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Oxalate-assisted Fe₂O₃ surface functionalization of nanosized MgMn₂O₄ and α -MnO₂ cathodes for rechargeable magnesium batteries†

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Mn-based transition metal oxide nanoparticles are promising candidates as cathode active materials for rechargeable magnesium batteries, but their high catalytic activity for oxidative electrolyte decomposition and large surface area deteriorate their cycle performance. A recent study [Yagi et al., J. Mater. Chem. A, 2021, 9, 26401-26409] demonstrated that the catalytic activity was less prominent in Fe-based oxides than in other transition metal oxides, containing Mn. Fe-based oxides show low catalytic activity for oxidative electrolyte decomposition compared with Mn-based congeners. The strong capability of oxalate ions for bridging transition metal ions was utilised to form thin, uniform Fe₂O₃ layers on nanoparticles of MgMn₂O₄ and α -MnO₂. The resulting Fe₂O₃ layers effectively suppressed side reactions during insertion and extraction of the Mg²⁺ ions and improved the capacity retention and cycle performance.

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Introduction

Rechargeable magnesium batteries (RMBs) have attracted increasing attention as advanced rechargeable batteries, because magnesium is an abundant element and magnesium metal anodes have high theoretical capacities (volumetric: 3833 mA h cm⁻³, gravimetric: 2205 mA h g⁻¹). Promising cathode active materials for 3 V-class RMBs include hollandite-type $\alpha\text{-MnO}_2,^{1-4}$ spinel-type MgM $_2\text{O}_4$ (M: transition metal), $^{5-7}$ and ZnMnO $_3.$ $^{8-10}$ Electrolytes compatible with both magnesium metal anodes and transition metal oxide cathodes are presently limited to glyme-based electrolytes.9 Despite the high catalytic activity of transition metal oxide cathodes containing Mn or Co, the oxidative decomposition of the glyme-based electrolytes on their surfaces during the extraction of Mg2+ ions is a crucial problem.11,12 Such side reactions can be suppressed by passivating the surface active sites, where coating the surface of the cathode active materials with inert oxides (e.g., V₂O₅ (ref. 13) or ZrO₂ (ref. 14)) or electroconductive polymers^{15,16} has been reported to

The diffusion of divalent Mg²⁺ ions in the transition metal oxides is sluggish because of the strong electrostatic interactions between the Mg²⁺ ions and the oxide sublattice. To minimize the diffusion length of Mg²⁺ ions and facilitate their insertion and extraction, oxide-based RMB cathode materials with nanosized dimensions and large surface areas, such as 3D-open channel nanostructures (structured MgMn₂- O_4 (ref. 18)) and ultrasmall (<2.5 nm) cubic MgMn₂ O_4 , have been utilized. Surface functionalization via self-organization of reagents on the surface of transition metal oxides is an ideal way to form thin, uniform, and dense layers on transition metal oxides with large surface areas. As an example of such a self-organizing process, we developed the phenyl phosphonate surface functionalization of structured MgMn₂O₄ by utilizing the strong binding of the phenylphosphonate groups to the surface of transition metal oxides.19

Herein, we present another self-organizing process to form a thin Fe₂O₃ layer on nanosized transition metal oxide-based RMB cathode materials by employing oxalate ions, which are small polydentate ligands commonly used to form uniform

improve their electrochemical properties. On the other hand, recent studies have shown that oxidative electrolyte decomposition is less prominent on MgFe₂O₄ than on MgMn₂O₄ and MgCo₂O₄, indicating that the catalytic activity of Fe for oxidative electrolyte decomposition is lower than that of Mn and Co.11 The partial substitution of Fe into Mn and Co sites (e.g., Mg(Co_{0.4}Fe_{0.6})O₄ (ref. 12)) and coating of the surfaces of MgMn₂O₄ with Mg-Fe binary oxides¹⁷ also improved the electrochemical properties.

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precipitates of multicomponent metal ions for the precursors of multicomponent ceramics. 20,21 The cathode active materials functionalized with oxalate ions were formed by treating in a solution of oxalate ions. They were converted to ones functionalized with Fe₂O₃ through treatment in a solution of iron(III) ions and subsequent heat treatment. The electrochemical properties of the resulting functionalized cathode materials are presented.

