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# Green synthesis of ZnO coated hybrid biochar for the synchronous removal of ciprofloxacin and tetracycline in wastewater†

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Preparation of biochar from kaolinite and coconut husk (KCB) and further activated with HCl (KCB-A) and KOH (KCB-B) via a microwave technique for the remediation of ciprofloxacin (CIP) and tetracycline (TET) from water was carried out. Characterization using scanning electron microscopy, energy dispersive X-ray, Fourier transform infrared spectroscopy and X-ray diffraction showed the successful synthesis of functionalized biochars. Batch adsorption experiments demonstrated the potential of the adsorbents for fast and efficient removal of CIP and TET from solution. The adsorption capacities were found to be 71, 140 and 229 mg g<sup>-1</sup> for CIP and 118, 117 and 232 mg g<sup>-1</sup> for TET removal on KCB, KCB-A and KCB-B, respectively. For KCB, KCB-B and KCB-B, CIP adsorption best followed the pseudo second order kinetic model (PSOM), pseudo first order kinetic model (PFOM) and intraparticle diffusion (IDP) respectively. TET adsorption followed PSOM for KCB, IPD for KCB-B and PFOM for KCB-A. CIP adsorption on KCB, KCB-A and KCB-B best fit the Temkin, Langmuir and Brouers–Sotolongo isotherms, respectively, and TET adsorption on KCB best fit Brouers–Sotolongo while KCB-A and KCB-B best fit Langmuir–Freundlich. Adsorption of both contaminants was thermodynamically feasible showing that these materials are excellent adsorbents for the treatment of pharmaceuticals in water.

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## 1. Introduction

Antibiotics are indispensable in the treatment of infectious diseases of humans and animals. Of the various antibiotics currently in use, ciprofloxacin, a member of the fluoroquinolone family, and tetracycline, a broad spectrum antibiotic, are the most commonly used in the treatment of bacterial infections and as animal growth promoters<sup>1,2</sup> and both have been classified as high priority emerging organic pollutants by the World Health Organisation (WHO).<sup>3,4</sup>

As a result of their widespread use, coupled with inadequate treatment of wastewater and their discharge into the environment, they have been detected in various systems such as soil, sediment, surface, ground and drinking waters including living tissues and this has raised concerns about their toxicity.<sup>5,6</sup> Even at low concentrations they have resulted in increase in antibacterial-resistant strains of organisms along with other adverse health effects such as stomatitis, leukopenia. In

addition, their presence in the environment results in aquatic toxicity, genotoxicity.<sup>6,7</sup> Increase in population and continued usage coupled with indiscriminate discharge into the environment is expected to cause further rise in these species. Once these pollutants reach water sources, they are able to transfer to other non-point sources. Antibiotics especially are difficult to degrade and 30–90% of administered doses remain undegraded in an organism.<sup>6</sup> The widespread use of antibiotics as promoters of good health ensures that they are continually used and discharged into the environment either in the form in which they were used or in their sometimes more toxic metabolite form. Their toxic effects coupled with their incessant input into the environment have led to policy makers, government agencies and the scientific community to promote technologies and strategies for the treatment of waters polluted by these substances.<sup>8</sup>

Wastewater treatment plants using conventional treatment methods have been found to be inadequate in the complete removal of these pollutants.<sup>6</sup> Advanced techniques have shown promising success in the removal of these pollutants. These methods are as varied as the physicochemical characteristics of each antibiotic compound. Oxidation processes have the disadvantages of high cost, incomplete mineralization and the production of even more toxic metabolites.<sup>6</sup> Low pressure membranes such as microfiltration, ultrafiltration and loose nanofiltration are limited due to their pore sizes while reverse

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osmosis, nanofiltration techniques have been found to have incomplete rejection as high as 95%.<sup>9</sup> Adsorption however does not result in generation of toxic by-products; is efficient; versatile; offers ease of operation and employs the use of low-cost materials as adsorbents<sup>10</sup> and is easily incorporated into current wastewater treatment processes.<sup>11</sup>

Biochar has attracted the attention of researchers as an adsorbent material due to its high porosity, surface area and adsorption capabilities,<sup>12</sup> the availability of diverse feedstock and ease of production,<sup>13</sup> low-cost and environmentally friendly nature.<sup>4</sup> Several materials have been utilized as starting precursors for biochar manufacture including mangosteen shells,<sup>14</sup> poplar wood,<sup>15</sup> sewage sludge,<sup>16</sup> corn straw<sup>17</sup> and sugarcane bagasse.<sup>18</sup>

In this study, biochar material prepared from a hybrid of two low-cost precursors was functionalized *in situ* followed by further activation to improve the physicochemical properties of the adsorbents while their efficiency for the simultaneous removal of ciprofloxacin and tetracycline under a range of experimental parameters was evaluated.

## 2. Materials and method

### 2.1. Materials

Kaolinite clay was obtained from Federal Institute of Industrial Research Oshodi (FIRO), Lagos State, Nigeria. Coconut husks were obtained from local markets in Ibadan, Oyo State, Nigeria. Ciprofloxacin (CIP), tetracycline (TET) and zinc chloride were obtained from Sigma-Aldrich. All chemicals were used as received.

### 2.2. Synthesis of hybrid biochar (KCB)

Firstly, the clay was purified according to the method reported by ref. 19 (see ESI†). Coconut husks were washed with distilled water, sun-dried and milled. Kaolinite clay, biomass and ZnCl<sub>2</sub> were thoroughly mixed in a mass ratio of 1 : 1 : 2 respectively. 10 mL of 0.1 M NaOH was added with constant stirring for 12 h after which it was transferred to an oven at 150 °C for 24 h to allow for impregnation. Then, the impregnated mixture was heated under microwave irradiation at 540 W for 10 min after which it was washed several times with distilled water until pH 7 to remove excess NaOH and ZnCl<sub>2</sub>. The obtained sample was dried at 105 °C, cooled and stored in an airtight container labeled KCB.

