Sodium Rechargeable Batteries with Electrolytes Based on Nafion Membranes Intercalated by Mixtures of Organic Solvents

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Received: 17 October 2018; Accepted: 31 October 2018; Published: 1 December 2018

Abstract: The possibilities of manufacturing batteries with Nafion 117 membranes in the Na\(^{+}\)-form intercalated by mixtures of non-aqueous organic solvents used both as an electrolyte, separator, and binder were investigated. Electrochemical stability of various organic solvent mixtures based on \(N,N\)-dimethylacetamide, ethylene carbonate, propylene carbonate, and tetrahydrofuran were characterized. It was shown that a sodium battery based on a Nafion-Na membrane intercalated by mixture of ethylene carbonate and propylene carbonate with a Na\(_3\)V\(_{1.9}\)Fe\(_{0.1}\)(PO\(_4\))\(_3\)/C positive electrode is characterized by a discharge capacity of \(\approx 110 \text{ mAh} \cdot \text{g}^{-1}\) (current density of 10 mA \cdot g\(^{-1}\)) at room temperature and shows the ability to cycle without degradation during 20 cycles. Batteries with Nafion membrane electrolytes, containing \(N,N\)-dimethylacetamide, were characterized using capacity fading during cycling, which is due to the interaction of \(N,N\)-dimethylacetamide and a negative sodium electrode.

Keywords: sodium rechargeable battery; polymer electrolyte; Nafion; cycle stability; electrochemical stability

1. Introduction

Of late, sodium-ion batteries have attracted keen attention [1–3]. In such batteries, the positive and negative electrodes are made of materials capable of reversibly inserting sodium without the release of a free metallic phase. This is what ensures their safe functioning. Sodium rechargeable batteries differ from sodium-ion counterparts in that sodium metal is used as a negative electrode in such batteries. The theoretical specific capacity of metallic sodium during its anodic dissolution amounts to 1165 mAh \cdot g\(^{-1}\), which is much higher than that of intercalation compounds of the negative electrode of the sodium-ion batteries. For example, sodium titanate (Na\(_4\)Ti\(_3\)O\(_7\)), sodiated hard carbon (Na\(_8\)C), sodiated germanium (Na\(_{1.6}\)Ge), and sodiated phosphorus (Na\(_3\)P) have a theoretical specific capacity for sodium extraction of 154, 282, 390, and 804 mAh \cdot g\(^{-1}\), respectively [4]. In addition, the potential of sodium metal at its anodic dissolution is noticeably more negative than that of intercalation electrodes during discharge, which contributes to an increase in the discharge voltage of the battery. At the same time, it is known that metallic sodium (as well as metallic lithium) is not capable of prolonged cycling
in liquid electrolytes due to encapsulation and dendrite formation. The latter can lead to a shorting and ignition of the battery with metallic sodium.

The use of a solid polymer electrolyte, in principle, can create a battery with a metallic sodium negative electrode, the energy density of which will be higher than that of the sodium-ion battery. Another advantage of using a solid polymer electrolyte consists in its flexibility, which makes it possible to obtain a large area of electrolyte, and to design batteries of different geometries. In addition, the use of polymer electrolytes reduces the known effect of the volume changes of electrode material during sodium intercalation/deintercalation [5,6].

To date, quite a lot of work has been published on the creation and investigation of sodium-ion solid polymer electrolytes [7–10]. The most widespread class of polymer electrolytes are true solid polymer electrolytes [7]. They are based on polymer matrices in which sodium salts with a bulky anion are dissolved. In most cases, the polymer basis of such electrolytes is polyethylene oxide [11]. Such electrolytes are characterized by low ionic conductivity at room temperature ($\approx 10^{-5}$ S·cm$^{-1}$) [12]. Replacement of salts with membrane ionogenic groups, such as $-\text{SO}_3^-$, $-\text{COO}^-$, etc., as well as the addition of a low-molecular plasticizer, allows higher conductivity values [9,13]. The most promising polymer electrolytes at present are Nafion-type membranes. Due to the perfluorinated polymer matrix, they have high thermal and electrochemical stability, as well as a high degree of dissociation of functional groups, which ensures high ionic conductivity [14–16].

