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Synthesis, Structure and Electrochemical Properties of $\text{Na}_3\text{MB}_5\text{O}_{10}$ (M = Fe, Co) containing M^{2+} in tetrahedral coordination

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Abstract

In the search for new cathode materials for sodium ion batteries, the exploration of polyanionic compounds has led to attractive candidates in terms of high redox potential and cycling stability. Herein we report the synthesis of two new sodium transition metal pentaborates $\text{Na}_3\text{MB}_5\text{O}_{10}$ (M = Fe, Co), where $\text{Na}_3\text{FeB}_5\text{O}_{10}$ represents the first sodium iron borate reported at present. By means of synchrotron X-ray diffraction, we reveal a layered structure consisting of pentaborate B_5O_{10} groups connected through M^{2+} in tetrahedral coordination, providing possible three-dimensional Na-ion migration pathways. Inspired by these structural features we examined the electrochemical performances versus sodium and show that $\text{Na}_3\text{FeB}_5\text{O}_{10}$ is active at an average potential of 2.5 V vs. Na^+/Na^0 , correlated to the $\text{Fe}^{3+}/\text{Fe}^{2+}$ redox couple as deduced from *ex situ* Mössbauer measurements. This contrasts with $\text{Na}_3\text{CoB}_5\text{O}_{10}$ which is electrochemical inactive. Moreover, we show that their electrochemical performances are kinetically limited as deduced by complementary AC/DC conductivity measurements, hence confirming once again the complexity in designing highly performant borate-based electrodes.

Introduction

Nowadays Li-ion batteries have become an essential device of our daily life, powering portable electronics such as laptops and mobile phones as well as electric- or hybrid vehicles. Despite a tremendous research effort over the last 25 years, they are still limited for large scale applications due to the high cost of stored energy per volume/ weight.^{1,2} Moving from Li- to Na-ion batteries could noticeably reduce the costs, considering the unlimited availability of sodium in earth's crust contrary to lithium. Similar to the Li-ion technology, the energy density of Na-ion cells is in principal limited by the performance of the positive electrode.³ Therefore strong research efforts have been focused on several positive electrode materials in the past, mainly based on layered sodium transition metal (TM) oxides Na_xMO_2 ^{4,5} or polyanionic type compounds such as $\text{Na}_3\text{V}_2(\text{PO}_4)_2\text{F}_3$ (NVPF)⁶ or $\text{Na}_2\text{Fe}_2(\text{SO}_4)_3$.⁷ The latter class of materials shows promising properties such as elevated redox potentials around 3-4 V vs. Na^+/Na^0 which are however negated by moderate gravimetric capacities due to the dead weight of the polyanionic group.^{7,8} Using borate BO_3^{3-} instead of phosphate PO_4^{3-} or sulfate SO_4^{2-} offers higher theoretical gravimetric capacities. It is well known that boron can adopt different oxygen coordination (BO_3^{3-} and BO_4^{5-}) capable of forming a large variety of polyborate anionic networks.⁹ Materials built upon such polyborate networks can offer open structures, with possible cationic migration pathways,¹⁰⁻¹² making them prospective candidates as insertion/deinsertion cathode materials for rechargeable Na-ion batteries.

This encouraged us to explore polyborate based compounds as cathode materials for Na-ion batteries. Besides that, to the best of our knowledge, transition metal borates have not been explored yet as cathode materials for Na-ion batteries. Looking for appropriate model compounds, recent publications about the synthesis and structure of compounds out of the family of sodium pentaborates $\text{Na}_3\text{MB}_5\text{O}_{10}$ ($\text{M} = \text{Mg}, \text{Ca}, \text{Zn}$)^{13,14} caught our interest, since it should be theoretical possible to reversibly desodiate these materials if M is a redox active TM. Based on this information we embarked in the synthesis of $\text{Na}_3\text{MB}_5\text{O}_{10}$ bearing 3d metals ($\text{M} = \text{Fe}, \text{Co}$). Herein we discuss our strategy to successfully prepare these new phases and describe their crystal structure as deduced by synchrotron X-ray diffraction. We also report their electrochemical and transport properties as deduced by means of galvanostatic- and AC/DC conductivity measurements. In spite of their rich crystal chemistry, these new phases are shown to present poor electrochemical performances due to kinetic limitations.

Experimental section

a) Synthesis

Stoichiometric amounts of NaOH (Sigma Aldrich, $\geq 97.0\%$), $\text{Fe}(\text{C}_2\text{O}_4)\cdot 2\text{H}_2\text{O}$ (Sigma Aldrich, 99.99%) or $\text{Co}(\text{OH})_2$ (Alfa Aesar, 99.9%) and H_3BO_3 (Alfa Aesar, 99+%) were mixed together for 15 min in argon using a high energy milling apparatus (SPEX). The resulting solid was reground with a mortar and pestle, filled into an alumina boat and heated up to 700°C for 1 h under argon flow. $\text{Na}_3\text{FeB}_5\text{O}_{10}$ was immediately transferred to the glove box for further use, since it undergoes degradation in ambient atmosphere. At the opposite, $\text{Na}_3\text{CoB}_5\text{O}_{10}$ is stable in air.

To prevent the precursors reacting with ambient atmosphere, all steps of the synthesis were carried out under inert atmosphere in an argon filled glove box or under argon flow in a tube furnace. All chemicals were stored under argon, sodium hydroxide NaOH and boric acid H_3BO_3 were dried prior to use at 200°C for 4 h in vacuum, and at 55°C for 24 h in air, respectively.

b) Physical characterization

Synchrotron XRD patterns were recorded in transmission mode ($\lambda=0.413965 \text{ \AA}$) at the 11-BM beam line at Argonne National Lab. The powder was placed into the provided sample holders, however in the case of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ it was filled in $\varnothing=0.7 \text{ mm}$ quartz capillaries before which were sealed under inert atmosphere and then introduced into the sample holder. All powder patterns were refined using the Rietveld method¹⁵ as implemented in the FullProf program.¹⁶

^{57}Fe Mössbauer spectra were collected using 20-30 mg/cm^2 of positive electrode powder. The spectrometer is operated in the constant acceleration mode and in transmission geometry. A $^{57}\text{Co}(\text{Rh})$ source with a nominal activity of 370 MBq was used. The source and the absorber were always kept at 295 K. All the isomer shifts are given relative to α -Fe standard at 295 K.

