Production and Characterization of Activated Carbon Derived from Costus Afer Leaves (C. afer) for the Adsorption of Methylene Blue Dye from an Aqueous Solution

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Abstract

In this work, Costus afer leaves (C. afer) was utilized as an agricultural waste material for the synthesis of activated carbon (AC) used for the adsorption of methylene blue dye from its aqueous solution. The raw material was prepared, chemically activated using KHCO$_3$ with an impregnation ratio of 1:3, and later carbonized at a temperature of 700$^\circ$C in an inert N$_2$ atmosphere for 1 hour to produce activated carbon. The proximate analysis of the biomass revealed the percentage of ash content, moisture content, volatile matter, and fixed carbon present in the waste biomass. The raw
Adsorption of methylene blue validated the Langmuir model with an R$^2$ of 0.999 and a Qm of 11.83 mg/g. These results show that activated carbon prepared from Costus afer leaves constitutes an effective low-cost material for the adsorption of Methylene dye from wastewater.

**ABBREVIATIONS**

| AC | Activated Carbon |
| CAL | Costus Afer Leaves |
| KHCO$_3$ | Potassium Bicarbonate |
| ACCAL | Activated Costus Afer Leaves from KHCO$_3$ |
| MB | Methylene Blue |
| ASTM | American Standard for Testing of materials |
| BET | Brunauer Emmett and Teller |
| EDX | Energy Dispersive X-ray |
| FTIR | Fourier Transform infrared Spectroscopy |
| SEM | Scanning Electron Microscopy |
| XRF | X-ray Fluorescence |

**1. INTRODUCTION**

One of the major global concerns is the discharge of toxic effluents into the environment, which poses hazardous threats to life and animals. These effluents contain dangerous contaminants like dyes, heavy metals, and organic compounds that are very harmful to lives when discharged without effective water treatment. Dyes are widely used in textiles, plastics, food, and photoelectrochemical industries. The removal of dyes from textile wastewater is crucial for mitigating environmental pollution by ensuring that wastewater complies with regulatory standards. According to research, Nigeria produces approximately 32 million tons of waste [1]. The textile, cosmetic, and clothing industries use dyes to color their products. The intensive use of organic dyes results in a high proportion of these compounds in wastewater, necessitating solutions for their removal and disposal. Dyes used in textile industries are acidic, basic, azo, and caustic dyes [2]. An important basic dye is Methylene Blue (MB), which is widely used as a coloring agent for wool, cotton, and silk in the textile industry. Methylene blue is a commonly used cationic dye for coloring, which can also cause eye burns in humans and animals, dyspnea, and skin irritation [3]. In humans, extended contact with these dyes can lead to organ failure, diarrhea, heartburn, and tachycardia. These dyes are hazardous to the environment because of their toxicity, posing a major threat to aquatic ecology and living organisms because they are extremely poisonous and carcinogenic [4]. In addition, these dyes are non-biodegradable and harmful to humans when discharged into the environment because of their chemical structure. MB dye has some adverse effects on human health, but it also has potential applications in various fields. Thus, MB concentrations in water bodies have been rigorously controlled by environmental and health protection authorities, and these contaminants are given priority for elimination [5].

Today, innovative and modern water treatment solutions are being developed to manage the release of dangerous contaminants from wastewater. Several techniques such as coagulation, adsorption, solvent extraction, ion exchange, chemical precipitation, evaporation, and membrane filtration technologies have been employed in wastewater treatment at different time points [6-9]. Most of these methods are feasible in large-scale industries: but among these methods, adsorption has been an economically viable approach that can remove multiple contaminants simultaneously using a wide variety of low-cost absorbents [10]. Adsorption is a waste treatment technique used in dye removal. A broad range of adsorbents remove common dyes from various sources.
Examples of these adsorbents include sawdust [11], fly ash [12], agricultural waste [13], graphene [14], and carbon nanotubes [15].

Activated carbon (AC) is widely used as a reliable adsorbent for the adsorption of color/dye from wastewater. AC is a carbonaceous material with an amorphous solid structure that has a high degree of porosity and a well-developed surface area with numerous oxygenated functional groups such as carboxylic acids, phenols, carbonyls, and lactones [16]. Owing to its unique properties, activated carbon, a porous carbonaceous material, has several applications in wastewater treatment, air purification, and desalination [17]. They are commonly prepared and produced from agricultural residues by carbonization and activation [18]. ACs are widely derived from agricultural wastes because of their low cost and economic abundance. These adsorbents can be defined as the substance on which the adsorbate is adsorbed. The adsorbate can be likened to the substance in the solution that is adsorbed by the adsorbent. Various adsorbents have been used in the production of AC from agricultural materials like peanut shells [19], coconut shells [20], cassava peeling [21], snail shells [22], and plantains [23]. Their high carbon content, low ash content, and high volatile matter make them suitable precursors for the production of AC [24].

