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Removal of Anionic and Cationic Dyes from Wastewater Using Activated Carbon from Palm Tree Fiber Waste

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Abstract: This study focuses on using a facile method for the green preparation of activated carbon (AC) from palm tree fiber (PTF) waste. The synthesized cost-effective AC was investigated for the removal of an anionic dye (Congo red, CR) and a cationic dye (Rhodamine B, RhB) from wastewater. The morphological and structural characterization of the synthesized AC were performed by scanning electron microscopy (SEM), transmission electron microscopy (TEM), surface area, Fourier transform infrared spectroscopy (FTIR), total pore volume, average pore diameter and pore size distribution, zeta potential, and zero-point charge. To investigate the adsorption efficiency, different parameters such as adsorbent dosage, solution pH, initial dye concentration, and duration were applied using the batch experiments. Various adsorption isotherm and kinetics models were applied to study the adsorption mechanism and dynamics. The results showed that chemical activation with a weak acid (H₃PO₄) at 400 °C for 30 min is a fast method for the activation of each precursor and produces a high yield. The result of analysis showed an increase in the adsorption capacity at pH 2. The maximum adsorption capacity was 9.79 and 26.58 mg g⁻¹ at 30 min for CR dye and RhB dye, respectively. The optimum adsorbent dosage for the activated carbon from palm tree fiber (PTFAC) was 0.15 g with a high percentage removal of CR (98.24%) and RhB (99.86%) dyes. The adsorption isotherm and kinetic studies were found to be favorable and feasible for assessing the adsorption of dyes with the Langmuir model and pseudo-second-order reaction, respectively. In addition, the AC showed reusability up to five cycles. The results showed that the synthesized AC was environmentally friendly and successfully removed dyes from wastewater.

Keywords: activated carbon; adsorption; anionic and cationic dyes; palm tree fiber wastes; recycles

1. Introduction

The continuous growing population and their requirements have seriously contaminated water with several undesirable materials, for example, artificial dyes [1]. Synthetic dyes obtained from organic or inorganic compounds are mainly composed of two compounds—chromophores and auxochromes. The chromophore produces the color of dye, whereas the auxochrome is responsible for the intensity of color [2,3]. Before the invention of artificial dyes in 1956, the natural dyes obtained from plant sources as roots, berries, bark, leaves, wood, fungi, and lichens were used [4]. After 1956, artificial dyes derived from petrochemicals were used as coloring agents for several products in the market. Additional engaging colors have been introduced to various advanced industries such as textiles, paper, animal skin tanning, food processing, plastic, cosmetic, and dye-producing industries [5]. Synthetic dyes are usually carcinogenic, cause serious damage to water resources, and contribute to environmental pollution. The environmental and health concerns associated with the wastewater effluents have led scientists to search for simple, inexpensive, and rapid solutions. The putrefaction of dyes from wastewater can be done using three methods, namely physical, biological, and chemical methods [6]. These include
precipitation, oxidation, coagulation, adsorption, and membrane separation [7–9]. Among these methods, adsorption is a cost-effective, simple, and feasible technique for the removal of organic dyes from wastewater. Many synthetic ingredients have been effectively used as adsorbents for the elimination of contaminants from water [10–12].

Among them, activated carbon (AC) is considered as the most commonly used adsorbent in the industry for the removal of contaminants [13]. The pore structure of the AC is quite developed with a high surface area and good adsorption capacity. The external chemical functional groups and structural properties of AC make it suitable for many applications used for the elimination of organic and inorganic contaminants from contaminated water [14,15]. However, the fabrication of marketable AC is costly; thus, research attention has been paid to develop a low-cost and competent AC using several sources. The production of AC from natural sources or agroindustry wastelands was considered because they are renewable, highly available, and cost-effective, and they have good properties. Thus, their disposal is required for a cleaner environment. In this regard, many researchers have reported on the preparation of AC from various industrial denim fabric waste [16] and agriculture waste (biomasses) such as coconut coir, apricot stones [17], Nigella sativa L [18], palm shell [19], pecan shell [20], pine cone [21], tapioca peel [22], macadamia nutshell [23], Mucuna pruriens and Manihot esculenta [24], snail shell [25], and tea leaves [26].

There are two methods to produce AC: physical and chemical activation. In physical activation, raw material is carbonized and then activated with steam and carbon dioxide. Chemical activation involves the impregnation of a chemical-activating agent in the precursor material followed by activation at temperatures between 400 and 700 °C under the nitrogen atmosphere [27]. The advantages of chemical activation are the following: (a) it occurs in a shorter time compared to that prepared by physical activation; (b) the produced AC has a high surface area with high yield; and (c) the activating chemical agent influences the breakdown of carbon and prevents the development of tar and volatile substance, thereby increasing the yield of AC. In addition, in the case of chemical activation, the process of dehydration and oxidation requires lower temperatures compared to that in the physical activation [28]. In the chemical activation method, ZnCl₂, KOH, and H₃PO₄ are extensively used as chemical-activating agents for the activation of lignocellulosic materials, which have not been carbonized. In addition, KOH is used for the activation of coal precursors or chars. The most common of activating agents include KOH and ZnCl₂ since they provide microporous features, which restricts these materials’ applications for the adsorption of high molecular weight molecules in aqueous solution [16]. Thus, attempts were made toward the use of H₃PO₄ as a proper and low-cost activating agent for AC as reported earlier for many biosorbent lignocellulosic material [16]. In the literature, various parameters were studied such as the concentration of activation agent, temperature, and activation time [29–31].

The present study focuses on the facile preparation of AC from biomass by chemical and thermal activation methods. The vast number of date palm trees in Saudi Arabia is about 23 million. Thus, palm tree fiber (PTF) is one of the greatest freely existing biomasses, and it is common agricultural waste in Saudi Arabia. Therefore, PTF has been chosen as the specific biomass in order to produce more economical AC and easy accessibility collection [32]. The parameters such as the activation agent, activation temperature, and activation time were investigated for PTF activation in this study. Another goal of the study was to use various analytical techniques to examine the efficiency of the prepared AC to remove anionic dye Congo red (CR) and cationic dye rhodamine B (RhB) from polluted water. The influence of initial concentration, duration, adsorbent dosage, and pH on the adsorption of CR and RhB were studied. Performing equilibrium isotherm and kinetics of adsorption for CR and RhB as well as exploring the re-generation of used activated carbon from palm tree fiber (PTFAC) were also considered as the main objectives.
2. Materials and Experimental Procedures

