


 Cite this: *RSC Adv.*, 2021, 11, 1261

Ultra-small Fe₃O₄ nanodots encapsulated in layered carbon nanosheets with fast kinetics for lithium/potassium-ion battery anodes†

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Iron oxides are regarded as promising anodes for both lithium-ion batteries (LIBs) and potassium-ion batteries (KIBs) due to their high theoretical capacity, abundant reserves, and low cost, but they are also facing great challenges due to the sluggish reaction kinetics, low electronic conductivity, huge volume change, and unstable electrode interphases. Moreover, iron oxides are normally prepared at high temperature, forming large particles because of Ostwald ripening, and exhibiting low electronic/ionic conductivity and unfavorable mechanical stability. To address those issues, herein, we have synthesized ultra-small Fe₃O₄ nanodots encapsulated in layered carbon nanosheets (Fe₃O₄@LCS), using the coordination interaction between catechol and Fe³⁺, demonstrating fast reaction kinetics, high capacity, and typical capacitive-controlled electrochemical behaviors. Such Fe₃O₄@LCS nanocomposites were derived from coordination compounds with layered structures *via* van der Waals's force. Fe₃O₄@LCS-500 (annealed at 500 °C) nanocomposites have displayed attractive features of ultra-small particle size (~5 nm), high surface area, mesoporous and layered feature. When used as anodes, Fe₃O₄@LCS-500 nanocomposites delivered exceptional electrochemical performances of high reversible capacity, excellent cycle stability and rate performance for both LIBs and KIBs. Such exceptional performances are highly associated with features of Fe₃O₄@LCS-500 nanocomposites in shortening Li/K ion diffusion length, fast reaction kinetics, high electronic/ionic conductivity, and robust electrode interphase stability.

 Received 6th October 2020
 Accepted 14th December 2020

DOI: 10.1039/d0ra08503k

rsc.li/rsc-advances

Introduction

In the last three decades, rechargeable lithium-ion batteries (LIBs) have been widely used in portable electronic devices, stationary energy storage stations, and electric vehicles due to their high energy density, high coulombic efficiency, and long cycle life.^{1–5} However, the gap between limited element reserves and vast demands for Li resources is getting larger because of the large-scale application of LIBs.^{6–9} Therefore, the development of an alternative energy storage system is quite urgent.¹⁰ Potassium ion batteries (KIBs) have attracted wide interest due to even distribution, low cost, and similar electrochemical behaviors.¹¹ At first glance, KIBs have no advantage over LIBs, as the cation radius of K (0.138 nm) is larger than that of Li (0.076 nm), and has higher redox potential (−2.93 V) compared to Li

(−3.04 V).^{12–14} However, the abundance of K (2.09 wt%) on Earth is thousands of times higher than that of Li (0.0017 wt%), and more even distribution.^{15,16} In the last few years, extensive efforts have been devoted to demonstrate the application possibilities for both LIBs and KIBs, achieving many important progresses.^{11,17–19} However, great challenges are remained to develop suitable and cheap anode materials for both LIBs and KIBs with features of high capacity, good rate capability, and long cycle life.

Transition metal oxides (TMOs, M = Mn, Fe, Ni, Co, Cu, *etc.*) have attracted many attentions due to their high capacities (around 1000 mA h g^{−1}), natural abundance, and environmental friendliness.^{12,20–23} Among them, Fe₃O₄ is regarded as one of the most promising anode candidates for both LIBs and KIBs due to the high theoretical capacity (926 mA h g^{−1}),⁴ natural abundance, low cost, and non-toxic feature.^{24,25} However, there are still many challenges remained for Fe₃O₄ materials in LIBs and KIBs on the following aspects: (1) large volume changes during conversion reactions with Li/K cations, resulting in severe electrode pulverization and contact loss from current collectors; (2) poor ionic and electronic conductivity, leading to the low ionic diffusion coefficient and high electrical resistance; (3) unstable solid electrolyte interface (SEI), giving low coulombic efficiencies; (4) large particles size, relating to

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ra08503k

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long durations of chemical diffusion.^{26,27} To overcome these drawbacks, several effective strategies are reported to enhance the electrochemical performances, including nanometerization,^{19,28} carbon coating,^{29,30} heteroatom-doping,^{31–34} and surface-engineering.^{35–37} Nanometerization is normally applied for minimizing particle size and shortening diffusion pathways, but Ostwald ripening inevitably happens during hydrothermal and annealing treatments, leading to particle aggregations.³⁸ Carbon coating is one kind of universal route to immensely enhance the electronic conductivity, providing fast charge transferring and robust structural stability.³⁹ Heteroatom-doping is adopted on enhancing ionic conductivity for better rate capability.^{32,40} Surface-engineering works on the SEI layer with better mechanical stability, delivering long cycle life.^{3,41} Metal coordination compounds with high surface area and porous feature are considered as good precursors to synthesize electrode materials.^{3,42} Various TMOs materials with mesoporous and graphitic carbon skeleton, derived from metal coordination compound's precursors, are synthesized for anodes of LIBs/KIBs.^{43,44}

To enhance the rate capability and cycling performance of Fe₃O₄-based material, layered iron coordination compound with the catechol-iron coordination were synthesized to further prepare ultra-small Fe₃O₄ nanodots (~5 nm in average particle size) encapsulated in thin layered carbon sheets (~4 nm in thickness, labeled as Fe₃O₄@LCS). Such Fe₃O₄@LCS nanocomposites have displayed high surface area, thin-layered, and mesoporous structure. Different Fe₃O₄@LCS nanocomposites, obtained at different temperatures, were applied as anodes in LIBs/KIBs, demonstrating high reversible capacity, good cycle stability, and rate performance. These exceptional electrochemical behaviors are highly ascribed to the elaborately designed material features on shortening Li/K ion's diffusion length, fast reaction kinetics, high electronic conductivity, and robust structure stability.^{29,45}

Experimental

Chemicals

Catechol (C₆H₄(OH)₂, >98%), iron(III) nitrate hexahydrate (Fe(NO₃)₃·9H₂O, >98.5%), *N,N*-dimethylformamide (C₃H₇NO, DMF, >99.5%), ethanol (ET, >95%) were purchased from Sino-pharm Chemical Reagent Co. Ltd. All the chemicals were used without further purification. Deionized water (DI) was obtained from a Milli-Q System (Millipore).

