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Co₃O₄ Nanoparticles Embedded Carbonaceous Fibre: Nanoconfinement Effect on Enhanced Lithium-Ion Storage

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Co₃O₄ nanoparticles embedded carbonaceous fibres were prepared from Co²⁺ coordinated regenerated cellulose fibres, which showed a high reversible capacity and excellent cycling stability as anode materials for Li-ion batteries.

Rechargeable lithium-ion batteries (LIBs) have long been considered as most promising reversible energy storage for a wide variety of application.¹ Graphite has been extensively applied as anode material for commercial LIBs because of their low cost and good stability. However, a theoretical capacity of only 372 mAh g⁻¹ significantly limits its LIBs performance.² Although transition metal oxides (TMOs) are promising alternatives for high-performance LIBs anode materials because of their high theoretical capacities, they suffer from the problems of poor electronic conductivity and a large volume change during charge/discharge process, resulting in poor cycling performance.³ To address the drawbacks of the TMOs anodes, one approach is to develop nanostructure TMOs/carbon composites,⁴ where the nanoconfinement effect of carbon matrix can efficiently alleviate the aggregation and mechanical strain generated by repeated volume expansion/contraction of TMOs nanoparticles (NPs) and enhance the electrochemical conductivity. For instance, the confinement of SnO₂ NPs inside few-walled carbon nanotubes can contribute to an excellent cycling performance since the volume restriction inside carbon nanotubes can efficiently inhibit breakage during lithium insertion/extraction.⁵ However, the traditional strategies to prepare these composites need either complicated synthesis or relatively high cost. The simple processes towards scaling up production of TMOs/carbon anodes are still highly desirable.

Cellulose is a linear homopolymer with high molecular weight composed of β-D-glucopyranose units connected through β-(1-4) linkages, which is a key source of sustainable and non-toxicity materials.⁶ It is a green strategy to produce highly functionalized carbonaceous materials by means of the thermal or hydrothermal carbonization of cellulose.^{7,8} Cellulose could be modified and functionalized to carboxymethyl cellulose (CMC) directly by replacing the hydroxyl groups (–OH) groups with carboxymethyl (–CH₂COO[–]) groups.⁹ The

negatively charged carboxylate (–COO[–]) groups in the two adjacent chains of CMC or intra-macromolecular could coordinate with metal ions (Mⁿ⁺), such as Co²⁺, Ni²⁺, and Fe³⁺, to form Mⁿ⁺-CMC through ion-exchange process, for the applications of biolabeling and flameresistant materials.^{10–12} After pyrolysis of the Mⁿ⁺-CMC in inert gas atmosphere, it is expected to obtain a TMOs NPs/carbon composite. In this communication, –COO[–] surface groups were firstly created on the regenerated cellulose microfibrils (RCFs) to bind Co²⁺ cations (Co-RCFs). After heat treatment at 800 °C in N₂ and subsequent oxidation in air, the Co-RCFs converted to a novel nano/micro hierarchical structure in which Co₃O₄ NPs (~50–100 nm) were embedded in the surface of the carbonaceous fibres (Co₃O₄/CF-X, X denotes postoxidation time). The nanoconfinement effect on the Co₃O₄ NPs constrained by the carbonaceous fibre matrix could efficiently enhance the cycling performance.^{13–15} So as expected, the composite anode shows a high reversible capacity of 730 mAh g⁻¹ at 89 mA g⁻¹ after 100 cycles as well as excellent rate capability. Apparently, this simple and environmentally friendly strategy is more feasible to synthesize a TMOs/carbon LIBs anode on a large scale compared with the conventional chemical vapor deposition and hydrothermal methods.

Scheme. 1 The synthesis of Co₃O₄/CF-X from surface modified RCFs.

As illustrated in Scheme 1, the synthesis of the $\text{Co}_3\text{O}_4/\text{CF-X}$ samples was involved in a three-step process. Firstly, the RCFs were fabricated via wet-spinning technique by using the spinning dope of aqueous solution of RC and sodium hydroxide (NaOH) (Fig. S1). Secondly, the obtained RCFs were soaked and vibrated in NaOH/ethanol solution, and then chloroacetic acid (ClCH_2COOH) was added to convert the surface $-\text{OH}$ of RCFs into $-\text{CH}_2\text{COO}^-$ via typical etherification reaction. The obtained surface modified RCFs were soaked in 5 wt% cobalt chloride (CoCl_2) ethanol solution to capture Co^{2+} cations electrostatically on the fibers (Co-RCFs), resulting in the white RCFs turn to light purple. The weight percentage of Co species in the Co-RCFs is determined to be about 2.17% by using thermogravimetric analysis (TGA) (Fig. S2). Finally, after calcination of 800°C for 1 h in N_2 atmosphere and subsequent oxidation in air at 400°C for 30, 60, and 90 min, the Co-RCFs converted to $\text{Co}_3\text{O}_4/\text{CF-X}$.