Thermogravimetric analysis coupled with differential scanning calorimetry (TGA-DSC) measurements were recorded with a STA 449C Netzsch apparatus between room

temperature and 900°C under air by applying a heating rate of 5°C/min under air using around 15-20 mg of the pristine material.

c) Electrochemical characterization

Electrochemical tests were carried out in Swagelok-type cells. The positive electrode material was prepared by mixing the active material with 20 wt.% carbon SP for 10 min using a mortar and pestle under argon. Sodium was used as the negative electrode separated by two sheets of glassy fiber (Whatman GF/D borosilicate) soaked with the electrolyte from the positive side (~10 mg powder). As electrolyte a 1M solution of NaClO₄ in a mixture of ethylene carbonate (EC) and dimethyl carbonate (DMC) 1/1 (w/w) with 1wt.% fluoroethylene carbonate (FEC) additive was used. The cells were assembled in an argon filled glove box (MBraun, Germany, O₂ and H₂O < 0.1 ppm). Galvanostatic tests were performed at 55°C at a C/40 (C-rate here is defined as the time needed to remove or insert 1 Na⁺ from Na₃MB₅O₁₀ in 1 h) using a VMP3 potentiostat/galvanostat (BioLogic S.A., Claix, France).

Conductivity measurements were carried out using a BioLogic MTZ-35 setup with ionically blocking platinum electrodes equipped with a HTF-1100 furnace. AC impedance measurements were recorded from 30 MHz to 0.01 Hz in the temperature range from 250 to 400 °C under argon flow, applying 200 mV voltage amplitude. The activation energy E_a was calculated from fitting the experimental values using the Arrhenius equation $\sigma T = \sigma_0 \cdot \exp(-E_a/k_B T)$, where σ is the conductivity, T the temperature, σ_0 is a pre-exponential factor and k_B the Boltzmann constant. DC measurements were done by applying a constant voltage of 0.5 V until the current response reached a constant value. The resistance was then calculated from the steady state current response using Ohm's law. A pellet suitable for the measurement was prepared by pressing around 300 mg of powdered sample in a 10 mm die. The pellet was following annealed for 124 hours at 650°C under argon, or for 124 hours at 600°C in a vacuum sealed tube for $M = \text{Co}$ and $M = \text{Fe}$, respectively. Note that the conductivity values are not corrected by the pellets porosities (around 65-75% of theoretical density).

Results and Discussion

a) Synthesis

A standard solid state synthesis, which consists in mixing stoichiometric amounts of Na_2CO_3 , M(II) carbonate/nitrate and H_3BO_3 followed by annealing the mixture at temperatures between 650–700°C in air for several days/weeks, had already been reported to prepare $\text{Na}_3\text{MB}_5\text{O}_{10}$ (M = Mg, Ca, Zn).^{13,14} Our first attempts by replacing the M(II)-carbonate/nitrate through Co/Fe(II)-oxalate and heating the mixture under argon to avoid oxidation of the transition metal were not successful. In contrast by using NaOH instead of Na_2CO_3 as sodium source, we succeeded in forming the $\text{Na}_3\text{FeB}_5\text{O}_{10}$ phase, but not the cobalt one. The cobalt phase could only be obtained if cobalt(II) hydroxide $\text{Co}(\text{OH})_2$ was used as Co-precursor, hence preventing the easy reduction to metallic Co pertaining to the use of Co(II)-oxalate.

After the synthesis, both powders are made of large particles with sizes ranging from 1 to 50 μm as shown in Figure 1. Note that a longer annealing time did not improve the crystallinity of the products.

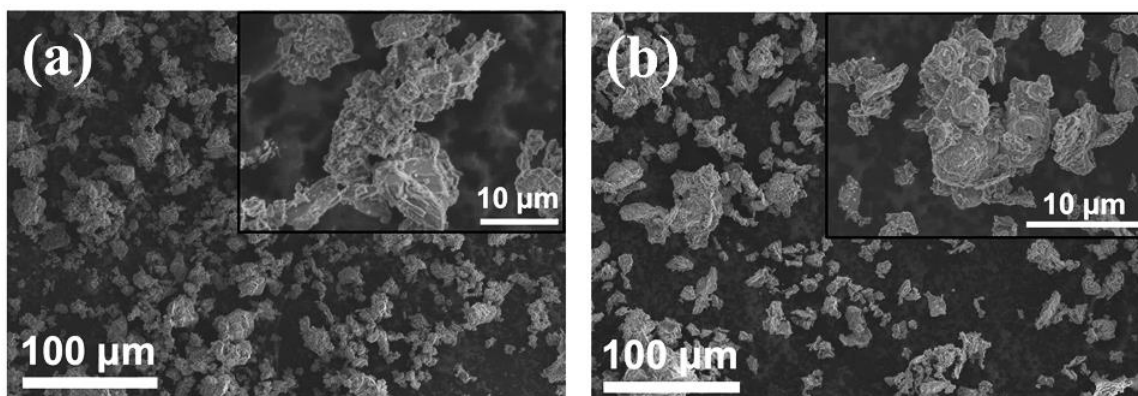


Figure 1: Representative SEM images of (a) $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and (b) $\text{Na}_3\text{CoB}_5\text{O}_{10}$ as prepared.

b) Structural characterization

The structure of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ was examined by means of synchrotron X-ray powder diffraction (XRD). Besides peaks attributed to Fe_3BO_5 impurity (ca. 4%), the XRD pattern can

be perfectly indexed in an orthorhombic unit cell having a *Pbca* space group with lattice parameters $a = 7.95493(1) \text{ \AA}$, $b = 12.30669(1) \text{ \AA}$ and $c = 18.03978(1) \text{ \AA}$ and a unit cell volume $V = 1766.073 \text{ \AA}^3$ (Table 1). We therefore refined the pattern with the Rietveld method starting from the isostructural model reported for $\text{Na}_3\text{MgB}_5\text{O}_{10}$,¹³ but replacing Mg with Fe. All atoms are placed in general positions 8c with one crystallographic site for iron, three for sodium, five for boron and ten for oxygen atoms. The Rietveld refinement is shown in Figure 2a and fully validates this approach. The final atomic positions are given in Table 1. Figure 3 shows the $\text{Na}_3\text{FeB}_5\text{O}_{10}$ structure. Iron atoms are located in the center of slightly distorted tetrahedra (Fe-O distances ranging from 1.97 to 2.12 \AA) with an average bond length of 2.02 \AA , in good agreement with reported Fe-O distances in tetrahedral oxygen coordination.^{17,18} All four oxygen ligands of the FeO_4 tetrahedra are also terminal oxygen atoms of distinct pentaborate anions $[\text{B}_5\text{O}_{10}]^{5-}$. The latter are built upon one BO_4 tetrahedra (average bond length 1.48 \AA) linked to four trigonal planar BO_3 groups by oxygen vertices forming $[\text{B}_5\text{O}_{10}]^{5-}$ units (Figure 3d). These pentaborate anions connect the FeO_4 tetrahedra to form 2D layers along the *ab*-plane. These layers are stacked along the *c*-axis with sodium atoms sitting between the layers as well as in channels along the *a*-axis.