Corn husk was investigated by Khodaie et al., [25] for the removal of methylene blue from wastewater using ZnCl₂ as an activating agent. The Langmuir isotherm model proved to be valid with an adsorption capacity of 298 mg/g. Groundnut shell chemically activated with H₂SO₄ was investigated by Melroy and Aravinda [26] for the removal of methylene blue from aqueous solution. The activated groundnut shell carbon results showed 95-99% of adsorption. Freundlich isotherm model was applied with an R² of 0.992. Oil palm wastes were investigated by Baloo et al., [27] for the adsorptive removal of methylene blue dye from an aqueous solution. The isotherm model was better represented by the Langmuir model with an R² of 0.979. Bush cane bark powder was investigated by Enenebeaku et al., [28] for the adsorption of methylene blue from an aqueous solution. The adsorption isotherm obeyed the Freundlich isotherm model with an R² of 0.9579 and an adsorption capacity of 23.49 mg/g.

Costus afer (C. afer) is a plant commonly known as ginger lily, spiral ginger, or bush cane. Various online searches have revealed that the plant has been used for traditional and medicinal purposes to treat diabetes, arthritis, stomach aches, and skin diseases [29]. It is abundant in the East region of Nigeria and is used for the aforementioned purposes. In Nigeria, it is known as Okpete in the Igbo language, mbitem in the Efik language, kakii-zuwa in the Hausa language, and tete-egun in the Yoruba language [30-32]. The utilization of C. afer as a precursor for the production of activated carbon remains scarce in literature but yields great potential for this course. Therefore, this study aimed to prepare and produce activated carbon from CAL using KHCO₃ as an activating agent for the adsorption of methylene blue dye from its aqueous solution. The physical and chemical properties of ACCAL were determined using different characterization techniques: and the effects of influencing parameters such as adsorbent dosage, contact time, and pH were studied. The process reported here is important for the conversion of waste into wealth.

2. MATERIALS AND METHODS

2.1 Apparatus and Equipment List


2.2 Raw Materials Collection

Costus afer leaves (C. afer) were used as precursors because of their physical properties and abundance at the Federal University of Technology, Owerri’s School region. These leaves were collected from bushes around Umuchima and transported to the laboratory. The leaves were washed carefully to remove dirt particles and impurities from their surfaces. After washing, the leaves were sun-dried for 3-4 days to eliminate moisture content and dried in an
oven for 4 hours at a temperature of 120°C. The leaves were crushed using a blender. Furthermore, the samples were sieved using a laboratory sieve to obtain very fine particles. The samples were placed in a safe place and labeled (Raw CAL) for further analysis.

2.3 Chemical Activation of CAL with KHCO₃

The chemicals used in this study for activation purposes were of analytical grade. The method applied in the chemical activation of CAL coincided with recent research by Liu et al., [33] with slight modifications. In a crucible, 4g of the dried raw sample was weighed using a weighing balance and mixed in 10g of KHCO₃ using an impregnation ratio of 1:3. The mixture was ground for 30 min to obtain a fine mixture, after which it was poured into a reactor, placed in a tube furnace, and heated to 700°C in an N₂ atmosphere in the absence of air for 1 h. As dark porous structures developed, the carbonized samples were allowed to cool to room temperature. To ensure that all contaminants and traces of leftover salts were removed from the activated samples and that the final products had a pH of 7 (neutral), they were thoroughly cleaned using distilled water (DI). Finally, the activated samples (ACCALKHCO₃) after washing were dried in an oven at 150°C for 3 h [34]. The activated samples were allowed to cool and stored in an airtight container labeled ACCAL-KHCO₃. Fig. 2. shows a schematic of the Costus afer activated carbon preparation.
2.4 Methylene Blue Dye Solution

The chemicals used to prepare the MB solution were of high analytical grade. Methylene Blue dye was purchased from the Zaria Chemical Laboratory, Nigeria. A stock solution of methylene blue dye was prepared by dissolving 0.5g of commercially available methylene blue in 1 L of distilled water to obtain a stock concentration of 50 mg/L. Experimental dye solutions of desired concentrations were prepared by appropriate dilution of the stock solution.

2.5 Proximate Analysis

When samples were heated and handled under specified circumstances, a set of tests known as proximate analysis was used to determine the composition of the accessible macronutrients (Ash content, moisture content, volatile matter, and fixed carbon content) for raw costus afer leaves. The ash content, moisture content, volatile matter and fixed carbon of the material samples was determined via the American Society for Testing and Materials standard method [35-37].

