Characterization and Application of Chitosan as a Natural Coagulant in Reducing Remazol Red Dyestuff Concentration and COD Value of Batik Liquid Waste

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Abstract

Research on chitosan application as a natural coagulant in reducing the concentration of remazol red dye from batik liquid waste has been carried out. This study aims to study the effect of the acidity of batik wastewater on the coagulant ability to reduce the remazol red dye concentration and the COD value of batik liquid waste. In this study, chitosan compounds before and after coagulation were characterized using Fourier Transform Infrared (FTIR) Spectroscopy. The pH range used in the coagulation process was pH 2–6. Post-coagulation chitosan FTIR spectra showed a shift in the wave number in the 3400 cm⁻¹ area, which indicated an interaction between the –OH group of chitosan and the dye remazol red. Remazol red dye was maximally coagulated by chitosan at pH 2. The percentage reduction in the dye concentration reached 100%, and the decrease in COD value at that pH was 71.69%.

1. Introduction

The development of the batik industry has had some positive and negative impacts. The negative impacts include the emergence of environmental problems due to batik liquid waste containing synthetic dye residue. The use of synthetic dyes such as remazol, naphthol, and indigosol produces synthetic dye waste that is non-biodegradable, carcinogenic, and can cause liver, kidney, anemia, and other cell disorders [1]. The treatment of dye waste before disposal needs to be done to avoid the risk of these hazards. Several researchers have reported using the adsorption method to reduce the dye contained in batik wastewater [2, 3, 4, 5, 6, 7]. Based on this research, the use of the adsorption method could not reduce the dye concentration in batik waste optimally.

In this study, the remazol red concentration in batik wastewater was reduced using the chitosan coagulation method. Remazol red dye is one type of dye that is often used by the batik industry in Yogyakarta. Remazol red dye is widely used because it has a reasonably strong color resistance. However, its photodegradation process in nature is prolonged [8], so it is necessary to find an appropriate and straightforward method to reduce the concentration of remazol red dye in batik liquid waste.

Coagulation is treating waste by adding a coagulant to the waste by stirring rapidly so that a homogeneous waste dispersion is obtained. Coagulation occurs due to the ions originating from the coagulant, which have a charge opposite the colloidal particles' charge. The addition of a coagulant causes instability of the colloidal particles and binds the colloidal particles to form a floc. Coagulants can neutralize the charge of colloidal particles and bind these colloidal particles to form floc [9]. The use of coagulants from synthetic chemicals can cause new problems in the environment due to synthetic byproducts. Therefore, it is necessary to look for alternative coagulants from natural materials that tend to be environmentally friendly and effective in reducing the dye content in batik wastewater.

Chitosan is poly-2-amino-2-deoxy-β-1,4-D-glucopyranose with the molecular formula C6H11N6O6 obtained from the chitin distillation process. Chitin comes from the skin, head, and tail of shrimp, which can be obtained from waste of vanamei shrimp, crab waste, and silage from tiger prawn heads [10]. Chitosan has a
good role in reducing environmental pollution, which is quite effective in purifying water and wastewater. Chitosan can reduce color levels, pH, BOD, COD, and can absorb heavy metals and dyes [10, 11]. Based on the properties and advantages of chitosan, in this study, chitosan was applied as a natural coagulant to reduce the dye concentration of remazol red and the COD value of batik liquid waste. As far as the author's knowledge, chitosan has never been used to process batik liquid waste dye. The effect of the acidity of the batik wastewater on the coagulant ability to reduce the remazol red dye concentration and the COD value of batik wastewater was studied in the pH range 2–6.

2. Methodology

2.1. Equipment and Materials

The equipment used in this study were: a set of standard laboratory glassware, Whatman 42 filter paper, universal pH, tweezers, analytical balance, teaspoon and glass stirrer, hotplate centrifuge, Shimadzu-8201 PC Fourier Transform Spectrophotometer (FTIR), and Spectrophotometer. UV-Vis 1800 Double Beam Shimadzu.

The primary materials used in this study include ready-to-use chitosan from the isolation of shrimp or crab shells produced by CV Chimultiguna (the structure of the chitosan compound is presented in Figure 1), batik liquid waste, and a standard solution of remazol red dye specifically for batik dyes (the structure of the remazol red compound is presented in Figure 2).

Meanwhile, this study’s supporting materials included HCl, NaOH, and distilled water, all of which were of analytical quality obtained from Merck.

![Figure 1. Structure of chitosan](image)

**Figure 1. Structure of chitosan**

![Figure 2. Structure of remazol red dye](image)

**Figure 2. Structure of remazol red dye**

2.2. Material characterization

The coagulant chitosan and standard remazol red dye used in this study were characterized by functional groups using an FTIR spectrophotometer.

