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Synthesis and performance evaluation of nanostructured $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ cathode materials in sodium ion batteries (SIBs)

Umair Nisar,^a ^{†a} Mona Hersi Gulied,^{†b} R. A. Shakoor,^c ^{*a} Rachid Essehli,^c Zubair Ahmad,^a Abdullah Alashraf,^a Ramazan Kahraman,^b Siham Al-Qaradawi^d ^d and Ahmed Soliman^e

This research work focuses on the synthesis and performance evaluation of $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($X = 0, 0.8$ and 1.0) cathode materials in sodium ion batteries (SIBs). The novel materials having a primary particle size of around 100–200 nm were synthesized through a sol–gel process by reacting stoichiometric amounts of the precursor materials. The structural analysis confirms the formation of crystalline, phase pure materials that adopt a monoclinic crystal structure. Thermal analysis indicates the superior thermal stability of $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ when compared to $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$. Galvanostatic charge/discharge analysis indicates that the intercalation/de-intercalation of a sodium ion (Na^+) into/from $\text{NaFe}(\text{SO}_4)_2$ ensues at about 3.2 V due to the $\text{Fe}^{2+}/\text{Fe}^{3+}$ active redox couple. Moreover, *ex situ* XRD analysis confirms that the insertion/de-insertion of sodium into/from the host structure during charging/discharging is accompanied by a reversible single-phase reaction rather than a biphasic reaction. A similar sodium intercalation/de-intercalation mechanism has been noticed in $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ which has not been reported earlier. The galvanostatic measurements and X-ray photoelectron spectroscopy (XPS) analysis confirm that the $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple is inactive in $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($X = 0, 0.8$) and thus does not contribute to capacity augmentation. However, suitable carbon coating may lead to activation of the $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple in these inactive materials.

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1 Introduction

Due to the technological advancements in portable electronics, the demand for secondary batteries has increased considerably in the last decade.^{1,2} To date, the lithium ion battery system is considered one of the best candidates for portable energy storage applications due to its high-energy and power density.^{2–5} However, due to rapid technological advancements, the demand for high energy, high power, safer and cheaper batteries is increasing at a great pace.^{1,6–8} Therefore, new battery systems with promising chemistries are also being explored as an alternative to lithium ion batteries.^{9–11}

In the last decade, much research was devoted to the development of sodium ion batteries (SIBs) due to their similar

electrochemical mechanism to lithium ion batteries. Also, the abundance of sodium resources in nature makes this technology much cheaper as compared to lithium ion batteries.^{12–14} Unfortunately, due to the larger size of sodium ion (Na^+) as compared to lithium (Li^+), it is difficult to identify new compounds capable of reversibly intercalating Na^+ at fast rates. Recently considerable efforts have been devoted to develop novel potential cathode materials for Na-ion batteries.^{14,15} Towards this direction, several cathode materials have been identified such as $\text{Na}_3\text{V}_2(\text{PO}_4)_3$, $\text{NaTi}_2(\text{PO}_4)_3$, $\text{Na}_4\text{Fe}_3(\text{PO}_4)_2(\text{P}_2\text{O}_7)$, NaNiO_2 , NaMnO_2 , NaFePO_4 , $\text{Na}_2\text{FeP}_2\text{O}_7$, $\text{Na}_3\text{V}_2(\text{PO}_4)_2\text{F}_3$, $\text{Na}_4\text{Co}_3(\text{PO}_4)_2\text{P}_2\text{O}_7$ demonstrating promising energy storage performance.^{16–24} However, to address the future needs, development of cheaper, safe and high-performance cathode materials is still challenging.^{14,15,25}

Recently, $\text{NaFe}(\text{SO}_4)_2$ has been reported having promising electrochemical performance. This material intercalates/de-intercalates sodium at (3.2 V) demonstrating charge/discharge capacity (80 mA h g^{-1}) with good rate capability and cyclability.¹² Being impressed with the electrochemical performance of $\text{NaFe}(\text{SO}_4)_2$ and promising merits of chromium (Cr) such as (i) reasonable high voltage and (ii) structural stability, we decided to synthesize an offshoot of $\text{NaFe}(\text{SO}_4)_2$ by substituting iron (Fe) with chromium (Cr). In this study, we report the synthesis,

^aCenter for Advanced Materials (CAM), Qatar University, P. O. Box 2713, Doha, Qatar.
E-mail: shakoor@qu.edu.qa