2.1. Instruments

The pH values of dyes and adsorbent powder were measured using a pH meter (MP220; Mettler Toledo). A cyclone mill (CT 193 CyclotecTM; Mill Collection) was used to ground the sample, and the sample was sifted by a KimLab ISO 3310 Std Test Sieve; the required size was a 0.43 mm mesh size. The mixtures were agitated using a reciprocating shaker (cat. no. 3006; GFL Shakers). The samples were carbonized in a muffle furnace (A-550; Vulcan), and after washing, they were dried in an oven (FN 055/120 Dry Heat Sterilizers). The spectroscopic reading was generated by using a UV-vis spectrophotometer (UV-1650PC, Shimadzu). Transmission electron microscopy (TEM) was performed using JEM-1230 (JEOL). The zeta potential of the PTFAC was measured by a Zetasizer (version 7.04, serial number MAL 1074157; Malvern Instruments). Scanning electron microscopy (SEM) was performed by Quanta FEG250. Fourier transform infrared spectroscopy (FTIR) was conducted by using a Spectrum Two FTIR Spectrometer (PerkinElmer) at the absorption spectra range of 4000–400 cm\(^{-1}\).

2.2. Materials and Chemical Reagent

Palm tree fiber waste was collected from the locally available palm trees on the campus of King Abdulaziz University, Jeddah, Saudi Arabia. The PTF was washed multiple times with hot distilled water to eliminate dust and impurities and then dried in sunlight for a few days. The dried palm fiber was crushed to powder form, sifted to the desired particle size, and used to prepare the AC. The obtained granular particle size of the pulverized fiber was approximately 0.43 mm. Raw materials and the prepared AC are shown in Figure 1.

![Figure 1](image_url)  
Figure 1. (a) Date palm tree, (b) Palm fibers, (c) Palm fibers powder, and (d) Activated carbon powder.

Congo red anionic dye (HiMedia) and RhB cationic dye (Sigma Chemical) were used as adsorbates (Table 1). Stock dyes solution (1000 mg L\(^{-1}\)) was made by dissolving a suitable amount of dye powder, which was accurately weighed on an electronic balance using deionized water. During the adsorption experiments, a diluted dye solution that had a concentration between 5 and 50 mg L\(^{-1}\) was prepared from the stock solution. The chemicals reagents such as H\(_3\)PO\(_4\), H\(_2\)SO\(_4\), HCl, and KOH were supplied by Fisher Scientific.

2.3. Chemical and Thermal Activation of Carbon from Palm Tree Fiber

The activation was carried out by impregnation of the PTF with various chemical agents as such as phosphoric acid (85%), sulfuric acid (98%), and potassium hydroxide (2 M) with an impregnation ratio of chemical/biomass being 3:1. The duration of the process was 24 h. The experiment was done under full safety precautions. Inside a fume hood, the chemical agent was added gradually to PTF to obtain a mixture. Then, the slurry was filtered, washed, and dehydrated in a dryer oven at 130 °C for 2 h. The designed
samples of chemically activated carbon PTF were termed as CAPTF-H₃PO₄, CAPTF-H₂SO₄, and CAPTF-KOH. The pyrolysis process was carried out to increase the efficiency of the fibers activated by phosphoric acid (85%). The fibers were carbonized at 200, 300, 400, 500, 600, and 700 °C several times (30, 60, 120, and 180 min) in a muffle furnace using a covered porcelain crucible for reduced oxygen. Then, they were cooled at room temperature and soaked in 1.0 M NaOH for 2 h. After soaking, the obtained materials were washed thoroughly with hot distilled water followed by cold distilled water to remove all free acid until they attained a neutral pH. Then, they were oven-dried at 130 °C for 2 h. Finally, the dried materials were milled into a fine powder and kept in a plastic bag until used. The samples produced by activated carbon palm tree fiber were termed as PTFAC, as shown in Figure 1.

Table 1. Properties of synthetic dyes.

<table>
<thead>
<tr>
<th>Dyes</th>
<th>Rhodamine B (RhB)</th>
<th>Congo Red (CR)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Molecular formula</td>
<td>C₂₈H₃₁N₂O₃Cl</td>
<td>C₃₂H₂₂N₄Na₂O₃S₂</td>
</tr>
<tr>
<td>Molecular structure</td>
<td><img src="image1" alt="RhB" /></td>
<td><img src="image2" alt="Congo Red" /></td>
</tr>
<tr>
<td>Molecular weight (g/mol)</td>
<td>479.02</td>
<td>696.66</td>
</tr>
<tr>
<td>Chemical/Dye class</td>
<td>Xanthene dye</td>
<td>Diazoy dye</td>
</tr>
<tr>
<td>λₘₐₓ (nm)</td>
<td>554 nm</td>
<td>496 nm</td>
</tr>
</tbody>
</table>

The adsorption process was performed by the batch-adsorption method. To select the fiscal chemical-activating agent (i.e., H₃PO₄, H₂SO₄, or KOH), the best pyrolysis temperature (200, 400, 600, or 800 °C), and suitable time (30, 60, 80, and 120 min) to remove CR and RhB dyes, the synthesized PTFAC were conducted under the following conditions: volume of dye solution = 20 mL, initial concentration = 25 mg L⁻¹, contact time = 120 min, temperature = 25 °C, adsorbent dose = 0.1 g, agitation speed = 300 rpm, pH = natural.

2.4. Characterization of AC

The specific surface area, total pore volume, average pore diameter, and pore size distribution of the prepared AC were examined by Brunauer–Emmett–Teller (BET) N₂ adsorption method. The presence of porosity, microanalysis, and surface morphology of the PTFAC carbonized at 400 °C were studied by using the SEM and TEM. The FTIR technique was used to study the effect of chemical treatment on biomasses and to recognize various functional groups present on the surface of the materials. This technique was used to examine the surface functional groups of the ACs.

