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6, 1152High-performance biphasic Na_xMnO₂ electrodes
for cost-effective and high-power aqueous
sodium batteries and capacitors†Andrii Boichuk,^a Tetiana Boichuk,^a Mahesh Eledath-Changarath,^a
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Aqueous sodium batteries and capacitors offer a low-cost and sustainable alternative to lithium-based energy storage systems, with their performance crucially dependant on the choice of electrode materials. Among the candidates commonly used in sodium-ion devices, various phase modifications of presodiated manganese oxide are considered promising. In this work, we synthesized biphasic (orthorhombic/monoclinic) NaMnO₂ using a cost-effective sol–gel technique and investigated its performance as an electrode material for aqueous sodium electrochemical systems. The performance was evaluated through cyclic voltammetry and galvanostatic charge–discharge measurements. The results demonstrated that NaMnO₂ electrodes were highly suitable for high-power energy devices, exhibiting a specific capacity of 103 mA h g^{−1} and high capacity retention, even under high current conditions (82% capacity retention as the current increased from 1C to 20C). The superior electrochemical performance, especially under high discharge current conditions, was attributed to the optimal combination of different pseudocapacitive mechanisms associated with the biphasic monoclinic-orthorhombic phase structure, which ensured both high capacity and stability during cycling, as well as the morphology of the samples. These results paved the way for the development of high-power, stable, and cost-effective aqueous sodium-ion storage devices.

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Introduction

Despite the dominance of lithium-based energy storage devices, cheaper, more abundant, safer, and efficient sodium-ion (Na) batteries^{1–4} and supercapacitors^{5–7} have garnered increasing attention. One of the most significant challenges in the intense research on these devices is to enhance their specific capacity, power, and energy, aiming to achieve competitive operational performance. This can be accomplished by selecting Faraday-active electrode materials that favor reversible Na ion intercalation.

Layered structures based on manganese oxides have emerged as promising candidates for such applications. For instance, high-voltage supercapacitors based on tetragonal NaMnO₂ have already been developed, exhibiting a calculated capacity of 89.6 F g^{−1} at a scan rate of 5 mV s^{−1} and demonstrating good cycle stability.⁸ In the presence of Na⁺ cations, the electrochemical potential of manganese oxide in electrolytes shifts, allowing for the achievement of higher specific energy storage. Furthermore, Na_xMnO₂ has been shown to exhibit beneficial electrochemical properties, especially in terms of cycle stability, when sodium doping is applied during synthesis.^{9–13}

Two phases of sodiated manganese oxide materials have been reported to date (Fig. 1a): monoclinic (α phase, space group C2/m) and orthorhombic (β phase, space group Pmmn). Additionally, β-NaMnO₂ systems have demonstrated capacity values of approximately 90 mA h g^{−1} at high currents, with a capacity retention of around 89% over the first 100 cycles.¹⁴ These results suggest that the α-phase is particularly effective in providing good rate capability, while the β-phase is more suited for maintaining long cycle life.¹⁴ Consequently, both phases exhibit distinct but complementary electrochemical properties, which can be leveraged through their combination

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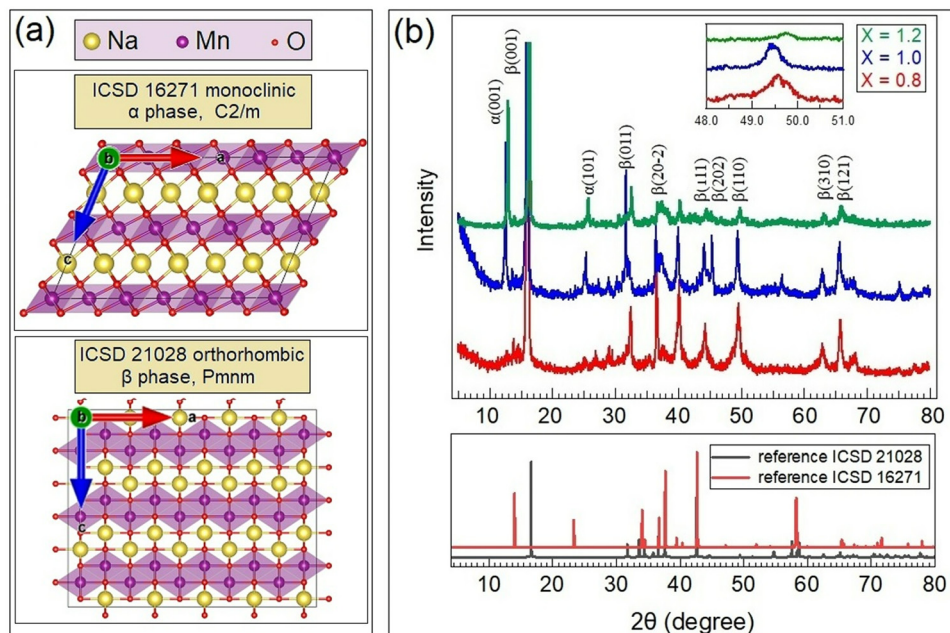


Fig. 1 (a) Orthorhombic and monoclinic phases of NaMnO_2 . (b) XRD patterns of as-synthesized $\text{Na}_{0.8}\text{MnO}_2$ ($x = 0.8$), NaMnO_2 ($x = 1.0$) and $\text{Na}_{1.2}\text{MnO}_2$ ($x = 1.2$).

in a biphasic composite. Such NaMnO_2 polymorphs were observed by Artem M. Abakumov *et al.*¹⁵ as quasi-periodic sequences with varying concentrations of $\alpha\text{-NaMnO}_2$ and $\beta\text{-NaMnO}_2$ phases within each NaMnO_2 layer of the composite. The electrochemical behavior of these biphasic materials has also been investigated.^{16,17} In particular, Debasis Nayak *et al.*¹⁶ reported discharge capacity values of $110.5 \text{ mA h g}^{-1}$ with a capacity retention of about 68% and Coulombic efficiency of 98.9% even after 120 cycles at 0.05C. The biphasic NaMnO_2 compounds described above have been successfully prepared using high-temperature solid-state reactions.^{15–17} However, owing to the lack of control over particle size, coarse-grained powders exhibit a small specific surface area, which hampers electrochemical performance at high currents (as is required for supercapacitors) and can be solved using a bottom-up approach, such as the sol-gel technique.^{18–20} Moreover, electrochemical testing of these micro-sized NaMnO_2 compounds was performed in non-aqueous electrolytes, which may increase the technological complexity of construction and, consequently, the final device cost.

In this work, we overcome these limitations by demonstrating the ability of simple and cheap sol-gel techniques to synthesize biphasic Na_xMnO_2 compounds, which appear to exhibit capacity values as high as 103 mA h g^{-1} at high scan rates with their preservation for 100 cycles in aqueous electrolytes.

Experimental

Synthesis

Several samples of sodium manganese oxide with different amounts of sodium were prepared using the sol-gel method. 0.2 M water solutions of manganese acetate tetrahydrate

$\text{C}_4\text{H}_6\text{O}_4\text{Mn} \cdot 4\text{H}_2\text{O}$ (Thermo Scientific, 99%) and sodium nitrate NaNO_3 (Panreac, 99%) were mixed by stirring for 5 hours. The pH was then adjusted to 8 by adding a 25% ammonia solution. The resulting solution was heated to 80°C and kept at low pressure (200 torr) for 24 h. The obtained product was manually ground, annealed at 750°C for 15 h at a heating rate of 3°C min^{-1} and finally quenched to room temperature. The choice of the final annealing temperature is determined by DTA (Fig. S1, ESI†) and is described under the literature conditions of α and β phase formation.^{21,22} We obtained three samples with different Na amounts and nominal stoichiometric compositions: $\text{Na}_{0.8}\text{MnO}_2$ ($x = 0.8$), NaMnO_2 ($x = 1.0$) and $\text{Na}_{1.2}\text{MnO}_2$ ($x = 1.2$). After cooling to room temperature, the samples were directly transferred to an Ar-filled glove-box and manually ground in an agate mortar.

