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Rational design of walnut-like ZnO/Co₃O₄ porous nanospheres with substantially enhanced lithium storage performance†

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Rational fabrication and smart design of multi-component anode materials to achieve desirable reversible capacities and exceptional cyclability are significant for lithium-ion batteries (LIBs). Herein, walnut-like ZnO/Co₃O₄ porous nanospheres were prepared by a facile solvothermal method, which were then applied as a mechanically stable anode material for LIBs. The rationally designed hybridized electrode brings favorable structural features, particularly ZnO/Co₃O₄ porous nanospheres with abundant vacant space and enhanced surface area, enhancing lithium/electron transport and relieving volumetric stresses during the cycling process. Moreover, several *in situ* hybridized anode materials with electrochemical cooperation further overcome the challenge of capacity decay and conductivity deficiency. The as-obtained ZnO/Co₃O₄ delivered a much better lithium storage performance compared with ZnO, Co₃O₄, and their physical mix. We believe that the novel design criteria will bring opportunities in exploration and promote the practical application of transition metal oxides.

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1. Introduction

To date, lithium-ion batteries (LIBs) are one of the major energy sources.^{1–3} With the rapid development of modern devices such as large-scale energy storage devices and electric vehicles, further improvement in energy/power densities, safety and lifespan are required for LIB.⁴ Graphitic materials are the main commercial anodes because of their chemical stability, low operating voltage *versus* Li/Li⁺ and low cost, but are still limited by their poor theoretical specific capacities (LiC₆, 372 mA h g⁻¹).⁵ Therefore, it is significant to explore advanced anode materials for LIBs.

Transition metal oxides (TMOs) hold substantial promise for LIBs since they can reversibly accommodate Li⁺ through conversion or alloying reactions, which involve the formation and decomposition of lithium oxide.^{6–8} Creating chemical bonds through chemical reactions between Li⁺ and active materials is conducive to high-performance anode materials, in which multi-electron transfer occurs and superior reversible capacity can be expected.⁶ However, their practical application in LIBs is limited by a short cycling life due to several intrinsic reasons: severe volume variation during lithiation/delithiation, pulverization of electrode active materials and slow ionic/electronic conductivity.^{6,9–11} Therefore, it is of great significance to surpass the above-mentioned issues and boost their electrochemical performance. For TMOs, nanostructure engineering and hybridization of different phases can be efficacious strategies.^{12–15}

Regarding nanostructure engineering, fabricating porous particles at the nano/microscopic level can effectively recover the lithium storage performance.^{16–23} For example, Wang *et al.* successfully prepared porous CoMoO₄ nanorods that exhibited a capacity of 603 mA h g⁻¹ at 400 mA g⁻¹.²⁴ Due to their extra void space and large surface area, these specially designed structures counteract structural strains upon cycling, enlarge electrode/electrolyte interface to offer more reactive sites, and promote Li⁺ diffusion. Besides nanostructure engineering, hybridization of several active materials has been reported as another promising strategy, since the heterojunctions are beneficial for electron/ion diffusion, mechanical stability and redox reaction kinetics.^{6,25} Both charge carrier transport and interfacial reaction kinetics can be enhanced by building heterostructures composed of several coupling components with different bandgaps.²⁶ In this regard, Fe₂O₃/MnO₂²⁷ and Co₃O₄/CoMoO₄²⁸ have been reported to show extraordinary lithium storage performance. Taken together, achieving synchronous combination of structural engineering at the nano/microscopic level and hybridization of TMOs will be an effective path towards LIBs with high lithium storage performance.

Among various TMO candidates, ZnO has a relatively high capacity (978 mA h g⁻¹) because Zn forms an alloy material with Li, indicating a larger amount of Li ions captured.^{6,9} It has been reported by Tarascon *et al.* that the “over capacity” was associated to the growth of SEI films and TEM was conducted to deduce this. In addition, they confirmed the catalytic role of Co in the generation of SEI films.²⁹ Therefore, it is promising to fabricate Co₃O₄-containing ZnO hybrid materials. However, fabricating *in situ* hybridized nanomaterials through one-pot synthesis while keeping an advantageous architecture is hard to achieve.

Herein, walnut-like ZnO/Co₃O₄ porous nanospheres were synthesized for the first time by a solvothermal route, possessing great structural advantages (Fig. 1). Specifically, the structure of porous nanospheres with ravines can benefit the transport of lithium ions, provide abundant electrochemically active

sites, and offer structural flexibility as it prevents the electrode from collision or deformation arising from huge volumetric change. Moreover, the *in situ* hybridization of ZnO and Co₃O₄ brings a synergistic effect, which endows the electrode with improved electronic conductivity and reinforced structural stability. The as-prepared ZnO/Co₃O₄ porous nanospheres exhibited enhanced electrochemical activity, superior cycling stability and high specific capability. The reported one-pot colloidal synthesis route is promising and can be extended to design other multi-component hierarchical TMO nanostructures.

2. Materials and methods

2.1. Synthesis

1.2 g Zn(NO₃)₂·6H₂O (Alfa Aesar), 0.8 g Co(NO₃)₂·6H₂O (J. T. Baker) and 1 g PVP (Alfa Aesar) were dissolved and stirred in methanol/DI water (86/10 mL) solution for 40 min, which was then heated in an autoclave at 180 °C for 6 h. The resulting precipitates were collected and washed with DI water five times, followed by drying and annealing in air at 450 °C for 40 min.

