ISSN: 2320-0294

STUDY ON ADSORPTION OF METHYLENE BLUE BY ACTIVATED CARBONDERIVED FROM IPOMOEA CARNEA

Dr. Shilpi Deep Mathur, Lecturer,

Department of Chemistry,

Rameshwai Devi Girls College, Bharatpur, Rajasthan (India)

ABSTRACT

Activated carbon has been prepared from stem of Ipomoea Carnea by acid treatment Surface structureinvestigation is carried out by scanning electron microscopy (SEM). Ipomoea Carnea stem waste has been evaluated as raw material for the preparation of activated carbon using HCL with the temperature ranging from 5000C to 8000C. The activated carbon prepared was characterized, by iodine value, moisture contain, ash content, pore volume andporosity. The BET surface areas between 864.653 m²/g and 1159.06 m²/g and micropores and mesopores with volumesbetween 0.004 and 0.358041 cc/g. Characterization studies such as SEM, BET, bulk density, moisture content, ashcontent, fixed carbon content, and surface area have been carried out to assess the suitability of derived carbons asadsorbents in the water and wastewater. The effects of factorssuch as contact time, adsorbent dose, pH and initialconcentration were investigated. The kinetics and equilibriumdata were confronted to several models. A study onadsorption of Methylene blue by activated carbon derived from Ipomoea Carnea was done.

Key words: Activated Carbon, Ipomoea Carnea, Carbonization, Activation, Methylene Blue.

INTRODUCTION

Solid waste disposal has become a majorproblem in India, either it has to be disposed safely or usedfor the recovery of valuable materials as agricultural wasteslike turmeric waste, ferronia shell waste, jatropha curcusseed shell waste, delonix shell waste and Ipomea Carniastem. Therefore these wastes have been explored for thepreparation of activated carbon by various techniques. Inorder to serious water pollution, we must understand theproblems and become part of the solution.

Activated carbon has since then been used in manyindustries. In particular, it has been commonly used for theremoval of heavy metal and organic dyes from textile wastewater. Many agricultural by products and waste materials usedfor the preparation of activated carbons. The plant IpomoeaCarnea belongs to family Convolvulaceae. Ipomoea Carnea, is a species of morning glory. Activated carbon is widely usedfor the purpose due to the large surface area available foradsorption or chemical reactions. Dyes are used in chemical, textile, paper, printing, leather, plastics and various foodindustries. Waste water passed out from the industry. So in this work Activated carbon used as an adsorbent for Methylene Blue dye.

MATERIAL AND METHOD

Collection of sample - The Ipomoea Carnea (morningglory), a low-cost and abundantly available plant, was usedfor removal of heavy metal from aqueous solutions. Ipomoea Carnea stems were collected in and around Neelbad Bhopal M.P., India. The stemof Ipomoea Carnea was dried at room temperature for afew days and then oven dried at 110^{0} C over night.



ISSN: 2320-0294

Activation with HCL-

Ipomoea Carnea stem wastematerial was treated with HCL for a period of 24 hours. Then the material was placed in the muffle furnace and carbonized at 400-800°C, for a period of 60 minutes. Afteractivation, the carbon obtained was washed sufficiently withplenty of water, dried and sieved then to desired particlesize.

Characterization

Moisture Content Determination-

A 1.0 g of the activated carbon sample was weighted and dried in an oven for fourhours at 150°C, until the weight of the sample becameconstant.

$$X_{0} = \frac{W_{1} - W_{2}}{W_{1}}$$

 W_1

Where

 X_0 = Moisture content on wet basis

 W_1 = Initial weight of sample, in gm

 W_2 = Final weight of sample after drying in gm

Particle Size

For the particle size determination, lots of samples were weighed and placed on top of a set of sievesranging from 75 to 1.4 × 103 μm. The sieves were shakenmanually for two minutes, after which the weight percent of the active carbon retained on the sieves and bottom panwas determined.

pH -

1 gm of the sample was weighed and dissolved in 3 ml of de-ionized water. The mixture was heated and stirredfor 3 minutes to ensure proper dilution of the sample. The solution was filtered out and its pH was determined using adigital pH meter.

Iodine Adsorption Number (IAN)-

1 g sample and 25 mlof standard iodine solution (0.023 M) added. 20 ml of this filtrate solution was titrated with the standard (0.1095 M)thio sulphate solution to the persistent of a pale yellow colour.5 ml of starch indicator was added and titration resumedslowly until a colorless solution appeared, the procedurewas carried out two more times. The titrations were also repeated with 20 ml portions of the standard iodine solutionnot treated with the sample to serve as the blank titration. The iodine number (IAN) was calculated from therelationship.

