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ORIGINAL ARTICLE

Synthesis of CNTs/CuO and its catalytic performance on the thermal decomposition of ammonium perchlorate

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Abstract Copper oxide (CuO) nanoparticles were successfully deposited on carbon nanotubes' (CNTs) surface via complex-precipitation method, the nanocomposite was characterized by transmission electron microscopy (TEM), scanning electron microscopy (SEM), X-ray photoelectron spectroscopy (XPS), X-ray powder diffraction (XRD), Raman spectroscopy, Fourier transform infrared (FT-IR) and Brunauer–Emmett–Teller (BET). The catalytic performance of CNTs/CuO on ammonium perchlorate (AP) decomposition was analyzed by differential thermal analyzer (DTA), the DTA results showed its excellent catalytic effect on AP decomposition, as 8 wt.% CNTs/CuO was added in AP, the second exothermic peak temperature decreased by 158 °C. Such composite may be a promising candidate for catalyzing the AP thermal decomposition.

1. Introduction

Ammonium perchlorate (AP), a white crystalline substance, is the most commonly used oxidizer in composite solid propellants (CSPs). Obviously, the thermal decomposition characteristic of AP affects the performance of CSPs. Generally speaking, the lower the decomposition temperature of AP, the shorter the delay time of propellant ignition, the higher the combustion rate and the better the performance of CSPs [1]. Over the past several decades, numerous catalysts have been employed to decrease the decomposition temperature of AP [2–5]. The results suggested that the decomposition process of AP was remarkably sensitive to the catalysts and the nanometer sized catalysts possessed better catalytic property as compared with their bulk size. Among the metal oxide catalysts, copper oxide (CuO) nanoparticles (NPs) showed particular catalytic effect due to their high concentration of dislocations and large surface areas [6]. However, nanoparticles are easy to aggregate due to their large surface energy, which will further affect their catalytic performance.

Extensive attention has been paid on CNTs since their first discovery [7] in many fields, such as catalyst [8], lithium ion batteries [9] and solar cells [10]. For its application in catalyst field, it should be noted that the perfect structure along its tube

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wall allows them to be the excellent dispersing-supporters for the NPs [11–13]. Our earlier work [14] showed that the CNTs/Cu exhibited better catalytic effect on AP thermal decomposition than the simple mixture of CuNPs and CNTs did, indicating CNTs can improve the catalytic performance of CuNPs as a supporter. However, CNTs/CuO has not been reported to catalyze the thermal decomposition of AP. Herein, in our work CNTs/CuO nanocomposite was prepared, compared with the simple mixture of CuONPs and CNTs, CNTs/CuO nanocomposite exhibited higher catalytic activity for AP thermal decomposition.

2. Experimental

2.1. Preparation of CNTs

Catalytic chemical vapor deposition (CCVD) method was employed to synthesize CNTs. In a typical experiment, the volume of benzene (carbon source) and thiophene (growth stimulant) was 150 and 1 ml, respectively. Catalyst (ferrocene, about 2 g) was uniformly spread on an alumina boat placed in a horizontal tube furnace. After the temperature of the furnace was raised to 1170 °C under a flow of nitrogen gas at a rate of 200 ml/min, hydrogen gas was introduced at a rate of 380 ml/min for 15 min. The as-prepared CNTs were functionalized by 100 ml concentrated mixed acids (H_2SO_4/HNO_3 = 3:1) under stirring condition at 75 °C for 2 h.

2.2. Synthesis of CNTs/CuO composite

Firstly, CNTs, SDS and copper chloride($CuCl_2$) were dispersed in 50 ml distilled water under ultrasound condition for 30 min, subsequently, superfluous aqueous ammonia(e) was added in the solution, then sodium hydroxide (NaOH) was added into the solution stirred at 50 °C drop by drop, after that, the solution was stirred at 50 °C for another 1 h. Finally, the sample was filtered, washed with distilled water several times, dried and calcinated in muffle furnace at 400 °C for 2 h. CuONPs were prepared following the same procedure above without adding CNTs at the beginning.

