Removal of Dyes from Dye Effluent by Using Sugarcane Bagasse Ash as an Adsorbent

Sachin M.Kanawade and R.W.Gaikwad

Abstract-Sugar cane bagasse ash, an agricultural byproduct, acts as an effective adsorbent for the removal of dyes from aqueous solution. Batch adsorption study was investigated for the removal of Acid Orange-II from aqueous solution. Adsorbents are very efficient in decolorized diluted solution. The effects of bed depth on breakthrough curve, effects of flow rate on breakthrough curve were investigated. The removal of dyes at different flow rate (contact time), bed height, initial dye concentration, column diameter, pH & temperature by Sugarcane Bagasse Ash as an adsorbent has been studied. It is found that percent adsorption of dyes increases by decreasing flow rate from 2 lit/hr to 1 lit/hr, by increasing bed height from 15cm to 45cm, by decreasing initial conc.150mg/lit to 100mg/lit, by increasing column diameter from 2.54cm to 3.5cm,by maintaining neutral pH & at temperature 45°c than 25°c & 35°c. The result shows that, bagasse ash is a good adsorbent for dye effluent treatment.

Index Terms—Adsorption, Dye, Effluent treatment, Environment, Sugarcane Bagasse Ash, Industry, India, Textile dye, Water pollution.

I. INTRODUCTION

Dyes production industries and many others industries which used dyes and pigments generated wastewater, characteristically high in colour and organic content. Presently, it was estimated about 10,000 of different commercial dyes and pigments exists and over 7×10^5 tones are produced annually world wide [1].

Dyes are widely used in industries such as textile, rubber, paper, plastic, cosmetic etc. Among these various industries, textile ranks first in usage of dyes for coloration of fiber. The convectional wastewater treatment, which rely on aerobic biodegradation have low removal efficiency for reactive and other anionic soluble dyes. Due to low biodegradation of dyes, a convectional biological treatment process is not very effective in treating a dyes wastewater. It is usually treated with either by physical or chemical processes. However, these processes are very expensive and cannot effectively be used to treat the wide range of dyes waste[2]. The adsorption process is one of the effective methods for removal of dyes from the waste effluent. The process of adsorption has an edge over the other methods due to its sludge free clean operation and completely removed dyes, even from the diluted solution. Activated carbon (powdered or granular) is the most widely used adsorbents because it has excellent adsorption efficiency for

Manuscript received May 21, 2011; revised June 15, 2011.

the organic compound. But, commercially available activated carbon is very expensive. Furthermore, regeneration using solution produced small additional effluent while regeneration by refractory technique results in a 10-15% loss of adsorbents and its uptake capacity [3]. This had lead to further studies for cheaper substitutions. Nowadays, there are numerous numbers of low cost, commercially available adsorbents which had been used for the dye removal (Table 1).

Properties of Acid orange-II

- ✤ CAS NO. 633-96-5
- ✤ EINECS NO. 211-199-0
- Formula -HOC10H6N=NC6H4SO3Na
- ✤ Molecular weight 350.32
- ◆ H.S. CODE 3204.12
- Physical state Yellow Powder
- Melting point 1640 C
- Solubility in water Soluble (116 g/lit)
- FASTNESS Light (3-4), Washing (4), Rubbing Wet (4-5), Rubbing Dry (3-4)
- NFPA RATINGS Health: 1; Flammability: 0; Reactivity: 0
- ✤ Stability Stable under ordinary conditions [11].

Structural formula of Acid orange - II



Fig 1.1 Structural formula for Acid orange-II

Sachin M.Kanawade is with Pravara Rural Education Society's Sir Visvesvaraya Institute of Technology, Chincholi, Chemical Engineering Department,Pune University,Tal-Sinnar, Dist- Nasik, M.S., India.

