Research Article

The Multiple Effects of Precursors on the Properties of Polymeric Carbon Nitride

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Polymeric graphitic carbon nitride (g-C$_3$N$_4$) materials were prepared by direct pyrolysis of thiourea, dicyandiamide, melamine, and urea under the same conditions, respectively. In order to investigate the effects of precursors on the intrinsic physicochemical properties of g-C$_3$N$_4$, a variety of characterization tools were employed to analyze the samples. The photocatalytic activity of the samples was evaluated by the removal of NO in gas phase under visible light irradiation. The results showed that the as-prepared CN-T (from thiourea), CN-D (from dicyandiamide), CN-M (from melamine), and CN-U (from urea) exhibited significantly different morphologies and microstructures. The band gaps of CN-T, CN-D, CN-M, and CN-U were 2.51, 2.58, 2.56, and 2.88 eV, respectively. Both thermal stability and yield are in the following order: CN-M > CN-D > CN-T > CN-U. The photoactivity of CN-U (31.9%) is higher than that of CN-T (29.6%), CN-D (22.2%), and CN-M (26.8%). Considering the cost, toxicity, and yield of the precursors and the properties of g-C$_3$N$_4$, the best precursor for preparation of g-C$_3$N$_4$ was melamine. The present work could provide new insights into the selection of suitable precursor for g-C$_3$N$_4$ synthesis and in-depth understanding of the microstructure-dependent photocatalytic activity of g-C$_3$N$_4$.

1. Introduction

Visible-light-active photocatalysts are attracting increasing interests for their potential application in the areas of environmental protection, material science, and solar energy conversion by directly utilizing nature sunlight and artificial indoor illumination [1–3]. During the past few decades, many efforts have been devoted to develop novel and efficient visible light photocatalytic systems, including inorganic photocatalysts (e.g., $\text{Fe}_2\text{O}_3$, $\text{BiOCl}$, $\text{Cu}_2\text{SnS}_3$, $\text{SrTiO}_3$, and $\text{WO}_3$/$\text{BiOCl}$) [4–7], organic photocatalysts (e.g., graphitic carbon nitride, g-C$_3$N$_4$), and elemental photocatalysts (e.g., Si, P, S, and Se) [8–10].

In particular, g-C$_3$N$_4$ material as a novel metal-free organic photocatalyst has triggered great attention in the fundamental and applied scientific researches due to its suitable electronic band structure, nontoxic nature, biocompatibility, high thermal and chemical stability, easily available at low cost, and amenability to chemical modification [11–16]. Properties mentioned previously make it a promising organic photocatalyst for solar energy converting, organic photosynthesis, drug delivery, and environment remediation under visible light irradiation [17–19].

g-C$_3$N$_4$ can be prepared by the direct pyrolysis of various organic precursors. Dong et al. prepared g-C$_3$N$_4$ from thiourea and urea at 550°C for 2 h [20, 21], respectively. Yan et al. obtained g-C$_3$N$_4$ from melamine at 550°C for 4 h [22]. Xu et al. prepared g-C$_3$N$_4$ by directly heating dicyandiamide first at 350°C for 2 h and then 550°C for another 2 h [23], respectively. The different pyrolysis conditions for g-C$_3$N$_4$ synthesis made the comparison of the precursors difficult. Until now, little information is known about the effects of precursors on intrinsic physicochemical properties of g-C$_3$N$_4$ under the same pyrolysis conditions, including morphology, band gap,
2. Experimental

2.1. Synthesis of g-C₃N₄. In a typical synthesis, 12 g of thiourea was put into a semi-closed alumina crucible with a cover and then heated to 550 °C in a muffle furnace for 3 h at a heating rate of 10 °C min⁻¹. Following the same procedure, g-C₃N₄ samples were prepared by directly treating dicyandiamide, melamine, and urea under the same condition (550 °C for 3 h in air), respectively. The precursors are easily available in the chemical industry at low cost (see Table 1). Various characterization tools were utilized to analyze the effects of precursors on the intrinsic physicochemical properties and photocatalytic activity of g-C₃N₄. The results showed that the precursors had significant effect on the properties of g-C₃N₄. Considering the cost, toxicity, and yield of the precursors and the properties of g-C₃N₄, the best precursor was confirmed.

2.2. Characterization. The crystal phases of the samples were analyzed by X-ray diffraction with Cu Kα radiation (XRD: model D/max RA, Rigaku Co., Japan). FT-IR spectra were recorded on a Nicolet Nexus spectrometer on samples embedded in KBr pellets. The morphology and structure of the samples were further examined by transmission electron microscopy (TEM: JEM-2010, Japan). The UV-vis diffuse reflection spectra were obtained for the dry-pressed disk samples using a Scan UV-vis spectrophotometer (UV-vis DRS: UV-2450, Shimadzu, Japan) equipped with an integrating sphere assembly, using BaSO₄ as the reflectance sample. Nitrogen adsorption-desorption was conducted on a nitrogen adsorption apparatus (ASAP 2020, USA). The thermal stability was detected by using thermogravimetry analysis (TG-DSC, Netsch STA 449F3) under N₂ gas atmosphere. All the samples were degassed at 150 °C prior to measurements.

