COPPER MODIFIED CARBON NITRIDE AS FLUORESCENCE SENSOR FOR NITRATE IONS

(Karbon Nitrida Terubah Suai Kuprum Sebagai Sensor Pendafluor Ion Nitrat)

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Abstract

In this study, newly developed copper modified CN composites were prepared and tested as a fluorescence sensor for detection of nitrate ions (NO$_3^-$). The structure and chemical properties of CN and copper modified CN composites were investigated via X-ray diffraction (XRD), Fourier transform infra-red (FTIR), diffuse reflectance ultraviolet-visible (DR UV-Vis) and fluorescence spectrophotometry. Three emission sites represented as C=N, C=O and C-N moieties were suggested to contribute as sensing sites in CN and copper modified CN composites. The sensing capabilities of CN and copper modified CN composites toward NO$_3^-$ in the range of 300 to 1800 μM were determined via a quenching technique. The quenching efficiencies ($K_{SV}$) of CN and copper modified CN composites were obtained from the Stern-Volmer plot. Among three emission peaks of CN, C=N sites were found to be the most sensitive site having the strongest interaction with NO$_3^-$ ion. By addition of Cu(0.5 mol%), the $K_{SV}$ of CN was improved from 2.11 x 10$^{-4}$ to 5.27 x 10$^{-4}$ μM$^{-1}$. This study showed that with the addition of copper as modifier, the performance of CN can be improved and the composite can be used as potential fluorescence sensor for the detection of NO$_3^-$.

Keywords: copper modified carbon nitride, fluorescence sensor, nitrate ion

Abstrak

Dalam kajian ini, karbon nitrida terubah suai kuprum yang baru telah disediakan dan diuji sebagai sensor pendafluor untuk mengesan ion nitrat (NO$_3^-$). Struktur dan sifat kimia bagi CN telah disiasat melalui pembelahan sinar-X (XRD), transformasian infra-merah Fourier (FTIR), spektroskopi pantulan serakan ultralembayung-cahaya nampak (DR UV-Vis) dan spektroskopi pendafluor. Tiga tapak pelepasan diwakili sebagai moiti C=N, C=O dan C-N telah dicadangkan untuk menyumbang sebagai tapak penderiaan CN dan komposit CN terubah suai kuprum. Keupayaan penderiaan CN dan komposit CN terubah suai kuprum terhadap NO$_3^-$ dalam lingkungan 300 hingga 1800 μM telah ditentukan melalui teknik pelindapan. Kecekapan pelindapan ($K_{SV}$) bagi CN dan komposit CN terubah suai kuprum telah diperoleh daripada plot Stern-Volmer. Di antara tiga tapak pelepasan CN, tapak C=N telah ditemui menjadi tapak yang paling sensitif dan mempunyai interaksi paling kuat dengan NO$_3^-$ ion. Dengan penambahan Cu(0.5 mol%), nilai $K_{SV}$ bagi CN telah bertambah baik daripada 2.11 x 10$^{-4}$ kepada 5.27 x 10$^{-4}$ μM$^{-1}$. Kajian ini menunjukkan bahawa dengan penambahan kuprum sebagai pengubahsai, prestasi CN boleh ditingkatkan dan komposit tersebut boleh digunakan sebagai potensi sensor pendafluor untuk mengesan NO$_3^-$.

Kata kunci: karbon nitrida terubah suai kuprum, sensor pendafluor, ion nitrat
**Introduction**

Nitrate (NO$_3^-$) is a widespread contaminant of groundwater which can be easily detected in the environment and cause hazard to human health. The NO$_3^-$ is largely used for food preservative, drinking water and fertilizing agents [1, 2]. Excessive consuming of food and drinking water containing high level of NO$_3^-$ can cause severe risk to human health especially infants. Human may metabolize NO$_3^-$ in blood and finally transform it into carcinogenic nitrosamine causing the condition known as methemoglobinemia or also called as baby blue syndrome [3, 4]. In addition, it has been reported that the healthy human urine range of NO$_3^-$ is in the range of 300 to 1800 µM [5]. Therefore, determination of NO$_3^-$ is an important aspect to human health and environment. Griess reagent has been used widely for detection of NO$_3^-$ ion. Basically, Griess reaction involved several steps starting from reduction of NO$_3^-$ to NO$_2^-$ ion by using vanadium (III) chloride [6]. The mixture was subsequently reacted with sulfanilamide and N-(1-naphthyl) ethylenediamine to produce pink coloured azo dye product which strongly absorbed light at 540 nm [5, 7-9]. However, it presents several disadvantages such as using high concentration of hazardous reagent in the reaction, complicated detection procedure and cannot be reused [10]. Thus, development of reusable and sensitive sensor for detection of NO$_3^-$ ion is highly required.