Zeta potential measurement was used for determining the surface charge of AC particles in the solution. In other words, the possible difference between the dispersion medium and the stationary layer of fluid attached to the particle was measured by zeta potential. The pH of the solution with distilled water was measured as follows: the mixture was prepared in a ratio of 10 mL water to 1.0 g of carbon; this mixture was stirred, and the pH was measured several times until a constant value was reached. The pH of zero-point charge (pHzpc) for AC was determined by the batch equilibrium method. During the measurement, the solid/liquid ratio was 1:1000; that is, 0.05 g AC was added to 50 mL of water. This suspension was left for 48 h, and the pH was measured and found to be 3. In another experiment, 0.05 g of AC was mixed with 50 mL of 0.1 M NaCl, and the pH of solution was fixed between 2 and 12 by 0.1 M hydrochloric acid or 0.1 M sodium hydroxide. The solution was stirred for 48 h on a shaker bath at an agitation speed of 120 rpm and at room temperature (25 °C). Thereafter, the final pH values were recorded, and the difference between the original and final pH values (ΔpH = pH_initial – pH_final) was calculated and compared with the initial ones.
2.5. Adsorption Processing

The adsorption of dyes onto PTFAC was assessed using batch adsorption experiments. The procedure was performed as follows: First, a series of dye solutions with initial concentrations of 5.0–50 mg L\(^{-1}\) was prepared. Second, dye solutions with different pH values (2.0–12) were prepared. Finally, 20 mL of dye solution was added into 50 mL volumetric flasks with different amounts of PTFAC adsorbent doses (0.05–0.25 g). Then, the mixtures were stirred in a shaker at an agitation speed of 250 rpm at different times (5.0–120 min) at room temperature. The solutions were filtered using the Whatman Qualitative Filter Paper Grade 2. The dye concentration was determined by using the UV-visible spectrophotometer at wavelength 554 nm and 496 nm for RhB and CR dyes, respectively. All the experiments were performed in triplicate, and the mean values were recorded. The amount of all dyes adsorbed by the PTFAC and the removal percentage was calculated by Equations (1) and (2), respectively:

\[
\text{Removal \%} = \frac{C_o - C_e}{C_o} \times 100 \quad (1)
\]

\[
q_e = \frac{(C_o - C_e)}{m} \times V \quad (2)
\]

where \(q_e\) is the quantity of adsorbed dye on 0.1 g of PTFAC (mg \(g^{-1}\)), \(C_o\) is the original dye concentration (mg \(L^{-1}\)), \(C_e\) is the residual concentration of the solution at equilibrium (mg \(L^{-1}\)), \(V\) is the volume of dye solution (L), and \(m\) is the amount of the adsorbent PTFAC (g).

2.6. Desorption Studies and Regeneration

To understand the sustainability of regenerating the AC and make the process more economical, desorption trials were performed to determine the efficiency of the AC. The traits of an excellent adsorbent should include not only high adsorption capacity but also reduction of the overall cost of the adsorbent by possessing high desorption capacity. Desorption studies of PTFAC were carried out using different concentrations of acetic acid \(\text{CH}_3\text{COOH}\), sodium hydroxide \(\text{NaOH}\), and different pH values of distilled water. The R-PTFAC is a sample after adsorption used to study the regeneration and recycle. Thereafter, 0.1 g R-PTFAC was added to 50 mL desorption solutions (0.1, 0.5, 1.0 M \(\text{NaOH}\); 0.1, 0.5, 1.0 M \(\text{CH}_3\text{COOH}\)) and distilled water and the pH was adjusted (2.5, 4.5, 6.5, 8.5, and 10.5). Then, it was stirred in a shaker at 250 rpm for 3 h at room temperature. After desorption, R-PTFAC was washed with distilled water until it became neutral and was reused for the removal of RhB and CR dyes from solution. The adsorption–desorption studies were performed for five sequential cycles. The dye concentration in the filtrate was examined by the UV–vis spectrophotometer. The effectiveness of the desorbed dye from AC was evaluated by Equation (2).

2.7. Application of Real Water Samples

The real water samples were collected from the water sewage treatment plant and tap water at King Abdulaziz University, Jeddah City. The well water and seawater samples were collected from Rabigh City. The samples were applied by the adsorption of dyes onto the PTFAC using batch experiment. Under the same condition of the adsorption process, the removal percentage was calculated by Equation (2).

3. Results and Discussion

3.1. Physical Properties, Characterization, and Morphology of Synthesized PTFAC

Surface area, total pore volume, and mean pore diameter of the prepared AC, as determined by adsorption/desorption isotherm of nitrogen, are shown in Figure 2a. As a result, the prepared AC has a 648.90 m\(^2\) g\(^{-1}\) (BET) specific surface area and 2.83 cm\(^3\) g\(^{-1}\) total pore volume. It is noteworthy to mention that the specific surface area for the raw material before activation was 9.0 m\(^2\) g\(^{-1}\) [33]. As per the results of adsorption/desorption
isotherms presented in Figure 2a,b, the AC carbon follows type V isotherm of mesoporous materials with a 0.79 nm mean pore diameter, and the process is proceeded with multilayer adsorption. The data found are in line with the surface area value reported in the literature based on the modification method such as chemicals, physical, or pyrolysis activation [17].

![Figure 2](image)

**Figure 2.** (a) Adsorption/desorption isotherm and (b) pore size distribution of activated carbon from palm tree fiber (PTFAC).

On the other hand, the zeta potential and zero-point charge of the prepared AC are presented in Figure 3. The prepared PTFAC shows a zeta potential value of −28.3 mV and has high surface electronegativity. The pH values for PTFAC in a solution and pHzpc were 3.56 and 3, respectively. This means that the surface of PTFAC behaves as positively charged under a pH value of 3 and negatively charged beyond a pH value of 3 [34]. In addition, similar values of pHzpc have been reported for white pine sawdust (pHzpc 3.65) [35] and Virgin Granular AC activation by HNO3 (pHzpc 3.0) [36], which are close to the value observed in this study.

Figure 4 shows the SEM image of the palm fiber carbonized at 400 °C and TEM image of the AC after grinding. Figure 4a show that after thermal treatment, the fiber is converted into a skeleton of carbon grains with a size range between 1 and 4 μm. The TEM image of PTF shows a rough surface and porosity (Figure 4a). Meanwhile, the SEM image of raw biomass PTF is reported in ref. [33]. The PTF has a rough surface with no evidence of porosity and is covered by arrays of protrusions, which can be Si particles. The porosity of carbon is due to the decomposition of lignin, cellulose, and hemicellulose during carbonization, resulting in the formation of micropores and mesopores [37,38]. The cell structure of the PTFAC can be seen clearly by TEM. After grinding, the particles were converted into rounded and edged grains with a size range of 20–50 nm (Figure 4b).
Furthermore, the TEM image of raw biomass PTF indicates that the cell wall structure of the fiber is almost round [36].