^bDepartment of Chemical Engineering, College of Engineering, Qatar University, P. O. Box 2713, Doha, Qatar

^cQatar Environment and Energy Research Institute, Hamad Bin Khalifa University, Qatar Foundation, 5825, Doha, Qatar

^dDepartment of Chemistry & Earth Sciences, College of Arts and Science, Qatar University, P. O. Box 2713, Doha, Qatar

^eGas Processing Center (GPC), Qatar University, P. O. Box 2713, Doha, Qatar

[†] Authors with equal contributions.

structural, thermal properties and electrochemical performance of $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$, where ($x = 0, 0.8$ and 1.0). To the best of our knowledge, this is a comprehensive report which describes the sodium insertion/de-insertion mechanism and structural alteration in $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ during the charging/discharging process.

2 Experimental

2.1 Materials preparation

$\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ where ($x = 0, 0.8, 1.0$) was synthesized by a sol-gel process using sodium nitrate, iron nitrate nonahydrate, chromium nitrate and ammonium sulfate (Sigma Aldrich) in stoichiometric ratios. Firstly, all the precursors were dissolved in 100 ml of distilled water with continuous stirring at around 50°C . After 2 hours of continuous stirring, the temperature was raised to 70°C and the solution was evaporated until a clear transparent gel was formed. The resulting gel was dried in a conventional oven at around 120°C for 6 hours. Later, the gel was homogeneously ground into fine powder and then calcined at 420°C for 6 hours in air to form pure $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($x = 0, 0.8, 1.0$). During the calcination process, the heating/cooling rates were set at 5°C per minutes to ensure the formation of phase pure crystalline materials. A schematic diagram of the synthesis process is presented in Fig. 1.

2.2 Structural characterization

Powdered XRD (Panalytical diffractometer) using $\text{Cu-K}\alpha$ radiations was used to identify the phase purity and structural details of the synthesized materials. The data were collected over the 2θ angular range of 10 – 90° with a step size of 0.01° . The morphology (particle size, shape, distribution) and elemental mapping of the synthesized materials were undertaken using scanning electron microscopy (SEM, NOVA NANOSEM 450). Thermal stability of the synthesized materials was studied using thermogravimetric analysis (Perkin Elmer, TGA 4000) from room temperature to 500°C at the heating rate of 10°C in a N_2 atmosphere. Fourier-transform infrared spectroscopy (Perkin Elmer Frontier FT-IR) was further used to confirm the phase purity using KBr pellet method in the range of 500 – 2500 cm^{-1} wave numbers. The activity of redox couples was studied through X-ray Photoelectron spectroscopy (XPS) (Thermo-Scientific-Sigma Probe).

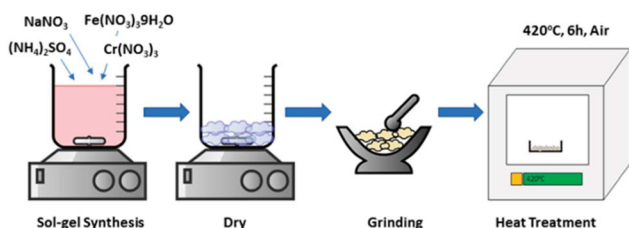


Fig. 1 Schematic process for sol-gel synthesis of $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($x = 0, 0.8, 1.0$).

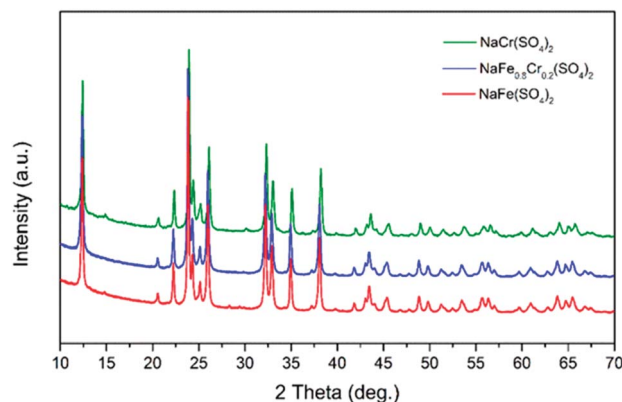


Fig. 2 XRD patterns of pristine $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$.

