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Probing the Electrochemical Behavior of {111} and {110} Faceted Hollow Cu₂O Microspheres for Lithium Storage

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Transition metal oxides with exposed highly active facets have become a rising issue as anode materials for lithium ion batteries, because more dangling atoms exposed at the active surface facilitate the reaction between the transition metal oxides and lithium. In this work, we probed the electrochemical behavior of hollow Cu₂O microspheres with {111} and {110} active facets on the polyhedron surface as anodes for lithium storage. Compared to commercial Cu₂O nanoparticles, hollow Cu₂O microspheres with {111} and {110} active facets show a rising specific capacity at 30 cycles which then decreases after 110 cycles during the cycling process. Via advanced electron microscopy characterization, we reveal that this phenomenon can be attributed to the highly active {111} and {110} facets with dangling "Cu" atoms facilitating the conversion reaction of Cu₂O and Li, where part of the Cu₂O is oxidized to CuO during the charging process. However, with the reaction proceeding, more and more formed Cu nanoparticles cannot be converted to Cu₂O or CuO. This leads to a decrease of the specific capacity. We believe that our study here sheds some light on the progress of the electrochemical behavior of transition metal oxides with respect to their increased specific capacity and the subsequent decrease via a conversion reaction mechanism. These results will be helpful to optimize the design of transition metal oxide micro/nanostructures for high performance lithium storage.

1. Introduction

Lithium ion batteries (LIBs) have been widely used in many portable electronic devices, such as mobile phones, laptops and cameras owing to the high energy and power density, combined with a long cycle lifetime. Due to extremely high reversible capacity and energy density, low cost, environment-friendly, transition metal oxides have been considered as ideal anodes to replace graphite for lithium ion batteries.¹⁻⁷

Generally, the lithiation and de-lithiation process for many transition metal oxides is a reversible conversion reaction.^{2, 3, 7} Therefore, the arrangement of the surface atoms of these

transition metal oxides structures are crucial to such conversion reaction and therefore closely related to their electrochemical performance. From a chemical reaction point of view, the surface structure of electrode materials with exposed high surface energy facets should demonstrate excellent electrochemical performances for LIBs, because more dangling atoms exposed at the surface will facilitate the reaction between transition metal oxides and lithium. For instance, Zhao and co-workers reported that special Fe₂O₃ octahedra with highly active {111} facets exhibit an increased capacity and excellent high rate cycling performance because of the active {111} facets ensuring electrons and Li⁺ effectively transferring between the electrolyte and the interface of the Fe₂O₃ octahedra.⁸ Sun and co-workers studied nanosized Co₃O₄ octahedra with enclosed {111} facets for lithium storage and found the capacity slightly increasing during the charge/discharge process, owing to the nanosize and the {111} facets.⁹ Indeed, this specific capacity increase is widely found for transition metal oxides as anode materials for lithium storage.^{10, 11} And the authors have usually attributed this to the decomposition of the solid electrolyte interface (SEI) and/or the reversible growth of polymeric gel-like layers resulting from kinetically activated electrolyte degradation.^{9, 12} However, from the conversion reaction itself, this explanation is not convincing enough. Recently, our group has synthesized Mn₃O₄ nano-octahedra, which show not only anomalous magnetic properties and superior photo decomposition

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[‡] Electronic Supplementary Information (ESI) available: SEM and HRTEM images, BET surface areas of commercial Cu₂O nanoparticles. See DOI: 10.1039/x0xx00000x

activity,¹³ but also excellent cycling performance and rate capability for LIBs. We proposed that the exposed {011} facets with more dangling Mn atoms facilitate more and more Mn²⁺ to be converted to Mn³⁺ during the discharge-charge process.¹⁴ This has been further evidenced on the unique walnut-shaped porous MnO₂/C nanospheres, which demonstrate a gradual increase of the specific capacity, owing to a deeper conversion reaction for further oxidation from Mn²⁺ to Mn³⁺ during the anodic process.¹⁵ Therefore, we argue that the capacity increase is mainly from the conversion reaction itself, namely, the enhanced reaction kinetics of Mn²⁺ to Mn³⁺ instead of coming from the SEI layer and/or the reversible growth of polymeric gel-like layers.