### 2.3. Activation of KCB

1 g each of KCB was added to 50 mL of either 0.1 M KOH or 2 M HCl in a beaker. The mixtures were stirred constantly for 1 h after which they were filtered and washed severally until neutral pH. This was followed by drying at 105 °C. The samples obtained were stored and labelled KCB-A and KCB-B for acid and alkali activated samples respectively.

### 2.4. Surface characterization

The mineralogical compositions and crystallinity of the adsorbents were determined *via* X-ray diffraction (XRD) with patterns

recorded on a BRUKER AXS (Germany) D8 Advance X-ray Diffractometer and operated in a continuous  $\Theta$ - $\Theta$  scan in locked coupled mode with Cu-K radiation. Surface characteristics and elemental compositions of the adsorbents were determined using scanning electron microscopy (SEM: BRUKER) with an operating voltage of 20 kV equipped with energy dispersive X-ray (EDX). Fourier transformed infrared spectroscopy (FTIR) spectra of the biochars were obtained using Shimadzu FT-IR 8400S (class 1, Laser product) Spectrophotometer. The pH at point of zero charge (pHpzc) was determined by using the salt addition method as described by ref. 20.

### 2.5. Batch adsorption experiments

Equilibrium adsorption experiments were performed in batch mode by agitating 50 mg of each adsorbent with 10 mL aqueous solution of either CIP or TET with concentrations ranging from 10–150 mg L<sup>-1</sup> for 2 h to determine the equilibrium adsorption capacity of the adsorbents. The solutions were filtered and the residual concentrations of the pollutants in solution were determined using a UV-vis spectrophotometer (Shimadzu UV-1650 pc) at wavelengths of 277.5 nm and 355 nm for CIP and TET respectively, determined after scanning. The adsorption capacity for each pollutant was estimated using eqn (1) below:

$$q_e = \frac{(C_o - C_e) \times V}{m} \quad (1)$$

where  $C_o$  and  $C_e$  are the initial and equilibrium concentrations of the CIP and TET (mg L<sup>-1</sup>),  $q_e$  is the amount of pollutant adsorbed (mg g<sup>-1</sup>),  $M$  is the mass of the dry adsorbent used (g) and  $V$  is the volume of the solution (litres). The non-linear forms of Langmuir (L), Freundlich (F), Dubinin–Radushkevich (D–R), Temkin (T), Langmuir–Freundlich (L–F) and Bruoers–Sotolongo (B–S) isotherm equations were applied to the data obtained to describe the nature and mechanism of adsorption.

Kinetic studies were carried out by reacting 500 mg of each adsorbent with 100 mL of CIP or TET. 10 mL of each solution was withdrawn using a syringe at predetermined time intervals (within 1 h). Results were analysed with five kinetic models: the pseudo first-order (PFOM), pseudo second-order (PSOM), Bruoers–Sotolongon fractal kinetics (BSF), intraparticle diffusion (IDP) and Elovich ( $E$ ) kinetic models. Details of both isotherm and kinetic models are given in Tables S1 and S2 (ESI†).

## 3. Results and discussion

### 3.1. Surface characterization

pHpzc is the pH that indicates the net surface charge on adsorbents in solution. The ease of adsorption of molecules or ions to the surface of adsorbents is dependent on the pHpzc of such material. The information provided by the pHpzc is useful in understanding and manipulating the best pH at which a particular adsorbate can be removed from solution by altering the net surface charge of a material to be used as an adsorbent in solution. KCB, KCB-A and KCB-B were found to have pHpzc of 6.8, 6.0 and 6.6 respectively (Fig. 1). Below these values, the



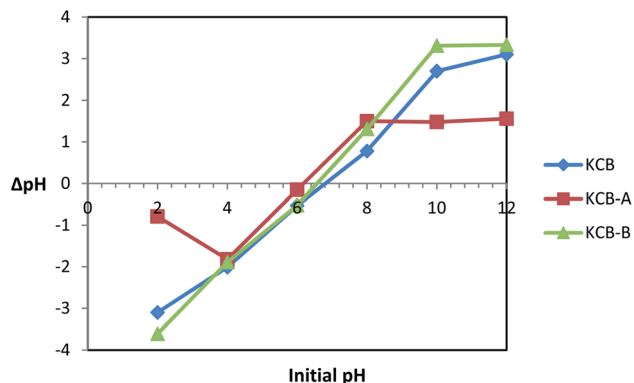


Fig. 1 pHpzc plot for KCB, KCB-A and KCB-B.

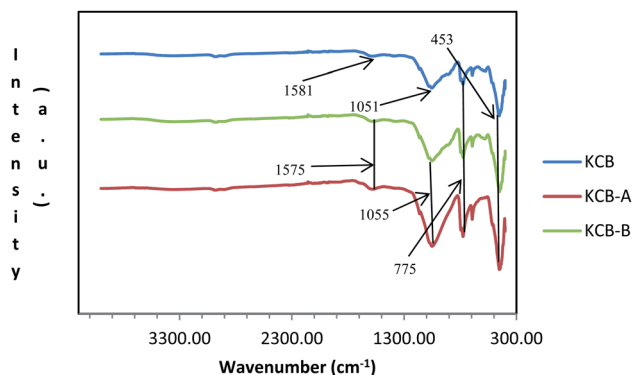


Fig. 2 FTIR spectra of KCB, KCB-A and KCB-B.

surface charge on the adsorbent in solution is positive and above this value, the charge on the adsorbent is negative.

