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Heterogeneity of biochars properties dependent on feedstock sources and production temperatures

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1	ABSTRACT: This study was conducted with a wide range of production temperatures
2	$(200^{\circ}\text{C}-650^{\circ}\text{C})$ and a series of feedstock sources (n=12) to quantify the influence of these
3	two factors on any given biochar property. The quantitative evaluation was completed using
4	two indices, feedstock-depended heterogeneity (H_F) and temperature-depended
5	heterogeneity (H _T), obtained from the statistic analysis of coefficient of variation. The
6	values of H_F or H_T were positively related to the heterogeneity and correspondingly to the
7	influence extent. Total organic carbon, fixed carbon, and mineral elements of biochars
8	varied greatly among different feedstocks but were less affected by temperature. Biochar
9	surface area and pH was less influenced by feedstock than by temperature, while pore
10	volume and CEC was more affected by feedstocks than temperature. Biochar recalcitrance
11	was mainly determined by production temperature, while potential total C sequestration
12	depended mainly on feedstocks. CP-MAS ¹³ C NMR and FTIR showed that alkyl-C,
13	aliphatic-C and aromatic-C was highly related to the production temperature. Raman
14	spectroscopy revealed that distribution and state of sp ² -bonded carbon remained stable with
15	feedstock and temperature. The results indicated that the two indices could be suitable for
16	assessing the effect extent of feedstock source or production temperature on biochar
17	properties.

Keywords: Biochar; Feedstock-depended heterogeneity; Temperature-depended

20 heterogeneity; Physiochemical properties; Chemical structure

21 **1. Introduction**

Biochar, pyrogenic organic material derived from incomplete combustion of biomass, has 22 recently received much attention due to its great potential in a wide range of environmental 23 applications. In addition to its ability to serve as a carbon sink for mitigation of global climate 24 change [1-3], biochar may be used as an effective contaminant sorbent [4-6] or soil nutrient 25 amendment [7, 8]. However, the utility of each specific biochar depends upon its inherent properties. 26 27 For example, biochars with high specific surface area may be used as sorbents, whereas the ones with high recalcitrance may function in carbon fixation [9]. Those rich in nutrients and minerals 28 29 would be better used as soil amendments to improve fertility [10].

It has been shown that biochar characteristics are influenced by production variables such as 30 feedstock source, heat temperature, heat duration, pyrolysis atmosphere, etc. Among these, feedstock 31 32 source and heat temperature are considered to be main controls on biochar characteristics [11, 12]. For example, increases of pH, CEC, and trace metals concentration occur with increasing production 33 temperature [13-15]. Biochars derived from wood biomasses often have higher surface area than 34 grass biochar [15, 16]. However, most previous studies focused on a few feedstock materials or those 35 falling into one or two categories such as crop biomasses, wood derivatives, or manures, or those 36 made at only a few production temperatures. For example, Cantrell et al. [17] studied the impact of 37 pyrolysis temperature and manure source on physicochemical characteristics of five manures biochar 38 39 made at only two temperatures. Pereira et al. [18] investigated the labile fraction of C in biochar 40 derived from three trees (pine, poplar and willow) at two temperatures. In general, biochars of the feedstock with the same category show similar properties compared to those made from parent 41 material of very different types. 42

If we are to make use of biochar to the fullest extent of its possible applications, we must develop an understanding of its physiochemical variations for a broader range of biochar types than has previously been examined. Optimizing biochar for a specific application may require selection of a

feedstock as well as pyrolysis production technique and conditions to produce biochars with specific 46 characteristics [19]. Thus, the objectives of this study are (i) to determine how the two main factors, 47 feedstock source and production temperature, affect the biochar properties and (ii) to evaluate which 48 one of the two factors dominates one property of biochar based on a series of temperatures from 49 200°C to 650°C and a variety of source materials (12 waste biomasses) representing 6 categories: 50 animal manure, wood waste, crop wastes, food wastes, aquatic plants, and municipal waste. Two 51 52 evaluation indices, feedstock-depended heterogeneity (H_F) and temperature-depended heterogeneity (H_T) are introduced to quantify the influence of feedstock source or production temperatures, 53 54 respectively, on any given biochar property and tell which one is dominance. In this way, production materials and conditions can be chosen to produce biochars optimized for any given application. The 55 comparison of H_F and H_T also gave a new insight to the origin and evolution of the variation 56 properties observed among biochars. 57

