Preparation and Characterization of ZnO-Zeolite Nanocomposite for Photocatalytic Degradation by Ultraviolet Light

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ABSTRACT

The increasing growth of the textile industry does not only provide benefits in the economic sector but also has the potential to damage the environment, because it generates the dye wastewater which is hard to eliminate. Procion red is one of the synthetic textile dyes that is toxic to the aquatic environment and it needs to be processed properly. The photocatalytic method of processing dye wastewater is the most effective, because it can remove the harmful pollutants in the dye wastewater. This study aimed to prepare and characterize the ZnO-Zeolite nanocomposites for photocatalytic applications tested with a 50 mg/L procion red dye sample. The nanocomposites consisted of the ZnO semiconductors and synthetic zeolite adsorbents prepared by using the sol-gel method. The dye degradation test was carried out under the irradiation conditions with ultraviolet (UV) lamp. Apart from the ZnO-Zeolite nanocomposite, testing was also carried out with the synthetic zeolite and ZnO. The results of SEM-EDX and XRD characterization proved that the nanocomposite forming components were ZnO and zeolite and could be seen from the resulting peaks. BET showed that the surface area value of the ZnO-Zeolite nanocomposite increased to 95.98 m²/g, the pore size of the ZnO-Zeolite nanocomposite was 4.42 nm, and the total pore volume was 0.08 cm³/g. The obtained average crystalline size of ZnO-Zeolite nanocomposite was 32.87 nm. The percentage of dye degradation using the ZnO-Zeolite nanocomposite for 120 minutes has reached 90.42%.

Keywords: nanocomposite, ZnO, synthetic zeolite, sol-gel, photocatalytic, procion red

INTRODUCTION

The textile industry plays an important role in the Indonesian industry and increases rapidly. The Ministry of Industry in Indonesia shows that the export value of the textile industry and textile products sector throughout 2019 reached US$ 12.9 billion and the growth performance of the textile industry was recorded to have increased by 19%. However, apart from contributing to the economy, the textile industry also has the potential to damage the environment. The color waste causes pollution that is toxic to the environment. The waste containing dyes usually consist of non-biodegradable organic compounds that can cause pollution in the environment, especially in the aquatic environment, so that the dye must be oxidized into simpler molecules (Viswanathan, 2018).

Procion red is one type of synthetic dye that is most often used in the textile industry, such as the fabric or batik home industry. This dye is very difficult to degrade. Consequently, it is necessary to have an effective treatment to reduce the color content and organic compounds in the textile wastewater. Several methods of processing the
textile wastewater have been developed by many researchers, namely by the processes of adsorption, chlorination, ozonation, and biodegradation (Amri and Utomo, 2017).

Some of the weaknesses of this methods are – in addition to the less than optimal degradation results (Naimah and Rahyani, 2014) – high operational costs, generation of secondary pollutants, and the fact that these methods are still relatively difficult to apply in Indonesia. The use of photocatalysts with the photodegradation method is the most effective alternative in the wastewater treatment. The photodegradation method utilizes photocatalysts and irradiation by UV light (Agustina et al., 2015). Photocatalysis is used to degrade the organic components with the help of ultraviolet light (Sharfan et al., 2018). The UV radiation required for the photocatalytic processes can be obtained from artificial sources or the Sun (Krzemincka et al., 2015).

\( \text{TiO}_2 \) and \( \text{ZnO} \) are semiconductors that are most widely used as photocatalysts in the photocatalytic degradation of dye wastewater. However, \( \text{ZnO} \) is more readily available, cheaper, and \( \text{ZnO} \) can absorb more of the solar spectrum than \( \text{TiO}_2 \), so that \( \text{ZnO} \) has a higher photocatalytic activity (Saravanan et al., 2013). \( \text{ZnO} \) is widely used as a photocatalyst, because it has good chemical stability, and is classified as a non-toxic material (Kusdianto et al., 2019).

