COMPARISON OF THE EFFECTIVENESS AND ECONOMY OF 850µM AND 590µM COCONUT SHELL ACTIVATED CHARCOAL PARTICLES IN WATER TREATMENT

¹ P. O. Dike, ² E. U. Igbonagwam and ³ O. G. Akana

(Department of Civil Engineering, Federal University of Technology Owerri, Imo State, NIGERIA)

ABSTRACT — This study investigates the effectiveness of coconut shell activated charcoal in water treatment process using common table salt or sodium chloride (NaCl) as the activation agent and water sample sourced from Otammiri River in Imo state as the test sample while comparing the effectiveness and economy of an 850µm and 590µm particle sizes of activated carbon in the water treatment processes. To form the carbon, coconut shell was heated in open air to form the carbon (charcoal) which is then grinded into a fine state. To activate the carbon, sodium chloride was added to the fine carbon and heated in an oven to a temperature of about 600°C. Two different particle sizes of activated carbon was produced, 850µm and 590µm. Three major parameters were analysed in accordance to WHO (World Health Organization) standard for the two different activated carbon sizes, these parameters include; Physical tests, Chemical tests, and Biological Analysis test. The results of the tests showed that while both particle size activated carbon where very effective in water treatment, the 590µm particle size of activated carbon is more effective as it yields better result within a short duration. But as regards to economy, the 850µm particle size of activated carbon is more effective as it can be easily re-used.

Index Terms — Activated carbon (Activated charcoal will be used interchangeably with activated carbon throughout this work, also written as AC), adsorption, E.Coli, Colonies, $L/M \rightarrow Litres/Minute$, sp \rightarrow Species.

_ _ _ _ _ _

1. INTRODUCTION

Water sources and systems, globally, have been faced with the problems of pollution, both by natural causes and man-made causes. Water treatment, being a means of quality improvement remains a valuable tool for making polluted water portable. In treatment, the sources and levels of pollution tend to determine the extent of treatment facility to be adopted (Kamrin, Michael, Nancy Hayden, 1990).

Most naturally occurring water are not free from contaminants, especially surface water and atmospheric water and therefore requires one form of treatment or

• Akana Osita is a Civil Engineer in Nigeria.

the other to meet acceptable quality standards for portability. Since often, suspended particles in the water constitute the pollution requiring treatment, introduction of chemical or treatment inputs that can be used to remove or separate these particles should be evaluated.

The turbidity, taste, odour, hardness, heavy metals and bacteria have placed a limit on the utility derived by the consumers of the water. Treatment to remove these therefore is pertinent for improving the quality of the water. These results to the use of chemical such as chlorine (Cl2), in water treatment but water treated chlorine can be harmful for consumption. Also, the use of chlorine in water treatment has been of great use but in excess can result to major health issues and it removes essential nutrients gotten from water such as calcium, etc.

Contamination of surface and atmospheric water, suspended particles in ground water and surface water has reduced the quality of drinking water which is why treatment is essential in order to make the water potable.

Author, Paul Osunna Dike is currently pursuing masters degree program in Advanced Structural engineering at Imperial College London, United Kingdom, E-mail: dike_paul@ymail.com

Co-Author, Ezinne Igbonagwam graduated from the Federal University of Technology, Owerri and is Currently working as an Environmental Engineer in Nigeria. E-mail: Zinnyigbonagwam1@gmail.com

Arbitrary choice of filtrates to be used for treatment is a bit lacking in that sand and gravel used in filtration cannot remove all contaminants.

The specific objective of this study is to check the efficiency and quality of activated carbon (charcoal) in water treatment. This objective will be pursued by, evaluation of the ability of activated carbon in removing some physical contaminants in water; ascertaining the efficiency of the activated charcoal in removing some chemical contaminants in water; evaluating the ability of activated charcoal in adsorbing microbial pathogens in water; and studying the adsorption efficiency of two different activated charcoal of different particle sizes (prepared from coconut shells and activated with sodium chloride, common salt) in the removal of pollutants in water.