The Fe-Mössbauer spectrum for the as prepared powder is shown in Figure 4, and the majority of iron can be fitted with a single doublet, confirming the occurrence of one single crystallographic site for iron. The isomer shift $\text{IS} = 1.02 \text{ mm/s}$ and a quadrupole splitting $\text{QS} = 2.10 \text{ mm/s}$ is typical of Fe^{2+} in tetrahedral coordination (Table S1).^{19,20} Note that the Mössbauer spectrum contains about 4% of Fe_3BO_5 impurity, consistent with the values obtained from the synchrotron XRD pattern (Figure 2a).

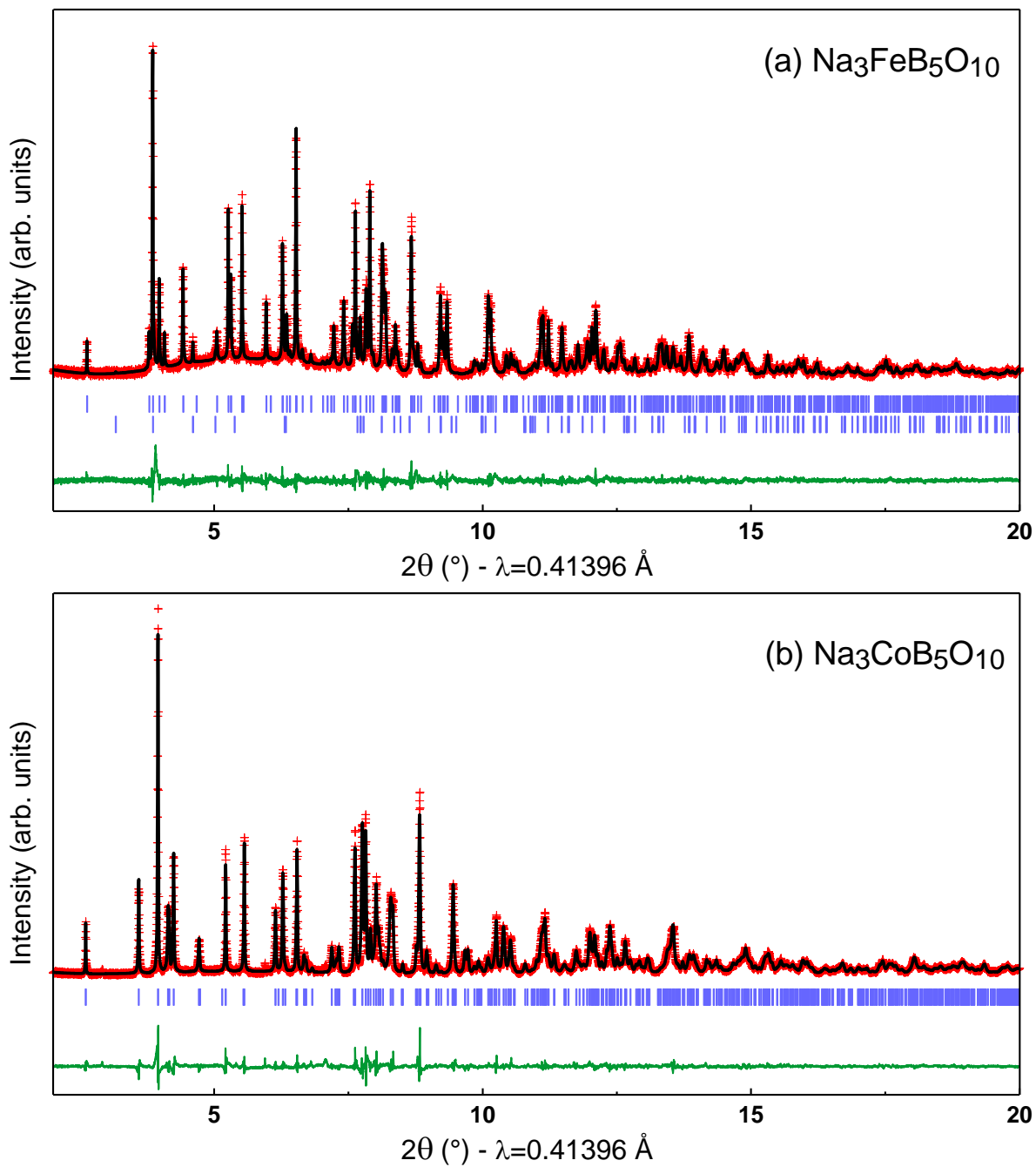


Figure 2: Rietveld refinement of synchrotron X-ray diffraction patterns for (a) $\text{Na}_3\text{FeB}_5\text{O}_{10}$, and (b) $\text{Na}_3\text{CoB}_5\text{O}_{10}$. The red crosses, black line, and green line represent the observed, calculated and difference patterns, respectively. The positions of the Bragg reflections are shown as vertical blue bars. For $\text{Na}_3\text{FeB}_5\text{O}_{10}$, the lower set of tick marks is Fe_3BO_5 (~4 w%).