2.3 Electrochemical characterization

The cathodes were fabricated by forming a slurry composed of 70 wt% active material $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$, 20 wt% conductive carbon (super-P) and 10.0 wt% polyvinylidene fluoride binder in 1-methyl-2-pyrrolidone (NMP). The slurry was homogeneously mixed for 10 hours and was later cast on an aluminum foil using a doctor blade. The cast electrode sheet was dried in a conventional oven for 6 hours to remove NMP. The dried cathode sheet was then rolled using a rolling machine for good compaction and then was dried in a vacuum oven for 12 hours to remove the last traces of water. The electrodes of 14.0 mm diameter were punched from vacuum dried cathode sheet and later shifted to the glove box for cell fabrication. Electrochemical measurements were performed by fabricating 2032-type coin cells in an Ar-filled glove box. Sodium metal was used as the negative electrode. The electrolyte was composed of 1 M NaClO_4 dissolved in ethylene carbonate (EC) and 2.0 wt% of fluoroethylene carbonate (FEC). Galvanostatic charge/discharge tests were performed using WonAtech battery cycler (WBSC 3000L, Korea) in the voltage range of 1.5–4.5 V at room temperature.

3 Results and discussion

The structural properties and phase purity of the synthesized phases was studied through XRD analysis. The XRD patterns of pristine $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$ are presented in Fig. 2. It can be noticed that the synthesized materials are crystalline, and the absence of any impurity peak (s) confirm the high purity of the developed materials.

Table 1 The calculated crystallite size for the developed $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$

Identity	2θ	K	λ	β	L (Å)
$\text{NaFe}(\text{SO}_4)_2$	23.8941	1.0747	1.547	0.1663	0.2138
$\text{NaCr}(\text{SO}_4)_2$	23.9995	1.0747	1.547	0.1663	0.2129
$\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$	23.9046	1.0747	1.547	0.1919	0.1852



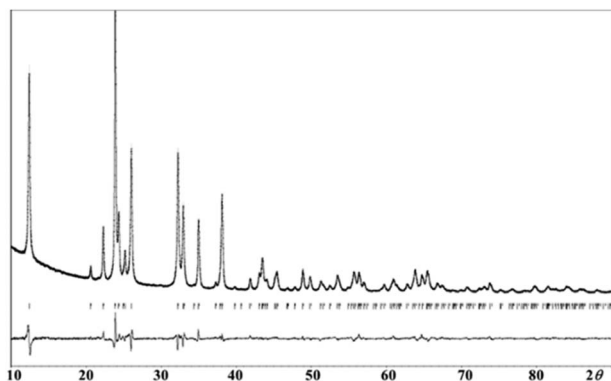


Fig. 3 Final observed, calculated, and difference plots for PXRD (Cu-K α radiation) Rietveld refinement of NaFe_{0.8}Cr_{0.2}(SO₄)₂. [R_p = 0.040, R_{wp} = 0.052, R_{exp} = 0.022].

The XRD data was used to calculate the crystallite size (L) using the well-known Scherrer equation. The calculated crystallite size for the developed NaFe(SO₄)₂, NaFe_{0.8}Cr_{0.2}(SO₄)₂ and NaCr(SO₄)₂ is presented in Table 1.

In order to have more insight of the structural details of the synthesized phases, Rietveld refinement was carried out. Our analysis confirms that NaFe_xCr_{1-x}(SO₄)₂ (x = 0, 0.8 and 1.0) adopts a monoclinic crystal structure. As an example, Fig. 3 shows a Rietveld refinement of pristine NaFe_xCr_{1-x}(SO₄)₂ where (x = 0.8) which shows good agreement between the experimental and calculated patterns.

The refined lattice parameters and other crystallographic information of NaFe_{0.8}Cr_{0.2}(SO₄)₂ is presented in Table 2. Moreover, atomic coordinates and isotropic displacement parameters for NaFe_{0.8}Cr_{0.2}(SO₄)₂ are also tabulated in Table 3.

The crystal structure for NaFe_{0.8}Cr_{0.2}(SO₄)₂ is shown in Fig. 4. As can be seen in this structure, the sodium (Na), iron (Fe) and chromium (Cr) ions occupy the octahedral sites in the a - b plane which are bridged together by (SO₄)⁻ polyanions and creates an interplanar space for 2D Na⁺ diffusion.