Activation was found to alter the surface chemistry of KCB. KCB-A showed reduced pHpzc values compared with KCB. This is attributed to acid treatment of the sample which lowered its pHpzc value due to the introduction of functional groups on the surface of the adsorbent that increased the acidic surface oxides.<sup>21</sup>

Fig. 2 shows the FTIR spectra of the adsorbents. The loss of the external hydroxyls usually associated with kaolinite can be due to the effect of heat during the modification process.<sup>22</sup> The absorption shown by the very weak band at around  $1580\text{ cm}^{-1}$

can be attributed to H–O–H bending of kaolinite. The presence of Zn–O absorption peak in the adsorbent is shown by absorption at around  $453\text{ cm}^{-1}$  which is attributed to the  $\text{ZnCl}_2$  used during the modification. The broad band at  $1055\text{ cm}^{-1}$  is due to the presence of Al–Si–O stretching of amorphous aluminosilicates which shifted to  $1051\text{ cm}^{-1}$  in KCB-A and KCB-B. The peak observed at about  $775\text{ cm}^{-1}$  indicates Si–O vibrations of the quartz while the peak appearing at about  $800\text{ cm}^{-1}$  indicates the presence of Al–O–H bond vibrations. Activation with either acid or alkali was found to increase the intensity of these peaks. Similar results were reported<sup>22</sup> for the synthesis of a kaolinite/pawpaw composite (HYCA).

The SEM images of KCB, KCB-A and KCB-B as shown in Fig. 3(a–c) show that the adsorbent pores have irregular shapes and sizes majorly ellipsoidal and are heterogeneously distributed. Their sharp edges also suggest good crystallinity.<sup>23</sup> The adsorbents are highly porous with pore sizes ranging from  $7\text{ }\mu\text{m}$  to  $20\text{ }\mu\text{m}$  which indicates the heterogeneous nature of the adsorbents. The presence of Zn is seen from the scattering of particles on the surface of the adsorbents showing its aggregation on the surface of the adsorbent rather than incorporation into the structure. Surface structure of KCB changed slightly after modification with both acid and alkali. Activation also appears to unblock some pores resulting in the generation of more visible pores than that seen for KCB.<sup>24</sup>

Elemental (EDX) analysis (Table 1) confirms the carbonization of the precursors<sup>25</sup> which resulted in a higher carbon content than kaolinite ( $0.16\%$ )<sup>22</sup> and coconut husk ( $20\%$ ).<sup>26</sup>

Table 1 Chemical elements present in KCB, KCB-A and KCB-B as determined by EDS analysis (atom%)

Atoms	Lines	KCB	KCB-A	KCB-B
C	K-series	43.20	89.27	80.58
Si	K-series	38.71	4.11	1.50
O	K-series	7.26	5.99	6.16
Zn	K-series	5.15	—	7.41
Fe	K-series	3.86	—	1.36
Ca	K-series	1.20	—	0.84
Cl	K-series	0.62	0.54	0.57
K	K-series	—	—	1.58
Al	K-series	—	0.09	—
	Total	100	100	100

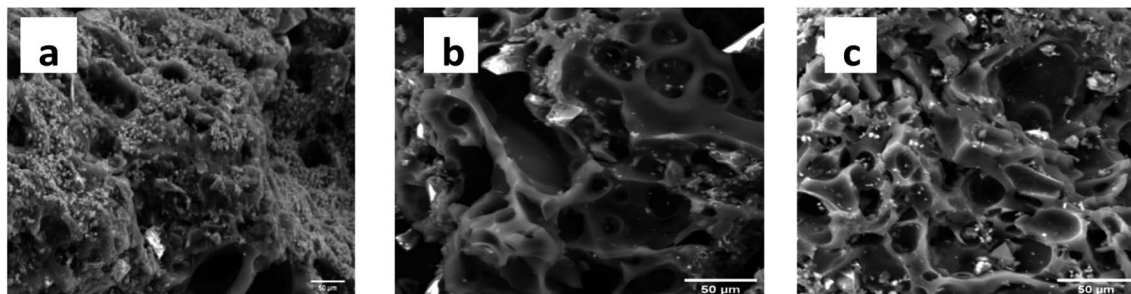


Fig. 3 SEM images of (a) KCB (b) KCB-A and (c) KCB-B.



Activation of KCB further increased the amount of carbon by approx. 100% to give 89% and 81% in KCB-A and KCB-B respectively which shows that activation improved upon the surface properties of KCB-A and KCB-B.

XRD further confirms the result of FTIR (Fig. 4). It displays characteristic diffraction peaks of kaolinite at  $2\theta$  of  $20.81^\circ$  and  $26.61^\circ$  for KCB and  $20.80^\circ$  and  $20.57^\circ$  for both KCB-A and KCB-B corresponding to the reflections from (110) and (002) plane respectively. This suggests that the crystal structure of kaolinite is not affected by modification with the biomass or the synthesis process. Activation of KCB with either acid or alkali resulted in an increase in the intensity of the peaks. This is attributed to structural disorder which affected the crystallinity of the sample.<sup>27</sup> The full width at maximum height (FWMH) is used to determine the  $L_c$  which is a measure of crystallite size and obtained using the Debye-Scherrer equation (eqn (2)).<sup>28</sup>

$$L_c = \frac{k\lambda}{\beta_{002} \cos \theta_{002}} \quad (2)$$

The values of 5.4, 5.4 and 6.1 nm were obtained as the crystallite sizes for KCB, KCB-A and KCB-B respectively (Fig. 5).

## 3.2 Adsorption performance

### 3.2.1 Effect of pH.

pH influences the characteristics of both the adsorbate and adsorbent as well as the interactions between them which changes as pH changes.<sup>29</sup>

The plots of the effect of pH (2–12) on the adsorption of CIP and TET with an initial concentration of  $40 \text{ mg L}^{-1}$  are shown in Fig. 6. For the 3 adsorbents, the removal of TET was higher in basic medium (pH 10) than in acidic medium. At this pH, the surfaces of these adsorbents are expected to be negatively charged. TET is negatively charged at this pH, occurring as a divalent anionic species. It is expected that under these conditions, repulsion mechanism will be activated leading to decrease in adsorption. However, there was an increase in adsorption showing a higher affinity of the adsorbents for this divalent form of TET. It is also a testament to the fact that electrostatic interactions is not at play between TET and the adsorbents. Further increase in pH from 10 to 12 resulted in