58

59 2. Materials and Methods

60 2.1. Biomass Collection and Biochar Production

Twelve common waste biomasses were collected from a farm in Shanghai, China and divided 61 into 6 categories including animal manure, wood waste, crop residue, food waste, aquatic plants, and 62 municipal waste. The biomasses were air-dried and then ground to less than 2 mm for biochar 63 production. Details on the production of biochar were described previously [20]. Briefly, to evaluate 64 the feedstock source effect, all 12 ground waste biomasses were heated at 500°C under O₂-limited 65 atmosphere for 4 h. To examine pyrolysis temperature effect, a wastes-based biochar (pig manure) 66 and plant-based biochar (wheat straw) were chosen and pyrolyzed at 200°C, 350°C, 500°C and 67 650°C. 68

69 2.2. Biochar characterization

70 Total C analysis of biochar was conducted on an element analyzer (Vario EL III, Elementar,

German). Ash content, volatile matters (VM), and fixed carbon (FC) were determined according to
standard ASTM methods [21-23]. The metal concentrations in biochars were measured in the
digestion solution using the inductively coupled plasma (ICP-AES, ICAP6000 Radial, Thermo,
English), following biochar digestion using the USEPA method 3050B [24]. The cation exchange
capacity (CEC) was determined according to a modified barium chloride compulsive exchange
method [25]. All analyses were conducted in duplicate.

The solid phase of biochar was characterized by thermogravimetry (TG) analysis (PerkinElmer

Pyris 1 TGA) with heating from 25°C to 900°C at a rate of 20°C per min. Surface functional group

distributions were determined by infrared (IR) spectroscopy (IR Prestige 21 FTIR, Shimadzu, Japan)

80 and nuclear magnetic resonance (CP-MAS ¹³C-NMR) spectra (AVANCE III 400, Bruker,

81 Switzerland), which were obtained at a frequency of 100.6 MHz using a Varian Unity Inova 400

82 NMR spectrometer. Specific surface area and pore size distribution of biochars were determined

using a BET-N₂ SA analyzer (JW-BK222, Jwgb, China). Raman spectroscopy analysis was

conducted using a visible Raman system (Bruker Senterra R200-L, American) with a 15 mW 532 nm He-Ne laser with excitation line set to $\lambda_0 = 532$ nm.

86 2.3. Calculations

Fixed Carbon (FC) of biochar was calculated as the sum of moisture, ash, and volatile matter
subtracted from 100 [23].

89
$$FC(\%) = 100 - Moisture(\%) - Ash(\%) - VM(\%)$$
 (1)

An index R₅₀ was used to evaluate the thermal recalcitrance of biochar and was obtained by TG
analysis, as recently proposed by Harvey et al [26]:

92
$$\mathbf{R}_{50,\text{biochar}} = \frac{\mathbf{T}_{50,\text{biochar}}}{\mathbf{T}_{50,\text{graphite}}}$$
 (2)

where $T_{50, \text{ biochar}}$ and $T_{50, \text{ graphite}}$ are the temperature values corresponding to 50% weight loss via oxidation/volatilization of biochar and graphite, respectively. Values are obtained directly from TG 95 thermograms that have been corrected for water and ash content.

Carbon sequestration potential (CS) was defined as the final carbon reserved in soil, which was calculated by subtracting the carbon lost during pyrolysis from the initial C in raw biomass, and multiplying by the recalcitrance (R_{50}) of biochar products. M was the weight of the feedstock.

99
$$\mathbf{CS(\%)} = \frac{\mathbf{M(g)} \cdot \mathbf{Biochar yield} \ (\%) \cdot \mathbf{C\% in Biochar} \cdot \mathbf{R_{50}}}{\mathbf{M(g)} \cdot \mathbf{C\% in feedstock}}$$
(3)

100 The feedstock-depended heterogeneity (H_F) and temperature-depended heterogeneity (H_T) of 101 biochars were calculated using the coefficient of variation (CV) in statistical method, and the larger 102 the H_F or H_T is, the more influenced by feedstock or production temperature the biochar property is:

103
$$H_F \text{ or } H_T = \frac{\text{Standard deviation}}{\text{mean value}}$$
 (4)