Characterization methods

X-ray diffraction (XRD) characterization of the samples was carried out using a Bruker D8 ADVANCE A25 diffractometer ($\text{Cu-K}\alpha$ radiation source with a wavelength $\lambda = 1.54 \text{ \AA}$) in the 2θ range $10\text{--}80^\circ$ with a step size of 0.00205° . Rietveld refinement procedures were carried out using FullProf software. The surface morphology of the powders was analyzed by SEM (Hitachi S4800 microscope at 10 kV of acceleration voltage). High-resolution X-ray photoelectron spectroscopy (XPS) measurements were performed using a SPECS GmbH system (base pressure $1.0 \times 10^{-10} \text{ mbar}$) equipped with an ASTRAIOS 190 2D-CMOS hemispherical analyzer. Photoelectrons were excited with the $\text{Al-K}\alpha$ line (1486.7 eV) of a monochromatic X-ray source $\mu\text{-FOCUS 500}$ (SPECS GmbH). Measurements were taken at room temperature with a pass energy of 50 eV. Morphological, compositional and structural analyses of Na_xMnO_2 samples



were also performed by high-resolution transmission electron microscopy (HRTEM) using a field emission gun TECNAI G² F20 microscope operating at 200 kV, with the capabilities of selected area electron diffraction (SAED) and energy dispersive X-ray spectroscopy (EDX) in the facilities of the Servei Central de Suport a la Investigació Experimental (SCSIE) at the University of Valencia. To prepare the TEM samples, the Na_xMnO₂ samples were scratched, removed from the substrate and deposited onto a carbon film supported on a copper grid.

Electrochemical measurement technology

The active mass of NaMnO₂ (80 wt%) was mixed with 15 wt% of carbon black and 5 wt% of PTFE binder and ground in a mortar. An electrode was prepared by repeated brush coating onto nickel foam current collectors. The electrode area was 0.3 cm², and the loading of the active mass was 8 mg independent of the active material and electrolyte. The cyclic voltammetry (CV) tests were performed (Gamry 5000P equipment) using a three-electrode cell with Pt wire and Ag/AgCl electrode as counter and reference electrodes in an aqueous 1 M solution of Na₂SO₄(NS), Na₂CO₃(NC) and NaNO₃(NN) (scan rate region of 0.5–50 mV s^{−1}). The specific capacitance of electrode (*C*) estimated from the cyclic voltammogram was calculated as follows:

$$C = S / (2\Delta U m V),$$

where *S* is the area between the CVA curves, *m* is the mass of the active material, and *V* is the scan rate. The cycleability of electrode materials in different electrolytes was investigated during 100 cycles at a scan rate of 50 mV s^{−1}.

Results and discussion

Structure, morphology and stoichiometry of Na_xMnO₂

The XRD patterns of the as-synthesized and annealed at 750 °C Na_xMnO₂ samples (*x* = 0.8, 1.0, and 1.2) are shown in Fig. 1b. The spectra obtained were compared with the data reported for the monoclinic and orthorhombic NaMnO₂ phases, revealing that the sol-gel samples studied here comprised a combination of β (ICSD 21028) and α (ICSD 16271) polymorphous phases of sodiated manganese dioxide in a proportion depending on the sodium content. Based on Rietveld refinement processes, low Na-content samples, such as the Na_{0.8}MnO₂ ones, show a single β phase structure. Samples *x* = 1.0 and *x* = 1.2 can be described as two-phase systems although stoichiometric NaMnO₂ samples tend to present a higher ratio of the α phase (rough estimation of the ratio of α-phase to β-phase is 69:31 for *x* = 1.0 and 76:24 for *x* = 1.2). The cell parameters of all prepared samples are summarized in Table 1. The orthorhombic phases present parameters compatible with those previously reported.²³ In the case of monoclinic α phase samples, the *c* lattice parameter results are larger than those reported in,^{24,25} which is attributed to the presence of a higher amount of sodium between MnO₂ layers (Fig. 1a). The diffraction peak shift has similar values ($\Delta 2\theta \approx 0.2^\circ$), and it leads to proportional changing of lattice

Table 1 Lattice constant values for the α and β phases obtained through Rietveld refinement

Sample	Phases	<i>a</i> , Å	<i>b</i> , Å	<i>c</i> , Å
Na _{0.8} MnO ₂	β	4.934	2.852	6.091
	α	4.462	2.654	7.023
NaMnO ₂	β	5.004	2.893	6.207
	α	4.457	2.647	7.012
Na _{1.2} MnO ₂	β	4.926	2.880	6.193

parameters for both phases. As shown in,¹⁵ such a correlation of lattice constant can be explained by quasi-periodic modulated sequences using the known α- and β-NaMnO₂ polymorphs as the two limiting cases.

Fig. 2(a)–(c) shows low magnification TEM micrographs of Na_xMnO₂ samples with different Na/Mn ratios together with selected area electron diffraction (SAED) patterns acquired in these samples. As revealed by TEM micrographs, the morphology of the synthesized product is made of relatively large crystals. Diffraction spots observed in the SAED patterns shown at the bottom of Fig. 2(a)–(c) are produced by the small monocrystals randomly orientated, giving an angular distribution of (*hkl*) spots at a distance proportional to 1/*d_{hkl}* from the (000) spot for each allowed reflection. The SAED pattern produced by Na_{0.8}MnO₂ can be indexed only by invoking an orthorhombic phase and is in good agreement with 00-025-0844 in JCPDS files (database). In contrast, the high-resolution TEM images acquired in NaMnO₂ and Na_{1.2}MnO₂ samples evidence the co-existence of both monoclinic and orthorhombic phases, in agreement with that observed by XRD (Fig. 1). In fact, interplanar distances and lattice parameters, as extracted from SAED patterns and HRTEM images of Na_xMnO₂ (see Fig. S2 and Table S1 of the ESI†), are also in good agreement with those obtained by XRD.

SEM micrographs (Fig. 2d, e and f) suggest a high degree of crystallinity, with the presence of layered particles of sodium manganese oxide clearly distinguishable in all cases, while samples with deviations from sodium stoichiometry present more irregular particles with more defect morphology. In fact, samples with *x* = 1.0 tend to be composed of regular hexagons with a surface size of about 1.5 × 1.5 microns. Energy-dispersive spectroscopy (EDS) measurements were performed to map the elemental distribution on the Na_xMnO₂ surface (Fig. S3, ESI†). Because we observed deviation from the homogeneous distribution of Na and Mn elements for non-stoichiometric samples (*x* = 0.8, *x* = 1.2), XPS measurements were performed in all Na_xMnO₂ samples to obtain additional information about their chemical composition and atomic bonding nature.

The XPS spectra of Na 1s, Mn 2p, and O 1s core levels acquired in these samples are depicted in Fig. 3a–c, respectively, which are deconvoluted by assuming Gaussian lineshape components. From this analysis, it appears that the Na 1s XPS spectra (Fig. 3a) are composed of different components with an intensity ratio that depends on the sample stoichiometry. A first component, centered at a binding energy of 1071.7 eV,



