REMOVAL OF MALACHITE GREEN FROM AQUEOUS SOLUTION BY ACTIVATED CARBON PREPARED FROM ALMOND SHELL

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ABSTRACT

The objective of this study is to assess the adsorption potential of activated carbon prepared from almond shell for removal of malachite green (MG) from aqueous waste. Almond shell based activated carbon was prepared chemically using ortho-phosphoric acid at 700-800°C in a modified muffle furnace. Prepared activated carbon was characterised for proximate and ultimate analysis, iodine number, N₂-BET surface area, Fourier-Transform Infrared spectroscopy (FT-IR) and Scanning electron microscopy (SEM). FT-IR & SEM results strongly support significant spectral and chemical changes occurring due to the activation of the almond shells by phosphoric acid treatment. Batch isotherm and kinetic study experiments were conducted to determine the adsorption of the malachite green from aqueous solution with almond based activated carbon (ABAC). It is inferred from the results that ABAC is very effective and economically viable adsorbent for removal of malachite green from aqueous solution.

Key words: SEM, FT-IR, Almond shells, Adsorption, Malachite green, Kinetics, Adsorption isotherms.

INTRODUCTION

The discharge of effluents from the industries is one of the potential sources of dyes contamination and pollution1-3. These industries discharge highly coloured effluents with very high Biological oxygen demand (BOD) as well as Chemical oxygen demand (COD)4. Malachite Green is a very popular dye and used extensively in dyeing of silk, leather, plastic & paper. The discharge of malachite green can cause serious harms to aquatic life. It can cause injuries to humans and animals by direct contact, inhalation and ingestion5. Various serious problems such as carciogesis, mutagenesis, tetratogensism, respiratory toxicity & reduced fertility in humans have been reported6. The removal of such dangerous pollutants to permissible limit is mandatory as per environment protection Laws. Agricultural wastes
are the chief raw materials being studied for this purpose, for they are renewable, mostly available in huge amounts & potentially less expensive\textsuperscript{7,8}. Till now, adsorbents such as neem, sawdust, bentonite clay, bottom ash, orange peel, algae, rice straw, coffee bean, Kapok hull, Pandanuts, have been evaluated for the removal of dyes from aqueous solution\textsuperscript{6,9-13}. This study reports the use of almond shell based activated carbon (ABAC) a low cost alternative adsorbent for the removal of malachite green from aqueous solution.

**EXPERIMENTAL**

**Materials and methods**

Malachite green dye GR grade ortho-Phosphoric acid (H\textsubscript{3}PO\textsubscript{4}) GR grade, hydrochloric acid (HCl) GR grade, Merck India, were procured from local market. Almond shells were collected from local market as waste material.

**Preparation of activated carbon**

The almond shells were washed & dried in open air for 2 days then crushed to 10-12 mm size. The crushed shell mass was dried in oven at 108 ± 2°C. For activation process, 10 g of dried raw material was impregnated with H\textsubscript{3}PO\textsubscript{4} in the ratio 1:1 for 24 hrs and carbonised at 700-800°C for 1 hr. The carbonized material was then cooled to room temperature and washed with 5 N HCl thoroughly. To remove excess acid the material was washed with double distilled water. After washing the material was dried at 108 ± 2°C in moisture oven for about 24 hrs. This carbon was kept in a dessicator for further use. A part of Sample was pulverized and passed through 325 mesh for physico-chemical analysis.

**Characterization of activated carbon**

The prepared activated carbon (ABAC) was characterised for proximate, ultimate, Iodine value, N2-BET surface area, pore volume, FT-IR and SEM, Table 1.

| Table 1: Physico-chemical properties of prepared activated carbons |
|-----------------------------------------|---|---|---|---|---|---|---|---|---|
| Carbon | M % | Ash % | V.M. % | F.C. % | C % | H % | N % | Iodine Value | N2-BET surface area | Pore volume |
| ABAC    | 17.06 | 6.24 | 12.4 | 64.3 | 78.4 | 3.5 | 0.78 | 1093 | 1492.3 | 1.14 |

**SEM morphology analysis**

The surface morphology of ABAC is shown in Fig. 1, which shows a very clear view of the different types of pores (micro, semi micro, macro) present, which are formed
during activation process. Some white crystals are also visible in the SEM that may be due to mineral matter present in it.