104

105 **3. Results and Discussion**

106 *3.1. Bulk physicochemical properties*

107 Concentrations of total carbon (TC) and fixed carbon (FC) in all 12 biochars ranged 24.2-75.8% and 3.84–72.9% with the H_F of 0.37 and 0.76 (Table 1). Increasing the temperature from 200°C to 108 650°C increased TC and FC (Table 1). In the temperature range, concentrations of TC and FC of the 109 110 pig manure biochar were 37.0-45.3% and 12.3-42.3%, with the H_T of 0.09 and 0.48, respectively, while wheat straw biochar contained 38.7-68.9% TC and 22.5-72.1% FC, with the H_T of 0.23 and 111 0.41, respectively (Table 1). All H_T were lower than the H_F, indicating that TC and FC of biochars 112 were more influenced by feedstock source than by production temperature. 113 Both volatile matter (VM) and weight yield were more sensitive to temperature, indicated by 114 their higher H_T (0.5–0.81) than H_F (0.27–0.36). Kloss et al. [16] reported a similar result that labile, 115 aliphatic compounds undergo a great loss during pyrolysis. Ash content was more sensitive to 116 feedstock due to its higher H_F (0.53) than H_T (0.33–0.34). As shown in Table 1, ash is higher in 117 manure and sludge biochar (18.1–42.9%), while crop residue biochar contained low ash 118

119	(2.10–7.49%). The higher ash in the manure biochar was due to richness of mineral constituents [20].
120	Biochars from different feedstocks had wide range minerals, while the mineral concentrations
121	changed little with production temperature from 200°C to 650° C (Table 2). The H _F for each mineral
122	element was higher (0.87–2.00) than H_T (0.40–0.51), indicating that mineral elements of biochars
123	were more influenced by feedstock source than temperature. Generally, manure contained more
124	nutrient P than crop residue and grass biochar, whereas nutrient K was higher in crop residue and
125	grass biochar than that in manure biochar (Table 2). Thus, the utility of biochars as a soil amendment
126	to improve soil fertility should be classified carefully according to different feedstock sources rather
127	than production temperature.
128	Biochar pH varied less among the different feedstocks (8.8–10.8) than among the production
129	temperature (5.43–10.8) (Table 1). Therefore, biochar was influenced more by temperature (H_T =0.19)
130	than by feedstock (H_F =0.05). By contrast, the CEC varied greatly among biochars of different
131	feedstocks ($H_F=0.9$) but relatively little with temperature ($H_F=0.52-0.65$). This may be explained that
132	CEC is related to cations (e.g., K, Ca, Mg) which vary greatly with feedstocks (Table 2).
133	The physical structure of biochars, such as surface area (SA), pore volume (PV), and average
134	pore size (APS) are typically related to its sorption and water holding capacity which, in turn, relates
135	to its effect on soil structure, contaminant mobility, and microbial interactions. The heterogeneities of
136	SA and APS were more related to production temperature (H_T =0.72–1.55) than feedstock
137	(H _F =0.58–1.09), while PV was more influenced by feedstock (H _F =1.11) than temperature
138	$(H_T=0.49-0.81)$ (Table 1). The influence of feedstock on PV was perhaps related to the relative
139	proportion of hemicelluloses, cellulose, and lignin fractions in biomasses. A dramatic rise in SA was
140	observed when the temperature was increased above 350°C, at which point, cellulose is known to
141	decompose and a phase transition from layered C to amorphous char occurs [27].
142	3.2. Recalcitrance and stability