Fig. 5(a-c) shows the morphology of NaFe(SO₄)₂, NaCr(SO₄)₂ and NaFe_{0.8}Cr_{0.2}(SO₄)₂ respectively. It can be clearly seen that the synthesized materials have irregular particle morphology containing primary particle size between 100–200 nm. The particle agglomeration can also be clearly noticed in NaCr(SO₄)₂ and NaFe_{0.8}Cr_{0.2}(SO₄)₂. Fig. 5(d-f) shows the elemental mapping for NaFe_{0.8}Cr_{0.2}(SO₄)₂. It can be clearly seen from

Table 2 Crystallographic and structure refinements data for NaFe_{0.8}Cr_{0.2}(SO₄)₂

Sr. #	Description	Crystallographic
1	Chemical formula	NaFe _{0.8} Cr _{0.2} (SO ₄) ₂
2	Crystal system, space group	Monocline, $C2/m$
3	Lattice parameter (a)	7.98494 (Å)
4	Lattice parameter (b)	5.12776 (Å)
5	Lattice parameter (c)	7.13229 (Å)
6	β	92.2166 (°)
7	Volume of unit cell	291.81 (Å ³)

Table 3 Atomic coordinates and isotropic displacement parameters (in Å²) for NaFe_{0.8}Cr_{0.2}(SO₄)₂

Atom	Wyck.	Occ.	X	Y	Z	U_{iso}
Na1	2c	1	0	0	1/2	0.0219
Fe1	2a	0.8	0	0	0	0.0119
Cr1	2a	0.2	0	0	0	0.0119
S1	4i	1	0.36107	0	0.22128	0.0096
O1	8j	1	0.46999	0.23434	0.20424	0.0090
O2	4i	1	0.24124	0	0.06535	0.0032
O3	4i	1	0.29052	0	0.40408	0.0080

Fig. 5(e and f) that iron and chromium are homogeneously distributed throughout the synthesized materials.

To investigate the thermal stability of NaFe(SO₄)₂, NaFe_{0.8}Cr_{0.2}(SO₄)₂ and NaCr(SO₄)₂, thermo-gravimetric analysis (TGA) was performed from room temperature to 500 °C in N₂ atmosphere (Fig. 6). The NaFe(SO₄)₂ and NaCr(SO₄)₂ retains about 98.58 and 97.98% of their original weights up to 500 °C respectively. On the other hand, NaFe_{0.8}Cr_{0.2}(SO₄)₂ retains around 99.21% of its original weight. The results clearly indicate that the NaFe(SO₄)₂ and NaCr(SO₄)₂ materials have inferior thermal stability as compared to NaFe_{0.8}Cr_{0.2}(SO₄)₂. The inferior thermal stability of NaFe(SO₄)₂ and NaCr(SO₄)₂ as compared to NaFe_{0.8}Cr_{0.2}(SO₄)₂ indicates their inferior structural stability with increasing temperature. Furthermore, small substitution of iron with chromium to form NaFe_{0.8}Cr_{0.2}(SO₄)₂ improves the thermal stability of the material as indicated by weight loss (99.21%). The better thermal stability of chromium doped NaFe_{0.8}Cr_{0.2}(SO₄)₂ may be attributed to stronger bonding between iron (Fe) and chromium (Cr) in the crystal structure. It has been reported earlier that chromium doping enhances the thermal stability of the materials which is due to the high excess stabilization energy of Cr³⁺ in the structure.^{26–28} The stabilized structure therefore, is also expected to improve the electrochemical stability of the material specially at higher temperature.

Fig. 7 shows the FTIR spectra of NaFe(SO₄)₂, NaFe_{0.8}Cr_{0.2}(SO₄)₂ and NaCr(SO₄)₂ recorded between 2500 and

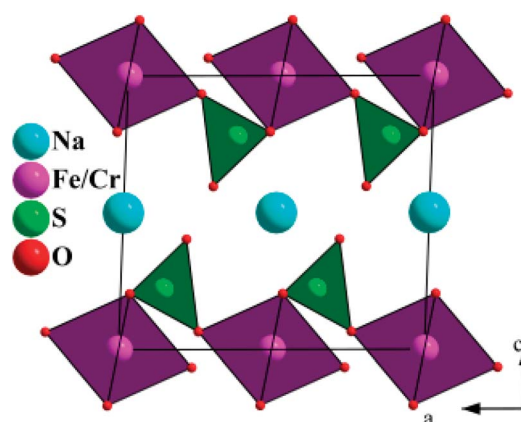


Fig. 4 The crystal structure of NaFe_{0.8}Cr_{0.2}(SO₄)₂ projected along [010].