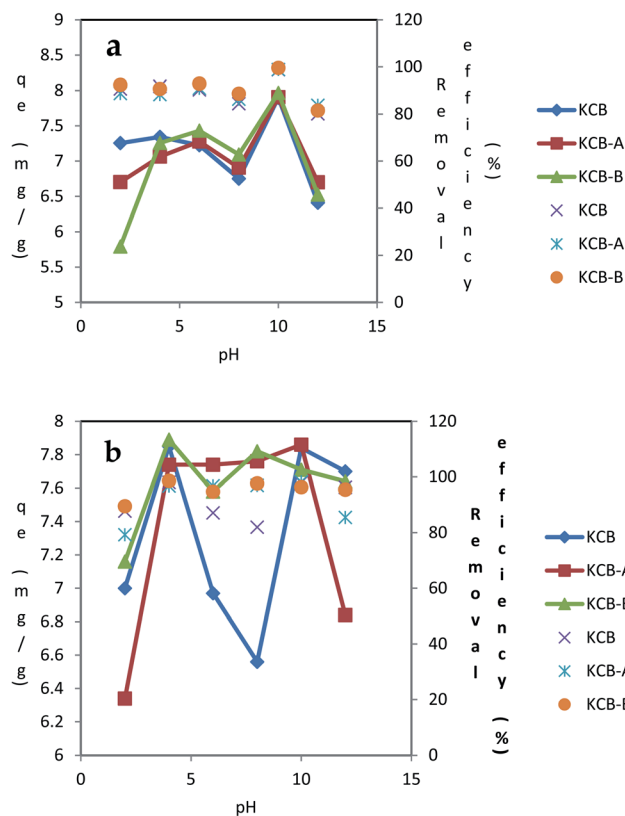


Fig. 5 Effect of pH on the adsorption of (a) TET (b) CIP using KCB, KCB-A and KCB-B (initial concentration =  $40 \text{ mg L}^{-1}$ ; contact time = 60 minutes; adsorbent dosage =  $0.05 \text{ g}$ ; temperature =  $303 \text{ K}$ ).

a decrease in adsorption. For CIP adsorption on KCB-A, there was an increase in adsorption up to pH 4. Between pH 4 and 10, there was no significant change in adsorption as the solution moved from being acidic to alkaline. At pH 12 however, there was a sharp decline in adsorption with further increase in pH. The  $\text{pH}_{\text{pzc}}$  of KCB-A is 6.0. Below this pH, the surface of this adsorbent is positively charged.

On the other hand, at pH below 6.1, CIP exists as a cation, as an anion beyond 9 and as a neutral molecule between 6.1 and 8.9. With this information, it is expected that there will be a decline in adsorption at pH below 6. Based on experimental observations, this was not the case. Rather than a decline in adsorption at pH less than 6, there was a steady rise in adsorption up to pH 4. This result shows that adsorption could not have been due to electrostatic interactions between the adsorbent and the adsorbate. This phenomenon has been suggested to be due to the suppression of electrostatic interactions by the hydrophobicity of CIP.<sup>30</sup> Between pH 6 and 8, the adsorption was steady, averaging 97%. Non-polar adsorbents are capable of adsorbing organic pollutants which exist in molecular form<sup>31</sup> which could explain the higher adsorption at those pH values. Also at pH beyond 9, CIP is anionic while the adsorbent is negatively charged. This should result in a decline in adsorption as a result of electrostatic repulsion. This led to a decrease in adsorption from pH 10. However, this decline in adsorption from pH 10 could be attributed to decrease in



Fig. 4 XRD spectra of KCB, KCB-A and KCB-B.