![Fig. 1: SEM of ABAC](image)

**FT-IR analysis of ABAC**

To resolve the functional groups and its wave numbers, spectral analysis was done for ABAC using Fourier transform infrared spectrometer (FT-IR) (Perkin-Elmer, PE-RXI) in the range of 450-4000 cm$^{-1}$. The FT-IR spectrum of ABAC as shown in Fig. 2 clearly show the presence of various functional groups on ABAC. The predominant presence of free phenollic –OH, C-H stretch vibrations, C=O in the quinine configuration are found in ABAC. The asymmetric O=C=O stretch vibration of CO$_2$ adsorbed in pores of ABAC are also a found.

![Fig. 2: FTIR of ABAC](image)
<table>
<thead>
<tr>
<th>Molecular formula</th>
<th>Molecular weight</th>
<th>(\lambda_{\text{max}}) (nm)</th>
<th>Melting point (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(\text{C}<em>{52}\text{H}</em>{54}\text{N}<em>{4}\text{O}</em>{12})</td>
<td>927.1</td>
<td>616</td>
<td>159°C</td>
</tr>
</tbody>
</table>

**Estimation of malachite green**

The concentration of malachite green in the supernatant solution after and before adsorption was determined using a double beam UV/VIS spectrophotometer (Lambda 35) at 616 nm. The properties of malachite green are shown in Table 2. It was found that the supernatant solution from the activated carbon did not exhibit any absorbance at this wavelength and also that the calibration curve was very reproducible and linear over the concentration range used in this work.

**Method for adsorption**

**Adsorption studies**

Adsorption experiments were performed in a set of 7 BOD bottles (250 mL) where 50 mL solution of malachite green dye with 50 mg L\(^{-1}\) concentration was placed in each bottle. 0.0025, 0.005, 0.01, 0.015, 0.02, 0.025, 0.03 g ABAC was added to these bottles and kept in an isothermal shaker (30 ± 1°C) for 24 hr to reach equilibrium of the solid-solution mixture. The pH of the dye was found to be 6.7. After shaking the flasks for 3 days, samples were withdrawn from the conical flasks and the MG solutions were separated from the adsorbent by centrifugation at 3000 rpm. Residual dye concentrations in the supernatant solutions were estimated by measuring absorbance at 616 nm with Perkin-Elmer UV/VIS spectrophotometer.

**Kinetic studies**

For kinetics study a cylindrical glass vessel of 5 L capacity fitted with 8 baffles was used. 0.4 g of accurately weighted activated carbon was introduced into 2 L of dye solution of known concentration with constant stirring. The adsorbate was taken out from the vessel at regular time intervals and concentration was determined by using UV/visible spectrophotometer.

**Adsorption isotherms**

The amount of adsorption at equilibrium, \(Q_e\) (mg g\(^{-1}\)), was calculated by:

\[
Q_e = \frac{(C_0 - C_e)V}{W} \quad \ldots(1)
\]
Where; $C_0$ and $C_e$ (mgL$^{-1}$) are the liquid-phase concentrations of dye at initial and equilibrium, respectively. $V$ (L) is the volume of the solution and $W$ (g) isthe mass of dry adsorbent used. In general, the adsorption isotherm describes how adsorbates interact with adsorbents as shown in Fig. 3 and therefore it is critical in optimizing the use of adsorbents.