One of the disadvantages of the photocatalytic process is the weak adsorption capacity. In order to overcome this problem, the photocatalyst can be combined with an adsorbent as a supporting material. Wismayanti et al., (2015) combined the \( \text{ZnO} \) photocatalyst and activated charcoal adsorbent used for the degradation of the methylene blue dye.

The combination with this adsorbent aims to optimize the photocatalytic contact with dye pollutants. In addition, the adsorbent does not need to be regenerated, because the pollutants that have stuck to the adsorbent will be degraded in situ by the photocatalyst, so that adsorbent saturation can be avoided. Good adsorbents have high adsorption capacity and will certainly result in a high absorption percentage (Darmansyah et al., 2016).

Zeolite is a porous material that has been widely used as a catalyst, adsorbent, and ion exchanger (Rahman et al., 2018). Synthetic zeolite was chosen to be the adsorbent because synthetic zeolite has the physical properties that much better than those of natural zeolite. Besides, synthetic zeolite has a uniform pore size and is more evenly distributed so that the adsorption results will also be maximized.

One of the most successful methods for preparing the nano-sized metal oxide materials is the sol-gel method; hence, this method is suitable for application in the preparation process of the \( \text{ZnO-Zeolite nanocomposites} \). The sol-gel method is used for the preparation of thin films and powder-shaped materials, where the alkoxides are hydrolyzed by alcohol to metal hydroxides (Rahman et al., 2018).

In this research, the characterization and preparation process of the \( \text{ZnO-Zeolite nanocomposites} \) for the photocatalysis was carried out by testing the degradation of the procion red dyes using the sol-gel method. The advantage of the \( \text{ZnO-Zeolite composite produced is that the ZnO-Zeolite composite is nano-sized so that its photocatalytic activity is maximized and is indicated by the high dye degradation results. The costs required are also lower due to the use of ZnO photocatalyst raw materials and zeolite adsorbents which are more affordable than other materials.}

**MATERIALS AND METHODS**

**Materials and preparation**

Precursor \( \text{Zn(CH}_3\text{COO)}_2\cdot2\text{H}_2\text{O} \) (Zinc Acetate), \( \text{ZnO} \), synthetic zeolite, 99% ethanol, NaOH, and HCl were supplied from Sigma Aldrich and synthetic procion red powder dyes were obtained from dye stuff store (Fajar Setia, Jakarta).

Zeolite is activated to increase its purity (Wulandari et al., 2019). Synthetic zeolites that were crushed with grinders and sifted with 400 mesh sieve were then activated by heating in an oven at 110°C for 2 hours, then washed with 0.4 M HCl for 1 hour, followed by washing with distilled water until neutral. The synthetic zeolite was filtered and dried in an oven at 110°C for 120 minutes.

**Materials characterization**

Zeolite before activation, zeolite after activation, and \( \text{ZnO-Zeolite Nanocomposite} \) were characterized by X-Ray diffraction (XRD), Scanning Electron Microscope-Energy Dispersive X-ray (SEM-EDX), and Brunauer-Emmett-Teller (BET). XRD analysis was carried out to see the crystal structure based on the information on the
peaks of the scattering angle (XRD Bruker D8 Advance), SEM-EDX analysis to see the morphology or surface image of the material (SEM-EDX Hitachi SU3500), and BET for specific surface area, mean pore size, and the pore volume in the sample (BET Nova 4200e).

**Preparation of the ZnO-Zeolite nanocomposite**

Zn \((CH_3COO)_2\cdot2H_2O\) precursor is added with synthetic zeolite that has been activated in the precursor and synthetic zeolite ratio of 2:1 (Salam et al., 2018), then dissolved in 80 mL of 99% ethanol. The precursor and zeolite mixture was heated to 76°C for 120 minutes while stirring in a reflux flask. Then, 225 mL of 2 M NaOH was added to the solution and stirred for 1 hour. The mixture was allowed to stand for 12 hours and was then filtered with a Whatman filter paper. The precipitate obtained was then heated in an oven at 60°C for 24 hours and then stored in a desiccator to keep it dry. The ZnO-Zeolite nanocomposites were analyzed by BET, SEM-EDX, and XRD.