2.3. Preparation of batik liquid waste solution at various pHs

Batik liquid waste (30 mL each) was poured into a 100 mL beaker, and the pH is adjusted to the range 2–6. The pH adjustment was made by adding a solution of HCl and/or NaOH. The initial content of remazol red dye in batik liquid waste of various pH was analyzed using a UV-Vis Spectrophotometer. The initial and final COD values of batik liquid waste were tested at the Testing and Calibration Laboratory, Center for Environmental Health and Disease Control Engineering (BBTKLPP), Yogyakarta.

2.4. Remazol red standard solution preparation

2.4.1. Preparation of remazol red mother liquor

The main solution of 1000 ppm remazol red dye was prepared by dissolving 1 g of remazol red in 1000 mL of distilled water. Next, the mother liquor was diluted into 5, 10, 15, 20, and 25 ppm of 30 mL each, which was then used as standard solutions.

2.4.2. Determination of the maximum wavelength of the remazol red dye

The maximum wavelength was determined by measuring the remazol red solution in the wavelength range of 400–750 nm using a UV-Vis spectrophotometer. The maximum wavelength obtained was then used for the measurement of the standard solution.

2.4.3. Determination of the standard curve of the dye remazol red

The mother liquors of remazol red dye with variations of 5, 10, 15, 20, and 25 ppm were measured their absorbance using a UV-Vis spectrophotometer at the maximum wavelength of remazol red. Furthermore, a graph of the relationship between concentration and absorbance of remazol red dye solution was made as a calibration curve.

2.5. Coagulation of batik wastewater using chitosan as a coagulant

0.300 g of chitosan were each put into a 100 mL beaker and mixed with 30 mL of batik liquid waste solution with pH variations of 2, 3, 4, 5, and 6. The mixture was stirred for 60 minutes with a variety of stirring, where the first 3 minutes of stirring was done quickly, and the next 30 minutes was done with slow stirring. Then, the mixture was left for 30 minutes. Then the mixture was filtered using Whatman filter paper. The dye concentrations of each filtrate were analyzed using a UV–Vis spectrophotometer with three repetitions followed by COD testing. Meanwhile, the precipitate obtained was dried and then characterized using an FTIR spectrophotometer.

2.6. Testing the COD (Chemical Oxygen Demand) value

COD testing is carried out by sending the coagulated filtrate to the Testing and Calibration Laboratory, Center for Environmental Health and Disease Control Engineering (BBTKLPP), Yogyakarta.

3. Results and Discussion

3.1. FTIR spectra of chitosan

Chitosan characterization was performed using an FTIR spectrophotometer. This characterization was carried out on chitosan before the coagulation process.
and chitosan after coagulation. The purpose of characterization was to determine the functional groups in chitosan and determine changes in the functional groups in chitosan before and after coagulation. The FTIR analysis spectra of chitosan before and after coagulation are presented in Figure 3.

![Figure 3. FTIR spectra of (a) remazol red dye, (b) chitosan before coagulation, (c) chitosan after coagulation](image)

Table 1. Wave numbers on chitosan FTIR spectra, before and after coagulation

<table>
<thead>
<tr>
<th>No</th>
<th>Functional groups</th>
<th>wavenumber (cm⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>before coagulation</td>
</tr>
<tr>
<td>1</td>
<td>(vb) O-H which overlaps with (vs) N-H</td>
<td>3417.86</td>
</tr>
<tr>
<td>2</td>
<td>(vb) aliphatic C-H</td>
<td>2924.09</td>
</tr>
<tr>
<td>3</td>
<td>(vs) Aliphatic C-H</td>
<td>2877.79</td>
</tr>
<tr>
<td>4</td>
<td>(vs) aromatic C-H</td>
<td>1604.77</td>
</tr>
<tr>
<td>5</td>
<td>(v) C=O [secondary amide]</td>
<td>1640.47</td>
</tr>
<tr>
<td>6</td>
<td>(v) C=O [secondary amide protonation]</td>
<td>1604.77</td>
</tr>
<tr>
<td>7</td>
<td>(α) CH₃</td>
<td>1381.03</td>
</tr>
<tr>
<td>8</td>
<td>(ω) β-1,4-glycosidic</td>
<td>894.97</td>
</tr>
</tbody>
</table>

Note: vb: stretching vibration, vs: symmetrical stretching vibration, vb: bending vibration, α: stretching vibration

Figure 3 (b) shows that the chitosan spectra before coagulation had several main peaks at wavenumber 894.97 cm⁻¹; 1381.03 cm⁻¹; 1604.77 cm⁻¹; 1651.07 cm⁻¹; 2368.59 cm⁻¹; 2877.79 cm⁻¹; 2924.09 cm⁻¹; and 3417.86 cm⁻¹. The characteristic of chitosan can be seen by the appearance of stretching vibrations in the area of wave number 894.97 cm⁻¹, which indicates the presence of β-1,4-glycosidic bonds. The emergence of absorption at the wave number 1604.77 cm⁻¹ indicates a stretching vibration of the N-H group from the amide [12]. The appearance of absorption at wave number 1651.07 cm⁻¹ indicates a stretching vibration of the amide's N-H group. Meanwhile, Figure 3 (c) shows that the chitosan spectra after coagulation did not give rise to new absorption but several shifts in the wave number in the wavenumber 3400 cm⁻¹, as shown in Table 1. It is suspected that the interaction that occurs between chitosan and remazol red is merely a physical interaction. The absorption peak shift is thought to be due to the electrostatic interaction between chitosan and remazol red dye, described in more detail in section 3.2.