2.5.1 Ash content

2.0 g of dry raw sample was weighed inside a crucible, oven-dried at 105°C, and later introduced into a tube furnace (Gallenhamp) that was then heated to 600°C at a heating rate of 10°C min\(^{-1}\) for 1 hr. The sample was further cooled and weighed in a weighing balance.

\[
\text{Ash content (\%)} = \frac{W_1 - W_2}{W_3} \times 100
\]

Where, \(W_1 = \text{Weight of Crucible + Sample (g)}\), \(W_2 = \text{Weight of Crucible + Sample after (g)}\), and \(W_3 = \text{Weight of Sample (g)}\).

2.5.2 Moisture content

A 2.0 g dried raw sample in a cleaned crucible was oven-dried at 110°C for 3 h until the weight of the samples remained constant. Samples were dried, cooled in a desiccator at room temperature, and then weighed again to calculate the moisture content using the following formula:

\[
\text{Moisture Content (\%)} = \frac{W_1 - W_2}{W_1} \times 100
\]

2.5.3 Volatile matter

A dry sample (1.0 g) was weighed inside a crucible, oven-dried at 105°C, and heated to 600°C for 1 h in a tube furnace (Gallenhamp) at a heating rate of 10°C min\(^{-1}\). The carbonized sample was then allowed to cool to ambient temperature, removed from the furnace, and stored in a desiccator.

2.5.4 Fixed carbon

The Fixed carbon content was obtained by subtracting the percentage of moisture, ash, and volatile matter content of the materials from a hundred percent according to the method of Alongamo et al., [31]:

\[
\text{Fixed carbon (\%)} = 100 - \text{moisture(\%)} - \text{ash content (\%)} - \text{volatile matter(\%)}
\]

2.6 Characterization of Adsorbents

The surface morphology of the adsorbent was analyzed using a scanning electron microscope ((Phenom-World) operated at a voltage of 15 kV. Scanning electron microscopy was used to study the morphology and surface structure of the adsorbent material [38]. SEM images of the raw and activated samples revealed the structure and morphology of the samples. Before scanning the adsorbent, the samples were sputtered on carbon tape and set on a fixed reduction sample holder, and SEM images were obtained. SEM analysis was performed using an energy-dispersive X-ray (EDX) spectrometer to reveal the elemental composition of the samples. BET analysis was used to determine the specific surface area of the raw and activated CAL. The raw and activated samples were degasified to remove gases and impurities. These samples were pretreated in a flow of pure nitrogen gas at 30 ml/min for 1 h at 250°C before surface area measurements [39]. XRF analysis was used to determine the elemental composition of the materials found in the adsorbent [20]. The functional groups of the samples were examined using FTIR Spectroscopy. FTIR spectroscopy was performed using Agilent Technologies (Cary 680 FTIR) [40].
2.7 Batch Adsorption Experiment

Batch adsorption studies were performed at different dosages, contact times, and pH values to evaluate the adsorption parameters of ACCAL-KHCO₃ for the removal of Methylene blue from its aqueous solution. The batch adsorption experiments were performed in a set of 100 ml conical flasks with varying amounts of adsorbent dosage, contact time, and pH. The percentage of methylene blue removal was calculated as follows:

\[
\% \, RE = \frac{C_o - C_i}{C_o} \times 100
\]

Where RE (%) is the MB removal percentage, C₀ is the initial MB concentration (mg/L) in the wastewater, Cᵢ is the MB concentration at time t, (mg/L) [41].

The amount of MB adsorbed per unit mass of on the adsorbents, qₑ, was determined using the equation (3.5) below:

\[
q_e = \frac{C_o - C_e}{m} \times V
\]

Where q (mg/g) is the adsorption capacity, V (L) is the volume of MB solution, C₀ (mg/L) and Cₑ (mg/L) are the initial and equilibrium concentrations of methylene blue, and m (g) is the weight of the adsorbents (g) [42].

2.8 Adsorption Isotherm

Adsorption isotherm models are vital for providing the mechanism of the adsorption process, which is pertinent for the design of the adsorption process. Adsorption isotherms are crucial for comprehending catalytic processes and the adsorption mechanism [43]. They are also employed to optimize the adsorbent mass during the adsorption process and to model experimental equilibrium data [44]. In this study, Langmuir and Freundlich models that express experimental isotherm data were used. The Langmuir isotherm is based on the equilibrium between adsorption kinetics and was originally proposed by Langmuir [45]. The adsorption isotherm is crucial in the determination and optimization of the adsorption system. Scientifically, the adsorption isotherm is a functional expression for the variation of adsorption with the concentration of the adsorbate in the bulk solution. The Langmuir isotherm model is based on the assumption of ideal adsorption in which uptake occurs on the adsorbent surface by monolayer adsorption.

