Coordination Polymers for High-Capacity Li-Ion Batteries: Metal-Dependent Solid-State Reversibility

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ABSTRACT: Electrode materials exploiting multielectron-transfer processes are essential components for large-scale energy storage systems. Organic-based electrode materials undergoing distinct molecular redox transformations can intrinsically circumvent the structural instability issue of conventional inorganic-based host materials associated with lattice volume expansion and pulverization. Yet, the fundamental mechanistic understanding of metal–organic coordination polymers toward the reversible electrochemical processes is still lacking. Herein, we demonstrate that metal-dependent spatial proximity and binding affinity play a critical role in the reversible redox processes, as verified by combined 13C solid-state NMR, X-ray absorption spectroscopy, and transmission electron microscopy. During the electrochemical lithiation, in situ generated metallic nanoparticles dispersed in the organic matrix generate electrically conductive paths, synergistically aiding subsequent multielectron transfer to π-conjugated ligands. Comprehensive screening on 3d-metal–organic coordination polymers leads to a high-capacity electrode material, cobalt-2,5-thiophenedicarboxylate, which delivers a stable specific capacity of ~1100 mA h g−1 after 100 cycles.

KEYWORDS: conversion, high capacity, lithium-ion batteries, organic ligand, reaction mechanism

1. INTRODUCTION

The reversible topotactic transition of two-dimensional inorganic compounds through the intercalation chemistry is the most extensively investigated redox process. However, the limited storage capacity cannot meet the emerging demands of the future society. Alloying process that enlists groups 14 or 15 elements (e.g., Si, Ge, Sn, and P) provides greater capacities by accommodating multiple Li2,3 However, it often accompanies the fundamental mechanistic understanding of metal–organic coordination polymers toward the reversible electrochemical processes is still lacking. Herein, we demonstrate that metal-dependent spatial proximity and binding affinity play a critical role in the reversible redox processes, as verified by combined 13C solid-state NMR, X-ray absorption spectroscopy, and transmission electron microscopy. During the electrochemical lithiation, in situ generated metallic nanoparticles dispersed in the organic matrix generate electrically conductive paths, synergistically aiding subsequent multielectron transfer to π-conjugated ligands. Comprehensive screening on 3d-metal–organic coordination polymers leads to a high-capacity electrode material, cobalt-2,5-thiophenedicarboxylate, which delivers a stable specific capacity of ~1100 mA h g−1 after 100 cycles.

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polymer frameworks through the comprehensive screening of the first-row transition-metal (3d-metal) cations and inverse-Wüster-type molecules. This approach leads to a tailored coordination polymer, cobalt-2,5-thiophenedicarboxylate, which maintains an enhanced reversible capacity of \( \approx 1100 \text{ mA h g}^{-1} \) at a high current density of 500 mA g\(^{-1}\).

2. EXPERIMENTAL SECTION

2.1. Sample Preparation and Characterization. Mn, Fe, Co, Ni, Cu, or Zn cation was coordinated with a terephthalate (TP) ligand in an aqueous solution via salt metathesis reaction (see the Supporting Information, section I). The crystal structure, number of hydrates, and the thermal stability were estimated by powder X-ray diffraction (XRD) and thermogravimetric/differential thermal analyses (TGA/DSC) (Figures S1 and S2, Supporting Information). Fourier transform infrared (FT-IR) spectra (Figure S3) revealed characteristic asymmetric carboxylate vibration peaks, aromatic and symmetric carboxylate vibration peaks, and thermogravimetric/differential thermal analyses (TGA/DSC) (Figures S1 and S2, Supporting Information). The crystal structure, number of hydrates, and the thermal stability were estimated by powder X-ray diffraction (XRD) and thermogravimetric/differential thermal analyses (TGA/DSC) (Figures S1 and S2, Supporting Information).

2.2. Electrochemical Tests. The activated material was mixed with carbon black (Super P) and binder in a 4:3:1 weight ratio. The electrochemical performance was evaluated using 2032 coin cells with a lithium metal anode and 0.8 M LiPF\(_6\) in ethylene carbonate/diethyl carbonate (1:1, v/v) electrolyte solution. Galvanostatic experiments were performed at 25 °C, and specific current densities of 100 or 500 mA g\(^{-1}\) were applied to electrode materials.