However, research on the role of the active facets in the electrochemical performance is still seldom reported. To verify our assumption, other transition metal oxides with active facets should be studied. For instance, Xue and co-workers found that the cubic Cu₂O with more {100} facets shows the highest capacity and the additional capacity of Cu₂O is due to the active facets made it easily oxidized to CuO at the high voltage with the presence of an organic electrolyte.¹⁶ However, they only provided CV curves to verify their point. More work should be carried out to probe the electrochemical behaviour of Cu₂O with such special active facets for lithium storage.

In this work, we studied the hollow microspheres of Cu₂O (HMs-Cu₂O) with exposed {111} and {110} active facets as anode materials for LIBs. Such special Cu₂O demonstrates an increased capacity during the Li⁺ insertion/extraction process. Compared to commercial Cu₂O nanoparticles without special active facets, HMs-Cu₂O delivers not only an enhanced capacity and stability, but also enhanced reaction kinetics with an advanced capacity increase during the Li⁺ insertion/extraction process. We successfully evidenced that {111} and {110} active faceted Cu₂O can be easily oxidized to CuO in the charging process using transmission electron microscope (TEM), high angle annular dark field-scanning transmission electron microscopy (HAADF-STEM) and electron energy loss spectroscopy (EELS). However, part of Cu nanoparticles cannot be further converted to Cu₂O and CuO during the cycling process. This leads to a capacity decrease in a prolonged cycling process.

2. Experimental Section

2.1 Synthesis of hollow Cu₂O microspheres

All chemicals are in analytical grade (Aladdin Industrial Corporation, China) and used as received. The reaction is the same as in our previous work.¹⁷ For a typical synthesis, 2 mmol of Cu(NO₃)₂·3H₂O (Sinopharm Chemical Reagent Co., Ltd.) is first dissolved in deionized water at a certain volume. Next, 40 mL ethylene glycol (Aladdin Industrial Corporation) is added into the above blue solution. After magnetic stirring for 2 h, the transparent pale blue solution is transferred into 50 mL Teflon-lined stainless steel autoclaves, sealed and transferred to the oven. Then the oven temperature is raised to 180 °C and maintained at that temperature for 1 h. The solid product is

filtrated and washed with ethanol, to remove the unreacted ethylene glycol in the final product, and dried at 40 °C in air.

For comparison, commercial cuprous oxide nanoparticles (Cu₂O-NPs) without specific morphology and active facets were purchased from Aladdin Industrial Corporation (China). From the SEM image shown in Fig. S1, the particles have a size of ~50 nm and are tightly aggregated together. HRTEM images shown in Fig. S2 reveal there is no amorphous or other phases in the Cu₂O-NPs.

2.2 Materials characterization

Structural and chemical analyses were performed by powder X-ray diffraction (XRD) employing a Bruker D8 Advanced diffractometer with Cu Kα radiation. The morphology of the samples was studied with a Hitachi S-4800 scanning electron microscope (SEM) equipped with a field emission gun (FEG) at an accelerating voltage of 5 kV. Transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM) were performed on a JEM-2100F transmission electron microscope at 200 kV. High angle annular dark field-scanning transmission electron microscopy (HAADF-STEM) and electron energy loss spectroscopy (EELS) were performed on a FEI Titan 80-300 "cubed" microscope fitted with an aberration-corrector for the imaging mode and the probe forming mode, a monochromator, a GIF Quantum energy filter for spectroscopy, operated at 300 kV. The specific surface area is measured using a Tristar II 3020 (Micromeritics, USA) and calculated by the Brunauer-Emmett-Teller (BET) method. The results are shown in Table S1, which shows the similar low BET surface area of HMs-Cu₂O and Cu₂O-NPs.