For comparison, ZnO and Co₃O₄ were prepared using 2.0 g Zn(NO₃)₂·6H₂O or 2.0 g Co(NO₃)₂·6H₂O. In addition, physically mixed ZnO/Co₃O₄-PM (weight ratio = 3 : 2) was prepared by mixing as-obtained ZnO and Co₃O₄.

2.2. Characterization

The morphologies and structural aspects were obtained by transmission electron microscopy (TEM, Hitachi HT7700), high-resolution transmission electron microscopy (HRTEM, Hitachi HD2700), and scanning electron microscopy (SEM, Hitachi SU8230). The crystal structures were determined by Bruker D8 Advanced Powder XRD. X-ray photoelectron spectroscopy (XPS) was achieved with the Thermo K-Alpha XPS system. The pore features and Brunauer–Emmett–Teller surface area were characterized by physisorption (Microtrac BEL Corporation).

2.3. Electrochemical measurements

Active materials, super P carbon black (MTI Corporation) and sodium carboxymethyl cellulose (*M_w* 250 000, Sigma Aldrich) were mixed with a weight ratio of 8 : 1 : 1 in DI water. The slurry was then cast on copper foil, followed by dehydration at 120 °C overnight. Cells were assembled utilizing Li metal as the counter electrode, 1 M LiPF₆ in EC/EDC (v/v = 1 : 1, Sigma Aldrich) as the electrolyte, and Celgard 2400 as the separator. Cyclic voltammetry (CV) was performed between 3.0 and 0.01 V at 0.2 mV s⁻¹. Electrochemical impedance spectroscopy (EIS) measurements were measured between 100 kHz and 0.01 Hz.

3. Results and discussion

The synthetic procedure of walnut-like ZnO/Co₃O₄ involved two steps, a solvothermal method using PVP as the surfactant,

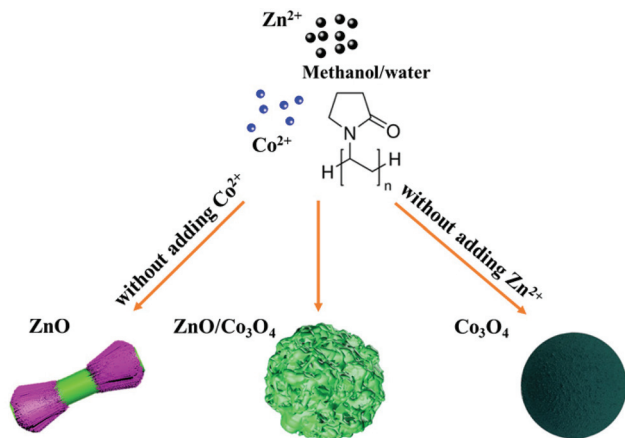


Fig. 1 Diagram for the fabrication of ZnO, Co₃O₄ and ZnO/Co₃O₄.

followed by an annealing process at 450 °C in the air for 40 min. Fig. S1† shows the SEM images of ZnO/Co₃O₄ samples obtained under different reaction time (1, 2 and 3 h), respectively. The formation of walnut-like ZnO/Co₃O₄ nanospheres goes through three stages: (1) the generation of spherical particles by the solvothermal reactions between metal nitrates and PVP; (2) the capping of PVP: sufficient PVP forms a continuous layer to warp the whole surface of nanospheres and promotes the growth in all direction. As the reaction progresses, relatively excessive metal nitrates and insufficient dosages of PVP are not able to form a continuous outer layer of the nuclei, resulting in the growth of exterior nanorods in the lateral directions and the walnut-like surface is generated; (3) the growth of walnut-like nanospheres *via* Ostwald Ripening and finally reach the best stabilization status. The annealing temperature was determined through TGA analysis (Fig. S2†). Here, 450 °C was chosen as the annealing temperature to ensure a complete decomposition of precursors. A mass loss (~5.6%) can be observed below 400 °C, which can be assigned to the decomposition of organic content and evaporation of the absorbed solvent.³⁰ The formation of ravines was observed after annealing, resulting in a walnut-like morphology. Besides, the surface became rugged and porous, arising from the elimination of organics during the annealing process.⁶ When the annealing temperature further increases to 550 and 650 °C, the collapse of 3D structure and formation of debris

could be observed (Fig. S3†). Thus, the annealing temperature was set at 450 °C. It is interesting to find that ZnO and Co₃O₄ displayed significantly different morphologies: dumbbell-like ZnO and sphere-like Co₃O₄, both featuring a relatively smooth surface, suggesting that both Zn²⁺ and Co²⁺ are significant to generate walnut-like ZnO/Co₃O₄ (Fig. 1).

In Fig. S4,† 0.4 g of ZnO/Co₃O₄ and graphite were shown. Compared with commercial graphite, ZnO/Co₃O₄ occupied less volume with a larger packing density, which is important for LIBs to achieve high volumetric energy density.^{6,31} Thus, ZnO/Co₃O₄ is promising to replace graphite when applied as anode for LIBs.