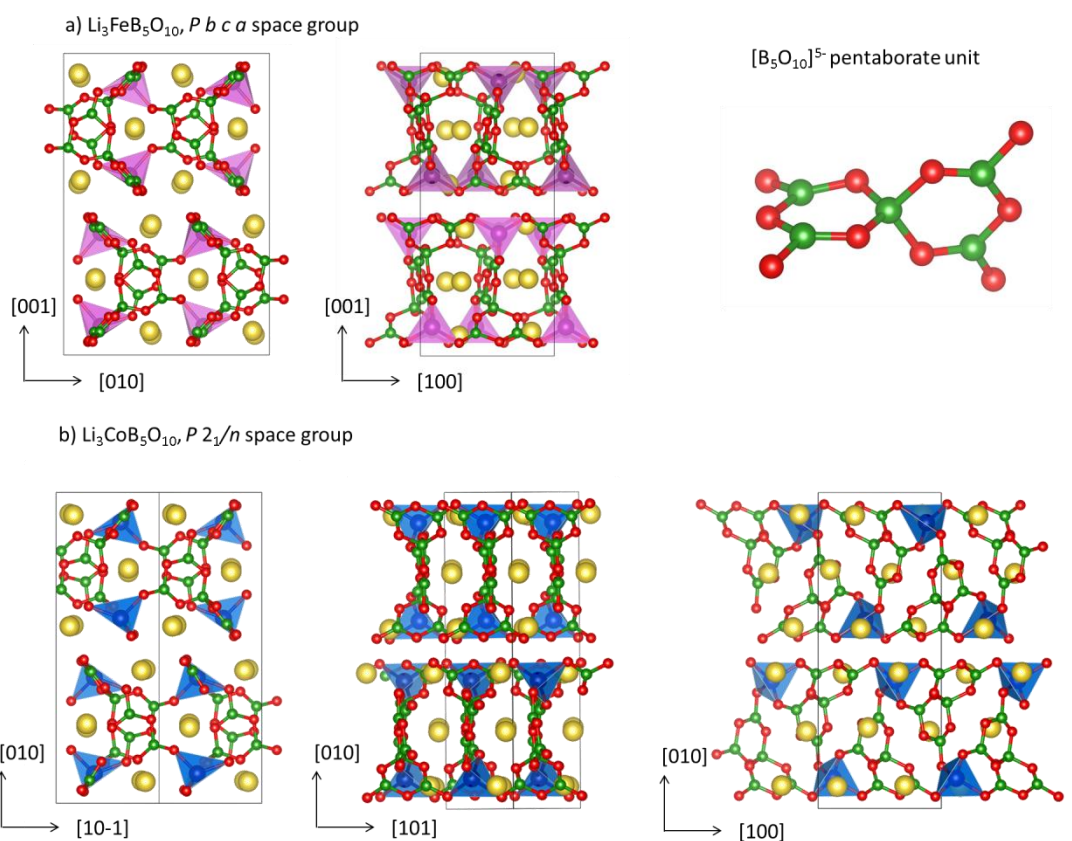


Figure 3: Structure of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ (a) and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ (b) viewed along different crystallographic directions, and illustration of the pentaborate unit $[\text{B}_5\text{O}_{10}]^{5-}$. Oxygen, boron, sodium are coloured in red, green and yellow respectively. Iron atoms are in pink in (a) and cobalt atoms are in blue (b), and both are shown in their tetrahedral environment.

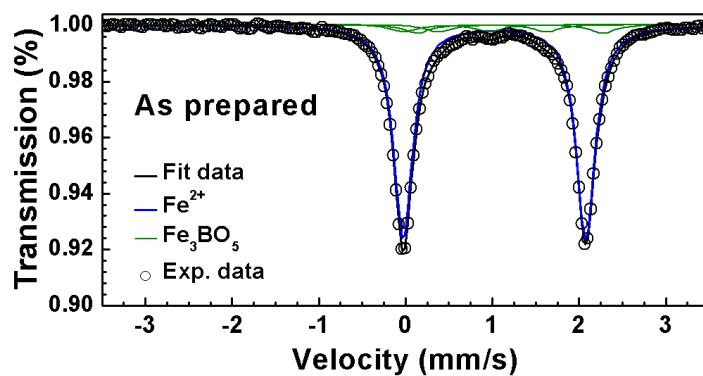


Figure 4: Room temperature Mössbauer spectrum of the as prepared $\text{Na}_3\text{FeB}_5\text{O}_{10}$.

Table 1: Structural parameters of Na₃FeB₅O₁₀ determined from Rietveld refinement of the synchrotron XRD pattern. Isotropic temperature factors have been imposed equal for a same chemical species.

Na ₃ FeB ₅ O ₁₀ <i>P b c a</i> space group					
a= 7.95493(5) Å, b= 12.30668(9) Å, c= 18.03983(13) Å					
V=1766.07(2) Å ³ Z=8					
Density = 2.548 g/cm ³					
Bragg R-factor=5.96%, χ^2 =1.02					
Atom	Wyckoff site	x	y	z	B (Å ²)
Na1	8c	0.8292(5)	0.4179(4)	0.0800(3)	1.97(7)
Na2	8c	0.6936(5)	0.6544(3)	0.2548(3)	1.97(7)
Na3	8c	0.7908(5)	0.4278(4)	0.4330(2)	1.97(7)
Fe1	8c	0.4135(3)	0.68078(13)	0.09531(9)	1.29(4)
B1	8c	0.414(2)	0.1491(10)	0.0758(7)	1.09(14)
B2	8c	0.7215(19)	0.1756(12)	0.0763(7)	1.09(14)
B3	8c	0.5457(17)	0.2864(10)	0.1604(7)	1.09(14)
B4	8c	0.5578(17)	0.3620(11)	0.3004(7)	1.09(14)
B5	8c	0.4911(16)	0.4776(10)	0.1874(7)	1.09(14)
O1	8c	0.2900(10)	0.1182(6)	0.0430(4)	1.53(6)
O2	8c	0.4238(12)	0.2192(5)	0.1266(3)	1.53(6)
O3	8c	0.5902(11)	0.1222(5)	0.0472(4)	1.53(6)
O4	8c	0.7243(9)	0.2498(6)	0.1277(3)	1.53(6)
O5	8c	0.8771(9)	0.1416(5)	0.0435(3)	1.53(6)
O6	8c	0.5657(8)	0.2707(5)	0.2427(4)	1.53(6)
O7	8c	0.5371(8)	0.3980(5)	0.1466(3)	1.53(6)
O8	8c	0.4969(8)	0.4514(5)	0.2724(4)	1.53(6)
O9	8c	0.5762(10)	0.3344(5)	0.3713(3)	1.53(6)
O10	8c	0.4102(9)	0.5690(5)	0.1844(3)	1.53(6)

Having successfully solved the structure of $\text{Na}_3\text{FeB}_5\text{O}_{10}$, we turned next to $\text{Na}_3\text{CoB}_5\text{O}_{10}$. The corresponding synchrotron XRD pattern immediately indicates that the structure of the Co-based sample is different in symmetry from the iron analog. The Bragg peaks can be rather indexed in a monoclinic space group $P 2_1/n$ with lattice parameters $a = 6.65143(1) \text{ \AA}$, $b = 18.20986(1) \text{ \AA}$, $c = 7.80513(1) \text{ \AA}$ and $\beta = 114.79220(1)^\circ$ (unit cell volume $V = 858.240 \text{ \AA}^3$). A Rietveld refinement was carried out from the structural model of $\text{K}_3\text{CdB}_5\text{O}_{10}$ ¹⁹, and by replacing K and Cd with Na and Co, respectively. The final Rietveld refinement is shown in Figure 2b and Table 2 gathers the refined structural parameters. All atoms are in general position $4e$ of space group $P 2_1/n$ with one crystallographic site for cobalt, three for sodium, five for boron and ten for oxygen atoms. Figure 3b presents the structure of $\text{Na}_3\text{CoB}_5\text{O}_{10}$. Similarly to the iron counterpart and despite a different space group, the structure is also built upon CoO_4 tetrahedra and $[\text{B}_5\text{O}_{10}]^{5-}$ units. The average Co-O bond length in the tetrahedral is 1.98 \AA , in agreement with values reported in the case of other compounds possessing Co^{2+} in tetrahedral environment.²⁰ Overall, the CoO_4 tetrahedra and $[\text{B}_5\text{O}_{10}]^{5-}$ arrangement resembles the one of the Fe counterpart, with similar 2D layers stacked along the b -axis.