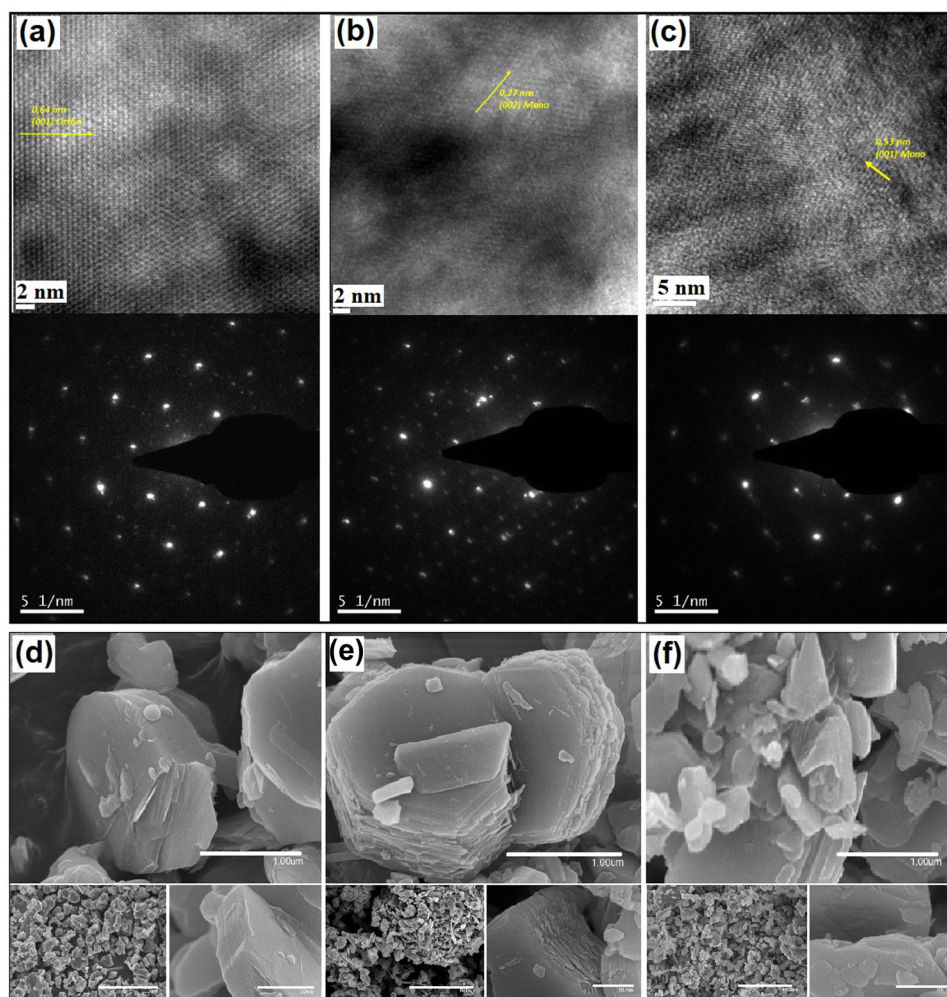


Fig. 2 TEM micrographs with corresponding selected area electron diffraction (SAED) patterns of $\text{Na}_{0.8}\text{MnO}_2$ (a), NaMnO_2 (b), $\text{Na}_{1.2}\text{MnO}_2$ (c). SEM images of $\text{Na}_{0.8}\text{MnO}_2$ (d), NaMnO_2 (e) and $\text{Na}_{1.2}\text{MnO}_2$ (f).

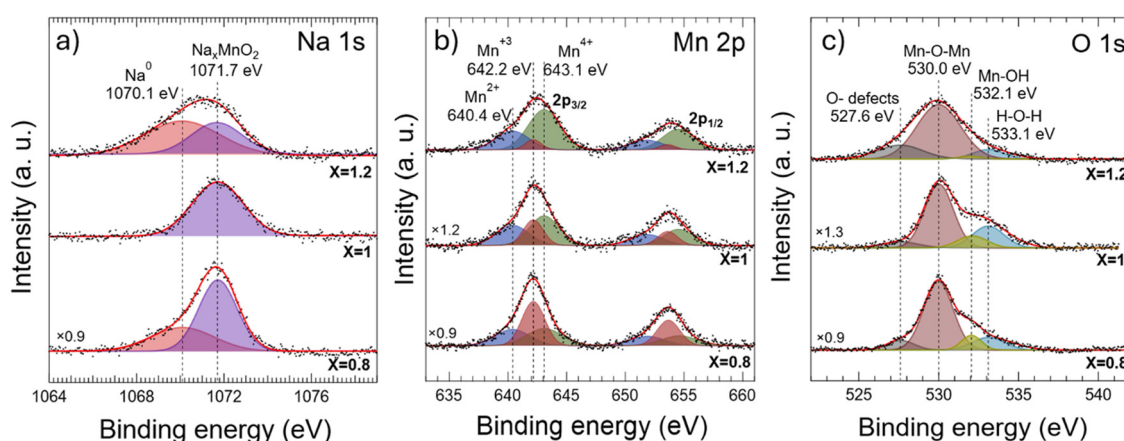


Fig. 3 Na 1s (a), Mn 2p (b) and O 1s (c) core-level spectra acquired by XPS for $\text{Na}_{0.8}\text{MnO}_2$, NaMnO_2 and $\text{Na}_{1.2}\text{MnO}_2$ powders. The fitting curves of the experimental data (red solid lines) were obtained through deconvolution, assuming Gaussian lineshape components. Shadowed Gaussian peaks correspond to the components identified through the deconvolution processes.

is observed in all analyzed powders, which is attributed to Na^+ cations intercalated between MnO_2 layers in orthorhombic

or/and monoclinic Na_xMnO_2 configurations.^{1,26} On the low energy side of the Na 1s spectra, an additional component



centered at 1070.1 eV in binding energies can be resolved in non-stoichiometric compounds Na_xMnO_2 (*i.e.*, $x = 0.8$ and 1.2), attributable to the presence of metallic Na^0 .² In the case of the $\text{Na}_{0.8}\text{MnO}_2$ sample, the formation of metallic sodium can be explained by complications of sodium migration inside the grains at the initial step of synthesis owing to stable orthorhombic phase formation, which causes the appearance of oxygen defects. For the sample $\text{Na}_{1.2}\text{MnO}_2$, the obvious reason is an excess of sodium and its accumulation in the intergranular space. These results show that Na_xMnO_2 powders with $x = 1$ sodium concentration exhibit a higher crystal quality, as observed by SEM measurements (Fig. 2i) and an efficient Na incorporation into the lattice.

Because in-site Mn cations are only surrounded by oxygen anions (in both orthorhombic and monoclinic structures), the observation of any XPS signal attributable to oxidation states different from the Mn^{3+} one expected in stoichiometric NaMnO_2 provides valuable information about the effects of Na incorporation into the host lattice. Deconvolution processes performed on the Mn 2p spectra (Fig. 3b) allow for the resolution of up to three different Mn 2p_{3/2} and Mn 2p_{1/2} spin-orbit doublets whose spin-orbit splitting is ≈ 11.5 eV.^{27–29} These three doublets, whose Mn 2p_{3/2} components are centered at 640.4, 642.2, and 643.1 eV, appear to come from Mn^{2+} , Mn^{3+} , and Mn^{4+} related species, respectively,^{28,30,31} which indicates the presence of MnO and MnO_2 in our samples, coexisting with the Na_xMnO_2 host lattice.³² Moreover, the evolution of the relative weight of each Mn 2p doublet in the Mn 2p XPS signal with the Na content reveals a continuous enhancement of the presence of Mn^{4+} -related species to the detriment of the Mn^{3+} -related ones, as sodium content increases in the Na_xMnO_2 powder, while the presence of Mn^{2+} -related species remain stable.

To obtain more insight into the chemical state of the elements of the compounds studied here, we show in Fig. 3c the O 1s XPS signal acquired in the Na_xMnO_2 powders. Deconvolution of the O 1s spectra (Fig. 3c and Table S2, ESI†) allows the resolution of at least four singlet peaks, which appear at different binding energy (BE) positions: A first singlet related to the presence of oxygen defects (O-defects) and impurities (BE = 527.6 eV),^{33,34} a second one attributable to the host Mn–O–Mn lattice (BE = 530.0 eV),^{27,30,35–37} a third one attributable to manganese hydroxide Mn–OH (BE = 532.1 eV),^{29,35,38} and a fourth one related to adventitious adsorbed H–O–H water (BE = 533.1 eV).^{29,35,36,38} From the XPS results shown in Fig. 3a and c, a clear correlation can be established between the presence of O-defects density, the linewidth of the O 1s peak coming from the Mn–O–Mn lattice, and non-ionic metallic sodium atoms (1070.1 eV) (see also Table S2, ESI†), indicating that NaMnO_2 powders prepared with $x = 1$ Na content exhibit a higher crystal quality (lower defect density and narrower O 1s linewidth of the Mn–O–Mn related peak) than the non-stoichiometric ones, which is in agreement with morphological SEM results reported here (Fig. 2). Moreover, the stoichiometric NaMnO_2 powders present the highest $\text{Na}^+/\text{Mn}^{3+}$ content ratio among the powders prepared (see Fig. S5d of the ESI†), which suggests that optimal

ionic sodium incorporation processes occur in the stoichiometric NaMnO_2 powders.

We calculated Na, Mn and O atomic ratios of the powder's surface composition by considering core-level cross-sections at the used photon energy (Fig. S5d, ESI†). The atomic percentage of total chemical composition (Fig. S5d upper panel ESI†) agrees with those measured by EDS (Fig. S3, ESI†). By considering just the chemical composition of ionic elements bounded in the Na_xMnO_2 powder crystal (Na^+ , Mn^{3+} and O^{2-}), *i.e.*, not including non-ionic elements, amorphous components and defects (see Fig. S5d bottom panel ESI†) show a decreasing amount of Mn^{3+} oxidation state with increasing sodium amount. The calculated atomic $\text{Na}^+/\text{Mn}^{3+}$ ratios depicted in Fig. S5f of the ESI† linearly increase with increasing sodium amounts. It is worth mentioning that the maximal amount of incorporated ionic sodium is found in powders prepared with $x = 1$ sodium concentration (see Fig. S5d bottom panel ESI†).

Electrochemical testing of aqueous electrolytes

As described in the experimental section, Na_xMnO_2 powders of different Na contents are introduced in aqueous electrolytes to analyze their electrochemical properties using the Ag/AgCl reference electrode. Cyclic voltammograms are measured in a voltage range from -0.25 V to 0.8 V. For all samples and electrolytes (Fig. 4), we obtain similar cyclic voltammograms with two parts of pseudocapacitance and corresponding pairs of peaks: intercalation/deintercalation and reversible redox reactions at or near the surface of the electrode material, as described below.