Initially electrolytes based on the Nafion membrane were proposed for lithium-ion batteries [17–24], and later for their sodium-ionic analogs [8,25,26]. Such electrolytes are obtained by transferring Nafion from a hydrogen to a sodium form (ion exchange in aqueous solutions of sodium salts), followed by removal of water and impregnation with an aprotic solvent. Thus, in fact, they are gel-polymer electrolytes. The principal possibility of creating a sodium battery is mentioned in Reference [25] using the example of the Nafion membrane in sodium form intercalated by a mixture of ethylene carbonate (EC) and propylene carbonate (PC). In that work, Na$_{0.44}$MnO$_2$ was used as the active material of the positive electrode. The conductivity of the resulting electrolyte was 0.35 mS·cm$^{-1}$ at room temperature and 1.52 mS·cm$^{-1}$ at 70 °C. The laboratory cell was tested at a temperature of 45 °C and demonstrated smaller values of the discharge capacity, but better cyclability than that with a liquid electrolyte (1 M NaClO$_4$ in an EC-PC mixture).

Recently, the influence of the nature of organic solvents impregnating the membrane in the sodium form on the ionic conductivity was investigated [26]. A certain correlation (but not proportionality) was found between the solvent uptake and the ionic conductivity. Membranes, impregnated with solvents containing N, N dimethylacetamide (DMA), namely EC-DMA, EC-diethyl carbonate (DEC)-DMA, and PC-DMA-tetrahydrofuran (THF), are characterized by the highest ionic conductivity ($\approx 4$ mS·cm$^{-1}$ at 30 °C).

The purpose of this work was to investigate the possibilities of manufacturing batteries with polymer electrolytes based on Nafion membranes in the Na$^+$-form. It was taken into account that not only their ionic conductivity, but also their electrochemical stability and interaction with electrode materials are important features of polymer electrolytes.

2. Results

2.1. The Composition of the Materials Obtained

The IR spectra of the obtained electrolytes (Figure S1 of the Supplementary Materials) are identical to the spectra presented in the Supplementary of the paper [26]. The presence of organic solvents in membranes was revealed by a set of vibrational frequencies inherent in these solvent molecules. In the frequency range 1600–1800 cm$^{-1}$, the C=O stretching vibrations characteristic of organic carbonates (EC, PC and DEC) were observed; at a frequency of $\approx 1640$ cm$^{-1}$, a band corresponding to the valence vibration of the C=O bond in the DMA molecule could be singled out; the presence of the THF molecule could be detected by the characteristic antisymmetric valence vibration of the ring
characteristic of cyclic ethers at $\approx 906 \text{ cm}^{-1}$. The absence of vibrations of OH groups at $\approx 3490 \text{ cm}^{-1}$ in the IR spectra of the obtained samples indicated that the membranes containing aprotic solvents and their mixtures contained no water or methanol. According to the X-ray diffraction data (Figure S2 of the Supplementary Materials), $\text{Na}_3\text{V}_{1.9}\text{Fe}_{0.1}(\text{PO}_4)_3/C$ is a rhombohedral modification of the NASICON structure ($R\overline{3}c$ space group). The carbon of the composite is X-ray amorphous. According to elemental analysis, the carbon content in $\text{Na}_3\text{V}_{1.9}\text{Fe}_{0.1}(\text{PO}_4)_3/C$ was 8.9% by weight. According to EDX data this composite contains Na, V, P, O, C and Fe. The ratio of the elements Na:V:P:Fe corresponded to the initial load.

2.2. The Electrochemical Stability of Polymer Electrolyte Liquid Phase

Figure 1 shows cyclic voltammograms (CVs) for liquid electrolytes based on different solvent mixtures in the potential range from 0.001 to 4.5 V versus Na/Na$^+$ at a potential scan rate of 10 mV·s$^{-1}$. For all electrolytes, the first two cycles of CV are shown. Attention must be drawn to the difference in scale of the ordinates in Figure 1a,b.

