PAPER • OPEN ACCESS

Photocatalytic removal of eosin dye from aqueous solution over titanium dioxide

To cite this article: Hazim Y Al-Gubury et al 2020 IOP Conf. Ser.: Mater. Sci. Eng. 871 012031

View the article online for updates and enhancements.

The First International Conference of Pure and Engineering Sciences (ICPES2020)

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031

Photocatalytic removal of eosin dye from aqueous solution over titanium dioxide

Hazim Y Al-Gubury*, Suad T Saad, Nour Abd Alrazzak and Ruqaia M Naif

Department of Chemistry, College of Science for Women, University of Babylon, Hilla, Iraq

suad_saad80@yahoo.com

Abstract. A Various concentrations of Eosin dye were irradiated using titanium dioxide as a catalyst. The irradiation was carried out using the catalyst (0.1gm/100ml), mercury lamp 125 Watts from external source and at Rt. The effect of TiO_2 on the photocatalytic degradation of Eosin was studied in various conditions such as, studying the effect of loaded mass of titanium dioxide, effect of eosin dye concentration and effect of inorganic anions.

1. Introduction

Environment including water, air and soil is subjected to threat by different kinds of pollutants where these pollutants affect the human life [1-2]. Water pollution is one of the serious problems which has received more attention [3]. Many reasons are contributed to this problem and one of them is dyes [4]. Dyes have received more attention due to their toxicity, strong color and stability [5,6]. However, they have been used in industry for coloring for example, in textiles industry and as a result of this use the discharged water is contaminated with these dyes [7-8].

Different techniques have been used to treat this problem and one of them is using the photocatalytic degradation [4,9]. This type of degradation can be achieved using various metal oxides (semiconductors) as catalysts and one of them is titanium dioxide [10,11]. TiO₂ has been extensively used in photocatalytic degradation because it's efficient, cheap, photocatalytic active, and non- toxic [12-14].

In addition, the work of this kind of degradation relies on the oxidation process which leads to the formation of the reactive free radicals [15-17]. These radicals include hydroxyl and superoxide which resulted from the irradiation of the catalyst in the presence of water and oxygen and as a result this leads to the oxidization and degradation of these dyes [18].

In this paper TiO_2 was used as a catalyst to degrade Eosin 'Figure 1'. Different parameters have been measured to study the degradation efficiency of Eosin.



Figure 1. Chemical Structure of Eosin Dye

2. Materials and Methods

2.1 Chemicals

TiO₂ was supplied by Fluka AG and Eosin was supplied by sigma – Aldrich.

The First International Conference of Pure and Engineering Sciences (ICPES2020)IOP PublishingIOP Conf. Series: Materials Science and Engineering 871 (2020) 012031doi:10.1088/1757-899X/871/1/012031

2.2. Photocatalytic degradation processes of Eosin dye using TiO2:

The degradation process of Eosin has been investigated using the photo reactor. This reactor consisted of two parts. The first part has been used for cooling the Eosin solution using the running water which passed through it. The second part was the reaction solution vessel with (100 ml) capacity. The removal of the studied dye has carried out using mercury lamp (125 W). All experiments were carried out by mixing (0.1 gm) of the catalyst with (10 mg/L) of Eosin solution. The dye suspended solution was kept under stirring for (20 min) and was bubbled with air (10 ml/min) during the irradiation process. A (3 ml) of the reaction mixture was withdrawn every (10 min) and then centrifuged at (4000 rpm) to remove the catalyst. All samples absorption was measured by UV-Vis spectrophotometer [19,20].

3. Result and Discussion

3.1 Effect of Titanium Dioxide Mass on the Photocatalytic Degradation of Eosin

To study the effect of TiO_2 on the degradation of Eosin, (10 ppm) of this dye was used with (10ml/min) flow rate of air, and at (298 K) RT. Figure 2 represents photocatalytic degradation processes of Eosin at different loaded masses of titanium dioxide. Photocatalytic degradation of Eosin gradually increased as the mass of TiO_2 increased until it reached (0.1gm /100 ml), then gradually decreased. Because at the higher dosage of the catalyst, the light will scatter and only the solution layers located on the top will receive the light [21,22]. Furthermore, when the mass of TiO_2 is below (0.1gm /100ml) the photo degradation of Eosin also decreased. Because the surface area decreased and that gave less light absorption by the catalyst that led to less degradation of Eosin.