2.3 Electrochemical measurements

The anodes are fabricated by mixing the active material, conductive carbon (super P), and polyvinylidene fluoride (PVDF) at a weight ratio of 70:20:10, respectively. N-methyl pyrrolidone (NMP) is used as a solvent. The resulting slurries are cast onto a copper current collector, and then dried at 120 °C under vacuum for 12 h. The electrode foils are cut into disks with 12 mm in diameter. CR2025 coin-type cells are assembled in an argon-filled glove box by stacking a microporous polypropylene separator containing a liquid electrolyte of LiPF₆ (1M) in ethylene carbonate (EC)/dimethyl carbonate (DMC) (1:1, v/v) between the Cu₂O anode and the lithium metal foil cathode. The loading of active material in the composite electrodes is ~1.0 mg/cm². The cells are charged and discharged galvanostatically using a battery test system (LAND CT2001A) in a potential range between 0.02 and 3.0 V at various current densities (1 C = 400 mA g⁻¹).

3. Results and Discussion

Fig. 1 shows the XRD pattern of the as-synthesized product. All peak positions and relative intensities match very well with cubic Cu₂O (JCPDS No. 065-3288) and no peak of impurities are found. The sharp narrow peaks also suggest that the products are well crystallized.

Fig. 1. The XRD result of the as-synthesized HMs-Cu₂O.

Fig. 2a presents the typical SEM image of the as-synthesized Cu₂O with a uniform diameter of ~700 nm. Fig. 2b shows one broken microsphere, indicating the hollow structure of the as-synthesized Cu₂O. Furthermore, the surface of the Cu₂O microspheres clearly reveals that the microspheres are composed of randomly cross-linked polyhedra. Close observation shows that the active facets of {111} and {110} are exposed at the surface. TEM confirms the hollow structure of HMs-Cu₂O (Fig. 2c). The lattice fringes observed by HRTEM (Fig. 2d) correspond to 0.245 and 0.302 nm, in agreement with the {111} and {110} planes of Cu₂O, respectively. This is consistent with the SEM observations, indicating the exposed highly active {110} and {111} facets at the surface of HMs-Cu₂O. It is known that in Cu₂O, each oxygen atom is at the center of a tetrahedron of copper atoms, which linearly coordinates with two oxide ions.^{17, 18} Fig. 2e and Fig. 2f demonstrate the surface atomic species and state of {110} and {111} facets respectively, which are composed by either copper cations or oxygen anions.^{19, 20} Our previous study on the absorption of different charged organic chemicals suggests that the {110} and {111} facets are mainly terminated by dangling Cu⁺ cations for HMs-Cu₂O.¹⁷ Therefore, the {110} and {111} faceted Cu₂O with

dangling Cu⁺ cations is beneficial for the electrochemical reaction of Cu₂O: Cu₂O + Li → Cu + Li₂O. Further, the hollow structure is also helpful for storing the electrolyte and buffering the volume change during the Li⁺ insertion/extraction process.^{21, 22}

Fig. 2. Electron microscopy images of HMs-Cu₂O. (a) SEM image at low magnification, (b) one hollow microsphere, (c) TEM image at low magnification and (d) HRTEM image, (e and f) the surface atomic configurations in the {111} and {110} planes of cubic Cu₂O.

Fig. 3. (a) CV response of the HMs-Cu₂O electrode at 0.1 mV s⁻¹ for the first four cycles. (b) discharge-charge voltage profiles for the 1st, 2nd, 10th, 50th, 100th and 200th cycles of HMs-Cu₂O at 0.2C. (c) cycling performance of the HMs-Cu₂O with the coulombic efficiency and commercial Cu₂O-NPs electrodes at 0.2C.

The electrochemical properties of the hollow Cu₂O microspheres are then evaluated. Fig. 3a shows the first four cyclic voltammetry curves of HMs-Cu₂O at a scan rate of 0.1 mV s⁻¹ from 0.02 to 3.0V. At the first scan, a strong cathodic peak at ~0.9-1.2V is displayed, corresponding to the chemical reduction reaction of Cu₂O → Cu.²² This peak shifts to ~1.6 V in the following scans, indicating a decreased polarization of the HMs-Cu₂O electrode and suggesting an easy electrochemical

reverse reaction during the Li⁺ insertion-extraction process. The weak peak located at ~0.6V and that further shifts to ~0.8V is attributed to the deep lithiation^{23, 24} and the formation of an SEI layer between Cu₂O and the electrolyte.¹⁴ At the anodic scan, there are two apparent peaks located at ~1.4 and ~2.4V, which are ascribed to the partial decomposition of SEI and the recombination of Cu and Li₂O, respectively.¹ These results are in agreement with the discharge-charge profiles at

