (6)$$

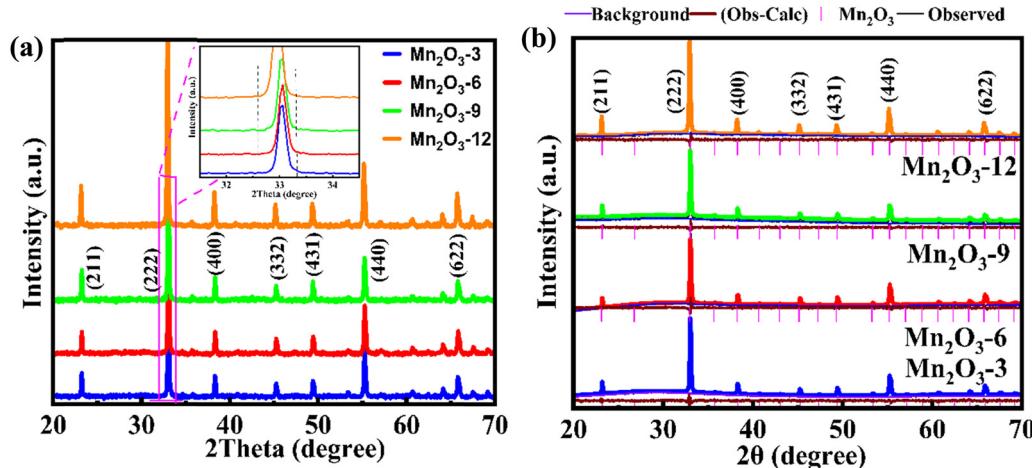


Fig. 2 (a) XRD patterns and (b) Rietveld refined patterns of  $\text{Mn}_2\text{O}_3$ .



Table 1 Structural characteristics of  $\text{Mn}_2\text{O}_{3-x}$ 

Samples	Average particle length ( $\mu\text{m}$ )	$a$ ( $\text{\AA}$ )	$V$ ( $\text{\AA}^3$ )	Crystallite size (nm)	Microstrain ( $10^{-3}$ )
$\text{Mn}_2\text{O}_{3-3}$	11.82	9.408	832.60	129.58	1.52
$\text{Mn}_2\text{O}_{3-6}$	9.94	9.405	831.89	124.91	1.63
$\text{Mn}_2\text{O}_{3-9}$	8.64	9.410	833.16	113.65	1.68
$\text{Mn}_2\text{O}_{3-12}$	8.32	9.412	833.85	87.20	1.73

where  $\beta$  represents the full-width-at-half-maximum (FWHM),  $\theta$  denotes the angle of diffraction,  $k$  is a constant (shape factor  $\sim 0.9$ ),  $\lambda$  represents the wavelength of the X-ray used (0.15406  $\text{\AA}$ ). The crystallite size and average microstrain were calculated using the W–H plot from individual phases of all main peaks at (211), (222), (400), (332), (134), (440), and (622) and presented in Table 1.

As listed in Table 1, the urea content is observed to influence the lattice parameter of  $\text{Mn}_2\text{O}_3$ . Urea can decompose during hydrothermal synthesis to release ammonia ( $\text{NH}_3$ ) and carbon dioxide ( $\text{CO}_2$ ). This may affect the oxidation state of manganese, leading to the creation of oxygen vacancies. An increase in oxygen vacancies can cause lattice expansion due to the relaxation of bonds in the crystal structure, increasing the lattice parameter. Further, upon hydrolysis, urea forms ammonia, raising the pH of the solution. Changes in pH could also lead to different degrees of hydration or hydroxylation at the surface, indirectly affecting the internal structure and lattice parameters.<sup>30</sup> On the other hand, both average particle and crystallite sizes are observed to decrease with the increase in urea content. The rise in pH upon the hydrolysis of urea can increase the supersaturation level of the metal cations, leading to a faster nucleation rate. When the nucleation rate is higher, smaller nuclei form instead of a few large particles, which leads to a decrease in average particle and crystallite size. The increase in pH also affects the surface charge of the growing particles. A higher surface charge can increase repulsive electrostatic interactions between particles, preventing their coalescence and resulting in smaller particles. The steric hindrance or surface passivation effect, and the adsorption of decomposition products such as  $\text{NH}_3$  and  $\text{CO}_2$  reduces the chance of small particles clumping together, contributing to smaller particle sizes.

**3.1.2 Morphological study.** The surface morphology of the synthesized  $\text{Mn}_2\text{O}_3$  was characterized by field emission scanning electron microscopy (FESEM) shown in Fig. 3(a)–(d). The morphology of the synthesized sample is cubic and porous. First, the particles' length was measured using ImageJ software from the FESEM images. Then, the average particle size was estimated by fitting the log-normal distribution as shown in eqn (7).<sup>31</sup>

$$f(L) = \left( \frac{1}{\sqrt{2\pi}\sigma L} \right) \exp \left[ -\frac{(\ln L - \mu)^2}{2\sigma^2} \right] \quad (7)$$

where  $f(L)$  is the function of length ( $L$ ), and  $\mu$  and  $\sigma$  represent the mean and standard deviation. The average length (arithmetic mean of a log-normally distributed variable  $L$ ) is calculated from the  $\exp(\mu + \sigma^2/2)$  expression. The calculated average length of the particles decreased with an increase in the concentration of urea

during the synthesis of  $\text{Mn}_2\text{O}_3$ , with values of 10.56, 9.44, 7.71, and 7.39  $\mu\text{m}$  for  $\text{Mn}_2\text{O}_{3-3}$ ,  $\text{Mn}_2\text{O}_{3-6}$ ,  $\text{Mn}_2\text{O}_{3-9}$ , and  $\text{Mn}_2\text{O}_{3-12}$ , respectively (Fig. 3(a')–(d')). The decrease in average particle length with increasing urea content during the synthesis of  $\text{Mn}_2\text{O}_3$  is primarily due to the influence of urea on nucleation and growth processes. High urea concentrations enhance the nucleation rate, limiting individual particle growth and resulting in smaller sizes.

**3.1.3 Surface adsorption/desorption.** The surface area and porosity of  $\text{Mn}_2\text{O}_3$  are measured using their  $\text{N}_2$  adsorption–desorption isotherms. Fig. 4(a)–(d) shows a type IV isotherm.<sup>32</sup> The pore size distribution of mesoporous  $\text{Mn}_2\text{O}_3$  was calculated from the isotherm (inset of Fig. 4) in the 5–15 nm range. Using a Brunauer–Emmett–Teller (BET) surface analysis, the specific surface area of mesoporous  $\text{Mn}_2\text{O}_{3-3}$ , 6, 9, and 12 are 7.33, 11.54, 9.54, and  $9.93 \text{ m}^2 \text{ g}^{-1}$  with an average pore size of 3.14, 3.10, 3.00, and 2.88 nm, and the total pore volume of 0.012, 0.018, 0.014, and  $0.014 \text{ cm}^3 \text{ g}^{-1}$ , respectively, as listed in Table 2.  $\text{Mn}_2\text{O}_{3-6}$  has the highest specific surface area.

**3.1.4. X-ray photoelectron spectroscopy study.** Fig. 5(a)–(c) shows the XPS plots of  $\text{Mn}_2\text{O}_3$  particles. The XPS survey shows the characteristic peaks for Mn and O in the porous  $\text{Mn}_2\text{O}_3$  particles. The high-resolution Mn 3s spectrum of porous  $\text{Mn}_2\text{O}_3$  cubes peaks at 640.8 eV and 652.7 eV correspond to  $2p_{3/2}$  and  $2p_{1/2}$  transitions, respectively (Fig. 5(a)). This agrees with the reported binding energy values of the oxidation states of  $\text{Mn}^{3+}$ .<sup>23,33</sup> Fig. 5(b) depicts the high-resolution spectrum of O 1s. The binding energy peaks at 528.9 and 530.7 eV indicate Mn–O–Mn formation, as these binding energy peaks are typically attributed to the  $\text{O}^{2-}$ .<sup>34</sup> The deconvoluted peaks of the Mn 2p spectra of  $\text{Mn}_2\text{O}_3$  samples are shown in Fig. 5(c), indicating the presence of  $2^+$ ,  $3^+$ , and  $4^+$  oxidation states. The Mn  $2p_{3/2}$  signal exhibits peaks at 640.5 eV for  $\text{Mn}^{2+}$ , 641.4 eV for  $\text{Mn}^{3+}$ , and 643.0 eV for  $\text{Mn}^{4+}$ . The Mn  $2p_{1/2}$  signal is divided into three peaks at around 652.0 eV for  $\text{Mn}^{2+}$ , 653.0 eV for  $\text{Mn}^{3+}$ , and 654.6 eV for  $\text{Mn}^{4+}$ .<sup>35</sup> These values are consistent with the reported values of  $\text{Mn}_2\text{O}_3$ .<sup>36</sup> The primary oxidation state detected for  $\text{Mn}_2\text{O}_3$  is  $+3$  (magenta), followed by  $+4$  (orange), and there is a very low presence of the  $+2$  oxidation state (cyan color). The oxidation peak ratios of  $\text{Mn}^{3+}/\text{Mn}^{2+}$  are 1.5, 1.7, 3.3, and 1.4, and for  $\text{Mn}^{3+}/\text{Mn}^{4+}$  the oxidation peak ratios are 1.3, 1.2, 1.6, and 1.4 for  $\text{Mn}_2\text{O}_{3-3}$ , 6, 9, and 12, respectively, indicating a predominance of  $\text{Mn}^{3+}$  states in the  $\text{Mn}_2\text{O}_{3-9}$  mM urea sample. The XPS data shows a noticeable oxygen deficiency in  $\text{Mn}_2\text{O}_3$ , as noted in a lower-than-expected O 1s peak intensity. The estimated composition from the XPS data is  $\text{Mn}_2\text{O}_{1.33}$ ,  $\text{Mn}_2\text{O}_{1.28}$ ,  $\text{Mn}_2\text{O}_{1.27}$ , and  $\text{Mn}_2\text{O}_{1.29}$  for  $\text{Mn}_2\text{O}_{3-3}$ , 6, 9, and 12, respectively. This suggests an oxygen vacancy ( $\text{Mn}_2\text{O}_{3-\delta}$ ) or surface reduction of the  $\text{Mn}_2\text{O}_3$  structure, contributing to altered electrochemical properties.

### 3.2. Electrochemical analysis

**3.2.1. Cyclic voltammetry.** The capacitive performance and electrochemical behavior of the supercapacitor's electrode have been determined by three-electrode systems' cyclic voltammetry (CV) measurements. In particular, CV measurements distinguish between capacitive (non-faradaic) and faradaic (redox)



