Adsorption-desorption of doxycycline using pyrophosphoric acid-modified

biochar derived from sesame stalk

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Abstract

The knowledge of the adsorption-desorption behavior of doxycycline on pyrophosphoric acid modified bio-24 char derived from sesame is limited. In this study, we examined the doxycycline sorption on pyrophosphoric 25 acid modified-sesame stalk biochar. The isotherm and kinetic sorption data showed pyrophosphoric acid 26 treatment enhanced the sorption of doxycycline on both modified and un-modified biochar; chemisorption 27 including EDA (π - π electron-donor-acceptor) interaction, pore filling and H-bonding might be the primary 28 mechanism. The maximum adsorption capacities for unmodified biochar were 87.6 mg g^{-1} , and 153.9 mg g^{-1} 29 for modified biochar. More than 90% of the adsorption capacity was retained after three successive adsorp-30 tion-desorption cycles. Besides, the strong electrostatic attraction between biochars and doxycycline might 31 largely explain the improved sorption capacity of doxycycline along with increasing pH from 5 to 9. The main 32 responsible mechanisms for the sorption of doxycycline included surface complexation, H-bonding, EDA 33 interactions, pore-filling effects. The results of the current study display that pyrophosphoric acid-modified 34 biochar has potential applications as an efficient, recyclable adsorbent for the removal of antibiotics from 35 wastewater for low-cost remediation. 36

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Keywords: Sorption, doxycycline, chemisorptions, modification, adsorbent

1. Introduction

Doxycycline (CAS no. 564-25-0) is a broad-spectrum bacteriostatic agent (antibiotic) synthetically 39 derived from a naturally occurring tetracycline produced by Streptomyces species bacteria known as 40 oxytetracycline; it is a member of the tetracycline class of antibiotics [1]. Doxycycline is one of the 41 widely used tetracycline antibiotics in the world. The occurrence of antibiotics in the natural envi-42 ronment can induce serious risk to human health and ecosystems by causing growth of antibi-43 otic-resistant bacteria, which can occur at low concentrations [2]. In recent times, the wide presence 44 of doxycycline in groundwater has attracted extensive attention [2, 22]. Thus, it is of great importance 45 to develop highly efficient and low-cost methods to eliminate antibiotics from contaminated-water. 46

There are several ways to eliminate antibiotics -which are often persistent compounds- from 47 waste-water, including bio-degradation, chemical oxidation, photo-degradation, and adsorption [3, 48 23]. Among them, adsorption techniques have numerous advantages over other techniques, such as 49 low energy consumption, high removal efficiency, environmental friendliness and easy operation [4]. 50 Recently, biochars have been manufactured by carbonizing the feedstock and then utilized as ad-51 sorbents for antibiotics [5]. Nonetheless, the adsorption capacities of unmodified biochars are usually 52 low, and their limitations include low density, low surface area and small particle size. To enhance the 53 adsorption capacity of biochar, several modification approaches, including chemical and physical 54 activation techniques have been suggested and used on biochars [6]. Most of the treatment methods 55 could efficiently alter the physico-chemical properties of the biochar surface, i.e. increased density of 56 functional groups, surface area, H/C, O/C ratios and altered porous structures [7, 8]. 57

Pyrophosphoric acid (aka diphosphoric acid, [(HO)₂P(O)]₂O) is a strong acid that can change phys-58 iochemical attributes of biochar. Therefore, pyrophosphoric acid was introduced in the current work 59 to modify the biochar, with the purpose of increasing the doxycycline sorption capacity. In this work, 60 sesame stalk biochar was manufactured via pyrolysis at 500°C. Then pyrophosphoric acid was ap-61 plied to modify the produced biochar. The adsorption-desorption properties of doxycycline on both 62 pyrophosphoric acid-modified and pristine biochar were examined. The main objectives of this study 63 were to (1) assess the adsorption-desorption capacity of doxycycline on pyrophosphoric ac-64 id-modified and pristine biochar (2) elucidate the physiochemical changes before and after modifi-65 cation. 66

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2. Materials and Methods

Materials and biochar production

Doxycycline hyclate (CAS no. 24390-14-5) was obtained from Asiatic chemical Karachi, Pakistan. 70 Sesame stalk was taken from an agricultural farm in Faisalabad, Pakistan. Sesame stalks were 71 air-dried and then used to produce the biochar by pyrolysis with a temperature at 500 °C for 4 h. 72