**Photocatalytic Degradation Test**

The ZnO-Zeolite nanocomposite was tested in a degradation process of 50 mg/L procion red synthetic dye. The photocatalytic test was carried out by using a UV lamp. The dye photodegradation was also applied with synthetic zeolite and Zinc Oxide (ZnO) as a comparison.

The nanocomposite of ZnO-synthetic zeolite was weighed in the amount of 0.1 g and then mixed with 25 ml of synthetic dye (Salam et al., 2018) with the concentration of 50 mg/L, the mixture was then stirred in a shaker at 320 rpm and placed in a reactor that has been assembled with ultraviolet lamp (Evaco 254 nm). The degradation time started when the UV lamp was turned on and the samples were taken for 5, 10, 15, 20, 25, 30, 60, 90, and 120 minutes to analyze the color degradation. The samples were taken after the specified time and then filtered by a Whatman filter before the dye concentration was analyzed using a UV-Vis Spectrophotometer.

**Analysis of degradation results with a UV-Vis spectrophotometer**

The degraded sample mixture was filtered to separate the ZnO-Zeolite nanocomposite from the procion red synthetic dye. The absorbance and final concentration result of the procion red’s dye which has been absorbed by the ZnO-Zeolite nanocomposite was measured using a UV-Vis spectrophotometer at the maximum wavelength. From these absorbance values, the concentration of procion red, as well as the percentage of procion red photodegradation were calculated.

\[
\text{Degradation Percentage} = \frac{(C_1-C_2)}{C_1} \times 100\% \quad (1)
\]

Eq. (1) shows the formula to calculate the degradation percentage of the dye, where \(C_1\) is the initial concentration of dye and \(C_2\) is the final concentration of dye that were measured by using a UV-Vis Spectrophotometer.

**RESULTS AND DISCUSSION**

**The characterization results of the ZnO-Zeolite nanocomposite**

The characterization of the ZnO-Zeolite nanocomposite was carried out by SEM-EDX, BET, and XRD. The XRD results of zeolite before activation and after activation are presented in Figure 1 (a) and (b).

The diffraction patterns of synthetic zeolites before and after activation show almost the same peaks, but the intensity at the peaks is different. Synthetic zeolite after activation has a decreased peak number. The XRD results of synthetic zeolite before activation showed that peaks were at 2θ = 10.01°, 11.5°, 14.01°, 22.0°, 24.99°, 25.5°, 26.01°, 29.99°, and 31.9°. After the activation process, the synthetic zeolite diffraction pattern showed that the peak was at 2θ, namely 10.01°, 12°, 14.99°, 20.11°, 22.08°, 26.01°, 27.11°, 29.98°, and 31.00°. This peak indicates the presence of zeolites.

This is similar to the research by Nyankson, (2018), who stated that this type A zeolite records peaks at 2θ: 7–8, 10–11, 22–24 and 29–31 is characteristic of zeolite A with cubic crystals.
at ZnO at 2θ (degree) showed the peak values of 31.95; 32.71; 34.61; 36.44; 47.75; 56.75; 63.02; 66.54; 68.10; 69.25; 72.71; 76.61 and 77.13. The XRD ZnO pattern results are identical to the hexagonal phase with the Wurtzite structure with a space group P63mc where, a = 3.247 Å and c = 5.198 Å (Alfarisa et al, 2018).

Figure 3 presents the XRD results for the ZnO-Zeolite Nanocomposite. The XRD results obtained were identical to the ZnO peak with diffraction peaks at 2θ of 31.43°, 34.51°, 36.37°, and 54.99°. This is in accordance with the XRD results for hexagonal ZnO (Mohan and Renjanadevi, 2016) and shows ZnO with a wurtzite
structure (Kang et al, 2010). The XRD of nano-composite showed that other peaks were also found to be around angles of $2\theta = 10.01^\circ$, $22.12^\circ$, $26.11^\circ$, and $29.98^\circ$, this indicates the presence of synthetic zeolites similar to the results of the study (Nyankson, 2018).