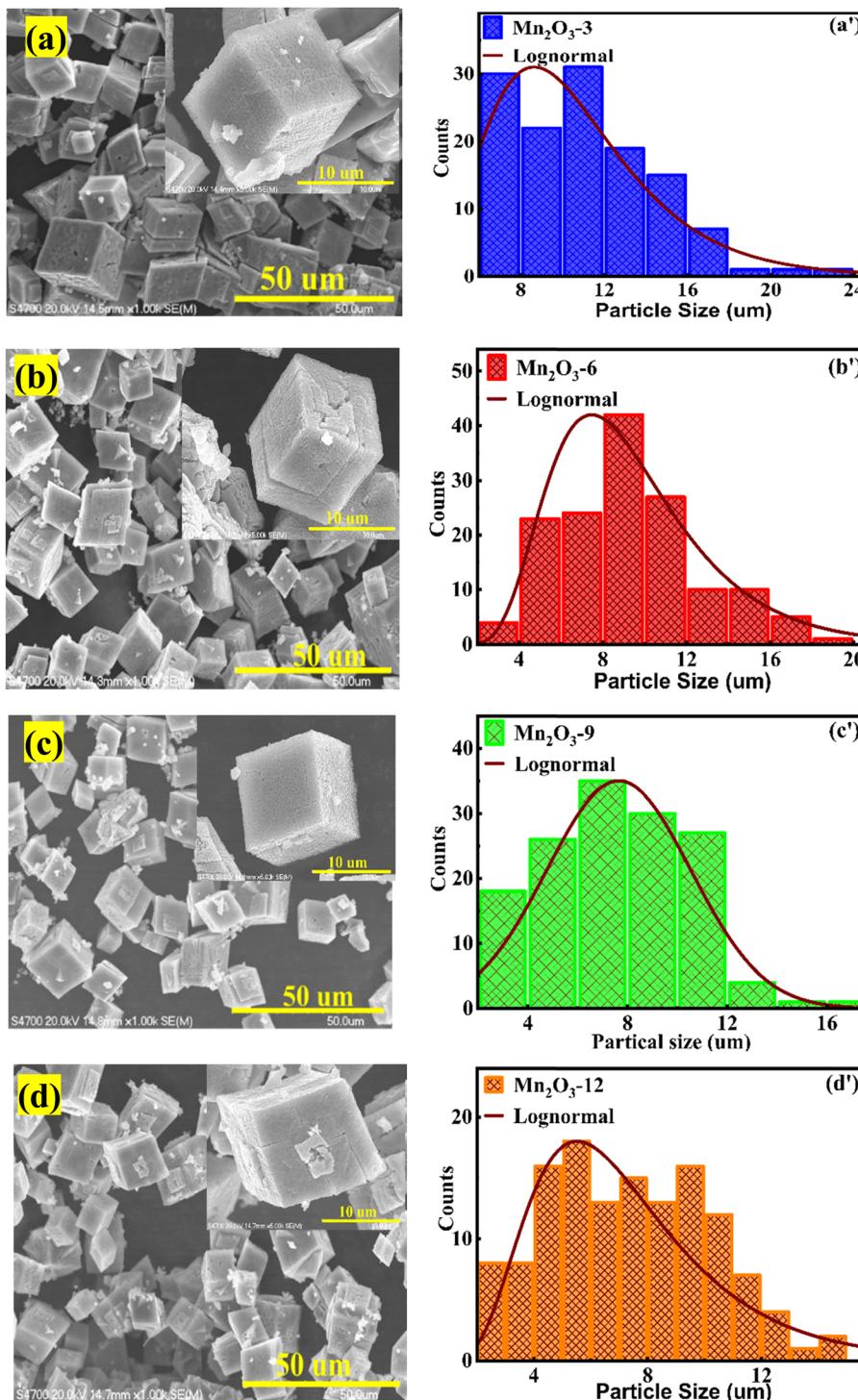


Fig. 3 FESEM images of (a)  $\text{Mn}_2\text{O}_3$ -3, (b)  $\text{Mn}_2\text{O}_3$ -6, (c)  $\text{Mn}_2\text{O}_3$ -9, (d)  $\text{Mn}_2\text{O}_3$ -12, and (a')–(d') histograms with log-normal distribution of length of  $\text{Mn}_2\text{O}_3$  particles.

processes, allowing for a better understanding of the electrode's behavior.

Fig. 6(a)–(d) shows the cyclic voltammetry measurement of  $\text{Mn}_2\text{O}_3$ - $x$  at different scan rates from 1 to 300  $\text{mV s}^{-1}$  in 1 M KOH electrolyte. The scan rate voltammetry provides insight into the rate capability and kinetics of the electrode material.

The noticeable oxidation and reduction peaks represent reversibility, and the shifting of oxidation and reduction peaks with increasing scan rate shows the battery-type behavior of electrodes.<sup>37,38</sup>

The specific capacitance ( $C_{\text{sp}}$ ) (Fig. 7) of  $\text{Mn}_2\text{O}_3$ - $x$  samples was calculated using eqn (14), revealing that  $\text{Mn}_2\text{O}_3$ -9 exhibited



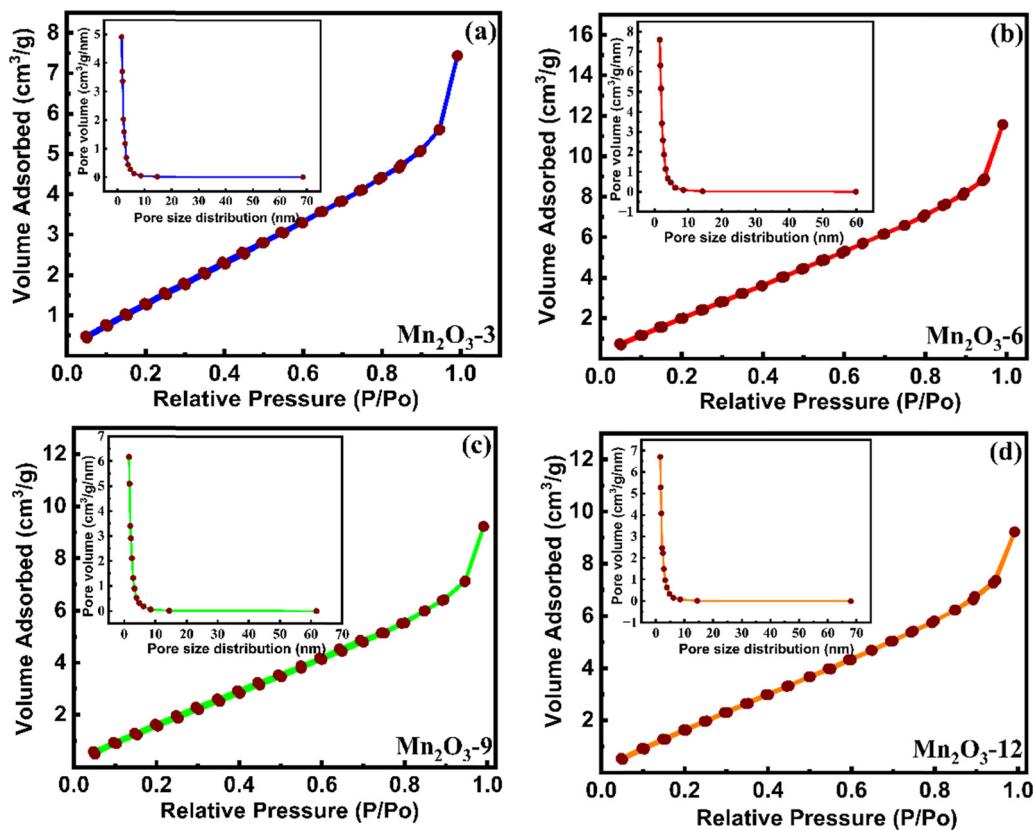


Fig. 4 (a)–(d)  $\text{N}_2$  adsorption and desorption plots of the  $\text{Mn}_2\text{O}_3$  particles measured at 77 K. The inset shows pore size distribution.

Table 2 BET surface area and pore size distribution of  $\text{Mn}_2\text{O}_3$

Compound ( $\text{Mn}_2\text{O}_3$ with different urea ratios)	Weight (g)	BET surface area ( $\text{m}^2 \text{g}^{-1}$ )	Average pore size (radius nm)	Total pore volume ( $\text{cc g}^{-1}$ )
$\text{Mn}_2\text{O}_3\text{-}3$	0.244	7.32	3.15	0.012
$\text{Mn}_2\text{O}_3\text{-}6$	0.185	11.53	3.11	0.018
$\text{Mn}_2\text{O}_3\text{-}9$	0.203	9.54	3.00	0.014
$\text{Mn}_2\text{O}_3\text{-}12$	0.223	9.93	2.88	0.014

a higher  $C_{\text{sp}}$ . The  $C_{\text{sp}}$  values at a scan rate of  $1 \text{ mV s}^{-1}$  were 224, 319, 881, 238, and  $10 \text{ F g}^{-1}$  for  $\text{Mn}_2\text{O}_3\text{-}3$ ,  $\text{Mn}_2\text{O}_3\text{-}6$ ,  $\text{Mn}_2\text{O}_3\text{-}9$ ,  $\text{Mn}_2\text{O}_3\text{-}12$ , and pure Ni form, respectively. Thus, pure Ni foam's specific capacitance is significantly lower than that of  $\text{Mn}_2\text{O}_3\text{-}x$ . Therefore, we can assume that the specific capacitance of

electrodes originates from the  $\text{Mn}_2\text{O}_3$  compound rather than Ni foam. Moreover, the superior  $C_{\text{sp}}$  of  $\text{Mn}_2\text{O}_3\text{-}9$  suggests enhanced electrochemical performance compared with the other concentrations. The specific surface area is not higher in  $\text{Mn}_2\text{O}_3\text{-}9$  than in others; however, most of the Mn ions are in

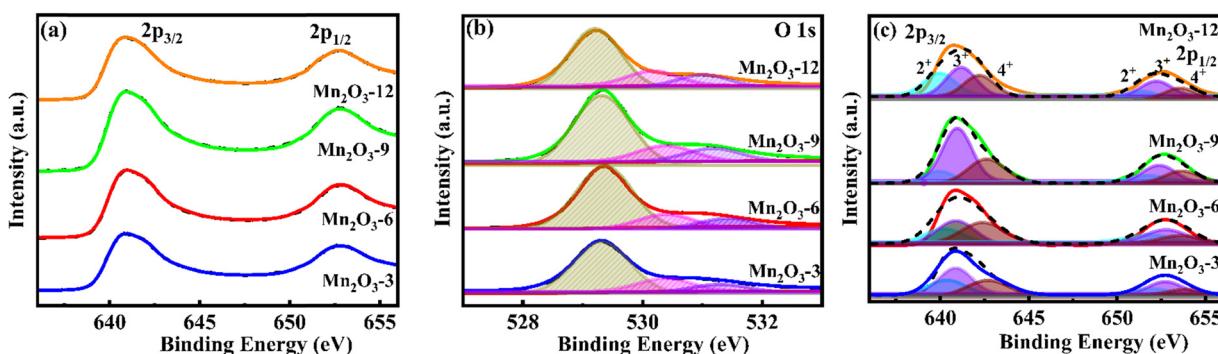


Fig. 5 X-ray photoelectron spectra. (a) High-resolution spectra of Mn 2p, (b) O 1s, and (c) deconvoluted peaks of Mn 2p of  $\text{Mn}_2\text{O}_3$  particles.

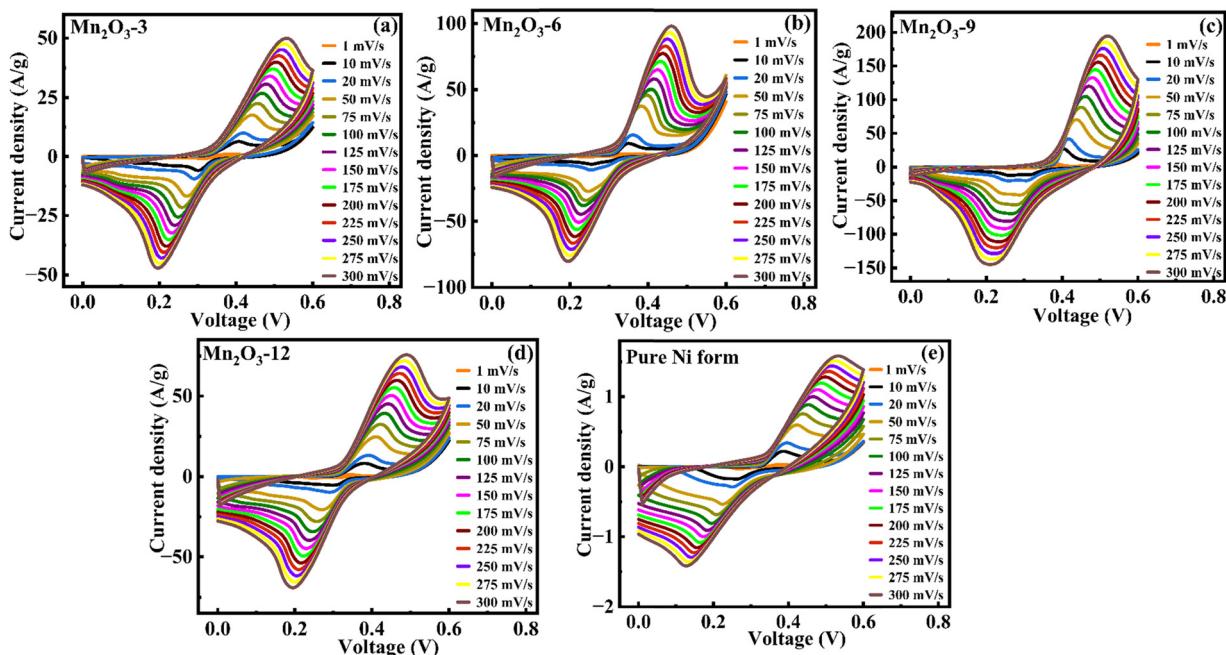


Fig. 6 Cyclic voltammetry curves of (a)–(d)  $\text{Mn}_2\text{O}_3\text{-}x$  and (e) pure Ni form at different scan rates from 1–300  $\text{mV s}^{-1}$  in 1 M KOH electrolyte.

