Hierarchical Ni-Fe layered double hydroxide/MnO₂ sphere architecture as an efficient noble metal-free electrocatalyst for ethanol electro-oxidation in alkaline solution

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Supporting information

1. Experimental

CO stripping experiments were performed as follows: after purging the solution with N_2 for 20 min, gaseous CO was bubbled for 15 min to form CO adlayer on catalysts while maintaining potential at 0.3 V. Then excess CO in solution was purged with N_2 for 20 min and CO stripping voltammetry was recorded in 1 M KOH solution at 50 mV s⁻¹.

2. Results and discussion





Fig. S1 (a) CO stripping curves on MnO₂ (a), Ni-Fe LDH (b) and LDH@MnO₂ (c) recorded in 1 M KOH solution

To have a deep insight into the synergistic effect between Fe-Ni LDH and the MnO₂, the CO stripping voltammograms were recorded, as shown in Fig. S1. As shown in Fig. S1(a), it is clear that there is no obvious difference between the first and second CV curves from CO stripping voltammograms. Therefore, it manifested that the CO cannot be adsorbed on MnO₂. Comparing the CO stripping CV curves from Ni-Fe LDH and LDH@MnO₂ catalysts, as shown in Fig. S1(b) and (c), no great differences was found no matter in magnitude of current, or in CV shapes. Furthermore, the synergistic effect between Fe-Ni LDH and the MnO₂ cannot be explained by CO stripping voltammograms. The main reason for the synergistic effect between Fe-Ni LDH and the MnO₂ cannot be assorbed to the facile adsorption of OH_{ads} species onto MnO₂ [1]. Therefore, it is inferred that MnO₂ could increase the

concentration of OH_{ads} species on Ni-Fe LDH surface, and these OH_{ads} can react with C_{1ad} intermediate species to produce CO_2 or water soluble products, releasing the active sites on LDH for further electrochemical reaction [2-5].

Reference

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