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## Manganese electrode for all-solid-state fluoride batteries\*

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We investigate MnF3 as an electrode material for all-solid-state fluoride batteries. The initial discharge capacity due to defluorination was 535 mA h g<sup>-1</sup>. Manganese was confirmed to be reduced and oxidized during charge-discharge measurements. Metallic Mn was also reversibly fluorinated and defluorinated as a starting material.

Fluoride batteries have attracted attention as next-generation batteries because of their high chemical stability and high theoretical energy density due to the flexible choice of electrode materials. Fluoride (F<sup>-</sup>) anions function as charge carriers in fluoride batteries; therefore, various metals are potential candidates for the conversiontype electrodes. In addition, there is a wide variety of solid electrolytes, including PbSnF<sub>4</sub>, BaSnF<sub>4</sub>, BaCaF<sub>4</sub>, and tysonite-type and BaF<sub>2</sub>based fluorites. 1-16 In 2011, A. Reddy et al. reported the first all-solidstate fluoride battery, and since then, there have been many recent reports of all-solid-state fluoride batteries. 17-32 We recently reported the application of a FeF3 cathode for fluoride batteries, where a large discharge capacity of 579 mA h g<sup>-1</sup> was observed at first discharge.<sup>29</sup> In addition, many groups reported CuF2 as a cathode material for fluoride batteries. 18,27 As such, transition metal fluorides have been shown to be promising cathode materials for large-capacity batteries. Among the possible electrode materials for fluoride batteries, MnF<sub>3</sub> has advantages of a large theoretical capacity (718 mA h  $g^{-1}$ ) and low cost. The theoretical gravimetric energy density for a full cell composed of a MnF<sub>3</sub> cathode and a Mg anode is 1127 W h kg<sup>-1</sup>, which is comparable with that for a FeF3 cathode. However, in the case of Li<sup>+</sup>-ion batteries, manganese fluoride shows a large overpotential.<sup>28</sup> On the other hand, the following simple reaction may occur in the fluoride battery:

$$MnF_3 + 3e^- \rightleftharpoons Mn + 3F^-$$
 (1)

This reaction does not form the LiF insulator, which is typically formed in lithium-conversion type batteries when metal fluorides are used as an electrode material. Therefore, the use of MnF3 as an electrode in fluoride batteries is considered to be reasonable with respect to advantages such as the low cost and large capacity. Here we report all-solidstate fluoride batteries based on a MnF3 electrode with Ba<sub>0.6</sub>La<sub>0.4</sub>F<sub>2.4</sub> (BLF) used as the electrolyte and a Pb-based counter electrode.

Fig. S1 (ESI†) show SEM images of the MnF<sub>3</sub> powder before and after ball-milling. Although the particle size before ballmilling was 100 nm-3 µm, that after mechanical ball-milling at 600 rpm for 12 h was more uniform at 100 nm-1 μm. The crystallinity of the MnF3 powder was significantly decreased after ball-milling (Fig. S2, ESI†). The electrochemical performance was evaluated at 433 K because of the large IR drop in the solid BLF electrolyte at room temperature. Considering high-temperature operation, the chemical stability of MnF<sub>3</sub> was evaluated at high temperatures. XRD patterns for MnF3 after heating at various temperatures under Ar are shown in Fig. S3 (ESI†). Peaks assigned to MnF3 were observed up to 433 K; however, MnF<sub>3</sub> was decomposed at 473 K and peaks assigned to MnF2 were newly formed. On the other hand, FeF3 is stable even at 473 K, as we previously reported.<sup>29</sup> Therefore, FeF<sub>3</sub> is more stable than MnF<sub>3</sub> at high temperature. Fig. 1(a) shows SEM images of the MnF3-based composite powder (MnF<sub>3</sub>, BLF, acetylene black(AB)). The primary BLF solid electrolyte particles were smaller than 1 µm and secondary particles were larger than 10 µm, which is good agreement with our previous study.<sup>29</sup> Fig. 1(b) and (c) show cross-sectional images of the BLF/MnF<sub>3</sub> electrode interface after heating at 433 K. Fig. 1(b) shows both a highly connected interface region and a region where separation has occurred. Fig. S4 (ESI†) shows a cross-sectional SEM image of the MnF3 electrode/BLF electrolyte interface after heating at 433 K. The thickness of the MnF<sub>3</sub> electrode was 88 μm. Fig. 2(a) shows discharge-charge curves for an all-solid-state fluoride battery with a Pb/PbF2-SnF2-AB/ BLF/MnF<sub>3</sub>-BLF-AB structure between -2 V and 2 V at 433 K