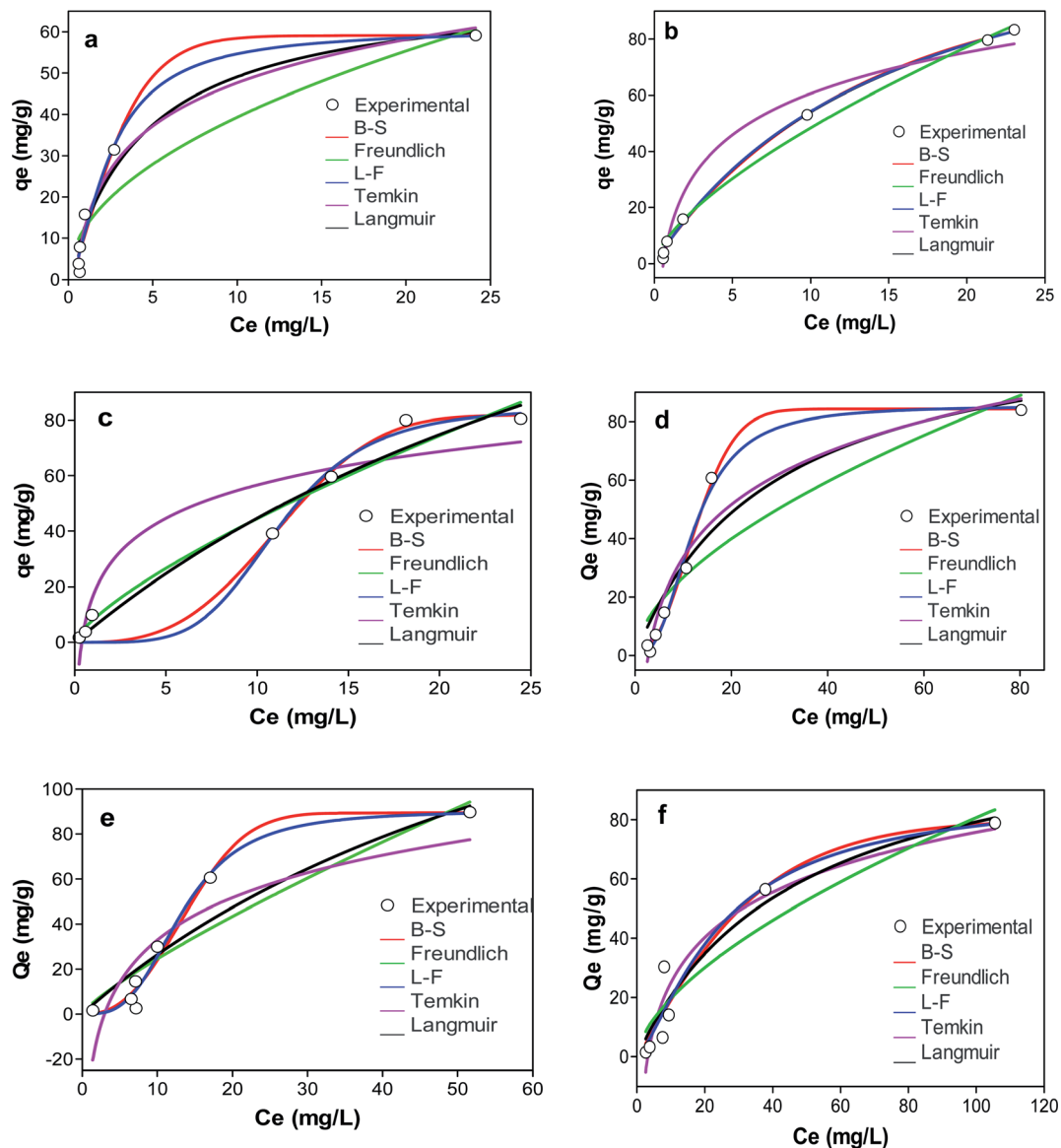


Fig. 6 Graphs showing the adsorption isotherms of (a) KCB (b) KCB-A (c) KCB-B on CIP (d) KCB (e) KCB-A and (f) KCB-B on TET.

hydrophobicity of CIP rather than electrostatic repulsions between the adsorbent and adsorbate as hydrophobicity has been found to reduce with the formation of anions. KCB-B on CIP showed an increase in adsorption up to pH 4. Between pH 4 and 8 where the CIP molecules are expected to be neutral, there was a decline in adsorption capacity. At that pH too, the surface of the adsorbent is positively charged which could be responsible for the decline in adsorption because beyond the pHPzc of the adsorbent where its surface becomes increasingly negative and deprotonation of the second amine group of CIP begins to occur, adsorption increases slightly. As soon as the negative charge on the adsorbent is fully established and CIP becomes fully deprotonated at pH beyond 8, adsorption capacity declines. KCB also shows a similar adsorption pattern to KCB-B except that the decrease extends from pH 4 to 8. There was increase in adsorption capacity from pH 8 to 10 followed by

a final decline in adsorption. Adsorption of CIP and TET onto these adsorbents have been attributed to take place *via* a series of mechanisms notably  $\pi$ - $\pi$  interactions, hydrophobic interactions, London dispersion forces, hydrogen bonding or van der Waals forces.<sup>32</sup>

**3.2.2 Equilibrium isotherm studies.** To optimise the design of an adsorption system, it is important to establish the most appropriate correlation for the adsorption equilibrium curves. Langmuir, Freundlich, Dubinin-Radushkevich and Temkin (two-parameter models); Langmuir-Freundlich (L-F) and Bruoers-Sotolongo (three-parameter models) were used to interpret the adsorption data. Their plots are shown in Fig. 6a-f and the values of their parameters are presented in Table S1.† Among the two parameter isotherms, with respect to the equilibrium data, Langmuir isotherm gave the best fit with the highest  $R^2$ , 0.999 and 0.985 for CIP adsorption on KCB-A and



KCB-B respectively followed closely by Freundlich ( $R^2 = 0.996$  and  $0.984$ ) while Temkin ( $R^2 = 0.989$ ) gave the best for adsorption on the inactivated composite followed closely by Langmuir ( $R^2 = 0.977$ ). This suggests that CIP adsorption followed a series of complex chemisorption mechanisms.<sup>7</sup> The Langmuir maximum monolayer CIP uptake by KCB, KCB-A and KCB-B are  $71.1$ ,  $139.9$  and  $228.7$   $\text{mg g}^{-1}$  respectively. KCB-B and KCB-A had better rates of adsorption than KCB for CIP as can be seen in their adsorption capacities ( $228.7$  and  $139.9$   $\text{mg g}^{-1}$ ). This can be attributed to an increase in the functional sites on the activated adsorbents brought about by an introduction of functional groups that can serve as adsorption sites. Adsorption of the TET on the other hand, showed the monolayer adsorption capacity to be  $118$ ,  $116.6$  and  $231.9$   $\text{mg g}^{-1}$  for KCB, KCB-A and KCB-B respectively. The best model among the two-parameter

isotherms to fit the experimental data for KCB and KCB-B is the D-R model (Fig. 7a-f) ( $R^2 = 0.974$  and  $0.971$ ) followed closely by Temkin. For adsorption of TET on KCB-A, Langmuir ( $R^2 = 0.965$ ) followed by Temkin ( $R^2 = 0.962$ ) best fit the data. Like CIP adsorption, the adsorption of TET can be said to involve complex chemisorption mechanisms.

The D-R isotherm is applied to estimate the porosity apparent free energy and the characteristics of adsorption. The value of  $E$ , which represents the free mean energy ( $\text{kJ mol}^{-1}$ ) was obtained (Table S1†). An  $E$  value  $< 8$   $\text{kJ mol}^{-1}$  is indicative of physical adsorption.<sup>33</sup> The value of  $E$  for the adsorption of TET on the adsorbents and KCB on CIP were less than  $8$   $\text{kJ mol}^{-1}$  showing that the mechanisms of adsorption were physical in nature. This was corroborated by the enthalpy values obtained for thermodynamics studies (Table S3†).