The constituent components of the ZnO-Zeolite nanocomposite can be seen from the peaks produced from the graph, where the degrees of the peaks are shown, which indicate that the constituent components of the nanocomposites are ZnO and zeolite. The morphology of synthetic zeolites before and after the physical and chemical activation processes showed different results.

On the basis of the XRD data, the calculation performed in accordance to the Scherrer method using X-Powder Software obtained an average crystalline size of the ZnO-Zeolite nanocomposite of 32.87 nm. This suggests that the resulting ZnO-Zeolite composite has reached the nano-size, so it can be concluded that the ZnO-Zeolite nanocomposites prepared by using the sol-gel method were successfully formed.

The results of zeolite characterization before and after activation with magnifications of 25 thousand times are presented in Figure 4. (a) and (b). The structure and shape of the activated zeolite look more regular and smoother when compared to the zeolite structure before the activation process. The zeolite size that has been activated is also more uniform than that of the zeolite before activation, although it is clear that there are large and small ones.

Figure 5 (a) shows the morphological results of ZnO with SEM magnification of 50 thousand times. The morphology of ZnO showed a fairly uniform shape; each particle is evenly distributed and homogeneous. This zinc oxide has a hexagonal phase with the wurtzite structure when viewed from the XRD analysis.

Figure 5 (b) presents the results of SEM characterization of the ZnO-Zeolite nanocomposites with magnification of 25 thousand times. The
particle size distribution of the nanocomposites looks more uniform, the distribution of particles is also more even and smoother. The particles also appear to be arranged more uniformly.

From the SEM images of the nanocomposites, it can be seen that there are two types of components in composite with different colors. The results of the SEM morphological analysis were marked by the sticking of ZnO particles on the surface of the zeolite layer. There is a combination of colors from the two different components, this can be seen from the dark colored zeolite and the light colored ZnO.

Table 1 presents the EDX results of the elements contained in zeolite before and after activation, ZnO, and ZnO-Zeolite nanocomposite. From the results of the EDX characterization, it can be seen that the chemical elements in the active zeolite are 43.99% wt of O element, 30.01% wt of Si, 20.58% of C, and 5.42% of Al.

The O, Al, and Si content of synthetic zeolites after the activation process were higher than those of synthetic zeolites before activation. The EDX results of ZnO show that ZnO contains zinc and oxygen components with a weight percentage of 65.89% Zn and 34.11% O. Zn has a greater percentage than oxygen. The ZnO-zeolite nanocomposite showed a Zn content with a weight percent of 56.13% and an oxygen content of 22.42%. The content of Zn has a greater percentage than other elements. The percentage of Zn and oxygen present in the ZnO-zeolite nanocomposite indicates that zinc and oxygen are present in the nanocomposite. The presence of Zn in the nanocomposites is due to the synthesis process made by a sol-gel process using a zinc acetate precursor.

There are other elements in the nanocomposite, such as C in 12.61%, Al in 2.75% and Si in 6.09%. The silica and aluminum contents indicate the presence of zeolite in the nanocomposites. These elements indicated that zeolite and ZnO are contained in nanocomposite and it proved that the ZnO-Zeolite nanocomposite has been successfully synthesized.

The results of BET characterization showed the values of the surface area, pore size, and pore volume of the ZnO-Zeolite nanocomposite sample. Zeolite before and after activation was also analyzed by BET.

The BET results of the three materials are presented in Table 2. The synthetic zeolite before activation has a lower surface area than the zeolite after activation. On the basis of the BET analysis, it is seen that the specific surface area of the synthetic zeolites before activation is 20.30 m²/g, while the surface area of the zeolites after activation is 47.19 m²/g. The surface area of the synthetic zeolite after the activation process showed a sufficient increase.

This increase in surface area is due to the opening of the synthetic zeolite pores which were originally covered by impurities. The release of these impurities can open the pores of the synthetic zeolite during heating in the zeolite activation process. The heating process will evaporate the water molecules and other impurities contained in the zeolite crystals so that a cavity with a surface that is larger than before activation is formed. Pore size and pore volume are inversely related to surface area.