At low scanning rates, a cathodic peak can be noticed at voltage values of 0.3 – 0.35 V irrespective of the powder and electrolyte employed, reflecting an incorporation of Na ions into the Na_xMnO_2 lattice, which is particularly more pronounced at slow voltage changes (Fig. S4, ESI†). Additionally, two peaks can be observed, one cathodic at 0.5 V and the other anodic at 0.6 V, whose intensities increase as the scan rates increase. The small voltage difference between the cathodic and anodic peaks indicates fast redox/oxidation reactions, mostly on the electrode surface. Particularly, there is a marked intensity in these peaks for all samples in Na_2CO_3 electrolyte as well as for samples with non-stoichiometric compositions in other electrolytes probably due to the presence of surface defects (as suggested in XPS and shown in Fig. 3). Additionally, in the particular case of sodium carbonate electrolytes in the three-electrode configuration, CV curves *versus* the current zero line are asymmetric, which implies the low reversibility of intercalation/absorption reactions.

Following one of the main objectives of the present work, the preparation of a stable biphasic structure with optimized morphology to operate under high current conditions with high reversibility, the specific capacity values are calculated based on the results obtained from cyclic voltammograms. Table 2 shows the calculated specific values and their retention after 100 CVA cycles (scan rate of 50 mV s^{-1}) for Na_xMnO_2 samples with $x = 0.8$, 1.0 and 1.2 in three different electrolytes selected. The maximum value (103 mA h g^{-1}) is obtained for NaMnO_2 sample



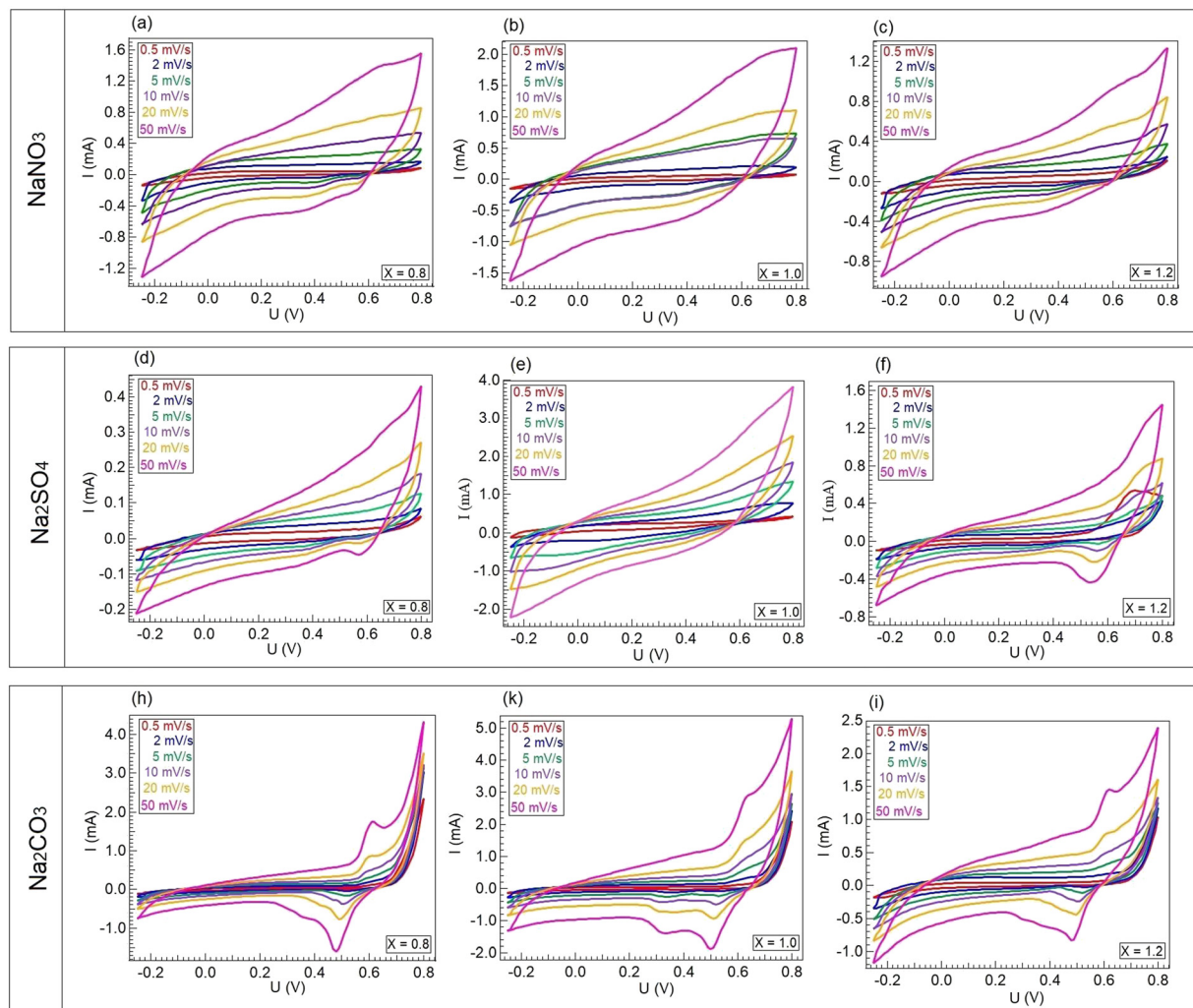


Fig. 4 Cyclic voltammetry (CV) curves of $\text{Na}_{0.8}\text{MnO}_2$ (a), (d) and (g), NaMnO_2 (b), (e) and (h) and $\text{Na}_{1.2}\text{MnO}_2$ (c), (f) and (i) in different electrolytes.

Table 2 CVA-based calculated values of specific capacity and their retention for $\text{Na}_{0.8}\text{MnO}_2$, NaMnO_2 and $\text{Na}_{1.2}\text{MnO}_2$ in different electrolytes

Sample	Electrolyte					
	NaNO_3		Na_2SO_4		Na_2CO_3	
	Capacity, mA h g^{-1}	Capacity retention, %	Capacity, mA h g^{-1}	Capacity retention, %	Capacity, mA h g^{-1}	Capacity retention, %
$\text{Na}_{0.8}\text{MnO}_2$	13	66	63	82	60	72
NaMnO_2	23	89	103	93	61	83
$\text{Na}_{1.2}\text{MnO}_2$	18	70	42	80	39	42

cycling in sodium sulfate-based electrolytes. In addition, the highest capacity retention rate (93 percent after 100 cycles) was found for the NaMnO_2 sample in a sodium sulfate solution.

The different behavior of the electrochemical performance of the devices prepared can be found in the structural and morphological properties of the electrode materials, which bend the balance towards one of the two possible pseudocapacitance mechanisms: intercalation (diffusion-controlled processes) and surface absorption. According to the discharge curves (Fig. 5a–c) obtained for cells based on Na_xMnO_2 in different electrolytes (discharge current 100 mA g^{-1}), charge

accumulation occurs more commonly *via* adsorption, but the presence of a weakly evident horizontal plateau at 0.3 volts (*versus* Ag/AgCl electrode) indicates Faraday reactions. For diffusion-controlled processes, the magnitude of cathodic and anodic currents, which was proportional to the mass transport rate, resulted in an increase in the scan rate. As shown in Fig. 5(d)–(f), we observe linear dependencies between measured based on CVA cathodic and anodic current peaks and the scan rate square, indicating a diffusion-controlled behavior.

