Poll Res. 42 (1) : 43-47 (2023) Copyright © EM International ISSN 0257–8050 DOI No.: http://doi.org/10.53550/PR.2023.v42i01.008

USE OF BI₂S₃ AS A PHOTOCATALYST FOR PHOTOCATALYTIC DEGRADATION OF TEXTILE AZO DYE REACTIVE BLUE 160 (RB160)

DIMPLE SHARMA¹ AND K.S. MEENA²

Department of Chemistry, M.L.V. Government College Bhilwara 311 001, Rajasthan, India

(Received 13 July, 2022; Accepted 21 September, 2022)

ABSTARCT

In apparel industries removal of the color of dye is one of the main problem. In the current work semiconductor photo-catalyst Bi_2S_3 utilized in photo catalytic decolorizing of textile azo dye Reactive Blue 160 under UV light. Spectrophotometric measurements of the reaction rate were made. It has been investigated how many variables, including dye concentration, photocatalyst quantity, and pH, affect reaction efficiency.

KEY WORDS: Reactive blue, Bi₂S₃, Photo-catalytic degradation

INTRODUCTION

Water is the most valuable material on the earth, without it life will not exit. It is crucial to avoid polluting these priceless resources due to the lack and pollution of water supplies. Textile industries produce enormous quantities of wastewater containing colourful dyes that are both poisonous and non-biodegradable (Reife, 1996). Synthetic dyes are often used nowadays in several products, such as textiles, wood, leather, and plastic. However, roughly 12 percent of these colours are lost in the process of dyeing, and about 20 percent of this waste ends up in the environment. (Rafi *et al.*, 2021). Dyes which are bright in color Water-soluble colours, such as reactive and acid dyes, are the most difficult to remove.

As a result of the accidental release of toxic wastewater a pollution problem arises in worldwide textile industries which have a big impact on the quality of water supplies. According to the World Bank report 17 to 20 percent of industrial water contamination rises from textile dyeing and treatment. Therefore, this is a significant environmental concern for the textile industry. There is a demand for ecologically acceptable technology to remove colours from industrial and municipal effluent, as environmental consciousness increases.

Classical ways to clean up dirty water that are

still used today are adsorption, (Nasuha, 2010; Nagawe, 2018), coagulation (Riera, 2010), biodegradation (Xu *et al.*, 2011) chlorination (Ge, 2008), ion flotation (Shakir, 2010), sedimentation (Zodi, 2010), membrane process (Jirankova, 2010) and solvent extraction (Juang, 2008). However, physical and biological treatments do not eliminate contaminants; they merely change their phase. The disadvantage of chemical approaches is that they utilise strong oxidants such as chlorine and ozone, which are themselves pollutants (Algarni *et al.*, 2022) Complete purification of these processes' final products requires further processing. Dye photodecolorization is a potential technique for treating of industrial effluent.

Due to the cheap cost of the procedure, the absence of secondary pollution, and the approach's environmental friendliness, it is a viable option. In recent years, the advanced oxidation method of photo-catalysis has attracted a significant amount of focus on the process of water purification. To make sure the environment is safe and clean, it is important to separate and break down these organic wastes. Photo-catalysis, the removal of colours from water using an advanced oxidation process (AOP) is now being done without the creation of any dangerous by-products (Iqbal *et al.*, 2018). AOPs are based on the production of extremely reactive species like hydroxyl radicals (•OH), which are second only to fluorine in terms of their substantial oxidation potential [E0 = + 2.80 V]. Numerous organic contaminants are quickly and non-selectively oxidised by hydroxyl radicals. (Daneshwar *et al.*, 2003). And it can be improved further with the use of UV-visible radiation that produces extra hydroxyl radicals.

Photocatalytic degradation of direct black155 and Reactive Blue 160 dye were examined by using UV light in photochemical reactor with photocatalyst SnO_2 (Meena and Meena, 2021). Under UV light, zinc oxide also takes part in the photo-catalytic degradation of Reactive Red 152. (Meena and Dadheech, 2019). In the presence of the heterogeneous photocatalyst CeFeO₃, the azo dye Reactive Blue 160 underwent photocatalytic degradation. (Pamecha *et al*, 2016). $SnO_2/Bi_2S_3/BiOC1/Bi_{24}O_{31}Cl_{10}$ Composites' Easy Synthesis Significantly Increases Photocatalytic Degradation of Rhodamine B under Visible Light (Fenelon *et al.*, 2020).

The usage of a paint-coated substrate for wastewater remediation through the photocatalysis experiment using visible light irradiation was both economical as well as free from separation and filtration processes to recover the catalyst. The $BiVO_4^-$ paint composite coated on an aluminium substrate was used for photocatalytic MB dye degradation (Kumar and Vaish, 2022).