 $IAN = 12.69 \text{ N } (V_2-V_1) \text{ mole iodine/g sample}$

Where:

N is the normality of thiosulphate solution

V₁ is the volume of the thiosulphate (ml) used for the titration of the sample-treated aliquot

V₂ is the volume of the thiosulphate (ml) use for the bank titration,

W is the mass of the sample used (gm).



ISSN: 2320-0294

Ash content-

2.0 grams of sample was placed into acrucible and reweighed with its content heated in a furnaceat 9000C for 3 hours. The sample was cooled to roomtemperature and reweighed. Ash content was calculated between the differences in weight.

Volatile matter content -

A 2 gm of sample was taken incylindrical crucible closed with a lid. It was then heated to 925 Úc for exactly 7 minutes in a muffle furnace. Then the crucible was cooled in desiccators and weighted volatilematter on dry basis.

VM = 100[100(B-F) - M(B-G)]

[(B-G)(100-M)]

B=Mass of crucible, lid and sample before heating

F=Mass of crucible, lid and contents after heating

G=Mass of empty crucible & lid

M=% of moistures determined above

Fixed carbon content-

Fixed carbon FC = 100 - (%moisture content+ % volatile matter + % ash content)

BET Test

(Brunauer, Emmett and Teller) - The results of BET surface area, macropore, mesopore and microporevolumes of the produced activated carbon. The surfaceareas were higher for HCL activated carbons and this isexpected because chemical activation normally developsmore porosity and it gives high surface area.

Scanning electron microscope analysis (SEM)

Thesurface morphology of the activated carbon was testedusing scanning electron microscopy. At such magnification, SEM micrographs clearly revealed that wide variety of poresare present in activated carbon along with fibrous structure. It is also found that there are holes and caves type openingson the surface of the adsorbent.

Adsorption studies of activated carbon-

Removal ofMethylene Blue dyes from aqueous solution by usingIpomoea Carnea stem waste.

Batch adsorption study-

The batch adsorption wascarried out at room temperature. The effects of contacttime, adsorbent dose, pH of solution and the initialconcentration of these adsorbates were studied. Thequantity adsorbed by a unit mass of an adsorbent atequilibriumand the adsorption percentage were calculated using the following relations:

$$\% R = \frac{C_o - C_e}{C_o} \times 100$$

$$Q_e = \frac{(C_o - C_e)}{C_o} \times V$$

Where is the concentration of adsorbate at equilibrium(mg.L-1); is the initial concentration of adsorbate (mg.L-1).

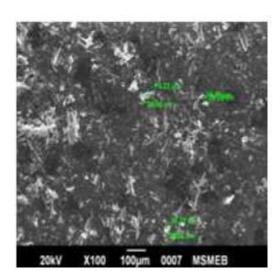
Result

IJESM

Activated carbon prepared from Ipomoea Caeneastem waste was found effective in adsorption of dye fromaqueous solution. The activated carbon was characterized pH, moisture content, bulk density, pore volume, porosity, and ash content. Iodine no. a measure of the microporecontent of the activated carbon is 254 mole iodine/g sample. Moisture content of activated carbon have less than 1%. Ash determination shows that increase in carbonization temperature reduces the ash content. It is known that materials with the lowest ash content are most active. ThepH of sample is 6.5. The result showed that the Activated carbons activated with HCL are neutral after washing. Anadsorption test has been carried out for Methylene Blue under different experimental conditions in batch mode. The adsorption of Methylene Blue was dependent on adsorbent surface characteristics, adsorbent dose, Methylene Blueconcentration, time of contact and temperature. The results indicated that the activated carbon is very useful for removing Methylene Blue from wastewater.

Table – 1: Characteristics of Activated Carbon

S.No.	Parameters	Observations
1	Moisture content in 2.0g	2.3%
2	pH 6.5	
3	Iodine adsorption number (IAS)	254
4	Ash content	1.40%
5	Fixed carbon %	81%
6	ccl adsorption capacity in %	64\$
7	Pore size/value A	1.89839 nm
8	BET Surface area in m2/g	$1159.06 \text{ m}^2/\text{g}$
9	Langmuir Surface area in m2/g	$2152.48 \text{ m}^2/\text{g}$
10	Volatile matter	15.3%



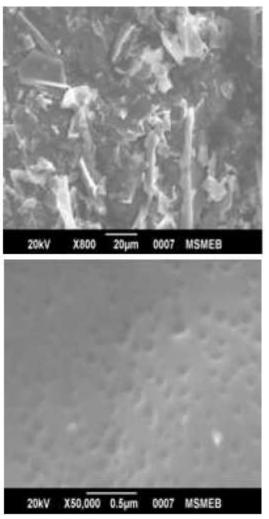


Fig. 1: Activated carbon from Ipomoea Carnea SEM

Adsorption result -

The highest adsorption of 95.1% occurred at pH 6. When increase adsorbent dose MB adsorption was increase. Figure-4 shows that the equilibrium was attained at 110 minute when the maximum adsorption onto activated carbon was reached. When Methylene Blue concentration was increase the rate of adsorption was decreases (Fig-5).