Among all samples studied, samples in NaNO_3 electrolyte showed the lowest capacity values (Table 2), which is attributable



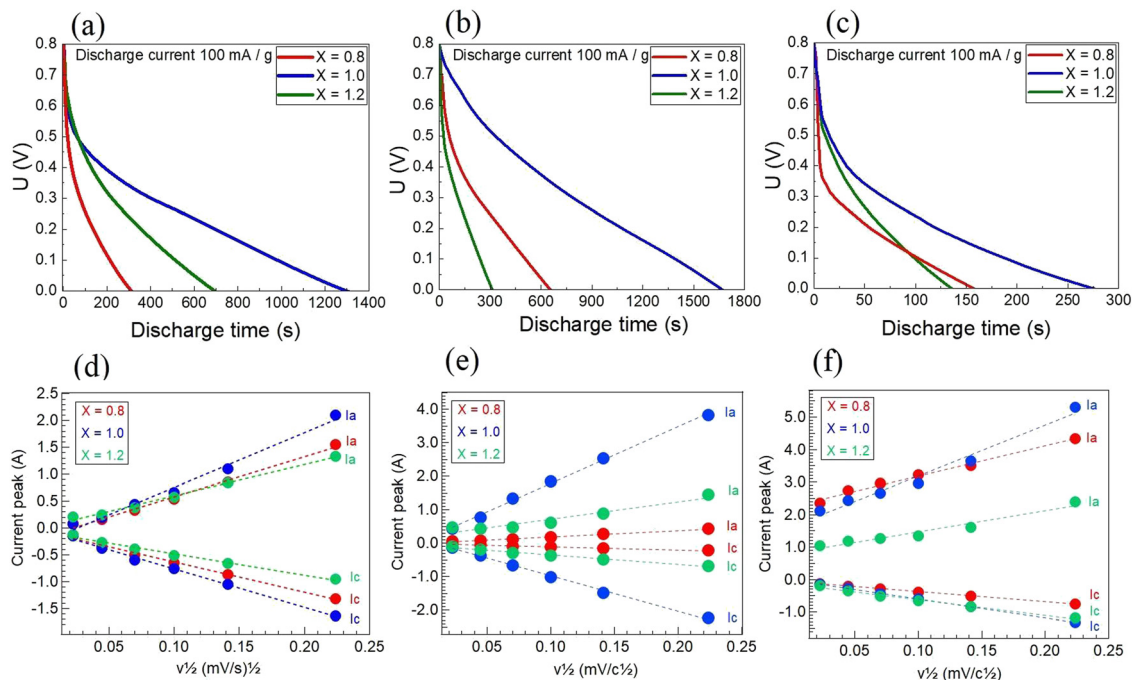


Fig. 5 Discharge curves of Na_xMnO_2 electrode in NaNO_3 (a), Na_2SO_4 (b), Na_2CO_3 (c). The relationship between the cathodic (I_c) and anodic (I_a) current peaks and square root of the scan rate for NaNO_3 (d), Na_2SO_4 (e), and Na_2CO_3 (f).

to the small difference found between anode and cathode current peaks and their low values (from -1 mA to -1.5 mA for cathodic and from 1.2 mA to 2 mA for anodic), even in a high scan rate region (Fig. 5d). Cycling in Na_2CO_3 electrolytes (Fig. 5f) demonstrates low cathodic peak values for all electrode materials employed here, as well as significantly high anodic peak values (up to 5 mA at high scan rates), which demonstrates low Coulombic efficiency and capacity retention. Fig. 5e illustrates that sample cycling in Na_2SO_4 electrolytes reflects a symmetrical behavior whose highest values of cathodic and anodic peaks occur for NaMnO_2 ($x = 1.0$).^{39,40}

Intercalation and absorption pseudocapacity mechanisms monitored by XPS

Typically, upon cycling, the solid-electrolyte interface (SEI) degrades upon unwanted reactions at the electrode/electrolyte interface defects and dislocations, blocking local Na^+ /electrons diffusion.⁴¹ To determine the stability degree of the devices prepared in this work, the XPS technique is employed to determine the effects of pseudocapacitive charge accumulation processes. Hereinafter, we focus on Na_xMnO_2 -based devices exhibiting the highest operation values observed here (*i.e.*, devices prepared with Na_2SO_4 electrolytes).

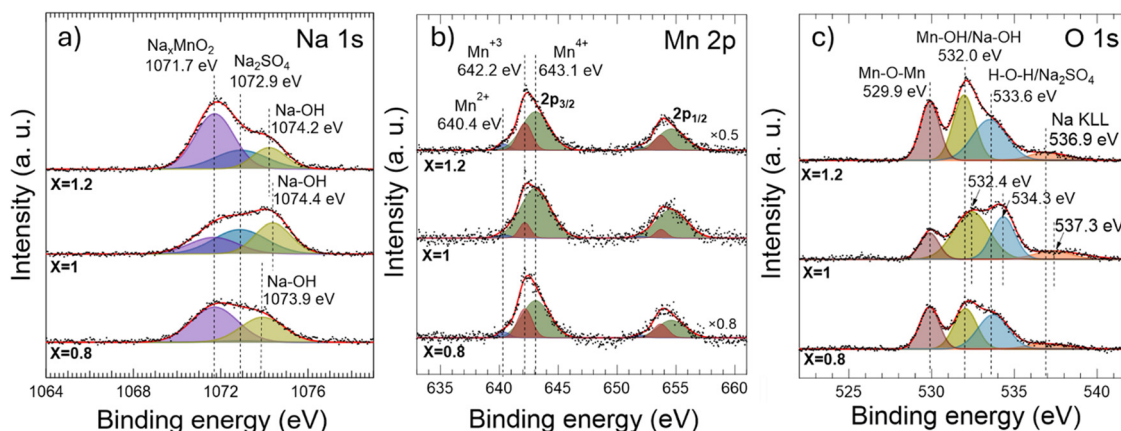


Fig. 6 High-resolution X-ray photon emission spectroscopy (XPS) analysis of $\text{Na}_{0.8}\text{MnO}_2$ ($x = 0.8$), NaMnO_2 ($x = 1.0$) and $\text{Na}_{1.2}\text{MnO}_2$ ($x = 1.2$) electrodes in Na_2SO_4 electrolyte after 100 CVA cycles at 50 mV s^{-1} , with corresponding Na 1s (a), Mn 2p (b) and O 1s (c) core-level spectra. Note the presence of the Na KLL sodium Auger peak (BE = $533.9 \text{ eV} - 537.3 \text{ eV}$)^{26,28,33,38} which was absent in the O 1s spectra of the powders. The detection of this Auger peak in the cycled samples suggests the accumulation of Na atoms at the electrode surface, likely originating from the Na_2SO_4 electrolyte.



Fig. 6 shows XPS spectra acquired in Na_xMnO_2 electrodes ($x = 0.8, 1.0$, and 1.2) after 100 operation cycles at 50 mV s^{-1} in Na_2SO_4 electrolytes. To unravel the chemical effects of cycling on electrodes, acquired XPS spectra are deconvoluted by following procedures similar to those carried out for the powder materials. Comparing the XPS results obtained in the powders (Fig. 3) with those obtained in the cycled electrodes (Fig. 6), it appears that, first, two new Na 1s components develop at the high binding-energy side of the XPS spectra of the cycled electrodes, which is detrimental to the Na^0 -related XPS signal detected in the powders. Note that the powders were not in contact with the electrolytes. However, the samples in Fig. 6 are cycled and, therefore, in contact with the electrolytes, a second source of Na. One of the new Na 1s-related peaks, the one centered at 1072.9 eV , stems from the Na_2SO_4 electrolyte,⁴² while the one centered at $\sim 1074 \text{ eV}$ can be attributed to residual NaOH surface layer formed during cycling.^{38,41} Regarding the XPS Mn 2p core-level spectra measured in the cycled electrodes (Fig. 6b), deconvolution processes performed in these spectra reveal a notable enhancement of the presence of Mn^{4+} species to the detriment of Mn^{2+} ones in the cycled electrodes compared to the powder samples. This can be confirmed by a closer examination of the atomic ratios of manganese oxidation states before and after cycling (see Fig. S5e of the ESI†, bottom panel). A ratio between both Mn oxidized states, $\text{Mn}^{4+}/\text{Mn}^{3+}$, is 2.7 for $\text{Na}_{0.8}\text{MnO}_2$ ($x = 0.8$) and $\text{Na}_{1.2}\text{MnO}_2$ ($x = 1.2$) cycled electrodes and 12.0 for NaMnO_2 ($x = 1$) cycled electrode (Fig. S5e in the ESI†). These facts (*i.e.*, the absence of a metallic Na component after cycling, attributable to its oxidation by interacting with the electrolyte,³⁷ and the relative enhancement of both the Mn^{4+} -like species and NaOH at the surface of the samples) suggest that cycling promotes the accumulation of NaOH at the surface of the samples, partly substituting for Na^+ ions in the starting Na_xMnO_2 powder. Deconvolution of the O 1s core-level XPS spectra acquired in the cycled Na_xMnO_2 electrodes (Fig. 6c) seems to support these observations. Besides the observation of the O 1s XPS signal coming from adsorbed H–O–H water and Na_2SO_4 electrolyte residuals at $\text{BE} \sim 533.9 \pm 0.4 \text{ eV}$,^{29,35,36,38} a notable enhancement of the XPS signal from Mn- and Na-hydroxides^{29,35,38} is produced.