0.2C (Fig. 3b). In the cycling discharge-charge profiles, we note that with the cycle number increasing, a new charge plateau appears at ~2.8V. This plateau may correspond to the further redox reaction between CuO and Cu₂O according to previous work²⁵ because the typical phase transformation between Cu₂O and Li can be expressed as:

(1st sloping part of the discharge curve: 1.4 to 1.0 V);

(for charge voltage > 2.2V).

Fig. 3c displays the cycling performance of HMs-Cu₂O and commercial Cu₂O-NPs with the cut-off voltages of 0.02-3.0V (versus Li/Li⁺) at a current density of 0.2C (1C = 400 mAh g⁻¹). The initial discharge capacity of HMs-Cu₂O is 834.7 mAh g⁻¹ at 0.2C, which is much higher than that of commercial Cu₂O-NPs (364 mAh g⁻¹). The coulombic efficiency of HMs-Cu₂O is also shown in Fig. 3c. It clearly shows the increased coulombic efficiency from the first cycle of 50% to almost 100% after the second cycle. It is interesting to note that the specific capacity slightly increases after 30 cycles for HMs-Cu₂O. This is quite different from the previous Cu₂O polyhedra with active facets, where no capacity increase is shown.¹⁶ After further cycling, the specific capacity gradually decreases from 110 cycles on. Comparing to HMs-Cu₂O, the capacity of commercial Cu₂O-NPs only exhibits very little increase after 80 cycles. Generally, smaller nanoparticles are beneficial for an enhanced capacity. However, in this experiment, the smaller nanoparticles from commercial Cu₂O-NPs demonstrate a lower capacity than that of HMs-Cu₂O in microsize. Most probably this is because of the exposed {111} and {110} facets, which can enhance the conversion reaction kinetics. This is similar to previous works on metal oxides with active facets.²⁶ This enhanced reaction kinetics in HMs-Cu₂O makes its capacity higher than that of commercial Cu₂O-NPs. And the capacity increase largely advances that of commercial Cu₂O-NPs. According to the previous work on Cu₂O polyhedral with active facets for LIBs¹⁶ and formula (2), we suppose that part of Cu₂O converts to CuO because of the high theoretical capacity of CuO (674 mA h g⁻¹). This further oxidation explains the capacity increase during the cycling process.

To verify the above speculation, we performed SEM and TEM analysis to follow the structural evolution of HMs-Cu₂O during the cycling process. We first selected HMs-Cu₂O after 50 cycles at 0.2C because of the improvement of capacity as shown in Fig. 3c. Fig. 4a shows the SEM image of HMs-Cu₂O after 50 cycles from the charged state at 0.2C. It clearly displays the well maintained shape and the exposed facets. Fig. 4b provides a close observation of the reacted HMs-Cu₂O, showing the SEI film on the surface of the Cu₂O microspheres. Fig. 4b inset displays the presence of many small nanoparticles at the surface of the polyhedron after the SEI layer is peeled off. Fig. 4c-f show TEM images of the reacted HMs-Cu₂O and the SEI layer, respectively. Fig. 4c clearly shows the presence of some nanoparticles at the surface of the polyhedron. The HRTEM image (Fig. 4d) of the area indicated in Fig. 4c and the corresponding FFT (Fig. 4d inset) reveal the existence of Cu nanoparticles, although the interplanar spacing of Cu (111)

($d_{\text{Cu}(111)} = 0.209 \text{ nm}$) is closed to that of CuO (200) ($d_{\text{CuO}(200)} = 0.212 \text{ nm}$) and Cu₂O (200) ($d_{\text{Cu}_2\text{O}(200)} = 0.213 \text{ nm}$), however the interplanar spacing of Cu (200) ($d_{\text{Cu}(200)} = 0.532 \text{ nm}$) can be easily detected. Meanwhile, the HRTEM image (Fig. 4f) of the SEI layer and the corresponding FFT (Fig. 4f inset) also demonstrate that some Cu nanoparticles are embedded in the SEI layer, suggesting that the generated Cu nanoparticles infiltrate into the SEI layer from the Cu₂O microsphere and cannot be fully converted to Cu₂O during the electric cycles. However, the composition and distribution of CuO and Cu₂O is still unclear from HRTEM or FFT, because they have very closed lattice spacings (CuO: cubic, $a = 4.25 \text{ \AA}$; Cu₂O: cubic, $a = 4.27 \text{ \AA}$).