Figure 3 shows the kinetic analysis for TiO₂. The graph of $\ln (A_0 / A_t)$ vs. time is a straight line through the origin. Therefore, the rate of photodegradation follows a first order law and the slope gives k (min⁻¹) [23].

The First International Conference of Pure and Engineering Sciences (ICPES2020)

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031



Figure 3. Relationship between $ln(A_o/A_t)$ and irradiation time on different dosages of titanium dioxide

While, Figure 4 shows the relationship between the rate constant of the reaction and the loaded masses of the catalyst. It was found that the reaction rate constant increased when the loaded mass of the catalyst was increased until the reaction reached the optimum condition (0.1 gm/100 ml) [24].



Figure 4. Relationship between rate constant and dosage of TiO_2

3.2 Effect of Eosin concentration on its photocatalytic degradation

Eosin with different concentrations (10 - 50) ppm were irradiated using TiO₂ (0.1gm / 100 ml), with the light intensity (8.22 mW/cm²) and at room temperature. 'Figure 5' shows that the rate of the photocatalytic degradation gradually decreased when the Eosin concentration increased. Also, it shows that the best concentration of the dye was (10ppm). At this concentration a largest area of TiO₂ will be saturated with Eosin and this resulted in the absorption of more exciting photons. Moreover, the presence of excess dye will prevent the light to penetrate through the suspended solution, and this leads to decrease the photocatalytic degradation of Eosin on the catalyst surface [25, 26].

The First International Conference of Pure and Engineering Sciences (ICPES2020)

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031



Figure 5. Relation between the change in (A / A0) with irradiation time using various concentrations of Eosin

Figure 6 shows the kinetic analysis for TiO_2 . The graph of $ln (A_0 / A_t)$ vs. time is a straight line. This figure also shows the photo degradation of Eosin dye which is in good agreement with the Langmuir-Hinshelwood model [27-29].



Figure 6. Relationship between $\ln (A_o/A_t)$ and irradiation time on different concentration of Eosin dye

The high photo degradation efficiency (83.54%) was achieved when the Eosin concentration was (10 ppm). Figure 7 illustrates the photocatalytic degradation efficiency (P.D.E) with different concentrations of Eosin.

The First International Conference of Pure and Engineering Sciences (ICPES2020) **IOP** Publishing

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031



Figure 7. P.D.E of Eosin at various concentrations of the dye

4. Role of inorganic anions in Eosin photocatalytic degradation

Sodium chloride was used to study the effect of inorganic ions on the degradation of Eosin. Without any additions of this salt the degradation rate of Eosin was (91.77%) while, the rate decreased to (72.47%) when chloride ions were added (10 ml). This decrease in the photocatalytic activity is due to the scavenging of •OH radicals by these ions [30]. 'Figures .8 and 9'. show respectively the effect of the inorganic ions on Eosin degradation and the photodegradation efficiency of the dye using various concentrations of these ions.



Figure 8. Effect of NaCl on the photocatalytic degradation of Eosin

 The First International Conference of Pure and Engineering Sciences (ICPES2020)
 IOP Publishing

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031



Figure 9. Photodegradation efficiency of Eosin with different concentrations of NaCl

5. Conclusion

The photocatalytic degradation process of Eosin depended on the mass of titanium dioxide where the optimum value was (0.1 gm/100 ml⁾. In addition, the photocatalytic destruction of Eosin obeyed the pseudo-first-order reaction where, the degradation process decreased. This dropping is accompanied with increasing the concentration of Eosin which due to the decrease of the concentration of OH^- which adsorbed on TiO_2 surface and the dye concentration was (10 ppm) as an optimum value. Furthermore, the degradation process of the dye increased with the increase of light intensity because of the increase of photoelectron in the conduction band and this led to an increase in the electron-hole pairs number and a decrease in recombination process. The percentage of dye degradation was (91.77%) and (72.47%) in the absence and the presence of the inorganic ions respectively.

Acknowledgement

Sincerely we would like to thank the University of Babylon, College of Science for Women, Department of Chemistry for providing the necessary infrastructural facilities during the current study.