![Figure 1. CVs at aluminum electrode in 1 M NaClO$_4$ in mixtures of solvents: (a) EC-DEC-DMA, EC-DMA, PC-DMA-THF and (b) EC-PC, EC-THF, EC-PC-THF-DMA at scan rate 10 mV·s$^{-1}$. Solvents compositions and cycle numbers are shown at the figures.](image)

The lowest electrochemical stability was noted for 1 M NaClO$_4$ in a three-component mixture of PC-DMA-THF (Figure 1a). In this case, the anode process proceeds at potentials somewhat more positive than 2 V. The nature of the anode process remains uncertain. Undoubtedly, a certain part of the current is spent for anodic dissolution of aluminum with the possible formation of any complex compounds, since after the registration of the CVs on the aluminum electrode, certain signs of local corrosion that were distributed unevenly over the electrode surface were noticeable. At the same time, an appreciable part of the anode charge was spent on oxidizing the components of the electrolyte, most likely DMA. Similar, but less intense, anode processes at potentials more positive than 2 V were observed for all DMA-containing solvents (Figure 1a,b). The cathode process, which took place in a 1 M solution of NaClO$_4$ in a three-component mixture of PC-DMA-THF at potentials more negative than 3 V, refers to the reduction of oxidation products of DMA in the previous anode semi-cycle, since at the CVs taken in such a solution in the potential range from 0 to 2 V, cathode currents of less than 0.002 mA·cm$^{-2}$ were recorded, i.e., it was almost three orders of magnitude smaller than that shown in Figure 1a.

The highest electrochemical stability from all the electrolytes studied refers to 1 M NaClO$_4$ in a four-component EC-PC-THF-DMA solvent, as well as to solutions in a binary EC-PC and EC-THF mixtures (Figure 1b).
The anode processes described above (Figure 1a) are conjugated to a certain passivation of the aluminum electrode. It is possible to trace the passivation of the aluminum electrode in various electrolytes by analyzing Figure 2, which shows CVs for the first six cycles with an increasing range of cycling potentials. As can be seen from Figure 2a, 1 M NaClO₄ in the EC-DMA mixture was susceptible to oxidation at potentials more positive than 2 V and caused passivation of the aluminum electrode during cycling. A noticeable electrolyte oxidation at potentials more positive than 2 V was also noted for 1 M NaClO₄ in mixtures of EC-DEC-DMA (Figure 2b). In the solution of NaClO₄ in the four-component EC-PC-DMA-THF mixture, the rate of anodic oxidation was much lower than in previous electrolytes (Figure 2c), and a gradual passivation of the electrode was also observed here. The smallest passivation of the aluminum electrode was noted for a 1 M NaClO₄ in a mixture of PC-DMA-THF (Figure 2d). When the aluminum electrode was cycled in NaClO₄ solutions in mixtures of EC-DMA, EC-DEC-DMA, and EC-PC-DMA-THF (i.e., in all solutions containing EC), passivation led to gradual inhibition of the anodic process. This did not occur in a solution of NaClO₄ in the PC-DMA-THF mixture (Figure 2d), therefore, when using such a solution, a permanent irreversible capacity loss appeared in the battery, which excludes the use of such a solvent in practical devices.

Figure 2. Aluminum electrode passivation at cyclic polarization in different electrolytes: (a) 1 M NaClO₄ in EC-DMA, (b) 1 M NaClO₄ in EC-DEC-DMA, (c) 1 M NaClO₄ in EC-PC-DMA-THF, and (d) 1 M NaClO₄ in PC-DMA-THF.

The oxidation rate of DMA depends significantly on which mixture it was in (Figures 1 and 2). The oxidation rate (currents at potentials more positive than 2 V) decreased in the series PC-DMA-THF > EC-DMA > EC-DEC-DMA > EC-PC-THF-DMA. This thesis was also confirmed by the results of a cyclic voltammetric study of a 1 M solution of NaClO₄ in plain DMA. The corresponding CVs are shown in Figure 3.
were tested. In the case of using a polymer membrane intercalated by an EC–PC mixture, it was of
interest to compare our results with the results given in Reference [25], where the electrolyte was also
formed. Interestingly, when other solutions containing DMA (solutions of NaClO$_4$ in a mixture of
PC-DMA-THF) come into contact with sodium, the color change happened at an enlarging potential
range. Cycle numbers are shown.

The rate of the anodic process in a solution of NaClO$_4$ in plain DMA was seen to be noticeably
lower than in a 1 M NaClO$_4$ in a three-component mixture PC-DMA-THF. It can be assumed that in
this case it was precisely THF that had a de-passivating effect on the aluminum surface.