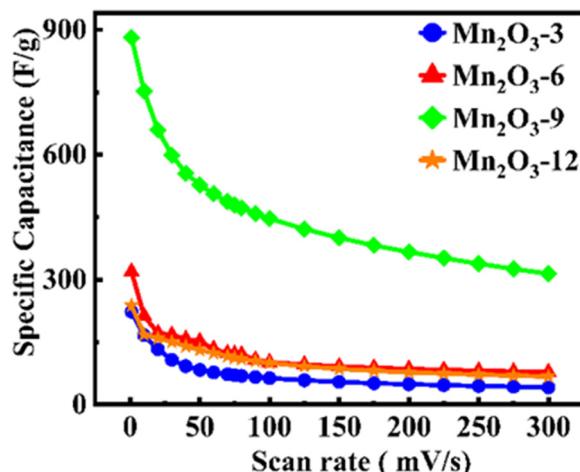


Fig. 7 Specific capacitance ( $C_{\text{sp}}$ ) vs. scan rate of  $\text{Mn}_2\text{O}_3\text{-}x$ .

the  $\text{Mn}^{3+}$  oxidation state, which helps to enhance the electrochemical behavior. The oxidation states of Mn play an important role in the redox rate.  $\text{Mn}^{2+}$  has a half-filled  $3\text{d}^5$  configuration, which is relatively stable due to the symmetry and exchange energy in a half-filled shell. This stability makes  $\text{Mn}^{2+}$  less prone to oxidation (losing an electron) to  $\text{Mn}^{3+}$ . Meanwhile,  $\text{Mn}^{4+}$  has a  $3\text{d}^3$  configuration and is generally more stable than  $\text{Mn}^{3+}$  because it is closer to a noble gas configuration when paired with ligands in a compound. Thus,  $\text{Mn}^{4+}$  tends to be less reactive in redox reactions than  $\text{Mn}^{3+}$ . Alternatively,  $\text{Mn}^{3+}$  has a  $3\text{d}^4$  configuration, which is less stable than  $\text{Mn}^{2+}$  because it lacks the symmetry and stabilization of the half-filled shell. Therefore,  $\text{Mn}^{3+}$  is more reactive in redox reactions, as it can either gain or lose an electron to achieve a more stable state. Also,

abundant oxygen vacancies ( $\text{Mn}_2\text{O}_3\text{-}\delta$ ) provide a fast diffusion pathway for the electrolyte ions and more active sites and help maintain the structural integrity, which is crucial to enhance the electrochemical behavior.<sup>39</sup> Consequently, the  $\text{Mn}_2\text{O}_3\text{-}9$  mM urea sample was selected for further electrochemical characterization in different electrolytes to explore its potential for energy storage applications.

Fig. 8(a) and (b) show the cyclic voltammetry measurement at a scan rate of 1  $\text{mV s}^{-1}$  in 1 and 3 M KOH, NaOH, and  $\text{NaNO}_3$  electrolytes.  $\text{Mn}_2\text{O}_3\text{-}9$  enclosed the highest surface area in 1 M KOH electrolytes compared with other electrolytes. The CV curves' highest enclosed surface area pointed towards the superior electrochemical performance in the KOH electrolytes.

The CV measurements are conducted at different scan rates to understand the reaction kinetics, diffusion processes, electrode behavior, and material characteristics. Fig. 9(a) and (b) show the CV plots of  $\text{Mn}_2\text{O}_3$  electrodes at different scan rates from 1–300  $\text{mV s}^{-1}$  between 0.0 and 0.6 V in 1 and 3 M of various electrolytes, respectively. A pair of redox peaks and non-rectangular shapes of CV curves suggest that the  $\text{Mn}_2\text{O}_3$  electrode stores energy significantly from the dominant faradaic process in different electrolytes. The distinct, broad, symmetric anodic and cathodic peaks of the CV curves indicate a reversible faradaic redox reaction in charge storage involving Mn oxidation states (e.g.,  $\text{Mn}^{2+} \leftrightarrow \text{Mn}^{3+} \leftrightarrow \text{Mn}^{4+}$ ) of  $\text{Mn}_2\text{O}_3$  in different electrolytes. In the electrochemical storage mechanism, the redox interaction of cations  $x^-$  ( $x^- = \text{OH}^-, \text{NO}_3^-, \text{SO}_4^{2-}$  ions from the KOH,  $\text{Na}_2\text{SO}_4$ , and NaOH electrolytes) with  $\text{Mn}_2\text{O}_3\text{-}9$  in different electrolytes can be described using eqn (8) and (9).<sup>40,41</sup>



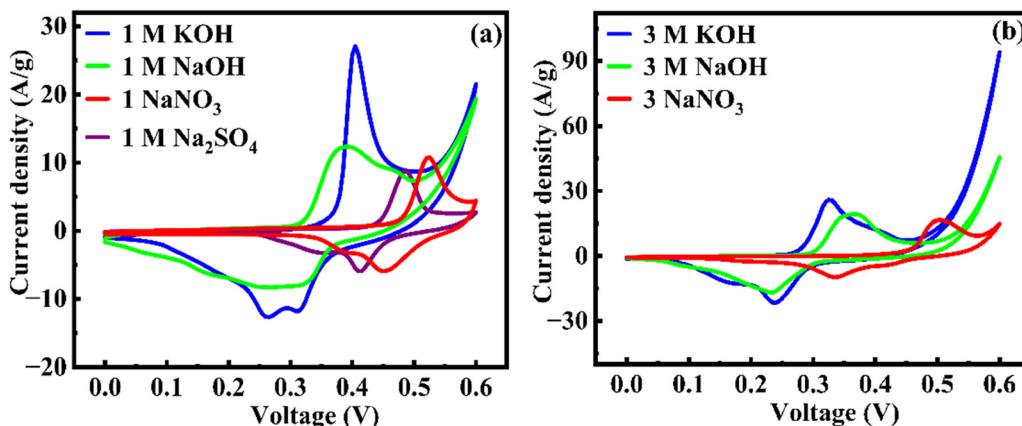


Fig. 8 Cyclic voltammetry curves of  $\text{Mn}_2\text{O}_3$ -9 in (a) 1 M and (b) 3 M concentrations of different electrolytes.

The redox intercalation/de-intercalation of cations ( $\text{M}^+ = \text{Na}^+$ ,  $\text{K}^+$ , and  $\text{H}^+$ ) in the electrode of  $\text{Mn}_2\text{O}_3$  can be expressed in eqn (9):<sup>42</sup>



The charge storage mechanism in electrochemical systems can be predominantly categorized into two types: non-faradaic contributions from the electric double layer (capacitive) and faradaic contributions from diffusion-controlled processes. To identify the dominant contribution to the charge storage mechanism, the relationship between current and scan rate is analyzed using the power law:<sup>43</sup>

$$i = av^b \quad (10)$$

where  $a$  and  $b$  are the adjustable parameters, and exponent  $b$  is derived from the slope of the logarithmic plot of eqn (10). The value of  $b$  typically ranges from 0.5 to 1. Values closer to 0.5 suggest a predominant contribution from diffusion-controlled processes, whereas values approaching 1 indicate a greater contribution from surface charge effects.  $\text{Mn}_2\text{O}_3$  has  $b$ -values of 0.56, 0.68, 0.73, and 0.71 in 1 M KOH, NaOH,  $\text{NaNO}_3$ , and  $\text{Na}_2\text{SO}_4$ , and 0.69, 0.59, and 0.73 in 3 M KOH, NaOH, and  $\text{NaNO}_3$  electrolytes, respectively (Fig. 10). Our observations show that  $b$ -values across all electrolytes are consistently closer to 0.5, indicating that diffusion-controlled contributions predominantly govern the charge storage mechanism in these systems.

The Randles–Sevcik equation is used in CV to analyze the linear relationship between the peak current and the square root of the scan rate ( $i_p \propto v^{1/2}$ ), which gives insight into reaction kinetics and diffusion-controlled processes.<sup>44</sup>

$$i_p = 2.69 \times 10^5 A C_0 n^{3/2} D^{1/2} v^{1/2} \quad (11)$$

where  $i_p$  is the peak current,  $C_0$  is the surface concentration,  $A$  is the surface area,  $D$  represents the diffusion coefficient, and  $v$  is the scan rate. The observed linear relationship between the square root of the scan rate and the current suggests that the predominant contribution to charge storage arises from

diffusion-controlled reactions. The slope of eqn (11) is directly proportional to the square root of the diffusion coefficient. It is observed that  $\text{Mn}_2\text{O}_3$ -9 demonstrated a more pronounced slope in 3 M electrolytes, reflecting enhanced electron transfer efficiency during redox reactions. This indicates that a significant portion of the charge storage is governed by a diffusion-controlled mechanism, as shown in Fig. 11(a) and (b). This implies that increasing the electrolyte concentration enhances the diffusion-controlled charge storage capacity, likely due to efficient ion transport.

As  $b$ -values are not exactly 0.5, the material exhibits a hydride charge storage mechanism: a combination of surface-controlled ( $k_1 v$ ) and diffusion-controlled ( $k_2 v^{1/2}$ ). The quantitative difference between the capacitive and diffusion-controlled contribution to the total charge storage was calculated according to:<sup>45</sup>

$$i(V) = k_1 v + k_2 v^{1/2} \quad (12)$$

For analytical purposes, this equation can be rearranged as:

$$\frac{i(V)}{v^{1/2}} = k_1 v^{1/2} + k_2 \quad (13)$$

where  $v$  is the scan rate. The  $k_1$  and  $k_2$  values are determined from the slope and the  $y$ -intercept of eqn (12), respectively. Eqn (12) and (13) analyze complex electrode processes involving surface-controlled (where  $k_2 = 0$  or  $b = 1$ ) and diffusion-controlled (where  $k_1 = 0$  or  $b = 0.5$ ) ion or electron transfer processes. The quantitative representation of the diffusion-controlled and surface-controlled contributions to total energy storage is presented in the bar graph (Fig. 12(a) and (b)). The diffusion-controlled contribution is more than 50% in all electrolytes, even at a scan rate of  $300 \text{ mV s}^{-1}$ .