TEM and SEM were performed on ZnO/Co₃O₄, ZnO, Co₃O₄ and their corresponding solvothermal precursors. TEM images in Fig. 2 confirmed the porous feature of ZnO/Co₃O₄, which could effectively alleviate volume change upon cycling and promote electrolyte penetration. As shown in Fig. 3, the panoramic overview demonstrated the successful synthesis of highly uniform and dispersed ZnO/Co₃O₄ consisting of walnut-like porous nanospheres with a diameter of around 250 nm. Compared with solvothermal precursors (Fig. S5 and S6†), the morphology of ZnO/Co₃O₄ was well-maintained, but with a relatively rougher surface and deeper ravines. The uniform distribution of Zn and Co elements in our ZnO/Co₃O₄ is important for the battery performance and was confirmed using scanning transmission electron microscopy (STEM)

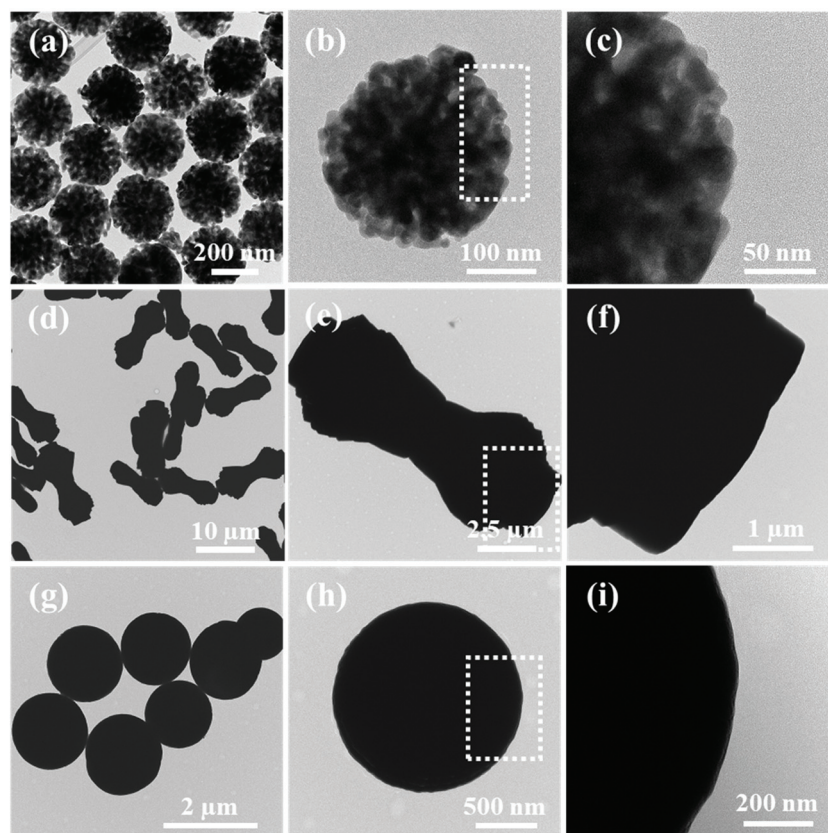


Fig. 2 TEM images of (a–c) ZnO/Co₃O₄, (d–f) ZnO and (g–i) Co₃O₄.

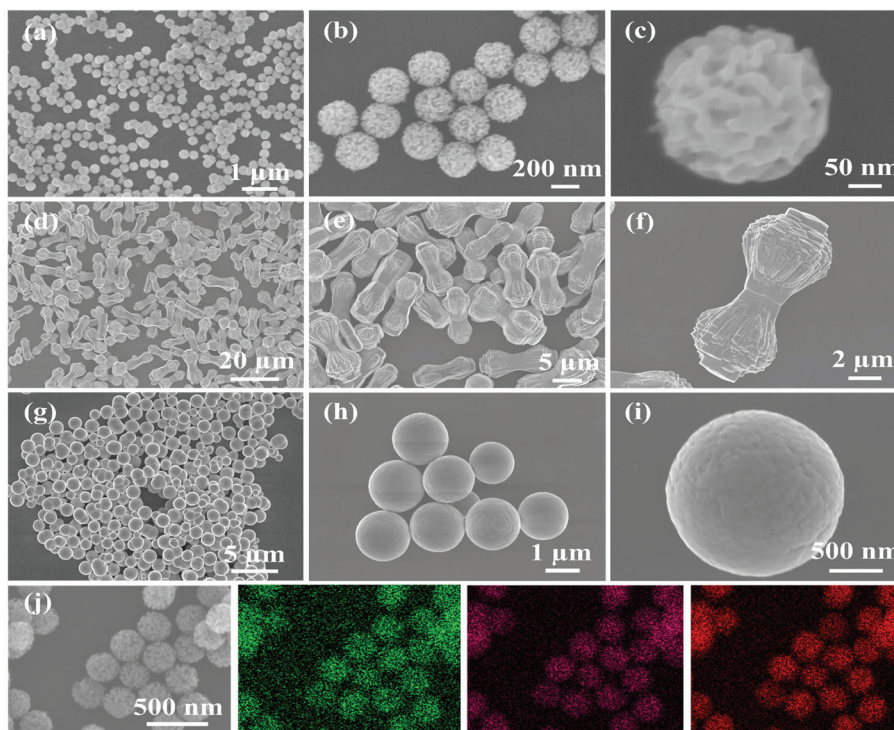


Fig. 3 SEM images of (a–c) ZnO/Co₃O₄, (d–f) ZnO and (g–i) Co₃O₄. (j) Spatial distributions of Zn, Co and O in ZnO/Co₃O₄.