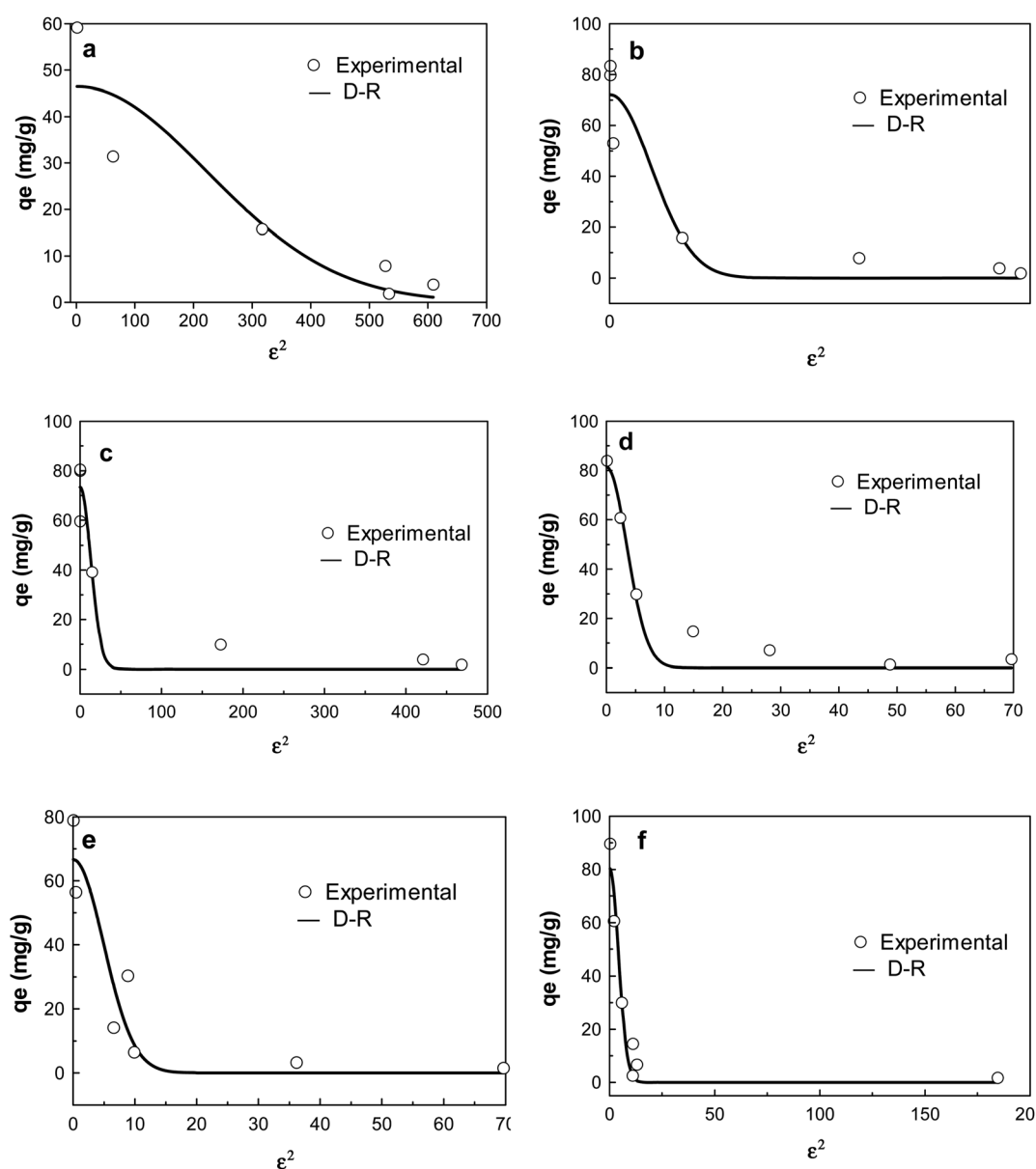


Fig. 7 D-R isotherm plots of adsorption of CIP on (a) KCB (b) KCB-A (c) KCB-B; TET on (d) KCB (e) KCB-A and (f) KCB-B.



The heterogeneity factor ( $n$ ) in the Freundlich model indicates the degree of heterogeneity and hence how favourable the adsorption is.<sup>33</sup> The value of Freundlich constant,  $n$  was greater than 1 for all adsorbents on both CIP and TET indicating that the adsorption was favourable at studied conditions. This is corroborated by the  $n$  factor predicted using L-F model and further buttressed by the value of the dimensionless constants,  $R_L$  which were all less than one. The surface heterogeneities of the adsorbents are confirmed by the presence of various functional groups present on the adsorbents (Fig. 2).

The experimental data were further modelled with three-parameter isotherm models in a bid to improve the precision of analysis. Comparison of the  $R^2$  showed better fitting when compared with the two parameter isotherm models. Of the three models however, L-F isotherm, a model that combines the features of Langmuir and Freundlich isotherms was the best fit for the adsorption of CIP ( $R^2 > 0.98$ ) while B-S isotherm was the best fit for adsorption of TET ( $R^2 > 0.997$  and  $R^2 > 0.969$  for KCB and KCB-A respectively) except by KCB-B whose adsorption best fit L-F ( $R^2$ ; error analysis). L-F is used for predicting the heterogeneous nature of the adsorption system while circumventing the limitation of high adsorbate concentrations of Freundlich.<sup>34</sup> This shows that both monolayer and multilayer are occurring simultaneously within the system, indicating that the adsorption could be driven by more than one force of attraction. Studies have shown that for most organic pollutants, a number of bonding mechanisms such as electrostatic interactions, ion-exchange,  $\pi$ - $\pi$  interactions, H-bonding or hydrophobic interactions occur in tandem during the adsorption process.<sup>34,35</sup>

The fit of the experimental data with B-S isotherm is used to predict the adsorption capacity of the adsorbents for complex, heterogeneous systems involving sorbent materials with different chemical characteristics.<sup>36</sup> The high  $R^2$  values ( $R^2 > 0.999$ ) observed for both CIP and TET for this model show that the adsorbents possess heterogeneous active sites. The adsorption capacities obtained for these adsorbents are seen to

be higher when compared with those obtained for other agro-genic materials in previous studies as can be seen in Table 2 indicating the higher potential of these adsorbents for CIP and TET removal.