The specific capacitance,  $C_{\text{sp}}$ , is estimated by integrating the area under the current–potential curve, as described in eqn (14):<sup>4</sup>

$$C_{\text{sp}} = \frac{1}{m(V_c - V_a)} \int_{V_a}^{V_c} I(V) dV \quad (14)$$

where  $C_{\text{sp}}$  ( $\text{F g}^{-1}$ ) is the specific capacitance,  $m$  is the mass of the active material in the electrode,  $v$  is the scan rate ( $\text{mV s}^{-1}$ ),  $V_c - V_a$  is a potential window, and  $I(V)$  is the response current. The

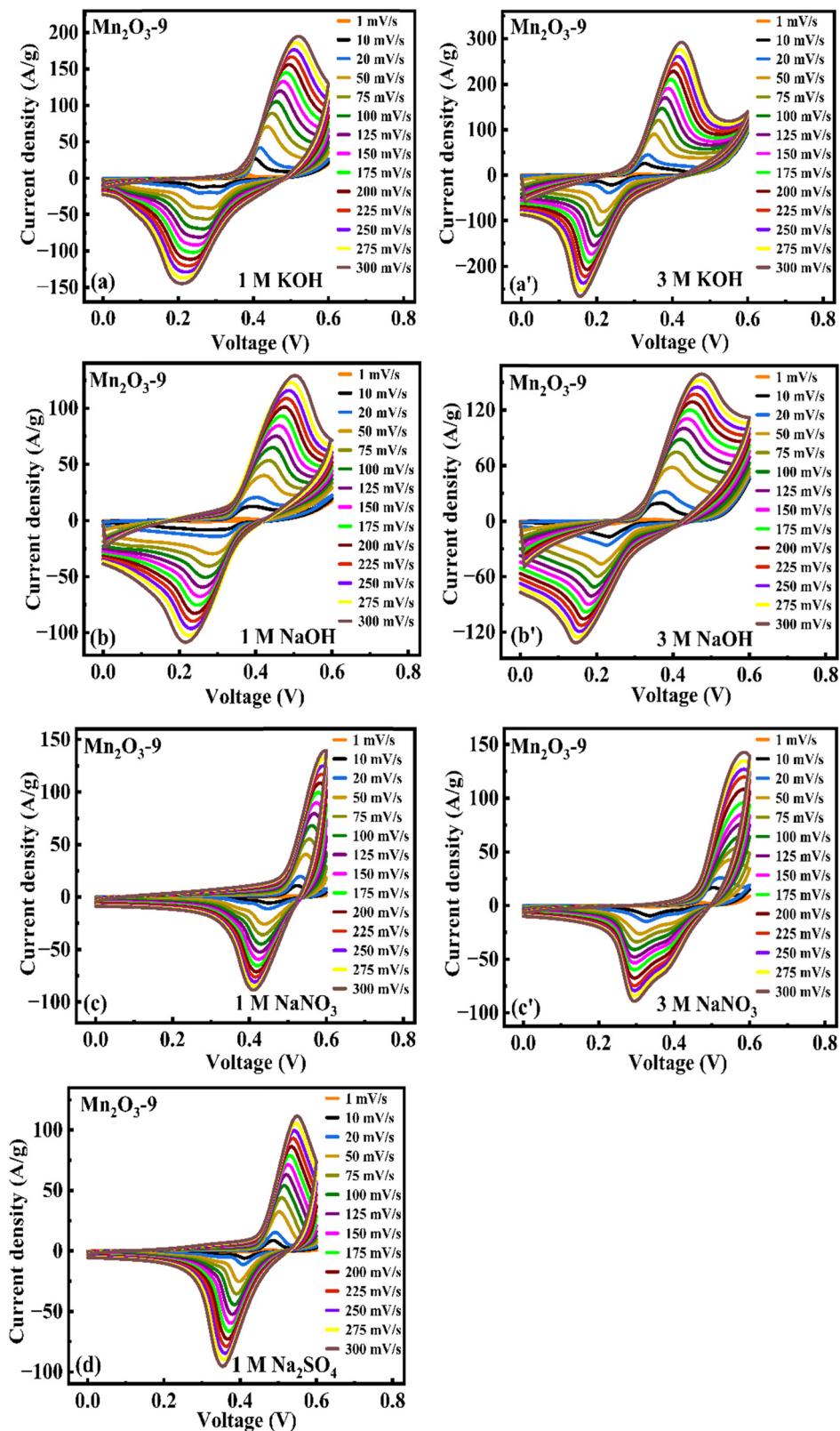


Fig. 9 Cyclic voltammetry curves of  $\text{Mn}_2\text{O}_3$ -9 at different scan rates in (a)–(d) 1 M and (a')–(c') 3 M concentrations of different electrolytes.

integral part gives the area under the current potential curve. As seen in Fig. 13, the mesoporous  $\text{Mn}_2\text{O}_3$ -9 electrode shows the  $C_{\text{sp}}$  of 641, 240, 588, and 300  $\text{F g}^{-1}$  in 1 M KOH,  $\text{Na}_2\text{SO}_4$ , NaOH, and

$\text{NaNO}_3$ , and 1043, 768, and 519  $\text{F g}^{-1}$  in 3 M KOH, NaOH, and  $\text{NaNO}_3$  electrolytes at a scan rate of  $1 \text{ mV s}^{-1}$ , respectively, presented in Table 3. The  $C_{\text{sp}}$  values of KOH electrolytes are

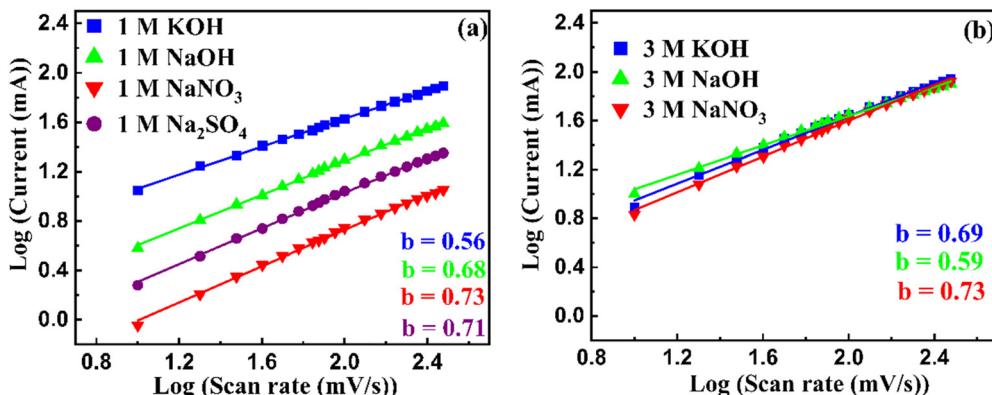
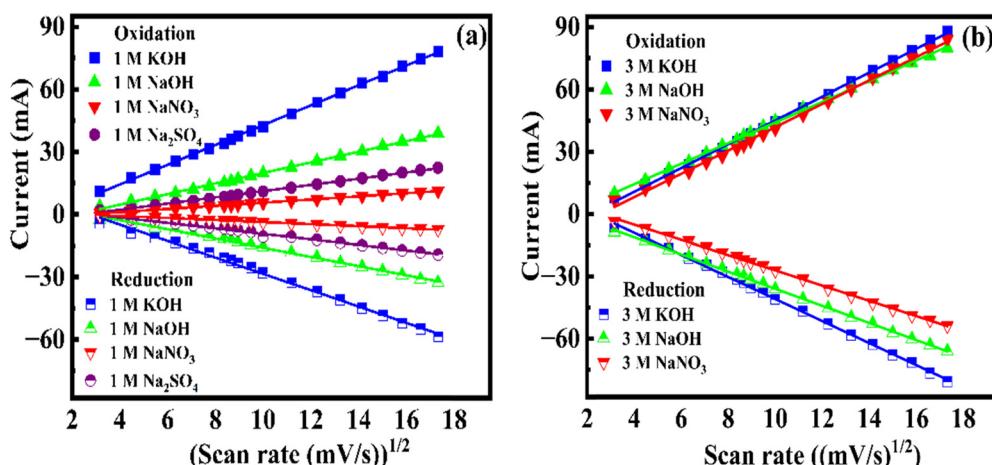
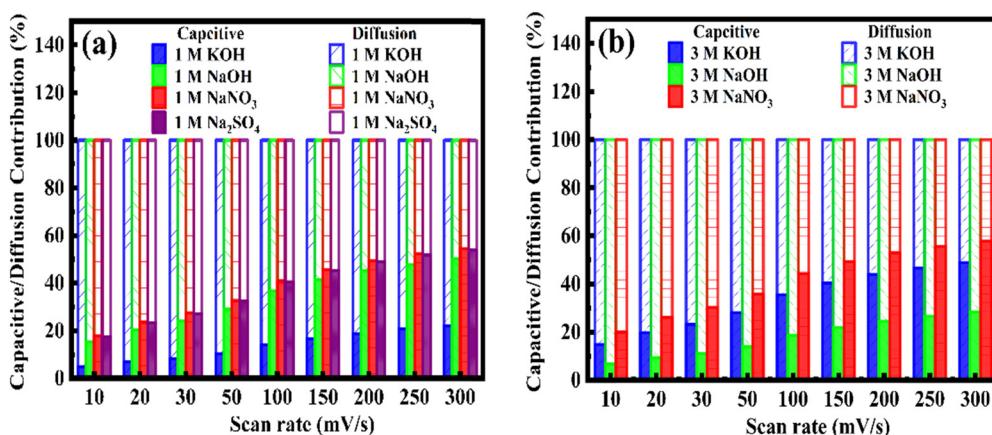
Fig. 10  $b$ -Values plots of  $\text{Mn}_2\text{O}_3$ -9 in (a) 1 M and (b) 3 M concentrations of different electrolytes.

Fig. 11 (a) and (b) Relationship between the peak currents in cyclic voltammetry and the square root of scan rates in 1 M and 3 M concentrations of different electrolytes.

Fig. 12 Percentage capacitive and diffusion contribution to the total charge storage in the  $\text{Mn}_2\text{O}_3$ -9 mM urea sample measured in (a) 1 M and (b) 3 M electrolyte concentrations at different scan rates.

higher than those of others owing to the hydrated cationic radius, cationic mobility, conductivity, and their effect on charge/ion exchange and diffusion. V Nithya *et al.* studied the electrochemical performance of  $\text{Bi}_2\text{WO}_6$  particles in various electrolytes, such as 1 M KOH, 6 M KOH, 1 M LiOH, 1 M NaOH, and 1 M  $\text{Na}_2\text{SO}_4$  solutions. They reported that the current response of  $\text{Bi}_2\text{WO}_6$  in

various electrolytes decreases in the order 6 M KOH > 1 M KOH > 1 M NaOH > 1 M LiOH > 1 M  $\text{Na}_2\text{SO}_4$ . This higher current response of KOH compared with the other electrolytes is due to the different hydration sphere radius of  $\text{K}^+$  (3.31 Å) and  $\text{Na}^+$  (3.58 Å). The cationic radius is higher in  $\text{K}^-\text{H}_2\text{O}^{\delta-} < \text{Na}^-\text{H}_2\text{O}^{\delta-}$ . The reported conductivity of  $\text{K}^+$  ( $73 \text{ cm}^2 \Omega \text{ mol}^{-1}$ ) and

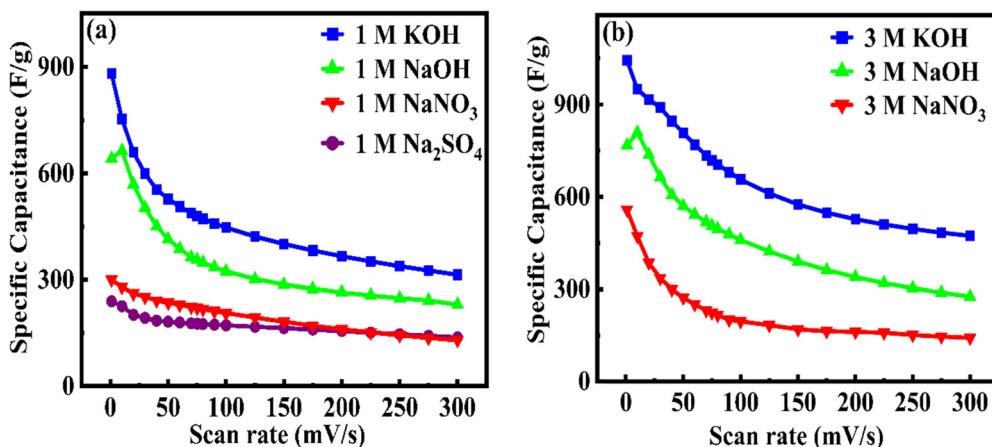


Fig. 13 Relationship between  $C_{sp}$  and the scan rates for  $\text{Mn}_2\text{O}_3\text{-}9$  measured in (a) 1 M and (b) 3 M concentrations of different electrolytes.

