Rechargeable magnesium-ion battery based on a TiSe$_2$-cathode with $d$-$p$ orbital hybridized electronic structure

Yunpeng Gu$^1$, Yukari Katsura$^2$, Takafumi Yoshino$^2$, Hidenori Takagi$^{1,3,4}$ & Kouji Taniguchi$^{5,6}$

Rechargeable ion-batteries, in which ions such as Li$^+$ carry charges between electrodes, have been contributing to the improvement of power-source performance in a wide variety of mobile electronic devices. Among them, Mg-ion batteries are recently attracting attention due to possible low cost and safety, which are realized by abundant natural resources and stability of Mg in the atmosphere. However, only a few materials have been known to work as rechargeable cathodes for Mg-ion batteries, owing to strong electrostatic interaction between Mg$^{2+}$ and the host lattice. Here we demonstrate rechargeable performance of Mg-ion batteries at ambient temperature by selecting TiSe$_2$ as a model cathode by focusing on electronic structure. Charge delocalization of electrons in a metal-ligand unit through $d$-$p$ orbital hybridization is suggested as a possible key factor to realize reversible intercalation of Mg$^{2+}$ into TiSe$_2$. The viewpoint from the electronic structure proposed in this study might pave a new way to design electrode materials for multivalent-ion batteries.

At present, rechargeable ion batteries have become one of the key systems for energy storage. In particular, lithium-ion batteries have been widely applied for most portable electronic devices as power sources. However, rechargeable batteries of further safety and low cost are recently being required for large scale applications, such as power sources of smart power grids and electric vehicles. Rechargeable magnesium-ion batteries are considered as one potential solution$^{1-5}$. Mg-metal, which might be used as a dendrite-free anode$^{6,7}$, is stable in ambient atmosphere and abundant in the earth crust compared with Li-metal. In addition, the specific volumetric capacity of Mg reaches 3833 mAh/cm$^3$, which is higher than that of Li (2046 mAh/cm$^3$), due to the bivalency of Mg-ion.

On the other hand, the development of Mg-ion batteries has been limited because of difficulties in selecting suitable cathode materials. The strong electrostatic interaction between bivalent Mg-ions and host lattices often causes slow solid state diffusion of Mg$^{2+}$ within the local crystal structure and prevents reversible insertion/extraction of Mg$^{2+}$. In the past years, various kinds of materials have been proposed as candidates for cathodes of Mg-ion batteries, such as Chevrel phases Mo$_6X_8$ ($X = S, Se$)$^{8,9}$, $\alpha$-Mo$_5Se_3^{10}$, TiS$_2$ nanotube$^{11}$, graphene-like MoS$_2^{12}$, WSe$_2$ nanowire$^{13}$, MgMnSiO$_4$ ($M = Fe, Mn, Co$)$^{14-20}$, MgFePO$_4$:F$^{21}$, FePO$_4$,$^{22}$ MnO$_2$,$^{23,24}$ V$_2$O$_3$ xerogel/thin film$^{25-27}$, organic compounds$^{28-30}$, Prussian Blue analogues$^{31,32}$. However, in most cases, Mg-ion batteries could only function by assistance of nanoscale morphology of electrode materials$^{33-35}$ or screening of the electrostatic interaction by solvent species such as...
aqueous ions\textsuperscript{31,32}. One exception is Chevrel phases, Mo$_6$X$_8$ (X = S, Se). Electrochemical performance of micro-crystal Mo$_6$X$_8$-electrodes is reported to be comparable with that of nano-crystal at ambient temperature\textsuperscript{9}. In the Chevrel phases, delocalized electronic orbitals of Mo$_6$X$_8$, in which two electrons are accommodated by cluster units of Mo$_6$ and the variation of formal charge on individual Mo atom are suppressed to only 1/3\textit{e}, are suggested to possibly moderate local structural deformation induced by the electrostatic interaction between Mg$^{2+}$ and the local lattice, in which extra electrons are accommodated\textsuperscript{4,33}. As a consequence of charge distribution over multiple atoms, the charge imbalance during the insertion of bivalent ions is supposed to be easily relieved in the Chevrel phase. In the recent studies, other compounds with clustered structures, such as o-Mo$_9$Se$_{11}$ and C$_6$0, are also reported to display reversible Mg-ion insertion reactions\textsuperscript{10,30}. Thus, systems with delocalized electrons over multiple atoms might be expected as good candidates for rechargeable cathodes of Mg-ion batteries.