Using chalcogens as photocatalysts for degradation has grabbed the interest of the scientific community in recent years. Due to its cost in comparison to TiO_2 and ZnO and its low band gap energy, which ranges from 1.3 to 1.7 eV, bismuth sulphide is an attractive material. (Zhu, 2017).

MATERIALS AND METHODS

Reactive Blue 160 was chosen for the current photocatalytic degradation investigations. This is a diazo dye with sulphonate groups. It was acquired from Bhilwara Textile Industries (Raj). The Structural formula of the dye is $C_{38}H_{23}C_{12}N_{14}Na_5O_{18}S_5$ and Molecular mass is 1309.86gm. The photocatalyst Bismuth sulphide (Bi₂S₃) was used as a photocatalyst. Bi₂S₃ is a semiconductor that has a band gap that is on the narrower side. It has been shown to be an exceptional light absorption material, which finds widespread use in photonic devices. The laboratory's reagents were all of analytical quality.

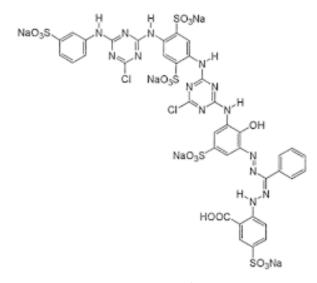


Fig. 1. Reactive Blue 160.

Bi₂S₃ was used as a photocatalyst while RB 160 degradation was examined at various pH levels, catalyst loadings, and dye concentrations. To make a dye solution that has a concentration of 1X10⁻³M, 1.309 grammes of dye were dissolved in a volume of 1000 millilitres of distilled water. In order to determine the initial absorbance of the dye solution, a UV-VIS spectrophotometer was used. At 560nm (max), the highest absorption value was noted. The reaction mixture was made up of three millilitres of Reactive Blue 160 solution (1x10⁻³ M) and 0.3 grammes of bismuth sulphide. The amount of the reaction mixture was brought up to 100 ml by the addition of double-distilled water. The reaction mixture included 3x10⁻⁵M dye molecules. (Dadheech, 2021) The reaction mixture was exposed to UV radiation using a photochemical reactor to do the photobleaching. Using a pH metre, the solution's pH was determined (Systronics 106). At certain time intervals, a spectrophotometer set at 560 nm (max) was used to measure absorbance in order to track the reaction's progress. The rate at which the colour faded over time was also continuously examined.

RESULTS AND DISCUSSION

At 554 nanometers, the photocatalytic degradation of RB160 was observed. The best conditions for dye photooxidation were dye concentration = 3×10^{-5} , pH = 8, and catalyst quantity = 0.3g/100 ml. Fig. 2 displays visually the effect of RB160's photodegradation.

Time (min)	abs	1+log abs
0	0.664	0.822
10	0.561	0.749
20	0.478	0.679
30	0.402	0.605
40	0.341	0.533
50	0.287	0.458
60	0.242	0.385
70	0.205	0.313
80	0.172	0.238
90	0.145	0.164

Table 1. Depicts a common run of photocatalyticdegradation of Reactive Blue 160.

Based on the theory of first order reaction, the straight line graph of 1+logA vs time. This phrase k=2.303xSlop was used to figure out the rate constant.

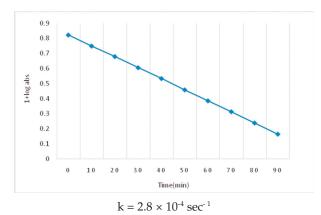


Fig. 2. Photocatalytic degradation of Reactive Blue 160

Effect of parameters

Variations in pH have the following effects

Variations in pH have an effect on the photocatalytic degradation of RB160. The pH has a significant impact in the formation of hydroxyl radicals.

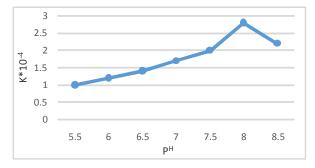


Fig. 3. A curve demonstrating the impact of changing hydrogen ion concentration on the speed at which the dye degrades.

Photocatalytic degradation was carried out at pH levels ranging from 3 to 9, while all other parameters remained constant. It was discovered that when pH increases, so does the rate of reaction. The rate of degradation was greatest at pH 8.0.

Variation in dye concentration has the following effects

The impact of dye concentration variation on photodegradation was studied for Bi_2S_3 with varying concentration of RB160 from 1.0×10^{-5} to 5×10^{-5} M, while all other variables were held constant. Up to a concentration of 3×10^{-5} M for Bi_2S_3 , it was shown that the degradation rate increases with increasing dye concentration. The surface of the dye catalyst becomes saturated as the dye concentration rises. As a consequence, the rate of decolorization slows.