3. RESULTS AND DISCUSSION

3.1. Electrochemical Properties of Conversion Materials. Coordination polymer architectures comprising divalent 3d-metal cations and TP ligand have been chosen because of the well-established synthetic protocols, cost effectiveness, and low environmental impact (see the Supporting Information, sections I & II). Figure 2 shows the potential profiles of anhydrous 3d-metal TP samples at a current density of 100 mA g\(^{-1}\). It is noteworthy that redox couples are located not only on the transition metal but also on the inverse-Wüster-type TP ligand, as depicted in Figure 1c. The potential profiles exhibited a multistage electrochemical lithiation behavior: (i) an abrupt potential drop at 1.0–1.4 V linked to the metal redox couple,\(^{12,23,24,33−35}\) (ii) a voltage plateau near 0.6–0.8 V attributed to the carboxylate-to-enolate conversion,\(^{12,22,36−38}\) and (iii) a subsequent sloping profile associated with the progressive vinyl-to-saturated bond conversion\(^{39−41} \) of the organic scaffold (see Figure 1b). The formation of SEI films or electrically isolated islands brings irreversible capacity loss,\(^{42,43}\) resulting in the low initial Coulombic efficiencies (CEs) (78.8% for CoTP, 77.0% for NiTP, 62.5% for FeTP, 50.4% for CuTP, 46.9% for MnTP, and 32.5% for ZnTP). However, the CE values reach close to 100% within a few cycles. Further cycling displays different profiles as compared to the first cycle, typically observed features for the conversion processes undergoing large structural rearrangement.\(^{1,44−46}\)

CoTP, FeTP, and NiTP delivered high reversible charge capacities of 1155, 1143, and 1092 mA h g\(^{-1}\) after 10 cycles, respectively, whereas MnTP, CuTP, and ZnTP carried relatively low specific capacities of 794, 604, and 511 mA h g\(^{-1}\), respectively. Most samples showed stable cycle performance (Figure 3a,b). Noticeably, lithiation-induced activation behavior occurred for MnTP with gradual capacity drop. The rate performance (Figure 3c) was further investigated, and the discharge capacities of the active materials were recorded by subtracting the Super P contribution (Figure S5). MnTP also displayed the poorest rate capability, with a 26% retained capacity with increasing current density from 100 to 2000 mA g\(^{-1}\). FeTP, CoTP, and NiTP exhibited good capacity retention ratios above 63%. Despite diminished capacities, CuTP and

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Figure 1. (a) Conventional conversion reaction of inorganic host materials. (b) Redox chemistry of inverse-Wüster-type molecules. (c) This study: metal-dependent redox reactions of metal−organic complexes.

Figure 2. Voltage profiles of 3d-metal TPs at a current rate of 100 mA g\(^{-1}\).
ZnTP showed good rate performance in the range of 1000–2000 mA g\(^{-1}\). To clarify the electrochemical behaviors, differential capacity (dQ/dV) measurements were also carried out. During the first lithiation step (Figures 3d and S6), FeTP, CoTP, NiTP, and CuTP were studied.
CuTP revealed characteristic M(II) reduction peaks at 1.02, 1.19, 1.26, and 1.30 V, respectively. Within the measured voltage range, reductive peaks for MnTP and ZnTP were not observed, presumably because of their shift to the lower potential region. Subsequent reductive peaks (0.8–0.9 V) are originated from the carboxylate-to-enolate transformation. During the delithiation process (Figure 3e), enolate-to-carboxylate reconversion was observed in a range of 0.74–1.04 V. The regeneration of M(II) cations for FeTP, CoTP, and NiTP was found in the range of 1.55–1.84 V. On the basis of the dQ/dV analysis, the polarization gap (40–260 mV) was much smaller than that of the M(II)/M(0) redox couple (540–580 mV), indicating favorable redox kinetics of the carboxylate-enolate conversion. It is noteworthy that additional capacity is observed in the range of ~1.5–3.0 V beyond the theoretical capacity estimated from the M(II)/M(0) redox couple, implying the concurrent involvement of the aromaticity recovery process (see also Figure 2). This asymmetrical redox route upon discharge/charge cycling is linked with voltage hysteresis, the characteristic feature for the conversion-type processes. However, metal oxidative peaks for MnTP, CuTP, and ZnTP could not be detected. To elucidate the origin of the metal-dependent reversible conversion process, high-capacity/reversible CoTP and low-capacity/irreversible CuTP were selected as representative models for comparison (Table S1, Supporting Information).