Table 3 Specific capacitance,  $C_{sp}$ , of  $\text{Mn}_2\text{O}_3\text{-}9$  in different electrolytes at a scan rate of  $1 \text{ mV s}^{-1}$

Electrolytes	$C_{sp}$ ( $\text{F g}^{-1}$ )	
	1 M	3 M
KOH	881.3	1043.2
NaOH	641.1	767.8
$\text{NaNO}_3$	299.7	558.4
$\text{Na}_2\text{SO}_4$	239.9	—

$\text{Na}^+$  ( $50 \text{ cm}^2 \Omega \text{ mol}^{-1}$ ) at room temperature implies that the mobility of ions is higher for  $\text{K}^+$  than  $\text{Na}^+$ . By changing the anionic species, higher mobility and conductivity were observed. The  $\text{OH}^-$  ion has higher conductivity ( $198 \text{ cm}^2 \Omega \text{ mol}^{-1}$ ) and mobility ( $2.06 \times 10^{-6} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ) compared with other  $\text{NO}_3^-$  and  $\text{SO}_4^{2-}$  ions.<sup>46</sup> Therefore, due to the smaller hydrated ions and higher mobility and conductivity of  $\text{K}^+$  ions,  $\text{Mn}_2\text{O}_3$  displays the highest  $C_{sp}$  in 1 M and 3 M KOH electrolytes. Additionally, owing to the smaller hydration sphere radius of  $\text{K}^+$  ions, they acquire a small charge density (weaker solvation interactions with water molecules, which favor easier polarization during the desolvation process), which further helps in easy transfer of  $\text{K}^+$  ions into the electrode during the redox reactions.<sup>47</sup>

The  $C_{sp}$  of the  $\text{Mn}_2\text{O}_3$  electrode in  $\text{NaOH}$ ,  $\text{Na}_2\text{SO}_4$ , and  $\text{NaNO}_3$  electrolytes has different values. The results show that electrodes display higher  $C_{sp}$  in KOH electrolytes. The  $C_{sp}$  follows  $\text{OH}^- > \text{NO}_3^- > \text{SO}_4^{2-}$  due to the increased anionic size of the nitrate and sulfate ions compared with hydroxyl ions.<sup>48</sup> The  $C_{sp}$  values increased with electrolyte concentrations (1 and 3 M), implying that higher-concentration electrolytes have more ions. The abundance of ions leads to more charge carriers, which can efficiently move between the electrode and the electrolyte, increasing the overall mobility and conductivity.

**3.2.2 Charge-discharge studies.** Further, charge-discharge experiments were conducted to test the ability of the capacitors. Fig. 14(a) and (b) show the charge-discharge curves at a current density of  $1 \text{ A g}^{-1}$  of  $\text{Mn}_2\text{O}_3\text{-}9$  in different electrolytes: KOH, NaOH,  $\text{NaNO}_3$ , and  $\text{Na}_2\text{SO}_4$  at 1 and 3 M (except  $\text{Na}_2\text{SO}_4$ ). In both cases, the charge/discharge characteristics of  $\text{Mn}_2\text{O}_3\text{-}9$  showed asymmetry between the nonlinear charge and discharge curves, further proving the strong faradaic behavior of the electrode.

The charge-discharge measurements were also carried out at different current densities from 1 to  $15 \text{ A g}^{-1}$  to understand rate capability, capacity retention, and charge storage kinetics. Fig. 15(a) and (b) show the charge-discharge curves of  $\text{Mn}_2\text{O}_3\text{-}9$  at different current densities.

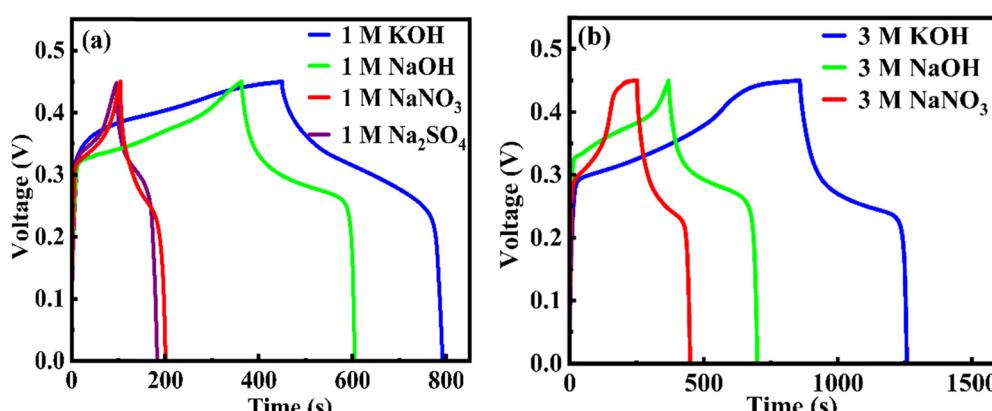


Fig. 14 Chronopotentiometry charge-discharge curves of  $\text{Mn}_2\text{O}_3\text{-}9$  at a current density of  $1 \text{ A g}^{-1}$  in (a) 1 M and (b) 3 M concentrations of different electrolytes.



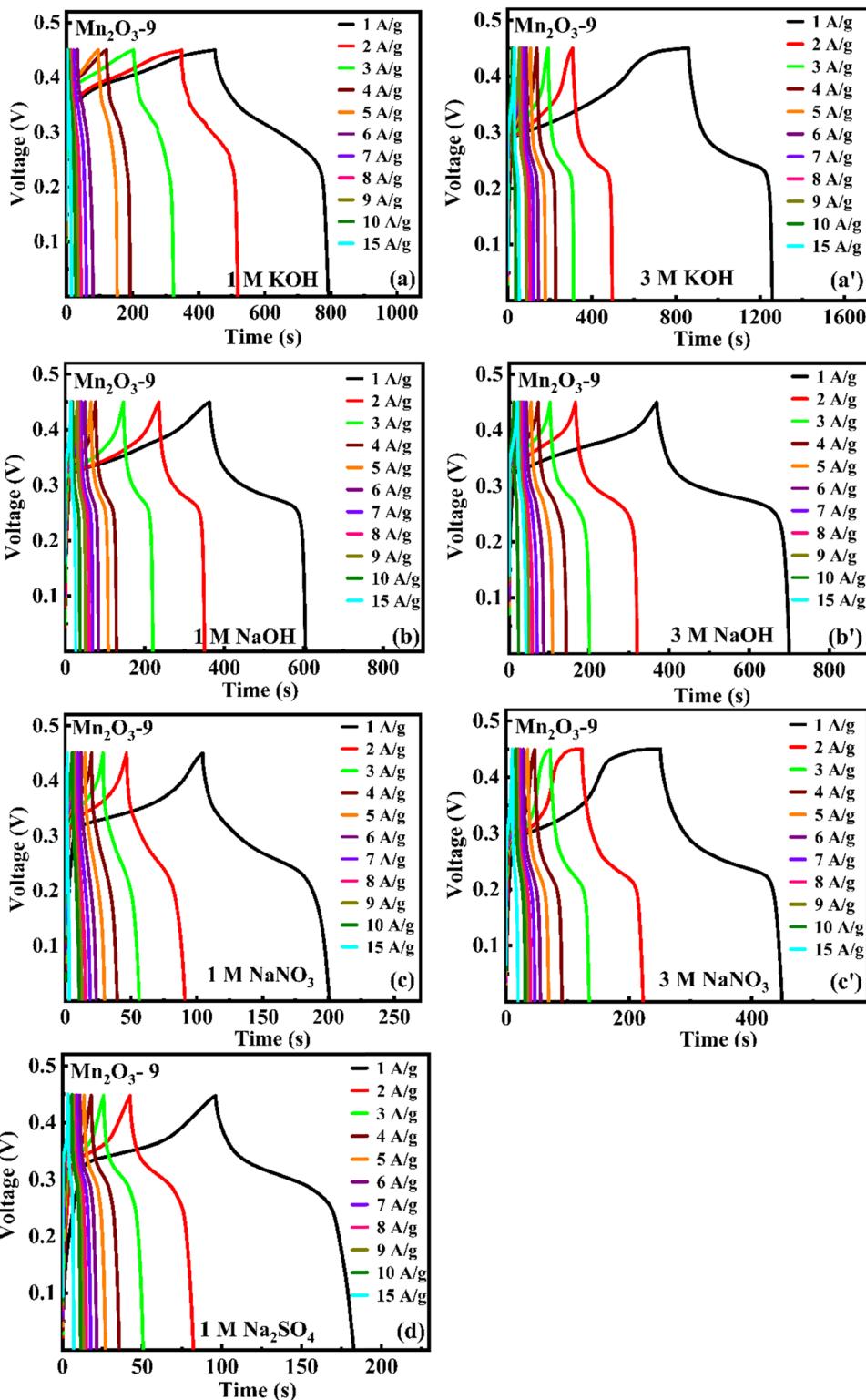


Fig. 15 Chronopotentiometry charge/discharge curves of  $\text{Mn}_2\text{O}_3$ -9 at different current densities in (a)–(d) 1 M and (a')–(c') 3 M concentrations of different electrolytes.

$\text{Mn}_2\text{O}_3$  undergoes pseudocapacitive behavior in alkaline KOH and NaOH electrolytes. In these electrolytes,  $\text{Mn}_2\text{O}_3$  participates in faradaic redox reactions, where it undergoes reversible changes in its oxidation states (e.g.,  $\text{Mn}^{3+} \leftrightarrow \text{Mn}^{4+}$ )

and forms manganese oxyhydroxides ( $\text{MnOOH}$ ) at the surface. This interaction allows the material to efficiently store charges, contributing to a higher capacitance and ultimately, a longer discharge time. Meanwhile, in neutral electrolytes ( $\text{Na}_2\text{SO}_4$  and

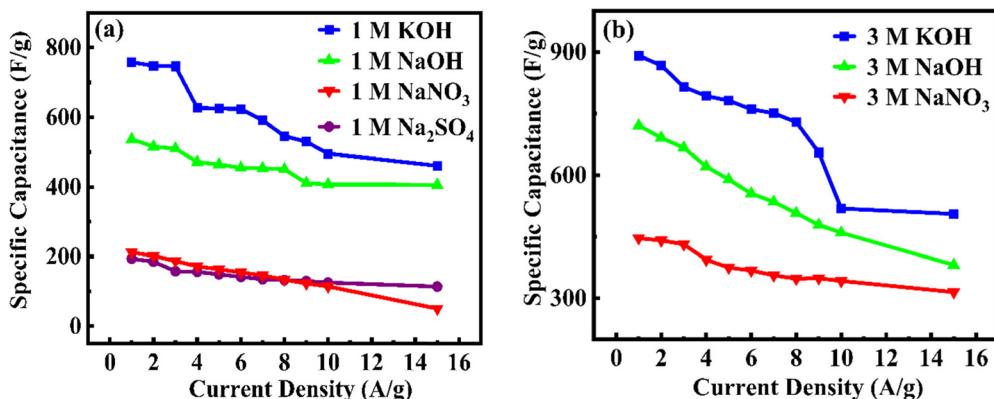


Fig. 16  $C_{sp}$  vs. current densities measured via chronopotentiometry in different electrolytes at (a) 1 M and (b) 3 M concentrations of electrolytes.

Table 4 Specific capacitance,  $C_{sp}$ , energy density, and power density of  $\text{Mn}_2\text{O}_3\text{-9}$  in different electrolytes measured at  $1 \text{ A g}^{-1}$  current density

$\text{Mn}_2\text{O}_3\text{-9}$	Electrolytes	$C_{sp}$ ( $\text{F g}^{-1}$ )	Energy density ( $\text{W h kg}^{-1}$ )	Power density ( $\text{W kg}^{-1}$ )
1 M	KOH	758.5	10.6	397.7
	NaOH	536.5	26.8	398.3
	NaNO <sub>3</sub>	211.9	10.6	397.7
	Na <sub>2</sub> SO <sub>4</sub>	193.8	9.7	400.6
3 M	KOH	891.4	44.6	398.1
	NaOH	721.0	36.0	394.7
	NaNO <sub>3</sub>	446.3	22.3	397.3

$\text{NaNO}_3$ ),  $\text{Mn}_2\text{O}_3$  does not undergo the same redox reactions or surface transformations. Thus, the material cannot take advantage of pseudocapacitive behavior to the same extent. The charge storage process is primarily an EDLC type, which is less efficient and provides lower charge storage capacity and a shorter discharge time in these neutral electrolytes.

The  $C_{sp}$  of  $\text{Mn}_2\text{O}_3\text{-9}$  (Fig. 16) was calculated from the charge-discharge curve using eqn (15):<sup>49</sup>

$$C_{sp} = \frac{I_m \times \Delta t}{\Delta V} \quad (15)$$

where  $I_m$  is the current density,  $m$  is the mass of the active material,  $\Delta t$  is the discharge time, and  $\Delta V$  is the potential window. The  $C_{sp}$  is highest at a current density of  $1 \text{ A g}^{-1}$  in all electrolytes and decreases with increasing current density. As the current density increases, the charge/discharge processes occur more rapidly, limiting the time available for ions to diffuse into deeper or less accessible parts of the electrode. This implies incomplete utilization of the active material, reduced ion adsorption, and decreased  $C_{sp}$ . The  $C_{sp}$  of  $\text{Mn}_2\text{O}_3\text{-9}$  is  $759 \text{ F g}^{-1}$  and  $891 \text{ F g}^{-1}$  in 1 and 3 M KOH electrolytes, respectively, which is higher than that in other aquatic electrolytes, presented in Table 4.