image and elemental mappings (Fig. S7†). Also, HRTEM image (Fig. S8†), shows close proximity between small domains of ZnO and Co₃O₄. As a result, a strong E-field was introduced by large energy bandgap difference between ZnO and Co₃O₄ (2.07 eV for ZnO, and 3.2 eV for Co₃O₄), thus electron/ion diffusion and surface redox reaction kinetics were facilitated.^{25,32} As shown in Fig. 2 and 3, ZnO exhibited a dumbbell morphology with a large aspect ratio, while Co₃O₄ exhibited a spherical morphology. Their relatively smooth surfaces limit their lithium storage performances.

As shown in Fig. 4a, peaks at 31.9°, 34.6°, 36.4°, 47.7°, 56.8° and 63.0° were assigned to (1 0 0), (0 0 2), (1 0 1), (1 0 2), (1 1 0) and (1 0 3) crystal planes of ZnO, and the remaining peaks located at 19.0°, 31.4°, 36.9°, 44.9°, 59.5° and 65.4° corresponded to the (1 1 1), (2 2 0), (3 1 1), (4 0 0), (5 1 1) and (4 4 0) planes of well-crystallized Co₃O₄.^{6,33} XPS was performed to determine elemental valence states. As can be seen from Fig. 4b, the wide XPS survey spectrum confirmed the existence of Zn, Co and O. Meanwhile, two peaks at 1044.5 and 1021.4 eV in the high-resolution Zn 2p spectrum (Fig. 4c) were related with Zn 2p_{1/2} and Zn 2p_{3/2}, implying the existence of Zn²⁺.⁶ Co 2p spectrum (Fig. 4d) showed two couples of peaks at 795.1 and 779.9 eV, corresponding to Co 2p_{1/2} and Co 2p_{3/2}, respectively, suggesting the formation of Co₃O₄.³³ These XPS analyses are consistent with the XRD result. Specific surface area and porous structural characteristics influence the lithium storage performance of electrode materials,³⁴ thus the porous nature of ZnO/Co₃O₄ was investigated by Brunauer–Emmett–Teller analysis (BET). As observed in Fig. 4e and f, ZnO/Co₃O₄ exhibi-

ted a porous feature, and the specific surface area was 17.55 m² g⁻¹, much larger compared with ZnO and Co₃O₄ (0.75 and 2.91 m² g⁻¹, respectively) (Table S1†). Large surface area and rich pores bring effective electrode/electrolyte interaction and offer effective pathways for ion/electron diffusion.²⁵

The electrochemical performance of ZnO, Co₃O₄ and ZnO/Co₃O₄ were evaluated. CV curves of ZnO/Co₃O₄ exhibited a rich redox reaction (Fig. 5a). Peaks from 0 to 0.4 V were caused by Li–Zn alloying process.^{35,36} The peak at 0.62 V was related to the formation of metallic Co, Li₂O and SEI film.^{33,35} Meanwhile, the peak at 1.71 V originated from the conversion reaction of Co to Co₃O₄.³⁵ Peaks at 2.11 V and 2.61 V corresponded to the reoxidation of Zn.³⁵ The shift of anode peaks arose from structural reorganization and polarization of electrode materials.³⁶ Evidently, the overlapped curves demonstrated extraordinary electrochemical reversibility. As indicated by the results of galvanostatic charge/discharge measurement shown in Fig. 5b, three obvious plateaus agreed well with CV curves. The well-overlapped discharge/charge profiles indicated good electrochemical stability. In contrast, as shown in Fig. 5c and d, both ZnO and Co₃O₄ delivered lower specific capacities compared with that of ZnO/Co₃O₄. Moreover, there was a severe capacity decay for ZnO, resulting from slow ion/electron transport.⁶

The cycling performances were evaluated at 0.1 A g⁻¹. As shown, discharge/charge capacities of ZnO/Co₃O₄ in the first cycle were 1352.3/1048.3 mA h g⁻¹ (Fig. 6a). The capacity decay arose from the generation of SEI and trapped Li-ions within

