## **Supporting Information**



**S. I. Figure 1.** As described in S. I. Figure 1, the Raman spectrum shows two characterized groups. After serious analyses, one of them is attributed to the crystallized  $Co_3O_4$  (peaks of  $E_g$ ,  $F^1{}_{2g}$ ,  $A_{1g}$  and  $F^2{}_{2g}$ )<sup>[1]</sup>. The other one can be assigned to amorphous carbon (D, 2D and D+G bands) and graphitized carbon (G band). D band is believed to result from the carbon's disorder and G band normally is ascribed to the graphitized carbon <sup>[2,3]</sup>. Both of the D band and G band for the peapod-like samples exhibit the strong intensities, implying the co-existence of amorphous and graphitized carbons in the samples. 2D and D+G bands also confirm the presence of the disordered and randomly arranged graphitized layers in the peapod-like  $Co_3O_4$ @carbon nanocomposites<sup>[2]</sup>.

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**S.I. Figure 2.** S. I. Figure 2a is the representative TEM image for the homemade  $Co_3O_4$  nanoparticles. For the preparation of these  $Co_3O_4$  nanoparticles, generally 12.5ml deionized water (D. I. water) and 12.5ml concentrated ammonia (fuming, 28wt%) are mixed homogeneously under strong magnetic stirring for 5 min. Then 5ml 1M aqueous  $Co(NO_3)_2$  solution will be introduced into the former solution to form the final solution in brown color after another 30 min stirring. Afterwards, the solution is transferred into 45ml Teflon-lined autoclave and heated at 160-180°C for 16 hrs. The precipitates in the liner are washed with 3 times of deionized water and 1 time of ethanol by centrifugation. S. I. Figure 2b is the corresponding HRTEM image for the as-collected  $Co_3O_4$  nanoparticles.