The pore size of zeolite before activation was 21.98 nm, decreased to 9.22 nm after the

<table>
<thead>
<tr>
<th>Component</th>
<th>Weight percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>ZnO</td>
</tr>
<tr>
<td></td>
<td>Synthetic zeolite before activation</td>
</tr>
<tr>
<td></td>
<td>35.24</td>
</tr>
<tr>
<td>O</td>
<td>34.11</td>
</tr>
<tr>
<td>Zn</td>
<td>65.89</td>
</tr>
<tr>
<td>Al</td>
<td>2.81</td>
</tr>
<tr>
<td>Si</td>
<td>18.00</td>
</tr>
</tbody>
</table>

Table 2. BET characterization results

<table>
<thead>
<tr>
<th>Measurement subject</th>
<th>ZnO</th>
<th>Synthetic zeolite before activation</th>
<th>Synthetic zeolite after activation</th>
<th>ZnO-zeolite nanocomposite</th>
</tr>
</thead>
<tbody>
<tr>
<td>Specific surface area (m²/g)</td>
<td>19.19</td>
<td>20.30</td>
<td>47.19</td>
<td>95.98</td>
</tr>
<tr>
<td>Pore size (nm)</td>
<td>4.27</td>
<td>21.98</td>
<td>9.22</td>
<td>4.42</td>
</tr>
<tr>
<td>Total volume in pores (cm³/g)</td>
<td>0.04</td>
<td>0.11</td>
<td>0.09</td>
<td>0.08</td>
</tr>
</tbody>
</table>
activation process. The total pore volume of synthetic zeolite decreased from 0.11 cm³/g to 0.09 cm³/g after activation. The process of heating during zeolite activation causes the release of the impurities that cover the pores of the zeolite and result in reduced pore size and volume. The pore volume shows the pore depth of the zeolite produced.

Meanwhile, the BET ZnO results show that ZnO has a surface area of 19.19 m²/g, a total pore volume of 0.04 cm³/g and a pore size of 4.27 nm. ZnO has a smaller surface area. BET characterization results for the ZnO-Zeolite nanocomposites showed the highest surface area value of 95.98 m²/g. However, the pore size and total pore volume decreased. The pore size of the ZnO-Zeolite nanocomposite is 4.42 nm, and the total pore volume is 0.08 cm³/g. This is also due to the ratio of zinc acetate precursor in nanocomposites that are larger than synthetic zeolite so that ZnO can cover zeolite pores and cause pore size and reduce the pore volume of the nanocomposite. This will have an impact on the degradation process, because the surface area and the pore volume can increase the dissolution rates and adsorption capacity.

The Degradation Results with a UV-Vis Spectrophotometer

The degradation process of the procion red dye was carried out using different materials, namely ZnO-Zeolite nanocomposite, synthetic zeolite, and ZnO. The percentage of dye degradation obtained shows the ability of the material to degrade procion red dyes. The best percentage of degradation results was achieved by using the ZnO-Zeolite nanocomposites than the other materials. The absorbance value of the procion red dye solution with a concentration of 50 mg/L was measured with a UV-Vis spectrophotometer at a wavelength of 470–570 nm with a range of 10 nm. The maximum wavelength value obtained is 534 nm. The degradation time testing was carried out for 120 minutes, and the change in the final concentration of procion red was measured every specified time.

The percentage results of the procion red degradation using different materials, namely the ZnO-Zeolite, ZnO, and synthetic zeolite nanocomposites are presented in Figure 6. The photocatalytic process using the ZnO-Zeolite nanocomposites resulted in the highest percentage of degradation compared to using ZnO and synthetic zeolites.

The percentage of degradation results obtained with a concentration of 50 ppm procion red and using a UV lamp. Figure 6 shows the highest degradation percentage value is at 120 minutes using the ZnO-zeolite nanocomposite, which is 90.42% compared to those using synthetic zeolite and ZnO only that reached 50.64% and 80.64%, respectively. Meanwhile, during the initial 5 minutes the dye degradation process has also occurred, but the percentage of degradation or reduction in the concentration of the dye was still low, below 20 percent.