Table 2: Structural parameters of $\text{Na}_3\text{CoB}_5\text{O}_{10}$ determined from Rietveld refinement of the synchrotron XRD pattern. Isotropic temperature factors have been imposed equal for all oxygen atoms.

$\text{Na}_3\text{CoB}_5\text{O}_{10}$ $P 2_1/n$ space group					
$a = 6.65143(6) \text{ \AA}$, $b = 18.20981(15) \text{ \AA}$, $c = 7.80512(7) \text{ \AA}$ and $\beta = 114.79223(3)^\circ$					
$V = 858.235(12) \text{ \AA}^3$ $Z = 4$					
Density = 2.647 g/cm^3					
Bragg R-factor = 6.56% , $\chi^2 = 1.47$					
Atom	Wyckoff	x	y	z	B (\AA^2)

	site				
Co1	4e	0.4938(2)	0.09529(6)	0.12100(16)	0.97(3)
Na1	4e	1.0215(5)	0.06927(15)	0.1537(4)	1.31(7)
Na2	4e	0.5356(5)	0.07253(15)	-0.2967(4)	2.25(9)
Na3	4e	0.2007(5)	0.24423(14)	-0.0962(4)	0.85(3)
B1	4e	0.7050(15)	0.2127(5)	-0.0092(14)	0.7(2)
B2	4e	0.1192(16)	0.0766(4)	-0.2204(12)	0.5(2)
B3	4e	0.1200(19)	0.1643(6)	-0.4532(14)	1.9(3)
B4	4e	0.2734(16)	0.1881(6)	0.3233(13)	1.7(3)
B5	4e	0.8089(15)	0.0738(5)	0.4833(12)	0.52(20)
O1	4e	0.3823(8)	0.3659(3)	-0.0583(6)	0.85(3)
O2	4e	0.6864(9)	0.1334(2)	0.0034(6)	0.85(3)
O3	4e	0.6234(8)	0.0399(2)	0.3567(6)	0.85(3)
O4	4e	0.2150(8)	0.1274(3)	-0.2604(7)	0.85(3)
O5	4e	0.2347(8)	0.1406(2)	-0.5576(7)	0.85(3)
O6	4e	0.2214(8)	0.0375(2)	-0.0479(6)	0.85(3)
O7	4e	0.1189(7)	0.2431(3)	-0.4250(6)	0.85(3)
O8	4e	1.0892(9)	-0.0428(2)	0.3357(7)	0.85(3)
O9	4e	0.3217(8)	0.1712(2)	0.1747(6)	0.85(3)
O10	4e	-0.2188(8)	0.2304(2)	-0.1428(7)	0.85(3)

This similarity does not come as a surprise since we noticed that both structure are actually structurally related. For instance, $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ lattice basis vectors are related by the following matrix:

$$\begin{pmatrix} \mathbf{a}_{Fe} \\ \mathbf{b}_{Fe} \\ \mathbf{c}_{Fe} \end{pmatrix} = \begin{pmatrix} 1 & 0 & 1 \\ -1 & 0 & 1 \\ 0 & -1 & 0 \end{pmatrix} \begin{pmatrix} \mathbf{a}_{Co} \\ \mathbf{b}_{Co} \\ \mathbf{c}_{Co} \end{pmatrix}$$

The thermal stability of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ was checked by the TGA-DSC measurements. The data were recorded between room temperature and 900°C by applying

a heating rate of 5°C/min under air (Figure S1). $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ are stable until 600 and 750°C, respectively. Above 600°C, an exothermic peak is observed for $\text{Na}_3\text{FeB}_5\text{O}_{10}$. This peak is most likely associated with the dissociation of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ into iron oxide Fe_2O_3 and $\text{Na}_2\text{B}_4\text{O}_7$, as confirmed by the corresponding XRD pattern taken after annealing a sample for 12h at 650°C (Figure S2). A weight gain around 2% is visible above 600°C, furthermore suggesting oxygen uptake through the formation of Fe_2O_3 . At higher temperatures we note the presence of large endothermic peaks at 710 and 770°C attributed to the melting point of the decomposition products in the case of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ ($\text{Na}_2\text{B}_4\text{O}_7$ and Fe_2O_3) and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ respectively.

c) Electrochemical characterization

As described in the crystallographic section, both $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ structures offer the possibility of 3D Na mobility, making it suitable as a possible cathode material for Na-ion batteries. Therefore we examined their electrochemical performance in half cells versus sodium in non-aqueous electrolyte using galvanostatic experiments.

A typical voltage-composition curve for $\text{Na}_3\text{FeB}_5\text{O}_{10}$ starting on oxidation with a C/40 rate at 55°C between 4.5 and 1.5 V is shown in Figure 5a. During the initial oxidation voltage increases rapidly and above 4.0 V a pseudo plateau is reached where it slopes slowly up to the cutoff potential of 4.5 V. Upon following reduction the potential drops steep down to 2.0 V reaching a sloping plateau down to the lower cutoff voltage of 1.5 V. Approximately ~ 0.6 Na^+ can be removed during the first charge and 0.3 Na^+ can be reinserted upon following discharge, leading to a reversible capacity of around 30 mAh/g (80 mAh/g theoretical capacity for the reversible removal of 1 Na^+). Interestingly the amount of reversible Na^+ reinserted during the second cycle is larger than for the initial one, pointing towards an irreversible Na^+ loss during the first oxidation. As the galvanostatic curve shows a large voltage gap between charge and discharge, a precise determination of the $\text{Fe}^{3+}/\text{Fe}^{2+}$ redox potential was impossible. Consequently galvanostatic intermittent titration technique (GITT) with open circuit voltage (OCV) periods each 0.1 Na^+ extracted/inserted was done, allowing the system to reach the true thermodynamic potentials. As shown in Figure 6b, after each positive or negative current pulse, strong alternative drops and increases in potential (from 0.5 to ~ 2 V) are observed, indicative of poor Na^+ /electronic transport within this phase.

