Data Article

Adsorption and photocatalytic scavenging of 2-chlorophenol using carbon nitride-titania nanotubes based nanocomposite: Experimental data, kinetics and mechanism

M.A. Barakat a,b, Rajeev Kumar a,∗, Jamiu O Eniola a

a Department of Environmental Sciences, Faculty of Meteorology, Environment and Arid Land Agriculture, King Abdulaziz University, Jeddah 21589, Saudi Arabia
b Central Metallurgical R & D Institute, Helwan 11421, Cairo, Egypt

A R T I C L E   I N F O

Article history:
Received 2 November 2020
Revised 10 December 2020
Accepted 11 December 2020
Available online 16 December 2020

Keywords:
C3N4/TiO2 nanotube
2-Chlorophenol
Adsorption
Photocatalysis
Kinetics
Mechanism

A B S T R A C T

Adsorption and interaction of pollutant species on surface of the catalyst materials play an important role on the photocatalysis process. Herein, experimental data on the adsorption behavior of 2-chlorophenol (2-CP) onto graphitic pure carbon nitride (C3N4), titania nanotubes (TiO2–NTs) and carbon nitride/titania nanotubes nanocomposite (C3N4/TiO2–NTs) from synthetic wastewater has been summarized. The data on photocatalytic degradation of the 2-CP under both ultraviolet (UV) and visible light irradiation is also presented. This work also evaluates the 2-CP scavenging efficiency of C3N4/TiO2–NTs nanocomposite prepared by calcination of 2 wt.% melamine with TiO2–NTs at 450 °C. The adsorption and photocatalysis experiments were conducted for 180 min at pH 7 with 100 mL solution of 2-CP (40 mg/L) and 0.05 g catalyst material. The acquired data can be valuable to identify the equilibrium time for 2-CP adsorption onto C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite. Moreover, the obtained data can be useful to identify the suitable light source for the decomposition of 2-CP in the aquatic environment. The evaluated kinetic data might be significant for identifying the adsorption and photocatalysis
reaction rate onto the applied catalyst materials. The obtained adsorption and photocatalysis data have been compared with that in literature to identify the adsorption and photocatalysis behavior of 2-CP on numerous catalysts at different experimental conditions.

© 2020 The Author(s). Published by Elsevier Inc. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/)

### Specifications Table

| Subject | Environmental science, material science |
| Specific subject area | Wastewater purification, material synthesis, adsorption, photocatalysis, | |
| Type of data | Tables, Figures |
| How data were acquired | The amount of 2-chlorophenol in the aqueous solution before and after adsorption and photocatalysis was analyzed by HACH DR6000 UV–visible spectrophotometer. UV-visible diffuse reflectance spectra of the C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite were recorded on VARIAN Cary 500, USA. |
| Data format | Raw and analyzed |
| Parameters for data collection | At different times, the experimental data were obtained in the dark and under the UV and visible light illumination to analyze the adsorption and photocatalytic efficiency of the synthesized pure and hybrid material. The equilibrium attainment time for 2-CP adsorption was studied in the dark while the photocatalytic decomposition under UV (112 W) and visible light irradiation (104 W). Moreover, the efficiency of the C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite for 2-CP adsorption and photocatalytic degradation were compared. |
| Description of data collection | The data related to adsorption and photocatalysis was collected in the form of the concentration of the 2-CP. A certain amount of 2-CP solution was drawn every 30 min and filtered by a 0.22 μm membrane syringe filter. The adsorption and photocatalytic experiments were performed between 0 and 180 min. The solution pH 7 was kept constant during the whole adsorption and photocatalysis process. |
| Data source location | King Abdulaziz University, Jeddah, Saudi Arabia |
| Data accessibility | Raw data are provided with the article in a supplementary file. Mendeley Data under identification number: https://data.mendeley.com/datasets/zwkrtdg85b/3 |
| Related research article | M Anjum, R Kumar, SM Abdelbasir, MA Barakat. Carbon nitride/titania nanotubes composite for photocatalytic degradation of organics in water and sludge: Pre-treatment of sludge, anaerobic digestion and biogas production. Journal of environmental management 223, 2018, 495–502. https://doi.org/10.1016/j.jenvman.2018.06.043 |