The Cu₂O and CuO can be differentiated through high resolution EELS analysis, due to the differences of Cu L_{2,3} edge between Cu¹⁺ and Cu²⁺.²⁷ Furthermore, by using an annular detector to collect the electrons scattered by the sample over large angles, image acquisition and the collection of energy loss scattered electrons using a GIF can be done simultaneously. This makes HAADF-STEM-EELS ideally suited for the study of Cu₂O/CuO/Cu hybrid as imaging can be directly combined with local chemical information. We performed HAADF-STEM-EELS spectrum analysis on reacted HMs-Cu₂O

Fig. 4. Electron microscopy images of HMs-Cu₂O after 50 cycles for the charge state at 0.2C. (a) SEM image and a Cu₂O microsphere with SEI layer (inset), (b) a Cu₂O microsphere with partially peeled off SEI layer and corresponding exposed Cu₂O microsphere core (inset). (c) TEM image of an exposed Cu₂O microsphere core and corresponding low magnification TEM image (inset). (d) HRTEM image of the area indicated in (c) and corresponding FFT pattern, (e) TEM image of the surface of SEI layer from a Cu₂O microsphere (inset), and (f) HRTEM image of the area indicated in (e) and corresponding FFT pattern.

after 50 cycles from the charge state at 0.2C. Fig. 5a shows a HAADF-STEM image of a typical HMs-Cu₂O microsphere after 50 cycles, with its corresponding EELS spectrum of the Cu L_{2,3} edge shown in Fig. 5c (blue colour). An EELS spectrum of the same L_{2,3} edge from pure CuO (black), pure Cu₂O (green) and pure Cu (red) are also shown in Fig. 5c as reference. Note that all spectra, shown in Fig. 5c, have been processed after background removal. The energy shift of the first major

maxima peak in energy loss near edge structure (ELNES) of CuO relative to Cu₂O and Cu is ~2.3 eV (Fig. 5c), consistent with a previous report.²⁸ The EELS spectrum of Cu-L₃ in HMs-Cu₂O after 50 cycles (zone1) presents two split peaks indicated by A and B, which also reveals the co-existence of CuO, Cu₂O and Cu. Although the maximum of the first ELNES peaks of Cu₂O and CuO occur at the same position, the existence of Cu has been verified by the HRTEM and FFT in Fig. 4. In order to understand

Fig. 5. (a-b) HAADF-STEM images of HMs-Cu₂O after 50 and 80 cycles for the charge state at 0.2C, respectively. (c) Copper L_{2,3} core-loss EELS spectra acquired from the indicated areas shown in (a) and (b) compared to referencing Cu compounds: CuO (black curve), Cu₂O (green curve) and Cu (red curve) and from different areas indicated in (a) zone1 (50 cycles sample) and in (b) zone2 (80 cycles sample), (d-h) and (i-m) EELS maps of HMs-Cu₂O after 50 and 80 cycles at 0.2C, respectively. (e and j) Cu₂O (green), (f and k) CuO (red), (g and l) Cu (blue), and (h and m) Cu₂O/CuO/Cu overlaid color mapping.

the distribution of these components (Cu/Cu₂O/CuO), the Cu-L₃ edges in the HAADF-STEM-EELS data are further fitted using the EELSModel program²⁹ and EELS mapping can be obtained for the different components (Fig. 5d-h). The distribution of Cu, Cu₂O and CuO is clearly presented, showing more aggregation of CuO than Cu and Cu₂O on the surface of the microsphere. Therefore, combining the TEM and HAADF-STEM-EELS results, we confirm that Cu₂O, CuO and Cu co-exist in the HMs-Cu₂O after 50 cycles. In addition, Cu nanoparticles appear in the SEI layer and more CuO nanoparticles aggregate on the surface of the polyhedron. Although we only characterized the charge state after 50 cycles, we believe that the Cu nanoparticles are unreversed to Cu₂O at the beginning of the reaction. The CuO generated on the surface can enhance the capacity of the electric cycles because of its high theoretical capacity.