143 The ability of biochars to resist abiotic and biotic degradation (herein referred to as recalcitrance)

is crucial to their success as a soil carbon sequestration. Harvey et al (2012) have developed an index 144 (R_{50}) to evaluate the recalcitrance of biochars, which uses the energy required for thermal oxidation 145 of the biochars (normalized to that for oxidation of graphite) as a measure of recalcitrance [26]. 146 The water and ash content-corrected thermogravimetry patterns of biochars are presented in Fig. 147 1. The temperatures at which 50% biochar weight loss occurred ranged within 614–727°C for all 148 feedstocks and within 338–767°C for all production temperature of pig manure biochar and wheat 149 150 straw biochar. The calculated R₅₀ for biochars from all feedstocks fell in a narrow range of 0.69–0.82, with H_F being 0.06, while R_{50} for biochars produced at 200°C–650°C was within a wide range of 151 152 0.38–0.87, with H_T being 0.29–0.34 (Table 1), indicating that the recalcitrance of biochar was mainly determined by production temperature, which was also expected from previous findings [9]. Biochar 153 recalcitrance is related to aromatic C which increased with increasing temperature, regardless of 154 155 nature of feedstocks (shown below). Fig. 1 also shows that all biochars produced at same temperature had similar R₅₀ and the gap among different feedstocks enlarged with the increase of production 156 temperature, further suggesting that temperature was the dominating control on recalcitrance. 157 Carbon sequestration potential (CS) was evaluated assisted by R_{50} as shown in equation 3. CS of 158 all 12 biochars ranged 23.7–54.0% with H_F being 0.27, while those for pig manure biochar and 159 wheat straw biochar at production temperature of 200°C–650°C were 33.1–38.4% and 34.3–44.6%, 160 respectively, with H_T being 0.07 and 0.11, respectively. The H_T was lower than H_F , indicating that 161 temperature had less influence on the carbon sequestration capacity. It is probably that low 162 163 production temperature could retain much C in biochar, but a considerable amount of these C would be abiotically or microbially mineralized [9, 28]; when the temperature increases, more C would lose 164 in pyrolysis, but more recalcitrant C would be produced [29]. The contradictory effects would keep 165 166 biochar-C less changed. Therefore, the C sequestration was mainly determined by the inherent molecular configuration of biomasses [30]. 167

168 *3.3. Biochar chemical structure*

The carbon cluster size and functional group distributions were identified by CP-MAS ¹³C NMR 169 and FTIR, and are shown in Fig. 2 and 3, respectively. The ¹³C NMR spectrograms of biochars from 170 12 feedstocks were very similar, whereas they varied greatly among those produced from a single 171 feedstock type across a range of temperatures (Fig. 2). Table 3 summarizes the relative proportion of 172 C in each chemical functional groups for the biochars examined, which were integrated in the 173 chemical shift (ppm) resonance intervals of 0-46, 46-65, 65-90, 90-108, 108-145, 145-160, 174 160–185, 185–225 ppm [31]. Clearly, aromatic C with chemical shift of 108–145 ppm was the main 175 C-containing functional group in all biochars (45.0-80.3%), with an H_F of being 0.15. The aromatic 176 C in biochars increased from 2.24% at 200°C to 62.9% at 650°C with H_T being 0.68 (Table 3). 177 Therefore, the aromatic C was mainly controlled by the production temperature, agreeing with the 178 179 recalcitrance shown above.

180 The control of aromatic content by production temperature, as opposed to feedstock type, was true for other C-containing functional groups. For example, the subdominant abundance of C was 181 alkyl C (mainly CH₂ and CH₃ sp³ carbons) at the chemical shift of 0-46 ppm accounted for 182 10.9–18.6% of the C-containing functional groups in biochars of different feedstocks at production 183 temperature 500°C ($H_F = 0.15$) and for 3.17–38.8% in biochars of different temperatures ($H_T = 0.90$). 184 The 200°C biochars retained properties like the raw materials. For example, the C within 46–65 ppm 185 and 65–90 ppm, representing methoxy and N alkyl C from OCH₃, C–N and complex aliphatic 186 carbons, respectively, as well as O-alkyl C was in high proportions. 187

The FTIR spectra also indicate a range of superficial functional groups among different biochars
(Fig. 3). The absorption peaks at 2916 cm⁻¹ are assigned to saturated C-H stretching vibration
(aliphatic C-H), and a wide absorption peak at 3200–3500 cm⁻¹ is attributed to –OH stretching [20].
These peaks existed in all biomasses, while disappeared above 350°C, which were influenced more
by temperature indicating the dehydration of cellulosic and ligneous components (Fig. 3 c and d).
The peaks at 1465–1340 cm⁻¹ are saturated C-H bending vibration and it is of great difference among

194 biochars of feedstocks, while less difference among biochars produced at different temperatures. The -COO anti-symmetric stretching of amino acids (1574 and 1600 cm⁻¹) appeared in wood and crop 195 waste biochars, which presented little change until the temperature rose to 650°C. The intensity of 196 C=O stretching of aromatic rings (1593 cm^{-1}) decreased with temperature rise and seemed similar in 197 all feedstocks. Peaks at 874 and 1034 cm⁻¹ were assigned to the bands of the out-of-plane bending for 198 $CO_3^{2^2}$, which exists more in biochars of wastes and manures and less in plant-based biochars, and 199 was less influenced by production temperature [32, 33]. The NMR and FTIR results all showed the 200 aromatization among different feedstocks and production temperature [34]. The recalcitrance and C 201 202 sequestration have close relationship with carbon configuration, which perhaps determines the breakdown of C-bond and re-aggregation of C cluster under heat treatment [35]. 203