### Value of the Data

- Data is valuable to develop new hybrid adsorbent and catalyst materials for the efficient removal of the contaminant from the wastewater.
- The adsorption data revealed that pure C3N4 is a better adsorbent than TiO2–NTs, and C3N4/TiO2–NTs nanocomposite for the removal of 2-CP.
- Kinetic data can be used to find the rate of 2-CP adsorption and photocatalytic degradation onto C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite.
- Data could be valuable to identify a suitable radiation source for photocatalytic applications.
- Data may be applicable to find the equilibrium time for the adsorption and photocatalysis of 2-CP onto C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite.
- Data could be used to identify the band gap energy, conduction band and valance band energy level of the C3N4, TiO2–NTs, and C3N4/TiO2–NTs nanocomposite.
1. Data Description

The data presented in the article explore the adsorption and photocatalytic properties of C$_3$N$_4$, TiO$_2$–NTs, and C$_3$N$_4$/TiO$_2$–NTs nanocomposite for scavenging of 2-CP from aqueous solution [1]. Adsorption plays a vital role in the photocatalysis process. It is assumed that good interaction between the pollutant species with the catalyst surface facilitates better photocatalytic decomposition [2-4]. Prior to starting the photocatalysis of the pollutant, adsorption was performed in the dark to identify the saturation of the catalyst and to determine the pollutant scavenging efficiency of the materials during adsorption and photocatalysis [5,6]. Adsorption kinetic analysis is important to find the rate of the 2-CP removal and to identify the nature of the process, i.e., chemisorption or physical sorption [7]. The liner plots and the adsorption kinetic parameters such as calculated adsorption capacity (q$_e$), values of the rate constant (k) and correlation coefficient (R$^2$) have been reported.

**Fig. 1** shows a schematic diagram for the synthesis of the C$_3$N$_4$/TiO$_2$–NTs nanocomposite in two steps. The adsorption proprieties of C$_3$N$_4$, TiO$_2$–NTs, and C$_3$N$_4$/TiO$_2$–NTs nanocomposite for 2-CP scavenging is shown in **Fig. 2**. The liner plots for adsorption equilibrium data fitted to the pseudo-first order and pseudo-second order kinetic models are shown in **Fig. 3** at 40 mg/L of 2-CP concentration. The values of the pseudo-first order and pseudo-second order kinetic parameters obtained from the liner plots in **Fig. 3** are included in **Table 1**.

**Fig. 4** shows the plot for the degradation of the 2-CP over C$_3$N$_4$ under UV (light intensity 112 W) and visible light irradiation (light intensity 104 W) for 180 min, and 2-CP concentration was 40 mg/L. **Fig. 5** illustrates the liner plots for the zero-order, first-order, and second-order

![Fig. 1. The schematic diagram for the synthesis of C$_3$N$_4$/TiO$_2$–NTs nanocomposite.](image)

**Table 1**
The values of kinetic parameters for 2-CP adsorption onto C$_3$N$_4$/TiO$_2$ NTs, C$_3$N$_4$ and TiO$_2$-NT.

|                | Pseudo-first order | Pseudo-first order Pseudo-second order |
|----------------|--------------------|---------------------------------------|
|                | qe$^{(exp)}$ (mg/g) | R$^2$ | K$_1$ (1/min) | qe$^{cal}$ | R$^2$ | K$_2$ (g/mg/min) | qe$^{cal}$ (mg/g) |
| C$_3$N$_4$/TiO$_2$ -NTs | 27.45              | 0.9581 | 0.018         | 36.54     | 0.8664 | 0.00011       | 53.48          |
| C$_3$N$_4$    | 34.62              | 0.914  | 0.002         | 57.59     | 0.9280 | 0.00018       | 53.76          |
| TiO$_2$-NTs  | 27.5               | 0.9567 | 0.020         | 34.7      | 0.9888 | 0.00030       | 40.0           |
Fig. 2. The adsorption of 2-CP onto C₃N₄, TiO₂–NTs, and g-C₃N₄/TiO₂–NTs nanocomposite.

Fig. 3. Plots for pseudo-first order and (a) pseudo-second order kinetic models for 2-CP adsorption onto C₃N₄, TiO₂–NTs, and g-C₃N₄/TiO₂–NTs nanocomposite.