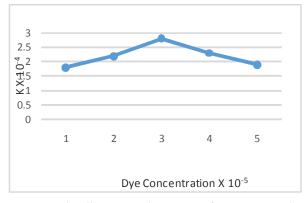


Fig. 4. A plot illustrating the impact of variations in dye concentration on the rate of dye decolorization.

Changing the Catalyst Concentration and Its Effects

By changing the quantity of Bi₂S₂ while keeping all other components constant, the influence of photocatalyst amount on the photo-decolorization of Reactive Blue 160 was studied. It was found that when catalyst amount is increased, the rate of degradation for Bi_2S_3 rises to 0.3g/100ml. The findings are shown in Fig. 5. Following an increase in catalyst concentration, the rate of reaction either decreases or remains almost constant. A possible explanation for this behaviour is that when the quantity of catalyst is increased, the surface area of the catalyst will expand and include more active sites. Because there are no longer any substrate dye molecules accessible for adsorption on semiconductor active sites beyond a certain quantity of catalyst, the rate of reaction decreases.

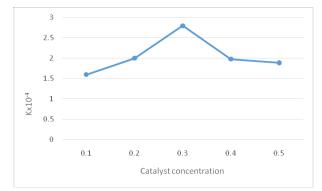


Fig. 5. A curve indicating how changes in catalyst dosage affect the dye decolorization pace.

Mechanism

The first step in the process of photocatalysis is the absorption of photons with an energy that is equal to or higher thanke semiconductor Bi_2S_3 band gap. The conduction band is reached by the electrons after they pass through the valence band. As a direct result of this, a hole, denoted by the symbol h+, is produced in the valence band. The photogenerated hole acts as a potent oxidising agent, while the conduction band electron performs the function of a reducing agent in a semiconductor.

 $\operatorname{Bi}_2 S_3 + hv \rightarrow \operatorname{Bi}_2 S_3^*$ $h^+ (vb) + e^- (cb)$

These pairs of electrons and holes move to the surface of the catalyst, which is the location where radicals are formed.

Hole (h^+) + OH \rightarrow OH Hole (h^+) + H₂O \rightarrow OH \rightarrow H $^+$ e⁻ + O₂ (ads) \rightarrow O₂ \rightarrow O₂ \rightarrow O⁻ (hydroxyl radical)

Superoxide radicals and peroxide radicals are powerful oxidising species. These radicals reacts with dye molecules to turn them into oxidised molecules. At the same time, a dye molecule absorbs the right amount of radiation and moves from its initial singlet state to its triplet state through a "intersystem crossing."

¹Dye_o + $hv \rightarrow$ ¹Dye₁ (single state) ISC ¹Dye₁ \rightarrow ³Dye₁ (triple state) ³Dye_o + radicals \rightarrow colourless end products

Propane-2-ol was used as a scavenger to verify the participation of the hydroxyl radical OH[•]. The pace of the reaction was significantly slowed down when 2-propanol was included in the mixture. The fact that degraded reaction mixture contains carbon dioxide (CO_2) , nitrate ion (NO_3^-) , and nitrite ion (NO_2^-) shows that this process is fully mineralized. The final goods are environmentally safe.

CONCLUSION

The photocatalyst Bi₂S₃ is utilised to degrade the dye Reactive Blue 160. According to the experimental findings, pH, dye concentration, and catalyst quantity all had an impact on the photodegradation of RB160. With increasing photocatalyst dose and initial azo dye concentration up to a certain point, the rate of dye photodegradation increased. Degradation is more likely to occur in basic situation. The ideal photodegradation conditions were found to be pH 8, 0.3gBi₂S₂/100ml, and 3 X 10⁻ ⁵M dye concentration. On Bi₂S₃, dyes degrade using first-order kinetics. The predominant reactive species in the degradation, according to experiments on radical scavenging, is the hydroxyl radical. Overall results indicate that this photocatalyst should be used to treat industrial effluent wastewater.

ACKNOWLEDGEMENT

The authors are grateful to the Principal and Head of the Department of Chemistry M. L. V. Government College Bhilwara for providing necessary facilities.