![Figure 3](image3.png)

**Figure 3.** (a) Zeta potential and (b) Point of zero charge of PTFAC.

![Figure 4](image4.png)

**Figure 4.** (a) SEM image of palm tree fiber carbonized at 400 °C and (b) TEM image of PTFAC.

The examination of the prepared AC before and after the adsorption of dyes by FTIR can give better insight into the expected attraction forces between the PTFAC and dyes. Figure 5 shows the FTIR spectra of the PTF and after the adsorption of CR and RhB dyes by PTFAC. As shown in Figure 5, the following function groups are detected in the spectra: the band observed at around 3335 cm\(^{-1}\) in the fiber and PTFAC spectra was attributed to the O–H stretching of the hydroxyl group of alcohol present in cellulose and lignin, and
the band at 1625 cm\(^{-1}\) was attributed to O–H bending of adsorbed water [39,40]. The band at 2900 cm\(^{-1}\) was ascribed to the unsaturated C–H stretching vibration of the alkene or aromatic groups [39]. The band observed at around 2925 cm\(^{-1}\) in the spectrum of fiber was attributed to C–H vibration of organic matters in the fiber, which had nearly vanished in the PTFAC, indicating that most of the elements such as hydrogen and oxygen are removed after activation [16,41]. The bands at around 1605–1620 cm\(^{-1}\) were attributed to the C=C stretching vibration in the aromatic rings [42]. The bands between 1238.81 cm\(^{-1}\) and 1032.31 cm\(^{-1}\) were attributed to the C–O stretching in carboxylic groups present in cellulose, typically originating from the oxidized carbon [18,20]. The FTIR spectrum of PTFAC demonstrates significant change by lowering peaks intensities when contrasted to the PTF spectrum. The bands between 1220 and 1118 cm\(^{-1}\) can also ascribed to phosphorous species, such as hydrogen-bonded P=O, stretching vibrations O–C in P–O–C of aromatic, P=OOH, and polyphosphate chain P–O–P [29,31]. In addition, the band around 1032.31 cm\(^{-1}\) assigned to the silicon atom initially attached to the oxygen atom Si–O [43]. The bands observed at around 470 cm\(^{-1}\), the spectra of carbons, were assigned to the siloxane Si–O–Si group [43]. When the surface of PTFAC was covered by dyes, most of its characteristic bands disappeared or mostly evanesced due to the adsorption of dye on its surface. For example, O–H, C=C, and C–O bands and all types of phosphorous species bands and Si–O–Si groups disappeared. This observation enhances the electrostatic attraction forces between the adsorbent and adsorbed dyes through noncovalent interactions, which may include dipole–dipole interaction, intermolecular forces, and hydrogen bonding interaction.

![Figure 5](image-url)

**Figure 5.** The comparison of Fourier transform infrared spectroscopy (FTIR) spectra of palm tree fiber (PTF) before and after the activation and after adsorption of Rhodamine B (RhB) and Congo red (CR) dyes by PTFAC.

### 3.2. Optimization of Preparation Conditions for PTFAC

To optimize the preparation conditions of PTFAC, adsorption trial tests were carried out using PTF activated with different chemical agents such as H\(_3\)PO\(_4\) (85%), H\(_2\)SO\(_4\) (98%), and KOH (2 M) with an impregnation ratio of 3:1 (chemical/biomass). In addition, adsorption trial tests were conducted on the AC prepared at different temperatures at various times. Figure 6 shows the removal percentage of CR and RhB dyes by PTF activated by H\(_2\)SO\(_4\), H\(_3\)PO\(_4\), and KOH. From the mentioned dyes, the fiber activated by H\(_2\)SO\(_4\) showed the highest removal percentage; hence, the H\(_2\)SO\(_4\) is considered a strong oxidizing
and dehydrating agent. As a result of the strong acid and base, H$_2$SO$_4$ and KOH exert a high corrosive effect on the precursor that produces weight loss of the precursor, which is less than that of the raw material. Indeed, the activation of AC by H$_3$PO$_4$ improves the pore structure during the thermal treatment, which improves the porosity in terms of macropores and mesopores of the PTFAC [8,44]. Thus, the activation by H$_3$PO$_4$ is highly recommended for functionalization of the surface of the used bioadsorbent that has an environmentally friendly in nature and easy to recover and recycle properties [17,45].

![Graph](image)

**Figure 6.** The removal percentage of CR and RhB dyes on the synthesized PTFAC with different (a) chemical activation reagents, (b) activation temperature of CA-PTF-H$_3$PO$_4$, and (c) activation times of CA-PTF-H$_3$PO$_4$ at 400 °C.

The removal percentage of dyes, adsorbed on PTFAC prepared at different temperatures and times, are shown in Figure 6b,c, respectively. Thermal treatment data revealed the chemical activation of AC with H$_3$PO$_4$ at 400 °C for 30 min in a facile and fast short analytical time process for AC activation. Thus, it can be concluded that the cellulose and hemicellulose in the PTF gradually decompose at 400 °C, resulting in a high specific surface area. On the other hand, a long activation time causes degradation and decreases
the surface area and porosity of carbons, damaging the pores because of the excessive reaction with other activated agents. Hence, chemical activation with H$_3$PO$_4$ at 30 min is adopted in the subsequent study for dyes removal in batch mode.

3.3. Influence of Parameters on the Adsorption Process (pH, Duration, Initial Concentration, Adsorbent Dosage)

The batch experiment was done to study the influence parameters using $C_0 = 25$ mg L$^{-1}$, PTFAC mass = 0.1 g, V = 20 mL, $t = 25^\circ$C. In the adsorption process, the charge on the adsorbent and the characteristics of adsorbate in solution are determined by the pH, which has a radical impact on the adsorption ability. Figure 7a shows the effect of pH between (2.0–12) on the removal percentage of CR and PhB dyes by PTFAC. Upon increasing the pH values, the removal percentage declined slightly from 98.3% to 95.2% in the case of CR and from 99.9% to 99.6% for RhB; apparently, the maximum adsorption is observed at pH 2.0. As the zero-point charge of the adsorbent is at pH 3, it means that the adsorption is high when pH is less than $p_{Hzpc}$ and low when pH is more than $p_{Hzpc}$. However, the cation dye (RhB) and anion dye (CR) at different pH values show high percentage removal, which makes the PTFAC significant in the removal of wastewater, because dyes in wastewater are usually in a wide pH range. It is well known that the adsorption of dyes by the AC can proceed by van der Waals, electrostatic, and H-bonding interactions. The CR is a dipolar molecule; it exists in an anionic form with red color at basic pH and in a cationic form with blue color at acidic pH. It is also observed that the surface of AC is covered by phenolic and carboxylic groups due to oxidation from the atmosphere and water medium [46]. At lower pH, protonation of phenolic (PhOH) and carboxylic acid (COOH) groups occurs on the surface of PTFAC, which leads to an increase in electrostatic interactions with the cationic CR dye; therefore, the adsorption of CR on the AC surface is higher. However, at a higher pH, the carboxylic and phenolic groups on the surface of AC are expected to be completely ionized to carboxylate (COO$^-$) and phenoxide (PhO$^-$) ions. Thus, the electrostatic repulsion between the anionic–CR and anionic–AC surface decreases the adsorption capacity. In the case of RhB, the pH has no significant effect on the removal percentage. The RhB may exist in an anionic form with red color at basic pH and in a cationic form with blue color at acidic pH. It is also observed that the surface of AC is covered by phenolic and carboxylic groups due to oxidation from the atmosphere and water medium [46]. At lower pH, protonation of phenolic (PhOH) and carboxylic acid (COOH) groups occurs on the surface of PTFAC, which leads to an increase in electrostatic interactions with the cationic CR dye; therefore, the adsorption of CR on the AC surface is higher. However, at a higher pH, the carboxylic and phenolic groups on the surface of AC are expected to be completely ionized to carboxylate (COO$^-$) and phenoxide (PhO$^-$) ions. Thus, the electrostatic repulsion between the anionic–CR and anionic–AC surface decreases the adsorption capacity. In the case of RhB, the pH has no significant effect on the removal percentage. The RhB may exist in cationic, laconic, or zwitterionic forms depending on the pH of the solution medium. At lower pH, the RhB dye ions are cationic and present in a monomeric molecular form that could easily enter the pore structure of the carbon, resulting in higher adsorption. At higher pH, the RhB dye ions are zwitterionic and present in a dimeric molecular form. The electrostatic interaction between the carboxylate ion and xanthene groups in zwitterionic results in the formation of dimers of bigger molecular forms that cannot easily enter the pores of carbon, resulting in lower adsorption. The adsorption at all pH values in both the dyes shows a very insignificant variation, indicating no ion exchange has occurred. The physical forces such as van der Waals, hydrogen bonding and dipole–dipole interactions might participate in this developed adsorption [30]. Similar studies on the effect of pH at acidic media in removing RhB and CR dyes by other adsorbents reported that RhB and CR dyes have maximum adsorption at pH 3.0 [47].