Modification of biochar

An aqueous solution of pyrophosphoric acid (H₄P₂O₇) was used for biochar modification for improving the sorption capability for the adsorption of doxycycline. Briefly, 15 g of biochar were 75 soaked in 30 mL of 20% H₄P₂O₇ solution for 24 h at room temperature. Afterwards, the pyrophos-76 phoric acid-modified biochar samples were washed with deionized (DI) water until the supernatant's 77 pH was stable. Thereafter, the supernatant was discarded and biochar samples were dried in an oven. 78

Characterization of unmodified and modified biochar

The elemental composition was used to measure polarity indexes of biochar before and after modi-80 fication. The H/C atomic ratios were used as a sign of aromaticity. Moreover, O/C ratios and polarity 81 indexes (O + N)/C ratios were calculated to assess polarities of manufactured biochar. Ash contents of 82 biochar were measured according to the ASTM (American Society for Testing and Materials) method 83 D1762-84. Additionally, Brunauer-Emmett-Teller (BET) was used to measure the surface area. 84

Sorption experiments

Kinetics sorption

15mg modified and unmodified biochar was added to 40mL glass tubes comprising 30mL doxycy-87 cline solution with 80mg L^{-1} concentration. All glass tubes were placed in shakers and agitated at 88 200rpm at room temperature from 10 minutes to 1440 minutes. Each sample was measured in trip-89 licate and also tubes comprising only 30mL doxycycline solution of the same concentration were 90 used for noticing doxycycline loss during this experiment. At pre-determined times, glass tubes were 91 extracted and centrifuged at 2800rpm for 25 minutes and then filtered via 0.45µm millipore mem-92 branes. Afterwards, the filtrate was measured via a spectrophotometer at 409 nm [9]. 93

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Sorption isotherms

Sorption isotherms for modified and unmodified biochars were measured at various initial concentration of doxycycline: 10, 20, 40, 60, 80, 100, 120, 150, 200, and 250mgL^{-1} . To each 40mL glass tube, 5mg of modified and unmodified sesame adsorbent and 30mL of doxycycline solution with different concentration were added, followed by shaking at 250rpm at room temperature to reach apparent equilibrium based on kinetic sorption.

pH effect on sorption isotherms

The doxycycline solutions with different concentrations were adjusted to pH 5, 7, 9 via using 0.1M 101 NaOH and HCl solution. Hereafter, isotherm sorption trials were performed at room temperature. 102

Data exploration

The adsorption capacity and removal efficiency (%) of doxycycline onto modified and un-modified 104 biochar were calculated using Eq. 1 and 2: 105

$$q_e = (C_i - C_e) \times V)/W \tag{1}$$

% removal doxycycline= $(C_i - C_e)/C_i \times 100$ (2)

Where $q_e (mg/g)$ signifies the adsorbed volume of doxycycline through the adsorbent, and C_i and C_e 108 are initial and equilibrium doxycycline concentrations (mg/L), respectively, while V represents the 109 total amount (L) of the solution and W stands for the weight (g) of the adsorbent. The kinetics adsorption data was analyzed through pseudo-1st-order and pseudo-2nd-order reactions: 111

Pseudo-first-order: 112

$$q_t = q_e(1 - \exp(-K_1 t))$$
 (3) 113

$$q_t = q e_2 k_2 t / 1 + q^* exp(-K_2 t)$$
(4) 115

To examine the reaction behaviour between doxycycline and biochars, data of adsorption isotherms ¹¹⁶ was fitted through two models applied [10]. ¹¹⁷

Freundlich:

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qe – Kičeli	(\mathbf{J})	119
Langmuir;		120
$q_e = q_{max}KLCe / 1 + KLCe$	(6)	121

(5)

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Desorption Study and Reusability of biochar

The modified and unmodified biochar was regenerated via extraction with 25mL of Milli-Q water. 123 After being washed and dried in an oven, the materials performed three adsorption-desorption cycles, 124 separately. This was followed through continuously for 24h at room temperature. The desorption 125 efficiency of modified and unmodified biochar was calculated using equation (7): 126

% desorption_doxycycline =	$C_{dec}/C_{adc} \times 100$	(7)	
, accorption, acceptine			

Where, C_{des} and C_{ads} are desorbed volume (mg/L) of doxycycline in the solution and adsorbed volume128of doxycycline through biochar, respectively.129