2. RESEARCH METHOD

2.1 Materials and Methods

Materials used during this experiment include the following: 25 pieces of 250mL and 3 pieces of 1000mL of glass bakers; Sodium chloride NaCl (common salt); Activated carbon (prepared from coconut shell and using sodium chloride as activating agent); pH and Conductivity Meter; turbidity meter; Colour meter; Distilled water; Domestic mortar and pistol; Stop watch; Funnel; Volumetric Flask; Sieve; Filter; Measuring Cylinder.

The 20 litres (or 4 and half gallons) of raw water sample was collected from Otammiri River. The filtrate used during this project is the carbonized coconut shells which will be activated with sodium chloride also known as common salt. The coconut shells were obtained from a farm in Obazu Mbieri Mbaitoli in Imo state, Nigeria.

2.1.1 Experimental Design

A completely Randomized Design was used for this experiment. Granular and pulverized carbonized coconut shell were concentrated and activated with sodium chloride (common salt). For the preparation of the activated carbon, a completely new and innovative method was adopted. This innovation has two parts;

1. Making good quality Coconut charcoal by improving the production process:

For this an iron pot was used to burn the coconut shell. The iron pot was used because when put under a very hot temperature it doesn't melt easily and it has a very high tendency of supporting the carbonization process by speeding up the heating and keeping t tightly covered. The pot was then put in an electric muffle furnace and set to a temperature of 600°C for 2 hours. But this was done after the coconut shells were broken into small pieces, washed thoroughly to remove particles and dirt on it and sun dried. This was done repeatedly till all the shells were carbonized. Now Water is poured to cool the charcoal. It is dried to get good quality coconut charcoal. The model can take about 8 coconut shells. But bigger chambers of can be fabricated to hold about 20 to 50 shells at a time.

2. Making of activated carbon from coconut shell charcoal:

The charcoal produced in the method given earlier is mixed with sodium chloride (table salt). The Sodium chloride is to be measured using weighing balance of 20g was mixed with a 10m of 1000mL distilled water respectively to form a paste. The paste is then dissolved into the remaining distilled water of 1000ml and stirred for 5min in a magnetic stirrer with a speed of 100rpm. This solution will be mixed with the carbon with iodine value up to 750.

The Activated carbon is then grinded to two particle sizes ($850\mu m$ and $590\mu m$), this was done so as to increase its surface area and to improve its adsorptive capacity.

2.2 Tests Conducted

A total of 22 tests was carried out on the water sample to ascertain the suitability and portability of the treated water in accordance to WHO standard. These tests can be grouped into Physical, Chemical, and Biological test. The tests to be conducted are summarized in Table 2.1 below while some of the test results as expected for a suitable and portable water according to WHO standards are summarized in Table 2.2 and Table 2.3 below;

Iron Total	Mg/l	Below 0.4	0.3	AAS
------------	------	-----------	-----	-----

*NTU= Nephelometric Turbidity Unit, *AAS= Atomic Absorption Spectrophotometer

2.2.1 Filtration Procedure

The filtration bed was set up and filtration was carried out in accordance to WHO standard. Also, results were completely analysed as specified by WHO.

3. RESULTS AND DATA ANALYSIS

3.1 Data Presentation

Two samples of filtered water were obtained by filtering the raw water with two different particle size of activated carbon. Sample 1 was obtained by filtering the raw water through the 850 μ m particle size of activated carbon while sample 2 was obtained by doing same through a 590 μ m particle size activated carbon. For each sample, 5 different sub-samples namely; Control, Sample A, Sample B, Sample C and Sample D, were obtained by collecting the filtered water at specific flow rates. The flow rates are; Sample A (At flow rate of 1L/15Mins), Sample B (At flow rate of 1L/30Mins), Sample C (At flow rate of 1L/45Mins), Sample D (At flow rate of 1L/60Mins) and the Control was the raw water sample before filtration (that is, 0L/Min).

1-litre of water being filtered was collected at 15minute intervals for the two different activated carbon particle size and tested. The various graphs of each physical and chemical parameter tested are plotted against the various collection flow rates for the two activated carbon particle sizes.

