# PAPER DETAILS

TITLE: Fabrication of Highly Efficient Fe3O4/SSIP/GO Composite Films for Removal of Methylene

Blue Dye

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PAGES: 2829-2837

ORIGINAL PDF URL: https://dergipark.org.tr/tr/download/article-file/3131199

#### Iğdır Üniversitesi Fen Bilimleri Enstitüsü Dergisi, 13(4), 2829-2837, 2023 Journal of the Institute of Science and Technology, 13(4), 2829-2837, 2023

#### ISSN: 2146-0574, eISSN: 2536-4618 DOI: 10.21597/jist.1294448

### Chemistry

**Highlights:** 

#### **Research Article**

Received: 09.05.2023

Accepted: 14.06.2023

**To Cite**: Nas, M. S., Calimli, M. H. & Demirbaş, Ö. (2023). Fabrication of Highly Efficient Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO Composite Films for Removal of Methylene Blue Dy. *Journal of the Institute of Science and Technology*, 13(4), 2829-2837.

#### Fabrication of Highly Efficient Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO Composite Films for Removal of Methylene Blue Dye

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#### **ABSTRACT:**

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- The composite film was synthesized by an extruder device.
- Tested optimum conditions for methylene blue removal
- Adsorption kinetics were performed using three different models

#### Keywords:

- Sunflower
- Adsorption
- Composite film
- Characterization

In this paper, we report the preparation of a new Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film for the purification of methylene blue dye from solution media. The preparation process of composite film was carried out to improve the interaction between the inner part of the sunflower stalk (SSIP) and surface matrix using magnetite/Graphene Oxide (Fe<sub>3</sub>O<sub>4</sub>/GO) minerals at weight ratios (0.2/4/0.2:w/w/w). The characterizations of as-developed Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film was successfully carried out by some advanced techniques such as FT-IR, SEM, and TGA analyses. Adsorption kinetic studies were evaluated in three different aspects as a pseudo-first-order model, pseudo-second-order model, and intraparticle diffusion-type model. Based on the R<sup>2</sup> results, it was seen that the Pseudo-second order kinetic model (0.999) acted more harmoniously than the pseudo-first-order (0.960) and intra-particle diffusion technique models (0.974). In the light of these findings, it can be said that the prepared Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film used for removal of methylene blue dye can be considered as a promising material

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#### **INTRODUCTION**

The sunflower plant, which is an annual and broad-leaved species, constitutes the majority of oil resources worldwide(P. et al. 2009) (Vaithanomsat et al., 2009). Parallel to the increase in the world population, the increase in the amount of oil consumption has resulted in a positive increase in sunflower cultivation areas. After the sunflower plant is harvested, an extremely high significant amount of product is produced as as the leaves, heads and stems of the sunflower plant (Follain et al. 2015). If sunflower stalks are not well controlled, a way is followed by burning them. This method leads to environmental pollution with severe consequences and misuse of the source material (Shi and Yang, 2012). However, a high amount of pectin and cellulose is obtained by extracting sunflower stalks (SSP), and a different alternative source is gained for the obtained pectin extraction, biofuel extraction, or cellulose-based materials (Xu et al. 2020), a very large specific surface area, and a special structure compared to lignocellulosic fiber materials (Sun et al., 2013). SSP can be considered an adsorbent source for the removal of polluted wastes in wastewater due to its unique physicochemical structural properties(Oguntimein 2015; Nas 2021), easy degradability (Jalali and Aboulghazi 2013), and costeffectiveness (Knapik and Stopa 2018). The rapid increase in industrial activities in recent times has led to the formation of wastes that leave serious negative traces on the environmental cycle and human health. Pollutants released into the aquatic environment from different industries are the main serious factors damaging the habitat. Toxic heavy metals, dyestuffs, drugs, organic acid products, phenolic substances, and halogenated compounds are some of the pollutants that negatively pollute the environment. Even in trace amounts, these products have very serious dangerous consequences for humans and aquatic organisms. E. Pollutants have a great role in environmental pollution due to the use of textiles, paper, cosmetics, printing, medicine, food, and leather (Tang et al. 2019). Dyestuffs in wastewater are very dangerous due to their high toxicity. In addition, since the dyes released into the environment give an undesirable color to the water, they prevent the sunlight from being taken sufficiently (Jiang et al. 2019). Before these waste materials are released into the environment, toxic and reactive dyes must be purified using appropriate processes (Nas and Kaya 2020). The adsorption process, membrane separation method, precipitation way, precipitation, ion exchange mechanism, flocculation process, electrochemistry way, photoelectrochemistry method, advanced oxidation process mechanisms (AOPs), and biological treatment method are among the most accepted methods to remove unwanted dyestuffs from the environment (Parvin et al. 2019; Wekoye et al. 2020). Among these treatment type methods, the adsorption method is the most preferred process approach due to its positive features such as being economical, easy to apply, and environmentally friendly (Cheng et al. 2017; Nas 2019; Bingül Reçber et al. 2022). In general, waste biomass products, clay-containing minerals, and activated carbon are used as a source of adsorbent material for dye removal in the adsorption process (Liu et al. 2019; Bayat et al. 2022). This study, the composite film obtained from a mixture of SSIP, Fe<sub>3</sub>O<sub>4</sub>, and GO was used as a vehicle for removing the MB dyestuff. The synthesized composite film samples were characterized by SEM, TGA, and FT-IR analysis techniques, respectively. As a result, we can say that it has been seen that it can be used effectively in the removal of pollutants such as methylene blue by using composite film in a water solution environment.