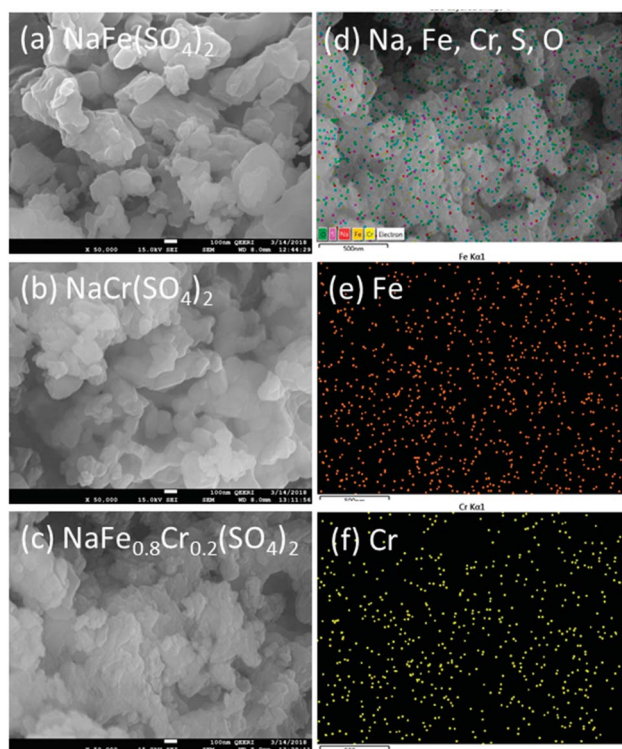


Fig. 5 SEM images for (a) $\text{NaFe}(\text{SO}_4)_2$, (b) $\text{NaCr}(\text{SO}_4)_2$ and (c) $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$, (d–f) elemental mapping for $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ for Fe and Cr.

500 cm^{-1} . Four main internal modes are observed, which can be described as (i) the symmetric stretching mode ν_1 , (ii) the symmetric bending mode ν_2 , (iii) the asymmetric stretching mode ν_3 , and (iv) the asymmetric bending mode ν_4 . As displayed in Fig. 7 the split of asymmetric stretching (ν_3 : 997 cm^{-1}) and bending (ν_4 : 688 cm^{-1}) modes at a low wavenumber can be ascribed to the peaks for the sulfate group which confirms the formation of sulfate-based materials. These findings are consistent with previous studies.^{29–31}

To confirm the electrochemical activity of $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$, differential capacity vs. voltage was plotted as shown in Fig. 8. It can be clearly seen from that $\text{NaFe}(\text{SO}_4)_2$ shows a peak at around 3.38 and 3.124 V

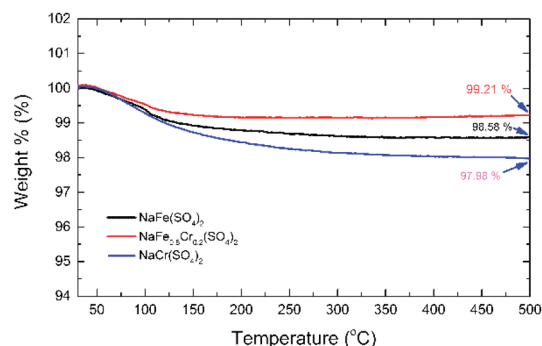


Fig. 6 TGA data for $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$.

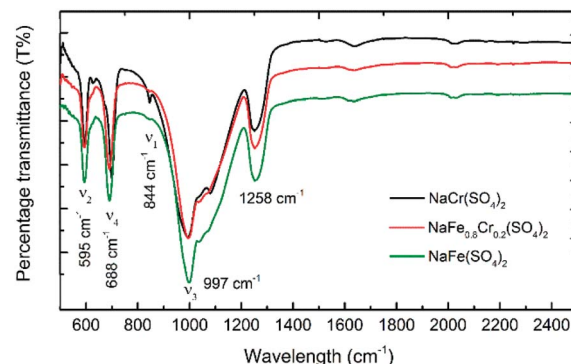


Fig. 7 FTIR spectra for $\text{NaFe}(\text{SO}_4)_2$, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and $\text{NaCr}(\text{SO}_4)_2$ for the range $500\text{--}2500\text{ cm}^{-1}$.