It should be noted, for the graphs below;

Sample 1 = water filtered through $850\mu m$ AC particle size,

Sample 2 = water filtered through 590 μ m AC particle size. Also, on the flow rate axis;

1= Raw water sample collected before filtration,

2 = water sample A collected after 15mins,

3 = water sample B collected after 30mins,

4 = water sample C collected after 45mins and

5 = water sample D collected after 60mins

Table 2.1: Parameters	Analysed
-----------------------	----------

PHYSICAL TEST	CHEMICAL TEST	BIOLOGICAL TEST
Conductivity	РН	Coliform: Other coliform -E. coli
Turbidity	Dissolved oxygen	Faecal streptococci
Total solids	Heavy metals (Cu, Zn, and Fe)	Salmonella
Total suspended solids	Nitrate	Vibrio
Total dissolved solids	Phosphate	Clostridia
	Sulphate	Yeast/moulds/parasites
	Biochemical oxygen demand (BOD)	Planktons
	Chemical oxygen demand (COD)	Total plate counts
	Chloride	

Table 2.2: Bacteriological Analysis

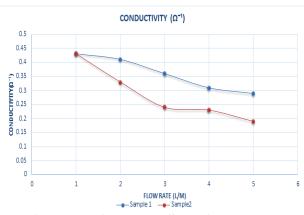
Parameter	Recommended raw water quality	Drinking water standards
	Acceptable value (mg/litre, unless otherwise stated)	Maximum acceptable value (mg/litre, unless otherwise stated)
Total coliform	5000MPN/100ml	0 in 100 ml
E.coli	5000MPN/100ml	0 in 100 ml
Coliform Bacteria		Absent

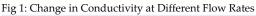
Table 2.3: Physical and Chemical Analysis

Parameter	Unit	Test Remarks	Requirem ent	Methods
Colour	Pt. Co- scale	3	15	Colorimetric
рН	Pt. Co scale	6.50	6.5-8.5	Electrometric
Turbidity	NTU	1	5	Turbidity meter
Copper	Mg/l	Below 0.3	1.0	AAS
Zinc	Mg/l	0.047	5	AAS

3.1.1 Physical Tests Result:

Physical tests are summarized in Fig 1-5 below;





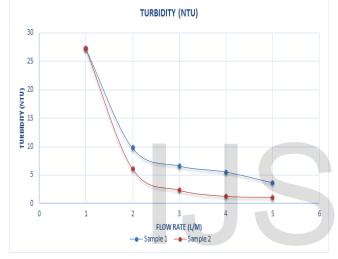


Fig 2: Change in Level of Turbidity

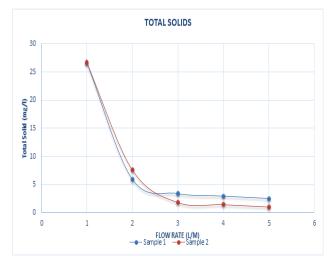


Fig 3: Change in Level of Total Solids

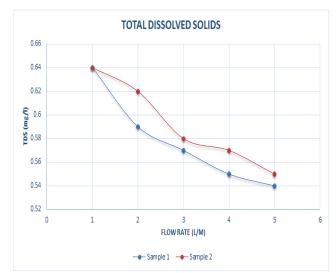


Fig 4: Change in Level of Total Dissolved Solids

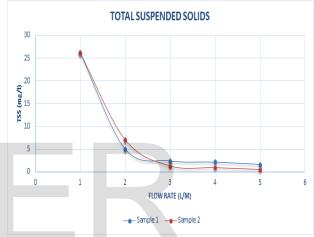


Fig 5: Change in Level of Total Suspended Solids

3.1.2 Chemical Tests Result:

Physical tests are summarized in Fig 6-17 below;

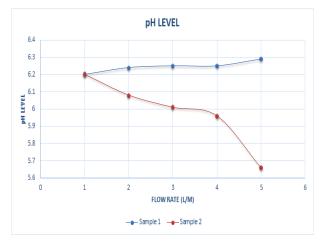
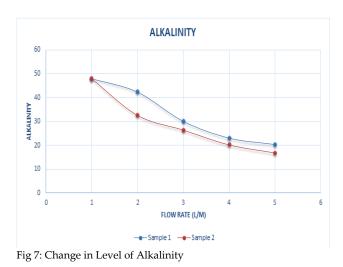