Table 2
The values of kinetic parameters for the degradation of 2CP by C₃N₄ in the presence of UV and visible light.

| Kinetic models  | Parameters       | C₃N₄ (Visible light) | C₃N₄ (UV)   |
|-----------------|------------------|----------------------|-------------|
| Zero-order      | k₀ (mg/L min)    | 113.6 x 10⁻³         | 72.1 x 10⁻³ |
|                 | R²               | 0.9587               | 0.9562      |
| First-order     | k₁ (1/min)       | 6 x 10⁻³             | 2.1 x 10⁻³  |
|                 | R²               | 0.9902               | 0.9608      |
| Second-order    | k₂ (L/mg. min)   | 3 x 10⁻⁴             | 6 x 10⁻⁵    |
|                 | R²               | 0.9871               | 0.9633      |

kinetic models. The values of zero-order, first-order, and second-order kinetic models are mentioned in Table 2.

The UV-visible diffuse reflectance spectra of C₃N₄, TiO₂–NTs, and g-C₃N₄/TiO₂–NTs nanocomposite is shown in Fig. 6a. The band gap energy calculated using the Tauc plot is shown in Fig. 6b. A schematic diagram shown in Fig. 7 indicates the active radical species’ production for
Fig. 4. Photocatalytic degradation of 2-CP over C$_3$N$_4$ under UV and visible light irradiation.

Fig. 5. Kinetics plots of 2-CP degradation over C$_3$N$_4$ under UV and visible light irradiation (a) zero-order kinetic (b) first-order kinetic and (c) second-order kinetic models.
Fig. 6. (a) UV-visible diffuse reflectance spectra and (b) Tauc plot for band gap energy calculation of C\textsubscript{3}N\textsubscript{4}, TiO\textsubscript{2}–NTs, and C\textsubscript{3}N\textsubscript{4}/TiO\textsubscript{2}–NTs nanocomposite.

Fig. 7. The schematic diagram of the photocatalysis mechanism of 2-CP degradation by C\textsubscript{3}N\textsubscript{4}/TiO\textsubscript{2}–NTs nanocomposite.

Table 3
Comparison of the adsorption efficiency of 2CP by different materials.

| Material                  | Removal  | Experimental conditions                | Ref   |
|---------------------------|----------|---------------------------------------|-------|
| Amberlite XAD-16 resin    | 2.27 mmol/g | pH-6, conc. – 11.68 mmol/L          | [8]   |
| (TNTs/ACF)                | 59.9 (mg/g) | pH- 5, conc- 20 mg/L, time –15 min   | [9]   |
| ZnO/Clay                  | > 16%     | pH-7, conc- 20 mg/L, mass-0.2 g      | [10]  |
| Clay                      | > 20%     | pH-7, conc- 20 mg/L, mass-0.2 g      | [10]  |
| ZnO                       | > 4%      | pH-7, conc- 20 mg/L, mass-0.2 g      | [10]  |
| C\textsubscript{3}N\textsubscript{4}/TiO\textsubscript{2} –NTs | 27.45 mg/g | pH-7, conc- 40 mg/L, mass-0.05 g     | This work |
| TiO\textsubscript{2}–NTs  | 27.5 mg/g | pH-7, conc- 40 mg/L, mass-0.05 g     | This work |
| C\textsubscript{3}N\textsubscript{4} | 34.62 mg/g | pH-7, conc- 40 mg/L, mass-0.05 g     | This work |

the decomposition of 2-CP into the mineral by-products. A comparison of the adsorption capacity and the photocatalytic efficiencies of various adsorbents and catalysts for 2-CP scavenging are shown in Tables 3 and 4, respectively.
Table 4  
Comparison of photolysis and photocatalytic efficiencies of various materials used for the degradation of 2-CP.

| Material           | Degradation (%) | Experimental conditions | Ref    |
|--------------------|-----------------|-------------------------|--------|
| ZnO/Clay           | 88              | pH-8.7, conc. – 20 mg/L, mass- 0.2 g | [10]   |
| ZnO                | 61              | pH-8.7, conc. – 20 mg/L, mass- 0.2 g | [10]   |
| 5% Ag-doped TiO2   | 74              | pH –10.5, conc. – 50 mg/L, mass- 0.005 g | [11]   |
| 0.2% Ru/TiO2       | 53              | pH – 6, conc. – 100 mg/L, mass- 0.002 g | [12]   |
| Co-doped TiO2      | 93.4            | pH – 9, conc. – 50 mg/L, mass- 0.01 g, time- 3 h | [13]   |
| ZnO                | 55.6            | pH – 9, conc. – 25 mg/L, time – 3 h | [14]   |

| Photolysis         | 27              | Time – 10 h              | [15]   |
| Photolysis         | 17.05           | pH-7, conc. – 40 mg/L, time – 3.5 h, mass- 0.05 g | This work |
| C3N4 (UV)          | 25.02           | pH-7, conc. – 40 mg/L, mass- 0.05 g | This work |
| C3N4 (visible light)| 70.25          | pH-7, conc. – 40 mg/L, mass- 0.05 g | This work |

2. Materials and methods

2.1. Materials

The 2-chlorophenol (2-CP) used as a model pollutant was supplied by Merck, Pvt Ltd. The powdered TiO2 used for the synthesis of TiO2 NT was provided by (P-25 Degussa Co.). C3N4 was synthesized by from melamine which was obtained from Sigma Aldrich.