R.W.Gaikwad is with Pravara Rural Education Society's Pravara Rural Engineering College, Loni, Tal-Rahata, Dist-A'nagar, M.S., India.

| Sr No | Adsorbont(s) | Dyo(s) | Deferences |
|--------|--------------------------------|---|----------------|
| 51.110 | Domboo dust acconst shall | Mathulana hive | Kelefences |
| 1. | Bamboo dust, coconut snen | Methylene blue | [4] |
| | enougherst shall give hugh | | [4] |
| | groundnut shell, rice husk | | |
| 2 | Sills aattan hull aaaanut tooo | Phodomine D. Conce and methodone | |
| ۷. | Slik cotton hull, coconut tree | Knodamine-B, Congo red, metnylene | [2,5] |
| | | http://www.doctoric.let.words.chite.com/w | [2,5] |
| | sawdust, sago waste, maize cob | blue, metnyl vlolet, malachite green | |
| 2 | | | [7] |
| 3. | Parthenium hysterophorus | Methylene blue, malachite green | [/] |
| | D: 1 1 | | [10] |
| 4. | Rice husk | Malachite green | [12] |
| | a · · · · | | |
| 5. | Coir pith | Acid violet, acid brilliant blue, | 510 1 0 |
| | | | [13,15] |
| | | methylene blue, Rhodamine-B | |
| | | | |
| 6. | Orange peel | Acid violet 17 | |
| | | | [16,20] |
| 7. | Indian Rosewood | Malachite green | [20] |
| | | | |
| 8. | Prosopic cineraria | Malachite green | [11,2] |
| | | | |
| 9. | Banana and orange peels | Methyl orange, methylene blue, | |
| | | | [19,17] |
| | | Rhodamine-B, congo red, methyl violet, | |
| | | | |
| | | acid black 10B | |
| | | | |
| 10. | Giant duckweed | Methylene blue | [21] |
| | | | |
| 11. | Banana pith | Congo red, Rhodamine-B, acid violet, acid | [17] |
| | * | brilliant blue | |
| 12. | Orange peel | Congo red, Rhodamine-B, procion orange | |
| | 0 | | [10] |
| 13. | Carbonized coir pith | Acid violet, Rhodamine-B | [4] |
| | I. | , | |
| 14. | Hardwood | Astrozone blue | [2] |
| | | | |
| 15 | Chitosan | Acid blue 25, basic blue 69 | [9] |
| | | | L* 1 |
| 16 | Mahogany sawdust rice husk | Acid vellow 36 | [14] |
| 10. | manogany sawaust, nee nusk | rea yenow 50 | [17] |
| 17 | Biogas residual slurry | Congo Red Rhodmine-B acid violet acid | |
| 1/. | Diogas residuai siurry | Congo Iccu, Knounnie-D, aciu violet, aciu | [3] |
| | | brilliant blue | [3] |
| | | ormant orac | |
| 10 | Dlum kernels | Pasia Pad 22 acid blue 25 | [7] |
| 18. | I IUIII KEIIIEIS | Dasie Keu 22, aciu Diue 25 | [/] |
| 10 | Rice buck | Safranina mathulana hlua | [19] |
| 19. | NICE HUSK | Sananne, meuryrene blue | [10] |

TABLE 1: SOME LOW COST MATERIALS FOR DYES REMOVAL FROM AQUEOUS SOLUTION

II. EXPERIMENTAL STUDIES

In the present study, bagasse fly ash (BFA) and wood ash have been utilized for the treatment of acid orange-II bearing aqueous solution. Experimental details of the study have been presented. This details include characterization of adsorbents, column studies etc.

A. Characterization of adsorbent

TABLE 2.1.1 PHYSICAL-CHEMICAL CHARACTERISTICS OF BFA & WA

| Surface area(m ² /g) | 205.96 |
|------------------------------------|---------|
| Pore volume(cm ² /g) | 0.115 |
| Avg pore diameter(A ⁰) | 19.38 |
| Heating value(KJ/g) | 7231.50 |
| Bulk density(Kg/m ³) | 97.35 |

TABLE 2.1.2 PARTICLE SIZE ANALYSIS OF (BFA & WA)

| Sieve | %wt |
|---------|-------|
| >600 | 0.40 |
| 600-500 | 0.17 |
| 500-425 | 4.65 |
| 425-300 | 2.12 |
| 300-212 | 5.30 |
| 212-150 | 20.50 |
| 150-125 | 3.20 |
| 125-90 | 36.75 |
| 90-63 | 11.25 |
| <63 | 15.65 |
| | |

Avg particle size = $130.52 \mu m$

TABLE 2.1.3 PROXIMATE ANALYSIS OF (BFA & WA)

| Characteristics | Value |
|------------------|-------|
| Moisture (%) | 5.10 |
| Volatiles (%) | 9.09 |
| Ash (%) | 35.58 |
| Fixed carbon (%) | 51.23 |

TABLE 2.1.4 ULTIMATE ANALYSIS OF (BFA & WA)

| Element | Value |
|---------|--------|
| C (%) | 61.560 |
| H (%) | 0.093 |
| N (%) | 3.235 |
| S (%) | 0.132 |

B. Column study

An acrylic column of 100 cm length and 2.54 cm internal diameter with 0.2cm wall thickness was used to contain the mixture of bagasse fly ash and wood ash as a fixed bed adsorber. A fine mesh was introduced at the bottom of the column over which a layer of ceramic beads was spread to prevent the escape of adsorbent. Acid orange –II was feed to the bed of BFA & Wood ash in up-flow mode to avoid channeling of effluent.