Results and discussion

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Biochars characterization

Table 1 exhibits the ash content, elemental compositions, aromatic ratio, surface area, and pore 132 volume of modified and unmodified biochar. The surface area significantly increased after modifi-133 cation (129.41 m² g⁻¹). It was much higher than unmodified biochar (13.64 m² g⁻¹). Pyrophosphoric 134 acid treatment could increase the biochar's surface area [11,24]. Pyrophosphoric acid treatment 135 generally increased the content of S, N and C but decreased O content as compared to pristine biochar. 136 The H content showed no obvious alteration between before and after modification. Moreover, the 137 ash content reduced after modification from 61.34 to 53.29%. Pyrophosphoric acid modification 138 could partially eliminate ash of pristine biochar [12,25]. A substantially negative association was 139 observed between ash and C contents in this work. Additionally, it was observed that the aromatic 140 ratios O/C, H/C, (O + N)/C of modified biochar were reduced compared to pristine biochar. This 141 decrease showed the biochar treated by pyrophosphoric acid turned into material with a less hydro-142 philic-nature with weak polar groups [12, 13,26]. 143

Table 1. Elemental composition (%), and surface area of unmodified and modified biochar
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Sample	С	Н	0	Ν	S	O/C	N/C	H/C	(O+N)/C	Ash %	SSA
											(m ² g ⁻¹)
Unmodified	32.16	0.64	11.61	1.04	0.11	0.19	0.04	0.01	0.24	61.34	13.64
Biochar											
Modified	34.15	0.66	6.32	1.45	0.14	0.33	0.03	0.02	0.22	53.29	129.41
Biochar										$\langle \rangle$	

Reaction Time and Kinetic Modelling

Figure, 1 and Table 2 represent the doxycycline removal percentage and sorption capacity by modified and 147 unmodified biochar at different times. After 120 minutes, there was an ambiguous difference between 180 and 148 1440 minutes, thus, 120 minutes was deemed to be a maximum equilibrium time for doxycycline sorbed onto 149 modified and unmodified biochar. Besides, it was observed that the sorption efficiency of treated-biochar was 150 higher than unmodified biochar. In this study, pseudo-1st-order and pseudo-2nd-order models were applied to 151 arrange the sorption data for understanding possible mechanism related with the doxycycline sorption on 152 modified and unmodified biochar. R² values of the pseudo-second-order model (0.902-0.913) were greater 153 than those of the pseudo-first-order model (0.804-0.842) (Table 2), comparable the most of antibiotics ad-154 sorption findings [14, 15, 16]. Furthermore, values of qe were closer to theoretical qe values considered 155 through the pseudo-second-order model; additionally proposing that kinetics doxycycline's sorption onto 156 modified and unmodified biochar could be preferably reported by the pseudo-2nd-order model. The pseu-157 do-2nd-order model exhibits chemisorption mechanism occurred between doxycycline and modified biochar 158 including valency forces through electrons exchange between biochar and doxycycline [17,27]. On the con-159 trary, the qe values of pyrophosphoric acid-treated biochar were greater than for unmodified-biochar. For 160 instance, theoretical q_e of doxycycline on modified biochar was 174.4mgg⁻¹, significantly greater than the 161 pristine biochar. It is evident that pore volume and SA of modified biochar significantly enhanced after py-162 rophosphoric acid modification (Table, 1), signifying pore-filling and surface adsorption might explicate the 163 greater qe of modified biochar than un-modified biochar (Figure 1 and Table 2). Moreover, pyrophosphoric 164 acid treatment could enhance functional groups number amount of adsorbent [12, 18, 28]., thus, chemical 165 interactions between functional groups of modified biochar and doxycycline molecules might also partly lead 166 to greater qe values of modified biochar compared to pristine biochar [18]. Therefore, possible cause for 167 greater qe values of modified biochar than pristine biochar might be explicated via the increased interactions 168

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between modified biochar and doxycycline owing to the improved number of functional groups, i.eOH and	169
-COOH [12].	170

Table. 2. Kinetic adsorption parameters of doxycycline onto unmodified and modified biochar

Sample	Pseudo-1 st -order			Pseudo-2 nd -order		
	qe (mg g ⁻¹)	$K_1 (h^{-1})$	R ²	qe (mg g ⁻¹)	K ₂ (h ⁻¹)	R ²
Unmodified	131.4	0.04	0.804	143.1	0.0003	0.902
BC				(
Modified BC	154.8	0.08	0.842	174.4	0.0006	0.913