2.3. Visible Light Photocatalytic Performance. The photocatalytic activity was investigated by removal of NO at ppb levels in a continuous flow reactor at ambient temperature. The volume of the rectangular reactor, which was made of poly-methyl methacrylate plastics and covered with quartz-glass, was 4.5 L (30 cm × 15 cm × 10 cm). A 100 W commercial tungsten halogen lamp (General Electric) was vertically placed outside and above the reactor. Four minifans were used to cool the flow system. Adequate distance was also kept from the lamp to the reactor for the same purpose to keep the temperature at a constant level. For the visible light photocatalytic activity test experiment, a UV cut-off filter (420 nm) was adopted to remove UV light in the light beam.

For each photocatalytic activity test, two sample dishes (with a diameter of 12.0 cm) containing photocatalyst powder were placed in the center of the reactor. The weight of the photocatalyst used for each dish was kept at 0.1 g. g-C₃N₄ sample was added into 30 mL of H₂O and sonicated for 10 min, and then photocatalyst samples were prepared by coating aqueous suspension of the samples onto the glass dishes. The coated dish was pretreated at 60 °C to remove water in the suspension and then cooled to room temperature before photocatalytic testing.

The NO gas was acquired from a compressed gas cylinder at a concentration of 100 ppm of NO (N₂ balance, BOC gas) with the National Institute of Standards and Technology (NIST) standard. The initial concentration of NO was diluted to about 600 ppb by the air stream supplied by a zero air generator (Thermo Environmental Inc., model 111). The relative humidity at indoor environmental condition is 40%–80%. The desired relative humidity in the present system is controlled at 50% in the gas flow which could simulate the indoor environmental conditions. The desired relative humidity (RH) level of the NO flow was controlled at 50% by passing the zero air streams through a humidification chamber. The gas streams were premixed completely by a gas blender, and the flow rate was controlled at 2.4 L min⁻¹ by a mass flow controller. After the adsorption-desorption equilibrium was achieved, the lamp was turned on. The concentration of NO was continuously measured by a chemiluminescence NO analyzer (Thermo Environmental Instruments Inc., model 42c), which monitors NO, NO₂, and NOx (NOx represents NO + NO₂) with a sampling rate of 0.7 L min⁻¹. The removal ratio (η) of NO was calculated as η (%) = (1 – C/C₀) × 100%, where C and C₀ are concentrations of NO in the outlet steam and the feeding stream, respectively. The kinetics of photocatalytic NO removal reaction is a pseudofirst order reaction at low NO concentration as ln(C₀/C) = kt, where k is the apparent rate constant.

Table 1: The cost, acute toxicity, and melting point of thiourea, dicyandiamide, melamine, and urea [16].

<table>
<thead>
<tr>
<th>Precursors</th>
<th>Cost ($/t)</th>
<th>Acute toxicity LD₅₀ (oral, rat)</th>
<th>Melting point (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thiourea</td>
<td>3906</td>
<td>125 mg/kg</td>
<td>182</td>
</tr>
<tr>
<td>Dicyandiamide</td>
<td>1538</td>
<td>4000 mg/kg</td>
<td>209</td>
</tr>
<tr>
<td>Melamine</td>
<td>2025</td>
<td>3248 mg/kg</td>
<td>345</td>
</tr>
<tr>
<td>Urea</td>
<td>450</td>
<td>8500 mg/kg</td>
<td>133</td>
</tr>
</tbody>
</table>
of graphitic materials, corresponding to the in-plane tri-triazine units which formed one-dimensional (1D) melon strands. Further observation implies (Figure 1(b)) that the typical (002) peak is 27.43° for CN-T, 27.47° for CN-D, 27.60° for CN-M, and 27.72° for CN-U, respectively, indicating that the interplanar distance tends to decrease and g-C₃N₄ structure becomes more compact. The dense structure can be ascribed to the localization of the electrons and stronger binding between the layers [27, 28]. The dominant (002) diffraction peaks shifted toward higher diffraction angles for CN-U compared with CN-T, indicating that O-containing precursors (urea) could improve the polycondensation of g-C₃N₄. In addition, the intensity and breadth of the XRD patterns (Figure 1) are different, due to the presence of different nanostructures and morphologies of the as-prepared g-C₃N₄ [29].