Preparation of metal coordination precursor

In a typical synthesis procedure, catechol (0.11 g) was added into mixed solvent (9 mL) of DI/ET/DMF (1 : 1 : 1, v/v/v) with continuous stirrings for 15 min as solution A. Iron(III) nitrate hexahydrate (0.808 g) was dissolved in mixed solvents with the same composition as solution B. Solution B was added drop by drop in solution A with continuous stirrings for another 30 min, and then treated at 150 °C for 1 h using a single-mode microwave (NOVA-2S, 1000 W) radiation reactor with digital-controlled temperature and pressure apparatus. The products

were collected by centrifugation, washed with deionized water and ethanol for 3 times, and then dried at 60 °C overnight.

Preparation of Fe₃O₄/LCS nanomaterials

The precursors were placed in a tube furnace and heated to 400, 500, 600 °C for 2 h in N₂ with a ramp rate of 2 °C min⁻¹ respectively. After cooling, ultra-small Fe₃O₄ nanodots were successfully encapsulated in porous carbon nanosheets and marked as Fe₃O₄@LCS-400/500/600. As a comparison, Fe₃O₄ bulk was prepared *via* similar procedures without the addition of catechol.

Materials characterizations

The structural information and morphology were characterized by an X-ray diffractometer (XRD, Rigaku D/max-2550V) and field emission scanning electron microscope (FE-SEM, JSM-6700F) as well as transmission electron microscope (TEM, JEM-2010F). Raman spectra were collected on a laser Raman spectrometer (Renishaw). Elemental analysis was measured by an Elemental Analyzer (Vario MICRO cube). Surface area and porous properties were measured on a Micromeritics ASAP 2460 analyzer. X-ray photoelectron spectrometer (XPS, PHI ESCA-5000C) was utilized to evaluate the surface compositions of composites before and after cycles.

Electrochemical measurement

CR2032 coin cells were assembled in an Ar-filled glove box where the concentrations of H₂O and O₂ are kept below 0.1 ppm. Active materials (Fe₃O₄@LCS), conductive agent (acetylene black), and binder (polyvinylidene difluoride, PVDF) were uniformly dispersed in *N*-methyl pyrrolidone (NMP) (8 : 1 : 1, w/w/w) and coated onto copper foil, then dried at 60 °C overnight. The mass loading of working electrodes was about 1 mg cm⁻². All capacities were calculated based on the whole weight of electrodes. For LIBs, lithium foil was used as the counter electrode. 1 M LiPF₆ was dissolved in the mixed solvent (1 : 1, v/v) of ethylene carbonate (EC) and diethyl carbonate (DEC) as the electrolyte, and the microporous polypropylene (Celgard 2400) film was applied as the separator. However, potassium ion batteries were composed of pure potassium foil (counter electrode), 0.8 M KPF₆ in a mixed solvent of EC/DEC (1 : 1, v/v) as the electrolyte, and glass microfiber filter membrane (Whatman, Grade GF/A) as the separator.

The galvanostatic charge/discharge and the galvanostatic intermittent titration technique (GITT) measurements were evaluated by LAND-CT2001A at different current densities (from 100 mA g⁻¹ to 2000 mA g⁻¹) with a voltage range of 5 mV to 3 V. Cyclic voltammetry (CV) was measured on a CHI 660D electrochemical working station. Electrochemical impedance spectroscopy (EIS) was performed on an Autolab PGDSTAT101 electrochemical workstation with a frequency range of 0.1–1 × 10⁶ Hz.



Results and discussion

Structure and morphology investigation

A series of $\text{Fe}_3\text{O}_4@\text{LCS}$ nanocomposites were successfully prepared *via* a two-step process, as shown schematically in Fig. 1a. The Fe-precursor was firstly prepared by chemical coordination assisted by the microwave radiation at 150°C , which can be detected by XRD and FTIR as shown in Fig. S2.† Iron cations performed as coordinator center binding with catechol as a ligand, forming a layered and stacked structure *via* van der Waals' force.²⁵ With the homogeneously distributed iron coordinator, ultra-small Fe_3O_4 nanodots were eventually isolated and encapsulated in porous carbon nanosheets, which were derived from carbonized catechol during the annealing process in N_2 at 400°C , 500°C and 600°C , avoiding the particle growth of Ostwald ripening. Moreover, there are many voids in carbon sheets containing Fe_3O_4 nanodots formed by adjacent

atom losses, providing pre-reserved spaces for volume expansions.

$\text{Fe}_3\text{O}_4@\text{LCS}$ -500 nanocomposites display a typical layered morphology with mutually perpendicular structures in Fig. 1b and c. No apparent particle aggregation is observed in the TEM image, demonstrating the ideal particle distributions of Fe_3O_4 nanodots. More morphology information is shown in the ESI (Fig. S1†) for a whole glance in different conditions. The elemental mapping images in Fig. 1d–g further indicate the homogenous distributions of C, Fe, and O in the $\text{Fe}_3\text{O}_4@\text{LCS}$ -500 sample, illustrating the elaborately inherited architecture from layered Fe-coordination compound. The atomic percentages of all elements in $\text{Fe}_3\text{O}_4@\text{LCS}$ (400/500/600) nanocomposites are shown in Table S1† for elemental comparison.