Fig 6: Change in pH Level



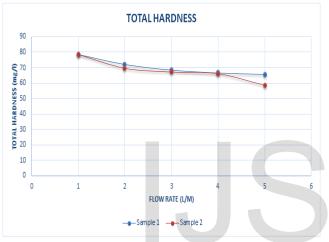


Fig 8: Change in Level of Total Hardness

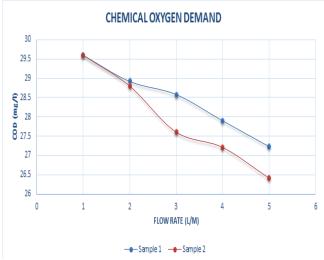


Fig 9: Change in Level of Chemical Oxygen Demand

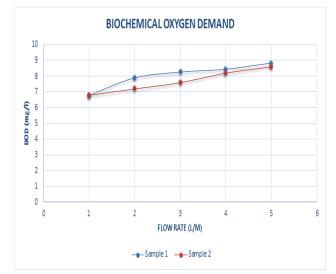


Fig 10: Change in Level of Bio- Chemical Oxygen Demand

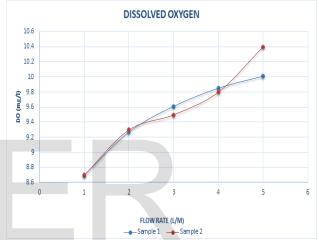


Fig 11: Change in Level of Dissolved Oxygen

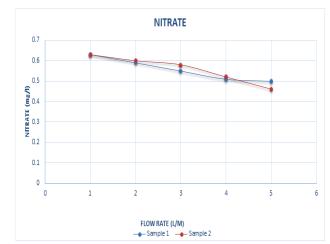


Fig 12: Change in Level of Nitrate

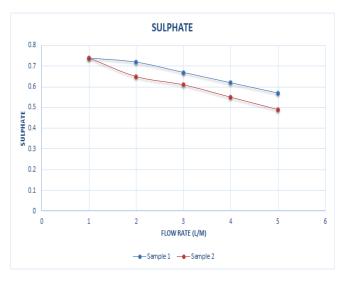


Fig 13: Change in Level of Sulphate

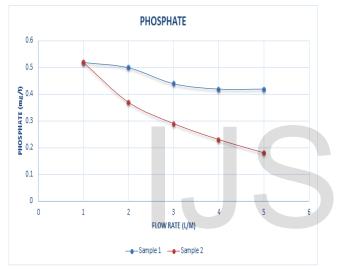


Fig 14: Change in Level of Phosphate

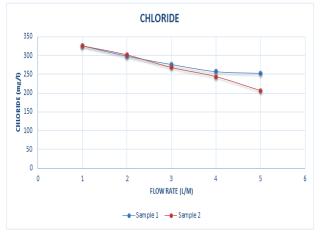


Fig 15: Change in Level of Chloride

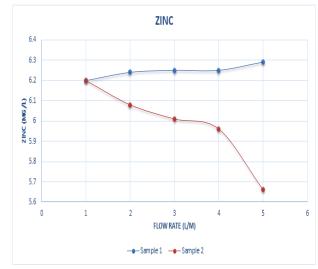


Fig 16: Change in Level of Zinc



Fig 17: Change in Level of Iron

3.1.3: Biological Tests:

For the first sample treated activated carbon of particle size $850 \mu m$

Table 4: Control Sample

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E. coli	35 colonies	35 colonies
Faecal streptococci sp	Nil	Nil
Salmonella sp	Nil	Nil
Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	50 colonies	5.0 x10 ²
COUNTS		

Table 5: Sample A

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	14 colonies	14 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast /molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	25 colonies	2.5
COUNTS		

Table 6: Sample B

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	12 colonies	12 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	45 colonies	4.5 x10 ²
COUNTS		

Table 7: Sample C		
MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	10 colonies	10 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	30 colonies	3.0 x10 ²
COUNTS		

Table 8: Sample D		
MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	7 colonies	7 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	28 colonies	2.8 x10 ²
COUNTS		

For the second sample treated with activated carbon of particle size of $590 \mu m;$