Fig. 1 (a) XRD patterns of RCF, Co-RCF, $\text{Co}_3\text{O}_4/\text{CF-30}$, $\text{Co}_3\text{O}_4/\text{CF-60}$, and $\text{Co}_3\text{O}_4/\text{CF-90}$. (b) FTIR spectra of the RCF, Co-RCF, $\text{Co}_3\text{O}_4/\text{CF-60}$, and commercial Co_3O_4 .

Fig. 1a shows the X-ray diffraction (XRD) patterns of RCFs, Co-RCFs, and $\text{Co}_3\text{O}_4/\text{CF-X}$. RCFs have the typical XRD pattern of cellulose II at 2θ value of $12.1^\circ, 20.1^\circ$, and 21.8° , and the crystalline polymorph of RCFs was formed during the wet-spinning process.¹⁶ The same diffraction peaks are also observed from the diffraction pattern of the Co-RCFs, indicating the cellulose II structure of the fibres is maintained. After calcination in N_2 atmosphere and subsequent oxidation in air, the Co^{2+} cations completely convert to Co_3O_4 species. The strong and sharp diffraction peaks at 2θ values of $31.3^\circ, 36.9^\circ, 44.9^\circ, 59.5^\circ$, and 65.3° are observed from the XRD pattern of $\text{Co}_3\text{O}_4/\text{CF-X}$ samples, corresponding to the diffraction of (220), (311), (400), (511), and (440) planes of cubic crystalline Co_3O_4 (JCPDS No. 42-1467). The peak intensity increases with the extension of postoxidation time from 30 to 60 and 90 min. In addition, the broad diffraction peak of amorphous carbon ($2\theta = 25.0^\circ$) is resulted from the pyrolysis of RCFs, which disappears after long time oxidation ($\text{Co}_3\text{O}_4/\text{CF-90}$). Fourier transformed infrared spectroscopy (FTIR) of the RCFs, Co-RCFs, $\text{Co}_3\text{O}_4/\text{CF-60}$, and commercial Co_3O_4 are displayed in Fig. 1b. For Co-RCFs, two characteristic peaks at 1750 cm^{-1} and 1235 cm^{-1} are ascribed to the stretching vibration of $\text{C}=\text{O}$ and COO^- ester functional groups,^{17, 18} respectively, indicating that the surface of RCFs has been modified by etherification reaction. After

pyrolysis and postoxidation, the Co-O stretching vibration bands of Co_3O_4 appear at 663 and 570 cm^{-1} in the spectrum of $\text{Co}_3\text{O}_4/\text{CF-60}$. The N_2 absorption-desorption isotherms and pore size distributions obtained from the Barrett-Joyner-Halenda (BJH) method are shown in Fig. S3. $\text{Co}_3\text{O}_4/\text{CF-60}$ exhibits a high specific surface area of $320\text{ m}^2\text{ g}^{-1}$, larger than those of $\text{Co}_3\text{O}_4/\text{CF-30}$ ($214\text{ m}^2\text{ g}^{-1}$), and $\text{Co}_3\text{O}_4/\text{CF-90}$ ($178\text{ m}^2\text{ g}^{-1}$). And $\text{Co}_3\text{O}_4/\text{CF-60}$ displays a multi-modal pore size distribution at $2.5, 9.0$, and 63.2 nm . The high specific surface and porous structure could increase the contact of active material with electrolyte, shorten Li^+ diffusion path and thus improve the electrochemical performance.¹⁹⁻²¹

The morphology of the RCFs, Co-RCFs and $\text{Co}_3\text{O}_4/\text{CF-X}$ was studied by field emission scanning electron microscope (FESEM). As shown in Fig. 2a, a single RCF is straight and smooth with a diameter of $\sim 15\text{ }\mu\text{m}$. After capture of Co^{2+} cations, the single Co-RCF possesses similar morphology with RCF (Fig. 2b), indicating that the introduction of the Co^{2+} cations does not influence the appearance of the fibres. After pyrolysis of Co-RCF in N_2 and oxidation in air, a large amount of Co_3O_4 NPs in the size range of $\sim 50\text{--}100\text{ nm}$ are homogeneously embedded on the surface of the fibres (Fig. 2c and Fig. S4). The magnified SEM image of $\text{Co}_3\text{O}_4/\text{CF-60}$ in Fig. 2d shows the highly porous structure close to the Co_3O_4 NPs, which can serve as the electrolyte pathway to ensure good electrolyte immersion with the Co_3O_4 NPs (Fig. 2d). In addition, elemental mapping by energy dispersive X-ray spectroscopy (EDS) was used to analyse the distribution of the C, Co, and O elements in the fibre. As shown in Fig. 2e, homogeneously distributed C, Co, and O elements can be detected in a single fibre of $\text{Co}_3\text{O}_4/\text{CF-60}$.