The phase information and purity of the as-prepared samples were shown in XRD patterns in Fig. 1h. The diffraction peaks of $\text{Fe}_3\text{O}_4@\text{LCS}$ -400/500/600 samples are almost in

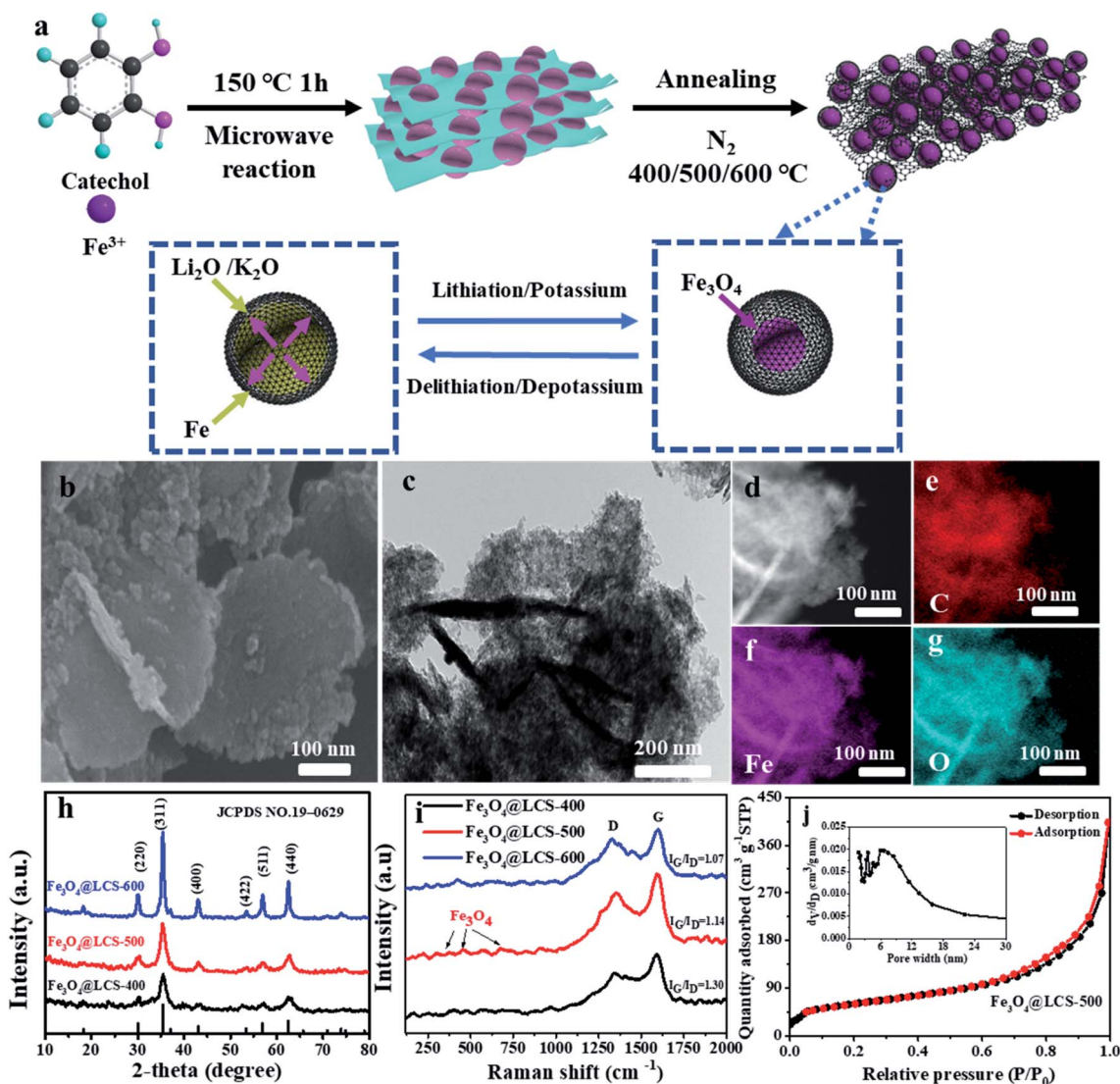


Fig. 1 (a) The synthesis schematic of the derived $\text{Fe}_3\text{O}_4@\text{LCS}$ (400/500/600) nanocomposites; (b) SEM images of $\text{Fe}_3\text{O}_4@\text{LCS}$ -500; (c) TEM images of $\text{Fe}_3\text{O}_4@\text{LCS}$ -500; (d–g) corresponding elemental maps of C, Fe, and O; (h) XRD patterns and (i) Raman spectra of $\text{Fe}_3\text{O}_4@\text{LCS}$ (400/500/600) nanocomposites; (j) nitrogen adsorption isotherm and the corresponding average pore size distribution of $\text{Fe}_3\text{O}_4@\text{LCS}$ -500.