Experimental

Synthesis of MgMn₂O₄

The structured MgMn₂O₄ powder was prepared by following a reported procedure. 18,19 Magnesium chloride hexahydrate (MgCl₂·6H₂O, 6 mmol, Fujifilm Wako Pure Chemical), manganese chloride tetrahydrate (MnCl₂·4H₂O, 12 mmol, Fujifilm Wako Pure Chemical), and citric acid (18 mmol, Fujifilm Wako Pure Chemical) were dissolved in 20 mL of ethanol, and propylene oxide (16 mL, Kanto Chemical) was added. The resulting metal-organic complex gel was maintained for 1 day at 25 °C, washed with ethanol and acetone to remove byproducts, and subjected to sequential solvent exchange with acetone for 1 day and cyclohexane 3 times in 3 days. The resulting wet gel was dried at 60 °C and heat treated at 350 °C for 5 h in a tube furnace in air.

Synthesis of α-MnO₂

The α-MnO₂ powder was prepared by following a reported procedure. 22-24 Manganese sulfate pentahydrate (MnSO₄·5H₂-O, 67 mmol, Fujifilm Wako Pure Chemical), ammonium sulfate ((NH₄)₂SO₄, 125 mmol, Fujifilm Wako Pure Chemical), and ammonium peroxydisulfate ((NH₄)₂S₂O₈, 67 mmol, Fujifilm Wako Pure Chemical) were dissolved in 100 mL of distilled water. The solution was then transferred to a Teflonlined stainless-steel autoclave and heated at 140 °C for 12 h in an oven. The product was washed with an acetonitrile solution of nitronium tetrafluoroborate (NO2BF4, Fujifilm Wako Pure Chemical).

Fe₂O₃ functionalization

Ammonium oxalate monohydrate ((NH₄)₂ox·H₂O, 0.25 mmol, Fujifilm Wako Pure Chemical) was dissolved in 3 g of distilled water. MgMn₂O₄ or α-MnO₂ (0.625 mmol) was added to the solution, and the suspension was stirred at room temperature for 3 h. The solid powder was separated by centrifugation, washed with water, and dried at 60 °C for 12 h. The dried solid powder was redispersed in a solution prepared by dissolving iron(III) nitrate nonahydrate (Fe(NO₃)₃·9H₂O, Fujifilm Wako Pure Chemical) in 3 g of methanol (Fujifilm Wako Pure Chemical) at a molar ratio of $MgMn_2O_4$ or α - MnO_2 : $Fe(NO_3)_3 = 1:x$, and the suspension was stirred at room temperature for 3 h. The solid powder was separated by centrifugation, washed with methanol, dried at 60 °C for 12 h, and heat treated at 350 °C for 5 h in a tube furnace in air.

Characterization

The resulting powder samples were evaluated by powder X-ray diffraction (XRD, SmartLab, Rigaku), Fouriertransform infrared (FT-IR) spectrometry (FT/IR-4600, JASCO) using an attenuated total reflection (ATR) unit with a diamond prism, scanning electron microscopy with energydispersive X-ray spectroscopy (SEM-EDS, PhenomPro, Thermo Fisher Scientific), and scanning transmission electron microscopy with EDS (STEM-EDS, JEM-ARM200F NEOARM, JEOL).

Electrochemical analysis

Dry composite cathodes were prepared by mixing the powder of pristine or Fe₂O₃-functionalized MgMn₂O₄ or α-MnO₂, acetylene black (AB, Denka; electrically conductive support), poly(tetrafluoroethylene) (PTFE, Du Pont-Mitsui Fluorochemicals; binder) in a mass ratio of 60:30:10, and pressing ~2 mg of the composite with ~1.2 mg of cathode active material onto an Al mesh. Electrochemical measurements of the composite cathode were conducted in an Ar-filled glovebox with a three-electrode cell (Fig. S1†) using a Mg ribbon (99.9%, Yoneyama Yakuhin Kogyo) as the counter electrode, and a Ag wire immersed in a triglyme (G3, Kanto Chemical) solution of 0.01 mol dm⁻³ AgNO₃ (Kanto Chemical) and 0.1 mol dm⁻³ magnesium bis(trifluoromethanesulfonyl) amide (Mg[TFSA]2, Kishida Chemical) as the reference electrode. The electrolytes used were 0.3 mol dm⁻³ [Mg(G4)] [TFSA]₂/[C₃mPyr][TFSA],^{25,26} prepared from tetraglyme (G4, Chemical), Mg[TFSA]₂, propylpyrrolidinium bis(trifluoromethanesulfonyl)amide ([C3mPyr [TFSA], Tokyo Chemical Industry), and 0.3 mol dm⁻³ G3 solution of magnesium tetrakis(hexafloroisopropyloxy)borate (Mg[B(HFIP)₄]₂).²⁷⁻²⁹ Galvanostatic charge-discharge and galvanostatic intermittent titration technique (GITT) tests were carried out using electrochemical analyzers (HZ-Pro and HJ1020mSD8, Hokuto Denko) at 10 mA g⁻¹ in the potential range from -1.6 to 0.9 V vs. Ag/Ag+ (from 1.0 to 3.5 V vs. Mg/ Mg²⁺). The test was initiated from the discharge step, and the charge capacity was restricted to 180 mA h g⁻¹ for MgMn₂O₄ and 200 mA h g⁻¹ for α-MnO₂ (approximately two-thirds of the theoretical capacities of MgMn₂O₄ (270 mA h g⁻¹)⁵ and α -MnO₂ (308 mA h g⁻¹)³⁰). The rest period of the GITT measurements was 2 h.