during oxidation and reduction respectively which corresponds to active $\text{Fe}^{2+}/\text{Fe}^{3+}$ redox couple. On the other hand, $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ shows peaks at around 3.169 and 3.005 V during oxidation and reduction which also corresponds to $\text{Fe}^{2+}/\text{Fe}^{3+}$ redox active couple. $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ demonstrates lesser polarization during charging and discharging whereas $\text{NaFe}(\text{SO}_4)_2$ experiences more polarization as evident in dQ/dV curves in Fig. 8. It can be noticed that the substitution of iron with chromium reduces the redox potential of $\text{Fe}^{2+}/\text{Fe}^{3+}$ which may be attributed to multicomponent effect as reported previously.^{29–31} Finally, $\text{NaCr}(\text{SO}_4)_2$ shows no oxidation and reduction peaks which clearly indicated the inactive behaviour of $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple in $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($x = 0, 0.8$).

Fig. 9(a–d) shows the electrochemical analysis of $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ under different C-rates. It is pertinent to

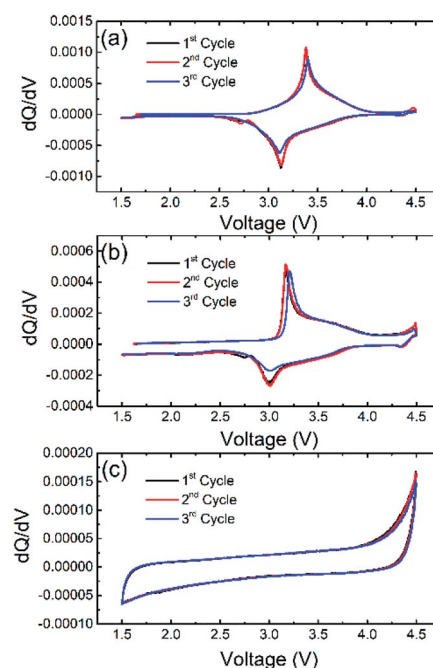


Fig. 8 dQ/dV curves for (a) $\text{NaFe}(\text{SO}_4)_2$, (b) $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ and (c) $\text{NaCr}(\text{SO}_4)_2$.



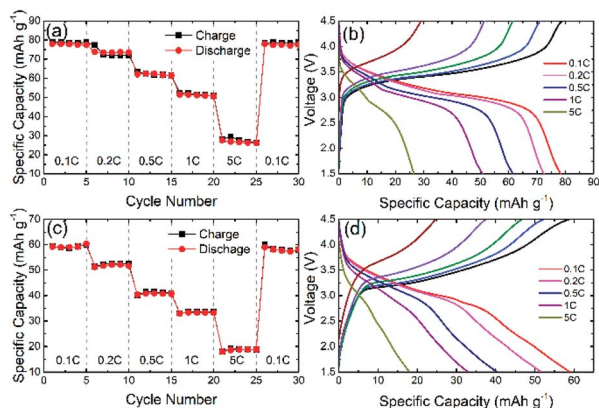


Fig. 9 Rate capability data, and galvanostatic charge–discharge curves at different C-rates for (a and b) $\text{NaFe}(\text{SO}_4)_2$, (c and d) $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$.

mention $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple is inactive and thus $\text{NaCr}(\text{SO}_4)_2$ has not included in the further electrochemical discussion. Fig. 9(a) shows the rate capability data for $\text{NaFe}(\text{SO}_4)_2$ at different C-rates. The initial discharge capacity at 0.1C is around 78 mA h g^{-1} which gradually decreases with increasing C-rate *i.e.* the discharge capacity at 1C is 50 mA h g^{-1} which reduces to 24 mA h g^{-1} and at 5C. Fig. 9(b) shows the galvanostatic charge/discharge curves for $\text{NaFe}(\text{SO}_4)_2$ which shows a slanting discharge plateau corresponding to $\text{Fe}^{2+}/\text{Fe}^{3+}$ redox couple. It can be noticed that intercalation/de-intercalation of sodium into/from the host structure takes place at about 3.2 V due to active $\text{Fe}^{2+}/\text{Fe}^{3+}$ redox couple which is consistent with dQ/dV analysis shown in Fig. 8.