Raman spectroscopy has been widely used to evaluate the microstructure of carbon materials, 204 particularly the distribution and state of sp²-bonded (aromatic) carbon [36], which is embedded in a 205 disordered and amorphous matrix of both sp³ and sp² carbon. The G-band centered at 1580 cm⁻¹ 206 arises from the in-plane vibrations of the sp²-bonded crystallite carbon and has been observed for 207 single crystal graphite, while another peak denoted as the "disorder" peak (or D-band) centered at 208 1357 cm⁻¹ is typically observed in polycrystalline graphite. The D-band is attributed to in-plane 209 vibrations of sp²-bonded carbon within structural defects. For disordered carbon materials the ratio of 210 the integrated intensities I_D/I_G is often reported to be inversely proportional to the lateral extension 211 La of the graphene materials [37]. 212

As shown in Fig. 4a and b, both G-band and D-band appeared in all 12 biochars with production temperature of 500°C and had the similar I_D/I_G (0.804–1.51), with low H_F (0.31) (Table 1), implying that ratio of disordered or strongly distorted structure of turbostratic carbon to ordered graphite crystals was less determined by feedstocks than production temperature. For biochars produced at a range of production temperature, bands were found to develop at 350°C, indicating the beginning of aromatization. The increase of I_D/I_G with temperature increasing from 350°C to 650°C was also not obvious ($H_T < 0.36$) since the temperature used in this study was in a relatively low range and their influence on biochar microstructure could be negligible.

4. Conclusions

Biochars of different physical and chemical properties will be more suited for one application or 222 another, e.g. soil amelioration, C sequestration, contaminant remediation, etc. The biochar properties 223 have been shown to be mainly controlled by feedstock source and production temperature. The 224 relationship between H_F and H_T for a range of properties in the biochar examined is depicted in Fig. 225 5. Biochar yield, pH, recalcitrance, and volatile matter plotted above the 1:1 line, indicating that 226 227 these properties are controlled more strongly by production temperature. Thus, any application of biochar related to these properties would call for greater attention to the production temperature. For 228 example, if a biochar is produced for carbon sequestration purpose, high temperature is required 229 230 since it increases recalcitrance. If a biochar is intended for use as adsorption sorbent, increasing temperature (>500°C) would improve the surface area and micropore volume. However, feedstock 231 should be also considered, since H_F was also high (Fig. 5). Biochar C, CEC, fixed C, carbon 232 sequestration, mineral concentrations, and ash content plotted below the 1:1 line (Fig. 5), indicating 233 that these properties are controlled more strongly by feedstock sources. Therefore, any application of 234 biochar related to these properties should focus on raw materials selection. If a biochar is prepared as 235 soil amendment, biomass rich in minerals would be advisable. For example, pig manures and aquatic 236 plant biochars contain abundant P, K, Ca, Mg, etc (Table 2). 237

Overall, the results obtained from this study indicate that feedstock source or production
temperature affect biochar properties to different degrees and consideration of production conditions
guide the development of 'optimum' biochars for different environmental applications

241

242

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Table 1

Compositions, physico-chemical properties, and structural characteristics of biochars derived from 12 waste biomasses produced at 500°C and biochars produced