Fig. 17(a) and (b) show the Ragone plot of  $\text{Mn}_2\text{O}_3\text{-9}$ , which is the relation between the log of energy density and power density. Energy density and power density are important parameters for understanding the electrochemical behavior of the materials. The energy density ( $E$ ,  $\text{W h kg}^{-1}$ ) and power density ( $P$ ,  $\text{W kg}^{-1}$ ) are calculated from eqn (16) and (17):<sup>50</sup>

$$E = \frac{0.5 \times C_{sp} \times \Delta V^2}{3.6} \quad (16)$$

$$P = \frac{3600 \times E}{\Delta t} \quad (17)$$

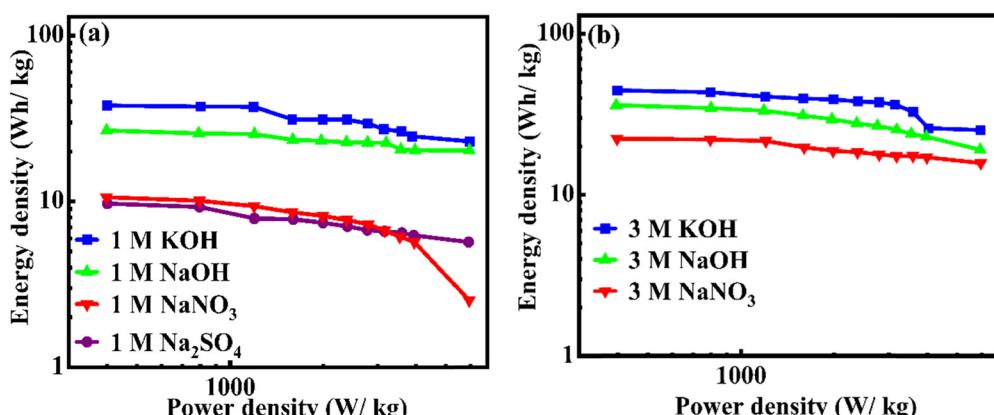


Fig. 17 Ragone plot of  $\text{Mn}_2\text{O}_3\text{-9}$  in (a) 1 M and (b) 3 M concentrations of electrolytes measured at a current density range from 1 to  $15 \text{ A g}^{-1}$ .



where  $E$  is the energy density ( $\text{W h kg}^{-1}$ ),  $C_{\text{sp}}$  represents specific capacitance ( $\text{F g}^{-1}$ ), and  $\Delta V$  is the voltage window (V);  $P$  is the power density ( $\text{W kg}^{-1}$ ), and  $t$  is the discharge time (s).

In Fig. 17(a) and (b),  $C_{\text{sp}}$  in 1 and 3 M KOH exhibits the highest energy densities across all power densities, demonstrating superior performance to other electrolytes.

The stability test of  $\text{Mn}_2\text{O}_3$ -9 electrode was evaluated for 5000 cycles at  $10 \text{ A g}^{-1}$  current density. Fig. 18(a) and (b) present the cyclic stability in different electrolytes. The electrode demonstrates excellent stability throughout the cycling process across all electrolytes and concentrations (1 and 3 M). After 5000 cycles, the  $C_{\text{sp}}$  retention is 89.5, 99.4, 81.0, and 86.1% in 1 M of KOH, NaOH,  $\text{NaNO}_3$ , and  $\text{Na}_2\text{SO}_4$  electrolytes, respectively. Moreover, the  $C_{\text{sp}}$  retention is 93.0, 94.0, and 89.5%, in 3 M of KOH, NaOH, and  $\text{NaNO}_3$  electrolytes, respectively. The inset graphs in Fig. 18(a) and (b) show the Coulombic efficiency of  $\text{Mn}_2\text{O}_3$ -9 electrode.

The Coulombic efficiency (CE) is calculated as the ratio of the charge released during the discharge ( $t_{\text{D}}$ ) to the charge input during charging ( $t_{\text{C}}$ ), expressed as a percentage using the formula:  $\text{CE} = \frac{t_{\text{D}}}{t_{\text{C}}} \times 100\%$ . The CE values are almost 100% for the electrode for every electrolyte used until 5000 cycles. The  $C_{\text{sp}}$

and energy density are higher in our work than in previously reported electrodes, as presented in Table 5. The inset in Fig. 18 shows that  $\text{Mn}_2\text{O}_3$ -9 has a higher Coulombic efficiency in NaOH,  $\text{NaNO}_3$ , and  $\text{Na}_2\text{SO}_4$  than KOH electrolytes, which might be because  $\text{Na}^+$  ions are smaller than  $\text{K}^+$  ions. The smaller the ions, the more efficiently they can migrate through the electrolyte and participate in the electrochemical reactions.<sup>51</sup>

**3.2.3. Electrochemical impedance spectroscopy (EIS).** To understand the electrochemical and electrical properties of the electrode, EIS was carried out on the synthesized  $\text{Mn}_2\text{O}_3$ -9. Fig. 19(a) and (b) show the EIS plots conducted in the frequency range of 0.1 to 100 kHz. The inset is the fitted equivalent circuit using “Gamry Echem Analysis” software, which consists of solution resistance ( $R_1$ ), constant phase element ( $\text{CPE}_1$ ,  $\text{CPE}_2$ ), charge-transfer resistance ( $R_2$ ), and Warburg impedance ( $W_{01}$ ). The fitted data of the equivalent circuit are presented in Table 6.

The Nyquist plot is divided into three parts. In the high-frequency region, the X-axis touching the intercept of curves represents the internal resistance ( $R_1$ ), which includes the intrinsic resistance of the active material's contact resistance between the active materials and the current collector interface, and the ionic resistance of the electrolyte. The straight line in the low-frequency region is the Warburg impedance, known as

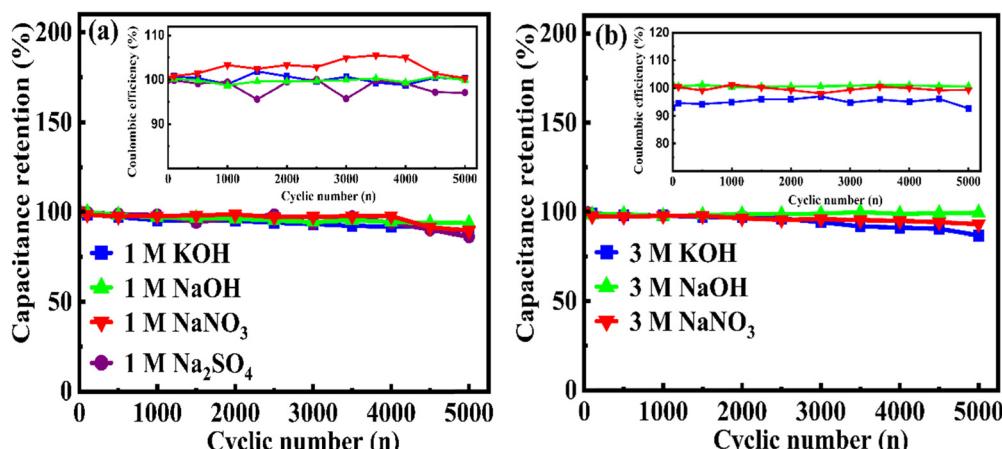


Fig. 18 Cyclic performance of  $\text{Mn}_2\text{O}_3$ -9 in (a) 1 M and (b) 3 M concentrations of electrolytes. The inset displays the Coulombic efficiency (CE) measured at  $10 \text{ A g}^{-1}$  current density.

Table 5 Comparison of the electrochemical performance of  $\text{Mn}_2\text{O}_3$  with previously reported electrodes

Material	Current density or scan rate	Electrolyte	$C_{\text{sp}}$ ( $\text{F g}^{-1}$ )	Energy density ( $\text{W h kg}^{-1}$ )	Power density ( $\text{W kg}^{-1}$ )	Ref.
$\text{Mn}_2\text{O}_3$ nanoparticle	$10 \text{ mV s}^{-1}$	6 M KOH	460	—	—	52
$\text{Mn}_2\text{O}_3$ -piperazine	$1 \text{ A g}^{-1}$	6 M KOH	809.0	40.4	365	24
$\text{Mn}_2\text{O}_3$	$1 \text{ A g}^{-1}$	1 M $\text{Na}_2\text{SO}_4$	209.1	—	—	53
$\text{Mn}_2\text{O}_3$ @nickel foam	$5 \text{ mV s}^{-1}$	LiCl	781	—	—	54
$\text{Mn}_2\text{O}_3$ -carbon aerogel	$1 \text{ mV s}^{-1}$	6 M KOH	368	—	—	55
$\text{Mn}_2\text{O}_3$ -9	$1 \text{ A g}^{-1}$	3 M KOH	891.4	44.6	398.1	Present work
$\text{Mn}_2\text{O}_3$ -9	$1 \text{ A g}^{-1}$	1 M $\text{Na}_2\text{SO}_4$	193.8	9.7	400.6	Present work
$\text{Mn}_2\text{O}_3$ -9	$1 \text{ A g}^{-1}$	3 M $\text{NaNO}_3$	446.3	22.3	397.3	Present work
$\text{Mn}_2\text{O}_3$ -9	$1 \text{ A g}^{-1}$	3 M NaOH	721.0	36.0	394.7	Present work
$\text{Mn}_2\text{O}_3$ nanofiber	$0.5 \text{ A g}^{-1}$	0.5 M $\text{Na}_2\text{SO}_4$	358	46.4	4.42	56
Porous $\text{Mn}_2\text{O}_3$ nanobars	$0.2 \text{ A g}^{-1}$	3 M KOH	249.7	—	—	57
$\text{Mn}_2\text{O}_3$ -cable-like nanofibers	$0.5 \text{ A g}^{-1}$	6 M KOH	216	—	—	58
$\text{Mn}_2\text{O}_3$ -carbon	$5 \text{ mV g}^{-1}$	6 M KOH	651	—	—	59



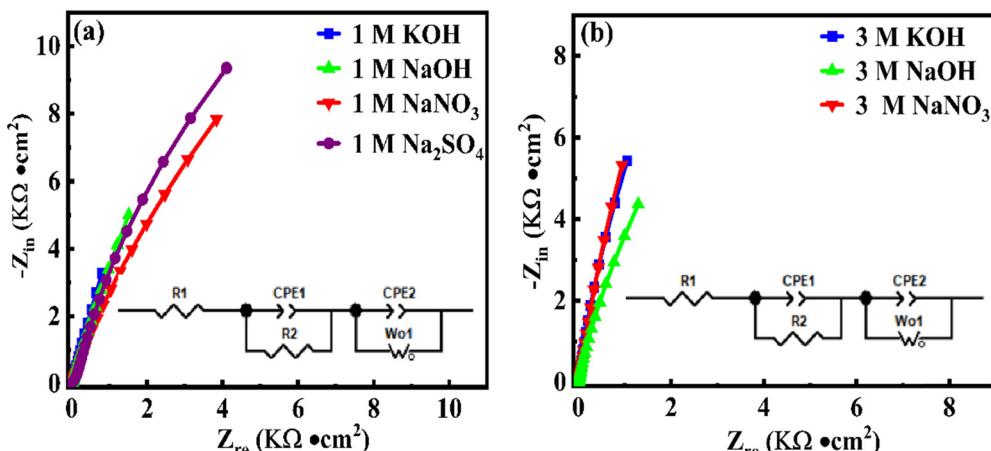


Fig. 19 Nyquist plots of  $\text{Mn}_2\text{O}_3$ -9 in (a) 1 M and (b) 3 M concentrations of different electrolytes. The insets are the equivalent circuits.