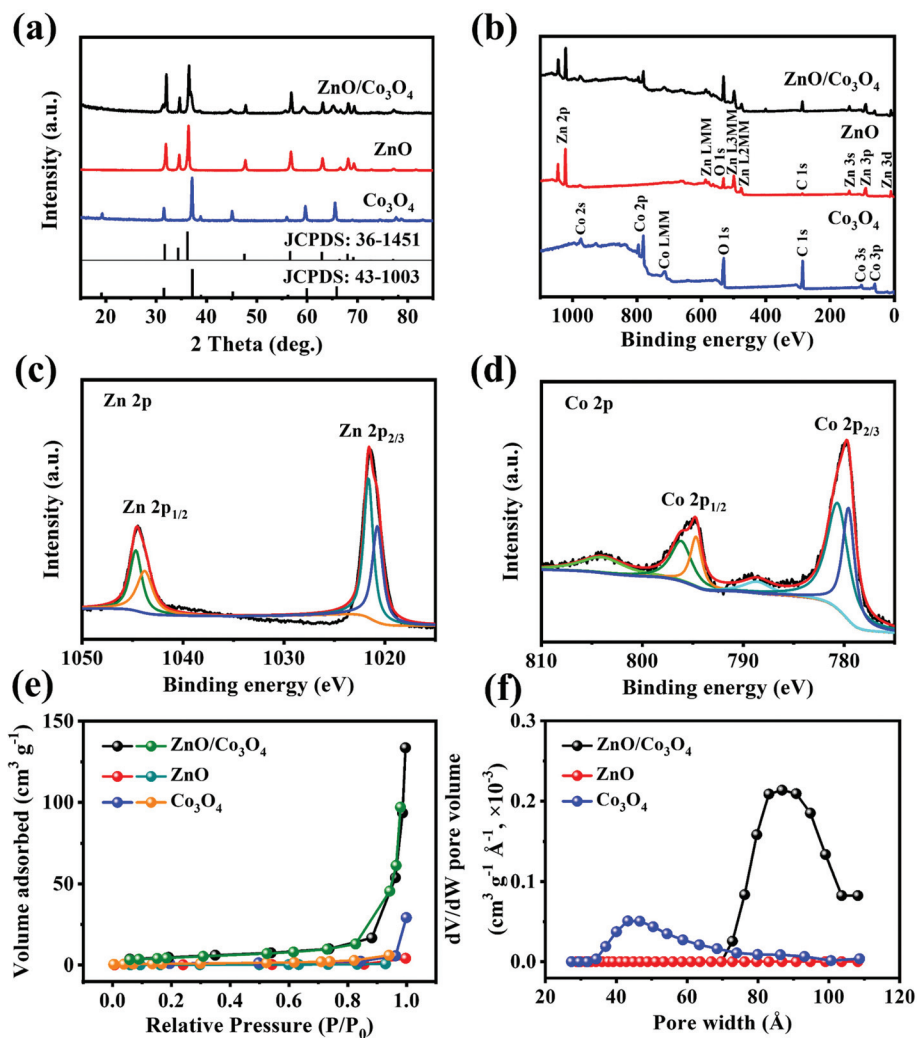


Fig. 4 (a) XRD and (b) XPS spectra of ZnO/Co₃O₄, ZnO and Co₃O₄. XPS spectra of elemental components (c) Zn and (d) Co elements in ZnO/Co₃O₄. (e) BET curves and (f) pore width distribution of ZnO/Co₃O₄, ZnO and Co₃O₄.

the electrode.⁶ After 10 cycles, the coulombic efficiency recovered to over 97%, demonstrating excellent reversibility. The alloying process of Zn with Li in ZnO/Co₃O₄ might aggravate the issue of low coulombic efficiency in the initial cycles. Thus, the volume ratio of ZnO was decreased to see if the low coulombic efficiency in the first cycles can be solved. As the ratio of ZnO and Co₃O₄ decreased from 3 : 2 to 1 : 8, the initial coulombic efficiency increased from 80.11% to 94.49%. When controlling the ratio of ZnO and Co₃O₄, mesoscale pore structure was observed to be adjusted. As shown in Fig. S11,[†] the composite exhibited a smoother surface with a less porous feature when there is more Co₃O₄, which was further confirmed by BET analysis (Table S2[†]). As shown in Fig. S12,[†] the cycling performance of composites with different ratio of ZnO and Co₃O₄ indicates that composites with larger surface area and richer pores exhibited higher discharge/charge capacities. In addition, it should be noticed that all ZnO/Co₃O₄ samples exhibited a gradual increase in capacity. This is a common

phenomenon for oxide electrodes, which can be attributed to several reasons: (1) the polymeric gel-like films generated from electrolyte decomposition participate in lithium storage and provide extra capacity;^{9,37} (2) the activation process improves the diffusion of lithium ions;¹³ (3) defects are introduced to electrode materials upon repeated lithiation/delithiation, bringing more active sites and promoting the reaction activity.^{6,33,38,39} After 100 cycles, the capacity retained at 1187.4 mA h g⁻¹. Unlike ZnO/Co₃O₄, the ZnO electrode exhibited a sharp capacity attenuation from 941.9 to 151.9 mA h g⁻¹, while the Co₃O₄ electrode delivered a relatively poor capacity of 746.9 mA h g⁻¹. The lithium storage performances of physically mixed ZnO and Co₃O₄ were also investigated. After 100 cycles, the capacity could only retain 244.1 mA h g⁻¹, highlighting the significance of *in situ* hybridization of ZnO and Co₃O₄. The presence of heterostructures generated from *in situ* hybridization of different components greatly enhanced interfacial reaction kinetics and charge carrier transport. Even at