The duration plays an important role in the adsorption process of RhB and CR dyes by PTFAC. Figure 7b shows that the removal percentage increases with the increase in the duration, with high values of percentage between the range (5.0–120 min). The dyes recorded a high absorption rate; their values were between 95.1% at 5 min and 99.5% at 120 min for CR dye, whereas they were 99.7% at 5.0 min and 99.9% at 120 min for RhB dye. It indicated that the PTFAC has a high efficiency in removing both adsorbed dyes. This is because of the very rapid adsorption that was found during the first 5.0 min, and even then, the amount of the adsorbate increased with time and reached a constant value of the maximum adsorption at 120 min. However, the availability of vacant active sites on the surface of the PTFAC is the result of the rapid increase in adsorption during the initial stage. Eventually, the process slows down due to the diffusion of dyes into the pores of the PTFAC, because the external sites were fully occupied [8]. A similar study reported an increase in the quantity of adsorption dyes with increased contact time. Using an initial concentration of 1.0 g L$^{-1}$, it was found that Raphia hookerie fruit epicarp could adsorb
312 mg/g RhB dye at acidic pH, and equilibrium was attained at 40 min for an initial dye concentration of 400 mg L$^{-1}$ [47].

![Figure 7. Influence of (a) pH, (b) contact time, (c) initial concentration, and (d) adsorbent dosage on elimination the RhB and CR dyes from solution by PTFAC (test conditions: $T = 298$ K, $V = 20$ mL).](image)

The effect of the initial concentration parameter is significant in determining the effectiveness of the adsorbent. The influence of initial dye concentration 5.0 to 50 mg L$^{-1}$ on the adsorption capacity of PTFAC is shown in Figure 7c. The adsorption capacities ($q_e$) were increased with the increase in the initial dye’s concentration from 5.0 to 50 mg L$^{-1}$. The adsorption capacities were between 0.99 and 9.99 mg g$^{-1}$ and 9.51 and 26.83 mg g$^{-1}$ for CR and RhB days, respectively. After 25 mg L$^{-1}$ initial concentration, no significant change was observed in the adsorption capacity; thus, the adsorption process reached the equilibrium. Initially, the adsorption process starts from the boundary layer; then, it diffuses into the adsorbent surface, and finally diffuses into the porous structure of the adsorbent [8]. In a similar study, an increase in the quantity of the dye adsorbed was observed on increasing the initial concentration of 1.0 g L$^{-1}$ of R. hookerie fruit epicarp for adsorption capacity with 43.5 mg g$^{-1}$ of RhB dye for an initial dye concentration of 50 mg L$^{-1}$, while it can have a high adsorption of 312 mg g$^{-1}$ of RhB dye for an initial dye concentration of 400 mg L$^{-1}$ [47].

The influence of adsorbent dosage (0.05 to 0.25 g) is shown in Figure 7d. The figure shows that the removal amount of RhB and CR dyes by PTFAC increased with the increase in the adsorbent dosage. Then, the equilibrium was attained after 0.1 g of PTFAC. There was not much increase in the removal percentage of the dye. The percentage adsorbed of CR dye increased from 97% (9.69 mg g$^{-1}$) for the adsorbent dosage of 0.05 g to 99% (1.99 mg g$^{-1}$) for the adsorbent dosage of 0.25 g. The adsorption capacity of RhB dye increased from 99.6% (9.96 mg g$^{-1}$) to 99.9% (1.99 mg g$^{-1}$) with an increase in the adsorbent dosage of 0.05 to 0.25 g, respectively. The high removal efficiency with a high amount of adsorbent (0.05–25 g) is due to the availability of binding sites for adsorption. The adsorption capacity ($q_e$) of the adsorbent was found to decrease with an increase in the adsorbent dosage, which could be due to the interaction of adsorbent particles such as aggregation or agglomeration, resulting in the decrease of the total surface area [8]. A similar study reported the increase
in the percentage of adsorbed dye on increasing the adsorbent dosage [48]. Another study reported the RhB dye concentration to be 100 mg L\(^{-1}\); at acidic pH, the percentage adsorption of RhB dye increased from 85.19% to 88.88% as the adsorbent dosage increased from 1.0 to 5.0 mg L\(^{-1}\) [47].