### 3.2. CoTP Versus CuTP: Spatial Proximity

Synchrotron-sourced ex situ X-ray absorption near-edge structure (XANES) analysis was applied to trace the valence state changes of CoTP and CuTP on the basis of the redox peaks from the dQ/dV plots (Figure 4a–d). The initial divalent state of CoTP was determined from a weak pre-edge peak at 7710 eV, assigned to the 1s → 3d transition. This peak arises from either noncentrosymmetric metal site or 4p–3d orbital mixing. In the reconversion process approaching 3.0 V, regenerated cobalt(II) oxidation states was observed with slight changes in absorption-edge and pre-edge peaks, implying disordered local environment. In sharp contrast, CuTP displayed irreversible behavior under the same electrochemical conditions.

Next, ex situ EXAFS study was conducted to unravel their local coordination environments (Figure 4e). The radial distance is given by radial distribution function, which is uncorrected for phase shifts. CoTP initially displayed a strong Co–O bond. When the redox potential reached 1.1 V, a strong intermetallic Co–Co bond appeared together with the Co–O bond. This indicates that metallic Co is still in close proximity to the oxygen atom from carboxylates (i.e., homogeneously embedded Co particles in organic ligand domains). No pronounced variations were observed until the potential reached 0.0 V. Upon full delithiation up to 3.0 V (reconversion step), the Co–Co metallic bond completely disappeared and a similar spectrum resembling the pristine stage was recovered. In contrast, the EXAFS pattern of CuTP upon cycling was distinctly different, as shown in Figure 4f. The intensity of the Cu–O bond became weaker, whereas a strong Cu–Cu correlation appeared after passing through the first reductive peak. At 0.0 V, the intensity of the Cu–O correlation weakened, suggesting that metallic copper clusters were segregated from the organic ligand matrix. During delithiation up to 3.0 V, the spectrum was found to be largely irreversible, where the Cu–Cu bond was mostly intact together with partial recovery of the Cu–O peak. These EXAFS data unequivocally indicate that the spatial arrangement of metallic clusters and organic ligand plays an essential role in determining the molecular structural reorganization and recovery.

To provide information on the spatial distribution of metallic clusters, high-resolution transmission electron microscopy (HRTEM) analyses (Figures 5 and S8) were conducted for both CoTP and CuTP. After full lithiation to 0.0 V, the average size of the metallic cobalt clusters (yellow circles) is about 2 nm. These nanoparticles are well dispersed in the organic matrix. In contrast, fully reduced metallic copper clusters show segregated structures (yellow circles, ~20 nm in size). This compositional inhomogeneity can disconnect segregated particles from the organic ligand matrix. To evaluate the role of metal redox process decoupled from organic redox chemistry, a parallel comparison between CoTP and CuTP was conducted within the potential range of 3.0–1.0 V (Figure S7). Although overall capacity dramatically diminished, the cobalt complex showed better reversibility than the copper complex. Additionally, the metal-dependent binding energies were evaluated by applying
the density functional theory (DFT) model system on the basis of metal cluster(s) and single TP ligand (for the details, see the Supporting Information, section I-3). It is noteworthy that cobalt clusters strongly coordinate with TP ligand, while stabilizing the overall surface energy. In contrast, copper clusters show less effective binding with the ligand, which can result in their segregation of the clusters from the organic matrix and solid-state irreversibility.

3.3. Classification of 3d-Metal TP Complex Systems.

After the initial observations on CoTP and CuTP, the remaining metal–organic complexes (MnTP, FeTP, NiTP, and ZnTP) were further explored. HRTEM images (Figures 5a and S10 in the Supporting Information) of CoTP, FeTP, and NiTP at 0.0 V typically show fine particles mostly below ca. 3 nm. On the other hand, CuTP, MnTP, and ZnTP samples at 0.0 V generally exhibit bigger particle sizes and larger size distributions (Figures 5b and S11). To unravel the valence state changes of 3d-metals, XANES analyses were then performed on a series of 3d-metal TP complexes. During the first discharge cycle (Figure S12), FeTP and NiTP were fully reduced to metallic iron and nickel. In the reconversion process approaching to 3.0 V, they were reconverted into divalent phase with slight variation, sharing the solid-state reversibility features with CoTP.