Recently, carbon nitride (CN) has become a current interest because of its unique properties such as non-toxicity, low cost, high photoluminescence intensity [11] and good photostability [12]. CN is a graphite like layered material where s-triazine (C$_3$N$_3$) or tri-s-triazine (C$_9$N$_7$) units connected with amino groups (-NH/NH$_2$) are held together through hydrogen bonds between the amino groups [13, 14]. The principal application of the CN in the environment is mainly in photocatalysis for photodegradation of environmental pollutants [15, 16], while only few studies have been reported on its application in optical detection of environmental pollutant [17-22]. For instances, the CN has been reported to act as fluorescence sensor for detection of Cu$^{2+}$ and Fe$^{3+}$ ions [16-18] and N'-Nitrosopyrrolidine [19]. Recently, metal-free CN was used as a fluorescence chemical sensor for detection of NO$_3^-$ [23]. Unfortunately, CN alone still has low sensitivity towards the detection of the NO$_3^-$ ion. Therefore, further investigation is still required to improve the performance of the CN.

The objective of this study is to design a sensitive sensing material based CN by improving the sensing sites of the CN for detection of the NO$_3^-$ ion. In order to improve the sensing sites of the CN, copper species was used as the modifier. Copper based material has attracted a great interest due to its low cost as compared to noble metals such as Au, Pt, or Pd. It has been reported that the amino groups from the CN can interact with copper ion to form a stable complex due to the strong coordination force between the amino group and copper ion [24]. Herein, this study reported the modification of the CN by using copper species prepared by impregnation followed by a reduction method. The synthesized composites were demonstrated to show better sensing performances towards the NO$_3^-$ ion.

**Materials and Methods**

**Material preparation**

CN was prepared by using urea (CON$_2$H$_4$, Sigma-Aldrich, 98%) as precursor via thermal polymerization technique at 823 k for 4 hours [21, 25]. As for the preparation of composites, certain amount of Cu(acac)$_2$ (Cu(C$_3$H$_7$O$_2$)$_2$), Merck, 99.99%) was dissolved in ethanol and mixed with CN via an impregnation process. The mixture was heated up to 363 K until all the ethanol was evaporated. The obtained solid was underwent thermal hydrogenation process at 473 K for 2 hours under hydrogen flow of 5 ml/min to produce copper species modified CN composites. Composites were denoted as Cu(x)/CN composites where x showed the mol ratio of Cu to CN, which were fixed at 0.1 and 0.5 mol%. As a reference, the Cu(acac)$_2$ was heated under thermal hydrogenation process at 473 K or 2 hours to obtain Cu metal.

**Characterizations**

Properties of CN and Cu(x)/CN composites were investigated by several instruments. The structure of the CN and its composite was identified by using powder X-ray diffractometer recorded with a Bruker D8 Advanced using Cu Kα irradiation (λ = 1.5406 Å). The surface area of CN and its composites was confirmed by using Brunauer-Emmett-Teller (BET) equation obtained from a Quantachrome NOVA TOUCH LX$^4$. The functional groups present in the CN and its composites were recorded by using a Nicolet iS50 spectroscopy by mixing the samples with potassium bromide as a pellet. The diffuse reflectance UV-Visible (DR UV-Vis) spectra were recorded by using
Shimadzu UV-Vis spectrophotometry (UV-2600) in the range of 250 – 800 nm. The emission spectra were determined via JASCO FP-8500 fluorescence spectrophotometer at room temperature.

Quenching test
The interactions between the CN and its composites with NO$_3^-$ were studied via fluorescence spectroscopy at room temperature. The quenching tests were carried out by exposing various concentrations of NO$_3^-$ in the range of 300 to 1800 µM onto the CN and its composites. The various concentrations of NO$_3^-$ (10 µL) were introduced into the sample (0.05 g) and the emission spectra were measured at excitation wavelengths of 277, 320 and 370 nm. The interaction between the samples and the NO$_3^-$ can be observed from the changes in the emission intensity measured at each excitation wavelength and evaluated from the Stern-Volmer plot.

Results and Discussion
Characterization of CN and Cu(x)/CN composites
To investigate the chemical structure and optical properties of CN and its composites, XRD, FTIR, DR UV-Vis and fluorescence spectroscopies measurements were performed. The XRD patterns of CN and its composites are given in Figure 1. The XRD pattern of the prepared Cu metal is also shown as a reference. Cu metal gave three obvious peaks at 2θ of 44.3, 50.4 and 74.1° as the characteristics of copper (0) species (JCPDS no. 00-004-0836), which pattern was virtually the same as the reported one [26]. On the other hand, the CN exhibited two diffraction peaks at 2θ of 13.10 and 27.30°, corresponding to in-planar repeating units and the distance between the nitride pores, respectively [20, 23, 27-30]. The Cu(x)/CN composites showed similar diffraction patterns to those of the CN, suggesting that the added Cu might be too low to be detected by XRD or the Cu was highly dispersed on the CN.