Results and discussion

Structural characterization

Fig. 1 shows a schematic illustration of the Fe₂O₃ functionalization process. First, the surface of a cathode active material was modified with oxalate ions by suspending them in an aqueous ammonium oxalate solution. Fe³⁺ ions were anchored to the surfaces of the resulting samples by treatment with a methanol solution of iron(III) nitrate. Finally, oxalate ions and residual nitrate ions were

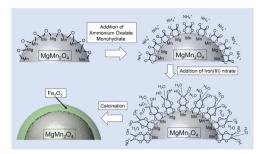


Fig. 1 Schematic illustration of the Fe₂O₃ functionalization process in the oxalate-assisted method.

decomposed by heat treatment to form thin uniform Fe₂O₃ layers on the structured MgMn₂O₄.

Fig. 2(a) shows the powder XRD patterns of the pristine and Fe₂O₃-functionalized MgMn₂O₄. The observed pattern of the pristine sample was essentially identical to those reported previously. 18,19,31,32 The similarity of the patterns before and after Fe₂O₃ functionalization indicated that the growth of MgMn₂O₄ crystallites and other crystalline phases during heat treatment was insignificant. Fig. 2(b) shows the ATR-FT-IR spectra of the MgMn₂O₄ powders during the course of Fe₂O₃ functionalization. The shoulder at ~650 cm⁻¹ was attributed to the Mn-O stretching mode of the MgMn₂O₄.³³ After treatment with the ammonium oxalate solution (green line), absorption bands attributed to the oxalate ions were observed at ~1320, ~1375, and ~1650 cm^{-1} , $^{34-36}$ and the ammonium ions were observed at ~ 1450 cm⁻¹.37 After treatment with the iron(III) nitrate solution (blue line), absorption bands attributed to the nitrate ions

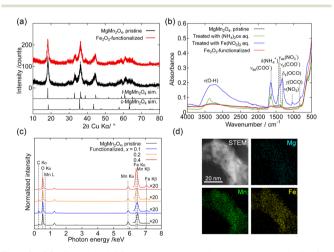


Fig. 2 (a) Powder XRD patterns of the pristine and Fe_2O_3 functionalized structured MgMn₂O₄ powders. Simulated patterns were calculated using RIETAN-FP38 and the structure parameters of MgMn₂O₄ reported in ref. 31. (b) ATR-FT-IR spectra of the pristine structured MgMn₂O₄ powder after treatment with ammonium oxalate or iron(III) nitrate solution, and after heat treatment at 350 °C for 5 h in air (Fe₂O₃-functionalized sample). (c) SEM-EDS spectra of pristine and Fe₂O₃-functionalized structured MgMn₂O₄ powders. (d) STEM and EDS elemental mapping (Mg, Mn, and Fe) images of Fe₂O₃-functionalized structured MgMn₂O₄.

appeared at ~1040 and ~1350 cm⁻¹, 37 where the broad absorption band peaked at ~3300 cm⁻¹, originating from the O-H stretching mode of FeOH groups and adsorbed or coordinated water molecules, became prominent. In the spectrum of the Fe₂O₃-functionalized sample after heat treatment (red line), the absorption bands attributed to the oxalate, nitrate, and OH groups disappeared almost completely. Fig. 2(c) shows the SEM-EDS spectra of the pristine and Fe₂O₃-functionalized MgMn₂O₄. The Fe Kβ line was observed only in the spectrum of Fe₂O₃-functionalized $MgMn_2O_4$, and intensified with increasing x. Fig. 2(d) shows the STEM-EDS elemental mapping images of the Fefunctionalized MgMn₂O₄, verifying the uniform distribution of Fe on the particles.