In this study, we have focused on the possible effect of electronic delocalization in metal-ligand units through orbital hybridization in transition-metal chalcogenides. When the atomic orbitals hybridize strongly between transition-metals and chalcogens, the electronic wave function of transition-metal chalcogenide spreads on both constituent atoms\textsuperscript{34} (See Figure S1a); The charge density of the introduced electrons should distribute over metal-ligand units as schematically shown in Fig. 1a. On the other hand, in a system with weak orbital hybridization, the electrons should be accommodated only in the transition metal orbitals as displayed in Fig. 1b (See also Figure S1b). In this study, taking into account that the degree of orbital hybridization is enhanced in the case that the difference between energy levels of atomic orbitals...
orbitals is small\textsuperscript{35}, layered TiSe\textsubscript{2} (Fig. 1c) has been selected as a candidate material based on the energy diagram for atomic d-orbitals of transition metals and p-orbitals of chalcogens (Fig. 1d)\textsuperscript{36}. As displayed in Fig. 1d, since the energy levels of valence atomic orbitals in TiSe\textsubscript{2}, 3d-orbital of Ti and 4p-orbital of Se, are close to each other, the hybridized electronic structure is expected around Fermi energy. In fact, the conducting properties, which could be ascribed to the delocalized electronic structure by orbital hybridization, have been confirmed in TiSe\textsubscript{2} around room temperature\textsuperscript{37}. Here, we report the rechargeable performance of TiSe\textsubscript{2} with d-p orbital hybridization as a cathode of Mg-ion batteries.

**Results**

The powder X-ray diffraction (XRD) result indicates that TiSe\textsubscript{2} used in this study takes a 1T-type structure (\(a = 3.5393(2)\) Å and \(c = 6.0103(3)\) Å) with a space group \(P\overline{3}-m1\) (Fig. 1e) as reported in the previous study\textsuperscript{38}. As shown in Fig. 1c, the 1T-type structure consists of successive Se-Ti-Se sandwich slabs, which are separated by the van der Waals gap. The sample was also checked by Scanning Electronic Microscopy (SEM), showing that TiSe\textsubscript{2} has a particle size around 10\(\mu\)m (Fig. 1e).

The orbital hybridization in TiSe\textsubscript{2} was confirmed by first principle calculation. The energy band dispersion and density of states (DOS) are shown in Fig. 2a, b, respectively. The calculated electronic band structure indicates that 1T-TiSe\textsubscript{2} is a semimetal with a small overlap between the valence band maximum at the Brillouin zone (BZ) center, \(\Gamma\)-point, and the conduction band minimum at the BZ boundary, \(L\)-point (Fig. 2a). This electronic band structure is consistent with the previous studies by \textit{Ab initio} band structure calculation\textsuperscript{39} and angle resolved photoemission measurements\textsuperscript{40,41}. The orbital components of valence and conduction bands indicate the existence of d-p orbital hybridization. As shown in the partial DOS of TiSe\textsubscript{2} (see Fig. 2b), the electronic states around the Fermi level \((E_F)\) consist of both 3d-orbital of Ti and 4p-orbital of Se. In this situation, since the wave function spreads over both Ti- and Se-atoms as shown in Fig. 1a, introduced electrons in the discharge process of TiSe\textsubscript{2} are expected to be accommodated into a “cluster-like” electronic state.