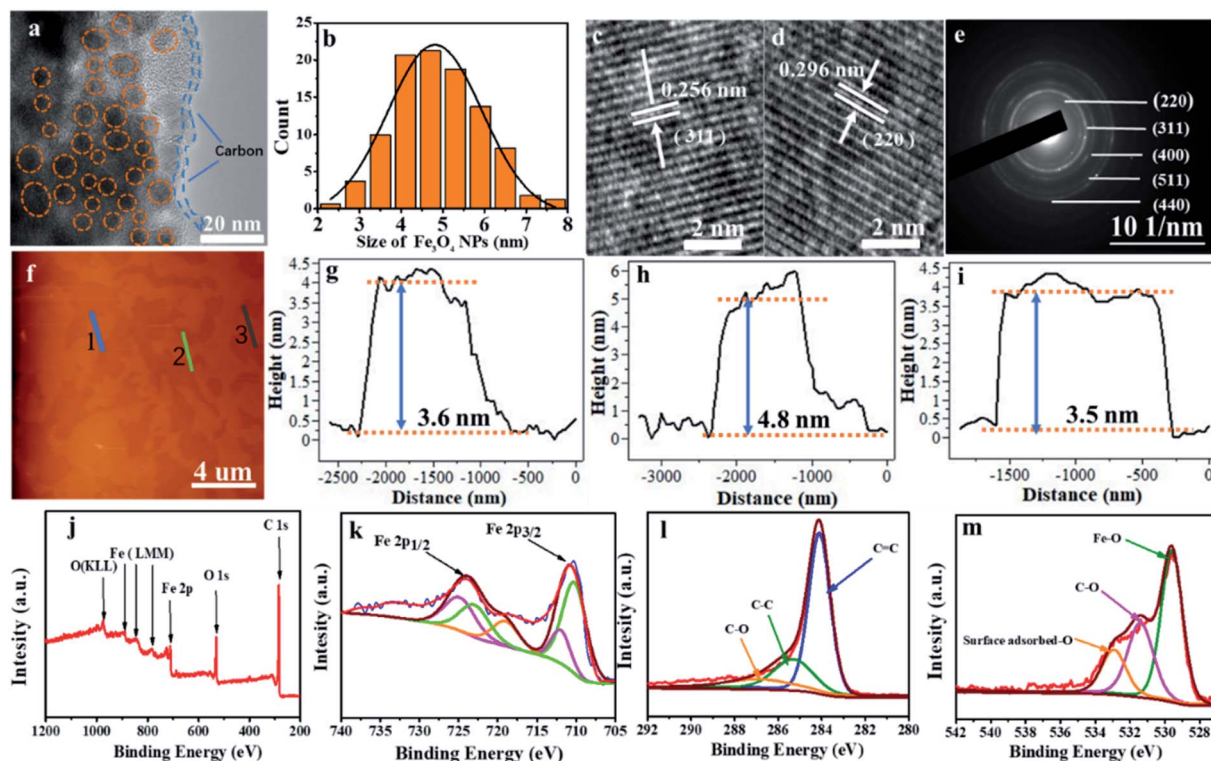


Fig. 2 HRTEM images (a), showing an apparent amorphous carbon layer and tiny Fe₃O₄ nanodots. The size distribution (b), and the crystal fringes (c and d), selected area electron diffraction (SAED) pattern (e); and the AFM (f) and the thickness profiles (g–i); and XPS spectra (j): Fe 2p (k); C 1s (l) and O 1s (m) of the Fe₃O₄@LCS-500 composites.

the same positions. Specifically, all characteristic diffraction peaks at 30.1°, 35.6°, 43.3°, 53.4°, 57.3°, 62.6° are well assigned to the (220), (311), (400), (422), (511) and (220) planes of Fe₃O₄ (JCPDS: 19-0629), which implies the full phase transformations of the precursor into Fe₃O₄-based materials. Meanwhile, the XRD patterns of Fe-coordination compound and Fe₃O₄ bulk are shown in Fig. S2a and b.† The Raman spectra of all composites are shown in Fig. 1i. Typical peaks at 383, 460, and 684 cm⁻¹ are assigned to E_g, T_{2g} and A_{1g} of Fe₃O₄.^{25,41} And peak positions of D band and G band are at ~1351 cm⁻¹ and 1601 cm⁻¹, in which D band stands for the disordered structure or defects in carbon and G band represents for the graphitic structure. Besides the intensity ratios of I_D/I_G of Fe₃O₄@LCS (400/500/600) samples are 1.30, 1.14, and 1.07, respectively. As a higher I_D/I_G value means a higher defect level, a lower value will result in a better electronic conductivity and higher mechanical stability. Therefore, a higher annealing temperature will give a lower I_D/I_G ratio, relating to a better electronic conductivity and good mechanical stability as well as less defects.⁴⁶ Furthermore, nitrogen absorption isotherms with pore size distributions of Fe₃O₄@LCS (400/500/600) nanocomposites were investigated by the BET (Brunauer, Emmett and Teller) method. As shown in Fig. 1j, the curve is coincided to the typical type IV H4 isothermal adsorption curve with an apparent hysteresis loop, displaying a high surface area of 213.84 m² g⁻¹ for Fe₃O₄@LCS-500 nanocomposites. Simultaneously, the relative average pore diameter was around 11.65 nm, leaving sufficient voids for

volume changes during cycles. In contrast, the surface areas of Fe₃O₄@LCS (400/600) in Fig. S3a and b† were 158.22 m² g⁻¹ and 250.50 m² g⁻¹, respectively. There is an apparent trend between specific surface area and annealing temperature, namely a higher surface area at higher annealing temperature.

The fine structure of Fe₃O₄@LCS-500 nanocomposites was further analyzed by high-resolution TEM (HRTEM) in Fig. 2a, displaying good crystallization of Fe₃O₄ nanodots. The evenly distributed Fe₃O₄ nanodots with an average particle size of around 5 nm, verified by the statistical plot in Fig. 2b, are homogeneously encapsulated by porous carbon matrix. The interplanar *d*-spacings (0.256 nm, and 0.296 nm) are corresponding to the (311) and (220) lattice planes of Fe₃O₄ in Fig. 2c and d. Other phase planes of (400), (511), and (440) are shown in the SAED pattern in Fig. 2e, exhibiting a polycrystalline nature. The average thickness of the Fe₃O₄@LCS-500 nanocomposites is around 4 nm based on atomic force microscopy (AFM) images in Fig. 2f–i, displaying an encapsulation structure because of the existence of single carbon shell with Fe₃O₄ nanodots. XPS technique was used to analyze the surficial valence state of Fe₃O₄@LCS-500 nanocomposites in Fig. 2j, illustrating the presence of Fe, C and O elements. The Fe 2p spectrum can be divided into five major peaks in Fig. 2k. The peaks of binding energy at 711.91 and 710.2 eV are to 2p_{3/2} of Fe³⁺ and Fe²⁺. And peaks at ~724.9 eV and 722.2 eV are referring to 2p_{1/2} of Fe³⁺ and Fe²⁺, respectively. Meanwhile, the peak at 719.02 eV is regarded as a satellite peak for the above four peaks,