Figure 1. Sorption kinetics of doxycycline on modified and unmodified biochar

Influence of Initial doxycycline -Concentration and Isothermic sorption

Since the pyrophosphoric acid-modified biochar exhibited greater sorption capacity compared to unmodified 182 biochar (Table, 2). Sorption data of doxycycline onto modified and unmodified biochar are represented in 183 Figure, 2. The sorption data exhibited significantly greater sorption capacity compared to unmodified biochar. 184 The data of doxycycline sorbed to modified and unmodified biochar fitted thru Freundlich and Langmuir 185 models (Table 3). Both the Freundlich and Langmuir models presented isothermic sorption well because R² 186 value of Langmuir (0.961–0.983) was same as R² of Freundlich model (0.982–0.989). Conforming to, q_{max} 187 fitted via Langmuir model, the q_{max} of doxycycline via modified biochar (153.9mgg⁻¹) was significantly 188 higher than untreated biochar (87.6mgg⁻¹). According to Freundlich, Kf of modified biochar (89.5mg⁻¹-nLn 189 g⁻¹) were also higher compared to untreated biochar (63.2 mg⁻¹-n Ln g⁻¹). These findings suggested that 190 modified biochar had hefty sorption capability and greater sorption ability as compared to untreated biochar, 191 which might be mainly explicated by the higher O/C ratio and surface area of pyrophosphoric acid-modified 192 biochar than pristine biochar. In this work, Langmuir and Freundlich both models could fit for adsorption data 193 of doxycycline on modified and unmodified biochar well, signifying the doxycycline adsorption on biochar 194 might be influenced via several processes. [13] Summarized primary sorption process of antibiotic on ad-195 sorbent surface, such as EDA interaction, surface complexation, cation exchange, electrostatic interaction, and 196 Hydrogen bonding. Pyrophosphoric acid-treatment removed the cations, for example, Ca²⁺, Na⁺ and Mg²⁺, 197 signifying the surface complexation and cation exchange should not be primary mechanisms for doxycycline 198 adsorption on modified and unmodified biochars [18]. Simultaneously, Table 2 and Figure 1 exhibited that 199 doxycycline was sorbed mainly via chemisorption mechanism including EDA-interaction and H bonding. In 200 comparison, O/C contents of modified biochar with unmodified biochar (Table 1), greater O/C ratio of mod-201 ified biochar proposed that modified biochar might have additional oxygen-enrich functional groups which 202 could aid as hydrogen-bond acceptors and therefore doxycycline adsorption on modified biochar was greater 203 raw adsorbent [19]. Contrastingly, physical adsorption might also partly bestow to greater adsorption ability 204 of modified biochar due to SA of modified biochar was greater than pristine adsorbent (Table 1), though 205 chemical adsorption was most influenced in this work. Nonetheless, it is difficult to value the support of π - π 206 EDA-interaction, H-bonding and physical adsorption for doxycycline sorption on modified and unmodified 207 biochars ([12]. 208

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Sample	Langmuir			Freundlich				
	qmax (mg g ⁻¹)	$K_L(L mg^{-1})$	R ²	Kf (mg ⁻¹⁻ⁿ	n	R ²		
				g ⁻¹)				
Unmodified	87.6	0.12	0.961	63.2	0.11	0.982		
BC								
Modified BC	153.9	0.26	0.983	89.5	0.14	0.989		
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		120			-			

Table. 3. Isothermic adsorption parameters of doxycycline onto unmodified and modified biochar



80 100 120 150 200 250

Effect of pH

Removal % 60

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0

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Effect of different pH values on adsorption of doxycycline on pyrophosphoric acid treated biochar was 217 examined because doxycycline form and adsorption mechanism of doxycycline on biochar was signif-218 icantly affected via pH. Figure 3 displays the impact of pH in sorption of doxycycline on modified and 219 unmodified biochar at 25 °C. Increasement in adsorption capacity with increased pH was noticed on 220 modified and unmodified biochar, and highest adsorption of doxycycline onto modified and unmodified 221 biochar was observed at 9 pH. The adsorption data of doxycycline on modified and unmodified biochar 222