The intercalation of NaOH into the electrode structure after cycling explains the increase in the interlayer distance between the nanosheets, as observed elsewhere.^{8,9} This widening of the interlayer distance explains the improvement in capacity with cycling (Table S2 of the ESI†) as a consequence of the better circulation of electrolyte ions within the crystal. The high capacity and its retention with enhanced circulation properties of NaMnO_2 ($x = 1$) cathode in Na_2SO_4 electrolyte can be attributed to the transformation of $\alpha\text{-NaMnO}_2$ crystal phase into the stable hydrated Na-birnessite phase,^{10,8,9} containing interlayer structural water between separated individual nanosheets without significant change in their initial crystal structure. The interlayer water mediates the interaction between the intercalated cation and the birnessite.^{8,9} This structural modification allows for the enhancement of cation absorption

not only on the electrode planar surface but also between the interlayers of individual electrode nanosheets, thus boosting cathode electrochemical properties.

Sample operation at different currents: results and their representation

The XPS and electrochemical results obtained here for NaMnO_2 -based devices indicate the coexistence of the two types of pseudocapacitance. Our samples can be used under conditions of low operating current values and devices with fast charge–discharge processes. For this purpose, we performed a more detailed study of CVA at different scanning rates and galvanostatic discharges at different currents. In CV measurements, the total current measured under a potential sweep rate can be interpreted as the sum of the current related to the slow diffusion-controlled process (i_d) and the current required to charge the double layer at the electrolyte interface or to initiate fast faradaic reactions on the exposed electrode surface (i_c).³⁶ An empirical description of the charge/discharge processes can be summarized as follows:

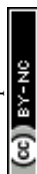
$$i(v) = i_c + i_d = av^b,$$

$$\log i(v) = \log a + b \log(v),$$

where a and b are adjustable parameters. Parameter b is determined from the slope of the linear plot of $\log i(v)$ versus $\log(v)$ and is used to provide kinetic information about the electrochemical reactions. In the case of cycling of the three samples in the Na_2SO_4 electrolyte (Fig. 7d), the calculated b values for the cathodic peaks are within 0.63–0.74 and anodic 0.59–0.69. The smallest among cathodic values (0.63) was observed for NaMnO_2 . This means that for this sample, we have a charge accumulation mechanism closer to the diffusion control than for the other samples. Consequently, we achieve higher values of capacity and discharge time (Fig. 5a–c).

For high power device applications, it is important to have high capacity values and save these values under higher discharge current. Based on the galvanostatic discharge results recorded here (Fig. S6 ESI†), the capacity rate (in percent compared with the capacity at 1C) on the discharge current for all electrolytes is shown in Fig. 7a–c. Even at 20C, cells based on NaMnO_2 demonstrate capacity retention at the level of 84% (NaNO_3) and 82% (Na_2SO_4). From these results, it appears that the best opportunities for application in aqueous sodium devices are biphasic NaMnO_2 with the best stoichiometry, larger lattice constant and optimized morphology based on an increase in interlayer spacing, enhancing cathode performance, especially using these structures as an electrode material for high power devices.

Ragone plots are employed to quantify the energy–power relationship of the electrode material or full electrochemical systems.^{43,44} Usually, based on measurements of charge–discharge characteristics, these plots show the dependency existing between specific power and energy. In our case, comparing the results of CVA and discharge characteristics of samples at different currents, we noted the relationship between CVA



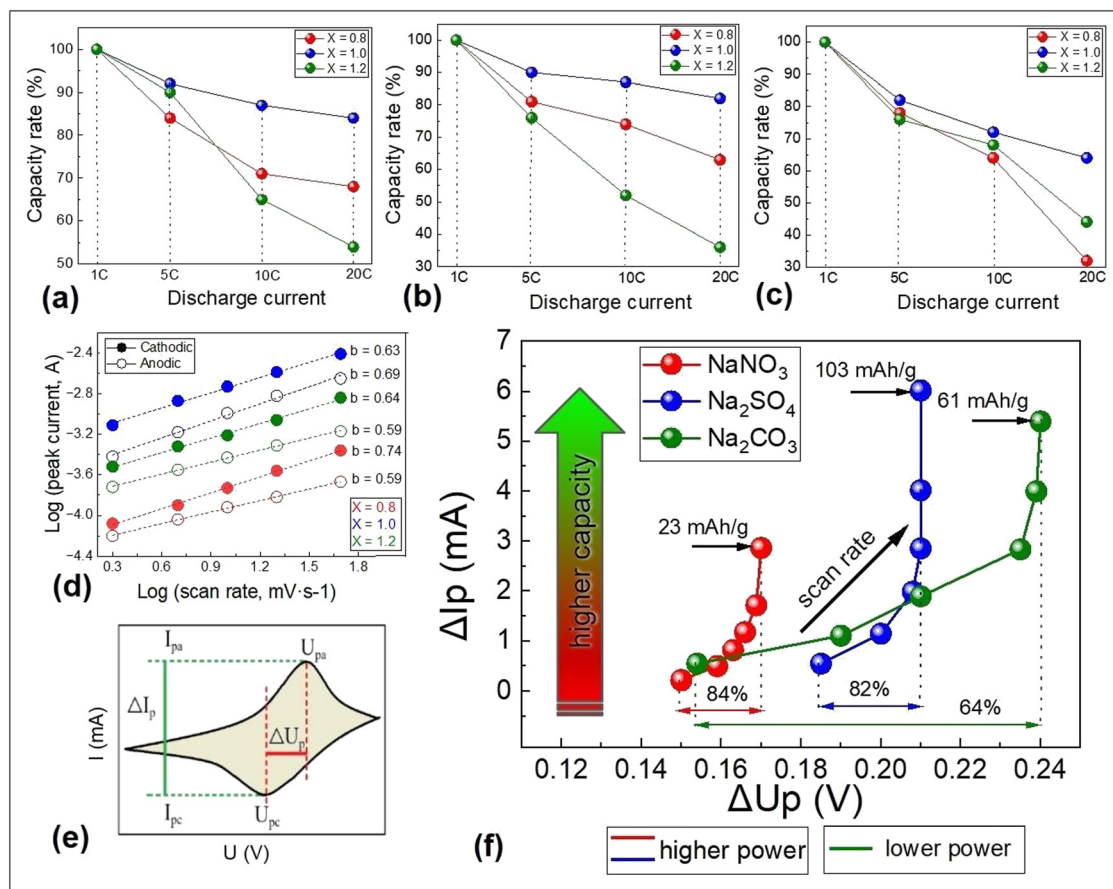


Fig. 7 Capacity rate (in percent compared with capacity at 1C) at different currents (5C, 10C, and 20C) for cells based on Na_xMnO_2 in NaNO_3 (a), Na_2SO_4 (b), and Na_2CO_3 (c). Linear fitting of the relationships between cathodic and anodic peak currents and scan rate for Na_xMnO_2 in Na_2SO_4 electrolyte (d). Dependencies between current peak differences and peak voltage differences (f) for NaMnO_2 at different scan rates (0.5, 2, 5, 10, 20 and 50 mV s^{-1}) and explanation of each elements (e).

curve shapes at different scan rates and discharge curves. This became the basis for our approach to constructing an analogue of the “power-capacitance” diagram based only on cyclic voltammetry, which can quickly provide information on specific capacities and their values under high current conditions. This plot (Fig. 7f) represents the relationship between ΔI_p and ΔU_p that could be taken exclusively from cyclic voltammetry at different scan rates. A detailed explanation of this approach is presented in Fig. S7 (ESI†). ΔI_p is the difference between anodic I_{pa} and cathodic I_{pc} current peaks for each scan rate, and ΔU_p is the difference between cathodic U_{pc} and anodic current peak U_{pa} voltage positions (Fig. 7e).

In the case of a cell based on NaMnO_2 electrode (Fig. 7f), the maximal value of ΔI_p (responsible for transferred charge amount during the cycle) is observed for Na_2SO_4 , and it is completely consistent with the calculated specific capacity based on CVA (Table 2). The ability to work under high currents can be estimated using ΔU_p : systems with better power ability show smaller values of ΔU_p with a gradual increase in the scan rate (For example, NaMnO_2 (Fig. 7a–c) has capacity retention of 84% (NaNO_3), 82% (Na_2SO_4) and 64% (Na_2CO_3) at a current of 20C, which correlates with the change in ΔU_p with increasing

scan rate (Fig. 7f)). It is important to note that this approach does not provide real values of capacity or power but can be successfully used in the case of studying the electrochemical properties of a certain cathode material in different electrolytes or their concentrations, or in symmetrical or asymmetrical systems with different anodes.