**3.2.3 Kinetic studies.** Kinetic models are mathematical equations that help to interpret the kinetic data obtained to describe the adsorption mechanism. The parameters of the kinetic models used in studying the experimental data are presented in Table S2 (see ESI†). As seen in Fig. 8 the adsorptions of CIP and TET on the adsorbents were fast with most adsorption occurring within the first 15 minutes. This is particularly important in industrial applications as it reduces adsorption time thereby reducing operation costs.<sup>43</sup> The residual root mean square error given by eqn (3) below was used in evaluating the best fitted model to the kinetic data. Using the error function, decreasing values of RSE indicates better fitting to the model.<sup>44</sup>

$$\sqrt{\frac{1}{n-2} \sum_{i=1}^N (q_{e(\text{exp})} - q_{e(\text{cal})})^2} \quad (3)$$

$q_{e(\text{exp})}$  and  $q_{e(\text{cal})}$  represent the experimental and calculated  $q_e$  values,  $N$  is the number of observations in the experimental data.

Of the kinetic models studied, the PFOM best described the adsorption of KCB and KCB-B for CIP adsorption and KCB-A and KCB-B for TET adsorption. Pseudo first order kinetic model works best at high concentrations showing that a high concentration of pollutants can be successfully removed.<sup>7</sup>

Though PFOM fits only over the initial 20–30 minutes of adsorption,<sup>31</sup> it is suitable for the removal of these pollutants as most of the pollutants were removed within that frame of time. On the other hand, PSOM gave the best fit for the data on CIP adsorption by KCB-A as well as TET adsorption by KCB. This indicates that the adsorption worked well over the whole range of adsorption time and follows the chemisorption process. It can also be said that the adsorption capacity was positively related to the amount of active sites on the adsorbent. Higher

Table 2 Comparison of adsorption capacities for CIP and TET uptake by agro-genic based adsorbents

Adsorbent	Pollutant	$q_{\text{max}}$ ( $\text{mg g}^{-1}$ )	Ref.
Pistachio shell powder coated ZnO nanoparticles	CIP	98.72	37
Eu doped SrAl <sub>2</sub> O <sub>4</sub> composite	CIP	7.10	38
Reduced GO/magnetite composite	CIP	18.98	16
Poly(1-trimethylsilyl-1-propyne)	CIP	33.10	39
Aluminum-pillared kaolin sodium alginate beads	CIP	68.36	40
Clay/coconut husk biochar	CIP	71	This study
HCl-activated clay/coconut husk biochar	CIP	140	This study
KOH-activated clay/coconut husk biochar	CIP	232	This study
Pistachio shell powder coated ZnO nanoparticles	TET	92	37
Eu doped SrAl <sub>2</sub> O <sub>4</sub> composite	TET	26	38
KOH-activated reed biochar	TET	146	41
Wood biochar	TET	120	7
$\alpha$ -Fe <sub>2</sub> O <sub>3</sub> /reduced graphene oxide	TET	39	42
Clay/coconut husk biochar	TET	118	This study
HCl-activated clay/coconut husk biochar	TET	117	This study
KOH-activated clay/coconut husk biochar	TET	232	This study