Here we used saline tube to control up-flow motion of solution from tank to column, which is carried out by difference in potential head .Also we provide four outlets at height of 15,30,45cm. A sample of treated effluent was collected periodically and analyzed for (C_t) by colorimeter. The flow through the tested column was continued until the acid orange- II concentration of column effluent approached 0.99, C_t / C_0 , which indicate the exhaustion point (C_x). The curve of C_t / C_0 vs. volume treated between concentration at 10% breakthrough (C_b) and the exhaustion (C_x) is called the breakthrough curve.

C. Experimental set-up



Fig.2.1 Experimental set-up of adsorption column

III. RESULT AND DISCUSSION

A. Effect of bed depth on break through curve

Fig.3.1, 3.2 shows the breakthrough curve for different BFA bed depth between 15 to 45 cm , at a flow rate of 1.0 and 2.0 lit/hr. The time required for the effluent to reach breakthrough concentration, t_b increased with increase in bed depth. This may be attributed to the increase in binding sites on the adsorbent. Although an increasing bed depth increases t_b , Very high bed depth is not useful for single column. The shape and gradient of the breakthrough curves were slightly different with the variable bed depth. As the bed height increases from the 15 cm to 45cm the curve changes their profile steep concave to flatter concave. The breakthrough curve of longer beds tended to be more

gradual, meaning that the column was difficult to be completely exhausted. Here, breakthrough time is taken as time of operation at which the ratio of effluent to the inlet Acid orange –II concentration becomes 0.1 i.e. the effluent concentration gives the value of 10 mg/lit.

In the present study, the flow rate of outlet was unstable when the bed depth was too high due to higher flow resistance, resulting from tighter packing of the longer bed containing larger amount of BFA and wide size distribution. Since the treated effluent was not properly drawn of at the top of the packed bed higher than the 45 cm for the diameter of 2.54 cm, the optimum depth to diameter ratio of the column containing BFA should not be higher than 18 (45 cm/2.54 cm). Actually this ratio can be applied for the scale up of the column adsorber [5].



Fig. 3.1 Effect of bed height with respect to time on removal of Acid orange – II by BFA (C0 = 100 mg/lit, flow rate = 1.0 lit/hr)



Fig. 3.2 Effect of bed height with respect to time on removal of Acid orange– II by BFA (C0 = 100 mg/lit, flow rate = 2.0 lit/hr

B. Effect of flow rate on breakthrough curve

Fig. 3.3 shows the breakthrough profile for the adsorption of Acid orange -II removal in the continuous flow fixed column of 15cm bed depth with BFA at room temperature. The flow rate was varied from 1.0 to 2.0 lit/hr while the inlet Acid orange -II concentration in the feed was held constant at 100 mg/lit. When the adsorption zone moves upwards and upper edge of this zone reaches the top of the column, the effluent concentration start to rise rapidly. This point is called breakthrough point, which indicate the tb. Result shows that a decrease in the flow rate at constant bed depth increases the tb, and therefore the Vb (volume of solution treated at breakthrough point), due to an increase in empty bed contact time (EBCT). Using a smaller flow rate, the front of the adsorption zone reaches the top of the column later, there by giving higher tb . An increase in the flow rate appears to increase the sharpness of the breakthrough curve. These results indicates that as the flow rate increases the shape of the breakthrough curve drastically changes from S- shaped to

that of downwardly concave shape. The curves exhibit a sharp leading edge and a broad trailing edge. The broadness of the trailing edge is most likely due to slow intra-particle diffusion within the pores of BFA.

As the flow rate through the bed increases, the depth of the adsorption zone increases and the time of the contact of the solution with adsorption zone decreases. In designing a column, the length of the adsorption zone represents the minimum bed depth needed to produced a low effluent Acid orange –II concentration this results suggest that low flow rate or longer contact time may required for Acid orange-II adsorption by column field BFA and Wood ash[5].