Almasi [46] described the adsorption isotherm as the relationship between the mass of adsorbed dye per unit mass of adsorbent. The linear form of Langmuir isotherms is expressed in given as:

\[
\frac{C_e}{q_e} = \frac{1}{K_L Q_m} + \frac{C_e}{Q_m}
\]

Where Ce is the equilibrium concentration (mg/L), Qₑ is the amount adsorbed at equilibrium (mg/g), Qₘ (mg/g) is the maximum adsorption capacity, and KL (L/mg) is the Langmuir constant. The values of Qₘ max and KL were calculated from the intercept and slope of the linear plot of Cₑ/qₑ against Cₑ using Origin Pro 9.

The Freundlich isotherm was the first adsorption equilibrium equation proposed in the literature (Freundlich [47]. Freundlich isotherm revealed the interaction of dye adsorption on the heterogeneous surface. Freundlich linear equation is expressed as follows:

\[
\ln Q_e = \ln K_F + \frac{1}{n} \ln C_e
\]

Where Ce is the equilibrium concentration (mg/L), Qₑ is the amount adsorbed at equilibrium (mg/g), and KF and n are Freundlich constants.

3. RESULTS AND DISCUSSION

3.1 Proximate Analysis

The results of the proximate analysis for the determination of macronutrients of costus afer leaves are shown in Fig. 3., which reveals, 1.5% ash content, 36.6% moisture content, 29.2% volatile mate, and 36.7% fixed carbon. From the results obtained, CAL exhibited characteristics of a good precursor for the production of activated carbon [48]. These findings were compared to other works of literature in terms of the proximate analysis as tabulated in Table 1.
Fig. 3. The proximate analysis of Costus afer leaves

Table 1. The proximate composition of various adsorbents

<table>
<thead>
<tr>
<th>Raw Adsorbent</th>
<th>Ash Content (%)</th>
<th>Volatile Matter (%)</th>
<th>Moisture Content (%)</th>
<th>Fixed Carbon (%)</th>
<th>References</th>
</tr>
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<tbody>
<tr>
<td>Cassava Peeling</td>
<td>8.0</td>
<td>13.0</td>
<td>1.0</td>
<td>78</td>
<td>Alongamo et al., [21]</td>
</tr>
<tr>
<td>Rumex abyssinicus plant</td>
<td>9.82</td>
<td>18.74</td>
<td>2.95</td>
<td>68.94</td>
<td>Fito et al., [49]</td>
</tr>
<tr>
<td>Coconut Shell</td>
<td>7.78</td>
<td>63.76</td>
<td>12.86</td>
<td>15.58</td>
<td>Khuluk et al., [50]</td>
</tr>
<tr>
<td>Coconut Husk</td>
<td>5.3</td>
<td>41.2</td>
<td>14.2</td>
<td>39.3</td>
<td>Asadu et al., [20]</td>
</tr>
<tr>
<td>Pure Lignin</td>
<td>11.6</td>
<td>36.6</td>
<td>7.9</td>
<td>43.9</td>
<td>Bedmohata et al., [39]</td>
</tr>
<tr>
<td>Costus afer</td>
<td>1.5</td>
<td>29.2</td>
<td>36.6</td>
<td>36.7</td>
<td>This work</td>
</tr>
</tbody>
</table>

Table 2. BET-surface area of different adsorbents

<table>
<thead>
<tr>
<th>Adsorbents</th>
<th>Activating Agent</th>
<th>BET-Surface Area (m²/g)</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Groundnut shell</td>
<td>KOH</td>
<td>691.69</td>
<td>Kumari et al., [52]</td>
</tr>
<tr>
<td>Spathodea campanulata</td>
<td>H₃PO₄</td>
<td>1054</td>
<td>Dimbo et al., [51]</td>
</tr>
<tr>
<td>Rice husks</td>
<td>H₃PO₄</td>
<td>244.479</td>
<td>Sawasdee et al., [53]</td>
</tr>
<tr>
<td>Costus afer leaves</td>
<td>KHCO₃</td>
<td>600</td>
<td>(This work)</td>
</tr>
</tbody>
</table>

3.2 Specific Surface Area

AC produced from CAL (600 m²/g) using KHCO₃ as an activating agent is higher than that of raw CAL (400 m²/g). Activated carbon with a higher surface area and a well-developed pore structure provides more sites for adsorption, leading to increased adsorption capacity [51]. Table 2 compares the specific surface areas of different activated adsorbents with the findings of this work. The increased surface area indicates that the adsorbent has more active sites on its surface, which may improve the large-scale industrial adsorption of contaminants from textile effluent. Therefore, the result of the BET-surface area proves that ACCALKHCO₃ can be commercially applicable for industrial water purification.

