Microscopy Study of the Morphology of Carbonized ZIF-67 Tailored by CTAB

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Metal-organic frameworks (MOFs) have been widely used in gas storage, catalysis, biomedical and electrochemical studies [1]. Among them, the carbonized ZIF-67 exhibits excellent performance in catalytic oxygen reduction reactions (ORR) [2], and the addition of cetyl trimethyl ammonium bromide (CTAB) has a great influence on its ORR performance. Therefore, it is crucial to study the relationship between the structure and performance of the carbonized ZIF-67 with the addition of CTAB. In this study, field emission scanning electron microscopy (FESEM) is employed to investigate effects of CTAB on the morphology and electrocatalytic properties of carbonized ZIF-67.

In a typical experiment, 1.43 g of Co(NO₃)₂·6H₂O and 3.24 g of 2-methylimidazole were dissolved in 50 mL methanol with or without 0.2 g/L CTAB in the solution. Then, the solution was mixed under magnetic stirring for 20 min and stood at 30 °C for 24 h. Solid product was separated by centrifugation and washed with methanol for three times, followed by vacuum drying at 40 °C for 48 h. The as-prepared ZIF-67 above was carbonized under N₂ atmosphere at 600 °C for 3 h with a heating rate of 2 °C/min and naturally cooled down to room temperature. The morphology of ZIF-67 and carbonized products was characterized by FESEM (JSM-6700F, JEOL), and electrochemical tests were conducted at a rotating ring-disk electrode (RRDE-3A, ALS Co.) on a CHI760D electrochemical work station with Ag/AgCl as reference electrode and Pt wire as counter electrode.

As shown in Figure 1, urchin-like carbon material is obtained from carbonizing ZIF-67 without CTAB, and a plurality of elongated carbon nanotubes (CNTs) appear on the surface of main structures. The nanotubes destroy original angular structure. In contrast, with CTAB, the carbonized ZIF-67 contains a small amount of CNTs (Figure 2), and the structure is clear and angular to retain the sodalite topology of the precursor (Figure 3a). FESEM characterizations clearly indicate that the microstructure of the carbonized ZIF-67 is affected by the addition of CTAB. Electrochemical performances of the two samples are given in Figure 3b. By comparison, the carbonized ZIF-67 without CTAB has better electrochemical performance. The onset potential (E_{onset}), half wave potential ($E_{1/2}$) and limiting current density (I_{JM}) of the carbonized ZIF-67 without CTAB are -0.067 V, -0.135 V and -6.39 mA cm⁻², respectively, positively shifted by 43 mV, 55 mV and 0.67 mA cm⁻² compared with the carbonized ZIF-67 with CTAB (E_{onset} = -0.11 V, $E_{1/2}$ = -0.19 V and I_{JM} = -5.72 mA cm⁻²). This means, the formation of CNTs on the surface of carbonized ZIF-67 can serve as electron transport channels and provide more active sites for electrocatalytic reactions.

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References

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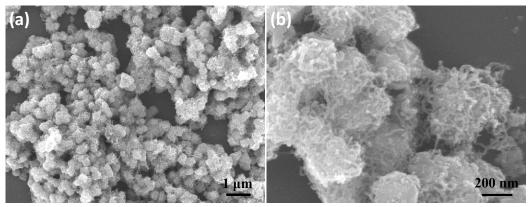


Figure 1. SEM images of carbonized ZIF-67 without CTAB.

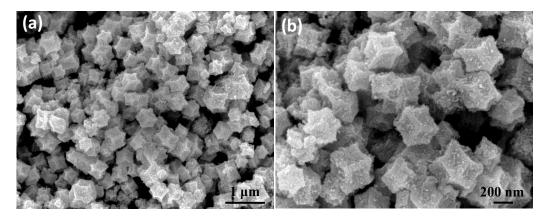


Figure 2. SEM images of carbonized ZIF-67 with the addition of CTAB.

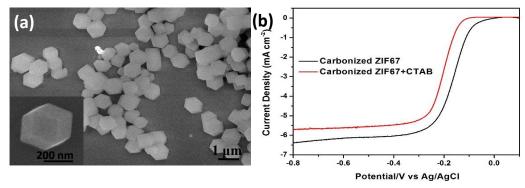


Figure 3. (a) SEM images of the ZIF-67 precursor, (b) linear sweep voltammetry (LSV) curves of carbonized ZIF-67 with/without CTAB at 1600 rpm in O₂-saturated 0.1 M KOH solution.