2.2. Synthesis

Herein, thermal methods were used to synthesize the C3N4, TiO2 NTs, and C3N4/TiO2–NTs nanocomposite, as previously reported [1]. The C3N4/TiO2–NTs nanocomposite used in this study was synthesized by calcination of 2 wt% of melamine with TiO2 NTs.

2.3. Adsorption and photocatalysis experiment

Adsorptive removal of 2-CP was investigated using the batch adsorption process, and all the photocatalytic experiments were conducted in the UV/visible light photochemical reactor (Luzchem, L2C 4V, Canada). The batch adsorption process and photocatalysis experiments were conducted in the 250 ml beaker containing the appropriate dose of catalyst/adsorbent (0.05 g) and 100 ml of 40 mg/L 2-CP solution. The pH of the 2-CP solution was adjusted to 7, using 0.1 M HCl or 0.1 M NaOH. The mixture containing catalyst was agitated at a speed of 200 rpm on a magnetic stirrer in the dark for adsorption experiments and in the presence of UV or visible light radiation and aeration for photocatalysis experiments. Samples were extracted at 30 mins intervals (30 – 180 min) using the pre-rinsed syringe and filtered through the 0.22 µm membrane filter. The concentration of 2-CP after adsorption and photocatalysis was analyzed using the UV-visible spectrophotometer (LANGE DR–6000, HACH, Germany) at a wavelength of 274 nm. The adsorption capacity and percentage of photocatalytic degradation were calculated using the following equation:

Adsorption capacity (qe) mg/g,

\[ q_e = (C_0 - C_e)V/m \]  \hspace{1cm} (1)

\[
\% \text{ degradation} = \frac{(C_0-C_e)}{C_0} \times 100 
\]  \hspace{1cm} (2)

Where, \( C_0, C \) and \( C_e \) represent the initial concentration, concentration at reaction time and concentration at adsorption equilibrium (mg/L), \( V \) is the volume (L) of 2-CP solution and \( m \) (g) is the weight of the adsorbent.
The adsorption kinetic study was performed by fitting the obtained time-dependent data to pseudo-first order and pseudo-second order. The linear equations are represented as follows:

Pseudo – first order kinetic : \( \log(q_e-q_t) = \log q_e - k_1 t/2.303 \)  \( (3) \)

Pseudo – second order kinetic : \( \frac{t}{q_t} = \frac{1}{k_2 q_e^2 + t/q_e} \)  \( (4) \)

where \( q_e \) and \( q_t \) represent the adsorption capacity (mg/g) at equilibrium and time \( t \) (min), respectively. \( k_1 \) is the pseudo-first order rate constant and \( k_2 \) (g/min) is the pseudo-second order rate constant.

The rate of 2-CP photocatalytic degradation by C\(_3\)N\(_4\) was also analyzed to investigate the photocatalytic behavior of the catalysts. The zero-order, first-order, second-order kinetic models were applied to analyze the experimental data at different contact times. The linear equations of zero-order, first-order, and second-order kinetic models are represented, respectively, as follows:

Zero – order kinetic : \( C = C_0 - k_0 t \)  \( (5) \)

First – order kinetic : \( \ln(C/C_0) = -k_1 t \)  \( (6) \)

Second – order kinetic : \( 1/C = (1/C_0) + k_2 t \)  \( (7) \)

where \( C_0 \) and \( C \) are 2-CP concentration at initial and reaction time \( t \) (min). \( k_0 \) (mg/L min), \( k_1 \) (1/min) and \( k_2 \) (L/mg.min) are zero-order, first-order and second-order rate constants, respectively.

**Declaration of Competing Interest**

The authors declare that there is no conflict of interest.

**Acknowledgment**

This project was funded by the Deanship of Scientific Research (DSR) at King Abdulaziz University, Jeddah, under the grant no G: 1291-155-1440. The authors, therefore, acknowledge with thanks DSR for technical and financial support.