Conclusions

In this work, we describe the synthesis of biphasic sodiated manganese oxide using the sol-gel technique. Based on XRD, XPS and TEM, it is shown that it is possible to obtain stoichiometric compounds with both polymorphous monoclinic and orthorhombic phases and a layered structure that is stable during electrochemical sodium insertion and extraction. Based on the CVA in aqueous electrolytes at different scan rates, it has been established that the total capacity is provided mostly by a diffusion-controlled process. Technologically, promising values of the specific capacity (103 mA h g^{-1}) are observed for stoichiometric biphasic systems in Na_2SO_4 electrolytes with high retention during the first 100 cycles (93%). Moreover,



these samples appear to exhibit a high ability to operate at high currents (82% capacity retention when the current increases from 1C to 20C). These results open new opportunities for using these materials in high power, highly stable and cheap aqueous sodium storages due to the possibilities of surface optimization during synthesis.

Author contributions

AB: conceptualisation, investigation, methodology, validation, visualization, writing (original draft preparation), writing (review and editing). TB: conceptualisation, investigation, methodology, validation, visualization, writing (original draft preparation). MEC: investigation, validation, data curation. MK: investigation, data curation, visualization. RA: methodology, validation. PPB: methodology, review and editing. MCA: formal analysis, funding acquisition, review and editing. SA: investigation, data curation, visualization, writing. JFSR: conceptualisation, supervision, funding acquisition, writing (review and editing).

Data availability

All data generated or processed during this study are available from the corresponding author upon reasonable request.

Conflicts of interest

The authors declare no conflict of interest.

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References

- 1 L. Zhao, *et al.*, Engineering of Sodium-Ion Batteries: Opportunities and Challenges, *Engineering*, 2023, **24**, 172–183, DOI: [10.1016/j.eng.2021.08.032](https://doi.org/10.1016/j.eng.2021.08.032).
- 2 N. Yabuuchi, K. Kubota, M. Dahbi and S. Komaba, Research Development on Sodium-Ion Batteries, *Chem. Rev.*, 2014, **114**(23), 11636–11682, DOI: [10.1021/cr500192f](https://doi.org/10.1021/cr500192f).
- 3 R. Usiskin, *et al.*, Fundamentals, status and promise of sodium-based batteries, *Nat. Rev. Mater.*, 2021, **6**(11), 1020–1035, DOI: [10.1038/s41578-021-00324-w](https://doi.org/10.1038/s41578-021-00324-w).
- 4 K. M. Abraham, How Comparable Are Sodium-Ion Batteries to Lithium-Ion Counterparts?, *ACS Energy Lett.*, 2020, **5**(11), 3544–3547, DOI: [10.1021/acsenergylett.0c02181](https://doi.org/10.1021/acsenergylett.0c02181).
- 5 H. Wang, C. Zhu, D. Chao, Q. Yan and H. J. Fan, Nonaqueous Hybrid Lithium-Ion and Sodium-Ion Capacitors, *Adv. Mater.*, 2017, **29**(46), 1702093, DOI: [10.1002/adma.201702093](https://doi.org/10.1002/adma.201702093).
- 6 J. Ding, W. Hu, E. Paek and D. Mitlin, Review of Hybrid Ion Capacitors: From Aqueous to Lithium to Sodium, *Chem. Rev.*, 2018, **118**(14), 6457–6498, DOI: [10.1021/acs.chemrev.8b00116](https://doi.org/10.1021/acs.chemrev.8b00116).
- 7 T. Boichuk, A. Boichuk, V. Vashchynskiy, M. E. Changarath, M. Krečmarová, R. Abargues, J. P. Martínez-Pastor and J. F. Sánchez-Royo, Highly porous activated carbon derived from plant raw material as high-performance anode for aqueous sodium-ion supercapacitors, *J. Power Sources*, 2024, **622**, 235356, DOI: [10.1016/j.jpowsour.2024.235356](https://doi.org/10.1016/j.jpowsour.2024.235356).
- 8 K. S. Kumar, D. Pandey and J. Thomas, High Voltage Asymmetric Supercapacitors Developed by Engineering Electrode Work Functions, *ACS Energy Lett.*, 2021, **6**(10), 3590–3599, DOI: [10.1021/acsenergylett.1c01484](https://doi.org/10.1021/acsenergylett.1c01484).
- 9 M. A. Khan, D. Han, G. Lee, Y.-I. Kim and Y.-M. Kang, P2/O3 phase-integrated Na_{0.7}MnO₂ cathode materials for sodium-ion rechargeable batteries, *J. Alloys Compd.*, 2019, **771**, 987–993, DOI: [10.1016/j.jallcom.2018.09.033](https://doi.org/10.1016/j.jallcom.2018.09.033).
- 10 S. Wang, *et al.*, Enhancing the capacitive deionization performance of NaMnO₂ by interface engineering and redox-reaction, *Environ. Sci.: Nano*, 2019, **6**(8), 2379–2388, DOI: [10.1039/C9EN00545E](https://doi.org/10.1039/C9EN00545E).
- 11 M. Palluzzi, L. Silvestri, A. Celeste, M. Tuccillo, A. Latini and S. Brutti, Structural Degradation of O3-NaMnO₂ Positive Electrodes in Sodium-Ion Batteries, *Crystals*, 2022, **12**(7), 885, DOI: [10.3390/cryst12070885](https://doi.org/10.3390/cryst12070885).
- 12 Q. Ding, W. Zheng, A. Zhao, Y. Zhao, K. Chen, X. Zhou, H. Zhang, Q. Li, X. Ai, H. Yang, Y. Fang and Y. Cao, W-Doping Induced Efficient Tunnel-to-Layered Structure Transformation of Na_{0.44}Mn_{1-x}W_xO₂: Phase Evolution, Sodium-Storage Properties, and Moisture Stability, *Adv. Energy Mater.*, 2023, **13**, 2203802, DOI: [10.1002/aenm.202203802](https://doi.org/10.1002/aenm.202203802).
- 13 Y. Lai, H. Xie, P. Li, B. Li, A. Zhao, L. Luo, Z. Jiang, Y. Fang, S. Chen, X. Ai, D. Xia and Y. Cao, Ion-Migration Mechanism: An Overall Understanding of Anionic Redox Activity in Metal Oxide Cathodes of Li/Na-Ion Batteries, *Adv. Mater.*, 2022, **34**, 2206039, DOI: [10.1002/adma.202206039](https://doi.org/10.1002/adma.202206039).