![Fig. 3: Adsorption isotherm of ABAC-MG System](image)

Fig. 3 clearly indicates favourable adsorption Isotherm for ABAC-MG system. Langmuir isotherm can be used as a model to describe the adsorption isotherm. The Langmuir equation is given as-

$$\frac{C_e}{q_e} = \frac{1}{Q_0 \times b} + \frac{C_e}{Q_0} \ldots(2)$$

Where $C_e$ is the concentration of adsorbate solution at equilibrium (mgL$^{-1}$), $q_e$ is the amounts of adsorbate adsorbed per mass of adsorbent (mg g$^{-1}$), $b$ is the equilibrium constant related to the sorption energy between the adsorbate and adsorbent (dm$^3$mg$^{-1}$) and $Q_0$ is limiting amount of adsorbate that can be taken up per mass of adsorbent. The calculated Langmuir constants are given in Table 3. As shown in the Table 3, the resulting $Q_0$ value is 333.3 mg g$^{-1}$.

**Table 3: Adsorption isotherm and kinetic data of ABAC-MG systems**

<table>
<thead>
<tr>
<th>Adsorbent</th>
<th>Langmuir adsorption constants</th>
<th>Freundlich adsorption constants</th>
<th>Langmuir kinetic constants</th>
</tr>
</thead>
<tbody>
<tr>
<td>Activated carbon</td>
<td>$Q_0$</td>
<td>b</td>
<td>$K_f$</td>
</tr>
<tr>
<td>ABAC</td>
<td>333.3</td>
<td>3.00</td>
<td>174.98</td>
</tr>
</tbody>
</table>

The result indicates that ABAC shows higher adsorption capacity for malachite green removal. The plot of $1/q_e$ vs $1/C_e$ in Fig. 4 for ABAC gave straight line implies that the
adsorption for adsorbent is well fitted to Langmuir adsorption model. The well-known logarithmic form of Freundlich model is given by the following equation:

$$\log q_e = \log K_F + \left(\frac{1}{n}\right) \log C_e$$  

Where $q_e$ is the amount adsorbed at equilibrium (mg g$^{-1}$), $C_e$ the equilibrium concentration of the adsorbate (MG) and $K_F$ and $n$ are Freundlich constants, $n$ giving an indication of how favorable the adsorption process and $K_F$ is the adsorption capacity of the adsorbent. The data, Table shows that Freundlich Isotherm model is not well fitted in the present study as shown in Fig. 5 for ABAC.

![Langmuir adsorption of ABAC-MG System](image1)

**Fig. 4: Langmuir adsorption of ABAC-MG System**

![Freundlich isotherm of ABAC-MG System](image2)

**Fig. 5: Freundlich isotherm of ABAC-MG System**

**Adsorption kinetics**

Following simplified expression based on Langmuir theory used to analyse the kinetic data.
\[
\ln\left[\frac{(C_t-C_e)}{(C_t+a)}\right] = -kC_t + \ln\left[\frac{(C_0-C_e)}{(C_0+a)}\right] \quad (4)
\]

Where, \( a = \frac{(C_0/kC_e)}{k} \) and \( K = \frac{k_a}{k_d} \)

\( K \) is Langmuir adsorption constant, \( k_a \) adsorption rate and \( k_d \) is desorption rate constant and \( t \) is time in min. The adsorption and desorption rate constants were thus evaluated by plotting \( \ln\left[\frac{(C_t-C_e)}{(C_t+a)}\right] \) vs \( t \), Table 3 reports the values of adsorption constants. From the results it is evident that the rate of adsorption on ABAC is More at initial time intervals as shown in Fig. 6. The value of \( K_d \) shows that ABAC is a good adsorbent for the removal of MG from aqueous solutions. They are very well fitted with Langmuir kinetic model as shown in Fig. 7.

![Fig. 6: Kinetic study of ABAC-MG System](image1)

![Fig. 7: Langmuir kinetic study of ABAC-MG System](image2)
CONCLUSION

In the present study, it has been concluded that;

(i) Phosphoric acid is a suitable activating agent for the preparation of ABAC.

(ii) Adsorption isotherms of malachite green dye on prepared ABAC are favourable.

(iii) Prepared almond based activated carbon has potential application for the removal of malachite green from aqueous solutions.

(iv) SEM analysis shows that ABAC has full of cavities with different pore diameter.

(v) Adsorption capacity increases with the increase of the adsorption dose.

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