References

- [1] Mahmoud A M , Poncherib A, Badrc Y and Abd El Waheda M G 2009 S. Afr. J. Sci. 105 299-303.
- [2] Kumar A and Pandey J 2017 IJMSE 1 1-10.
- [3] Mai DF, Chen CC, Chen L J, Liu C S 2008 J. Chromatogr. A, 1189 355-365.
- [4] Fosso-Kankeu E, Waanders F and Geldenhuys M 20157th International Conference on Latest Trends in Engineering & Technology 84-89.
- [5] Reza M K, Kurny A, Gulshan F 2017 Appl Water Sci. 7 1569-78.
- [6] Šíma J and Hasal P 2013 Chem. Eng. Trans. 23 79-84.
- [7] Tahir H, Hammed U, Jahanzeb Q and Sultan M, 2008 Afr. J. Biotechnol. 7 3906-11.
- [8] Hilal N 2011 Der Chemica Sinica 2 262-273.
- [9] AMETA R, PUNJABI B P and AMETA CS 2011 J. Serb. Chem. Soc. 76 1049-55.
- [10] Sacco O, Stoller M, Vaiano V, Ciambelli P, Chianese A and Sannino D 2012 INT J PHOTOENERGY 2012 1-8.
- [11] Wang C, Wang X, Xu B, Zhao J, Mai B, Peng P, Sheng G and Fu J 2004 J. photochem photobiol A 168 47-52.
- [12] Mukhlish B Z M, Najnin F, Rahman M M, Uddin J M 2013 J. Sci. Res. 5 301-314.
- [13] Al-Shamali S S 2013 Aust. J. Basic & Appl. Sci., 7, 172-176.
- [14] Mesheshaa S D, Tirukkovalluria R S, Chandraa R M, Bojjab S 2016 IJER 6 110-119.
- [15] Brahmia O 2016 IJACEBS 3 225-227.
- [16] Karim A S, Al-Gubury Y H and Abd Alrazzak N 2019 J. Phys. : Conf. Ser. 1294 1-7.
- [17] Silva C L, Barrocas B, Melo Jorge E M and Sério S 2018 In proceedings of the 6th International Conference on Photonics, Optics and Laser Technology (PHOTOPTICS) 334-340.

The First International Conference of Pure and Engineering Sciences (ICPES2020) IOP Publishing

IOP Conf. Series: Materials Science and Engineering 871 (2020) 012031 doi:10.1088/1757-899X/871/1/012031

- [18] Chen X, Wu Z, Liu D and Gao Z 2017 Nanoscale Res. Lett. 12 1-10.
- [19] SAAD T S, AL-GUBURY Y H and ABD ALRAZZAK N 2018 AJC 30 2334-36.
- [20] Saad T S, Abd Alrazzak N, Al-Gubury Y H and Essa E 2019 IOP Conf. Ser: Mat. Sci. Eng. 571 1-5.
- [21] Qin L, Liu M, Wu Y, Xu Z, GuoX and Zhang G 2016 Appl. Catal. B 194 50-60.
- [22] Al-Gubury Y H, Hassan F A, Alteemi S H, Alqaragully B M, Bennecer A and Alkaim F A 2017 JGPT 12 290-295.
- [23] Algubury Y H 2016 Malaysian Journal of Science 35 319-330.
- [24] Rattan K V, Purai A, Singh H and Manoochehri M 2008 Carbon Lett. 9 1-7.
- [25] Pei CC and Leung F WW 2013 Sep. Purif. Technol. 114 108-116.
- [26] Subramonian W, Wu Y T and Chai P S 2017 J. Environ. Manage. 187 298-310.
- [27] Purnawan C, Wahyuningsih S and Kusuma P P 2016 Indones. J. Chem. 16 347-352.
- [28] Patil N P, Bote D S and Gogate R P 2014 Ultrason. Sonochem. 21 1770-77.
- [29] Gosu V, Gurjar R B, Surampalli Y R and Zhang CT 2014 J. Environ. Chem. Eng. 2 1996-04.
- [30] Al-Gubury Y H, Alteemi S H, Saad M A and Al-Shamary R R 2019 Indones. J. Chem 19 292-297.