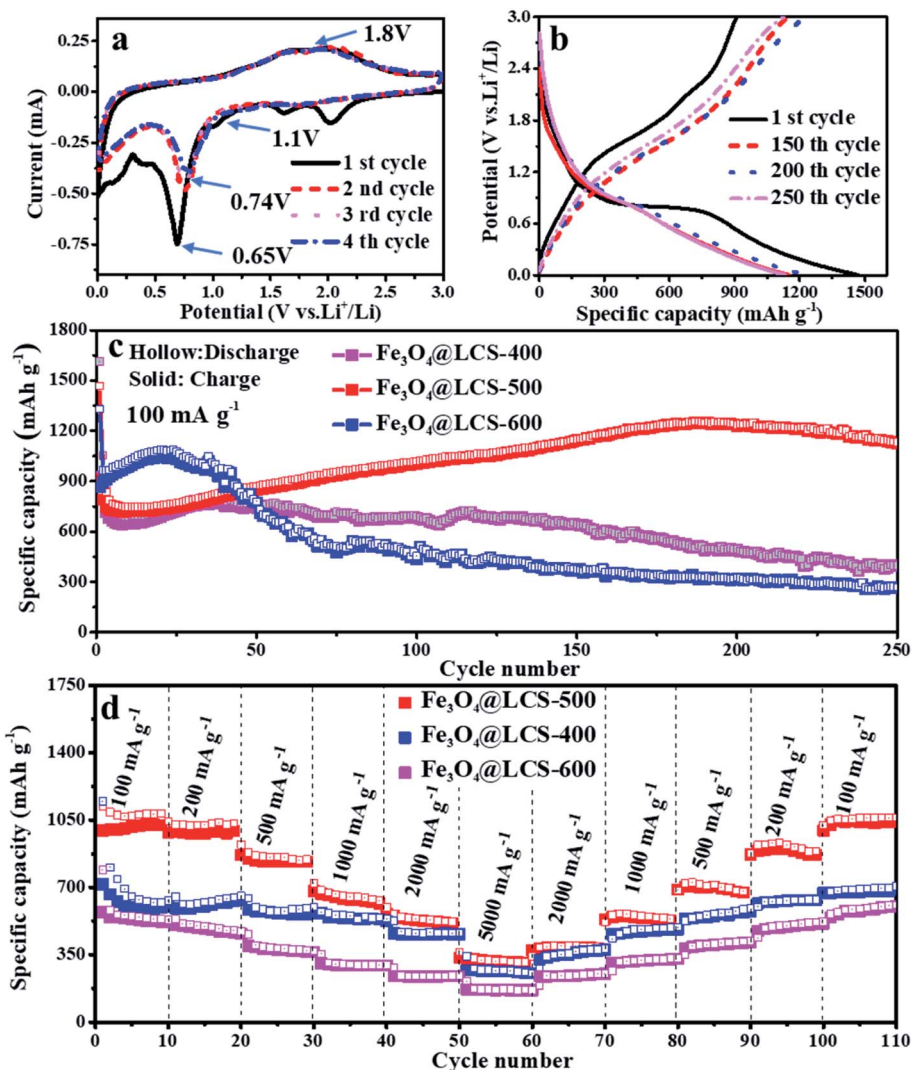
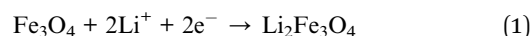


Fig. 3 Lithium storage performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ -(400/500/600) electrodes: (a) CV curves in the first four cycles at a scan rate of 0.1 mV s^{-1} ; (b) charge/discharge profiles at 100 mA g^{-1} ; (c) cycle performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ -(400/500/600) at the current density of 100 mA g^{-1} for 250 cycles; (d) the rate performance of $\text{Fe}_3\text{O}_4@\text{LCS}$ -(400/500/600) at various current densities.

demonstrating the formation of Fe_3O_4 .^{24,25,41,46} The C 1s spectrum is shown in Fig. 2l, divided into three sub-peaks at about 284.1, 285.6, and 286.9 eV. Those peaks are assigned to the covalent bonds of C=C, C-C, and C-O, respectively,^{3,47} indicating the existence of defects, in-plane structure, and surficial oxygen absorption. The O 1s spectrum in Fig. 2m shows peaks at about 529.6 and 531.6 eV, which are assigned to the oxygen states in Fe_3O_4 and C-O, and the peak at $\sim 533.7 \text{ eV}$ is attributed to the surface adsorbed oxygen.

The electrochemical performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ composites were investigated as anodes of LIBs. The CV curves in the first four cycles are presented in Fig. 3a. The small peaks above 1.1 V in the first reduction (cathodic) scan are associated with the formation of $\text{Li}_2\text{Fe}_3\text{O}_4$, which are disappeared in the following cycles because of the thermodynamics reason and the good stability of SEI layer. A special *in situ* XRD cell was assembled by directly coating the slurry on an ultra-thin Be foil to illustrate the generation of $\text{Li}_2\text{Fe}_3\text{O}_4$ phase. The diffraction

peaks of Fe_3O_4 could be observed obviously at the initial state, and then gradually disappeared in Fig. S10.† Diffraction peaks at 36.75° and 42.7° , corresponding to the (222) and (400) planes of $\text{Li}_2\text{Fe}_3\text{O}_4$, appeared when the voltage dropped to $\sim 1.1 \text{ V}$, and then disappeared in following discharge process, confirming $\text{Li}_2\text{Fe}_3\text{O}_4$ was truly generated in the reduction process. Those following current may come from the lithium storage behavior of porous carbon layers. The anodic processes occurred at about 1.8 V due to the reformation of Fe_3O_4 and the delithiation of carbon layers. Apparently, in the following cathodic sweeps, the reduction peak is slightly shifted to $\sim 0.74 \text{ V}$, which is mainly ascribed to the formation of the SEI film and slightly increased inner resistance. Obviously, the almost overlapped curves in the subsequent scanning cycles imply highly reversible capability. The reversible reactions are given as follows:



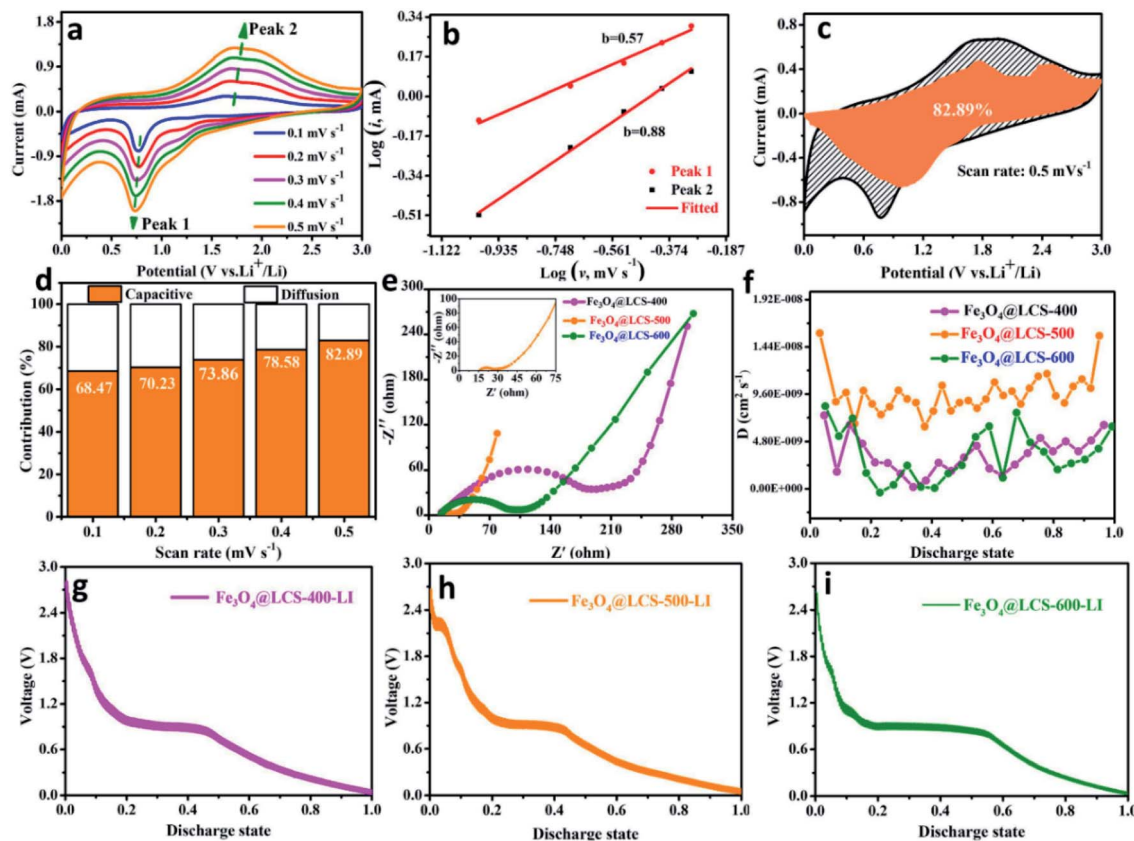
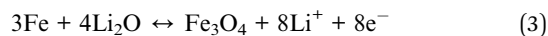
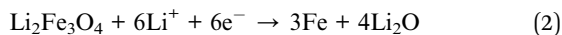


Fig. 4 Capacity contribution and lithium diffusion coefficients: (a) CV curves of the $\text{Fe}_3\text{O}_4@\text{LCS-500}$ electrode at different scan rates from 0.1 to 0.5 mV s^{-1} ; (b) b value with linear relationship between $\log(i)$ and $\log(v)$; (c) capacitive contributions of $\text{Fe}_3\text{O}_4@\text{LCS-500}$ electrode at the scan rate of 0.5 mV s^{-1} ; (d) contribution ratios of capacitive capacities at various scan rates; (e) the Nyquist plots of the $\text{Fe}_3\text{O}_4@\text{LCS-400/500/600}$ electrodes; (f) the calculated Li chemical diffusion coefficients; (g–i) GITT curves of $\text{Fe}_3\text{O}_4@\text{LCS-400/500/600}$ electrodes.



$\text{Fe}_3\text{O}_4@\text{LCS-500}$ nanocomposites delivered a high reversible capacity of 908 mA h g^{-1} with a coulombic efficiency of 65.76% at 100 mA g^{-1} in Fig. 3b. The irreversible capacity loss can be ascribed to the formation of the SEI layer and the decomposition of the electrolyte.⁴ Fig. 3c shows long-term cycling performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ (400/500/600) electrodes at 100 mA g^{-1} for 250 cycles. The $\text{Fe}_3\text{O}_4@\text{LCS-400/600}$ electrodes delivered a reversible capacity of 929 mA h g^{-1} and 868 mA h g^{-1} , respectively. Those electrodes have shown a similar capacity trend of a slight decrease in the first several cycles and gradual increase afterward, which is normally seen in many reports about the activation process of transition metal oxides during cycles.^{42,48} However, the Fe_3O_4 bulk electrode only delivered a reversible capacity of $729.2 \text{ mA h g}^{-1}$ and no such phenomenon as shown in Fig. S8.† $\text{Fe}_3\text{O}_4@\text{LCS-500}$ electrode maintained a reversible capacity of $1126.2 \text{ mA h g}^{-1}$ after 250 cycles. It exhibited anomalously high capacities beyond their theoretical value, which was probably related to the strong surface capacitance around metal nanoparticles and reversible storage

of spin-polarized electrons in the already-reduced iron nanodots.⁴ By contrast, other electrodes, including $\text{Fe}_3\text{O}_4@\text{LCS-400}$, $\text{Fe}_3\text{O}_4@\text{LCS-600}$, and bulk Fe_3O_4 , only displayed reversible capacities of 395.5 , $265.9 \text{ mA h g}^{-1}$, and $257.5 \text{ mA h g}^{-1}$ after 250 cycles, respectively. That is mainly associated to features of $\text{Fe}_3\text{O}_4@\text{LCS-500}$ nanocomposites on (i) the higher specific surface area and fewer defects in carbon layers, offering more active sites for reversible cation's accommodations, (ii) its lower inner resistance, facilitating to charge transfer and fast ionic diffusion coefficient. $\text{Fe}_3\text{O}_4@\text{LCS-500}$ electrode exhibited better rate performance than the others in Fig. 3d, delivering rechargeable capacities of 996, 988, 848, 666, 438 and 210 mA h g^{-1} at stepwise current densities of 100, 200, 500, 1000, 2000 and 5000 mA g^{-1} , respectively. When the current density was returned to 100 mA g^{-1} , the reversible capacity was recovered to around 1000 mA h g^{-1} , implying a fast lithium diffusion coefficient and kinetics. On the contrast, the capacities of $\text{Fe}_3\text{O}_4@\text{LCS-400/600}$ and bulk Fe_3O_4 electrodes (Fig. S8b†) were stable at low current densities, but decreased at high rates, displaying poor rate performances. Moreover, electrochemical performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ nanocomposites as anodes for KIBs were also measured and presented in Fig. S4, S5 and S9,† demonstrating the application possibility in KIBs.