When cyclic voltammetric measurements were carried out, one feature of a 1 M solution of sodium per-
chlorate in a mixture of PC-DMA-THF solvents was disclosed. When this solution was brought
into contact with the sodium electrode, the solution became red-brown, and the staining intensity
increased with time. With enough contact time, the solution viscosity increased up to solidification.

It was found by separate experiments that the active component of the solvent in this case was DMA.
According to References [27–29], we can suppose that some complex between DMA and Na$^+$-ion
was formed. Interestingly, when other solutions containing DMA (solutions of NaClO$_4$ in mixtures
EC-DMA, EC-DEC-DMA, and EC-PC-DMA-THF) come into contact with sodium, the color change
of the solution was much slower. This phenomenon is most likely explained by the fact that all
these solutions contained EC, and when sodium was in contact with such solutions, a fairly strong
passive film (solid electrolyte interphase—SEI) formed on its surface, which significantly inhibited
the interaction of sodium with the solution. A separate publication will be devoted to the mechanism
of chemical interaction of sodium with DMA with a detailed analysis of the products formed.

2.3. The Rechargeable Battery with a Polymer Electrolyte

As was already mentioned above, the ionic conductivity of the polymer electrolyte was an
important but not the determining parameter in deciding the applicability of the electrolyte in
sodium batteries. Also, it was essential to operate the positive electrode in contact with a certain
polymer electrolyte. Finally, this issue could be solved only by testing the real batteries. In this
work, coin cells with polymer membranes intercalated by solvents EC-PC and EC-PC-THF-DMA
were tested. In the case of using a polymer membrane intercalated by an EC–PC mixture, it was of
interest to compare our results with the results given in Reference [25], where the electrolyte was also
a Nafion membrane in sodium form intercalated by an EC-PC mixture, and Na$_{0.44}$MnO$_2$ was used
in a positive electrode. It was found via separate experiments that the membrane intercalated by the
four-component mixture of EC-PC-THF-DMA had almost an order of magnitude higher electrical
conductivity (more than 4 mS·cm$^{-1}$ at 30 °C) compared to the membrane intercalated by EC-PC
($\approx$0.7 mS·cm$^{-1}$ at 30 °C) [26], with close electrochemical stability (Table 1).

<table>
<thead>
<tr>
<th>Electrolyte</th>
<th>Thickness, μm</th>
<th>Solvent Uptake (n(Solvent/n(SO$_3^-$)))</th>
<th>Conductivity at 30 °C, mS·cm$^{-1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nafion 117-EC-PC</td>
<td>240</td>
<td>19.8</td>
<td>0.7</td>
</tr>
<tr>
<td>Nafion 117-EC-PC-THF-DMA</td>
<td>260</td>
<td>24.3</td>
<td>4.1</td>
</tr>
</tbody>
</table>

![Figure 3. CVs in 1 M NaClO$_4$ in DMA at enlarging potential range. Cycle numbers are shown.](image-url)
Besides, the selected membranes were characterized by a minimum decrease in conductivity with decreasing temperature.

Figure 4 shows the charge–discharge curves for coin cells with polymer electrolytes based on a Nafion 117 membrane intercalated by EC-PC (Figure 4a) and EC-PC-DMA-THF (Figure 4b) mixtures. The current density was 10 mA·g⁻¹ of active cathode material (Na₃V₁.₉Fe₀.₁(PO₄)₃/C). For comparison, the similar coin cell with a liquid electrolyte (1 M NaClO₄ in a mixture of EC-PC) was also tested. The positive electrode in such a cell was made of Na₃V₁.₉Fe₀.₁(PO₄)₃/C with a polyvinylidene difluoride binder.

The charge–discharge curves of coin cells with polymer electrolyte were generally similar to the charge–discharge curves of a battery with a liquid electrolyte, namely, the charge proceeded at a voltage of about 3.5 V and discharged at a voltage of about 3.3 V.

A slight difference appeared in the shape of a charging curve on which for a cell with a polymer electrolyte intercalated by the EC-PC mixture, the second plateau was recorded at a voltage higher than 3.9 V. The nature of this plateau was not clear, but we can suppose that its appearance may be due to the difficulty of extracting the last portions of sodium from Na₃V₁.₉Fe₀.₁(PO₄)₃/C in contact with the polymer electrolyte and consequently an increase in concentration polarization.