- 14 J. Billaud, *et al.*, β -NaMnO₂: A High-Performance Cathode for Sodium-Ion Batteries, *J. Am. Chem. Soc.*, 2014, **136**(49), 17243–17248, DOI: [10.1021/ja509704t](https://doi.org/10.1021/ja509704t).
- 15 A. M. Abakumov, A. A. Tsirlin, I. Bakaimi, G. Van Tendeloo and A. Lappas, Multiple Twinning As a Structure Directing Mechanism in Layered Rock-Salt-Type Oxides: NaMnO₂ Polymorphism, Redox Potentials, and Magnetism, *Chem. Mater.*, 2014, **26**(10), 3306–3315, DOI: [10.1021/cm5011696](https://doi.org/10.1021/cm5011696).
- 16 D. Nayak, J. K. Dora, S. Ghosh and V. Adyam, Superior diffusion kinetics and electrochemical properties of α/β -type NaMn_{0.89}Co_{0.11}O₂ as cathode for sodium-ion batteries, *J. Solid State Electrochem.*, 2022, **26**(5), 1231–1239, DOI: [10.1007/s10008-022-05157-3](https://doi.org/10.1007/s10008-022-05157-3).
- 17 D. Nayak, *et al.*, Biphasic α/β -Type NaMn_{0.89}Fe_{0.11}O₂ as a Cathode for Sodium-Ion Batteries: Structural Insight and High-Performance Relation, *ACS Appl. Energy Mater.*, 2022, **5**(1), 116–125, DOI: [10.1021/acsami.1c02576](https://doi.org/10.1021/acsami.1c02576).
- 18 M. Borlaf and R. Moreno, Colloidal sol-gel: A powerful low-temperature aqueous synthesis route of nanosized powders and suspensions, *Open Ceram.*, 2021, **8**, 100200, DOI: [10.1016/j.oceram.2021.100200](https://doi.org/10.1016/j.oceram.2021.100200).
- 19 O. M. Uhorchuk, V. U. Uhorchuk, M. V. Karpets, L. S. Kaykan, B. J. Deputat, A. M. Boychuk and M. I. Hasyuk, Lithium ferrite as the cathode of the electrochemical power sources: the perspectives of sol-gel synthesis method, *J. Nano-Electron. Phys.*, 2015, **7**, 02012[Google Scholar].
- 20 R. Lisovsky, B. Ostafiyshuk, I. Budzulyak, V. Kotsyubynsky, A. Boychuk and B. Rachiy, Nanostructured Iron-Substituted Lithium-Manganese Spinel as an Electrode Material for Hybrid Electrochemical Capacitor, *Acta Phys. Pol. A*, 2018, **133**, 876–878, DOI: [10.12693/APhysPolA.133.876](https://doi.org/10.12693/APhysPolA.133.876).
- 21 L. Zhang, Y. Zhang, Z. Su, H. Tian and L. Wang, Synthesis and Electrochemical Characterization of α -NaMnO_i as a Cathode Material for Hybrid Na/Li-Ion Batteries, *Int. J. Electrochem. Sci.*, 2019, **14**, 2422–2429, DOI: [10.20964/2019.03.82](https://doi.org/10.20964/2019.03.82).
- 22 Q. Zhao, A. Song, S. Ding, R. Qin, Y. Cui, S. Li and F. Pan, Preintercalation Strategy in Manganese Oxides for Electrochemical Energy Storage: Review and Prospects, *Adv. Mater.*, 2020, **32**, 2002450, DOI: [10.1002/adma.202002450](https://doi.org/10.1002/adma.202002450).
- 23 A. A. Nechikott, H. D. Yoo and P. K. Nayak, A mixed dual-ion electrolyte for high-rate performance and remarkable cycling stability of NaMnO₂ in hybrid supercapacitors, *J. Power Sources*, 2024, **591**, 233825, DOI: [10.1016/j.jpowsour.2023.233825](https://doi.org/10.1016/j.jpowsour.2023.233825).
- 24 T. Sato, K. Yoshikawa, W. Zhao, T. Kobayashi, H. B. Rajendra, M. Yonemura and N. Yabuuchi, Efficient Stabilization of Na Storage Reversibility by Ti Integration into O'3-Type NaMnO₂, *Energy Mater. Adv.*, 2021, 2021/9857563, DOI: [10.34133/2021/9857563](https://doi.org/10.34133/2021/9857563).
- 25 Z. Guo, X. Li, Y. Lyu and S. Xu, Improved electrochemical performance of B doped O'3-NaMnO₂ for Na-ion battery, *Electrochim. Acta*, 2022, **430**, 141079, DOI: [10.1016/j.electacta.2022.141079](https://doi.org/10.1016/j.electacta.2022.141079).
- 26 T. Sun, *et al.*, Micron-sized Na_{0.7}MnO_{2.05} as cathode materials for aqueous rechargeable magnesium-ion batteries, *Ionics*, 2019, **25**(10), 4805–4815, DOI: [10.1007/s11581-019-03057-7](https://doi.org/10.1007/s11581-019-03057-7).
- 27 J. Li, *et al.*, Na–Mn–O@C yolk-shell nanorods as an ultra-high electrochemical performance anode for lithium ion batteries, *J. Mater. Chem. A*, 2017, **5**(35), 18509–18517, DOI: [10.1039/C7TA06046G](https://doi.org/10.1039/C7TA06046G).
- 28 Z. Huang, *et al.*, High performance of Mn-Co-Ni-O spinel nanofilms sputtered from acetate precursors, *Sci. Rep.*, 2015, **5**(1), 10899, DOI: [10.1038/srep10899](https://doi.org/10.1038/srep10899).
- 29 Z. Yang, *et al.*, Facile Synthesis of Coaxial CNTs/MnOx-Carbon Hybrid Nanofibers and Their Greatly Enhanced Lithium Storage Performance, *Sci. Rep.*, 2015, **5**(1), 17473, DOI: [10.1038/srep17473](https://doi.org/10.1038/srep17473).
- 30 X. Feng and D. F. Cox, Oxidation of MnO(100) and NaMnO₂ formation: Characterization of Mn²⁺ and Mn³⁺ surfaces via XPS and water TPD, *Surf. Sci.*, 2018, **675**, 47–53, DOI: [10.1016/j.susc.2018.04.022](https://doi.org/10.1016/j.susc.2018.04.022).
- 31 G. Xu, C. Xie, Y. Wen, A. Tang and H. Song, Mn(OH)₂ electrodeposited on secondary porous Ni nano-architecture foam as high-performance electrode for supercapacitors, *Ionics*, 2019, **25**(7), 3287–3298, DOI: [10.1007/s11581-018-2824-8](https://doi.org/10.1007/s11581-018-2824-8).
- 32 D. Delgado, M. Minakshi, D.-J. Kim and C. Kyeong, Influence of the Oxide Content in the Catalytic Power of RANEY[®] Nickel in Hydrogen Generation, *Anal. Lett.*, 2017, **50**(15), 2386–2401, DOI: [10.1080/00032719.2017.1300806](https://doi.org/10.1080/00032719.2017.1300806).
- 33 W. Wei, *et al.*, Subsurface oxygen defects electronically interacting with active sites on In₂O₃ for enhanced photo-thermocatalytic CO₂ reduction, *Nat. Commun.*, 2022, **13**(1), 3199, DOI: [10.1038/s41467-022-30958-5](https://doi.org/10.1038/s41467-022-30958-5).
- 34 M. Han, *et al.*, Oxygen Defects in β -MnO₂ Enabling High-Performance Rechargeable Aqueous Zinc/Manganese Dioxide Battery, *iScience*, 2020, **23**(1), 100797, DOI: [10.1016/j.isci.2019.100797](https://doi.org/10.1016/j.isci.2019.100797).
- 35 J. D. A. Pereira, *et al.*, Tuning the morphology of manganese oxide nanostructures for obtaining both high gravimetric and volumetric capacitance, *Mater. Adv.*, 2020, **1**(7), 2433–2442, DOI: [10.1039/D0MA00524J](https://doi.org/10.1039/D0MA00524J).
- 36 G. Kucinskis, *et al.*, Enhanced Electrochemical Properties of Na_{0.67}MnO₂ Cathode for Na-Ion Batteries Prepared with Novel Tetrabutylammonium Alginate Binder, *Batteries*, 2022, **8**(1), 6, DOI: [10.3390/batteries8010006](https://doi.org/10.3390/batteries8010006).
- 37 X. Feng and D. F. Cox, Synthesis of a planar, multicomponent catalytic surface of Na₂CO₃/MnO, *Surf. Sci.*, 2021, **707**, 121807, DOI: [10.1016/j.susc.2021.121807](https://doi.org/10.1016/j.susc.2021.121807).
- 38 F. Xiao and Y. Xu, Pulse Electrodeposition of Manganese Oxide for High-Rate Capability Supercapacitors, *Int. J. Electrochem. Sci.*, 2012, **7**(8), 7440–7450, DOI: [10.1016/S1452-3981\(23\)15795-6](https://doi.org/10.1016/S1452-3981(23)15795-6).
- 39 V. Hiremath, M. Cho and J. G. Seo, Self-assembled Mn₃O₄ nano-clusters over carbon nanotube threads with enhanced supercapacitor performance, *New J. Chem.*, 2018, **42**(24), 19608–19614, DOI: [10.1039/C8NJ04850A](https://doi.org/10.1039/C8NJ04850A).
- 40 J. Liu, *et al.*, Advanced Energy Storage Devices: Basic Principles, Analytical Methods, and Rational Materials Design, *Adv. Sci.*, 2018, **5**(1), 1700322, DOI: [10.1002/advsc.201700322](https://doi.org/10.1002/advsc.201700322).



- 41 Y.-F. Liu, *et al.*, Layered oxide cathodes for sodium-ion batteries: From air stability, interface chemistry to phase transition, *InfoMat*, 2023, 5(6), e12422, DOI: [10.1002/inf2.12422](https://doi.org/10.1002/inf2.12422).
- 42 N. Zhao, C. Li and X. Guo, Long-life Na-O₂ batteries with high energy efficiency enabled by electrochemically splitting NaO₂ at a low overpotential, *Phys. Chem. Chem. Phys.*, 2014, 16(29), 15646, DOI: [10.1039/c4cp01961j](https://doi.org/10.1039/c4cp01961j).
- 43 T. Christen and C. Ohler, Optimizing energy storage devices using Ragone plots, *J. Power Sources*, 2002, 110(1), 107–116, DOI: [10.1016/S0378-7753\(02\)00228-8](https://doi.org/10.1016/S0378-7753(02)00228-8).
- 44 I. Beyers, A. Bensmann and R. Hanke-Rauschenbach, Ragone plots revisited: A review of methodology and application across energy storage technologies, *J. Energy Storage*, 2023, 73, 109097, DOI: [10.1016/j.est.2023.109097](https://doi.org/10.1016/j.est.2023.109097)-Part D.