3.3 Characterization of Adsorbent

3.3.1 Fourier-transform infrared spectroscopy (FTIR)

The location of the wavenumber (cm⁻¹) in the range 650 to 4000 cm⁻¹ obtained from the results of the Fourier-Transform Infrared (FTIR) spectra for the raw and activated samples is shown in Fig. 4, and their major bands are tabulated in Table 3 From the results obtained from the
spectral analysis of raw CAL, a broad absorption band centered around 3000-3500 cm\(^{-1}\), was observed at 3261.4 cm\(^{-1}\) corresponding to O-H bond stretching characteristic of carboxylic acids, alcohol, or water. The pattern observed in this study is somewhat comparable to that of previous research on the production of activated carbon [54-55]. The raw samples showed peaks between 2800 – 3000 cm\(^{-1}\), with maximum peaks at 2922.2 cm\(^{-1}\) and 2851.4 cm\(^{-1}\) for raw CAL, which indicates the presence of C-H (CH3 and CH2) stretch of the aliphatic hydrocarbon [56]. It is noticed that this broadband decreases after carbonization and leads to the absence of water molecules on the surface of the material. The disappearance of these peaks is visible after carbonization because of the loss of the hydroxyl group and absorbed water [57]. Bands around 2150 cm\(^{-1}\)- 2250 cm\(^{-1}\) correspond to the presence of C≡C Stretching. These peaks were located in the activated samples at 2109.7 cm\(^{-1}\) for ACCAL\(_{KHCO3}\). Sharp peaks were observed in the activated CAL sample spectra at 1900.4 cm\(^{-1}\) indicating cyanide and thiocyanide ion stretches. Broad peaks of the raw CAL at 1636.3 cm\(^{-1}\) are due to the conjugated C=C stretching vibration, which reveals the presence of aromatic carbon-carbon double bonds. It can be observed that the raw samples had more peaks than the activated ones, which implies that they have more functional groups. This agrees with the literature that carbonization and activation could lead to the disappearance of some functional groups [40].

3.3.2 Scanning electron microscopy (SEM) of raw and activated CAL

Scanning electron micrographs of raw and activated CAL are presented in Figs. 5(a), 5(b), 5(c) and 5(d). From the SEM images at low and high magnifications, it can be noticed that the surface morphology of the raw CAL has fine and well-arranged structures. It also contains a smooth and well-packed microstructure with fistulous crystallites, which indicate minerals. These growing crystallites indicate calcium, aluminum, and iron. The SEM micrograph of the activated CAL in Figs. 5(c-d) reveals that the sample is highly porous. The pore structure of the activated CAL is more developed than that of the raw CAL, which is consistent with previous studies showing that activation increases the pore structure of a material [58]. From the images of the activated CAL, as shown in Fig. 5(c-d), the activated samples have increased their porous network, which in turn increases their surface area, making them suitable for adsorption.

3.3.3 Energy dispersive X-ray spectroscopy (EDX) of raw and activated CAL

EDX spectral analysis was used to probe the elemental composition of the raw and activated samples. The energy dispersive X-ray Spectroscopy (EDX) spectral results for Costus af\(\)er (CAL) leaves are presented in Tables. 4 and 5 EDX analysis generated data comprising spectra with peaks corresponding to various components, as shown in Figs. 6(a) and 6(b). EDX spectra taken at 15 Kv, FOV: 537 µm, and a BSD full detector.

3.3.4 X-ray fluorescence (XRF) analysis of raw CAL

X-ray Fluorescence (XRF) analysis for the raw CAL was performed as shown in Fig. 7, confirming that the sample consisted of a good amount of carbon. XRF analysis was performed on the raw sample (CAL) to determine the chemical compounds and concentrations present in the sample. Fig. 7 reveals the dominant compounds and their concentrations in CAL, namely, potassium oxide (51.383%), calcium oxide (23.695%), alumina (5.915%), and silica oxide (4.659%), while other components were found in nominal amounts.

<table>
<thead>
<tr>
<th>S/No</th>
<th>Wavenumber/Frequency (Cm(^{-1})) of Absorption band</th>
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<tbody>
<tr>
<td>1</td>
<td>3261.4</td>
</tr>
<tr>
<td>2</td>
<td>2922.2, 2851.4</td>
</tr>
<tr>
<td>3</td>
<td>2109.7</td>
</tr>
<tr>
<td>4</td>
<td>1636.3</td>
</tr>
<tr>
<td>5</td>
<td>3852.8, 3693.8, 2325.9, 2076.1, 1207.7, 1028.7, etc.</td>
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Fig. 4. FTIR Spectra of: (a) Raw CAL (b) Activated CAL