Comparison of the charge–discharge curves of three cells (Figure 5) shows that when cycling in such a low rate (10 mA·g⁻¹), the difference in the voltage of the cells did not exceed the experimental discrepancy. This was not surprising. The ohmic voltage drop in the polymer electrolyte with thickness is 0.4 mm, at a current density of 10 mA·g⁻¹ was estimated as 8 mV for a cell with a polymer electrolyte based on an EC-PC mixture and about 1 mV for a cell with a polymer electrolyte intercalated by a four-component solution.

Figure 4. Charge–discharge curves of coin cells with polymer electrolytes based on Nafion 117, intercalated by EC-PC (a) and EC-PC-DMA-THF (b) mixtures. Current density of 10 mA·g⁻¹.

Figure 5. Charge–discharge curves (a) and cycling stability (b) for coin cells with liquid (1) and polymer electrolytes (2,3). (1)—1 M NaClO₄ in EC-PC, (2)—Nafion 117 membrane, intercalated by EC-PC mixture, (3)—Nafion 117 membrane, intercalated by a EC-PC-DMA-THF mixture. The capacity was normalized by a weight of cathode active material current density of 10 mA·g⁻¹.
3. Discussions

It is worth noting that a battery with a polymer electrolyte based on a Nafion 117 membrane intercalated by an EC-PC mixture is fully functional. From the battery described in Reference [25], it is advantageously distinguished by the practical constancy of the discharge voltage, which is explained by the features of the positive electrode from sodium vanadium-iron-phosphate. Besides, in the battery in the present work, a polymer electrolyte was used as a binder of the positive electrode, which ensured a more complete utilization of the entire volume of the active material. The initial capacity of a battery with a polymer electrolyte based on a Nafion 117 membrane intercalated by an EC-PC mixture was slightly lower than the corresponding value for a liquid electrolyte battery. However, with further cycling, almost identical values of the discharge capacity were recorded. The specific capacity of the electrode from the composite of sodium vanadium-iron-phosphate with carbon, registered in this work when cycling a battery with polymer electrolyte on the basis of a mixture of EC-PC at room temperature at current density of 10 mA·g⁻¹ was about 100 mAh·g⁻¹. Taking into account the content of carbon (8.9 wt%) in the Na₃V₁.₉Fe₀.₁(PO₄)₃/C composite, the discharge capacity was close to the theoretical value. Figure 5a,b shows the data on coin cells with liquid and polymer electrolytes cycling.

The discharge capacity of a coin cell with a polymer electrolyte based on the Nafion117 membrane intercalated by the EC-PC-DMA-THF mixture at the initial cycles was slightly higher than that of the cells with a liquid electrolyte and EC-PC-based polymer electrolyte. However, with a further cycling for the coin cell with the Nafion 117 membrane intercalated by the EC-PC-DMA-THF mixture, a noticeable capacity fading was recorded, which was due to the chemical interaction of DMA and metallic sodium. The stable operation of a battery with a polymer electrolyte based on the Nafion 117 membrane intercalated by a EC-PC mixture for 20 cycles demonstrated the principal possibility of manufacturing a sodium battery with such a polymer electrolyte and a positive electrode based on Na₃V₁.₉Fe₀.₁(PO₄)₃/C, capable of cycling without passivation and excessive degradation. It is worth noting that one of the important challenges in the design of a solid electrolyte battery is to ensure a good contact at the electrode/electrolyte interface. In the case of inorganic solid electrolytes, this is achieved by making composite electrodes (active material/electrolyte/carbon) followed by high-temperature sintering [30–32]. However, this does not completely solve the problem in the case of a thick layer of electrode material and with multiple intercalation/deintercalation, which is accompanied by a change in volume. The use of a polymeric binder and polymer electrolyte promotes good cycling by reducing the effect of changes in the volume of electrode material during intercalation/deintercalation. However, in the case of a non-conductive binder, the capacity of such cells is inferior to that of similar cells with liquid electrolyte. Thus, the initial capacity of the electrode based on Na₀.₄₄MnO₂ at 45 °C in the C/20 mode was inferior to the values obtained with the use of a liquid electrolyte and did not exceed 60 mAh·g⁻¹. In this work, the perfluorinated membrane in the Na⁺ form was used both as an electrolyte and as a binder. As a result, the capacity of the battery with polymer electrolyte based on the EC-PC mixture is comparable with the corresponding value for a battery with liquid electrolyte. The obtained result indicates a good contact of the polymer electrolyte and particles of the electrode material over the entire thickness of the electrode.