Fig. 9(c) shows the rate capability data for $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ at different C-rates. As compared with $\text{NaFe}(\text{SO}_4)_2$ the discharge capacity delivered by $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ is inferior due to the presence of inactive $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple as indicated in dQ/dV analysis. With the addition of chromium, the amount of active iron decrease, therefore, the discharge capacity decreases. It can be noticed that the discharge capacity delivered by $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ at 0.1C is around 60 mA h g^{-1} and it also decreases with increasing C-rate similar to $\text{NaFe}(\text{SO}_4)_2$ *i.e.* the capacity at 1C and 5C are around 30 and 15 mA h g^{-1} respectively. Fig. 9(d) shows the galvanostatic charge/discharge curves for $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ at different C-rates. The shape of the discharge curves for $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ is slightly different as compared $\text{NaFe}(\text{SO}_4)_2$ which is mainly due to inactive chromium. The average voltage plateau due to active $\text{Fe}^{2+}/\text{Fe}^{3+}$ redox couple is comparatively lower as compared to $\text{NaFe}(\text{SO}_4)_2$ which is consistent with our dQ/dV results and may be attributed to multicomponent transition metals effect.^{29–31} We believe that suitable carbon coating technique to deposit a thin conductive carbon layer on $\text{NaCr}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ may result in active $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple leading to better energy storage performance.

To further investigate the activity of the redox couples in $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$, X-ray photoelectron spectroscopy (XPS) was carried out. Fig. 10(a and b) reveals that the Fe is present as Fe^{3+} (binding energy around 725.06 and 712.39

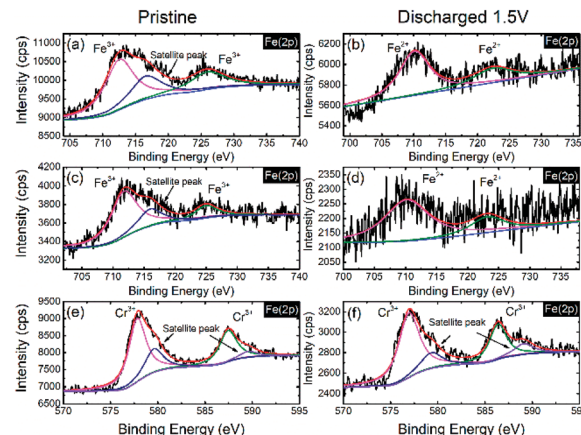


Fig. 10 XPS spectra for Fe in $\text{NaFe}(\text{SO}_4)_2$ (a) as prepared electrode, (b) discharged to 1.5 V, Fe in $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ (c) as prepared electrode, (d) discharged to 1.5 V and Cr in $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ (e) as prepared electrode, (f) discharged to 1.5 V.

eV) in the as-prepared electrode (before discharge) in $\text{NaFe}(\text{SO}_4)_2$, whereas it is reduced to Fe^{2+} (binding energy around 723.5 and 710.3 eV) after the cell is discharged to 1.5 V. The results for Fe in $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ are similar to $\text{NaFe}(\text{SO}_4)_2$ as shown in Fig. 10(c and d). Furthermore, the oxidation state for Cr in $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ is Cr^{3+} (binding energy around 577.73 and 587.48 eV) in the as-prepared electrode (before discharge) which is not changed after discharging to 1.5 V which clearly shows that $\text{Cr}^{3+}/\text{Cr}^{2+}$ redox couple is inactive and is consistent with the dQ/dV curves (Fig. 8) and charge-discharged curves as shown in Fig. 9(b and d).

Interestingly, the slanting voltage plateau in the charge/discharge curves of $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ indicates the intercalation/de-intercalation of sodium in these materials through a single-phase reaction as reported in previous studies for another cathode materials.²⁴ To confirm the interaction/de-intercalation mechanism of sodium into/from $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ during charging and discharging, the *ex situ* XRD was employed. For this purpose, the cells were charged/discharged at different cut-off voltages, immediately disassembled in a glove box, washed and dried. Finally, XRD spectra were recorded to study the structural variations during intercalation/de-intercalation of sodium into/from the host structure. Fig. 11(a and b) shows the *ex situ* XRD data for $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ respectively. It can be seen from that the XRD patterns that discharging the $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ to 1.5 V results in shifting of the XRD peaks to lower 2θ values which is due to the expansion of the unit cell due to intercalation of sodium into the host lattice. Furthermore, later charging the cells to 3.0 V and 4.5 V, shifts the XRD peaks to higher 2θ values which may be ascribed to the contraction of the unit cell due to de-intercalation of sodium from the host lattice. It is important to notice that there is only peak shifting during intercalation/de-intercalation and no new peak is generated which reflects the property of single-phase reaction. These results are consistent with the galvanostatic charge/discharge curves with slanting plateau which is