The chemical structures of dicyandiamide and melamine contain a C=N bond which plays a key role in the formation of g-C₃N₄. However, both the S-containing thiourea and O-containing urea do not contain a C=N bond. The fact indicates that the formation mechanisms and condensation degrees of thiourea and urea are different from that of dicyandiamide and melamine [20–22, 30].

Figure 2(a) shows that the weak absorption at the 700–800 cm⁻¹ region is assigned to the bending vibration mode of CN heterocycles and the characteristic out of plane bending vibration mode of the triazine units at 810 cm⁻¹ for all the samples [31]. All the samples reveal several bands in the 1200–1650 cm⁻¹ region, which corresponds to the typical stretching vibration modes of the heptazine heterocyclic ring (C₆N₇) units [32].

The stretching vibration modes of N–H and O–H at 3000–3500 cm⁻¹ are also observed (Figure 2(b)), indicating the existence of uncondensed amino groups and absorbed H₂O molecules in all the samples [33]. It can be seen that the peak intensity of CN-U at 400–4000 cm⁻¹ is stronger than that of CN-T, CN-D, and CN-M, indicating that O-containing
precursors could improve the polycondensation of g-C₃N₄ [34].

The morphologies and microstructures of CN-T, CN-D, CN-M, and CN-U were observed by TEM. As shown in Figure 3, all the samples consist of large amounts of packing layers with different sizes of nanosheets and nanoparticles, which exhibit obviously wrinkles and irregular shapes. Figure 3(a) shows that CN-T is composed of smooth and flat layers. Numerous large buckle nanosheets with aggregated structures which formed smooth layers in the CN-D sample can be seen in Figure 3(b). By observing carefully for CN-M (Figure 3(c)), the layers consist of dense and thick nanosheets with irregular shape. Figure 3(d) shows that CN-U is composed of smooth and thin layers with typical porous morphology and loose structure. The result clearly shows that CN-U prepared from urea favors the formation of typical porous structure. The fact indicates that the morphologies and microstructures of the resultant g-C₃N₄ strongly depend on the different heteroatom-containing precursors.

As shown in Figure 4(a), all the samples exhibit excellent visible light absorption, and the absorption edges of the samples shift apparently to longer wavelengths from CN-U to CN-T. The band energies (E_g) which can be estimated from the intercept of the tangents to the plots of (Ahv)^1/2 versus photo energy (Figure 4(b)) are 2.51, 2.58, 2.56, and 2.88 eV for CN-T, CN-D, CN-M, and CN-U, respectively. The fact indicates that the precursors could affect the band gap and absorption edge of g-C₃N₄, which may be caused by the different local structures, defects, and degrees of condensation during the pyrolysis [8, 11].

Figure 5 shows the nitrogen adsorption-desorption isotherms and corresponding curves of the pore size distribution for CN-T, CN-D, CN-M, and CN-U. The isotherms of CN-T (Figure 5(a)), CN-D (Figure 5(c)), and CN-M (Figure 5(e)) can be classified to type IV, which indicates the presence of mesopores. The CN-U exhibits type III behavior (Figure 5(g)), which can be ascribed to the weak adsorbent-adsorbent interaction [35]. The specific surface areas and pore volumes of CN-T (23 m²/g and 0.14 cm³/g) and CN-U (153 m²/g and 0.40 cm³/g) are significantly higher than that of CN-D (18 m²/g and 0.09 cm³/g) and CN-M (14 m²/g and 0.06 cm³/g). The data illustrates that the heteroatoms of sulfur and oxygen play a key role in increasing the specific surface area and enlarging the pore volume during the condensation. The formation of H₂S and CO₂, release of NH₃, and generation of additional H₂O vapor during the pyrolysis of thiourea and urea favor the expansion of the packing layers and porous structure [32, 35]. Further observation implies that all the samples have micropores and mesopores. The interconnected porous network could mainly contribute to the formation of micropores and mesopores of CN-T (Figure 5(b)) and CN-U (Figure 5(h)), the aggregation of nanosheets and nanoparticles could result in the formation of the micropores and mesopores of CN-D (Figure 5(d)) and CN-M (Figure 5(f)) [19, 20].
In order to understand the thermal stability of CN-T, CN-D, CN-M, and CN-U, the thermal analysis was carried out by using TG-DSC and the heating rate of 20°C/min under N₂ gas atmosphere. It can be found that the CN-T (Figure 6(a)), CN-D (Figure 6(b)), and CN-M (Figure 6(c)) became unstable when the heating temperature was above 550, 563, and 575°C, respectively. The exothermic peaks can be seen at 672.4 for CN-T, 683.1 for CN-D, and 649.5°C for CN-U, which should be attributed to the decomposition for the samples. Figure 6(d) shows that no significant weight loss of CN-U is recorded when the temperature is below 530°C, and complete decomposition of CN-U occurred at 750°C. Further observation reveals that there are two strong exothermic peaks at 550.1 and 682.4°C of CN-U, which can be ascribed to the sublimation and thermal decomposition [36], respectively. It should be noted that the thermal stability of the samples is different, due to the different degrees of condensation and the packing between the layers during the polymerization [37, 38].