## **MATERYAL VE METOT**

### Chemicals

FeCl<sub>3</sub>.6H<sub>2</sub>O, FeCl<sub>2</sub>.4H<sub>2</sub>O, sodium hydroxide compound, ethanol, graphite material, and methylene blue were used from Sigma Aldrich Chemical Company. In this trial study, the inner part of the

sunflower stalk, which is used as a precursor material for dye removal, was obtained from the province of Iğdır in Turkey.

## Preparation of Fe<sub>3</sub>O<sub>4</sub>

 $0.2 \text{ g FeCl}_3$  and  $0.08 \text{ g FeCl}_2$  were mixed in 36 mL deionized water at 50 °C for 15 minutes. Then, NaOH solution (0.1 M) prepared to the reaction medium was added dropwise until the black color disappeared (pH=9), and the solution mixture was stirred vigorously for 1 hour at room temperature. The final product was separated by a supermagnet material by thorough washing with deionized water and ethanol and allowed to dry for 2 hours in an oven at 60 °C (Wei et al. 2012).

## Preparation of graphene oxide

Graphene oxide synthesis was prepared with reference to the Hummers process (Benzait et al. 2021). The oxidation was carried out by adding sulfuric acid (360 mL) and phosphoric acid (40 mL) to the graphite flake (3 g) materials, respectively. After stirring for 30 minutes, potassium permanganate (18 g) was added to the medium and stirred for a day so that the oxidation process was efficient. Then 500 ml of water and 10 ml of hydrogen peroxide were added to this mixture. The final mixture was allowed to cool in an ice bath. Then the mixture was washed with copious amounts of HCI solution (1 M) and distilled water and left to dry in an oven at 70 °C for 12 hours.

## Preparation of Fe<sub>3</sub>O<sub>4</sub>/ SSIP /GO composites film

Before mixing, the Fe<sub>3</sub>O<sub>4</sub>, SSIP, and GO samples were dried at 105 °C for 2 hours. The products were then cooled to room temperature in a desiccator. After mixing the Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film at weight ratios (0.2/4/0.2:w/w/w), they were combined in a single screw extrusion apparatus. Composite film was made at a temperature range of 140-170 °C and a screw speed of 15 rpm.

### Characterization of prepared composite films

Fourier transforms infrared (FTIR) spectrum measurements of the materials prepared with SSIP, GO, and Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO, a Perkin Elmer Spectrum brand device was used by taking the wavelength range of 2500-400 as reference. Thermogravimetric measurements and differential thermal analyzes (TG/DTA) of the prepared samples were performed by taking the 25–800 °C temperature range as a reference, taking into account the 15°/min heating rate and nitrogen atmosphere. The morphological structures of the targeted products were analyzed using a scanning electron microscope (SEM) device.



Figure 1. FTIR spectrum of Fe $_3O_4$ , GO, SSIP and Fe $_3O_4$ / SSIP/GO



Figure 2. SEM pattern of Fe<sub>3</sub>O<sub>4</sub>/ SSIP/GO

#### Adsorption test process of methylene blue

Adsorption test analyses are performed by setting up a three-necked laboratory-type glass balloon setup. Methylene blue stock solutions were prepared using the specified amount of distilled water as solvent. Experimental run analyzes were performed at room temperature at a stirring speed of 400 rpm (pH 9) at initial concentration conditions of 5 x  $10^{-6}$  M methylene blue. The effect of the prepared composite films on the dye removal was tested in a time interval of 1440 minutes and at 400 rpm. It was done using NaOH solution (0,1 M) and HCl solution (0,1 M) to adjust the pH of the methylene blue solution. 3 mL sample was taken from the solution to interpret each adsorption assay. These samples were centrifuged at 5000 rpm (Cary 1E UV-Vis spectrophotometer) for 5 minutes. Adsorption data were instantaneously followed and determined using Equation (1) to test the amount of methylene blue absorbed on the composite film surface as shown below.

$$qt = \frac{(co-ce)}{m} \times V \tag{1}$$

where qt is the initial amount of the adsorbent dyestuff type, Co is the amount of the initial MB concentration value used, Ct is the amount of the MB concentration value at any point, m is the mass of the supporting material type used and V is the volume of the solution used in the process. (Demirbaş et al. 2002).