Fig. 2 SEM images of (a) RCF, (b) Co-RCF, (c, d) $\text{Co}_3\text{O}_4/\text{CF-60}$. (e) SEM image of $\text{Co}_3\text{O}_4/\text{CF-60}$ and corresponding EDS mapping for C, Co, O elements.

Fig. 3 (a) Low-magnification TEM, (b) high-magnification TEM, and (c) HRTEM image of $\text{Co}_3\text{O}_4/\text{CF}-60$.

Transmission electron microscope (TEM) and high-resolution TEM (HRTEM) images of $\text{Co}_3\text{O}_4/\text{CF}-60$ are presented in Fig. 3. As shown in Fig. 3a, a large number of Co_3O_4 NPs are distributed throughout a single fibre, and macropores are formed via the aggregation of large Co_3O_4 NPs. More importantly, it is notable that carbon matrix resulting from the carbonization of the $-\text{CH}_2\text{COO}^-$ groups is adjacent to the Co_3O_4 NPs (Fig. 3b), and such Co_3O_4 NPs-carbon nanoconfinement structure is efficient to increase the conductivity of the Co_3O_4 NPs and to counteract the pulverization of the Co_3O_4 NPs that takes place during charge/discharge cycles. Fig. 3c further shows a HRTEM image of the Co_3O_4 NPs. Apparently, the plane of $\{110\}$ of cubic Co_3O_4 is exposed, which corresponds to the $[011]$ zone-axis of Co_3O_4 NPs. The $\{110\}$ plane has a high density of atomic steps which is helpful to enhance electrochemical performance.^{22, 23}

Cyclic voltammetry (CV) and galvanostatic cycling analysis were used to examine the electrochemical properties of $\text{Co}_3\text{O}_4/\text{CF}-X$ as LIBs anodes. Co_3O_4 exhibits high theoretical capacity according to reduction/oxidation reactions ($\text{Co}_3\text{O}_4 + 8\text{Li}^+ + 8\text{e}^- \leftrightarrow 4\text{Li}_2\text{O} + 3\text{Co}$). Fig. 4a shows the CV curves of $\text{Co}_3\text{O}_4/\text{CF}-60$ at a scan rate of 0.1 mV s^{-1} . In the first cathodic scan, two cathodic peaks were observed at 0.67 and 0.85 V, corresponding to the formation of solid-electrolyte-interphase (SEI) film and electrochemical reduction (lithiation) reaction of Co_3O_4 with Li^+ . The broad anodic peak at 2.07 V can be ascribed to the oxidation (delithiation) reaction of Co_3O_4 . In the second cycle, the main oxidation peak shifts to 2.18 V, and the cathodic peak is observed at 1.16 V. The peak intensity and integral areas of the third cycle are close to that of the second one for $\text{Co}_3\text{O}_4/\text{CF}-60$, indicating the electrochemical reversibility is gradually built after the initial cycle. Fig. 4b shows the charge/discharge curves of the $\text{Co}_3\text{O}_4/\text{CF}-60$ at 89 mA g^{-1} in the voltage range of 0.01–3.0 V (vs. Li^+/Li) at the 1st, 2nd, 50th and 100th cycle, respectively. The first discharge and charge specific capacities of $\text{Co}_3\text{O}_4/\text{CF}-60$ are 1246 and 733 mAh g^{-1} with a Coulombic efficiency (CE) of 58.8%. The restricted CE during the first cycle is assigned to the decomposition of electrolyte, forming a SEI on the electrode surface, and to the irreversible insertion of Li ions into pores of the fibers. Nevertheless, the CE value rises to 99.1% in the fifth

Fig. 4 (a) CV curves of $\text{Co}_3\text{O}_4/\text{CF}-60$. (b) The $\text{Co}_3\text{O}_4/\text{CF}-60$ cycled at the 1st, 2nd, 50th, and 100th between 0.01 and 3.0 V (vs. Li^+/Li) at a current density of 89 mA g^{-1} . (c) Comparison of the cycling performance of $\text{Co}_3\text{O}_4/\text{CF}-30$, $\text{Co}_3\text{O}_4/\text{CF}-60$, $\text{Co}_3\text{O}_4/\text{CF}-90$, and the pure CF. (d) The reversible specific capacity of the $\text{Co}_3\text{O}_4/\text{CF}-60$ at different rates.

cycle and then remains above 98% in the following cycles (as shown in Fig. S5). In addition, after 50th and 100th charge/discharge cycles, it exhibits a high reversible capacity of $\sim 730 \text{ mAh g}^{-1}$, indicating good cycling stability.