However at the end of each OCV period an equilibrium potential of ~ 2.5 V vs. Na^+/Na^0 is reached, implying a biphasic Na insertion/deinsertion process. The difference in the amount of Na extracted and reinserted during the GITT measurement compared to galvanostatic cycling can be most likely correlated a pronounced electrolyte decomposition for the former one, as it is much more time consuming hence the electrolyte is much longer exposed to elevated temperature and voltage.

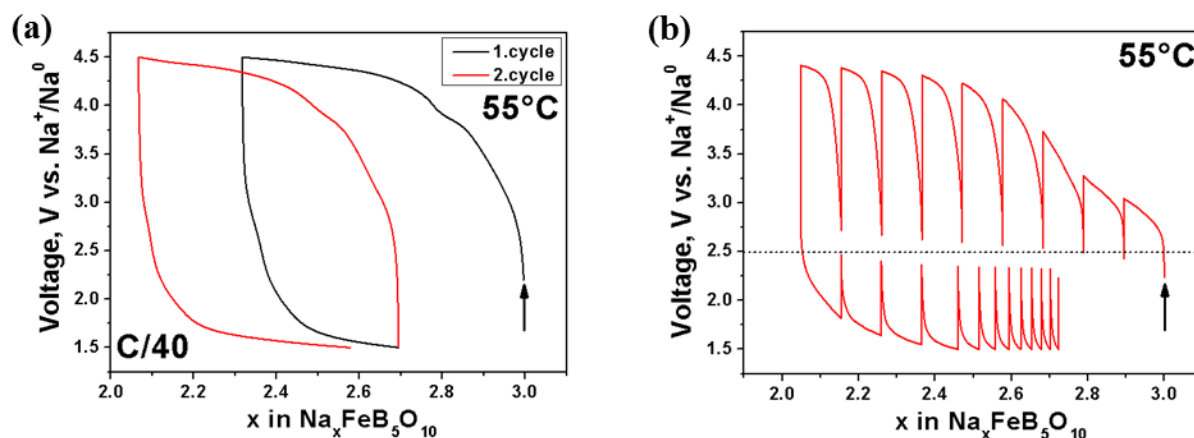


Figure 5: (a) Voltage-composition and (b) GITT curve of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ versus sodium between 4.5 and 1.5 V at 55°C using a C/40 rate. In the case of (b) OCV periods of 12h alternate with 4h of current pulse.

To confirm the $\text{Fe}^{3+}/\text{Fe}^{2+}$ redox process, *ex situ* Mössbauer spectra for charged (4.5 V) and discharged (1.5 V) samples of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ were recorded (Figure 6). To recover the active material, cells were disassembled inside the dry box, the powder was recovered and washed two times with DMC and dried under vacuum prior to examination. At the end of charge, the Fe^{2+} doublet is reduced to the benefit of an additional contribution arising from Fe^{3+} which is diminished upon following reduction, however the reduction of Fe^{3+} to Fe^{2+} is not fully reversible. The variations of the total amounts of $\text{Fe}^{3+}/\text{Fe}^{2+}$ are given in Table S1 and are in agreement with the amount of sodium extracted and reinserted as seen in the electrochemical charge/discharge curve.

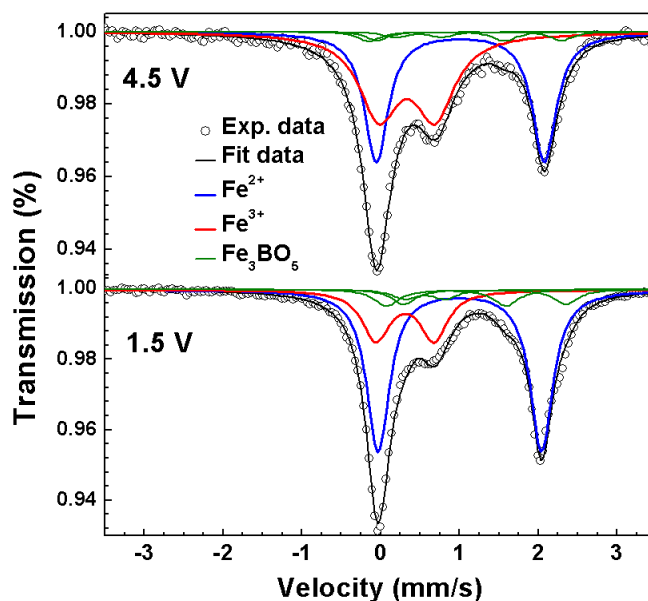


Figure 6: Room temperature *ex situ* Mössbauer spectra recorded for $\text{Na}_3\text{FeB}_5\text{O}_{10}$ at 4.5 V (charged) and at 1.5 V (discharged).

Lastly, we tested the Co-based phase by assembling a $\text{Na}/\text{Na}_3\text{CoB}_5\text{O}_{10}$ cell under a similar protocol as for $\text{Na}/\text{Na}_3\text{FeB}_5\text{O}_{10}$ and observed no electrochemical activity till 4.5 V. This does not come as a total surprise since we know from DFT calculations that in a well-defined family of 3d polyanionic compounds, the $\text{M}^{3+}/\text{M}^{2+}$ redox potential increases from Fe to Mn, Co and Ni with differences that can be near 1.5 V between $\text{Fe}^{3+}/\text{Fe}^{2+}$ and $\text{Co}^{3+}/\text{Co}^{2+}$ in olivine-type LMPO_4 compounds for instance. Thus, electrolyte with higher stability against oxidation is sorely needed to further explore such high potential redox materials.²¹

At this stage, a legitimate question deals with the origin of the poor electrochemical performances of these phases which can be due to either the inactivity of the cationic redox couples over the explored potential or to some kinetic limitations associated to poor ionic and electronic transport.

To first sense the aptitude of such compounds towards ionic conduction, bond valence energy landscapes (BVEL) were generated according to the method developed by S. Adams.²² Figure 7 shows the calculated BVEL maps representing the Na^+ conduction pathways in $\text{Na}_3\text{MB}_5\text{O}_{10}$ ($\text{M} = \text{Fe}$ (a), Co (b)). These maps were calculated considering anionic neighbours up to 8 Å for Na. For $\text{Na}_3\text{FeB}_5\text{O}_{10}$, infinite 1D Na^+ conduction paths, involving Na1 and Na3 sites and not Na2 one that is sterically hindered, appear along the a-axis at 1.00 eV above the minimum energy. When applying the same calculations to $\text{Na}_3\text{CoB}_5\text{O}_{10}$, we note that a percolation energy of 1.09 eV above the minimum is required to get fully connected

pathways for Na migration. For both compounds both percolation energies are in the same order of magnitude, moreover the Na^+ conduction pathways are lying in between the $\text{MO}_4/[\text{B}_5\text{O}_{10}]$ layers, obviously implying an anisotropic conduction.