3.2. The effect of pH variation of batik wastewater on the ability of chitosan coagulant to reduce the concentration of remazol red

Many factors influence the coagulation process, one of which is pH. The pH variation of batik wastewater was carried out to determine the optimum pH conditions for the coagulation of remazol red dye using chitosan coagulant. The variation in pH in the coagulation process can determine the interaction model between the coagulant and the dye remazol red.

![Figure 4. The relationship between pH and the percentage of coagulated remazol red (%)](image)

Figure 4 shows that the chitosan coagulant’s ability to coagulate remazol red dye decreases with increasing pH. The highest coagulation pH was obtained at pH 2. This can be explained that chitosan is a compound that is rich in −NH₂ and −OH groups. In the acidic conditions, the functional groups undergo protonation of −NH₂ and −OH⁻ so that the chitosan surface becomes positive [2, 13, 14]. Meanwhile, remazol red is an anionic dye which contains a sulfonic acid group (RSO₃⁻). Under acidic conditions, the sulfonic groups of the remazol red are negatively charged, and with increasing pH, the charge becomes neutral [15]. This causes the coagulation process to run well at pH 2-3, due to the interaction between the coagulant and remazol red dye, which is suspected through electrostatic interactions. As the pH increases, H⁺ ions decrease so that the −NH₂ and −OH groups in the chitosan are not properly protonated. The interaction between coagulant chitosan and remazol red is getting weaker at pH > 3. This is because at pH > 3, the −NH₂ and −OH groups have begun to deprotonate into −NH⁻ and −O⁻ so that the surface becomes negative and makes electrostatic interactions more difficult to occur. This coagulation strengthens the chitosan FTIR spectra results before and after coagulation, which does not produce new absorption. It only shows a shift in the wave number in the 3400 cm⁻¹ absorption area, which is the O-H absorption of the N-H overlap. This shows that the interaction between chitosan coagulant and remazol red dye is through electrostatic interaction. An illustration of the interaction between chitosan and RR dye is presented in Figure 5.
3.3. The performance of chitosan coagulant in reducing COD value of batik liquid waste

One of the parameters of wastewater quality that is often tested is the COD (Chemical Oxygen Demand) value. Chemical Oxygen Demand is the amount of oxygen required to oxidize compounds present in the water through a chemical reaction. Organic compounds are oxidized by potassium bichromate to CO₂ and H₂O gases and chromium ions. The higher the COD value in a sample of wastewater, the lower the quality and vice versa. Organic compounds in batik liquid waste other than dyes are waxes, so the concentration needs to be lowered before being discharged into the environment. Based on Table 2, the percentage reduction in COD value in this study was 71.69%. This value is lower than the chitosan coagulant's ability to reduce the concentration of the remazol red. This is presumably because chitosan coagulant has not interacted strongly with other organic compounds in the batik liquid waste. So, it is necessary to find the optimum conditions that are more appropriate to treat other organic compounds in the batik liquid waste other than remazol red.

<table>
<thead>
<tr>
<th>Table 2. COD value of batik liquid waste at pH 2 conditions</th>
</tr>
</thead>
<tbody>
<tr>
<td>before coagulation (mg/L)</td>
</tr>
<tr>
<td>COD</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Table 3. The performance of chitosan coagulant in reducing COD value of several types of waste</th>
</tr>
</thead>
<tbody>
<tr>
<td>Types of waste</td>
</tr>
<tr>
<td>-------------------------------------</td>
</tr>
<tr>
<td>drainage water in the Singosari sewer in Semarang</td>
</tr>
<tr>
<td>laundry liquid waste</td>
</tr>
<tr>
<td>Batik liquid waste</td>
</tr>
</tbody>
</table>

4. Conclusion

The coagulation method using chitosan can reduce remazol red concentration and the COD value of batik liquid waste. Obtained coagulation optimum pH conditions = 2 with a percentage reduction in the dye concentration by 100%. The chitosan coagulant is very good at reducing the concentration of the dye remazol red in acidic conditions, increasing the pH, decreasing the ability of chitosan. The decrease in COD value at the optimum coagulation condition showed a decreased percentage of 71.69%. Further research is needed regarding other parameters that can affect the coagulation process, such as coagulant dose, stirring time, and settling time.

Acknowledgments

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References


Figure 5. Illustration of the interaction between chitosan and remazol red