During the initial 25 minutes, degradation with the ZnO-Zeolite nanocomposite also reached a degradation percentage of 53.08%, and over 30 minutes it achieved more than 70%. This shows that the ability of the ZnO-Zeolite nanocomposites under UV light is more effective when compared to using only synthetic zeolite or ZnO-only degradation media, and the ZnO-Zeolite nanocomposite has high ability when exposed

![Figure 6. The photodegradation results of procion red 50 mg/l by using ultraviolet lamp](image-url)
to UV light. The smaller the size of the nanocomposites the greater the interaction of the reactants on the particle surface, enabling to achieve maximum degradation.

The correlation between the duration of degradation and the degradation percentage of the procion red is that the percentage of degradation will increase along with the degradation time used. Extending the dye degradation time will increase the formation of hydroxyl radicals as pollutant oxidator, so it gives more impact to the final degradation percentage (Agustina et al., 2015). The ZnO-Zeolite nanocomposite is the most effective material in degrading the procion dyes compared to synthetic zeolite and ZnO.

The degradation test with synthetic zeolite resulted in the lowest percentage of degradation. This is because the dye absorption process is only carried out by the synthetic zeolite adsorbents and only relies on the adsorption process on the zeolite surface. The greater the surface area, the more procion red dye can be absorbed (Indar and Kartikasari, 2017).

The ZnO-Zeolite nanocomposites produce the largest percentage of degradation compared to degradation using only ZnO or synthetic zeolites. This is because the ZnO-Zeolite nanocomposites do not only rely on the absorption of dyes by zeolite adsorbents, but the degradation process of dyes is also assisted by the presence of photocatalysts. ZnO in nanocomposites will increase the degradability of the dye with the help of a UV lamp. The percentage of dye degradation with an ultraviolet lamp will increase due to the use of a photocatalytic method which will break down the harmful components in the dye and produce safer final waste.

The photocatalytic method produces hydroxyl radicals which will become strong oxidizers in degrading waste, the process is simpler and rapid. Photocatalysis with the combination of the ZnO photocatalyst and synthetic zeolite adsorbent is more effective and the result of pollutant degradation is maximized. The common chemical feature in the photocatalytic process is the hydroxyl radicals formation as an oxidizer of organic pollutants (Zawadzki et al., 2018). The photodegradation method is cheaper and easy to apply (Agustina et al., 2020). Photocatalysis is a combination of a catalyst and photochemical processes. The photocatalytic process begins with the formation of pairs of positive electron holes (e⁺, h⁺) in semiconductor particles.

Riyani et al. (2015) explained that semiconductors have an empty valence band and conduction band so that exposure to the Sun or photon energy will cause excitation from the valence band to the conduction band (producing e⁺) which causes a void or hole in the valence band. Furthermore, the hole will react with H₂O in the solution to form hydroxyl radicals (•OH) which can degrade organic compounds into CO₂ and water. The electrons on the semiconductor surface can react with the electron catcher in the solution, namely O₂ to form superoxide radicals (•O₂⁻) which can also degrade organic compounds in solution. The radicals of (•OH) and (•O₂⁻) will continue to form as long as light rays still hit the photocatalyst and will degrade organic compounds into CO₂ and H₂O.

CONCLUSION

The ZnO-Zeolite nanocomposite that has been prepared by using the sol-gel method has surface area value of 95.98 m²/g, the pore size of 4.42 nm, and the total pore volume of 0.08 cm³/g. ZnO and zeolite components were also proven to be contained in nanocomposite, as showed in XRD peaks and SEM-EDX. The obtained average crystalline size of the ZnO-Zeolite nanocomposite amounted to 32.87 nm. The percentage of dye degradation using the ZnO-Zeolite nanocomposite for 30 minutes has reached 70% and after 120 minutes – 90.42%.

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