Table 9: Sample A

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	11 colonies	11 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	22 colonies	2.2 x10 ²
COUNTS		

Table 10: Sample B

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	3 colonies	3 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	12 colonies	1.2 x10 ²
COUNTS		

Table 11: Sample C

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	15 colonies	15 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	8 colonies	8.0 x10 ²
COUNTS		

Table 12: Sample D

MICROBIAL GROUP	PLATE	UNIT
	READING	(CFU/ML)
PATHOGENS		
Coliform: Other coliform	Nil	Nil
E.coli	7 colonies	7 colonies
Faecal streptococci sp	Nil	Nil
Salmonella & Vibrio sp	Nil	Nil
Clostridia sp	Nil	Nil
Yeast/molds/parasites	Nil	Nil
Plankton	Nil	Nil
TOTAL-PLATE	6 colonies	6.0 x10 ²
COUNTS		

3.2. DISCUSSION OF RESULTS

3.2.1. Discussion on the Physical Test

Turbidity: in sample 1, activated carbon of 850µm size, the water became clear at sample D (flow rate of 1L/60min) with turbidity of 3.64NTU, while the second sample (590µm AC size) became clear at sample B (flow rate of 1L/30mins) with turbidity of 2.34NTU as shown in FIG 2 above. Also, from FIG 1 above, sample 2 has a lower conductivity compared to sample 1. This clearly shows that while both particle sizes of the activated carbon were effective in clearing the water (WHO Standard for turbidity in 5.0NTU), the smaller AC particle size was more effective, reducing the turbidity to a minimum at a very short time interval.

Total solid, total dissolved solids and total suspended solids: these gave positive results to how effective the activated carbon of both particle sizes responded to the removal of these contaminants as from FIG 3, FIG 4 and FIG 5 above. But it can be clearly seen that the smaller particle size activated carbon was more effective.

3.2.2. Discussion on the Chemical Test

The Ph is an indicator of relative acidity or alkalinity of water. Values of 9.5 and above indicate high alkalinity while values of 3 and below indicates high acidity level. Both samples were appropriate and fell between the given standard of 5.5 - 8.5 for all flow rates as seen in FIG 6 and FIG 7.

The heavy metals (Fe and Zn) were both impressive with their outcomes, copper was not found in the raw water that is why no result was given. Iron and Zinc, from FIG 16 and FIG 17, both fell drastically below WHO standard after the treatment for sample A (1L/15min) for both samples which showed a huge progress on how activated carbon is at removing certain heavy metals.

Nitrates, phosphate, sulphate, and chloride also yielded wonderful result as from FIG 12, 14, 13 and 15 respectively, and they all fell below standards except for chloride treated with activated particle size of 850µm (FIG 15) which will eventually yield a better result if left for longer time period. for sample D, the level of chloride in the water after treatment was 3.49mg/l out of 253.49mg/l while its standard is 250.0mg/l. The BOD, COD and DO gave reasonable results as seen from FIG 10, 11 and 9 respectively.

3.2.3. Discussion on the Biological Test

Bacteriological tests show the presence of bacteria, characteristic of faecal pollution, coliform (E.coli), pathogens (salmonella, vibrio, clostridia), parasites, planktons, etc. which improved with satisfactory results after treatment (Tables 4 - 12 above). The raw water which contained faecal coliform was completely removed and the rest of also produced satisfactory result according WHO allowable limit for potable drinking water which stated that the following microorganisms listed above should be below 100 colonies but in these test the total plate count for the first sample was 25colonies for sample A, 45 colonies for sample B, 30 colonies for sample C, and 28colonies for sample D.

3.3. CHARACTERIZATION OF ACTIVATED CARBON

Adsorptive characteristics: Iodine number was determined to access the absorptive capacity of prepared activated carbon. Iodine adsorption is easy and commonly used methods to characterize activated carbon performance. Iodine is a small sized molecule, so it indicates the capacity of carbon to adsorb smaller molecules. However, the higher the iodine number the better its adsorptive capacitance. For the first activated carbon its iodine number is 510mg/gm while that of the second is 764mg/gm.