Table 4. EDX spectrum of CAL

<table>
<thead>
<tr>
<th>Element Number</th>
<th>Element Symbol</th>
<th>Element Name</th>
<th>Atomic Conc.</th>
<th>Weight Conc.</th>
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<td>C</td>
<td>Carbon</td>
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<td>Nitrogen</td>
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<td>K</td>
<td>Potassium</td>
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<td>15</td>
<td>P</td>
<td>Phosphorus</td>
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<td><strong>Total</strong></td>
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Fig. 5. SEM Images of (a) CAL at low magnification (b) CAL at high magnification (c) ACCAL\textsubscript{KHCO₃} at low magnification (d) ACCAL\textsubscript{KHCO₃} at high magnification

<table>
<thead>
<tr>
<th>Element Number</th>
<th>Element Symbol</th>
<th>Element Name</th>
<th>Atomic Conc.</th>
<th>Weight Conc.</th>
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<tr>
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<td>Phosphorus</td>
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<td>S</td>
<td>Sulfur</td>
<td>0.16</td>
<td>0.39</td>
</tr>
<tr>
<td>17</td>
<td>Cl</td>
<td>Chlorine</td>
<td>0.09</td>
<td>0.25</td>
</tr>
<tr>
<td>22</td>
<td>Ti</td>
<td>Titanium</td>
<td>0.02</td>
<td>0.09</td>
</tr>
</tbody>
</table>

| Total          |               | 100          |

3.4 Adsorption Experiment Parameters

3.4.1 Effect of adsorbent dosage

The effect of the adsorbent dosage of activated carbon on the percentage removal of MB is shown in Fig. 8. The adsorbent dosage was varied from 0.1 g to 0.5 g for 10 mL of 50 mg/L solution at a constant time of 10 min and pH of 7. As observed from the graph in Fig. 8, an increase in the dosage increases the percentage removal of MB. The optimum dose of ACCAL is
0.2g with a percentage removal rate of 70%. After the optimum dose is reached, the percentage removal efficiency declines thereafter. These may result from active sites induced by agglomeration and particle clustering. Clustering reduces the number of adsorption sites accessible to an adsorbate from the solution [59].

### 3.4.2 Effect of contact time

The effect of contact time is a vital factor in the percentage removal of MBD from its aqueous solution. For this experiment, an adsorbent dosage of 0.2g and a pH of 7 were kept constant while the time was varied from 5 min to 25 min at a constant MB solution of 10 ml of 50 mg/L. From Fig. 9, it can be seen that an increase in contact time leads to an increase in the removal of MB from its aqueous solution. The optimum time was 10 min, and thereafter, a reduction in the percentage removal rate of MB was observed. The highest percentage of removal (69.9%) was recorded at 10 min. Beyond the optimum time, a gradual decrease in the percentage removal was observed, which may be due to the lack of active sites or dislodged MB molecules [60-62].

### 3.4.3 Effect of pH

Fig. 10 shows the effect of pH on the percentage removal of methylene blue dye at a constant time of 10 min, constant solution of 10 ml of 50 mg/L, and constant adsorbent dosage of 0.2g. The effects of initial pH on the percentage removal of MB dye solution at a concentration of 100 mg/L were studied by varying the pH from 2 to 10. As the pH increases, it tends to increase the percentage of methylene blue removal. The percentage removal of the methylene blue dye from its aqueous solution by ACCAL\textsubscript{KHCO3} was optimum at pH 8, with a percentage removal rate of 66.5%. As shown in Fig. 10, at pH 2–4, there was a slightly constant rate of percentage removal of MB and a steady increase from pH 4 to 6. Above the optimum pH of 8, there was a gradual decrease in the percentage removal of the MB dye [63].

![EDX spectrum of (a) CAL (b) ACCAL\textsubscript{KHCO3}](image_url)
CrossRoads Scientific XRF–FP Analysis Report

File: C:\Users\Tênngườiطف\3D Objects\New folder (49)\RAW COSTUSR.str

1:55:52 AM 14-Nov-23

Comment line

Layer Table ===================================
# Thick Type Error Units Density Norm. Total
1  100 BULK 0.00 mg/cm2 0.00%  On  100.00