Table 6 Fitted parameters derived from fitting EIS curves using the equivalent circuit method for electrodes of  $\text{Mn}_2\text{O}_3$ -9 (Fig. 19)

$\text{Mn}_2\text{O}_3$ -9		$R_1$ ( $\Omega$ )	$R_2$ ( $\Omega$ )	$W_0$ ( $\Omega \times 10^{-3}$ )	CPE <sub>1</sub> ( $\alpha$ )	CPE <sub>1</sub> ( $Q; \Omega^{-1} \text{s}^\alpha$ )	CPE <sub>2</sub> ( $\alpha$ )	CPE <sub>2</sub> ( $Q; \Omega^{-1} \text{s}^\alpha$ )	Goodness of fit ( $\times 10^{-3}$ )
1 M	KOH	1.878	14.98	0.211	0.719	0.004	0.914	0.003	0.171
	NaOH	1.207	9.801	0.066	0.719	0.004	0.856	0.002	0.140
	NaNO <sub>3</sub>	2.217	83.73	0.254	0.732	0.002	0.951	0.001	0.656
	Na <sub>2</sub> SO <sub>4</sub>	3.427	129.8	0.179	0.736	0.001	0.958	0.001	0.682
3 M	KOH	1.182	9.941	0.126	0.638	0.01	0.960	0.002	0.166
	NaOH	0.900	5.221	0.213	0.701	0.007	0.926	0.002	0.183
	NaNO <sub>3</sub>	6.606	38.75	0.153	0.634	0.005	0.979	0.002	0.614

the ion diffusion resistance from the electrolyte to the electrode interface. The middle part is the charge-transfer resistance  $R_2$ , which is the diameter of the semicircle.<sup>60,61</sup> The absence of a semicircular curve indicates that the  $R_2$  values are small and that the ions or charge can easily transfer between the electrode and electrolyte. The  $R_2$  values of  $\text{Mn}_2\text{O}_3$ -9 are around 15, 10, 84, 130  $\Omega$  for 1 M KOH, NaOH, NaNO<sub>3</sub>, Na<sub>2</sub>SO<sub>4</sub>, and 10, 39, and 56  $\Omega$  for 3 M KOH, NaOH, NaNO<sub>3</sub> electrolytes, respectively. The frequency dispersion is generally attributed to a phenomenon known as “capacitance dispersion,” which is described using a constant phase element (CPE) and used in modeling the non-ideal capacitive behavior in electrochemical systems, particularly at the electrode–electrolyte interface marred with surface roughness, inhomogeneities, or complex geometries. The impedance of the CPE is expressed as  $Z_{\text{CPE}} = \frac{1}{Q(j\omega)^\alpha}$ , where  $Q$  ( $\Omega^{-1} \text{s}^\alpha$ ) is the CPE parameter and  $\alpha$  is the CPE exponent. The exponent  $\alpha$  ( $0 \leq \alpha \leq 1$ ) indicates the deviation from ideal capacitive behavior. The CPE behaves as a capacitor for  $\alpha = 1$ , a resistor for  $\alpha = 0$ , an inductor for  $\alpha = -1$ , and the Warburg impedance for  $\alpha = 0.5$ .<sup>62</sup> This result indicates that the NaOH electrolyte exhibits better conductivity for  $\text{Mn}_2\text{O}_3$ -9.

As evident from the above discussion, the  $C_{\text{sp}}$  of  $\text{Mn}_2\text{O}_3$  is notably higher in KOH electrolytes than in other electrolytes. The superior performance of electrodes in KOH electrolytes can be understood in terms of the molecular size of the hydrated ions and the molar conductance of electrolytes. As per the literature, the  $\text{Na}^+$  ionic radius of 0.95  $\text{\AA}$  is smaller than the  $\text{K}^+$

ionic radius of 1.33  $\text{\AA}$ .<sup>63,64</sup> This implies that  $\text{Na}^+$  can easily intercalate compared to  $\text{K}^+$ . However,  $\text{Mn}_2\text{O}_3$  displays the highest  $C_{\text{sp}}$  in KOH, followed by NaOH. Contrary to the expectation, this result is explained based on the ionic radii of hydrated ions.<sup>51,65</sup> In fact, the smaller ions form larger hydrated ions. For example, the hydrated cation of  $\text{Li}^+$  has an ionic radius of 3.82  $\text{\AA}$ , which is larger than the hydrated cation ionic radius of  $\text{Na}^+$  of 3.58  $\text{\AA}$  and the hydrated cation ionic radius of  $\text{K}^+$  of 3.31  $\text{\AA}$ .<sup>59</sup> Also, Ho *et al.*’s conductivity measurements report the conductance of KOH > NaOH.<sup>66</sup> Thus, smaller hydrated ionic radii of  $\text{K}^+$  ions contribute to the higher ionic conductivity of the KOH electrolyte than that of NaOH.<sup>67</sup> Further,  $\text{OH}^-$  ions exhibit superior conductivity and mobility compared with  $\text{NO}_3^-$  and  $\text{SO}_4^{2-}$ , contributing to a higher capacitive performance. The hydroxide ion is relatively small, and it forms a less extensive hydration shell when it is hydrated in water, compared with larger, more complex anions like  $\text{NO}_3^-$  and  $\text{SO}_4^{2-}$ . Due to their smaller size and less bulky hydration shell,  $\text{OH}^-$  ions move more easily through the solution, resulting in higher mobility. In contrast, nitrate ions are larger and more symmetrical, forming a more extensive hydration shell because their charge is spread over a larger area, which increases resistance to movement and lowers their mobility compared with  $\text{OH}^-$ . The sulfate ion is even larger and carries a higher charge (−2), causing a very strong interaction with water molecules and creating a highly hydrated, bulky complex. This large hydration shell significantly hinders its mobility and reduces its conductivity. Overall, the electrochemical performance of supercapacitors in aqueous electrolytes is strongly influenced by the



types of cations and anions, ion concentrations, hydrated ion sizes, conductivity, and ion mobility.

## 4. Conclusion

We have successfully synthesized  $\text{Mn}_2\text{O}_3$  using different concentrations of urea by a hydrothermal method. Among the four samples ( $\text{Mn}_2\text{O}_3$ -3, 6, 9, and 12), the  $\text{Mn}_2\text{O}_3$ -9 mM urea sample with relatively higher  $\text{Mn}^{3+}$  content displayed a higher  $C_{\text{sp}}$  in 1 M KOH electrolytes. Consequently, a detailed electrochemical analysis of  $\text{Mn}_2\text{O}_3$ -9 was conducted in different base and neutral electrolytes. The measured  $C_{\text{sp}}$  for  $\text{Mn}_2\text{O}_3$ -9 is higher at  $881 \text{ F g}^{-1}$  and  $1043 \text{ F g}^{-1}$  at a scan rate of  $1 \text{ mV s}^{-1}$  in 1 and 3 M KOH electrolytes, respectively. Similarly, the  $C_{\text{sp}}$  exhibited  $891 \text{ F g}^{-1}$  with an energy density of  $45 \text{ W h kg}^{-1}$  and  $398 \text{ W kg}^{-1}$  power density in 3 M KOH at a  $1 \text{ A g}^{-1}$  current density. The measured  $C_{\text{sp}}$ , energy, and power densities are superior in  $\text{KOH} > \text{NaOH} > \text{NaNO}_3 > \text{Na}_2\text{SO}_4$  electrolytes. The capacitive retention after 5000 cycles was higher in base electrolytes compared with neutral ones. This work introduces innovative strategies to enhance the  $C_{\text{sp}}$ , energy density, and power density of  $\text{Mn}_2\text{O}_3$ -based battery-type electrode materials by optimizing their crystal structure and choice of electrolytes. The findings demonstrate the potential of these materials as promising candidates for large-scale supercapacitor applications. The study conclusively shows that the electrochemical performance of supercapacitors is greatly affected by factors such as the type of electrolytes, including the types of cations and anions, salt concentration, hydrated ion size, conductivity, and ion mobility. These properties play a crucial role in determining ion transport dynamics, charge storage capacity, and the overall efficiency and stability of the device.

## Author contributions

Alisha Dhakal: synthesis, methodology, formal analysis, data curation, and writing – original draft. Dr Felio Perez: XPS measurement and analysis. Dr Sanjay R Mishra: project supervision, conceptualization, review, and editing.

## Data availability

Data for this article, including BET, electrochemical analysis, XPS, and XRD, are available at Zenodo at <https://zenodo.org/uploads/14866457>.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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## References

1. A. Evans, V. Strezov and T. J. Evans, Assessment of utility energy storage options for increased renewable energy penetration, *Renewable Sustainable Energy Rev.*, 2012, **16**, 4141–4147.
2. C. Liu, F. Li, M. Lai-Peng and H. M. Cheng, Advanced Materials for Energy Storage, *Adv. Mater.*, 2010, **22**, E28–E62.
3. J. R. Miller and P. Simon, Electrochemical capacitors for energy management, *Science*, 2008, **321**, 651–652.
4. P. Simon and Y. Gogotsi, Materials for electrochemical capacitors, *Nat. Mater.*, 2008, **7**, 845–854.
5. D. Wei, M. R. J. Scherer, C. Bower, P. Andrew, T. Ryhänen and U. Steiner, A nanostructured electrochromic supercapacitor, *Nano Lett.*, 2012, **12**, 1857–1862.
6. B. Conway, *Electrochemical supercapacitors: scientific fundamentals and technological applications*, Springer Science & Business Media, 2013.
7. H. Chen, T. Cong, W. Yang, C. Tan, Y. Li and Y. Ding, Progress in electrical energy storage system: a critical review, *Prog. Natl. Sci.*, 2009, **19**, 291–312.
8. A. G. Pandolfo and A. F. Hollenkamp, Carbon properties and their role in supercapacitors, *J. Power Sources*, 2006, **157**, 11–247.
9. S. Sarangapani, B. V. Tilak and C. P. Chen, Materials for electrochemical capacitors: theoretical and experimental constraints, *J. Electrochem. Soc.*, 1996, **143**, 3791–3799.
10. A. Dhakal, F. A. Perez and S. R. Mishra, Synergetic effect towards high electrochemical performance in  $\text{LaMnO}_3$ – $\text{Co}_3\text{O}_4$  composites, *Energy Adv.*, 2025, **4**, 162–175.
11. S. Rudra, K. Janani, G. Thamizharasan, M. Pradhan, B. Rani, N. K. Sahu and A. K. Nayak, Fabrication of  $\text{Mn}_3\text{O}_4$ – $\text{WO}_3$  nanoparticles based nanocomposites symmetric supercapacitor device for enhanced energy storage performance under neutral, *Electrochim. Acta*, 2022, **406**, 139870.
12. J. Chmiola, G. Yushin, Y. Gogotsi, C. Portet, P. Simon and P. L. Taberna, Anomalous increase in carbon capacitance at pore sizes less than 1 nanometer, *Science*, 2006, **313**, 1760–1763.
13. R. Kötz and M. Carlen, Principles and applications of electrochemical capacitors, *Electrochim. Acta*, 2000, **45**, 2483–2498.
14. B. E. Conway, V. Birss and J. Wojtowicz, The role and utilization of pseudocapacitance for energy storage by supercapacitors, *J. Power Sources*, 1997, **66**, 1–14.
15. Y. Jiang and J. Liu, Definitions of pseudocapacitive materials: a brief review, *Energy Environm. Mater.*, 2019, **2**, 30–37.
16. J. P. Zheng, P. J. Cygan and T. R. Jow, Hydrous ruthenium oxide as an electrode material for electrochemical capacitors, *J. Electrochem. Soc.*, 1995, **142**, 2699–2703.
17. A. Dhakal, F. A. Perez, S. Karna and S. R. Mishra,  $\text{LaMnO}_3$ – $\text{Mn}_3\text{O}_4$  nanocomposite: Synergetic effect towards high electrochemical performance, *J. Alloys Compd.*, 2024, **1008**, 176262.
18. H. Y. Lee, S. W. Kim and H. Y. Lee, Expansion of active site area and improvement of kinetic reversibility in electrochemical pseudocapacitor electrode, *Electrochim. Solid-State Lett.*, 2001, **4**, A19.



19 A. L. M. Reddy, M. M. Shajumon, S. R. Gowda and P. M. Ajayan, Multisegmented Au–MnO<sub>2</sub>/Carbon Nanotube Hybrid Coaxial Arrays for High-Power Supercapacitor Applications, *J. Phys. Chem. C*, 2010, **114**, 658–663.

20 S. C. Pang, M. A. Anderson and T. W. Chapman, Novel electrode materials for thin-film ultracapacitors: comparison of electrochemical properties of sol-gel-derived and electrodeposited manganese dioxide, *J. Electrochem. Soc.*, 2000, **147**, 444.