Fig. 6. TEM of HMs-Cu₂O in the charged state after 80 cycles at 0.2C. (a) Low TEM image of a typical microsphere, (b) HRTEM image of a Cu nanoparticle in the SEI film. (c) HRTEM image of the area indicated by a red box in (a). (d) corresponding FFT of the whole area in (c). (e-g) inverse FFT filtered image of CuO/Cu₂O (110) and (111) (e, green), Cu (200) (f, red), and overlaid color map (g).

Fig. 3c shows that the capacity of HMs-Cu₂O is stable after 70 cycles. We have therefore chosen the charge state after 80 cycles at 0.2C for TEM analysis. After 80 cycles, the microspheres maintain their pristine morphology (Fig. 6a). The Cu nanoparticles in the SEI layer can be easily observed in TEM (Fig. 6b). The edge of the microsphere with a thin SEI layer (amorphous surface, indicated by arrows) is presented in Fig. 6c and the corresponding FFT (Fig. 6d) confirms the co-

existence of Cu/Cu₂O/CuO. In order to investigate the distribution of the Cu nanoparticles, the inverse FFT filtered image of the Cu (200) ring is presented in Fig. 6f and the location of Cu nanoparticles are indicated by circles in Fig. 6f and c. The corresponding inverse FFT filtered image of CuO/Cu₂O (110) and (111) (Fig. 6e) and the overlaid color map reveal the distribution of Cu and CuO/Cu₂O on the edge of the microsphere. The Cu (111) ring is not used for the inverse FFT filtering image because of the narrow separation between the $d_{\text{Cu}(111)}$ and $d_{\text{CuO/Cu}_2\text{O}(200)}$. In order to confirm the existence of CuO and investigate the distribution of the Cu/Cu₂O/CuO components, EELS analysis is performed similar with the HMs-Cu₂O after 80 cycles. The EELS spectrum (Fig. 5c, zone2) and EELS mapping (Fig. 5 i-m) demonstrates still more aggregation of CuO on the surface of microsphere. These results reveal that three different valance states of Cu⁰, Cu⁺ and Cu²⁺ can be observed through TEM and EELS at the charging state after 80 cycles. Compared to reacted HMs-Cu₂O for 50 cycles, more Cu nanoparticles appear on the surface of the polyhedron after 80 cycles, while more CuO nanoparticles are still located on the surface of the microsphere. The copper on the surface of the active facets may enhance the conductivity while it is harmful to the specific capacity because the existence of copper may prevent the reverse of more Cu₂O in the charging process. However, the presence of CuO is beneficial for the specific capacity because of its high theoretical specific capacity (674 mAh g⁻¹). Our results verify our hypothesis that copper can be further oxidized to CuO via a two-step reaction according to formula 2: Cu converts to Cu₂O, which is further converted to CuO. This leads to the specific capacity increase during the cycling process. During this process, more and more CuO nanoparticles ensure the capacity increase although some Cu nanoparticles cannot be reversed to Cu₂O and/or CuO.

After 110 cycles, the specific capacity obviously decreases (Fig. 3c). TEM characterization on the electrode after 120 cycles at the charge state (Fig. 7a) shows that the microspheres still maintain their original appearance. A closer view of the surface of the microspheres (Fig. 7b) clearly shows that the compact and smooth active facets have transformed into smaller nanoparticles, similar to the ones after 50 and 80 cycles (Fig. 4c and Fig. 6a). Fig. 7c presents a HRTEM image of the surface of the microsphere. The corresponding FFT (Fig. 7c inset) shows a strong intensity ring of Cu (200) and a weak intensity of CuO/Cu₂O (111), indicating that more Cu nanoparticles occupy the surface of the microsphere. Fig. 7d displays a HRTEM image of a Cu nanoparticle in the SEI film. The size is similar to that of the Cu nanoparticles after 80 cycles (Fig. 6b). In an intensive HRTEM study, we find more and more Cu nanoparticles in the SEI film and at the surface of the polyhedron. This means that more and more Cu nanoparticles cannot be further oxidized to Cu₂O and CuO, which naturally leads to the specific capacity decrease due to more and more loss of active material.