The theoretical energy density (based on the weight of active materials only) of sodium battery with a polymer electrolyte and a positive electrode based on the Na₃V₁.₉Fe₀.₁(PO₄)₃/C composite amounts to 360 Wh·kg⁻¹. With due account for the weight of structural materials and polymer electrolyte of smaller thickness (less than 0.1 mm), one can expect to achieve a practical energy density of 180–200 Wh·kg⁻¹, which seems quite optimistic. The use of positive electrodes in a sodium battery with an active material having a higher specific capacity than Na₃V₁.₉Fe₀.₁(PO₄)₃ will result in a corresponding increase in the energy density.
4. Materials and Methods

4.1. Preparation of Electrolytes Based on a Nafion Membrane

The membranes were made by the same procedure as described in Reference [26]. The commercial Nafion 117 membrane manufactured by DuPont Co. (Wilmington, DE, USA) was conditioned according to scheme [33], then it was kept for two days in a 2 M aqueous NaCl solution with constant stirring. The membrane was then washed several times with deionized water until the chloride ions were completely removed and dried under vacuum at a temperature of 70 °C for 12 h. After that, the membrane was aged for 6 h in methanol at a temperature of 60 °C and again dried under vacuum. Such treatment with methanol contributed to an appreciable expansion of the pores in the membrane. The final operation was the soaking of membranes in mixed solvents. This operation was carried out in a glove box with an argon atmosphere containing less than 1 ppm of oxygen and water vapor. In the present study, the following mixtures of aprotic solvents containing equal volumes of components have been studied: EC-DMA, EC-PC, EC-THF, EC-DEC-DMA, PC-DMA-THF, and EC-PC-THF-DMA.

4.2. The Binder Preparation

A membrane-based binder was prepared by the following procedure: a membrane was casted from a 5 wt% solution of a Nafion in a proton form in a mixture of lower aliphatic alcohols and water (55:45). Then this membrane was transferred to the Na+ form by ion exchange in a manner like the one described above. The resulting Nafion-Na membrane was dried in vacuum at 70 °C. The dry polymer was placed in the calculated amount of dimethylformamide (DMFA) to prepare a 10 wt% solution. Dissolution took place at room temperature for 1 day with constant stirring.

4.3. The Cathode Material Synthesis

The Na$_3$V$_{1.9}$Fe$_{0.1}$(PO$_4$)$_3$/C nanocomposite was synthesized by the modified Pechini method according to the procedure described in References [34,35] using V$_2$O$_5$, Fe(NO$_3$)$_3$·9H$_2$O, NaNO$_3$, NH$_4$H$_2$PO$_4$, oxalic acid (a chelating agent and a reducer), ethylene glycol, and citric acid. The polycondensation of ethylene glycol and citric acid led to the formation of a polymer matrix that inhibited the growth of the composite particles.

4.4. The Materials Characterization

The infrared (IR) spectra of the obtained electrolytes based on Nafion membranes were recorded with a Nicolet iS5 IR spectrometer with a SpecacQuest attachment (Thermo Fisher Scientific, Waltham, MA, USA) in the attenuated total reflection mode with a diamond crystal in the wavenumber range of 500–4000 cm$^{-1}$. The phase composition of the synthesized cathode material was characterized using a Rigaku D/MAX diffractometer (Rigaku Corp., Tokyo, Japan), CuK$_\alpha$ radiation in the range of 10–60° 2θ. The carbon content of the composite was determined using a CHNS analyzer EuroEA 3000 (Evro Vector S.r.l., Pavia, Italy). The chemical composition of the cathode material was determined using a scanning electron microscope Carl Zeiss NVision 40 (Zeiss AG, Oberkochen, Germany) equipped with an energy dispersive X-ray (EDX) analyzer Oxford X-Max. Ionic conductivity was measured by impedance spectroscopy using an Elins Z-1500J AC bridge (1 kHz–2 MHz) (Elins Co., Chernogolovka, Russia) on symmetric carbon/membrane/carbon cells.