2.3. Characterization

X-ray powder diffraction (XRD) analysis of the samples was carried out with a German D8ADVANCE X-ray diffractometer with Cu Kα radiation (k = 1.54056 Å). X-ray photoelectron spectroscopy (XPS) was performed with an American Thermo ESCALAB 250 electron spectrometer using Al K irradiation. Morphology of the samples was investigated using a transmission electron microscopy (TEM) on a FEI instrument (T-12 TENcai) subjected to an acceleration voltage of 120 kV and a scanning electron microscopy (SEM) using a JEOL 35. Raman spectra of the samples were recorded in the frequency range of 200–2000 cm^{-1} using a Raman spectrometer (JY HR-800 type) with a laser excitation line at 532 nm. Fourier transform infrared (FT-IR) spectra were recorded on a NicdeT 740 spectrometer using pressed KBr pellets to test the chemical bonding of the samples from 500 to 3750 cm^{-1}. The Brunauer–Emmett–Teller (BET) surface area of as-synthesized samples was determined by using an instrument of the Beckman Coulter Co. Ltd., USA.

2.4. Catalytic analysis

Thermal decomposition study of pure AP and AP with catalysts was performed with the differential thermal analyzer (DTA, TA instrument SDT-Q600) under a heating rate of 20 °C/min in a static N_2 atmosphere with α-Al_2O_3 as reference material. Sample of approximately 1.5 mg was taken. The mass content of CNTs/CuO composite added in AP was 4 wt.%, the catalytic performance of the mixture of CNTs and CuONPs was also carried out in the same way as comparison. At the same time, CNTs/CuO with mass content of 1 wt.% and 8 wt.% was selected to study the effect of the mass content of CNTs/CuO on AP thermal decomposition.

3. Results and discussion

Fig. 1a shows the SEM image of CNTs/CuO, in which CuONPs are well dispersed along the CNTs surface. Fig. 1b shows the TEM image of CNTs/CuO, which further proves the uniform disperse of CuONPs on CNTs, also indicates the uniform thickness of the loading layer. It should also be mentioned that although the TEM specimen suffers from long time sonication, perfect loading of CuONPs on CNTs can still be seen clearly, indicating the strong force between CuONPs and CNTs [15].

Fig. 2 illustrates the XRD patterns of CNTs (Fig. 2a) and the synthesized CNTs/CuO composite (Fig. 2b). For the XRD pattern of CNTs, three characteristic peaks at 26.41°, 43.6° and 53.12° correspond to the (002), (101) and (004) inter-planar spacing of CNTs, respectively. For the XRD pattern of CNTs/CuO, the intensity of the three characteristic peaks of CNTs are weakening or disappearing, maybe caused by the fine loading of CuONPs on CNTs surface. Besides the diffraction peaks of CNTs, other strong diffraction peaks could be assigned to CuO (JCPDS Card No. 80-1916). From the Scherrer formula, it can be calculated that the average size of CuO on CNTs is 25 nm approximately. To further confirm the Cu species and its content on CNTs surface, XPS analysis was carried out (Fig. 3). Fig. 3a shows the XPS result of CNTs/CuO, from which C, O and Cu can be easily detected, according to the surface element analysis, the content of C, O and Cu is 45.07%, 34.57% and 20.36%, respectively. Fig. 3b shows the XPS result of Cu 2p, the two main peaks at about 933.6 and 953.6 eV are associated with the binding energy of Cu $2p_{3/2}$ and Cu $2p_{1/2}$, respectively, which confirm the formation of Cu^{2+}. Fig. 3c shows O 1s XPS curves, where the non-lattice oxygen species (e.g., hydroxyl group, adsorbed oxygen species) with peak position at 531.2 eV (marked as I) is evident. The peak at 529.6 eV (marked as II) corresponds to the surface lattice oxygen [16]. The content of non-lattice oxygen and lattice oxygen is 39.01% and 60.99% in the detection layer of the XPS analyses, respectively, which indicate that 20.36% Cu exists in the form of CuO.

To have a far better knowledge of the surface structure of the composite, Raman spectra of CNTs (Fig. 4a) and CNTs/CuO (Fig. 4b) are presented. As shown in Fig. 4b, the peak at 292 cm^{-1} is attributed to the Ag mode and the peaks at 342 and 627 cm^{-1} are attributed to the Bg mode of CuO. Both the spectra display the peak at about 1350 cm^{-1} (D-band) associated with the vibrations of carbon atoms in the disordered graphite structure and the peak at about 1585 cm^{-1}