**Supplementary Materials**

Supplementary material associated with this article can be found in the online version at doi: 10.1016/j.dib.2020.106664.

**References**

[1] M. Anjum, R. Kumar, S.M. Abdelbasir, M.A. Barakat, Carbon nitride/titania nanotubes composite for photocatalytic degradation of organics in water and sludge: pre-treatment of sludge, anaerobic digestion and biogas production, J. Environ. Manag. 223 (2018) 495–502.

[2] W. Zou, G. Gao, S.Y. Ok, L. Dong, Integrated adsorption and photocatalytic degradation of volatile organic compounds (VOCs) using carbon-based nanocomposites: a critical review, Chemosphere 218 (2019) 845–859.

[3] T. Fazal, A. Razaq, F. Javed, A. Hafeez, N. Rashid, S.U. Salma Amjad, S.M. Rehma, A. Faisal, F. Rehman, Integrating adsorption and photocatalysis: a cost effective strategy for textile wastewater treatment using hybrid biochar-TiO\(_2\) composite, J. Hazard. Mater. 390 (2020) 121623.

[4] N. Almoisheer, A. Alseroury, R. Kumar, T. Almeelbi, M.A. Barakat, Synthesis of graphene oxide/silica/carbon nanotubes composite for removal of dyes from wastewater, ESEV 3 (2019) 651–659.
[5] H. Atout, A. Bouguettoucha, D. Chebli, J.M. Gatica, H. Vidal, M.P. Yeste, A. Amrane, Integration of adsorption and photocatalytic degradation of methylene blue using TiO₂ supported on granular activated carbon, Arab. J. Sci. Eng. 42 (2017) 1475–1486.

[6] R. Kumar, M.A. Laskar, I.F. Hewaidy, M.A. Barakat, Modified adsorbents for removal of heavy metals from aqueous environment: a review, ESEV 3 (2019) 83–93.

[7] K.L. Tan, B. Hameed, H Insight into the adsorption kinetics models for the removal of contaminants from aqueous solutions, J. Taiwan Inst. Chem. Eng. 74 (2017) 25–48.

[8] K. Abburi, Adsorption of phenol and p-chlorophenol from their single and bisolute aqueous solutions on Amberlite XAD-16 resin, J. Hazard. Mater. B 105 (2003) 143–156.

[9] J. Duan, H. Ji, T. Xu, F. Pan, X. Liu, W. Liu, D. Zhao, Simultaneous adsorption of uranium(VI) and 2-chlorophenol by activated carbon fiber supported/modified titanate nanotubes (TNTs/ACF): effectiveness and synergistic effects, Chem. Eng. J. 406 (2021) 126752.

[10] A.H. Zyoud, S. Asaad, S.H. Zyoud, S.H. Zyoud, M.H. Helal, N. Qamhieh, A. Hajamohideen, H.S. Hilal, Raw clay supported ZnO nanoparticles in photodegradation of 2-chlorophenol under direct solar radiations, J. Environ. Chem. Eng. (2020) 104227.

[11] S.P. Onkani, P.N. Diagboya, F.M. Mtunzi, M.J. Klink, B.I. Olu-Owolabi, V. Pakade, Comparative study of the photocatalytic degradation of 2-chlorophenol under UV irradiation using pristine and Ag-doped species of TiO₂, ZnO and ZnS photocatalysts, J. Environ. Manag. 260 (2020) 110145.

[12] R.A. Elsalalomy, S.A. Mahmoud, Preparation of nanostructured ruthenium doped titania for the photocatalytic degradation of 2-chlorophenol under visible light, Arab. J. Chem. 10 (2017) 194–205.

[13] M.A. Barakat, H. Schaeffer, G. Hayes, S. Ismat-Shah, Photocatalytic degradation of 2-chlorophenol by Co-doped TiO₂ nanoparticles, Appl Catal. B Environ. 57 (2005) 23–30.

[14] A. Abdel Aal, M.A. Barakat, R.M. Mohamed, Electrophoreted Zn–TiO₂–ZnO nanocomposite coating films for photocatalytic degradation of 2-chlorophenol, Appl. Surf. Sci. 254 (2008) 4577–4583.

[15] I. Ilisz, A. Dombi, K. Mogyorósi, A. Farkas, I. Dékány, Removal of 2-chlorophenol from water by adsorption combined with TiO₂ photocatalysis, Appl. Catal. B Environ. 39 (2002) 247–256.