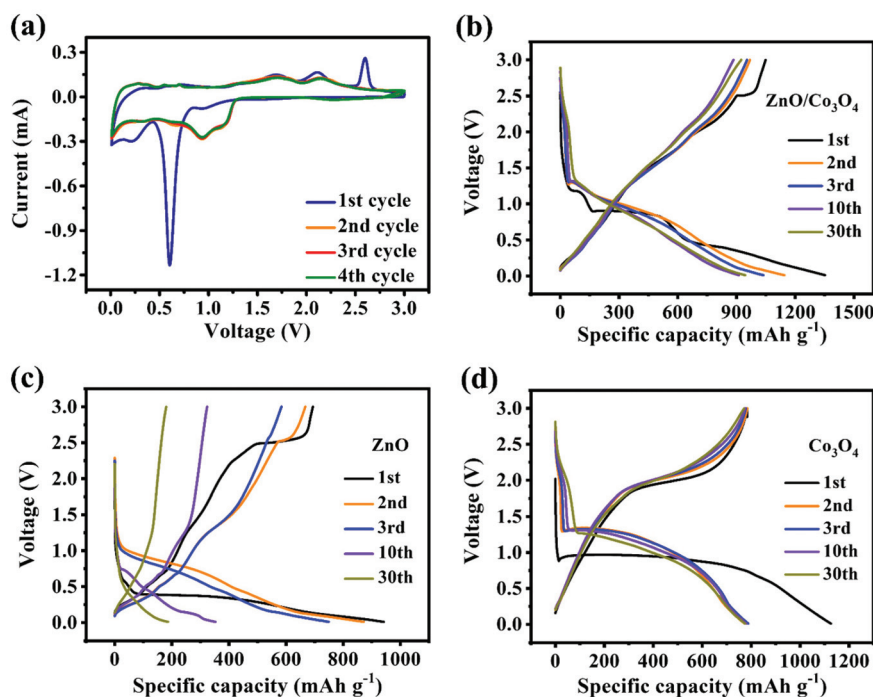


Fig. 5 (a) CV curves of ZnO/Co₃O₄ at 0.2 mV s⁻¹. Charge/discharge profiles at 0.1 A g⁻¹ for (b) ZnO/Co₃O₄, (c) ZnO and (d) Co₃O₄.

0.5 A g⁻¹ and 1 A g⁻¹, ZnO/Co₃O₄ electrode could still exhibit reversible capacities of 821.5 and 618.2 mA h g⁻¹ after 400 and 800 cycles, respectively (Fig. 6e). In contrast, the performances of ZnO, Co₃O₄ and ZnO/Co₃O₄-PM were unsatisfactory, arising from low surface area and large charge transfer resistance (Fig. S15†).

EIS analysis also confirmed the electrolyte/electrode activation process of ZnO/Co₃O₄. As shown in Fig. 6c and Fig. S16a,† there was an appreciable decrease in the interfacial resistance from 57.2 Ω to 53.9 Ω, indicating optimized electrochemical kinetics. The activation was related to several reasons: (1) the unique walnut structure is favorable for electron transport and ion diffusion; (2) the electrochemical reaction promotes electrolyte infiltration, thus reducing the electrode/electrolyte interfacial impedance.⁴⁰ ZnO/Co₃O₄ exhibited a smaller diameter of 53.9 nm, indicating lower charge transfer resistance, further confirming its structural advantage. As exhibited in Fig. S17 and S18,† intact nanospheres without any collapse and shedding could be detected for ZnO/Co₃O₄ after 100 cycles, suggesting extraordinary tolerance of lithiation/delithiation. These advantages contribute to enhanced reaction kinetics and electrochemical performance.

The lithium-ion diffusion coefficient was obtained from the equation below:⁴¹

$$D_{\text{Li}}^+ = R^2 T^2 / 2A^2 n^4 F^4 C^2 \sigma^2 \quad (1)$$

In the Nyquist plot, the linear curve is related with Li⁺ diffusion.^{30,42} σ was obtained to be 19.16 Ω s^{-1/2} from the fitted lines of Z' and $\omega^{-1/2}$ for ZnO/Co₃O₄ electrode after

cycling (Fig. 6d). The larger D_{Li}^+ of ZnO/Co₃O₄ indicates faster transport of electrons and ions, which was conducive to achieve excellent lithium storage performance.

The reversibility of ZnO/Co₃O₄ was further proved by *ex situ* XRD measurements (Fig. 6f). The peaks at 43.5°, 50.6° and 74.3° originated from Cu.⁴³ When the ZnO/Co₃O₄ was discharged to 1.2 V, peaks of ZnO and Co₃O₄ could be observed. No peaks related to ZnO and Co₃O₄ could be observed when further discharged, indicating their consumption during the discharge process. However, no diffraction peaks related to discharge products showed up, arising from their low crystallinity or amorphous states. This phenomenon has been reported in previous literature.⁶ At a fully charged state, the diffraction peaks of ZnO and Co₃O₄ reappeared, demonstrating their reversible generation.

The rate performance was examined at different current densities (Fig. 6b). ZnO/Co₃O₄ showed the most fascinating capacity retention ability compared with single components and their physical mix. The charge/discharge profiles were exhibited in Fig. S20.† As shown, ZnO/Co₃O₄ retained the voltage plateau well at all current densities. The excellent lithium storage performance of as-prepared walnut-like ZnO/Co₃O₄ porous nanospheres was superior to most reported ZnO, Co₃O₄ and ZnO/Co₃O₄ composites for LIBs (Fig. S21†).