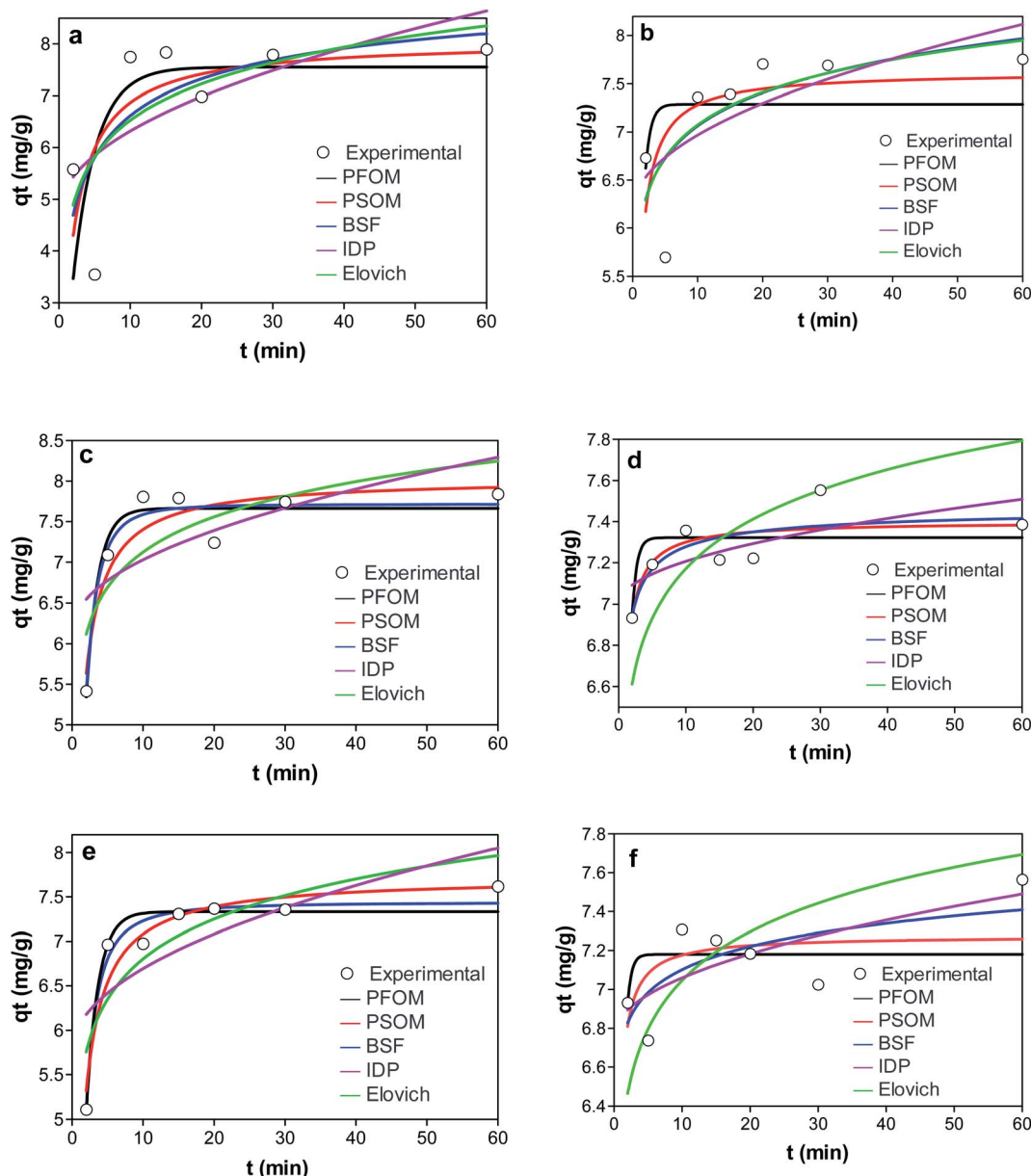


Fig. 8 Graphs showing the adsorption kinetics of (a) KCB (b) KCB-A (c) KCB-B on CIP (d) KCB (e) KCB-A and (f) KCB-B on TET.

values of  $h$  observed for the activated adsorbents are attributed to additional adsorption sites created by increasing functional groups brought about by chemical activation. This is further confirmed by the fitting with intraparticle diffusion (IPD) kinetic model. The rate constants as an indication of how fast the adsorption is show that the rates of adsorption by the activated adsorbents were faster than the inactivated adsorbent. It has been shown that if  $K_1$  is greater than  $K_2$ , the adsorption is restricted by internal diffusion.<sup>45</sup> The results in Table S2† suggest that internal diffusion occurred in addition to surface adsorption. If intraparticle is the sole limiting step, a plot of  $q_t$  versus  $t^{0.5}$  should pass through the origin. The line of fit did not pass through the origin, showing that intraparticle diffusion alone is not the sole rate limiting step and that surface diffusion

could also have contributed to the process<sup>46</sup> making the mechanism all the more complex. The high  $K_{IPD}$  values obtained for the adsorbents could be proportional to the number of functional sites produced by activation and this is supported by the  $h$  values reported earlier. The values of  $C$  obtained are an indication of the boundary layer thickness between the adsorbate and adsorbent.<sup>47</sup> The low value of  $C$  obtained for the adsorption shows that they are less affected by boundary layer with low resistance to external mass transfer. KCB-A and KCB-B were least affected by boundary layer when compared to KCB showing that activation had an impact on the mechanism of transfer of adsorbate to adsorbent.<sup>44</sup> Brouers–Sotolongo fractal kinetic model, BSF is used to describe complex adsorption systems by predicting that the adsorption followed a first order





mechanism. This closeness of the  $q_e$  values obtained from this model to the experimental  $q_e$  suggests that adsorption onto these adsorbents is *via* a complex mechanism involving various forces of attraction including the weak van der Waal forces.<sup>48</sup> This suggests that the adsorption sites of these adsorbents are heterogeneous in nature. On the other hand, Elovich model presented the worst prediction. This model does not propose a definite mechanism for the adsorption process.<sup>49</sup> This finding strengthens the prediction that the adsorption mechanism occurred *via* a series of complex mechanisms.

**3.2.4 Thermodynamics studies.** To examine the energy changes involved during the sorption process, the thermodynamic parameters of the Gibbs free energy ( $\Delta G^\circ$ ), the enthalpy change ( $\Delta H^\circ$ ), and the entropy change ( $\Delta S^\circ$ ) of the adsorption of CIP and TET were calculated using: Van't Hoff equation (see ESI†). The thermodynamic parameters as shown on Table S3† indicate that the adsorption of CIP and TET were thermodynamically feasible and spontaneous with negative  $\Delta G$  values. Lower temperature appeared to facilitate adsorption of TET better on the three adsorbents as the  $\Delta G$  value became increasingly positive at higher temperature. Increasing temperature provides energy which is needed to overcome the repulsive forces which could exist between solute molecules on both the solid and in solution thereby leading to increased adsorption at higher temperature. Also, there could be obstructions within the pores on the adsorbent which require sufficient kinetic energy to overcome. The opposite is true for adsorption of CIP giving the indication of an endothermic reaction. The resultant negative  $\Delta H$  value obtained for both CIP on KCB and KCB-A indicates the exothermic nature<sup>50</sup> of the adsorption on these materials.  $\Delta S^\circ$  value shows the degree of randomness of the system.

The positive  $\Delta S$  obtained for adsorption of CIP on KCB and KCB-A suggested an irreversible process and is indicative of structural changes happening within the system, an increase in randomness at the solid/solution interface upon adsorption. This reflects greater affinity of the adsorbates for the adsorbents with increase in temperature.<sup>33</sup>

## 4. Conclusion

This study reports the utilization of adsorbents prepared by microwave technology from low-cost, underutilized biomass in the successful removal of antibiotics pollutants from simulated wastewater. Pseudo first order kinetic model (PFOM) provided a better fit to the kinetic data for CIP adsorption KCB-B and TET adsorption on KCB-A. Pseudo second order kinetic model (PSOM) explained best the adsorption of CIP and TET on KCB while intraparticle diffusion (IDP) provided the best fit for CIP adsorption on KCB-A and TET adsorption on KCB-B. The adsorption isotherm models showed that the adsorbents prepared were very effective in the removal of CIP and TET with the chemically activated adsorbents showing greater affinity. Thermodynamics analyses indicate that the adsorption was spontaneous and favourable. These results demonstrate that these adsorbents provide alternative readily available, low-cost

and reusable materials that can successfully replace activated carbon in wastewater treatment.

## Author contributions

Abisola Egbedina: writing – original draft, methodology, investigation, formal analysis. Kayode Adebawale: conceptualization, supervision, project administration, writing – review & editing. Bamidele Olu-Owolabi: conceptualization, supervision, writing – review & editing. Emmanuel Unuabonah: conceptualization, methodology, writing – review & editing. Morenike Adesina: investigation.

## Conflicts of interest

The authors declare that they do not have any conflict of interest.

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