3.4. Adsorption Isotherms

Adsorption isotherm studies are very important as they describe how the adsorbed molecules are distributed between the PTFAC and the CR and RhB dyes in the solutions at equilibrium. They offer essential designing of the adsorption process by evaluating the adsorption effectiveness of the adsorbent. In this work, adsorption equilibrium data of CR and RhB dyes on PTFAC were analyzed by Langmuir and Freundlich adsorption isotherm models. The Langmuir isotherm model is the most extensively used system, which is based on a perfect hypothesis that adsorption occurs on the adsorbent at exact homogeneous sites and can be useful for monolayer adsorption [49]. The linear equation of the Langmuir isotherm model is presented below Equation (3).

\[
\frac{C_e}{q_e} = \frac{1}{q_{K_l}} + \frac{1}{q_{\text{max}}}C_e
\]  

(3)

where \(q_{\text{max}}\) is the maximum adsorption capacity and \(K_l\) (L/mg) is the Langmuir constant. \(q_{\text{max}}\) and \(K_l\) can be calculated by drawing \(C_e/q_e\) (specific adsorption) against \(C_e\) (Figure 8a). An additional important factor of Langmuir isotherm is the dimensionless separation factor \(R_l\), which provides an understanding on the favorability of the adsorption process. This has been given in Equation (4).

\[
R_l = \frac{1}{1 + K_l C_0}
\]  

(4)

In the adsorption process, the separation factor \(R_l\) specifies whether the adsorption isotherm is unfavorable \((R_l > 1)\), satisfactory \((0 < R_l < 1)\), irreversible \((R_l = 0.0)\), or linear \((R_l = 1.0)\) [6,15]. The Freundlich isotherm model measures the adsorption capacity and intensity of the adsorbate on heterogeneous surfaces, but it does not predict the maximum adsorption [50]. The linear form of the Freundlich isotherm is expressed by Equation (5).

\[
\log (q_e) = \log (K_f) + \frac{1}{n} \log (C_e)
\]  

(5)

As \(K_f\) and \(1/n\) are the Freundlich constants, they help calculate the capacity and strength of adsorption, respectively. \(K_f\) and \(n\) can be obtained from the intercept and slope by drawing \(\log q_e\) versus \(\log C_e\), respectively (Figure 8b). The \(n\) value gives the degree of nonlinearity between the concentration of the solution and adsorbent. When \(n = 1.0\), the adsorption is linear, whereas \(n < 1\) suggests that the adsorption process is chemical. However, if \(n > 1\), it means that the physical adsorption is satisfactory [15].

The Langmuir and Freundlich isotherms selected to explicate the interaction of CR and RhB dyes with PTFAC are shown in Figure 8. Figure 8a shows the Langmuir isotherm; as the linear correlation coefficient value (\(R^2\)) represents the accuracy of the model, it is found near the unity. The \(R^2\) values for the CR were 0.996, and for RhB, these were 0.980. The Freundlich isotherm in Figure 8b shows \(R^2\) values for CR and RhB to be 0.8575 and 0.8541, respectively. Thus, this suggests that the adsorption values of CR and RhB dyes are the best fit to the Langmuir adsorption model [49]. Based on the calculated data, the coefficients of Langmuir and Freundlich isotherms models are illustrated in Table 2. The Langmuir model shows a maximum adsorption capacity for CR dye at 10.4 mg g\(^{-1}\) and for RhB dye at 26.5 mg g\(^{-1}\) by PTFAC. The dimensionless separation factor \(R_l\) value found for CR dye was 0.04 and that for RhB dye was 0.034, indicating a favorable adsorption [6,15]. In addition, the highest value of the constant \(K_f\) for CR was 81.91 L mg\(^{-1}\), and for RhB, it was 1264.5 L mg\(^{-1}\), which indicates a strong bonding of dyes onto PTFAC [30]. In this...
study, the value of n for RhB adsorption was above unity, indicating favorable physical adsorption, but the value of n for CR was below unity, indicating chemical adsorption.

![Figure 8. Adsorption isotherms: Langmuir model of (a) CR dye; (b) RhB dye and Freundlich model for the adsorption of (c) CR dye; (d) RhB dye by PTFAC (reaction conditions: T= 298 K, pH = 2.0, $C_o = 25$ mg L$^{-1}$, mass = 0.15 g, V = 20 mL, shaking time = 120 min).]

Table 2. Adsorption isotherm parameters calculated for RhB and CR dyes adsorbed by PTFAC$^*$.  

<table>
<thead>
<tr>
<th>Adsorption Isotherm</th>
<th>Parameters</th>
<th>CR</th>
<th>RhB</th>
</tr>
</thead>
<tbody>
<tr>
<td>Langmuir</td>
<td>$q_{max}$ (mg g$^{-1}$)/$C_o$ (25 mg L$^{-1}$)</td>
<td>10.4</td>
<td>26.5</td>
</tr>
<tr>
<td></td>
<td>$K_L$ (L mg$^{-1}$)</td>
<td>81.91</td>
<td>1264.5</td>
</tr>
<tr>
<td></td>
<td>$R_L$</td>
<td>0.04</td>
<td>0.034</td>
</tr>
<tr>
<td></td>
<td>$R^2$</td>
<td>0.996</td>
<td>0.980</td>
</tr>
<tr>
<td></td>
<td>$\chi^2$</td>
<td>0.013</td>
<td>$1 \times 10^{-3}$</td>
</tr>
<tr>
<td>Freundlich</td>
<td>$K_f$ (mg g$^{-1}$) (mg L$^{-1}$)</td>
<td>0.24</td>
<td>$3.07 \times 10^2$</td>
</tr>
<tr>
<td></td>
<td>$1/n$</td>
<td>-1.43</td>
<td>1.357</td>
</tr>
<tr>
<td></td>
<td>$R^2$</td>
<td>0.857</td>
<td>0.854</td>
</tr>
<tr>
<td></td>
<td>$\chi^2$</td>
<td>4.131</td>
<td>$3 \times 10^{-3}$</td>
</tr>
</tbody>
</table>

* Reaction conditions: T= 298 K, pH = 2.0, $C_o = 25$ mg L$^{-1}$, mass = 0.15 g/ 20 mL, shaking time = 120 min.

In addition, the chi-square test was considered to calculate the best fitted model and validation of adsorption kinetics [51], using Equation (6).

$$\chi^2 = \sum \left( \frac{q_{e,exp} - q_{e,cal}}{q_{e,cal}} \right)^2$$  \hspace{1cm} (6)

where $q_{e,exp}$ and $q_{e,cal}$ are the experimental and calculated data of adsorption capacity at equilibrium (mg g$^{-1}$) for each model. This error equation is used to express the distributed error among the calculated and experimental values of the isotherm model. When the error is minor, it proves that the isotherm model is well fitted. According to the $\chi^2$ analysis, it was found that the Langmuir isotherm model showed a lower $\chi^2$ value for CR dye (0.013) and RhB dye (0.001), as depicted in Table 2, suggesting that the adsorption of both dyes
was monolayer type. The previous study reported that the acid-activated red mud can adsorb the CR dye, and the Langmuir isotherm model is the best fit because the chi-square was low [52].