Table 13: Compila	tion of characteristics	of the two different
activated carbon		

CHARACTERISTICS	FIRST ACTIVATED CARBON	SECOND ACTIVATED CARBON
Particle sizes	850µm (0.850mm)	590µm (0.590mm)
Mesh	No. 20	No.30
Inch	0.0331	0.0234
Bulk density g/ml	0.731	0.743
Conductivity (Ω ⁻¹)	0.01	0.02
РН	7.08	7.31
Ash content %	2.05	2.21

4. CONCLUSION AND RECOMMENDATION

4.1 CONCLUSION

Activated carbon adsorption is an effective means for reducing organic chemicals, chlorine, and unpleasant tastes and odors in water, (Guo J and Lua A.C. 2001).

This treatment can produce water of more desirable quality than that from some public or private supplies. Units ranging from simple, manually operated devices to complex, automatic ones are designed to ensure the reduction of specific contaminants. The 590µm particle size activated carbon is more effective as it yields better result within a short duration. But as regards to economy, the $850\mu m$ particle size activated carbon is advised as it can be easily re-used.

The effect of activating agents on the properties of activated carbon prepared from coconut shell by chemical activation has been investigated. Activating agents had no significant effect on the nature of surface functional group. Activated carbon impregnated with sodium chloride showed well developed pores structure. Iodine Number adsorption also indicates that sodium chloride is the best activating agent among the various activating agents tested. Hence this study shows that coconut shell can be used as a source of lignocellulosic material for the preparation of low cost, high surface area activated carbon with well-developed porosity using NaCl as an activating agent.

4.2. RECOMMENDATION

Based on the findings of this research work, the use of chemically activated carbon is very much effective in water treatment processes. The particle sizes in this research also showed that the smaller/finer the particle size of the activated carbon the better it is at giving the water a better quality within a short time frame, but using a smaller particle sized activated carbon would require changing the filter after use every 2-3 weeks, the cost of using smaller particle sized activated carbon will then be high. Lager particle size filters are advisable to use in water treatment because it can easily be reactivated and re-used.

In terms of speed, the 590 μ m particle size activated carbon was quicker in the water treatment process and more effective overall. To obtain a similar purity with the 850 μ m particle size activated carbon, the water will have to be treated more than once.

Overall the $850\mu m$ particle sized activated carbon is recommended due to its relative cheap cost on a mass scale and ability to provide relatively clean water for a large community at a reasonable cost.

ACKNOWLEDGMENT

The authors are grateful to Engr. Prof. Ezeh, J. C, Engr. Dr. Osuagwu, Engr. Dr. Owuse, and Engr. Dr. Arimanwa, J. I, for their help in this research. The Management of Federal University of Technology, Owerri, Nigeria is also appreciated for providing an enabling environment for the study and research.