4.5. The Electrochemical Studies

Cyclic voltammetric measurements in liquid electrolytes, i.e., in 1 M solutions of NaClO$_4$ in the above-mentioned mixed solvents (Section 4.1), were made to assess the electrochemical stability of the liquid phase of polymer electrolytes based on Nafion membranes. The measurements were carried out in three-electrode cells with an aluminum working electrode (foil 50 μm thick, “Rusal” Co., Irkutsk, Russia). The auxiliary and the reference electrodes were made from sodium metal, rolled on a stainless
steel mesh. To register cyclic voltammograms (CVs), a potentiostat-galvanostat R-20X8 from “Elins” (Elins Co., Chernogolovka, Russia) was used. The water content in solvents and electrolyte, determined by coulometric titration using the Karl Fischer method with a coulometric titrator 917 Ti-Touch (Metrohm, Switzerland) did not exceed 20 ppm. The potential scan rate was 10 mV s\(^{-1}\).

To evaluate the feasibility of using the polymer electrolytes under investigation in real sodium-ion batteries, coin cells with a polymer electrolyte were tested. For the sake of comparison, similar cells with liquid electrolyte (1 M solutions of NaClO\(_4\) in the mixture of EC-PC) were tested as well.

For coin cells with a polymer electrolyte, positive electrodes contained 80 wt% Na\(_3\)V\(_{1.9}\)Fe\(_{0.1}\)(PO\(_4\))\(_3\)/C, 10% acetylene black as a conducting additive, and 10% of Nafion-Na dissolved in DMFA as a binder. For coin cells with liquid electrolyte, the positive electrodes contained 80% Na\(_3\)V\(_{1.9}\)Fe\(_{0.1}\)(PO\(_4\))\(_3\)/C, 10% acetylene black, and 10% PVDF dissolved in N-methylpyrrolidone. The electrodes were dried, pressed with a pressure of 1 ton cm\(^{-2}\) and further dried at a temperature of 120 °C for 8 h. The amount of active material on the substrate (Al foil 50 µm thick, “Rusal” Co., Irkutsk, Russia) was about 5 mg·cm\(^{-2}\). The coin cells were of the 2016 type. Cyclic galvanostatic tests were carried out using a computerized charge–discharge measuring and computing complex AZRVRIK 50 mA–10 V (NTT Buster, St. Petersburg, Russia). Coin cells were assembled in a glove box with a dry argon atmosphere (JSC Spectroscopic Systems, Moscow, Russia). The water and oxygen content in the box did not exceed 1 ppm.

5. Conclusions

In the present work, an estimation of the possibility of creating a sodium battery with a polymer electrolyte based on a Nafion membrane intercalated with a mixture of aprotic organic solvents and a positive electrode based on the Na\(_3\)V\(_{1.9}\)Fe\(_{0.1}\)(PO\(_4\))\(_3\)/C composite was made. It has been established that an important factor in the choice of the organic solvent, which impregnates the Nafion membrane in the sodium form, was the electrochemical window. It was shown that sodium batteries with a Nafion membrane electrolyte containing dimethylacetamide were characterized by capacity fading during cycling, which was due to the interaction of DMA and a negative sodium electrode. A suitable electrolyte for a sodium battery, in terms of high conductivity and electrochemical stability, was a Nafion membrane in sodium form intercalated by binary solvent ethylene carbonate-propylene carbonate. A battery with this electrolyte showed a discharge capacity of \(\approx 110\) mAh·g\(^{-1}\) (current density of 10 mA·g\(^{-1}\)) at room temperature and the ability to cycle without degradation during 20 cycles.

Supplementary Materials: The following are available online at http://www.mdpi.com/2313-0105/4/4/61/s1. Figure S1: IR spectra of the obtained electrolytes. Figure S2: The XRD patterns of Na\(_3\)V\(_{1.9}\)Fe\(_{0.1}\)(PO\(_4\))\(_3\)/C.

Author Contributions: Material synthesis and preparation S.N. and D.V.; investigation, D.V., T.K., and A.C.; data curation, A.C.; writing—original draft preparation, S.N. and T.K.; writing—review and editing, A.S. and A.Y.

Funding: This research was funded by Russian Science Foundation, grant number 16-13-00024.

Conflicts of Interest: The authors declare no conflict of interest.

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