The Mg-ion battery with a TiSe\textsubscript{2} cathode shows reversible electrochemical charge/discharge performance at ambient temperature. Figure 3a displays a typical voltage profile of the coin cell at 25°C, which consists of a TiSe\textsubscript{2}-based cathode, a magnesium metal anode and 0.25M Mg(AlCl\textsubscript{3}-EtBu\textsubscript{2})/THF electrolyte solution, in which formation of the passivation film is suppressed thus Mg is reversibly deposited/dissolved on the anode\textsuperscript{42}. The derivative of the capacity (Q) with respect to the voltage (V), \(dQ/dV\), is plotted in Fig. 3b. The redox peaks are observed in the cathodic process (at 0.9 V vs. Mg/Mg\textsuperscript{2+}) and anodic process (at 1.2 V vs. Mg/Mg\textsuperscript{2+}), respectively. This result indicates that TiSe\textsubscript{2} can work as a rechargeable electrode material in the Mg-ion battery system.

Figure 3c shows cyclability of the TiSe\textsubscript{2} cathode in the first 50 cycles at 25°C. The specific capacity is kept at the values around 110 mAh/g during the whole cycling. Assuming that the redox species is Ti\textsuperscript{4+}/Ti\textsuperscript{3+}, the average specific capacity observed, 108 mAh/g, is about 83% of the theoretical capacity.

![Figure 2. (a) Dispersion of the energy band of 1T-TiSe\textsubscript{2}. (b) Total and partial density of states (DOS) for 1T-TiSe\textsubscript{2}.](image-url)
The specific capacity of TiSe$_2$ is comparable with that of Mo$_6$S$_8$ (~100 mAh/g), which is the prototype cathode material of Mg-ion batteries.

Structural variation after the discharge process was traced by measuring *ex situ* XRD patterns of Mg$_x$TiSe$_2$. XRD measurements were performed for selected compositions ($x = 0, 0.2, 0.4, 0.5$). As shown

![Graph showing charge/discharge curves](image)

Figure 3. (a) The charge/discharge curve (on the second cycle) of the Mg-ion battery cell with TiSe$_2$ measured at 25°C. (b) The derivative of the capacity ($Q$) in Fig. 3(a) with respect to the voltage ($V$), $dQ/dV$. (c) Cycle performance of the Mg-ion battery cell with TiSe$_2$ for capacity. The inset shows the charge/discharge curves on each cycle.
in Fig. 4a, the XRD pattern of TiSe$_2$ is maintained after the discharge process. In the XRD pattern of discharged cathode, a small broad peak appears around 47°, which are likely to be assigned to those of Se. Selenium phase seems to be produced by some decomposition of partial cathode, which might be the reason of capacity fading during the cycling. Focusing on the 00$l$ peaks ($l = 1, 2, 3$ and 4), all of them gradually shift to the lower angles. This result indicates that Mg$_2^+$ is intercalated into the van der Waals gap between TiSe$_2$ layers in the crystal lattice. During the Mg-ion intercalation, the lattice constant of c-axis, which was estimated from the XRD patterns in Fig. 4a, increases from 6.0103(3) Å ($x = 0$) to 6.072(3) Å ($x = 0.5$) (Fig. 4b). The structural reversibility was also confirmed by the ex situ XRD measurements. Figure 4c displays the XRD patterns after discharge/charge process. The 004 peak shifts to

Figure 4. (a) Powder XRD of Mg$_x$TiSe$_2$ ($x = 0$, 0.2, 0.4 and 0.5). The ex situ XRD measurements were carried out after the discharging processes. (b) Mg-content ($x$) dependence of unit cell parameter of c-axis. (c) 004 peak shift after discharging ($x = 0.48$; orange line) and charging processes ($x = 0.05$; red line). The peak in the initial state ($x = 0$) is represented by the blue line.
Discussion

So far, materials research on cathodes of Mg-ion batteries has been independently performed with emphasis on the crystal structure of each target material in most cases. On the other hand, by introducing the viewpoint based on electronic structures as in this study on TiSe\(_2\), a common feature seems to appear in the previously reported cathode materials of Mg-ion batteries, such as Mo\(_6\)X\(_6\) (X = S, Se)\(^{18}\), TiS\(_2\),\(^{13}\), MoS\(_2\),\(^{12}\) and WSe\(_2\)\(^{19}\). The common feature is that orbital hybridization is expected between the valence atomic orbitals of constituent atoms in these materials, which are d-orbital of transition metal atoms and p-orbital of chalcogen atoms, due to the close energy levels (Fig. 1d). Intriguingly, such small energy difference between transition metals’ d-orbital and ligands’ p-orbital can be found not only in the cathodes of Mg-ion batteries but also in the representative ones of Li-ion batteries; Li energy range. In addition, since selenium is a rare element that may be toxic, developing cathodes of Mg-ion batteries with transition metal oxides, such as Li\(_2\)MnO\(_4\) by X-ray spectroscopy\(^{49,50}\).

