

# Supplementary Information for

## Controlled Synthesis of Nanorods/Nanorings of a Novel Co-Cu Complex in Microemulsion at Room Temperature

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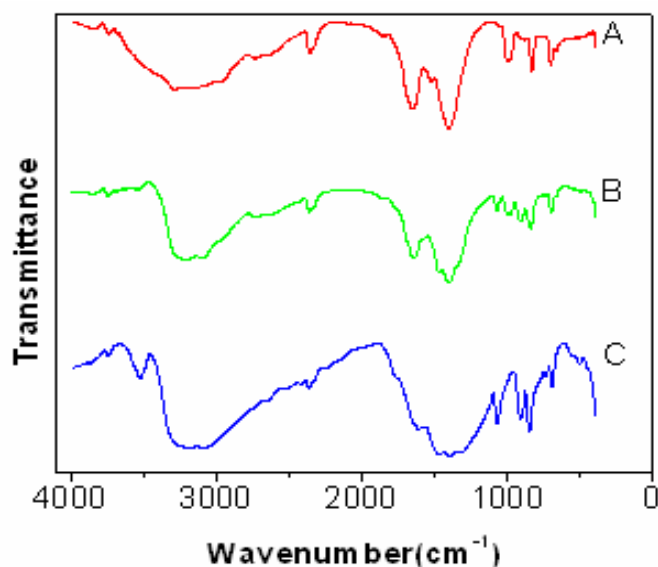
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## 1. Additional analyses of Infrared Absorption Spectra showing the composition of the Co-Cu complex nanorods and nanorings.

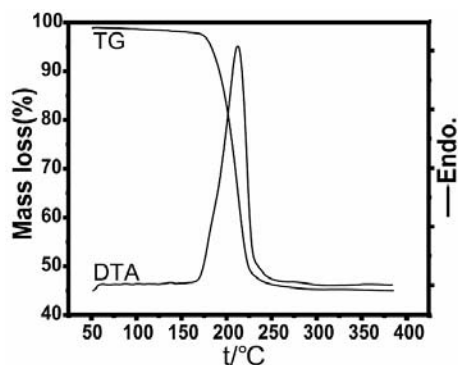


**Figure S1** FT-IR spectra of the nanorods(A), nanorings(B) and the bulk



The Fourier Transform Infrared (FTIR) spectra of the Co-Cu complex nanorods, nanorings, and the bulk  $[\text{Co}(\text{NH}_3)_6]_3[\text{Cu}_4(\text{OH})(\text{CO}_3)_8]\cdot 2\text{H}_2\text{O}$  are shown in Figure S1. The spectral peaks provide vibrational information for the functional groups in the samples, which is very important for characterizing the structure and composition of the materials. From Fig S1, it can be clearly seen that there is nearly a one-to-one correspondence of the major peaks among the spectra of the Co-Cu complex nanorods, nanorings and the bulk Co-Cu complex. In addition, the peak positions are also close in spite of small shifts. This indicates that the functional groups and structures are similar in the three materials. The small spectral shifts may be caused by the change from an fcc structure to a nanostructure. We conclude that  $[\text{Co}(\text{NH}_3)_6]_3[\text{Cu}_4(\text{OH})(\text{CO}_3)_8]\cdot 2\text{H}_2\text{O}$  has six characteristic peaks: the stretching vibration of O-H ( $3700\text{-}3650\text{ cm}^{-1}$ ), the stretching vibration of O-H in the crystallization water (appearing at  $3540\text{ cm}^{-1}$ ), the symmetric stretching vibration of N-H ( $3500\text{-}2600\text{ cm}^{-1}$ ), the stretching vibration of C=O ( $2400\text{ cm}^{-1}$ ), the bending vibration of O-H ( $1600\text{ cm}^{-1}$ ), and the bending vibration of N-H ( $1500\text{-}1300\text{ cm}^{-1}$ ).

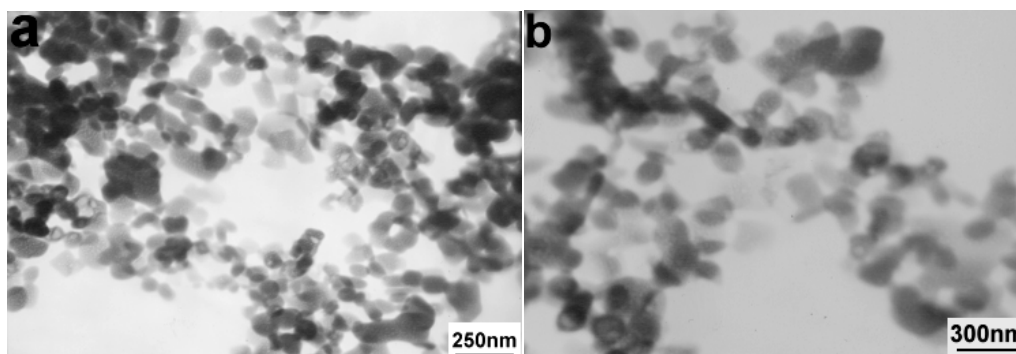
## 2. The result of thermal analysis.