Figure 3. TGA curve of Fe<sub>3</sub>O<sub>4</sub>/ SSIP/GO

#### **BULGULAR VE TARTIŞMA**

#### Investigation of the morphological and chemical structure of the composite film

The FT-IR spectrum of Fe<sub>3</sub>O<sub>4</sub>, GO, SSIP, and Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film particles is imaged in Figure 1. The band peak at 1588 cm<sup>-1</sup> in the FTIR spectrum of SSIP indicates the C—O stretch

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vibration peaks of aromatic carbonyl groups C=O, and C=C groups in the raw spectrum. Band peak values close to 1425 cm<sup>-1</sup> represent the CH<sub>2</sub> asymmetric deformation state and the variation due to aromatic C-C stretching. The peak band values observed between 1390 and 1011 cm<sup>-1</sup> represent the peaks associated with carboxylic acids, alcohols, esters, and phenolic compounds in lignin (O-H bending, C-O stretching). In the structure of cellulose and hemicellulose, a peak state of C-O-C stretching appeared at 1050 cm<sup>-1</sup> (Podder & Majumder, 2015). Peak band status was observed between 900 and 600 cm<sup>-1</sup> due to antisymmetric out-of-plane aromatic ring stretching (Baysal et al., 2018). SEM device was used to define the surface morphology of the targeted materials. The variation in the morphology of the doped iron oxide compound, GO, and crude SSIP were explained using SEM images of samples at different nanoscales in Figure 2. TGA analysis results are given regarding the samples' thermal degradation and mass losses. TGA results were used with the Perkin Elmer Pyres model analyzer. The Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film was analyzed at 0–500 °C. As indicated in Figure 3, all TGA analysis results were performed in a nitrogen gas environment and at a heating rate of 10 °C/min. Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film was synthesized using weight ratios (0.2/4/0.2:w/w/w) forming a homogeneous mixture. According to the TG/DTA curves of the Fe3O4/SSIP/GP composite film, it is seen that a two-step reaction interaction occurs. The composite film is resistant to temperatures up to 482 degrees, and it was noticed that if it was above this degree, it lost its strength at a high rate and its structure deteriorated. Therefore, a high rate of mass loss was observed.



**Figure 4.** (a) Effect adsorption of pH to MB removal (b) Effect adsorption of initial dye concentration to MB removal (c) Effect adsorption of temperature to MB removal

### Effect of diferent pH on adsorption kinetics

The variation of the adsorption kinetics of the dye solution at different pH ranges is given in Figure 4(a). As the pH increased, the amount of adsorption increased. Generally, there are both negatively and positively charged functional groups on the surface of the composite film used in adsorption kinetic studies (Nandi et al. 2009). The effect of removing methylene blue (MB) from the water solution with the composite film was tested in the pH range of 5-9. The hydrophobic functional groups on the composite film surface interact with MB (Ghaedi et al. 2015). The datas obtained from this result show the presence of very strong electrostatic forces between the positively charged MB and the negative density composite film when the pH value of the solution increases (Li et al. 2010; Çalımlı et al. 2019; Demirbaş et al. 2019).

### Effect of dye concentration on adsorption results

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Determination of the adsorption effect on MB dye concentration was tested by preparing samples of different concentrations of 0.5.10<sup>-5</sup>-2.5.10<sup>-5</sup> M. Experimental analyses in the presence of a contact time of 1440 minutes, pH 7, and a temperature of 25 °C. Increasing the MB dye concentration from 0,5.10<sup>-5</sup> M to 2,5.10<sup>-5</sup> M resulted in a negative synergistic resulting in a decrease in adsorption efficiency from 90% to 75%. The density of the pore areas available on the composite film is sufficient to adsorb a certain of dyestuff in concentration. The increase in dyestuff concentration above this value is not sufficient for adsorbing. In this case, it causes a slight decrease in the MB removal (Figure 4(b)).