To demonstrate the versatility of this method, Fe₂O₃ functionalization was applied to α-MnO2. Fig. 3 shows the powder XRD patterns and SEM-EDS spectra of α-MnO₂ before and after Fe₂O₃ functionalization. Fe₂O₃ functionalization preserved the XRD patterns, whereas the Fe K\beta line appeared only in the spectrum of the Fe₂O₃-functionalized sample, similar to the structured MgMn₂O₄ shown in Fig. 2.

Electrochemical characterization

Fig. 4 shows the galvanostatic charge-discharge curves of the dry composite cathodes of the pristine and Fe₂O₃functionalized MgMn₂O₄ powders. The current density was normalized to the mass of MgMn₂O₄ powder, including that of Fe₂O₃. For the pristine sample (Fig. 4(a)), the 1st charge cycle ended at ~2.9 V vs. Mg/Mg²⁺, whereas the charge potential notably increased with cycle number. This potential was probably due to oxidative electrolyte decomposition and the simultaneous increase in the with the accumulation overpotential associated decomposed products on the MgMn₂O₄ surface. Such an increase in the charge overvoltage with the number of cycles was significantly suppressed by surface functionalization with Fe_2O_3 .

The initial discharge capacity was comparable for the pristine sample (~230 mA h g⁻¹) and those functionalized with Fe₂O₃ (\sim 210, \sim 230, and \sim 240 mA h g⁻¹ at x = 0.1, 0.2,

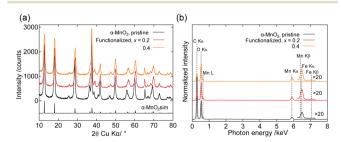


Fig. 3 (a) Powder XRD patterns of pristine and Fe₂O₃-functionalized $\alpha\text{-MnO}_2$ powders. Simulated pattern was calculated using RIETAN-FP, ³⁸ and the structure parameters of α -MnO₂ reported in ref. 39. (b) SEM-EDS spectra of pristine and Fe₂O₃-functionalized α-MnO₂ powders.

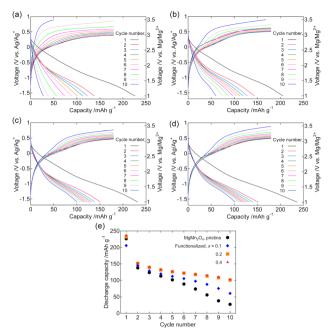


Fig. 4 Galvanostatic charge-discharge curves of dry composite cathodes of pristine MgMn₂O₄ (a) and those functionalized with Fe₂O₃ at x = 0.1 (b), 0.2 (c), and 0.4 (d) in 0.3 mol dm⁻³ [Mg(G4)][TFSA]₂/ [C₃mPyr][TFSA] at 100 °C. (e) Discharge capacity retention of samples shown in panels (a)-(d).

and 0.4, respectively). The decay in the discharge capacity between the 1st and 2nd cycles was mainly due to the incomplete extraction of Mg2+ ions during the 1st charge cycle. In the pristine sample, the discharge capacity rapidly deteriorated with cycle number. This discharge capacity fading was also suppressed by the Fe₂O₃ functionalization. Fig. 4(e) and S2† show the discharge capacity retention and Coulombic efficiency. The discharge capacity after the 10th cycle was \sim 100 mA h g⁻¹ for the samples with x = 0.2 and 0.4, and was notably higher

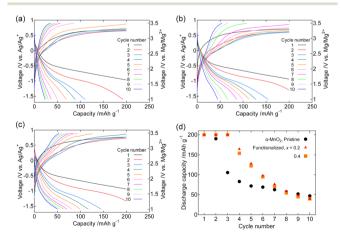


Fig. 5 Galvanostatic charge-discharge curves of dry composite cathodes of the pristine α -MnO₂ (a), and those functionalized with Fe_2O_3 at x = 0.2 (b) and 0.4 (c) in 0.3 mol dm⁻³ [Mg(G4)][TFSA]₂/[C₃mPyr][TFSA] at 100 °C. (d) Discharge capacity retention of samples shown in panels (a)-(c).

than that of the pristine sample (~ 25 mA h g⁻¹). Coulombic efficiency derived as the ratio of the discharge capacity to the charge capacity of the previous cycle, was also improved and exhibited ~0.6 at the 10th cycle for samples with x = 0.2 and 0.4.