The electrochemical performance of the $\text{Co}_3\text{O}_4/\text{CF}-60$ was compared with $\text{Co}_3\text{O}_4/\text{CF}-30$, $\text{Co}_3\text{O}_4/\text{CF}-90$, pure CF obtained by pyrolysis of RCF, and commercial Co_3O_4 . As shown in Fig. 4c, $\text{Co}_3\text{O}_4/\text{CF}-60$ exhibits a high reversible specific capacity of 730 mAh g^{-1} at 89 mA g^{-1} after 100 cycles. This value is much higher than those of $\text{Co}_3\text{O}_4/\text{CF}-30$ (496 mAh g^{-1}) and pure CF (170 mAh g^{-1}). For $\text{Co}_3\text{O}_4/\text{CF}-90$ and commercial Co_3O_4 electrodes, their specific capacities fades seriously after 50 cycles (Fig. 4c and S6), although they exhibits high capacities at the initial stage. The best electrochemical performance of the $\text{Co}_3\text{O}_4/\text{CF}-60$ electrode results from the optimal content of carbon in the composite which is able to simultaneously insure both high specific capacity and cycling stability. As shown in Fig. S7, about 75.8, 32.1, and 9.9 % mass ratio of carbon are remained in $\text{Co}_3\text{O}_4/\text{CF}-30$, $\text{Co}_3\text{O}_4/\text{CF}-60$, and $\text{Co}_3\text{O}_4/\text{CF}-90$ after postoxidation, respectively.

The $\text{Co}_3\text{O}_4/\text{CF}-60$ electrode also exhibits the specific capacities of 632, 540, 484, 420, and 240 mAh g^{-1} at 178, 445, 890, 1780, to 4450 mA g^{-1} , respectively, indicating excellent cycling performance (Fig. 4d). In addition, a reversible capacity of 750 mAh g^{-1} was recovered, when the current density is decreased to 89 mA g^{-1} . This indicates that the integrity of the fibrous electrode has been preserved even after experiencing high rate charge/discharge. As shown in Fig. S8, there is no obvious change in morphology of the $\text{Co}_3\text{O}_4/\text{CF}-60$ electrode before and after 100 galvanostatic charge-discharge cycles. We also compared the electrochemical performance of the $\text{Co}_3\text{O}_4/\text{CF}-60$ electrode with the reported $\text{Co}_3\text{O}_4/\text{carbon}$ composite electrodes^{24,25} (Fig. S9). Besides the high specific

capacity, the $\text{Co}_3\text{O}_4/\text{CF}-60$ electrode displays better cycling performance.

The high reversible specific capacity, excellent cycling stability and rate performance should be attributed to the nanoconfinement effect on the embedded Co_3O_4 NPs by the carbonaceous fibre matrix. First, the unique one dimensional carbonaceous fibre matrix is employed as effective “matrix” material to release the mechanical strain induced during Li^+ charge/discharge process and prevent the aggregation of Co_3O_4 NPs, and can serve as the channels for fast electron transport to improve the electronic transport, thus enhance the cycle and rate performance. Second, the macropores and mesopores in the carbonaceous fibre can store the electrolyte as the “reservoirs” to increase the interface between active materials and electrolyte, thus shortening the diffusion distance of Li^+ ions.²⁶⁻²⁸

In summary, we have prepared novel $\text{Co}_3\text{O}_4/\text{Carbon}$ fibres by simple pyrolysis of wet-spun modified cobalt cellulose fibres. The composite possesses the unique hierarchical microstructure in which Co_3O_4 NPs are embedded in micro-sized carbonaceous fibrous matrix and exhibits excellent electrochemical performance when used as an anode for Li-ion batteries. $\text{Co}_3\text{O}_4/\text{CF}-60$ obtained under the optimum conditions (oxidized with 60 min) exhibits high initial reversible specific capacity, excellent cycling stability, i.e. 730 mAh g^{-1} after 100 cycles at 89 mA g^{-1} , and good rate performance. The excellent electrochemical performance is attributed to its unique nanoconfinement structure where the synergistic effect of Co_3O_4 NPs and bifunctional carbon fibre matrix (confining scaffold and conducting network), which greatly enhances the reversible capacity, cycling stability and rate performance. This work may pave a new way for scalable synthesis of nanoconfined TMO/Carbon anode materials.

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Notes and references

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[†]Electronic Supplementary Information (ESI) available. Experimental Section, wet spinning process, SEM images of $\text{Co}_3\text{O}_4/\text{CF}-30$ and $\text{Co}_3\text{O}_4/\text{CF}-90$, N_2 adsorption-desorption isotherms and BJH pore size distribution analysis, TGA measurement, and electrochemical properties of commercial Co_3O_4 electrodes. See DOI: 10.1039/b000000x/.

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