**Figure S2** TG and DTA curves of the Co-Cu complex nanorods

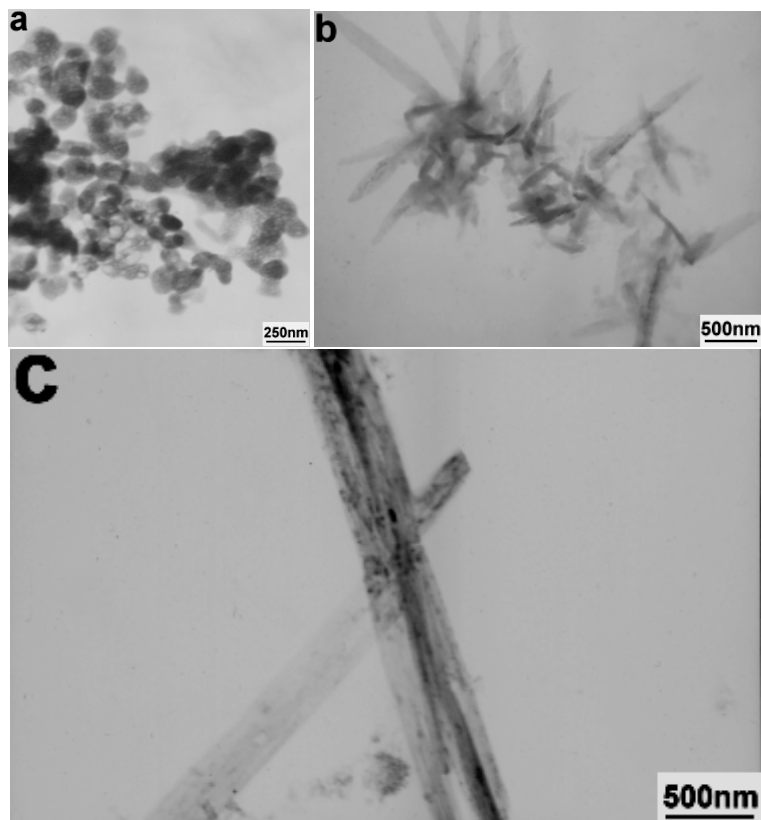
TG and DTA curves of the nanorings are similar to those of the nanorods. The molecular formula of this compound is  $[\text{Co}(\text{NH}_3)_6]_3[\text{Cu}_4(\text{OH})(\text{CO}_3)_8]\cdot 2\text{H}_2\text{O}$ , and its molecular weight is  $M_1=1269.79$ . Then from Figure S2 it can be clearly known that the crystallization water in the complex was stripped down at 120 °C and the WT (%) = -2.2%, in good agreement with the theoretic value. The process from 120 °C (entirely losing the water of crystallization) to 300 °C, the complex was oxidized and decomposed into  $\text{Co}_2\text{O}_3$  and  $\text{CuO}$ . This weight losing was about -55.1% and the theoretic value was -55.36%. Based on the analyses of XRD patterns, FT-IR spectra and TG-DTA curves, it is reasonable that the molecular formula of the complex is  $[\text{Co}(\text{NH}_3)_6]_3[\text{Cu}_4(\text{OH})(\text{CO}_3)_8]\cdot 2\text{H}_2\text{O}$ .

3. The effect of  $w$  and  $t$  on the morphologic formation of the Co-Cu complex nanomaterials (Here, the “ $w$ ” is  $[\text{H}_2\text{O}]/[\text{surfactant}]$  molar ratio in the system; “ $t$ ” is the time of the reaction).



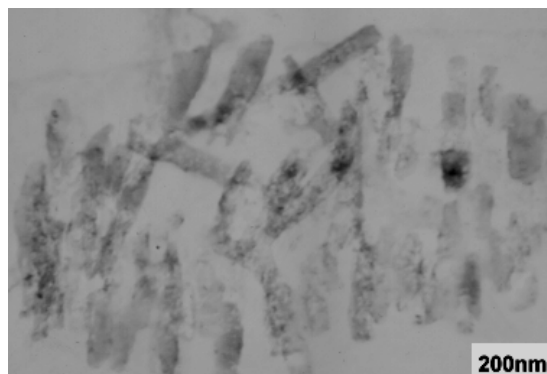
**Figure S3** The TEM images of the different parts of the products obtained with  $w = 35$  and  $t = 24$  h

Under this condition, only nanoparticles were obtained and their diameters were about 50 nm. This indicated that there was no micelle structures formed in the system, and the just microemulsion was round in shape.



**Figure S4** The TEM images of the products obtained with  $w = 40$  and (a)  $t = 1$  h, (b)  $t = 12$  h, (c)  $t = 24$  h

Keeping  $w = 40$  constant, we changed the reaction time as  $t = 1$  h (a), 12 h (b) and 24 h (c). The result (see figure S4) was shown as follows: Figure 4a ( $t = 1$  h) shows the morphology is spherical; Figure 4b ( $t = 12$  h) indicates a short needle-like morphology with a diameter of 80 nm and a length of 500 nm, respectively; from Figure 4c ( $t = 24$  h) nanorods can be observed with 100 nm in diameter and 2-3  $\mu\text{m}$  in length. It is assumed that when  $w = 40$ , the microemulsion may form a circular micro-reactor, and with the nucleation and growth of the crystal a circular micro-reactor can change into a rod-like structure because the required number of charge compensating cations remarkably increases as well.



**Figure S5** the TEM images of the products obtained with the condition is  $w = 50$  and  $t = 24$  h.

From the TEM images, we can find that the quality of the nanorods have some defects with low-grade crystallization.