![Figure 1. XRD patterns of Cu, CN and Cu(x)/CN composites](image)

The FTIR spectra of CN and the Cu(x)/CN composites are shown in Figure 2. The CN showed vibration bands at around 3300 – 3400 cm$^{-1}$ that can be attributed to the overlapping bands of primary and secondary amine, v(N-H) and v(O-H) groups [21-24, 31-35]. The stretching modes of v(C-N) heterocycles were observed at 1200 –1700 cm$^{-1}$, while the heterocyclic tri-s-triazine (C$_6$N$_7$) was found at around 809 cm$^{-1}$. Even though the presence of amino groups in the synthesized CN is difficult to be analysed by FTIR spectroscopy, it has been reported by XPS analysis [31]. It was obvious that the Cu(x)/CN composite exhibited almost similar peaks to those of the CN, indicating that the structure of CN was kept to be almost the same even after addition of copper species on the samples, similar to other reported literatures [24, 32].
Figure 2. FTIR spectra of CN and Cu(x)/CN composites

Figure 3 shows N$_2$ adsorption-desorption isotherms of CN and its composites. It was obvious that they exhibited type III isotherm, which referred to nonporous solids. There were no change occurred on the isotherm when the Cu species were added onto the samples, suggesting that the added copper species did not affect the physical properties of the CN. The BET specific surface areas of CN, Cu (0.1)/CN, and Cu (0.5)/CN were measured to be 91, 93 and 56 m$^2$/g, respectively. Addition of 0.1 mol% of Cu did not affect much the specific surface area, but the addition of 0.5 mol% of Cu significantly decreased the specific surface area. The decrease in the surface area has been also reported when Cu(acac)$_2$ was dispersed on the CN [32].

Figure 3. N$_2$ adsorption-desorption of (a) CN, (b) Cu(0.1)/CN and (c) Cu(0.5)/CN composites
The optical properties of CN and its composites were investigated by DR UV-Vis spectroscopy. As shown in Figure 4, CN and its composites exhibited three strong absorption bands at 277, 320 and 370 nm. The peak at 277 nm was corresponded to C=N groups (π-π*) in the conjugated aromatic triazine ring [20, 23, 31, 32], while the peak at 320 nm would be related to the presence of C=O moiety (π-π* and n-π*) due to the low condensation during the polymerization process of urea [30, 32, 36]. The peak at 370 nm was originated from the C-N terminal moiety (n-π*) [20, 23, 31, 32]. The Cu(x)/CN composites also gave similar absorption peaks to the CN, but with increased absorption peak intensities below 350 nm. The increase was more prominent for Cu(x)/CN composite with higher Cu loading amount. This would be the evidence of the presence of Cu on the samples since the peaks were contributed from the ligand to metal charge transfer (LMCT) transition between Cu and oxygen species in the composites [37].

The fluorescence properties of CN and Cu(x)/CN composites were studied by using fluorescence spectroscopy. Similar to the DR UV-Vis spectra, CN and its composites also shows three excitation peaks at 277, 320 and 370 nm, which referred to C=N, C=O and C-N terminal groups, respectively. On the other hand, only one emission peak was observed when CN and its composites were excited at 277, 320 or 370 nm [23]. As can be seen in Figure 5 (a), the intensity of the excitation spectra of the composites decreased with the increase in the loading amount of Cu species. This is believed due to certain interactions occurred between Cu species and CN. The same phenomenon also can be monitored from the emission spectra of CN and its composites at 455 nm for all excitation wavelengths, as can be seen in Figure 5 (b), (c) and (d), which illustrates the emission spectra of CN and its composites when excited at 277, 320 and 370 nm, respectively.

Figure 4. DR UV-Vis spectra of CN and Cu(x)/CN composites
Sensing performance of CN and Cu(x)/CN composites toward NO$_3^-$ ion

The fluorescence responses of CN and its composites to various concentrations of NO$_3^-$ (300-1800 µM) were studied by using a quenching technique. Since the CN and its composites have three emission sites, which are C=N, C=O and C-N terminal groups, it is important to determine which emission site has the strongest interaction to NO$_3^-$. Figure 6 shows the emission spectra of CN at 277, 320 and 370 nm before and after addition of various concentrations of NO$_3^-$. Based on the spectra, it is clearly seen that the intensity of CN decreased with the increase of the NO$_3^-$ concentration at all sensing sites, suggesting that the NO$_3^-$ ion interacted with C=N, C=O and C-N terminal sites and acted as a quenching agent to deactivate the emission sites of the CN. It is believed that there is an electrostatic interaction between the CN and its composites with NO$_3^-$ due to different charge between the sensing sites of the sensor samples and the NO$_3^-$ [38]. The same phenomena also can be observed for Cu(0.1)/CN and Cu(0.5)/CN samples, which spectra are shown in Figures 7 and 8, respectively.
Figure 6. Emission spectra of CN in the absence and presence of NO$_3^-$ with various concentrations, monitored at excitation wavelength of (a) 277, (b) 320 and (c) 370 nm.