## Effect of temperature parameter on adsorption Kinetics

In Figure 4(c), the adsorption kinetic change was tested under constant conditions determined at different temperatures. It has been observed that the temperature function is quite effective in the removal of MB dye adsorption on the composite film. Experimental studies were performed at 25, 38, 45, and 55 °C using pH 9 and  $20 \times 10^{-6}$  M initial methylene blue concentration constant conditions. It was determined that the best efficiency adsorption amount was obtained at 55 °C. This situation can be explained as follows. As the temperature increases, the adsorption value equivalent to the diffusion rate of the molecules on the composite film increases due to the increase in the kinetic energies of the molecules. The pores of the composite film increase in volume depending on the temperature increase. This creates a positive synergistic effect on the adsorption value of MB removal (Dahri et al. 2015). In addition, an image is shown in a lab-tested working example of MB dye removal with composite film (Figure 5) (Şen et al. 2018).



Figure 5.A lab-tested image of MB dye removal with Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film

## **Adsorption kinetics**

Adsorption kinetics were tested using the following formulas of pseudo-first-order, pseudosecond-order, and intra-particle diffusion model. Based on the  $R^2$  results, it was seen that the Pseudosecond order kinetic model (0.999) acted more harmoniously than the pseudo-first-order (0.660) and intra-particle diffusion model (0.974)

$$\ln(q_e - q_t) = \ln q_e - k_i t \tag{2}$$

$$\frac{t}{at} = \frac{1}{k_2 a_2^2} + \frac{1}{a_2} t \tag{3}$$

$$q_t = k_{int} t_{1/2} + C$$
 (4)

Where; the t,  $q_e$ ,  $q_t$ ,  $k_2$  and  $k_{int}$  (mg (g min<sup>-1/2</sup>)<sup>-1</sup> is time (min.), first amount of MB, last amount of MB, and the constant ratios, respectively (Ho and McKay 1999; Demirbaş et al. 2014).

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Isotherm Model	Model Equation	Parameters	Değişken sabitler
Freundlich	$\log(a_{2}) = \log KE + \frac{1}{\log C_{2}}$	R <sup>2</sup> =0.996	$\mathbf{K}_{\mathbf{F}}$ :Freundlich isotherm constant
	$log(qe) = log RF + \frac{-log Ce}{n}$		<b>n</b> :Adsorption intensity
Langmuir	Ce1_Ce	R <sup>2</sup> =0,948	<b>K</b> <sub>L</sub> :Langmuir isotherm cons.
	$\overline{qe} = \frac{1}{K_L(qmax.)} + \frac{1}{qmax.}$		qmax.:maximum adsoprtion capacity
Temkin	Temkin $RT (lm KT + lm Ca)$		<b>b:</b> Temkin isotherm cons.
	$qe = \frac{b}{b}(mkT + mce)$		<b>K</b> <sub>T</sub> :Temkin isotherm binding cons.

Tablo 1.	Isotherm	models	used	for	testing	adsor	ption	kinetics
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Langmuir, Freundlich, and Temkin adsorption isotherm models were tested to investigate adsorption isotherms. The isotherm parameters of the adsorption are detailed in Table 1. Based on the  $R^2$  results, it was found to be more compatible than the Freundlich (0.996), Langmuir (0.948), and Temkin (0.964) isotherm models.

## CONCLUSION

Composite film consisting of Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO was successfully prepared and applied to investigate the methylene blue removal efficiency. The kinetic parameters (temperature, initial concentration of methylene blue, and pH) of the adsorption between the Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film and methylene blue were analyzed. Experimental datas showed that increasing methylene blue concentration has a positive synergistic effect on adsorption efficiency. In addition, it was determined that the adsorption increased when the pH values of the prepared dye solution increased from 5 to 9. Adsorption kinetic studies were carried out at temperatures, contact times, and pH conditions, which determined the adsorption mechanism and isotherm model. As a result of the calculations, it was determined that there is a positive relationship consistent with the adsorption kinetic datas of MB on the Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film surface, Freundlich isotherm (R<sup>2</sup> =0.996) and the pseudo-second-order kinetic process model (R<sup>2</sup> =0.999). In this study, we can say that Fe<sub>3</sub>O<sub>4</sub>/SSIP/GO composite film can be preferred as an effective adsorbent for dye removal.

# TEŞEKKÜR

This study was funded by Iğdır University Scientific Research Projects Commission (No: MÜF0520A12).

# Çıkar Çatışması

The article authors declare that there is no conflict of interest between them.

# Yazar Katkısı

The authors declare that they have contributed equally to the article.

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