Sample Table ===================================

Layer  Component  Type Conc. Error  Units Mole% Error
1  SiO2  Calo  4.459 0.601 wt.%  5.994 0.773
1  V2O5  Calo  0.800 0.000 wt.%  0.000 0.000
1  Cr2O3  Calo  0.938 0.037 wt.%  0.026 0.019
1  MgO  Calo  0.087 0.030 wt.%  0.095 0.033
1  Fe2O3  Calo  1.134 0.049 wt.%  0.349 0.024
1  Co3O4  Calo  0.022 0.026 wt.%  0.007 0.008
1  NiO  Calo  0.011 0.021 wt.%  0.012 0.022
1  CuO  Calo  0.227 0.022 wt.%  0.220 0.022
1  Mn3O4  Calo  0.029 0.018 wt.%  0.040 0.026
1  MO3  Calo  0.911 0.084 wt.%  0.004 0.028
1  PO4  Calo  1.447 0.258 wt.%  1.006 0.140
1  SO3  Calo  4.448 0.749 wt.%  3.908 0.741
1  CaO  Calo  23.095 0.497 wt.%  32.662 0.871
1  MgO  Calo  0.000 0.000 wt.%  0.000 0.000
1  ZnO  Calo  53.393 0.501 wt.%  42.166 0.411
1  BeO  Calo  0.000 0.000 wt.%  0.000 0.000
1  Al2O3  Calo  5.915 1.915 wt.%  4.484 1.492
1  Ta2O5  Calo  0.000 0.000 wt.%  0.000 0.000
1  TiO2  Calo  0.002 0.067 wt.%  0.195 0.065
1  SmO  Calo  0.146 0.019 wt.%  0.138 0.019
1  Ag2O  Calo  0.000 0.092 wt.%  0.010 0.031
1  Cl  Calo  3.105 0.319 wt.%  6.770 0.259
1  SrO2  Calo  0.009 0.017 wt.%  0.006 0.011
1  Sm2O3  Calo  3.401 1.563 wt.%  1.744 0.802

Element Table ===================================

Elkex Line Cond Ratio Intensity Error Intensity Conc. Conc Calibration
Code  C(s)  (d/s)  Method  Coefficient
0  Ka  None  0.000  0.0000 Gaussian  25.491  None  0.000
Mg  Kx  None  0.800  1.7146 Gaussian  0.000  FP  0.000
Zn  Kx  None  10.382  3.9510 Gaussian  1.330  FP  0.000
Si  Kx  None  41.283  5.2070 Gaussian  2.176  FP  0.000
P  Kx  None  52.400  7.3146 Gaussian  0.826  FP  0.000
S  Kx  None  197.775  12.1785 Gaussian  1.621  FP  0.000
Cl  Kx  None  457.799  17.5180 Gaussian  3.155  FP  0.000
K  Kx  None  6336.789  61.8353 Gaussian  42.656  FP  0.000
Ca  Kx  None  1690.762  31.0965 Gaussian  16.935  FP  0.000
Ti  Kx  None  20.153  6.7598 Gaussian  0.123  FP  0.000
V  Kx  None  0.000  0.0000 Gaussian  0.000  FP  0.000
Cr  Kx  None  2.699  0.9031 Gaussian  0.071  FP  0.000
Fe  Kx  None  3.561  3.0315 Gaussian  0.067  FP  0.000
Mn  Kx  None  864.356  8.8031 Gaussian  0.783  FP  0.000
Co  Kx  None  8.827  10.4477 Gaussian  0.016  FP  0.000
Ni  Kx  None  5.672  2.8598 Gaussian  0.009  FP  0.000
Cu  Kx  None  159.716  13.7983 Gaussian  0.181  FP  0.000
Zn  Kx  None  3435.35  12.1460 Gaussian  0.302  FP  0.000
Sr  Kx  None  6.792  8.7062 Gaussian  0.007  FP  0.000
Ba  Kx  None  14.038  9.0679 Gaussian  0.003  FP  0.000
Ac  Kx  None  1.603  4.9255 Gaussian  0.008  FP  0.000

Fig. 7. XRF of raw CAL

Fig. 8. Effect of adsorbent dose on the percentage removal of MB
3.5 Adsorption Isotherm

Figs. 11 and 12 show the Langmuir and Freundlich isotherm plots. For the Langmuir and Freundlich isotherm models, the coefficients ($R^2$) were 0.999 and 0.9519, respectively. The data fit well with the Langmuir isotherm. Langmuir and Freundlich plots for MB adsorption were analyzed for adherence using linear regression. The adsorption was found to be accurately characterized by the Langmuir and Freundlich adsorption isotherm models, with validity from the Langmuir isotherm model, which showed a greater $R^2$ of 0.999. The values of $Q_m$, $R_L$, and $K_L$ were calculated from the slope and intercept of the plot and are presented in Table 6. $R_L$ is the separation factor that determines whether the adsorption process is favorable, unfavorable, or irreversible. If the value of $R_L$ is <0, it indicates an unfavorable adsorption process; $R_L$ between 0 and 1 ($0 \leq R_L < 1$) shows a favorable adsorption, while $R_L > 1$ shows an irreversible adsorption type. According to Malik [64], a value of $1/n < 1$ indicates the presence of new adsorption sites that will lead to an increase in adsorption capacity. In addition, $1/n > 1$ indicates weakness in the adsorption bond and a decrease in the adsorption capacity. Conversely, the Langmuir isothermal feasibility ($R_L$) and Freundlich isotherm constant associated with intensity ($1/n$) were 0.6163 and 0.5117, respectively, indicating a favorable adsorption process.