Biochar	Temper	OCb	FC ^c	CS ^d	Yield	VM ^e	Ash	pН	CEC ^f	SA ^g	PV ^h	APS ⁱ	$\mathbf{R}_{50}^{\mathbf{j}}$	I _D /I _G ^k
feedstock	ature ^a	(%)	(%)	(%)	(%)	(%)	(%)		(cmol·kg ⁻¹)	$(m^2 \cdot g^{-1})$	$(cm^{3} \cdot g^{-1})$	(nm)		
Cow manure	500°C	43.7	14.7	52.5	57.2	17.2	67.5	10.2	149	21.9	0.028	5.04	0.72	1.09
Pig manure		42.7	40.2	33.1	38.5	11.0	48.4	10.5	82.8	47.4	0.075	6.35	0.74	1.19
Shrimp hull		52.1	18.9	32.4	33.4	26.6	53.8	10.3	389	13.3	0.039	11.6	0.78	1.51
Bone dregs		24.2	10.5	28.3	48.7	11.0	77.6	9.57	87.9	113	0.278	9.86	0.82	1.15
Wastewater sludge		26.6	20.6	23.7	45.9	15.8	61.9	8.82	168	71.6	0.060	3.37	0.76	-
Waste paper		56.0	16.4	28.3	36.6	30.0	53.5	9.88	516	133	0.084	2.51	0.80	1.29
Sawdust		75.8	72.0	31.1	28.3	17.5	9.94	10.5	41.7	203	0.125	2.23	0.73	1.33
Grass		62.1	59.2	27.7	27.8	18.9	20.8	10.2	84.0	3.33	0.010	11.9	0.70	1.20
Wheat straw		62.9	63.7	34.4	29.8	17.6	18.0	10.2	95.5	33.2	0.051	6.10	0.71	1.10
Peanut shell		73.7	72.9	39.1	32.0	16.0	10.6	10.5	44.5	43.5	0.040	3.72	0.69	1.15
Chlorella		39.3	17.4	35.3	40.2	29.3	52.6	10.8	562	2.78	0.010	15.0	0.77	1.16
Waterweeds		25.6	3.84	54.0	58.4	32.4	63.5	10.3	509	3.78	0.009	9.52	0.78	0.80
	$\mathrm{H_{F}}^{\mathrm{l}}$	0.37	0.76	0.27	0.27	0.36	0.53	0.05	0.90	1.09	1.11	0.58	0.06	0.31
Pig manure	200°C	37.0	12.6	38.4	98.0	50.7	35.7	8.22	23.6	3.59	-	-	0.39	-
	350°C	39.1	34.7	33.6	57.5	27.4	37.2	9.65	49.0	4.26	0.024	12.8	0.55	0.56
	500°C	42.7	40.2	33.1	38.5	11.0	48.4	10.5	82.8	47.4	0.075	6.35	0.74	1.19
	650°C	45.3	19.2	34.4	35.8	10.7	69.6	10.8	132	42.4	0.062	5.80	0.78	0.90
	${\rm H_{T}}^{\rm m}$	0.09	0.48	0.07	0.50	0.76	0.33	0.12	0.65	0.97	0.49	0.47	0.29	0.36
Wheat straw	200°C	38.7	22.5	37.7	99.3	70.2	7.21	5.43	32.1	2.53	_	_	0.38	_
	350°C	59.8	53.2	44.6	52.5	31.3	14.7	8.69	87.2	3.48	0.010	11.3	0.55	1.24
	500°C	62.9	63.7	34.3	29.8	17.6	18.0	10.2	95.5	33.2	0.051	6.10	0.71	1.16
	650°C	68.9	72.1	41.5	26.8	11.1	16.2	10.2	146	182	0.093	2.05	0.87	1.32
	H_{T}	0.23	0.41	0.11	0.64	0.81	0.34	0.26	0.52	1.55	0.808	0.72	0.34	0.06

from pig manure and wheat straw at 200°C–650°C.

^a Biochar production temperature

^b Organic carbon

^c FC is fixed carbon (%, dry basis)

- $^{\rm d}$ CS is potential carbon sequestration (%) after pyrolysis and mineralization
- e VM is volatile matter (%, dry basis)
- f CEC is cation exchange capacity (cmol·kg⁻¹) g SA is BET-N₂ surface area ($m^2 \cdot g^{-1}$) h PV is pore volume (cm³ · g⁻¹)

- i APS is average pore diameter (nm)
- ^j R₅₀ is a novel index for evaluating biochar recalcitrance derived from thermogravimetric data (Harvey et al., 2011)
- k I_D/I_G, Ratio of D-band and G-band from Raman spectra
- 1 H_F, feedstock-dependant heterogeneity index (see text)

^m H_T , temperature-dependant heterogeneity index (see text)

Table 2

Mineral constituents (g·kg⁻¹) of biochars derived from 12 waste biomasses produced at 500°C and biochars