REFERENCES

- Akinyemi O.P & Taiwo E.A. (2004). "Production of activated carbon from agricultural wastes". Proceedings of Nigerian Society of Chemical Engineers held in Warri- Nigeria [p.1]. NSE.
- [2] Aleksandra M.K and Dick K. (2004). "Optimization and significance of ATP analysis for measuring active biomass in granular activated carbon filters used in water treatment
- [3] Babel B and Jurewicz C. (2004). "The use of microwaved carbonization irradiation system to produce activated carbon with coconut shells using potassium hydroxide (KOH) as an activating agent". IJPPT, Vol 12. Pg. 31-32.
- [4] Bellen, G., Anderson M. and Gottler R. (1984). "Management of Point-of-Use Drinking Water Treatment Systems" Final Report. U.S. Environmental Protection Agency: Water Engineering Research Laboratory, Office of Research and Development. Cincinnati. 1-2.
- [5] Bouchelta C, Medjram M.S, Bertrand O. and Bellat J.P. (2008). "Preparation and characterization of activated carbon from date stones by physical activation with steam". Journal of Analysis Applied Pyrolysis 82: 3-4.
- [6] Chandra TC, Mirna M.M, Sunarso J, Sudaryanto Y& Ismadji S. (2009). "Activated carbon from durian shell: preparation and characterization". Journal of Taiwan Institute Chemical Engineering 40: 3-4.
- [7] Crapper C.C and Miller N. et al. (1984). "Developments in Thermochemical Biomass Conversion". London: Blackie Academic &professional. Vol.1.
- [8] Cooney H.N, David O. (1999). "What to Expect from Carbon Cartridges". Water Technology 16 (11):76-79: 30-31.
- [9] Daud W.M & Ali W.S. (2004). "Comparison on pore development of activated carbon produced from palm shell and coconut shell". Bioresource Technology 93: 4.
- [10] Diebold, J.P. & Bridget, A.V. (2014). "Overview of Fast Pyrolysis of Biomass for The Production of Liquid Fuels in Bridgwater", A.V. & Boocock, D. G. B. (eds.): 6-7.
- [11] Diets V.R. (1990). "Bibliography of solid adsorption and United States cane sugar refineries and bone char manufactures". National Bureau Standards. Washington DC. 1.
- [12] Dileck C, Oznuh A.Y. (2008). "Production and characterization of activated carbon from bituminous coal through chemical activation". Afr. J. Biotechnology., 7(2): 2.
- [13] Elliott CG, Colby T, Kelly T.M, Hicks H.G. (1989). "Charcoal lung Bronchiolitis obliterans after aspiration of activated charcoal". Chest 96 Vol. (3): 672–4. doi:10.1378/chest.96.3.672. PMID 2766830. 14-15.
- [14] Elliot C, Colby T, Hicks H. (1989). "Charcoal lung bronchiolitis Obliterants after aspiration of activated charcoal". Vol. 1.
- [15] F. Rodriguez-Reinoso, Molina-Sabio M. (1998) "Textural and chemical characterization of micro-porous carbons: Adv. Colloid Interface Sci. Vol. 2.
- [16] Gadgil, Ashok. (1998). "Drinking Water in Developing Countries." Annual Review of Energy and the Environment 23: 6.
- [17] Guo J and Lua A.C. (2001). Characterization of adsorbents prepared from oil palm shell by thermal activation for removal of gaseous pollutants. Material Sci. 55: 2-3.

- [18] Guo J and Rockstraw D.A. (2006). Physicochemical properties of carbons prepared from pea nut by phosphoric acid activation. Bio resource Technology, 98: 14.
- [19] Haimour N.M, Emeish N. (2006). Utilization of date stones for production of activated carbon using phosphoric acid. Waste management. 26(6): 2.
- [20] Hasbrouck, S. (1986). "Removing Radon from Water Using Granular Activated Carbon Adsorption". Information Digest, University of Maine at Orono. 4.
- [21] Hu Z., Srinivasan M.P. (2001). "Mesoporous high surface area activated carbons". Microporous Mesoporous Mater, 34: 3.
- [22] Inamullah B, Khadija Q, Kazi RA, Abdul KA. (2008). "Preparation and characterisation of chemically activated almond shells by optimization of adsorption parameters for removal of chromium vi from aqueous solutions". World Academy of Science, Eng. Technol., 34: 1.
- [23] Kamrin M, Nancy H, Barry Christian, D.B and Frank D. (2016)."Home Water Treatment Using Activated Carbon".Cooperative Extension Service, Michigan State University. 4.

- [24] Ncibi M.C, Jeanne-Rose V, Mahjoub B, Jean-Marius C, Lambert J, Ehrhardt J.J, Bercion Y, Seffen M & Gaspard S. (2009). "Preparation and characterization of raw chars and physically activated carbons derived from marine Posidonia oceanica". (L.) fibres. Journal of Hazardous Material 165: 2.
- [25] Sang CK, In K.H, Kyung A.P. (1997). Preparation and performance of the Briquette type activated carbon based bituminous coal. J. Ind. Eng. Chem., 3(30): 1&2.
- [26] Srinvaskannan C, Mohammad Z.A. (2004). Production of activated carbon from rubber wood sawdust. Biomass Bioenergy. 27: 2.
- [27] Timothy W, Lambert Charles F.B, Holmes S.E. (1996).
 "Adsorption of microcystin-LR by activated carbon and removal in full scale water treatment". Vol.30 (6):1411–1422. R. E. Sorace, V. S. Reinhardt, and S. A. Vaughn, "High-speed digital-to-RF converter," U.S. Patent 5 668 842, Sept. 16, 1997.

IJSER