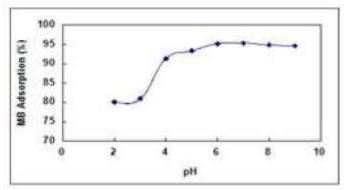


Figure -2 Effect of pH on adsorption of Methylene Blue

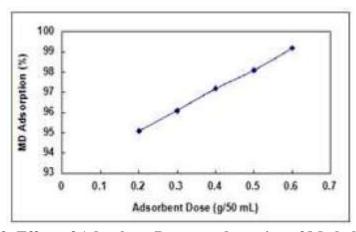


Figure-3: Effect of Adsorbent Dose on adsorption of Methylene Blue

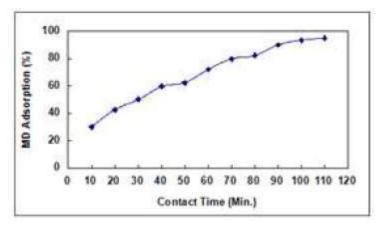
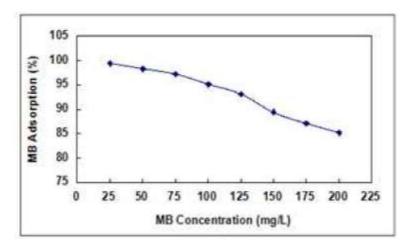


Figure -4. Effect of Contact Time on adsorption of Methylene Blue



CONCLUSIONS

This study indicates that efficient activated carbons can be obtained from Ipeomea Carnea by controlled activation with HCL for a number of industrial and residential applications. Due to the presence of high surface area, porosity, the activated carbon prepared from the agricultural waste it can be used for a variety of environmental application, dye removal, wastewater treatment and adsorption processtoo. The present study has shown the effectiveness of using AC in the removal of Methylene Blue dye from aqueous solutions. Acid Activated Ipomoea Carnea in different forms has a great role in modern life to clean environment.

REFERENCES

IJESM

- S. Karthikeyan and A. Babu Rajendran "Adsorption of Basic Dye (Rhodamine B) by a Low Cost Activated Carbon from Agricultural Solid Waste" An International Quarterly Scientific Journal Vol. 9 No. 3 pp. 461-472,2010.
- Barret, E.P., Joyner, P.B. and Hatenda, P. "The Determination of Pore Volume and Area Distribution in PorousSubstances. Computations from Nitrogen Isotherms." Journal of Analytical Chemistry, 73, 373-380.1951.
- Arockiaraj I and Renuga V "Sorption kinetics and dynamicstudies of basic dye by lowcost nanoporous activated carbon derived from Ipomoea Carnea stemwaste by sulphate process" Frontiers in Nanoscience and Nanotechnology 2016.
- Hameed, B.H, Din, A.T.M & Ahmad, A.L. Journal ofhazardous materials. 141(3), , 30-60, 2006.[24]
- Ademiluyi, T, Gumus, R, Adeniji, S.M & Jasem, O. Proceedings of the Nigerian society of chemical engineers, 34th Annual Conference Effurun, Nigeria, 2008.
- Madu P. C. and Lajide L. J "Physicochemical characteristics of activated charcoal derived from melon seedhusk". Chem. Pharm. Res., 5(5):94-98, 2013.[26]
- Ajayi OA, Olawale AS A "comparative study of thermaland chemical activation of Canarium SchweinfurthiiNutshell". J. Appl. sci. Res., 5(12): 2148-2152. 2009.
- Chandrakant d. Shendkar1, rasika c. Torane2, kavitas. Mundhe2, sangita m. Lavate1, avinash b. Pawar1,nirmala r. Deshpande. ''characterization and application of activated carbon prepared from wasteweed"international Journal of Pharmacy and PharmaceuticalSciences · March 2013.
- Lekene Ngouateu R. B., Kouoh Sone P M. A., NdiNsami J., Kouotou D., Belibi Belibi P. D., KetchaMbadcam J. "Kinetics and equilibrium studies of theadsorption of phenol and methylene blue onto cola nutshell based activated carbon" Int J Cur Res Rev | Vol 7Issue 9 M ay 2015
- Rengaraj, S., Moon, S.-H., Sivabalan, R., Arabindoo,B. and Murugesan, V. "Agricultural Solid Waste for theRemoval of Organics: Adsorption of Phenol from Waterand Waste Water by Palm Seed Coat ActivatedCarbon". Waste Management, 22, 543-548. 2002.