Fig. 7. TEM and HRTEM images of the charge state of HMs-Cu₂O after 120 cycles at 0.2C. (a) TEM image of the microspheres. (b) TEM image of the active facets. (c) HRTEM image of Cu nanoparticles and the SEI layer. (d) HRTEM image of a Cu nanoparticle.

To show the advantage of the high active facets on the surface of the microspheres, we further studied the cycle performance of HMs-Cu₂O at 1C. This capacity increase process should be similar to the process under 0.2C. The capacity increases after 200 cycles. The specific capacity increases from 150 mAh g⁻¹ at 300 cycle to 450 mAh g⁻¹ at 1500 cycles and then remains stable within a certain number of cycles (Fig. 8). The capacity of HMs-Cu₂O is largely over the theoretical capacity of Cu₂O (375 mA h g⁻¹). Compared with the highest specific capacity at 0.2C, it has a better performance at 1C. A possible reason is that at higher current density, the redox reaction of HMs-Cu₂O becomes deeper at 1C due to the exposed {111} and {110} active facets accelerating the conversion reaction. Namely, more and more Cu₂O will be oxidized to CuO during the cycling process at 1C because of the exposed {111} and {110} active facets.

On the basis of the results and analysis above, we can summarize the electrochemical behaviour of HMs-Cu₂O as follows (Fig. 9): (i) according to formula 1 and 2, due to the SEI layer formation and part of the formed Cu nanoparticles in the SEI layer are not reversed to Cu₂O, the specific capacity decreases very fast at the beginning of the reaction; (ii) with the reaction proceeding, Cu₂O can be further oxidized to CuO and some Cu nanoparticles appear at the surface of the polyhedron. Although part of the formed Cu nanoparticles cannot be reversed to Cu₂O, the formation of CuO dominates the reaction cycling process. This leads to the specific capacity increase; (iii) the formation of more and more CuO nanoparticles makes the specific capacity reach a maximum value; (iv) after the stable capacity stage, more and more CuO and Cu nanoparticles are formed, and the unreversed Cu to

Cu₂O and/or CuO dominates the final process, which leads to the specific capacity decrease.

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Fig. 8. Electrochemical properties of HMs-Cu₂O at 1C. (a) voltage profile for the 1st, 400th, 1000th and 1600th galvanostatic discharge and charge curves; (d) specific capacity as a function of cycle number.

Fig. 9. The schematic electrochemical behaviour process of the HMs-Cu₂O.

4. Conclusions

This work describes how hollow microspheres of Cu₂O with {110} and {111} active facets can be beneficial for lithium storage. The as-prepared Cu₂O electrodes display an enhanced reversible capacity and cycling stability because more dangling Cu atoms are exposed on the {111} and {110} facets and promote the conversion reaction between cuprous oxide and lithium. We systematically studied the morphological and chemical conversion of the Cu₂O microspheres upon cycling via SEM, HRTEM and EELS. At the capacity rising stage, there is the

formation of SEI layers, Cu nanoparticles in the SEI layer and CuO nanoparticles at the surface of the polyhedron. With the cycling proceeding, more and more Cu₂O is converted to CuO, resulting in a capacity maximum. Finally, more and more Cu nanoparticles cannot be reversed to Cu₂O and/or CuO, leading to a capacity decrease. This finding on the electrochemical behaviour for Cu₂O may be extended to other transition metal oxides with various phases based on a conversion reaction. We believe that our process presented here is helpful for the design of transition metal oxides based anode materials via a conversion reaction.

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Table of contents entry

The electrochemical behavior of hollow Cu₂O microspheres with {111} and {110} facets as anode for lithium storage is probed by TEM. The conversion of Cu₂O to CuO leads to capacity increase but more unreversed Cu nanoparticles result in capacity decrease during a prolonged cycling.