REFERENCES

- Algarani, T., Abduh, N., Kahtani, A. and Aouissi, A. 2022. Photocatalytic degradation of some dyes under solar light irradiation using ZnO nanoparticles synthesized from Rosmarinus officinalis extract. *Green Chemistry Letters and Review.* 15(2) : 460-473.
- Dadheech, A. and Kabra, B. 2021. Photocatalytic degradation of dye reactive blue 160 (RB160) in the presence of strontium chromate. *Poll Res.* 40 (1) : 93-96.
- Daneshwar, N., Salari, D. and Khatee, A.R. 2003. Photocatalytic degradation of azo dye acid red 14 in water:investigation of the effect of operational parameters. *J. Photochem. Photobiol. A: Chem.* 157 : 111-116.
- Fenelon, E., Bui, D., Tran, H., You, S., Wang, Y., Cao, T. and Pham, V. 2020. Straightforward Synthesis of SnO₂/Bi₂S₃/BiOCI-Bi₂₄O₃₁CI₁₀ Composites for Drastically Enhancing Rhodamine B Photocatalytic Degradation under Visible Light. ACS Omega. 32 : 20438-20449.

USE OF $\mathrm{BI}_2\mathrm{S}_3\mathrm{AS}$ A PHOTOCATALYST FOR PHOTOCATALYTIC DEGRADATION OF TEXTILE AZO ~~47

- Ge, F., Zhu, L. and Wang, J. 2008. Distribution of chlorination products of phenols under various pHs in water disinfection. *Desalination*. 225(1-3) : 156-166.
- Iqbal, M. 2018. Photocatalytic degradation of organic pollutant with nanosized cadmium sulphide. *Mater. Sci. Energy Technol.* 2(1): 41-45.
- Jirankova, H., Mrazek, J, Dolecek, P. and Cak, J. 2010. Organic dye removal by combined adsorption membrane separation process. *Desalination Water Treat.* 20 : 96-101.
- Juang, R., Huang, W. and Hsu, Y. 2008. Treatment of phenol in synthetic saline wastewater bysolvent extraction and two-phase membrane biodegradation. *J. Hazard. Mater.* 164 (1): 46-52.
- Kumar M. and Vaish R. 2022. Photocatalytic dye degradation using BiVO₄-paint composite coatings. *Mater Adv.* 3 : 5796-5806.
- Meena, K.S. and Meena, K. 2021. Photochemical bleaching of textile dye direct black 155 and Reactive Blue 160 by photocatalyst SnO_2 . *Ecology Research.* 1 : 48-55.
- Meena, K. S. and Dadheech, A. 2019. Photocatalytic degradation of textile dyes Methylene blue and Reactive Blue 160 by ZnO. *Poll. Res.* 38(1): 223-227.
- Nagawe, T., Rai, A.K., Ameta, R., Ameta, S.C.2018. Adsorption study for removal of crystal violet dye using MMT-MWCNTs composite from aqueous solution. J. Applicable Chem. 7(5) : 1252-1259.
- Nasuha, N., Hameed, B.H. and Din, A.T.M. 2010. Rejected tea as a potential low-cost adsorbent for the removal of methylene blue. *J. Hazard. Mate.* 175 : 126-132.
- Pamecha, K., Mehta, V. and Kabra, B.V. 2016.

Photocatalytic Degradation of commercial textile Azo Dye Reactive Blue160 by Heterogenous Photocatalysis. *Archives of Applied Science Research.*8(1):7-12.

- Rafiq, A. 2021. Photocatalytic degradation of dyes using semiconductor photocatalyst to clean industrial pollution. *Journal of Industrial and Engineering Chemistry*. 97 : 111-128.
- Reife, A., and Fremann, H.S.1996. *Enviornmental Chemistry of Dyes and Pigments.* ISBN: 978-0-471-58927-3
- Riera, T., Gutierrez-Bouzán, C. and Crespi, M. 2010. Combination of coagulation-flocculation and nanofiltration techniques for dye removal and water reuse in textile effluents. *Desalination*. 252 : 53-59.
- Shakir, K., Elkafrawy, A., Ghoneimy, H., Elrab Beheir, S. and Refaat, M. 2010. Removal of rhodamine B (a basic dye) and thoron (an acidic dye) fromdilute aqueous solutions and wastewater simulants by ion flotation. *Water Res.* 44 : 1449-1461.
- Xu, A., Li, X., Ye, S., Yin, G. and Zeng, Q. 2011. Catalyzed Oxidative Degradation of Methylene Blue by in situ Generated Cobalt (II)-Bicarbonate Complexes with Hydrogen Peroxide. *Appl.Catal., B.* 102 (1-2) : 37-43.
- Zhu, H., Jiang, R., Li, J., Fu, Y., Jiang, S. and Yao, J. 2017. Magnetically recyclable Fe₃O₄/Bi₂S₃ microspheres for effective removal of Congo red dye by simultaneous adsorption and photocatalytic regeneration. *Seperation and Purification Technology*. 179 : 184-193.
- Zodi, S., Potier, O., Lapicque, F. and Leclerc, J.P. 2010. Treatment of the industrial wastewaters byelectrocoagulation: Optimization of coupled electrochemical and sedimentation processes. *Desalination*. 261: 186-190.