Fig. 5 and S3† show the galvanostatic charge-discharge curves, discharge capacity retention, and Coulombic efficiency of the composite cathodes of pristine and Fe₂O₃functionalized α -MnO₂ (x = 0.2, and 0.4) in [Mg(G4)][TFSA]₂/ [C₃mPyr][TFSA] at 100 °C. Similar to structured MgMn₂O₄, Fe₂O₃ functionalization suppressed the increase in the charge potential with cycle number, and improved discharge capacity retention. The GITT profiles of the same composite cathodes are shown in Fig. 6. The amplitude of transient potential change was comparable in the 1st cycle, whereas smaller in the Fe₂O₃-functionalized sample in the 5th cycle. This observation indicates the reduction of overpotentials during charging and discharging by the functionalization. Fig. 7 and S4† show the galvanostatic charge-discharge curves, discharge capacity retention, and Coulombic efficiency of the composite cathodes of pristine and Fe₂O₃-functionalized α -MnO₂ in Mg[B(HFIP)₄]₂/G3 at 30 °C. In the pristine sample, the potential reached to the upper bound (3.5 V vs. Mg/Mg²⁺) during charging, indicating a large overpotential. In the Fe₂O₃-functionalized samples, the overpotential during charging was suppressed, and discharge capacity for up to the 5-6 cycles was significantly improved. The rapid fading of the discharge capacity thereafter in the Fe₂O₃-functionalized samples may be due to the reduced mobility of Mg²⁺ ions at 30 °C and the vulnerability of transition metal oxides to reduction in solvent (glyme)-rich electrolytes, both of which favour the destruction of α-MnO₂ crystallites rather than the reversible insertion and extraction of Mg²⁺ ions. Coulombic efficiency at the 10th cycle of the Fe₂O₃-functionalized samples with x = 0.4 was ~ 0.8 both in $[Mg(G4)][TFSA]_2/[C_3mPyr][TFSA]$ and $Mg[B(HFIP)_4]_2/G3$, and better than other two samples.

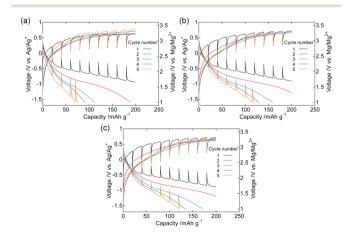


Fig. 6 GITT profiles of the pristine α -MnO₂ (a), and those functionalized with Fe_2O_3 at x = 0.2 (b) and 0.4 (c) in 0.3 mol dm⁻³ [Mg(G4)][TFSA]₂/[C₃mPyr][TFSA] at 100 °C.

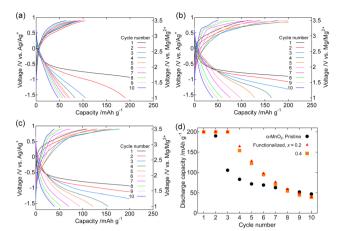


Fig. 7 Galvanostatic charge-discharge curves of dry composite cathodes of the pristine α -MnO₂ (a), and those functionalized with Fe_2O_3 at x = 0.2 (b) and 0.4 (c) in 0.3 mol dm⁻³ Mg[B(HFIP)₄]₂/G3 at 30 °C. (d) Discharge capacity retention of samples shown in panels (a)-(c).

Conclusions

An oxalate-assisted Fe₂O₃ functionalization technique was developed for the surface modification of nanosized transition metal oxides (MgMn₂O₄ and α-MnO₂) with large surface areas as cathode active materials for RMBs. Oxalate ions worked as efficient bridging agents between iron(III) ions and the surface of transition metal oxides. Fe₂O₃ functionalization suppressed side reactions including oxidative electrolyte decomposition, decreased overpotentials during charging and discharging, and improved discharge capacity retention. These observations demonstrate that the oxalate-assisted Fe₂O₃ functionalization is one of the powerful self-organizing surface functionalization techniques for the improvement of the electrochemical properties of cathode active materials for RMBs.

Data availability

The data of this study are available from the corresponding authors upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

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