Fig. 3.3 Effect of flow rate with respect to time on removal of Acid orange- II by BFA (C0 = 100 mg/lit, bed height =15 cm)

C. Effect of Flow rate (contact time) on Dye removal

From Fig.3.4,3.5 as the flow rate increases from 1 Lit/hr to 2 Lit/hr, the removal of Acid orange –II decreases for the same bed height. It is due to the minimum time of contact between adsorbent and adsorbate.



Fig.3.4 Effect of Bed height on Dye removal, flow rate=1lit/hr



Fig.3.5 Effect of Bed height on Dye removal, flow rate=2lit/hr

D. Effect of Bed height on Dye removal

From Fig.3.4,3.5 as the bed height increase from 15cm to 45cm, the removal of Acid orange- II increase for the same flow rate and initial concentration. It is due to the decrease in adsorbate to adsorbent ratio for the same initial concentration.

E. Effect of Initial Dye concentration on Dye removal

From Fig. 3.6 as we increase the Initial concentration of Acid orange –II from 100mg/lit to 150mg/lit, the Removal of Acid orange –II decrease for the same flow rate and Bed height. It is due the increase in adsorbate to adsorbent ratio for same flow rate.



Fig.3.6 Effect of initial Dye conc. on Dye removal, Bed height=15cm,flow rate=1lit/hr

F. Effect of Column Diameter (For same Contact time) -

Here we use two types of column of Diameter 2.54 cm and 3.5 cm. In both columns we fill adsorbent up to 15cm and having same initial concentration of 100mg/lit. From Fig.3.7 the rate of removal of acid orange _II is higher in larger dia.column than smaller Dia.column due to the increase in surface area for same contanct time.



Fig.3.7 Effect of Column Diameter on Dye removal, Flow rate=1lit/hr

G. Effect of pH on Dye removal

Here we use three different solution having pH range 3, 7, and 9. From Fig.3.8 the rate of removal of Acid orange –II is very higher at pH- 7 (Neutral), medium at pH-3 (Acidic range), lower at pH- 9 (Basic range)



Fig.3.8 Effect of pH on Dye removal, Flow rate=11it/hr,Bed height=15cm,Co=100mg/lit

H. Effect of Temperature on Dye removal

Temperature is one of the most important controlling parameter in adsorption. Adsorption is normally exothermic in nature and the extent and rate of adsorption in most cases decreases with increasing temperature of the system. Some of the adsorption studies show increased adsorption with increasing temperature.

The effect of temperature was investigated in the temperature range 25 to 45°C. From Fig. 3.9 the removal of acid orange-II increased from 95.12 to 99.95% by increasing the temperature from 25 to 45 °C Most of the adsorption processes are reported to be exothermic in nature, but present study has been found to be an example of endothermic adsorption.

This increase in adsorption is mainly due to increase in number of adsorption sites caused by breaking of some of the internal bonds near the edge of the active surface sites of the adsorbents



Fig.3.9 Effect of Temperature on Dye removal

IV. CONCLUSIONS

Sugar cane bagasse ash, an agricultural by-product, acts as an effective adsorbent for the removal of dyes from aqueous solution. Batch adsorption study was investigated for the removal of Acid Orange-II from aqueous solution. Adsorbents are very efficient in decolorized diluted solution. The effects of bed depth on breakthrough curve, effects of flow rate on breakthrough curve were investigated. The removal of dyes at different flow rate (contact time), bed height, initial dye concentration, column diameter, pH & temperature by Sugarcane Bagasse Ash as an adsorbent has been studied. It is found that percent adsorption of dyes increases by decreasing flow rate from 2 lit/hr to 1 lit/hr, by increasing bed height from 15cm to 45cm, by decreasing initial conc.150mg/lit to 100mg/lit, by increasing column diameter from 2.54cm to 3.5cm, by maintaining neutral pH & at temperature 45^{0C} than 25^{0C} & 35^{0C} . The result shows that, bagasse ash is a good adsorbent for dye effluent treatment.