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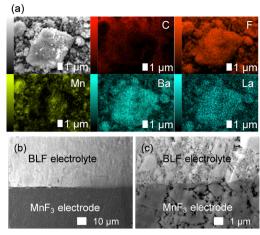


Fig. 1 SEM images of MnF<sub>3</sub>-based composite electrode and electrode/ electrolyte interface for fluoride battery cell. (a) SEM image and energy dispersive spectroscopy (EDS) elemental maps of as-synthesized electrode powder. (b) and (c) Cross sectional images of BLF/MnF<sub>3</sub> electrode interface after heating at 433 K ( $1000 \times$  and  $10000 \times$  magnification).

(defluorination of MnF<sub>3</sub>). The initial discharge capacity (starting from defluorination) was 535 mA h g<sup>-1</sup>, which was 75% of the theoretical capacity for MnF<sub>3</sub> (718 mA h g<sup>-1</sup>). Therefore, 2.2 F ions were shuttled from the MnF3 electrode during the initial discharge. This indicates that most of the MnF3 is decomposed to MnF<sub>2</sub> before discharge. The observed capacity is larger than that reported for BiF3 or CuF2 electrodes at 413 K in an all-solid-state fluoride battery. 17,27 However, this is still lower than that for an FeF<sub>3</sub> electrode (579 mA h g<sup>-1</sup> at first cycle), which was reported in our previous study.<sup>29</sup> The observed single potential plateau is in good agreement with the theoretical redox potential for MnF<sub>2</sub>/Mn (-0.991 vs. PbF<sub>2</sub>/Pb) and also

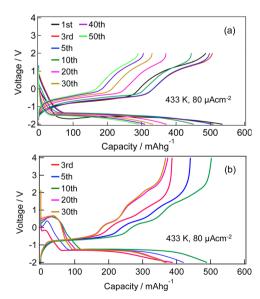
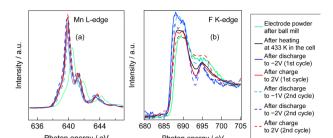


Fig. 2 Charge-discharge profiles for all-solid-state fluoride battery with  $MnF_3$  electrode in ranges of (a) -2 V to 2 V, and (b) -2 V to 4 V (starting from defluorination of MnF<sub>3</sub>). Cell structure is Pb/PbF<sub>2</sub>-SnF<sub>2</sub>-AB/BLF/ MnF<sub>3</sub>-BLF-AB. The capacity was calculated based on the weight of MnF<sub>3</sub>.