Figure 7. Emission spectra of Cu (0.1)/CN in the absence and presence of NO$_3^-$ with various concentrations, monitored at excitation wavelength of (a) 277, (b) 320 and (c) 370 nm.
In order to determine the most sensitive sensing sites on the CN and its composites, quenching efficiencies at each sensing sites of the samples need to be calculated. The quenching efficiencies of the sensor can be determined by the relationship between the reduced intensity and the added concentration of NO$_3^-$, which is reflected by the Stern-Volmer equation:

$$I_0/I = K_{SV} [Q] + 1$$  \hspace{1cm} (1)

where $I_0$ is the fluorescence intensity of sensor in the absence of quencher (NO$_3^-$) and $I$ is the fluorescence intensity of sensor in the presence of quencher (NO$_3^-$), $K_{SV}$ is the Stern-Volmer constant which measures the efficiency of quenching and $[Q]$ is the concentration of NO$_3^-$.

Among the three sensing sites of the CN and its composites, which are C=N, C=O and C-N terminal groups, the C=N showed the strongest interaction between the samples and the NO$_3^-$ ion. The Stern-Volmer plots of the CN and Cu(x)/CN composites for the C=N sites monitored at excitation wavelength of 277 nm are shown in Figure 9. The CN gave almost linear plot to sense NO$_3^-$ ion up to 1800 µM. On the other hand, the Cu (0.1)/CN composite and the Cu(0.5)/CN composites gave a linear plot up to 1500 and 900 µM, respectively. When the added concentration was higher than these concentration levels, saturation was observed and this was more prominently observed on the Cu(0.5)/CN. The addition of Cu was found to give lower concentration range detection, which may be caused by the blocking of the sensing sites by the added Cu species. However, it is clear that the addition of Cu improved the quenching efficiency of the CN. The calculated $K_{SV}$ values of CN, Cu(0.1)/CN and Cu(0.5)/CN composites at excitation wavelength of 277 nm were 2.11 × 10$^{-4}$, 2.50 × 10$^{-4}$ and 5.27 × 10$^{-4}$ µM$^{-1}$, respectively, as summarized in Figure 10. Although detection range on the Cu(0.5)/CN composite was lower, obviously the Cu(0.5)/CN composite showed almost twofold higher $K_{SV}$ value as compared to the unmodified CN. It is revealed that the presence of copper species improved the performance of CN for the detection of NO$_3^-$ ion. Therefore, it can be concluded that Cu(x)/CN composites were potential to be used as a fluorescence sensor for detection of NO$_3^-$ ion.
Figure 9. Stern-Volmer plots between the relative emission intensity of CN and Cu(\(\chi\))/CN and the concentration of NO\(_3^−\), monitored at excitation wavelength of C=\(\equiv\)N sites (277 nm)

Figure 10. The \(K_{SV}\) values for CN and Cu(\(\chi\))/CN for all the emission sites

**Conclusion**

The CN was successfully synthesized via a thermal polymerization technique, while the Cu(\(\chi\))/CN composites were prepared by impregnation of copper species, followed by thermal reduction process. XRD patterns and FTIR spectra did not show any changes, suggesting that the structural properties of CN were not influenced by the addition of copper species onto CN. On the other hand, the DR UV-Vis spectra showed the presence of copper species in the composites in the area below 350 nm, which was due to the LMCT between the copper species and CN. The CN and Cu(\(\chi\))/CN composites exhibited three emission sites, which are C=\(\equiv\)N, C=O and C-N sites. Among all these emission sites, the C=\(\equiv\)N site showed the most favorable sensing site for detection of NO\(_3^−\) ion. The quenching efficiencies were represented by the \(K_{SV}\) values, which values for CN, Cu(0.1)/CN and Cu(0.5)/CN composites were \(2.11 \times 10^{-4}\), \(2.50 \times 10^{-4}\) and \(5.27 \times 10^{-4}\) \(\mu\)M\(^{-1}\), respectively. Modification with copper species (0.5 mol%) onto the CN improved the quenching efficiency up to two times higher than the unmodified CN. This work clearly demonstrated that the Cu(\(\chi\))/CN composites can be used as a potential fluorescence sensor for detection of NO\(_3^−\) ion.
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