In the case of selenides, d-p orbital hybridization should be enhanced by high orbitals overlap due to the large 4p-orbital size of Se, compared with oxides or sulfides. Additionally, in TiSe\(_2\), the two-dimensionality of ion conducting channels in the van der Waals gap might contribute to suppress the effect of Coulomb repulsion between Mg-ions\(^{19}\). These factors in the TiSe\(_2\) cathode might contribute to its reversibility on electrochemical cycling and high specific capacity close to the theoretical value in the Mg-ion battery even with micro-sized electrode. In fact, the Mg-ion battery with another cathode material, 1T-VSe\(_2\), where the electronic and crystal structures are similar with those of 1T-TiSe\(_2\), also shows rechargeable cathode performance over 60 cycles with relatively high capacity of ~110 mAh/g at ambient temperature (Figure S2a, b). In these materials (TiSe\(_2\) and VSe\(_2\)) that can work as reversible Mg-ion battery cathodes in the form of micro-sized particles, further improvement in cyclability and rate capability might be expected by downsizing cathode materials to the nanometric scale as done in graphene-like MoS\(_2\)\(^{12}\) and WSe\(_2\) nanowire\(^{43}\).

The viewpoint from electronic structure of cathode materials could draw a guideline for developing high voltage Mg-ion batteries. In general, open circuit voltage of an ion-battery (\(V_{oc}\)) is given by difference in electrochemical potentials of electrons between anode (\(\mu_e^a\)) and cathode (\(\mu_e^c\)); \(V_{oc} = \mu_e^c - \mu_e^a\), where \(e\) is magnitude of electron charge\(^{51,52}\). Taking into account that the electrochemical potential of electrons is defined by Fermi level in vacuum and inner-potential, trend of ion-battery performance could be assessed by difference in Fermi levels between two electrodes in first approximation. Since the Fermi level of cathode is determined by the electronic structure of valence/conduction band, the voltage of battery using same anodes should reflect the energy levels of valence atomic orbitals, which constitute a valence/conduction band. In the case of TiSe\(_2\), because the energy levels of Ti 3d-orbital and Se 4p-orbital are located in the shallow energy range compared with late transition metals and oxygen as shown in Fig. 1d, the observed voltage, ~1 V vs. Mg/Mg\(^{2+}\), seems to be a lower value than those of Li-ion batteries with transition metal oxides, such as LiCoO\(_2\) and Li\(_2\)MnO\(_4\). In further material research for Mg-ion battery cathodes with improved performance, late transition metal oxides might be good candidates for reaching high voltages cells, because their valence bands consist of atomic orbitals located in the deep energy range. In addition, since selenium is a rare element that may be toxic, developing cathodes of oxides could be desirable from the standpoint of practical use and environmental concerns.

In summary, this study has demonstrated that the Mg-ion battery with a micro-sized TiSe\(_2\) cathode shows rechargeable performance at ambient temperature. In TiSe\(_2\), reflecting the close energy levels of 3d-orbital of Ti and 4p-orbital of Se, d-p orbital hybridization around the Fermi level was confirmed by the viewpoint based on electronic structures. The XRD measurements indicate that Mg-ions are reversibly intercalated into and deintercalated from the van der Waals gap between the TiSe\(_2\) layers. We have proposed one possible scenario that the charge delocalization in metal-ligand units by strong d-p orbital hybridization might be one of the key factors, which improve reversible performance of Mg\(^{2+}\)-intercalation/deintercalation. Further study based on the electronic structure could open a new way to design cathode materials for Mg-ion and other multivalent ion batteries.