Next, the valence states of MnTP were evaluated upon cycling. As shown in Figure S12b, Mn(II)/Mn(0) redox chemistry by the XANES analysis revealed mixed valence state and borderline features. Remarkably, ZnTP exhibited unique redox mechanism during electrochemical reaction; when fully...
lithiated to 0.0 V, reduction of divalent zinc cation did not stop at the metallic phase but progressively continued to form lithium−zinc (LiZn) alloy (Figure S12b). The LiZn phase was further confirmed by HRTEM and the corresponding fast Fourier transform pattern analysis (Figure S11). During the delithiation step approaching to 3.0 V, the white line was not shifted to the pristine stage exhibiting the solid-state irreversibility.

Overall, the 3d-metal TP complex systems are categorized into three groups according to the metal-specific electrochemical properties (Figure 6a): type I (CoTP, FeTP, and NiTP)—reversible/high performance group; type II (MnTP)—borderline group; and type III (CuTP and ZnTP)—irreversible/poor performance group. During the lithiation, type I compounds in situ generate metallic cobalt, iron, and nickel clusters in close proximity to the oxygen atom of the carboxylate group. The nanoparticles synergistically can assist successive lithiation to \( \pi \)-conjugated ligands. In contrast, type III compounds display the compositional inhomogeneity (i.e., disconnected metallic or alloy clusters from the organic ligand matrix). It results in an interfacial penalty hampering the multistage molecular rearrangement during the conversion/ reinversion reaction (see the schematic illustration in Figure 6b).

3.4. Cobalt\(^{1}\)-2,5-thiophenedicarboxylate. Having identified the 3d-metal-dependent solid-state reversibility of type I compounds, our attention was shifted to expand the repertoire of metal−organic coordination polymers. After the comprehensive screening of 3d-metal cations (Supporting Information, section I) and \( \pi \)-conjugated heterocyclic ligands (Supporting Information, section II), cobalt(II) and 2,5-thiophenedicarboxylate (TDC) moieties were assembled to yield CoTDC \( \cdot 1.5 \) H\(_2\)O via a facile hydrothermal reaction \(^{34,55}\) (for the details, see the Supporting Information, section III). Powder XRD patterns confirmed the formation of the desired product (Figure S18c). TGA/DSC data revealed the dehydration at \( \sim \)240 °C with \( \sim \)10 wt % loss, in good agreement with the corresponding number of hydrates (1.5 H\(_2\)O) (Figure S18d). The as-prepared sample was further investigated by FT-IR spectroscopy, and the characteristic dicarboxylate peaks were observed at 1515 and 1326 cm\(^{-1}\) assigned to \( \nu \)\(_{\mathrm{as}}\)(COOCo) and \( \nu \)\(_{\mathrm{s}}\)(COOCo), respectively (Figure S18e). The sample was annealed at 240 °C in vacuo, and the complete removal of the hydrates was confirmed by TGA/DSC and FT-IR spectrum (see Figure S19). Galvanostatic cycling tests were carried out using the annealed and amorphized CoTDC (Figures 7a and S20), displaying substantially improved electrochemical performances than aforementioned 3d-metal TPs. The reversible specific capacity of CoTDC maintained at 1420 mA h g\(^{-1}\) after 100 cycles under the current density of 500 mA g\(^{-1}\). After the removal of Super P contribution, the reversible capacity of anhydrous CoTDC was estimated to be 1192 mA h g\(^{-1}\) corresponding to \( \sim \)10 Li per cobalt metal (Figures S21 and S22).

Fully reduced state of CoTDC was then examined via ex situ XAS. The valence change was traced with normalized Co K-edge XANES spectra at various points (Figure 7b,c). After passing the first reductive peak at 1.25 V, the pre-edge peak gradually evolved and the white-line intensity concurrently decreased. The Co(0)/Li\(_{4+x}\)TDC phase (see also eqs 1–2a) during the charge step subsequently recovered divalent features of CoTDC (see also EXAFS patterns in Figure S23). HRTEM presents fully reduced nanoparticles in the size range of \( \sim 2\)–3 nm that are well dispersed in the organic matrix (Figures 7d and S24). Next, \(^{13}\)C-isotope-labeled CoTDC (CoTDC-\(^{13}\)C\(_1\)) was prepared to enhance the signal sensitivity of \(^{13}\)C NMR spectra (Scheme S2 and Figure S25, Supporting Information). The carboxylate \(^{13}\)C peak was observed in the chemical shift of 170–180 ppm (Figure S26). A broad peak \( \sim 100 \) ppm region is attributed to Super P. After lithiation and subsequent delithiation, the solid-state \(^{13}\)C NMR spectrum of CoTDC-\(^{13}\)C\(_1\) at 3.0 V provided a broad band centered at \( \sim 180 \) ppm, ascribed to the regenerated carboxylate \(^{13}\)C (Figure S26).