Biochar	Temper	Р	K	Ca	Mg	Cu	Zn	Al	Fe	Mn
feedstock	ature ^a									
Cow manure	500°C	0.646	1.021	3.795	1.569	0.013	0.052	0.506	0.616	0.044
Pig manure		4.386	3.560	3.474	2.801	0.078	0.101	0.455	0.696	0.123
Shrimp hull		2.585	1.896	21.03	0.590	0.013	0.015	0.024	0.023	0.006
Bone dregs		10.86	0.444	31.82	0.508	0.001	0.016	0.010	0.009	0.001
Wastewater sludge		1.702	0.525	6.573	0.645	0.038	0.152	1.929	2.209	0.045
Waste paper		0.124	0.079	22.84	0.584	0.001	0.010	0.361	0.455	0.008
Sawdust		0.061	1.189	2.290	0.348	0.001	0.010	0.097	0.168	0.009
Grass		0.590	5.151	5.236	0.530	0.003	0.023	0.109	0.152	0.011
Wheat straw		0.074	5.182	0.954	0.297	0.001	0.002	0.047	0.074	0.007
Peanut shell		0.166	1.733	1.338	0.458	0.002	0.003	0.218	0.256	0.018
Chlorella		0.717	13.67	17.50	0.779	0.003	0.012	0.547	0.409	0.912
Waterweeds		0.514	3.224	23.13	0.663	0.002	0.010	0.685	0.559	1.025
	H_F^{b}	1.66	1.19	0.93	0.87	1.78	1.37	1.27	1.27	2.00
Dia	2000	1 72	1.40	1 26	1 10	0.021	0.040	0.170	0 272	0.049
Fig	200 C	1.72	1.40	1.50	1.10	0.051	0.040	0.179	0.275	0.048
manure	500°C	2.94 1 30	2.30	2.33	1.00	0.032	0.008	0.303	0.400	0.082
	500 C	4.39	3.30	3.47	2.00	0.078	0.101	0.433	0.090	0.123
	UJU С Н _т с	-4.72	0.40	0.40	0.40	0.004	0.109	0.490	0.749	0.152
	11 ₁	0.40	0.40	0.40	0.40	0.40	0.40	0.40	0.40	0.40
Wheat straw	200°C	0.022	1.55	0.286	0.089	0.000	0.001	0.014	0.022	0.002
	350°C	0.042	2.94	0.540	0.168	0.001	0.001	0.027	0.042	0.004
	500°C	0.074	5.18	0.95	0.297	0.001	0.002	0.047	0.074	0.007
	650°C	0.082	5.75	1.06	0.329	0.001	0.002	0.052	0.082	0.008
	H_{T}	0.51	0.51	0.51	0.51	0.51	0.51	0.51	0.51	0.51

produced from pig manure and wheat straw at 200°C-650°C.

^a Biochar production temperature ^b H_F, Feedstock-depended heterogeneity (see text) ^c H_T, Temperature-depended heterogeneity (see text)

Table 3

Relative proportion (% of biochar-C) of chemical functional groups in biochars derived from 12 feedstocks at 500°C and biochars produced from pig manure and wheat straw at 200°C–650°C, determined by CP-MAS ¹³C NMR.

Biochar feedstock	Temper	Chemical shift (ppm), δ								
	ature ^a	0-46	46-65	65-90	90-108	108-145	145-160	160-185	185-225	225-250
Cow manure	500°C	10.87	-	-	3.59	61.4	7.68	4.79	6.48	6.58
Pig manure		11.7	2.9	16.0	4.20	45.0	2.80	18.6	0.40	-
Shrimp hull		12.7	1.9	2.50	6.00	75.0	4.10	-	-	1.90
Bone dregs		13.4	-	1.20	7.58	72.9	4.89	0.30	-	1.10
Wastewater sludge		18.6	1.5	5.10	5.90	63.4	5.30	-	0.70	1.90
Waste paper		13.7	-	0.20	2.49	75.0	5.28	-	-	5.18
Sawdust		12.4	-	1.00	2.29	78.3	2.89	-	-	5.08
Grass		11.0	0.00	0.00	2.50	80.3	2.79	0.30	-	4.29
Wheat straw		12.6	-	-	1.30	79.1	3.49	-	0.90	4.39
Peanut shell		12.5	0.80	-	1.60	79.6	3.90	-	-	3.90
Chlorella		13.4	0.60	1.30	3.70	77.3	2.80	-	-	3.90
Waterweeds		12.4	2.50	0.10	10.4	65.7	4.00	-	0.70	4.60
	H_{F}^{b}	0.15	1.85	2.20	0.63	0.15	0.35	5.62	8.37	0.65
Pig	200°C	38.8	20.6	27.8	5.37	2.79	0.60	7.56	-	-
manure	350°C	38.3	5.74	11.3	6.82	34.7	3.80	7.98	-	-
	500°C	11.7	2.9	16.0	4.20	45.0	2.80	18.6	0.40	-
	650°C	3.17	-	1.29	7.14	57.6	3.17	1.78	11.5	14.7
	${H_T}^c$	0.80	1.28	0.78	0.23	0.67	0.54	0.78	5.51	4.44
Wheat straw	200°C	3.77	13.3	66.9	14.0	1.69	0.89	_	_	_
	350°C	33.7	4.59	0.80	3.59	49.4	7.98	0.10	_	0.40
	500°C	12.6	-	-	1.30	79.1	3.49	-	0.90	4.39
	650°C	5.47	1.00	3.18	6.17	68.2	4.78	3.68	4.28	3.28
	H_{T}	0.99	1.33	1.87	0.88	0.69	0.69	3.05	1.96	1.10