3.5. Adsorption Kinetics

Kinetic investigations offer an important indication to realize the adsorption dynamics or adsorption mechanism using pseudo-first-order and pseudo-second-order models. The mathematical term of the pseudo-first-order model [53] is expressed in Equation (7).

\[
\log(q_e - q_t) = \log q_e - \frac{k_1}{2.303} t
\]

where \(q_t\) (mg g\(^{-1}\)) and \(q_e\) (mg g\(^{-1}\)) are the quantities of dyes adsorbed at time \(t\) and at equilibrium, respectively. \(k_1\) is the rate constant of the pseudo-first-order model (min\(^{-1}\)). The plot of \(\log(q_e - q_t)\) versus \(t\) for the adsorption of CR and RhB dyes on PTFAC results in linear graphs with a negative slope (Figure 9a); \(k_1\) and \(q_e\) are determined from the slope and intercept, respectively. In this case, the calculated \(q_e\) (CR 0.408, RhB 0.006 mg g\(^{-1}\)) values of both dyes were considerably different from the \(q_{exp}\) values. This suggests that the data do not fit well with the first-order model.

![Figure 9](image)

**Figure 9.** Adsorption kinetics: pseudo-first-order model of (a) CR dye; (b) RhB dye and pseudo-second-order model for the adsorption of (c) CR dye; (d) RhB dye by PTFAC (reaction conditions: \(T = 298\) K, pH = 2.0, \(C_o = 25\) mg L\(^{-1}\), mass = 0.15 g, \(V = 20\) mL).

The pseudo-second-order model describes that the adsorption process is a chemical process. It describes the proportion between the rate constant and the engaged active sites. The linear equation of the pseudo-second-order rate model is given in Equation (8).

\[
\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{t}{q_e}
\]
where \( k_2 \) is the pseudo-second-order model adsorption rate constant. The drawing of \( t/q_t \) against \( t \) provides linear graphs. The values of \( q_e \) and \( k_2 \) can be determined from the slope and intercept of the PTFAC graph (Figure 9b).

The pseudo-first-order model in Table 3 shows that the calculated \( q_e \) value is 0.408 mg/g for CR dye and 0.006 mg g\(^{-1}\) for RhB dye. The values vary significantly lower than the experimental \( q_e \) values, which are 4.972 mg g\(^{-1}\) for CR dye and 4.999 mg g\(^{-1}\) for RhB dye. However, with the pseudo-second-order model, the calculated \( q_e \) values are 4.992 mg g\(^{-1}\) for CR dye and 4.998 mg g\(^{-1}\) for RhB dye, which are similar to the experimental \( q_e \) values. In addition, the obtained \( R^2 \) values are very close to ones for the pseudo-second-order kinetics adsorption process, which were 0.999 and 1.0 for CR dye and RhB dye, respectively. The \( \chi^2 \) test was used to validate the kinetics models [47,54]. As \( \chi^2 \) values are generally found to be low, the value for CR dye was \( 7.74 \times 10^{-5} \), and for RhB dye, it was \( 2.01 \times 10^{-8} \). This indicated that the adsorption kinetics data obey the pseudo-second-order model. This is similar to a report on the adsorption of RhB dye onto the area-modified R. hookerie epicarp [47].

### Table 3. Adsorption kinetics parameters calculated for RhB and CR dyes adsorbed on PTFAC*

<table>
<thead>
<tr>
<th>Kinetic Models</th>
<th>Parameters (mg g(^{-1}))</th>
<th>CR</th>
<th>RhB</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( q_e,\text{cal} )</td>
<td>0.408</td>
<td>0.006</td>
</tr>
<tr>
<td></td>
<td>( k_1 ) (1/min)</td>
<td>–0.031</td>
<td>–0.023</td>
</tr>
<tr>
<td></td>
<td>( R^2 )</td>
<td>0.90</td>
<td>0.72</td>
</tr>
<tr>
<td></td>
<td>( \chi^2 )</td>
<td>51.05</td>
<td>3732.3</td>
</tr>
<tr>
<td>Pseudo first order</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>( q_e,\text{cal} )</td>
<td>4.992</td>
<td>4.998</td>
</tr>
<tr>
<td></td>
<td>( k_2 ) (g mg(^{-1}) min(^{-1}))</td>
<td>( 1.35 \times 10^{-3} )</td>
<td>( 9.75 \times 10^{-3} )</td>
</tr>
<tr>
<td></td>
<td>( R^2 )</td>
<td>0.999</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>( \chi^2 )</td>
<td>7.744 \times 10^{-5}</td>
<td>2.011 \times 10^{-8}</td>
</tr>
<tr>
<td>Pseudo second order</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>( q_e,\text{exp} )</td>
<td>4.972</td>
<td>4.999</td>
</tr>
</tbody>
</table>

* Reaction conditions: \( T = 298 \) K, 0.15 g/20 mL, \( C_0 = 25 \) mg L\(^{-1}\).

The adsorption of CR and RhB dyes on PTFAC is an intricate process that involves physical forces acting through pores contained and chemical bonding derived from the active groups on the PTFAC adsorbent [43,54]. Activation by phosphoric acid also enhances the acidic-containing functional groups on PTFAC, resulting in better dye removal from the aqueous solution by the used adsorbent [33]. FTIR analysis revealed the existence of hydrogen bonds between CR, RhB, and the surface of PTFAC. The H bonding interaction occurred between hydroxyl groups at the PTFAC surface and nitrogen-containing functional groups of CR and RhB in acidic medium in addition to the specific attraction between negatively charged \( \text{COO}^- \) and \( \text{SO}_3^- \) groups of CR dye and the positively charged PTFAC surface [47,55]. Protonation of –OH and –COOH groups present at the PTFAC surface in addition to the van der Waals force also enhances the uptake of the used analytes [47,55].