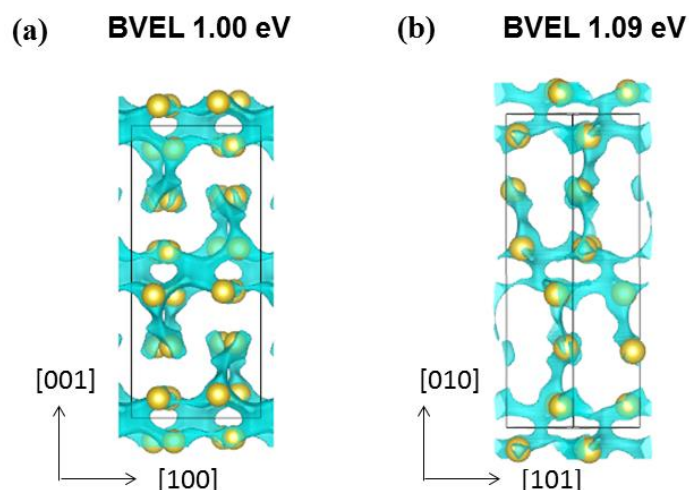


Figure 7: Bond valence energy landscape (BVEL) of (a) $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and (b) $\text{Na}_3\text{CoB}_5\text{O}_{10}$. The percolation energies are given in the figure. Iso-energy blue domains indicate paths for Na (yellow spheres) conduction.

In order to validate these results we measured material's conductivity by AC impedance spectroscopy. The Impedance spectra were recorded on sintered pellets between ionically blocking platinum electrodes for temperatures ranging from 250 to 400°C in argon flow. A depressed semicircle with a Warburg tail was measured for $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and $\text{Na}_3\text{CoB}_5\text{O}_{10}$ (Figure S3), suggesting the AC conductivity to be mostly ionic. Both spectra were fitted with the equivalent circuit shown as inset in Figure S4 including bulk and grain boundary contributions. The corresponding Arrhenius plots are shown in Figure 8 where for $M = \text{Fe}$, one can first note a linear increase of the bulk ionic conductivity from $1.5 \cdot 10^{-8}$ to $1.7 \cdot 10^{-7}$ S/cm at 250°C and 380°C respectively, with an activation energy of 0.65 eV. The grain boundary contribution also increases linearly from $1.9 \cdot 10^{-7}$ to $4.7 \cdot 10^{-7}$ S/cm for 250°C and 380°C respectively, exhibiting a smaller activation energy of 0.24 eV. Surprisingly, the grain boundary conductivity is higher than the bulk one. One possibility to account for this information is rooted in the existence of a minor Fe_3BO_5 impurity (4%) in our sample which is expected to affect the grain boundaries, since it is well known that impurities are typically

located between the grains. Therefore, they can lead to enhanced grain boundary conductivities so that our experimental values may be overestimated.²³

Turning to $M = \text{Co}$, the bulk ionic conductivity increases linearly from $6.9 \cdot 10^{-8}$ to $9.8 \cdot 10^{-7}$ S/cm between 280 and 390°C with an associated activation energy of 0.73 eV. The grain boundary ionic conductivity follows the same trend in a parallel way, however the absolute values are about one order of magnitude lower than the values obtained for the bulk conductivity ($5.9 \cdot 10^{-9}$ to $8.6 \cdot 10^{-8}$ S/cm), corresponding to an activation energy of 0.83 eV. Note that the linear extrapolation of the experimental values leads to RT conductivities of $4.0 \cdot 10^{-13}$ and $9.1 \cdot 10^{-17}$ S/cm for the bulk and grain boundary contribution respectively.

To clearly separate the ionic- from the electronic contribution, additional DC polarization measurements were conducted. After applying a potential of 0.5 V across the sample, the current response was followed and a sharp increase followed by a slow decay to a steady state current value was observed. It represents the pure electronic conduction of the material as we used ionically blocking Pt electrodes (Figure S4). Note that for $M = \text{Fe}$, the decay is approximately 4 times faster compared to $M = \text{Co}$, indicating faster Na^+ migration. For $M = \text{Fe}$ the electronic conductivity increases between 250 to 350°C from $6.3 \cdot 10^{-10}$ to $1.0 \cdot 10^{-8}$ S/cm and such an increase is correlated to an activation energy of 0.72 eV. For $M = \text{Co}$ an increase between 290 to 390°C from $1.1 \cdot 10^{-9}$ to $2.2 \cdot 10^{-8}$ S/cm is observed with a corresponding activation energy of 0.97 eV. Lastly, a linear extrapolation of the experimental values leads to RT electronic conductivities of $1.2 \cdot 10^{-14}$ and $5.4 \cdot 10^{-17}$ S/cm for $M = \text{Fe}$ and Co respectively.

For comparison, it should be recalled that the values for the RT bulk ionic conductivity of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ are one only order of magnitude lower than the one of LiFePO_4 ($\approx 2 \cdot 10^{-9}$ S/cm)²⁴ and two order of magnitudes higher than $\text{Li}_2\text{FeSiO}_4$ ($\approx 6 \cdot 10^{-14}$ S/cm)²⁵, both of which can reversibly insert one Li per unit formula. Having this in mind, the poor electrochemical performance of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ comes still somewhat as a surprise as it should not be ionically limited. Pushing further this comparison, it is worth mentioning that the electronic conductivity of LiFePO_4 is higher than that of $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and this could be a reason for the poor electrochemical properties as we need both ionic and electronic conductivity to electrochemically trigger insertion/ deinsertion reactions. Such a limitation can also be of structural nature with the under-stoichiometric $\text{Na}_{3-x}\text{FeB}_5\text{O}_{10}$ phase being thermodynamically unstable, the reason why for instance the de-insertion of Li from

$\text{Li}_2\text{FeSiO}_4$ could not be extended below one, as confirmed by DFT calculations. Turning back to (Figure 5b), one can note that the equilibrium potential reaches a plateau which is indicative of a two-phase Na insertion/ deinsertion reaction, hence one can assume the formation of an intermediate phase $\text{Na}_x\text{FeB}_5\text{O}_{10}$ which is most likely the limiting factor for electrochemical activity, similar to what was observed in the case of LiFeBO_3 .^{26,27} Distinguishing between electronic and structural limitations will require DFT calculations which have been planned in view of establishing the room temperature $\text{Na}_{3-x}\text{FeB}_2\text{O}_{10}$ phase diagram.