The photocatalytic performance of the as-prepared samples was further evaluated by removal of NO in gas phase in order to demonstrate their potential ability for indoor air purification under visible light irradiation at room temperature (Figure 7). There are four reactions of the photocatalytic materials, which involved that NO reacted with the photo-generated reactive radicals and produced HNO₂ and HNO₃ displayed in the following [39]:

\[
\begin{align*}
\text{NO} + 2\cdot\text{OH} & \rightarrow \text{NO}_2 + \text{H}_2\text{O} \\
\text{NO}_2 + \cdot\text{OH} & \rightarrow \text{NO}_3^- + \text{H}^+ \\
\text{NO} + \text{NO}_2 + \text{H}_2\text{O} & \rightarrow 2\text{HNO}_2 \\
\text{NO} + \cdot\text{O}_2^- & \rightarrow \text{NO}_3^-
\end{align*}
\] (1)

Figure 7(a) shows the variation of NO concentration (Cₐ/Cₐ₀ %) with irradiation time over the samples under visible light irradiation (λ > 420 nm). Here, Cₐ₀ is the initial concentration of NO, and C is the concentration of NO after photocatalytic reaction for time t. As shown in Figure 7(a), the concentration of NO for all samples decreased rapidly due to the photocatalytic degradation in 5 min. However, the reaction intermediates, and final products generated during irradiation may occupy the active sites of photocatalyst, which result in the decrease in activity. After 40 min irradiation, the removal rates and apparent rate constants of CN-T (29.2% and 0.310 min⁻¹) and CN-U (32.2% and 0.384 min⁻¹) are higher than that of CN-D (22.2% and 0.079 min⁻¹) and CN-M (26.2% and
Figure 5: N₂ adsorption-desorption isotherms of CN-T (a), CN-D (c), CN-M (e), and CN-U (g) and the corresponding pore size distribution curves of CN-T (b), CN-D (d), CN-M (f), and CN-U (h).
**Figure 6**: TG-DSC thermalgrams for CN-T (a), CN-D (b), CN-M (c), and CN-U (d).

**Figure 7**: Photocatalytic activity (a) and initial rate constants (b) of CN-T, CN-D, CN-M, and CN-U for removal of NO under visible light irradiation ($\lambda > 420$ nm) at room temperature.
0.298 min⁻¹), respectively. Generally, CN-T, CN-D, and CN-M with narrower band gaps than CN-U (see Table 2) are favorable for photocatalytic reaction, due to the enhanced visible light absorption with more available photogenerated electron hole. In fact, the high surface areas and large pore volumes can enhance the adsorption of reactants and diffusion of the reaction products and provide more efficient active sites for photochemical reaction [40, 41]. Thus, the surface area and pore volume of CN-U play a more significant role in NO degradation than the electronic properties. However, it is interesting to find that the CN-M exhibits higher photocatalytic activity than that of CN-D. Figure 4(b) shows that the band energy of CN-M is narrower than that of CN-D, which can enhance visible light absorption in CN-M sample. On the other hand, the enhanced condensation of CN-M can effectively reduce the number of structure defects (e.g., uncondensation groups of –NH₂ and –NH) which always capture the electrons or holes to prevent the following photoredox reaction [8]. The fact indicates that the photocatalytic efficiency strongly depends on the surface area, pore volume, band energy, and the degree of condensation of g-C₃N₄.

To ensure the large-scale application, four evaluation indicators were proposed for screening the most suitable precursors, including the cost, toxicity, and yield of the precursors and the removal rate for the contaminant. As shown in Tables 1 and 2, melamine exhibits the middle cost, toxicity, and removal ratio for the degradation of NO. However, it possesses the highest yield among the four precursors. Based on the advantages analysis previously mentioned, the best precursor is melamine.

4. Conclusions

In summary, g-C₃N₄ samples were prepared by a simple pyrolysis of thiourea, dicyandiamide, melamine, and urea under the same procedures, respectively. The systematic research results confirmed that the precursors have significant effect on the morphology, band gap, surface area, pore volume, thermal stability, and visible light photocatalytic activity of g-C₃N₄. Considering the cost, toxicity, and yield of the precursors and the properties of g-C₃N₄, the best precursor for preparation of g-C₃N₄ is melamine. The present work could provide new insights into appropriate selection of precursors for g-C₃N₄ synthesis as visible light photocatalyst in environmental protection.

Conflict of Interests

The authors declare that they have no conflict of interests.

Acknowledgments

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References