### 3.6. Desorption and Regeneration Studies

Desorption studies help to explain the adsorption mechanism in the recovery of the adsorbent and to investigate the possibility of regeneration of used PTFAC loaded with RhB and CR dyes. Table 4 shows that the desorption using distilled water by adjusting pH to 2.5, 4.5, 6.5, 8.5, and 10.5 has a high desorption efficiency and is very close (in percentage) to the CH\(_3\)COOH and NaOH (0.1, 0.5, and 1.0 M). Thus, the PTFAC loaded with RhB dye showed a high stability in the adsorption process. The desorption percentage at pH values of 2.5, 4.5, 6.5, 8.5, and 10.5 was 99.88%, 99.84%, 99.84%, 99.81%, and 99.77%, respectively, whereas in the desorption of CR loaded on PTFAC, the percentage decreased with increasing the pH from 2.5 to 10.5, which was 98.68%, 97.36%, 96.1%, 93.42%, and 93.42%, respectively. Hence, the optimum desorption percentage of CR and RhB was observed at pH 2.5 using distilled water as a green desorption reagent. This indicates that the adsorption of the dyes is mainly due to the Van der Waals, electrostatic, and H-bonding interactions, and
chemisorption mechanism [48]. In addition, the regeneration study of the adsorbent might yield an efficient economical treatment process. Figure 10 shows the regeneration cycle of PTFAC with CR and RhB dyes. It has no significant change of percentage removal in CR dye. The removal percentage was between 98.68 and 96.1%, which indicated that the use of PTFAC could be increased to more than five cycles. Thus, PTFAC showed high resistance to adsorption efficiency after five regeneration cycles, whereas there was a decrease in percentage removal in RhB dye with the increase of the number of cycles from the first cycle (99.88%) to the fifth cycle (54.49%). That was due to the effect of the repeated use of the PTFAC on the adsorption process. These dyes loaded on the PTFAC surface can change superficial structures and result in the loss or blockage of the adsorption sites in the AC. The magnetic lignosulfonate FCS studied two regeneration cycles for CR, and removal was 83% after five cycles [56]. It can be concluded that PTFAC is an efficient high-potential adsorbent for the removal of cationic and anionic dyes and can be recycled several times.

Table 4. Desorption percentage of CR and RhB dyes by PTFAC with different desorption reagents.

<table>
<thead>
<tr>
<th>Desorption Reagent</th>
<th>CR</th>
<th>RhB</th>
</tr>
</thead>
<tbody>
<tr>
<td>CH₃COOH (0.1 M)</td>
<td>90.78</td>
<td>99.69</td>
</tr>
<tr>
<td>CH₃COOH (0.5 M)</td>
<td>92.10</td>
<td>99.73</td>
</tr>
<tr>
<td>CH₃COOH(1 M)</td>
<td>89.47</td>
<td>99.65</td>
</tr>
<tr>
<td>NaOH (0.1 M)</td>
<td>93.42</td>
<td>99.77</td>
</tr>
<tr>
<td>NaOH (0.5 M)</td>
<td>92.10</td>
<td>99.81</td>
</tr>
<tr>
<td>NaOH (1 M)</td>
<td>92.10</td>
<td>99.84</td>
</tr>
<tr>
<td>H₂O (pH 2.5)</td>
<td>98.68</td>
<td>99.88</td>
</tr>
<tr>
<td>H₂O (pH 4.5)</td>
<td>97.36</td>
<td>99.84</td>
</tr>
<tr>
<td>H₂O (pH 6.5)</td>
<td>96.05</td>
<td>99.84</td>
</tr>
<tr>
<td>H₂O (pH 8.5)</td>
<td>93.42</td>
<td>99.81</td>
</tr>
<tr>
<td>H₂O (pH 10.5)</td>
<td>93.42</td>
<td>99.77</td>
</tr>
</tbody>
</table>

Figure 10. Regeneration cycle of PTFAC after adsorption by RhB and CR dyes.

3.7. Environmental Water Treatment

This study investigated the efficiency of PTFAC as an environmental application by determining the highest percentage removal of CR and RhB dyes. The environmental water samples were filtrated through No. 2 Whatman Qualitative filter paper; then, the spike method was used by obtaining a 15 mg L⁻¹ dyes concentration. Table 5 shows the percentage removal of CR and RhB dyes from contaminated water. Thus, the percentage removal values for RhB dye were 99.8% for tap water, 99.3% for well water, 99.4% for seawater, and 98.3% for sewage water from the treatment plant. The percentage removal values for CR dye were 99.8% for tap water, 94.4% for well water, 93.3% for seawater,
and 92.3% for sewage water from the treatment plant. This indicates the PTFAC has high
stability to remediate water from CR and RhB dye. The removal percentages from the
highest to the lowest are tap water > well water > seawater > sewage water from the
treatment plant. These results confirmed that PTFAC could be used to treat RhB and CR
wastewater in the environmental application without adjusting the pH of the wastewater.
Thus, the PTFAC has high removal efficiency in the remediation of contaminated water
from anion and cation dyes.

Table 5. Adsorptive percent removal of CR and RhB dyes from real water samples by PTFAC.

<table>
<thead>
<tr>
<th>Dyes</th>
<th>Tap Water</th>
<th>Well Water</th>
<th>Seawater</th>
<th>Sewage Water Treatment Plant</th>
</tr>
</thead>
<tbody>
<tr>
<td>% Removal of RhB</td>
<td>99.8</td>
<td>99.3</td>
<td>99.4</td>
<td>98.3</td>
</tr>
<tr>
<td>% Removal of CR</td>
<td>99.8</td>
<td>94.4</td>
<td>93.3</td>
<td>92.3</td>
</tr>
</tbody>
</table>

4. Conclusions
In the current study, palm fiber was successfully used as solid waste for the production
of AC activated by H$_3$PO$_4$ at 400 °C after 30 min. The established method is a facile and fast
method that can be used as an efficient adsorbent for the elimination of anionic and cationic
dyes from wastewater. The texture and surface characteristics of the prepared PTFAC
showed 648.90 m$^2$g$^{-1}$ surface area, 2.83 cm$^3$g$^{-1}$ total pore volume, and 0.79 nm mean pore
diameter with a zeta potential value of $-28.3$ mV and point of zero charge (pHzpc) equal
to 3. The FTIR study indicated the presence of various functional groups. The results of
the adsorption process of the prepared PTFAC showed that pH, adsorbent dosage, contact
duration, and initial dye concentration have a dominant effect on the adsorption of CR and
RhB dyes. The Langmuir isotherm showed a maximum monolayer CR and RhB adsorption
capacity of 10.4 mg g$^{-1}$ and 26.5 mg g$^{-1}$, respectively, of an initial concentration 25 mg L$^{-1}$
at pH 2.0, 25 °C, 120 min, and 0.1 g/20 mL adsorbent dose of AC. The kinetic studies
showed that the adsorption process fits in with the pseudo-second-order kinetics. The
desorption and regeneration studies of adsorbent were found to be effective until the fifth
regeneration cycle by using distilled water an eco-friendly desorption reagent at pH 2.5,
suggesting that the adsorbent could be regenerated and reused. Hence, the investigation
concludes that the PTFAC prepared from PTF is a cost-effective and promising adsorbent
toward dyes from environments.

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review & editing; S.A., methodology, formal analysis, writing—original draft; E.A.B., supervision,
validation; M.F.Z., investigation, formal analysis. All authors have read and agreed to the published
version of the manuscript.

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