Similar methods have also been used by other researchers to perform adsorption isotherms 44, [65-66]. The capacity of ACCAL$_{KHCO_3}$ was found to be 11.83 mg/g in the Langmuir model for the removal of methylene blue dye.
Fig. 11. Langmuir modelling plot for adsorption

Fig. 12. Freundlich modelling plot for MB onto ACCAL

Table 6. Adsorption Isotherm plots parameters for MB adsorption onto ACCAL-KHCO3

<table>
<thead>
<tr>
<th></th>
<th>Langmuir - ACCAL</th>
<th>Freundlich - ACCAL</th>
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<tr>
<td>$R^2$</td>
<td>0.999</td>
<td>0.9519</td>
</tr>
<tr>
<td>$Q_{max}$ (mg/g)</td>
<td>11.83</td>
<td>3.506</td>
</tr>
<tr>
<td>$K_L$ (L/mg)</td>
<td>0.062</td>
<td>0.5117</td>
</tr>
<tr>
<td>$R_L$</td>
<td>0.6163</td>
<td></td>
</tr>
</tbody>
</table>

Table 7. Comparison of adsorption capacity of activated carbons (Adsorbent)

<table>
<thead>
<tr>
<th>Adsorbents</th>
<th>Maximum Adsorption Capacity (Qm (mg/g))</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rice husks</td>
<td>9.8</td>
<td>Sharma et al., [66]</td>
</tr>
<tr>
<td>Fig leaves</td>
<td>41.7</td>
<td>Al-Asadi et al., [67]</td>
</tr>
<tr>
<td>Orange peel</td>
<td>18.6</td>
<td>Annadurai et al., [68]</td>
</tr>
<tr>
<td>Cola nut shells</td>
<td>87.12</td>
<td>Ndi et al., [69]</td>
</tr>
<tr>
<td>Coconut shells</td>
<td>15.2</td>
<td>Khuluk et al., [50]</td>
</tr>
<tr>
<td>Bush cane bark powder</td>
<td>23.49</td>
<td>Enebeaku et al., [28]</td>
</tr>
<tr>
<td>Rice husks</td>
<td>26.3</td>
<td>Sawasdee et al., [53]</td>
</tr>
<tr>
<td>Costus afer leaves</td>
<td>11.83</td>
<td>This work</td>
</tr>
</tbody>
</table>

3.6 Comparative Analysis of Other Adsorbents

The comparison of a few adsorbents, including CAL, and their maximum MB adsorption capacity is shown in Table 7. This initial investigation into ACCAL’s adsorption capacity shows that it has great potential for eliminating coloring materials from aqueous solutions. The material’s significant cost reduction and ease of accessibility outweigh its adsorption capabilities, which are somewhat lower than those of many other known adsorbents. These early results call for more research, including the use of acids or bases to activate the adsorbent (CAL), higher impregnation ratio, isotherm studies, and continuous flow operations to assess the adsorption capabilities of ACCAL.

4. CONCLUSION

This work underscores the potential of Costus afer leaves (CAL) waste as a cost-effective, sustainable, and eco-friendly adsorbent for the removal of methylene blue dye from aqueous solutions. The proximate analysis of CAL precursors revealed favorable characteristics, such as high fixed carbon content and low ash content, indicating their suitability for activated carbon production. Characterization of the activated carbon using Fourier transform infrared spectroscopy (FTIR), Scanning electron microscopy (SEM), and Energy dispersive X-ray (EDX) corroborated previous research and provided a comprehensive understanding of the material’s structure and composition. Batch studies demonstrated that the adsorption of methylene blue was influenced by parameters...
such as dosage, contact time, and pH. Optimal conditions for maximum adsorption efficiency were determined to be an adsorbent dose of 0.2g, contact time of 10 minutes, and pH of 8. Under these conditions, activated Costus afer leaves (KHCO$_3$) achieved a remarkable percentage removal rate of 70.58% with a maximum adsorption capacity of 11.83 mg/g for ACCAL$_{KHCO_3}$. These findings highlight the potential of Costus afer leaves as a promising adsorbent for water treatment applications. Further research could explore scaling up the production of activated carbon from CAL and investigating its performance with increasing impregnation ratio, and temperature. Overall, this work contributes to the growing body of knowledge on sustainable solutions for wastewater remediation.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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