^a Biochar production temperature

^b H_F, Feedstock-depended heterogeneity

^c H_T, Temperature-depended heterogeneity

Note: The spectra were integrated in the chemical shift (ppm) resonance intervals of 0–46 ppm (alkyl C, mainly CH₂ and CH₃ sp³ carbons), 46–65 ppm (methoxy and N alkyl C from OCH₃, C–N and complex aliphatic carbons), 65–90 ppm (O-alkyl C, such as alcohols and ethers), 90–108 ppm (anomeric carbons in carbohydrate-like structures), 108–145 ppm (aromatic and phenolic carbon), 145–160 ppm (Oxygen aromatic carbon and olefinic sp² carbons), 160–185 ppm (carboxyl, amides and ester) and 185–225 ppm (carbonyls).

Figure Captions

- FIGURE 1. Corrected thermogravimetry patterns of biochars derived from 12 feedstocks at 500°C (a) and biochar produced from pig manure and wheat straw at 200°C–650°C (b).
- FIGURE 2. CP-MAS ¹³C NMR spectrogram of biochars derived from 12 feedstocks at 500°C (a, b) and biochar produced from pig manure (c), and wheat straw at 200°C–650°C (d). 1. Cow manure, 2. Pig manure; 3. Shrimp hull; 4. Bone dregs; 5. Wastewater sludge; 6. Waste paper; 7. Sawdust; 8. Grass; 9. Wheat straw; 10. Peanut shell; 11. Chlorella; 12. Waterweeds.
- FIGURE 3. FTIR spectra of biochars derived from 12 feedstocks at 500°C (a, b) and biochars produced from pig manure (c) and wheat straw at production temperature ranging 200°C–650°C (d).
- FIGURE 4. Raman spectra of biochars derived from 12 feedstocks at 500°C (a, b) and biochars produced from pig manure (c) and wheat straw at production temperature ranging 200°C–650°C (d).
- FIGURE 5. Comparison of feedstock-depended heterogeneity (H_F) and temperature-depended heterogeneity (H_T) for different properties of biochar. See Table 1 for abbreviation.



Fig. 1. Corrected thermogravimetric curves for biochars derived from 12 feedstocks at HTT of 500° C (a) and biochar produced from pig manure and wheat straw at HTT 200° C– 650° C (b).



Fig. 2. CP-MAS ¹³C NMR spectrogram for biochars derived from 12 feedstocks at HTT 500°C (a, b) and for biochar produced from pig manure (c), and wheat straw at HTT ranging 200°C–650°C (d). 1. Cow manure, 2. Pig manure; 3. Shrimp hull; 4. Bone dregs; 5. Wastewater sludge; 6. Waste paper; 7. Sawdust; 8. Grass; 9. Wheat straw; 10. Peanut shell; 11. Chlorella algae; 12. Waterweeds.



Fig. 3. FTIR spectra of biochars derived from 12 feedstocks at 500°C (a, b) and biochars produced from pig manure (c) and wheat straw (d) at production temperature ranging 200°C–650°C.



Fig. 4. Raman spectra of biochars derived from 12 feedstocks at 500°C (a, b) and biochars produced from pig manure (c) and wheat straw (d) at production temperature ranging 200°C–650°C.



Fig. 5. Comparison of feedstock-depended heterogeneity (H_F) and temperature-depended heterogeneity (H_T) for different properties biochar.See Table 1 for abbreviation.