The superior lithium storage performance of walnut-like ZnO/Co₃O₄ arises from several aspects: (1) the nanometer size enhances the reaction kinetics due to facilitated electron/ion transport, and eases the mechanical strain caused by volume variation. (2) The large specific surface area provides abundant reactive sites, which is more beneficial with capacitive behav-

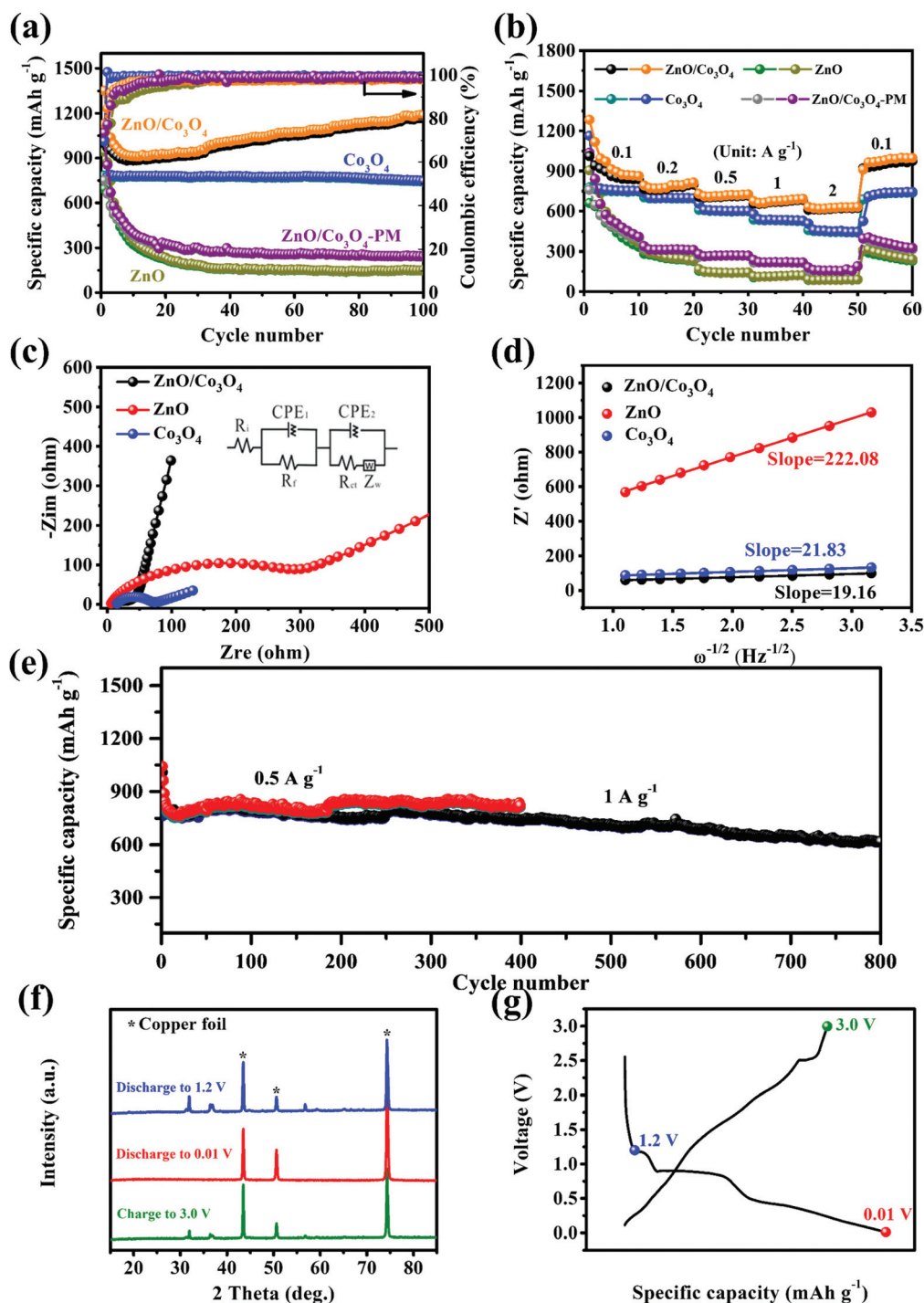


Fig. 6 (a and b) Cycling and rate performance for ZnO, Co_3O_4 , ZnO/ Co_3O_4 , and their physical mix. (c) Nyquist plots of ZnO/ Co_3O_4 , ZnO and Co_3O_4 after 100 cycles at 0.1 A g^{-1} and (d) fitted plot results of Z' versus $\omega^{-1/2}$. (e) Cycling performance of ZnO/ Co_3O_4 at large current densities. (f and g) Patterns of XRD analysis of ZnO/ Co_3O_4 at different states in the initial cycle.

ior, enhancing lithium storage and migration. (3) The rational hybridization of ZnO and Co_3O_4 endows the composite with high conductivity and structural stability, guaranteeing long-term cycling durability. (4) The abundant channels provided by the walnut-like structure facilitate ion/electron diffusion, thus ensuring fast charge transfer. (5) The space alleviates

huge volume variation throughout the cycling process, achieving excellent structural integrity and cycling durability.