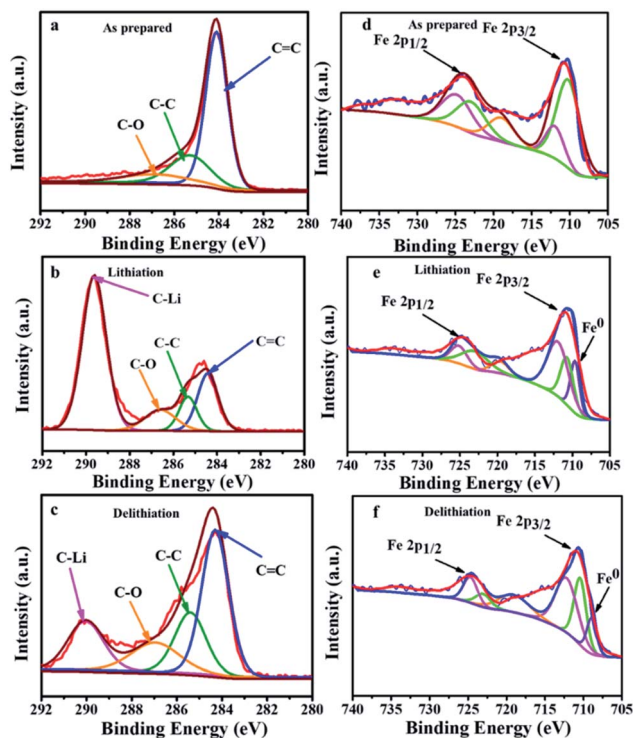


Fig. 5 *Ex situ* XPS spectra of carbon and iron in the different states of Fe_3O_4 @LCS-500 electrodes in LIBs. (a) Carbon spectrum; (b) carbon spectrum in the fully lithiated state; (c) carbon spectrum in the fully delithiated state; (d) iron spectrum; (e) iron spectrum in the fully lithiated state; (f) iron spectrum in the fully delithiated state is close to 1.0, it is inclined to a capacitive controlled process.^{25,51}

To measure the specific values of the lithium diffusion coefficient and quantify the capacitive characterization, we utilized the CV method cycled at different scan rates to measure

lithium cation's behaviors and diffusion coefficients. Fig. 4a depicts CV curves at different scan rates (from 0.1 to 0.5 mV s^{-1}). During scans, the intensities of redox peaks are increased with scan rates, which might be assigned to the different storage behaviors (the pseudo-capacitive contribution or cation-diffusion contribution). That can be estimated by the following equation:^{39,49,50}

$$i = k_1v + k_2v^{1/2} = av^b \quad (4)$$

where i is the peak current and v is the scan rate, a and b values are constants. The capacitive-contribution is related to k_1v , and diffusion-type is associated with $k_2v^{1/2}$.

When the value of the slope (b) approaches 0.5, the storage behavior is dominated by diffusion-type reactions; when it is close to 1.0, it is inclined to a capacitive controlled process.^{25,51}

Fig. 4b exhibits the relationship between $\log(i)$ and $\log(v)$, where peak 1 (cathodic peak) and peak 2 (anodic peak) of the slope b were calculated to be 0.57 and 0.88, respectively. That means it was dominated by capacitive-controlled behavior. Fig. 4c presents the portion of the capacitive contribution (82.92%) against the whole scan area at 0.5 mV s^{-1} . Contribution ratios of capacitive-controlled behaviors in Fig. 4d were calculated to be 68.47%, 70.23%, 73.86%, 78.56% and 82.92% at different scan rates. It is noted that the capacitive contribution is increased with the increase of scan rates. Furthermore, the electrochemical impedance spectra of Fe_3O_4 @LCS-(400/500/600) electrodes are shown in Fig. 4e, S7 and Table S2,† in which the depressed semi-circle in the high and middle-frequency region is related to charge-transfer, and the inclined line in the low-frequency region is associated to ionic diffusion. Apparently, the semi-circle's radius of Fe_3O_4 @LCS-500 is much smaller than other electrodes, implying a lowest electrochemical resistance and best electronic conductivity. The

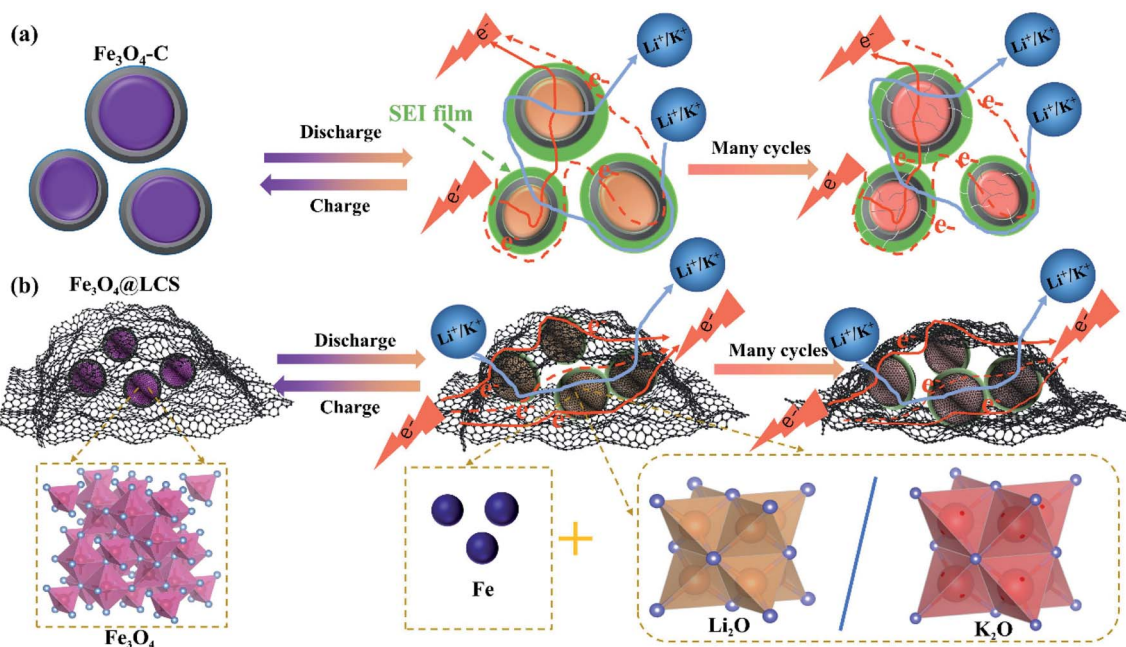


Fig. 6 Schematic illustration of the morphology evolutions of Fe_3O_4 -C (a) and Fe_3O_4 @LCS (b) materials.