indicates that the initial state before discharge is mainly MnF<sub>2</sub>. A discharge capacity of 290 mA h g<sup>-1</sup> was retained at the 50th cycle. The open-circuit voltage gradually increased with cycling, which indicates that the oxidation state of manganese is gradually increased, but is mainly charged to the Mn<sup>2+</sup> state. Fig. 2(b) shows discharge-charge curves for the all-solid-state fluoride battery with a MnF<sub>3</sub> electrode in the range of -2 V to 4 V at 433 K (starting from defluorination). The cell structure is the same as that in Fig. 2(a). Unlike the case for -2 V to 2 V (Fig. 2(a)), a discharge plateau is observed between 0 and 1 V and a charge plateau between 1 and 3 V. The theoretical potentials for the MnF<sub>3</sub>/MnF<sub>2</sub> and MnF<sub>2</sub>/Mn couples are 1.217 V (vs. PbF<sub>2</sub>/Pb) and -0.991 V (vs. PbF<sub>2</sub>/Pb), respectively. The observed potential plateaus are reasonably matched with these theoretical potentials, as shown in Fig. S5 (ESI†), which indicates that manganese is reduced and oxidized. By comparison with Fig. 2(a), oxidation of Mn<sup>3+</sup>/Mn<sup>2+</sup> should occur between 1 V and 2 V, and also above 2 V. The potential and capacity of the Mn3+/Mn2+ region gradually increased with cycling, which may be due to stabilization of the Mn<sup>3+</sup> species. The discharge capacity of 489 mA h  $g^{-1}$  at the 10th cycle is larger than that for FeF<sub>3</sub> (461 mA h g<sup>-1</sup> at the 10th cycle) within the same potential range.29 On the other hand, the hysteresis of the discharge-charge curve was dependent on the voltage range. The voltage difference for the 10th discharge-charge cycle at 350 mA h  $g^{-1}$  (see the point in Fig. S6, ESI†) is 0.945 V between -2 V and 2 V and 0.641 V between -2 V and 4 V. The decrease in overpotential after charging to 4 V may be related to the formation of Mn<sup>3+</sup> species; however, this has yet to be verified. On the other hand, the voltage difference for the 10th discharge-charge cycle at 350 mA h g<sup>-1</sup> for the FeF<sub>3</sub> and MnF<sub>3</sub> electrodes are 0.753 V and 0.641 V, respectively. Therefore, the overpotential of the MnF3 electrode is much lower than that of the FeF3 electrode when the voltage range of the chargedischarge measurement is between -2 V and 4 V. The IR drop for the BLF solid electrolyte is 0.06 V during operation (433 K and 80 μA cm<sup>-2</sup>); therefore, the internal resistance between 2 V and 4 V, excluding the IR drop, is 0.58 V. This is much lower than that for a manganese fluoride electrode for a lithium battery.<sup>33</sup> These results suggest that the charge transfer resistance differs significantly depending on the discharge-charge mechanism, although the operating temperature is different. In the case of lithium battery, insulating LiF is formed during conversion reaction. The effect of operating temperature on the all-solid-state fluoride battery with a MnF3 electrode was investigated (Fig. S7, ESI†). The cell was heated once to 433 K, then cooled to each respective temperature for measurement (413 K and 393 K). The initial defluorination capacities at 413 K and 393 K were 215 mAhg<sup>-1</sup> defluorination capacities at 413 K and 393 K were 215 mA h  $g^{-1}$  and 41 mA h  $g^{-1}$ , respectively, showing a substantial decrease compared to the performance at 433 K. A significant factor contributing to this high resistance is the decreased ionic conductivity of the electrolyte, indicating that applying a solid electrolyte with high conductivity at lower temperatures is essential for reducing the operating temperature. Ex situ XAS measurements were conducted to clarify the

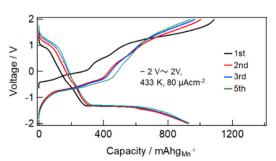


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Fig. 3 (a) Mn L-edge and (b) F K-edge XAS spectra of MnF<sub>3</sub> electrode obtained from current collector side.

discharge-charge mechanism for the MnF<sub>3</sub> electrode. Fig. 3(a) and (b) show Mn L-edge and F K-edge X-ray adsorption spectra of the MnF<sub>3</sub> electrode before and after the discharge-charge measurements. Peaks associated with L3 and L2 can be observed in this region of the Mn L-edge spectra.<sup>34</sup> Following heating of the constructed cell, the XAS peak positions shifted to lower energy. This indicates that MnF<sub>3</sub> is decomposed or reacts with the BLF electrolyte or AB in the composite electrode due to the high temperature or high pressure during construction of the cell. Therefore, the MnF3 composite electrode is not sufficiently stable in this structure. The adsorption peak at around 640 eV associated with MnF3 decreased after discharging and was then recovered after charging. Therefore, the Mn-based species is active for the redox reaction. As shown in Fig. S8 (ESI†), The F K-edge XAS spectra for the MnF3 electrode charged at different voltages (yielding MnF3 and MnF2) revealed the formation of Mn3+ by the peak at around 684 V observed after charging to 4 V, which is in contrast to the electrode charged to 2 V (no adsorption due to MnF2 was observed around this region, Fig. S8, ESI†), i.e., Mn3+ is formed only after charging to 4 V. We have reported that the BLF of the solid electrolyte shows two peaks around 689 eV in our previous work.<sup>29</sup> In the charged state of manganese fluoride electrode (Fig. 3(b)), contributions from the fluorine in manganese fluoride overlap, so clearly split peaks assigned to BLF electrolyte are not observed. On the other hand, after discharge in the second cycle, the spectrum is very similar to that of BLF, suggesting that metallic Mn is formed. From these results, the changes in the F K-edge spectrum indicate that a reversible redox reaction is occurring between manganese fluoride and metallic manganese. The XRD pattern after charging to 2 V did not indicate any formation of manganese fluoride; the species formed was thus amorphous (Fig. S9, ESI†). Fig. S10 (ESI†) shows SEM images of the MnF<sub>3</sub> electrode before and after 58 cycles between -2 V and 2 V. A comparison of these figures indicates that heating at 433 K produced a denser morphology. In addition, the morphology became denser after discharge-charge cycling, which may be due to long-term heating and pressing. EDS maps of the MnF<sub>3</sub> electrode after discharge-charge cycling are shown in Fig. S11 (ESI†). The particle size for the active manganese material is ca. 30 μm, which is much larger than that for the pristine material (ca. 1 µm, see Fig. 1). Therefore, degradation of the active electrode material is mainly caused by agglomeration during discharge-charge cycling. When the counter electrode for the