Methods

Sample preparation and electrochemical measurements. The purchased TiSe\(_2\) powder (>99%, High Purity Chemicals) was used as a cathode material. Cathodes were fabricated using TiSe\(_2\), acetylene black (AB) and polytetrafluoroethylene (PTFE) in a mass ratio of 81:9:10. The mixed active material was
pressed on a current collector of copper net and dried at 60 °C in vacuum. The amount of active material was approximately 3 ~ 4 mg per cathode. Tests of Mg-ion batteries were performed by 2032-type coin cells, which were assembled in an Ar-filled glove box. These cells consisted of TiSe₂ cathodes, magnesium metal anodes and the Mg(AlCl₃EtBu)₂ electrolyte dissolved at 0.25 mol/dm³ in tetrahydrofuran (THF). The galvanostatic charge/discharge tests were conducted using a potentiogalvanostat (Solartron, 1470E) at 25 °C. Current density dependence of capacity was tested for 5 mA/g, 10 mA/g, 20 mA/g and 50 mA/g, respectively (Fig. S3(a)). Rate capability, which is defined as the capacity ratio to the discharge capacity at 5 mA/g, decreases to ca. 50% at high current density of 50 mA/g (see Fig. S3(b)). The cell voltage in the galvanostatic charge/discharge measurements was cycled between 1.8 V and 0.2 V (vs. Mg/Mg²⁺) for 50 times by applying a constant current of 5 mA/g. Electrochemical characteristic of TiSe₂ in a electrolyte solution other than Mg(AlCl₃EtBu)₂/THF was examined through Cyclic voltammetry (CV) measurements with a three-electrode cell. As an electrolyte for CV measurements, Mg(ClO₄)₂ was dissolved in acetonitrile (AN) at 1 mol/dm³. A magnesium metal and a silver wire, which was put into a solution of 0.01 mol/dm³ AgNO₃ and 0.1 mol/dm³ tetrabutylammonium perchlorate in acetonitrile, were employed as a counter electrode and a reference electrode, respectively. In the CV measurement, the potential was scanned between 0 V and −1.5 V (vs. Ag/Ag⁺) at 0.5 mV/s. In the voltammogram, one peak was clearly observed for both cathodic and anodic processes, respectively (Fig. S4).

Material characterization. Structural characterization of samples was performed to analyze the change of TiSe₂ upon cycling by ex situ powder X-ray diffraction (XRD) using a graphite monochromatized Cu Kα radiation source. The particle size of sample was measured by scanning electronic microscopy (SEM). The variation of the Mg concentration in the TiSe₂ cathodes after discharge/charge process was traced by quantitative chemical analysis. The atomic ratio of Mg to Ti was determined quantitatively by inductively coupled plasma atomic emission spectrometry (ICP-AES). The electrodes were removed from the cells after the charge/discharge cycling process and rinsed with THF to dissolve out the electrolyte salt in an Ar-filled glove box before the measurements. In the ex situ structural analysis of electrodes, electrochemically prepared electrode samples were measured immediately after being transferred out of the glove box, or sealed with polyethylene-nylon films in an Ar-filled glove box before XRD measurements.

First-principle calculation. First-principle electronic structure calculations were performed by Full-potential Linearized Augmented Plane Wave (FLAPW) method implemented in WIEN2k code, under Tran-Blaha modified Becke-Johnson (TB-mBJ) exchange-correlation potential. Muffin-tin radii for Ti and Se were set to 2.50 and 2.25 a.u., respectively. The first Brillouin zone was divided into 21 × 21 × 10 k-point mesh with 436 irreducible points. No spin polarization was considered. Obtained density of states (DOS) was rescaled to the primitive unit cell.

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Y. G. performed the sample preparation, characterization and electrochemical measurements. Y. K. contributed to the first principle calculation. T. Y. assisted in the experiments. H. T. co-supervised the project. K. T. conceived the experiments, carried out the data analysis and supervised the project. All authors discussed the results and commented on the manuscript.

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Author Contributions

Y.G. performed the sample preparation, characterization and electrochemical measurements. Y.K. contributed to the first principle calculation. T.Y. assisted in the experiments. H.T. co-supervised the project. K.T. conceived the experiments, carried out the data analysis and supervised the project. All authors discussed the results and commented on the manuscript.

Additional Information

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