thermodynamic or kinetic behaviors were further analyzed by the galvanostatic intermittent titration technique (GITT) in Fig. 4f–i. The diffusion coefficient of Li^+ ion is acquired by the equation as follows:^{35,52}

$$D_{\text{Li}} = \frac{4}{\pi} \left(\frac{mV_m}{MA} \right)^2 \left(\frac{\Delta ES}{\Delta E\tau} \right)^2 \quad (5)$$

In which m and M mass and molecular weight of the active substance; V_m : active substance's molar volume; τ : constant titration time; A : electrode surface area; ΔES : steady-state voltage variation under current disturbance; $\Delta E\tau$: voltage variation under constant current titration.⁵³ The diffusion coefficient values of lithium (D_{Li}) of $\text{Fe}_3\text{O}_4@\text{LCS}$ -(400/500/600) electrodes are shown in Fig. 4f. During the whole discharging process, we can obviously observe that the diffusion coefficients of the $\text{Fe}_3\text{O}_4@\text{LCS}$ -500 was the highest ($\sim 1.04 \times 10^{-8} \text{ cm}^2 \text{ s}^{-1}$), which is much higher than others ($\text{Fe}_3\text{O}_4@\text{LCS}$ -400: $3.49 \times 10^{-9} \text{ cm}^2 \text{ s}^{-1}$; $\text{Fe}_3\text{O}_4@\text{LCS}$ -600: $6.29 \times 10^{-9} \text{ cm}^2 \text{ s}^{-1}$). The information of capacity contribution and diffusion coefficients of KIBs was shown in Fig. S5,† giving similar electrochemical behaviours.

The *ex situ* XPS patterns of $\text{Fe}_3\text{O}_4@\text{LCS}$ -500 sample, acquired at different reaction states, were examined to verify the redox reaction mechanism in LIBs. The main peak of carbon in Fig. 5a can be divided into three peaks at ~ 284.1 , 285.6 , and 286.9 eV, corresponding to C=C, C–C, and C–O, respectively. After the full lithiation, a new peak of 289.62 eV appears in Fig. 5b, which is assigned to the C–Li bond, demonstrating porous carbon nanosheets have contributed some capacity. When it is fully delithiated, the peak intensity is obviously decreased in Fig. 5c, indicating the partially reversible behavior of porous carbon nanosheets.^{11,47} The XPS spectrum of iron can be divided into five major peaks in Fig. 5d. The configuration of $\text{Fe } 2p_{3/2}$ is composed by peaks at ~ 711.91 and 710.2 eV, relating to a mixed valence state of +3 and +2. The peak at about 724.9 eV is referring to $\text{Fe } 2p_{1/2}$ (a valence state of +3). Meanwhile, the peaks at 722.2 eV is referring to $\text{Fe } 2p_{1/2}$ (a valence state of +2). After the full lithiation process, a new peak at 709.02 eV appears in Fig. 5e, which is ascribed to the formation of Fe^0 . In the fully delithiated process, the peak intensity is obviously weakened in Fig. 5f, indicating the reversible conversion reactions between Fe_3O_4 and Li .⁵⁴ The spectra of carbon and iron in different valence states of $\text{Fe}_3\text{O}_4@\text{LCS}$ -500 electrodes for KIBs were measured by *ex situ* XPS as shown in Fig. S6.†

The superior electrochemical performances of $\text{Fe}_3\text{O}_4@\text{LCS}$ electrodes for LIBs/KIBs are attributed to their unique features of the thin carbon layer, pre-reserved space, and ultra-small particle size as well as flexible capability. Carbon coated Fe_3O_4 ($\text{Fe}_3\text{O}_4\text{-C}$) bulk in Fig. 6a, perplexed by Ostwald ripening, are normally experiencing the growth of SEI film, and cracks on carbon shells after cycles. Even though electron and ion can easily transfer among particles.

Without the pre-reserved space, the rigid carbon shell is tended to crack after many cycles due to large volume changes and side reactions. Unlike $\text{Fe}_3\text{O}_4\text{-C}$ bulk, $\text{Fe}_3\text{O}_4@\text{LCS}$ materials

are composed of ultra-small Fe_3O_4 nanodots and layered carbon nanosheets with the flexible feature and pre-reserved spaces in Fig. 6b. Layered carbon sheets are functioned as an elastic buffer with pre-reserved spaces to prevent pulverization or aggregation of encapsulated ultra-small Fe_3O_4 nanodots during cycles, maintaining stable and unobstructed channels for the fast electronic and ionic conductivity.^{54–56} The thin and layered carbon nanosheets not only enable full utilization of active ultra-small Fe_3O_4 nanodots and abundant accessibility of electrolyte, but also accommodate the mechanical stress from the volume variations, as shown in Fig. S11a–i† in both LIBs and KIBs.^{57,58} Therefore, the unique features of the ultra-small particle size, less defects and stable layered structures in $\text{Fe}_3\text{O}_4@\text{LCS}$ composites are jointly contributed to the high capacity, good rate capability, and long cycle life as well as high diffusion coefficients.

Conclusion

In summary, $\text{Fe}_3\text{O}_4@\text{LCS}$ nanocomposites were derived from metal coordination compound with stacked layered structure, avoiding the Ostwald ripening phenomenon. This kind of materials has unique features of ultra-small particle size, high surface area, pre-reserved spaces, flexibility. When applied as anodes for LIBs/KIBs, $\text{Fe}_3\text{O}_4@\text{LCS}$ -500 nanocomposites with less defects and short ionic diffusion length, and high electronic conductivity have delivered exceptional electrochemical performances, which are largely ascribed to the unique properties of high diffusion coefficients, fast reaction kinetics, pre-reserved volume variation spaces, flexible carbon sheets and capacitive-controlled electrochemical behaviours. Such material design strategy and characterization methods can be extended to other anodes for rechargeable Li/Na/K ion batteries.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgements

The authors gratefully acknowledge the National Natural Science Foundation of China (21975154, 52073170), Shanghai Municipal Education Commission (Innovation Program (2019-01-07-00-09-E00021)) and Innovative Research Team of High-level Local Universities in Shanghai. Research is also supported by The Program for Professor of Special Appointment (Eastern Scholar) at Shanghai Institutions of Higher Learning and Shanghai Key Laboratory of Materials Protection and Advanced Materials in Electric Power. The authors thank Laboratory for Microstructures, Instrumental Analysis, and Research Center of Shanghai University for offering access to material characterizations.



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