ACKNOWLEDGEMENT

We are gratefully acknowledgements to Environmental Chemical Engg.Department of Pravara Rural Engg. College, Loni for experiment facility & their guidance, help in completion of this paper

REFERENCES

- Gong, R., Sun, Y., Chen, J., Liu, H. & Yang, C. (2005). Effect of chemical modification on dye adsorption capacity of peanut hull. *Dyes Pigments*, 67, 175–181.
- [2] Nigam, P., Armour, G., Banat, I.M., Singh, D. & Marchant, R. (2000). Physical removal of textile dyes from effluents and solid-state fermentation of dye-adsorbed agricultural residues. *Bioresource Technol.*,72, 219–226.
- [3] Mane, V.S., Mall, I.D. & Srivastava, V.C. (2007). Use of bagasse fly ashas an adsorbent for the removal of brilliant green dye from aqueous solution. *Dyes Pigments*, 73, 269–278.
- [4] Gong, R., Jin, Y., Chen, J., Hu, Y. & Sun, J. (2007). Removal of basic dyes from aqueous solution by sorption on phosphoric acid modified rice straw. *Dyes Pigments*, 73, 332–337.
- [5] Kavitha, D. & Namasivayam, C. (2007). Recycling coir pith, an agricultural solid waste, for the removal of procion orange from wastewater. *Dyes Pigments*, 74, 237–248.
- [6] Figueiredo, S.A., Loureiro, J.M. & Boaventura, R.A. (2005). Natural waste materials containing chitin as adsorbents for textile dyestuffs: Batch and continuous studies. Water Res., 39, 4142–4152.
- [7] Robinson, T., Chandran, B. & Nigam, P. (2002). Effect of pretreatments of three waste residues, wheat straw, corncobs and barley husks on dye adsorption. *Bioresource Technol.*, 85, 119–124.
- [8] Gong, R., Jin, Y., Sun, J. & Zhong, K. (2008). Preparation and utilization of rice straw bearing carboxyl groups for removal basic dyes from aqueous solution. *Dyes Pigments*, 76, 519–524.
- [9] Ong, S.T., Lee, C.K. & Zainal, Z. (2007). Removal of basic and reactive dyes using ethylenediamine modified rice hull. *Bioresource Technol.*, 98,2792–2799.
- [10] Laszlo, J.A. (1996). Preparing an ion exchange resin from sugar cane bagasse to remove reactive dye from wastewater. Text. Chem. Color,28, 13–17.
- [11] O'Connell, D.W., Birkinshaw, C. & O'Dwyer, T.F. (2008). Heavy metal adsorbents prepared from the modification of cellulose: A review.*Bioresource Technol.*, 99, 6709–6724.
- [12] Jústiz-Smith, N.G., Virgo, G.J. & Buchanan, V.E. (2008). Potential of Jamaican banana, coconut coir and bagasse fibres as composite materials.*Mater. Charact.*, 59, 1273–1278.
- [13] Banerjee, S.S., Joshi, M.V. & Jayaram, R.V. (2006). Effect of quartenary ammonium cations on dye sorption to fly ash from aqueous media. J. Colloid Interf. Sci., 303, 477–483.
- [14] Özacar, M. & Şengil, İ.A. (2003). Adsorption of reactive dyes on calcined alunite from aqueous solutions. J. Hazardous Mater., B98, 211–224.
- [15] Chiou, M.S. & Li, H.Y. (2003). Adsorption behavior of reactive dye in aqueous solution on chemical cross-linked chitosan beads. Chemosphere, 50, 1095–1105.
- [16] Malik, P.K. (2003). Use of activated carbons prepared from sawdust and rice-husk for adsorption of acid dyes: A case study of Acid Yellow 36.Dyes Pigments, 56, 239–249.
- [17] Tor, A. & Cengeloglu, Y. (2006). Removal of congo red from aqueous solution by adsorption onto acid activated red mud. J. Hazardous Mater., B 138, 409–415.
- [18] Ho, Y.S. & McKay, G. (1999). Pseudo-second order model for sorption processes. Process Biochem., 34, 451–465.
- [19] Lata, H.,Garg,V.K. & Gupta, R.K. (2007). Removal of a basic dye from aqueous solution by adsorption using Parthenium hysterophorus:An agricultural waste. *Dyes Pigments*, 74, 653–658.
- [20] Özdemir, Y., Doğan, M. & Alkan, M. (2006). Adsorption of cationic dyes from aqueous solutions by sepiolite. Micropor Mesopor Mat.,96, 419–427.
 Ho, Y.S. & McKay, G. (2003). Sorption of dyes and copper ions onto biosorbents.Process Biochem.,38,1047–1061.

Sachin M.Kanawade, photograph and biography not available at the time of publishing.

R. W.Gaikwad, photograph and biography not available at the time of publishing.