Stage 1: M(II)/M(0) redox couple.
**3.5. Role of Nanoparticles.** Pristine Li$_2$TDC in the absence of cobalt nanoparticles exhibited larger polarization, poor cyclability, and rate capability upon cycling compared with CoTDC (see Figure S27). To understand the enhanced redox kinetics of CoTDC, a galvanostatic intermittent titration technique (GITT) was employed to reveal the overpotential ($\eta$) differences (Figure 8a). The initial $\eta$ value of Li$_2$TDC was approximately 95 mV owing to the kinetically favorable carboxylate-to-enoate reduction. In contrast, the CoTDC electrode at the early lithiation (~15% SOC) exhibited a larger $\eta$ value linked to the nucleation of metallic clusters and the corresponding rearrangement of local coordination structures. After the formation of Co(0) phase, however, the $\eta$ value of CoTDC abruptly diminished to 94 mV, attributed to the corresponding carboxylate-to-enoate conversion (i.e., Co(0) + Li$_2$TDC + 2Li $\rightarrow$ Co(0) + Li$_2$TDC). Then, the $\eta$ values of both materials increased, where vinyl group reduction started. Notably, CoTDC exhibited a smaller $\eta$ value than Li$_2$TDC; the measured values of CoTDC were 162 and 159 mV at 50% SOC, whereas those of Li$_2$TDC were 235 and 211 mV (see the Figure 8a inset). To determine the dominant parameter affecting the reduced $\eta$ values, charge-transfer resistance was evaluated from the semicircle of electrochemical impedance spectroscopy (see also Figures S28 and S29 in the Supporting Information). The Arrhenius plots for charge transfer were obtained by preparing symmetric cells at 50% SOC, while changing the temperature from 253 to 283 K (see Figure 8b). Yet, slight differences in the activation energies were found for CoTDC (65.0 kJ mol$^{-1}$) and Li$_2$TDC (68.2 kJ mol$^{-1}$).

We speculated that the enhanced electrochemical performance of CoTDC might be attributed to the enhanced electrical conduction path from in situ generated metallic Co(0) nanoparticles (eqs 1-2a). To verify our hypothesis, pristine CoTDC was treated with reactive lithium metal assisted by a vortex mixer (see the Supporting Information, section I-1). This chemically lithiated sample showed XANES patterns similar to those of the electrochemically lithiated sample (Figure 8c). No lithium metal peak was detected in the ex situ XRD pattern afterward (Figure S30), indicating its reaction with CoTDC. The electrical conductivity of the chemically lithiated sample, Co(0)/Li$_2$TDC, was 2.48 $\times$ 10$^{-5}$ S cm$^{-1}$ comparable to the physical mixture of Li$_2$TDC and cobalt nanoparticles (Figure 8d), which suggests an electrical conduction network from the reduced metal domains. Improved electrical conductivity in the presence of interconnected metallic cobalt nanoparticles may underlie the improved electrochemical performance (see also Figure 6b).

**4. CONCLUSIONS**

In summary, we have successfully demonstrated that the metal-dependent solid-state redox reversibility of complex systems is governed by spatial proximity and binding affinity between 3d-metal and organic ligand matrix. Metallic cobalt nanoparticles in the size range of ~2–3 nm, well dispersed in the matrix, exhibit facile coordination with organic ligands. They can also offer electrical conduction paths that allow successive multielectron transfer to $\pi$-conjugated redox molecules. In stark contrast, the compositionally inhomogeneous copper clusters induce limited reconversion owing to an interfacial penalty. Comprehensive screening of coordination complexes aided by synthetic chemistry and electrochemical investigations leads to the high-capacity molecular complex, cobalt-2,5-thiophenedicarboxylate. We anticipate that this study has implications in the field of organic-based rechargeable batteries by elucidating the metal-dependent solid-state multistage redox behaviors.

**ASSOCIATED CONTENT**

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsami.8b04678.

Complete experimental procedures, reaction schemes, electrochemical performance, product characterizations, and DFT calculations (PDF)

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**Notes**

The authors declare no competing financial interest.

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