To further understand kinetics characteristics of ZnO/ Co_3O_4 , ZnO and Co_3O_4 towards lithium storage, CV curves were collected at scan rates of 0.4, 0.6, 0.8 and 1.0 mV s^{-1} (Fig. 7a–c). In general, the current intensity increases as the

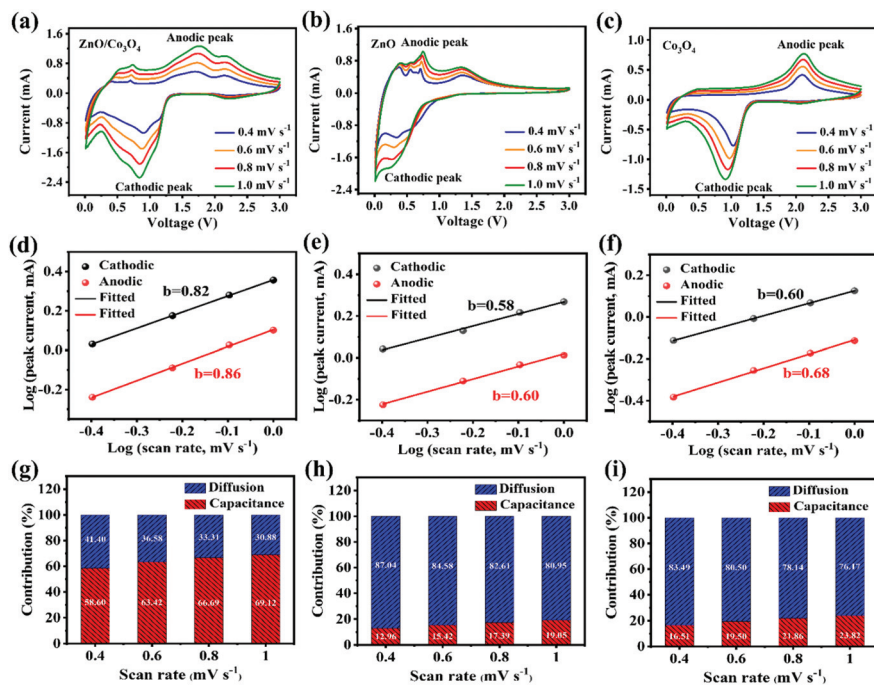


Fig. 7 CV curves of (a) ZnO/Co₃O₄, (b) ZnO and (c) Co₃O₄ at scan rates of 0.4, 0.6, 0.8 and 1.0 mV s⁻¹. Corresponding log *i* versus log *v* of (d) ZnO/Co₃O₄, (e) ZnO and (f) Co₃O₄. Diffusion/capacitance contribution ratio for (g) ZnO/Co₃O₄, (h) ZnO and (i) Co₃O₄.

scanning rate raises. The charge storage kinetics were evaluated by *b* from the equation below:⁴¹

$$i = av^b \quad (2)$$

For a capacitor-type electrode, the *b* value is 1.0, and for a battery-type electrode, the *b* value approaches 0.5. As revealed in Fig. 7d, the *b* values calculated based on eqn (1) were 0.82 and 0.86 for cathodic and anodic peaks, respectively, suggesting a capacitive dominant charge storage mechanism. As expected, the larger *b* value of ZnO/Co₃O₄ confirms that the 3D porous walnut-like structure was more favorable to Li⁺ storage occurring near the surface.

$$i(v) = k_1v + k_2v^{1/2} \quad (3)$$

Furthermore, the diffusion/capacitance contribution ratio can be quantitatively investigated according to eqn (3).⁴⁴ Fig. S22† reveals *k*₁ and *k*₂ values from *i(v)/v^{1/2}* versus *v^{1/2}* plot. As shown in Fig. 7, the capacitance contributions of ZnO/Co₃O₄ were 58.60, 63.42, 66.69 and 69.12 at different scan rates. The kinetics analysis demonstrated that diffusion-controlled and capacitive behaviors occurred simultaneously in ZnO/Co₃O₄. The capacitive process involves surface redox reactions, which are related to rapid electron diffusion, fast mass transport, and large accessible active sites. The amazing pseudocapacitive behavior of ZnO/Co₃O₄ arises from its unique walnut-like porous nanostructure, favorable conductive pathways, and large surface area, which is conducive to exert superior lithium storage performance.

4. Conclusions

In summary, this work developed an *in situ* fabrication of hybridized ZnO/Co₃O₄ porous nanospheres (*ca.* 200 nm in size) with mesoporous ravines on the surface. Such advantageous nanoarchitecture can provide several excellent properties: abundant pores to shorten the distance for Li⁺ diffusion, ultrafine ravines to accommodate the volume change and large surface area to facilitate surface redox reactions. Besides, the *in situ* fabrication of ZnO and Co₃O₄ brought a synergistic effect, and ZnO-Co₃O₄ heterostructure promoted redox kinetics and enhanced structural stability. Thus, assembled anode based on rationally designed walnut-like ZnO/Co₃O₄ composite exhibited high reversible capacity, excellent rate capability and impressive long-term cyclability. We believe that the bi-component ZnO/Co₃O₄ with walnut-like architecture is promising as a commercial electrode material for LIBs, and the facile strategy can be extended to design new TMO-based materials for energy storage applications.

Conflicts of interest

The authors declare no conflicts.

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