21 W. Li, J. Shao, Q. Liu, X. Liu, X. Zhou and J. H. E. Acta, *et al.*, Facile synthesis of porous Mn<sub>2</sub>O<sub>3</sub> nanocubics for high-rate supercapacitors, *Electrochim. Acta*, 2015, **157**, 108–114.

22 Z. Y. Li, M. S. Akhtar, P. T. M. Bui and O. B. Yang, Predominance of two dimensional (2D) Mn<sub>2</sub>O<sub>3</sub> nanowalls thin film for high performance electrochemical supercapacitors, *Chem. Eng. J.*, 2017, **330**, 1240–1247.

23 R. N. Reddy and R. G. Reddy, Synthesis and electrochemical characterization of amorphous MnO<sub>2</sub> electrochemical capacitor electrode material, *J. Power Sources*, 2004, **132**, 315–320.

24 A. Dhakal, F. A. Perez and S. R. Mishra, Electrochemical assessment of tailored Mn<sub>2</sub>O<sub>3</sub> cuboidal hierarchical particles prepared using urea and Piperazine, *Electrochim. Acta*, 2024, **507**, 145169.

25 K. Xu, Nonaqueous liquid electrolytes for lithium-based rechargeable batteries, *Chem. Rev.*, 2004, **104**, 4303–4317.

26 Y. Fang, Y. Huang, S. Zhang, W. Jia, X. Wang, Y. Guo, D. Jia and L. Wang, Synthesis of unique hierarchical mesoporous layered-cube Mn<sub>2</sub>O<sub>3</sub> by dual-solvent for high-capacity anode material of lithium-ion batteries, *Chem. Eng. J.*, 2017, **315**, 583–590.

27 N. F. M. Yusoff, N. H. Idris, M. F. M. Din, S. R. Majid, N. A. Harun and M. M. Rahman, Investigation on the Electrochemical Performances of Mn<sub>2</sub>O<sub>3</sub> as a Potential Anode for Na-Ion Batteries, *Sci. Rep.*, 2020, **10**, 9207.

28 D. Lide, *CRC handbook of chemistry and physics: a ready-reference book of chemical and physical data*, 1995.

29 V. D. Mote, Y. P. Purushotham and B. N. Dole, Williamson-Hall analysis in estimation of lattice strain in nanometer-sized ZnO particles, *J. Theor. Appl. Phys.*, 2012, **6**, 1–8.

30 A. Cihan, P. Zarzycki and Z. Hao, Surface Hydroxylation-Induced Electrostatic Forces Thicken Water Films on Quartz, *Langmuir*, 2024, **40**, 15099–15106.

31 S. Y. Park and A. K. Bera, Maximum entropy autoregressive conditional heteroskedasticity model, *J. Econometrics*, 2009, **150**, 219–230.

32 R. Bardestani, G. S. Patience and S. Kaliaguine, Experimental methods in chemical engineering: specific surface area and pore size distribution measurements—BET, BJH, and DFT, *Can. J. Chem. Eng.*, 2019, **97**, 2781–2791.

33 H. W. Nesbitt and D. Banerjee, Interpretation of XPS Mn(2p) spectra of Mn oxyhydroxides and constraints on the mechanism of MnO<sub>2</sub> precipitation, *Am. Miner.*, 1998, **83**, 305–315.

34 W. Lu, Y. Li, M. Yang, X. Jiang, Y. Zhang and Y. Xing, Construction of Hierarchical Mn<sub>2</sub>O<sub>3</sub>@MnO<sub>2</sub> Core–Shell Nanofibers for Enhanced Performance Supercapacitor Electrodes, *ACS Appl. Energy Mater.*, 2020, **3**, 8190–8197.

35 J. X. Flores-Lasluisa, F. Huerta, D. Cazorla-Amorós and E. Morallon, Manganese oxides/LaMnO<sub>3</sub> perovskite materials and their application in the oxygen reduction reaction, *Energy*, 2022, **247**, 123456.

36 M. A. Langell, C. W. Hutchings, G. A. Carson and M. H. Nassir, High resolution electron energy loss spectroscopy of MnO(100) and oxidized MnO(100), *J. Vac. Sci. Technol., A*, 1996, **3**, 1656–1661.

37 H. B. Li, M. H. Yu, F. X. Wang, P. Liu, Y. Liang and J. Xiao, *et al.*, Amorphous nickel hydroxide nanospheres with ultrahigh capacitance and energy density as electrochemical pseudocapacitor materials, *Nat. Commun.*, 2013, **4**, 1894.

38 L. Fan, L. Tang, H. Gong, Z. Yao and R. Guo, Carbon-nanoparticles encapsulated in hollow nickel oxides for supercapacitor application, *J. Mater. Chem.*, 2012, **22**, 16376–16381.

39 Z. Fang, S. ur Rehman, M. Sun, Y. Yuan, S. Jin and H. Bi, Hybrid NiO–CuO mesoporous nanowire array with abundant oxygen vacancies and a hollow structure as a high-performance asymmetric supercapacitor, *J. Mater. Chem. A*, 2018, **6**, 21131–21142.

40 Q. T. Qu, B. Wang, L. C. Yang, Y. Shi, S. Tian and Y. P. Wu, Study on electrochemical performance of activated carbon in aqueous Li<sub>2</sub>SO<sub>4</sub>, Na<sub>2</sub>SO<sub>4</sub> and K<sub>2</sub>SO<sub>4</sub> electrolytes, *Electrochim. Commun.*, 2008, **10**, 1652–1655.

41 V. Srinivasan and J. W. Weidner, Studies on the capacitance of nickel oxide films: effect of heating temperature and electrolyte concentration, *J. Electrochem. Soc.*, 2000, **147**, 880.

42 P. Y. Chen, A. Adomkevicius, Y. T. Lu, S. C. Lin, Y. H. Tu and C. C. Hu, The ultrahigh-rate performance of alkali ion-pre-intercalated manganese oxides in aqueous Li<sub>2</sub>SO<sub>4</sub>, Na<sub>2</sub>SO<sub>4</sub>, K<sub>2</sub>SO<sub>4</sub> and MgSO<sub>4</sub> electrolytes, *J. Electrochim. Soc.*, 2019, **166**, A1875–A1883.

43 T. Brezesinski, J. Wang, S. H. Tolbert and B. Dunn, Ordered mesoporous  $\alpha$ -MoO<sub>3</sub> with iso-oriented nanocrystalline walls for thin-film pseudocapacitors, *Nat. Mater.*, 2010, **9**, 146–151.

44 R. S. Nicholson, Theory and application of cyclic voltammetry for measurement of electrode reaction kinetics, *Anal. Chem.*, 1965, **37**, 1351–1355.

45 T. C. Liu, W. G. Pell, B. E. Conway and S. L. Roberson, Behavior of molybdenum nitrides as materials for electrochemical capacitors: comparison with ruthenium oxide, *J. Electrochim. Soc.*, 1998, **145**, 1882–1888.

46 H. Wu, X. Wang, L. Jiang, C. Wu, Q. Zhao, X. Liu, B. Hu and L. Yi, The effects of electrolyte on the supercapacitive performance of activated calcium carbide-derived carbon, *J. Power Sources*, 2013, **226**, 202–209.

47 B. Pal, S. Yang, S. Ramesh, V. Thangadurai and R. Jose, Electrolyte selection for supercapacitive devices: a critical review, *Nanoscale Adv.*, 2019, **1**, 3807–3835.

48 Y. U. Jeong and A. Manthiram, Nanocrystalline manganese oxides for electrochemical capacitors with neutral electrolytes, *J. Electrochim. Soc.*, 2002, **149**, A1419.

49 Y. Wang, Y. Song and Y. Xia, Electrochemical capacitors: mechanism, materials, systems, characterization and applications, *Chem. Soc. Rev.*, 2016, **45**, 5925.



50 A. Bard, L. Faulkner and H. White, *Electrochemical methods: fundamentals and applications*, John Wiley & Sons, 2022.

51 Z. Chang, Y. Yang, M. Li, X. Wang and Y. Wu, Green energy storage chemistries based on neutral aqueous electrolytes, *J. Mater. Chem. A*, 2014, **2**, 10739–10755.

52 Y. H. Son, P. T. M. Bui, H. R. Lee, M. S. Akhtar, D. K. Shah and O. B. Yang, A Rapid Synthesis of Mesoporous  $Mn_2O_3$  Nanoparticles for Supercapacitor Applications, *Coatings*, 2019, **9**, 631.

53 X. Chen, L. Li, X. Wang, K. Xie and Y. Wang, Effect of manganese valence on specific capacitance in supercapacitors of manganese oxide microspheres, *Chem. – Eur. J.*, 2021, **27**, 9152–9159.

54 M. S. Javed, H. U. Shah, N. Shaheen, R. Lin, M. Qiu, J. Xie, J. Li, R. Raza, W. Mai and C. Hu, High energy density hybrid supercapacitor based on 3D mesoporous cuboidal  $Mn_2O_3$  and MOF-derived porous carbon polyhedrons, *Electrochim. Acta*, 2018, **282**, 1–9.

55 X. Wang, L. Liu, X. Wang, L. Yi, C. Hu and X. Zhang,  $Mn_2O_3$ /carbon aerogel microbead composites synthesized by in situ coating method for supercapacitors, *Mater. Sci. Eng., B*, 2011, **176**, 1234–1238.

56 M. S. Kolathodi, S. N. H. Rao, T. S. Natarajan and G. Singh, Beaded manganese oxide ( $Mn_2O_3$ ) nanofibers: preparation and application for capacitive energy storage, *J. Mater. Chem. A*, 2016, **4**, 7883–7891.

57 S. Maiti, A. Pramanik and S. Mahanty, Electrochemical energy storage in  $Mn_2O_3$  porous nanobars derived from morphology-conserved transformation of benzenetricarboxylate-bridged metal-organic, *CrystEngComm*, 2016, 1–3.

58 J. Liang, L. T. Bu, W. G. Cao, T. Chen and Y. C. Cao, Facile fabrication of coaxial-cable like  $Mn_2O_3$  nanofiber by electrospinning: application as electrode material for supercapacitor, *J. Taiwan Inst. Chem. Eng.*, 2016, **65**, 584–590.

59 L. L. Zhang, T. Wei, W. Wang and X. S. Zhao, Manganese oxide–carbon composite as supercapacitor electrode materials, *Microporous Mesoporous Mater.*, 2009, **123**, 260–267.

60 M. E. Orazem and B. Tribollet, Electrochemical impedance spectroscopy, *New Jersey*, 2008, **1**, 383–389.

61 B.-Y. Chang and S.-M. Park, Electrochemical impedance spectroscopy, *Ann. Rev. Anal. Chem.*, 2010, **3**, 207–229.

62 J. B. Jorcin, M. E. Orazem, N. Pébère and B. Tribollet, CPE analysis by local electrochemical impedance spectroscopy, *Electrochim. Acta*, 2006, **51**, 1473–1479.

63 J. Zhi, Y. Wang, S. Deng and A. Hu, Study on the relation between pore size and supercapacitance in mesoporous carbon electrodes with silica-supported carbon nanomembranes, *RSC Adv.*, 2014, **4**, 40296–40300.

64 R. Wang, Q. Li, L. Cheng, H. Li, B. Wang, X. S. Zhao and P. Guo, Electrochemical properties of manganese ferrite-based supercapacitors in aqueous electrolyte: the effect of ionic radius, *Colloids Surf., A*, 2014, **457**, 94–99.

65 M. M. Vadiyar, S. C. Bhise, S. K. Patil, S. S. Kolekar, J. Y. Chang and A. V. Ghule, Comparative Study of Individual and Mixed Aqueous Electrolytes with  $ZnFe_2O_4$  Nano-flakes Thin Film as an Electrode for Supercapacitor Application, *ChemistrySelect*, 2016, **1**, 959–966.

66 P. C. Ho, D. A. Palmer and R. H. Wood, Conductivity measurements of dilute aqueous LiOH, NaOH, and KOH solutions to high temperatures and pressures using a flow-through cell, *J. Phys. Chem. B*, 2000, **104**, 12084–12089.

67 K. Fic, A. Platek, J. Piwek and E. Frackowiak, Sustainable materials for electrochemical capacitors, *Mater. Today*, 2018, **21**, 437–454.