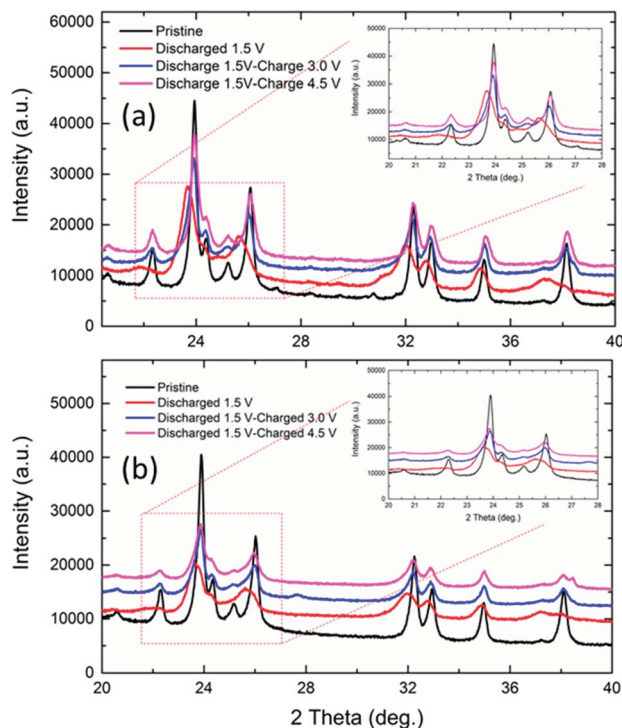


Fig. 11 *Ex situ* XRD patterns for (a) $\text{NaFe}(\text{SO}_4)_2$, (b) $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ at different voltages during charge and discharge.

associated with the single-phase mode of intercalation/de-intercalation in cathode materials.

Fig. 12 shows the cycling performance of $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ at 0.1C. The initial discharge capacity for $\text{NaFe}(\text{SO}_4)_2$ is around 80 mA h g^{-1} whereas for $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ it is around 60 mA h g^{-1} . After cycling for 30 cycles, the capacity for $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ is around 73 mA h g^{-1} and 59 mA h g^{-1} respectively. It is noticed that $\text{NaFe}(\text{SO}_4)_2$ shows slow capacity decay with successive cycling. Inset in Fig. 12 shows the capacity retention of $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ after 30 cycles. $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ retains 92.8% and 100% of its initial capacity after 30 cycles.

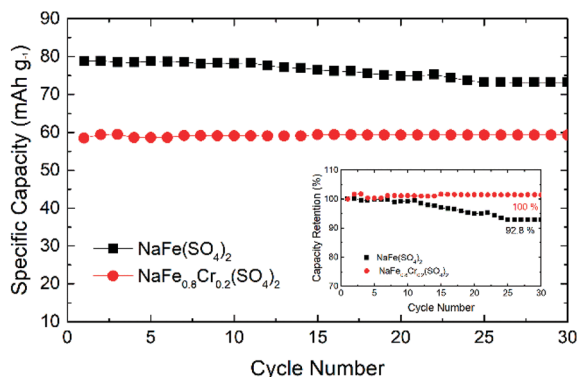


Fig. 12 Cycling performance of $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$, inset showing percentage capacity retention.

4 Conclusions

Phase pure $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ cathode materials (where $x = 0, 0.8, 1$) were synthesized through a sol-gel process and their structural, thermal and electrochemical properties are evaluated. The structural analysis indicates that $\text{NaFe}_x\text{Cr}_{1-x}(\text{SO}_4)_2$ ($x = 0, 0.8$ and 1.0) adopts a monoclinic crystal structure. It is noticed that $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ demonstrates better thermal properties as compared to $\text{NaCr}(\text{SO}_4)_2$ and $\text{NaFe}(\text{SO}_4)_2$. However, substitution of iron (Fe) with chromium (Cr) leads to inferior electrochemical performance which can be essentially ascribed to inactive $\text{Cr}^{2+}/\text{Cr}^{3+}$ redox couple. The *ex situ* XRD shows that the intercalation/de-intercalation mechanism in $\text{NaFe}(\text{SO}_4)_2$ and $\text{NaFe}_{0.8}\text{Cr}_{0.2}(\text{SO}_4)_2$ is governed by single-phase reaction.

Conflicts of interest

There are no conflicts to declare.

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