Different conc. mg L⁻¹

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Unmodifed BC

Modified BC



under several pH solutions were arranged by Freundlich and Langmuir models (Table 4). The R² values 223 showed the Freundlich and Langmuir could fit the adsorption data well, particularly Freundlich model. 224 Generally, Kf of doxycycline adsorption on biochars reduced but n enhanced with enhancing pH. Be-225 sides, the initial pH enhancing from 5-9, the fitted Kf of unmodified biochar reduced from 65.2 to 32.4 226 mg⁻¹-n Ln g⁻¹ and modified biochar decreased from 79.1 to 26.4 mg⁻¹-n Ln g⁻¹. As well as, q_{max} of 227 doxycycline on modified biochar enhanced from 139.8 mg g^{-1} to 562.1 mg g^{-1} with pH elevating from 228 5-9 which was much higher than unmodified biochar (119.2 to 358.8 mgg^{-1}). These findings evidently 229 proposed that adsorption ability of doxycycline on modified biochar was improved thru raised pH 5 to 9. 230 In general, the doxycycline properties and biochar were changed through solution pH, and subsequently 231 inducing sorption volume of doxycycline on the biochar. pHPZC of biochar and pKa of doxycycline was 232 suggested as key reason for alterations of adsorption capability of doxycycline onto biochar under var-233 ious pH values [20, 28, 29, 30]. Moreover, biochar surface charge was positive, when solution pH was 234 lower compared to their pHPZC and vice versa. At lower pH, doxycycline was cationic and the biochar 235 surface is generally positive, thus, the π - π electron (electron-donor- acceptor) interactions were weak. As 236 well as pH elevating, doxycycline turned zwitterionic and electron-donor-acceptor interactions between 237 biochar and doxycycline was strengthened. Therefore, increased adsorption of doxycycline was noticed. 238 In brief, changed charges of biochar and doxycycline should reasonably elucidate the variations of ad-239 sorption volume of doxycycline on modified and unmodified biochar under different pH values. Thus, 240 the increased sorption of doxycycline on modified biochar at increased pH could be mainly enlightened 241 by the strengthened electrostatic attraction between negative charges of doxycycline and positive surface 242 charges of biochar. 243

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Table. 4. Sorption isotherm parameters of doxycycline onto unmodified and modified biochar at differentinitial pH of solution.

Sample	Langmuir				Freundlich			
	qmax (mg g ⁻¹)		$K_L(L mg^{-1})$	R ²	Kf (mg ⁻¹⁻ⁿ	n	R ²	
					g ⁻¹)			
Initial pH=5						Δ		
Unmodified B	С	119.2	0.22	0.929	65.2	0.1	0.949	
Modified BC 139.8		139.8	0.57	0.941	79.1	0.1	0.973	
Initial pH=7								
Unmodified B	С	166.8	0.21	0.901	68.6	0.2	0.964	
Modified BC 2		258.7	0.07	0.984	50.4	0.3	0.996	
Initial pH=9	Initial pH=9							
Unmodified B	С	358.8	0.03	0.967	32.4	0.4	0.982	
Modified BC		562.1	0.02	0.993	26.4	0.5	0.998	



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Figure 3. Effect of various pH solutions on doxycycline removal

Desorption analysis

After successful doxycycline adsorption, the modified and unmodified biochars were exposed to regeneration 261 experiments in order to measure the stability and reusability of biochar. The experiments conducted based on 262 three adsorption desorption cycles to scrutinize repetitive use of biochar under the optimum condition. 263 Findings exhibited that sorption capabilities of modified biochar somewhat decreased after three adsorp-264 tion-desorption cycles (Figure 4). Very low desorption amount was noticed with Milli-Q water for modified 265 biochar (12.36-1.25 %) (Figure 4). Results proposed that pyrophosphoric acid-modified biochar was effi-266 ciently recycled and can be used in repeated doxycycline sorption batches at least three times with minimum 267 loss in their sorption volumes, using Milli-Q water as ideal desorption material [21, 32, 33, 34, 35]. 268





5. Conclusions

Pyrophosphoric acid-modification improved the doxycycline sorption on modified biochar, (about 30% en-272 hancement). The doxycycline sorption on pyrophosphoric acid-modified sesame adsorbent was greatly en-273 hanced along with increasing pH from 5 to 9. The π - π EDA interactions and H-bonding might be the primary 274 adsorption mechanism for doxycycline adsorption onto treated and un-treated biochar. With respect to the 275 pristine adsorbent, modified biochar revealed higher sorption capacities and volumes of doxycycline and 276 might be a virtuous adsorption agent for doxycycline elimination from the water, though the doxycycline 277 adsorption on unmodified biochar was low compared to modified biochar. The findings of the current study 278 demonstrated that sorption was spontaneous and multi-layered as evidenced by Langmuir and Freundlich 279 models, and endothermic as elimination efficacy can be enhanced by raising the temperature. 280

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