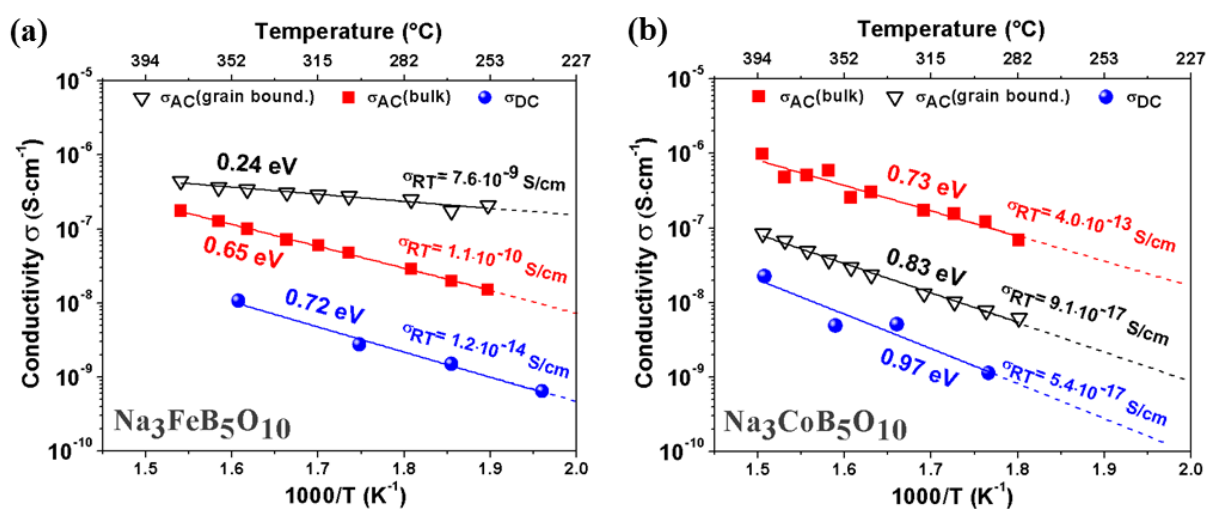


Figure 8: Arrhenius plot of the AC and DC conductivity of (a) $\text{Na}_3\text{FeB}_5\text{O}_{10}$ and (b) $\text{Na}_3\text{CoB}_5\text{O}_{10}$ derived in the temperature range from around 250 to 400°C during heating in argon flow.

Conclusion

We have reported the successful synthesis of two new sodium transition metal pentaborates $\text{Na}_3\text{MB}_5\text{O}_{10}$ ($M = \text{Fe}, \text{Co}$) and solved their crystal structures from synchrotron XRD. Related structures of both compounds show layers consisting of pentaborate B_5O_{10} groups connected through M^{2+} in tetrahedral coordination. Complementary galvanostatic- and *ex situ* Mössbauer measurements have confirmed a $\text{Fe}^{3+}/\text{Fe}^{2+}$ redox activity centered around 2.5 V vs. Na^+/Na^0 for $\text{Na}_3\text{FeB}_5\text{O}_{10}$. In contrast no electrochemical activity was

detected for $\text{Na}_3\text{CoB}_5\text{O}_{10}$. Combined BVEL calculations and AC/DC conductivity measurements reveal strong kinetic limitations for $\text{Na}_3\text{CoB}_5\text{O}_{10}$, reflected in the low RT ionic/electronic conductivities of $4.0 \cdot 10^{-13}$ and $5.4 \cdot 10^{-17}$ S/cm respectively. In contrast $\text{Na}_3\text{FeB}_5\text{O}_{10}$ shows ionic conductivity of $1.1 \cdot 10^{-10}$ S/cm comparable to LFP, however the RT electronic conductivity ($1.2 \cdot 10^{-14}$ S/cm) is much lower. Besides the poor electronic conductivity, the limited electrochemical performance could equally be due to the structural/thermodynamic instability of the intermediate phase $\text{Na}_x\text{FeB}_5\text{O}_{10}$ formed during the two-phase (de)sodiation reaction.

Although a reversible electrochemical activity could be demonstrated for $\text{Na}_3\text{FeB}_5\text{O}_{10}$, this material cannot be considered for practical applications. Attempts to combat the electronic limitation by coating the particles with a conducting carbon layer have failed in increasing its electrochemical activity. Obviously, these results call for further exploration of polyanionic borate based materials in terms of their ionic mobility, despite difficulties encountered during the design of new compounds out of this family.

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Notes

The authors declare no competing financial interest.

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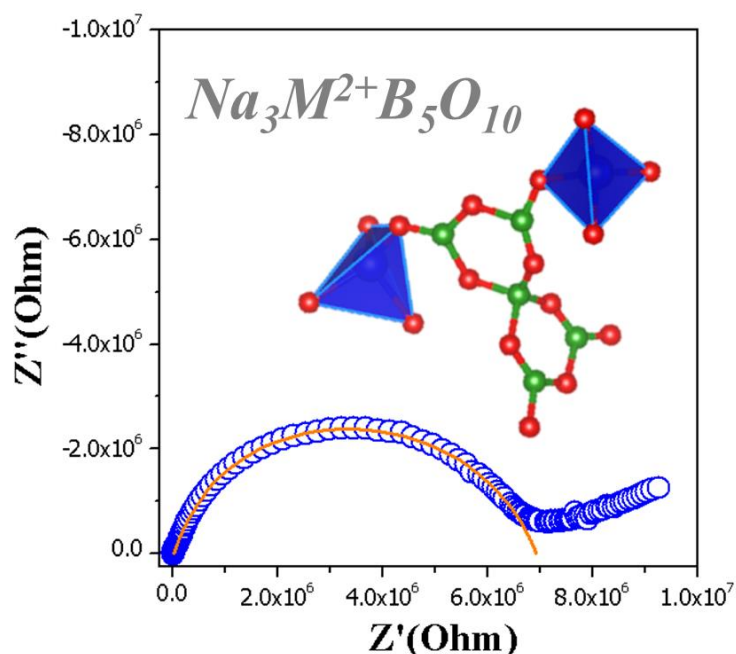
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TOC graphic



Representative Nyquist plot of the AC impedance of $\text{Na}_3\text{MB}_5\text{O}_{10}$ containing M^{2+} in tetrahedral coordination schematic view of two M^{2+}O_4 tetrahedra connected through one pentaborate B_5O_{10} unit.