full cell is a fluoride, the manganese electrode must accept fluoride ions during the first charge process. Therefore, charge-discharge cycling was also evaluated from the fluorination of metallic Mn. The metallic manganese powder consisted of particles with two different sizes, small (<1 μm) and large (ca. 7 μm) as shown in Fig. S12 (ESI†). Fig. 4 shows discharge-charge curves for the all-solidstate fluoride battery with a metallic Mn electrode between -2 V and 2 V at 433 K. The cell had a Pb/PbF2-SnF2-AB/BLF/Mn-BLF-AB structure. The initial charge capacity (fluorination of the cathode) was 1087 mA h g<sup>-1</sup>, which was 74% of the theoretical capacity (1463 mA h g<sup>-1</sup>) for the Mn electrode. The first discharge (fluorination) capacity was 931 mA h g<sup>-1</sup>, which was 63% of the theoretical capacity. This indicates slightly lower utilization efficiency compared with that starting from defluorination (75%, see Fig. 2(a)), although the difference is not significant. The opencircuit voltage before the initial charge was -0.648 V, which was higher than the theoretical potential for a MnF<sub>2</sub>/Mn couple  $(-0.991 \text{ V } \text{vs. PbF}_2/\text{Pb})$ . This is considered to be due to the oxidized layer formed on the metallic Mn. The second charge started from around -2 V, which indicates that the oxidized layer was removed during the initial charge and discharge process. This cell was prepared on the same day as the electrode powder. However, after the powder was stored for 11 days in a glovebox, the first defluorination capacity decreased to 690 mA h g<sup>-1</sup>. This indicates that metallic manganese is highly sensitive and the oxidation layer decreases the reversible capacity. In order to compare the difference in profiles between the cases from the MnF3 and Mn, chargedischarge profiles for all-solid-state fluoride battery with MnF<sub>3</sub> electrode and Mn electrode are shown in Fig. S13 (ESI†). During the initial defluorination, starting from Mn, the overall potential is high. This is influenced by the surface oxidation layer. However, from the second fluorination cycle onward, the fluorination curves show a profile similar to that of electrodes starting from MnF<sub>3</sub>, while only the defluorination curve for the electrode starting from MnF<sub>3</sub> showed a short plateau for the Mn<sup>3+</sup>/Mn<sup>2+</sup> redox couple appeared, even though the voltage range was same. This could be due to the lower charge transfer resistance due to the high electrical conductivity of metallic Mn as a starting material. These results suggest that a manganese electrode is a promising active material for both directions, i.e., starting from defluorination or



fluorination.

Fig. 4 Charge-discharge profiles for all-solid-state fluoride battery with metallic Mn electrode in ranges of -2 V to 2 V (starting from fluorination of metallic Mn). Cell structure is Pb/PbF2-SnF2-AB/BLF/Mn-BLF-AB. The capacity was calculated based on the weight of Mn

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In summary, the electrochemical reversibility of MnF<sub>3</sub> and metallic Mn was demonstrated in an all-solid-state fluoride battery. The MnF<sub>3</sub> electrode exhibited a defluorination capacity of 535 mA h g<sup>-1</sup> during the initial cycle and retained a defluorination capacity of 290 mA h g<sup>-1</sup> at the 50th cycle. The XAS results suggested that manganese was reduced and oxidized during the discharge-charge process. Metallic Mn was also reversibly fluorinated and defluorinated. A manganesebased electrode is thus a promising active material for fluoride batteries with a large energy storage capacity.

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### Data availability

The data supporting this article have been included as part of the ESL†

### Conflicts of interest

There are no conflicts to declare.

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