

Electronic Supplemental Information

## Triclosan adsorption using wastewater biosolids-derived biochar

Yiran Tong<sup>a</sup>, Brooke K. Mayer<sup>a</sup>, Patrick J. McNamara<sup>a\*</sup>

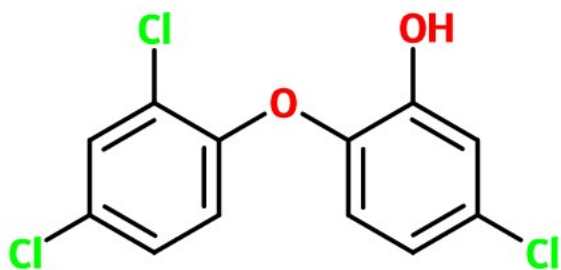
<sup>a</sup> Department of Civil, Construction and Environmental Engineering, 1637 West Wisconsin Avenue, Marquette University, Milwaukee, WI, USA

\*Corresponding author e-mail: [patrick.mcnamara@mu.edu](mailto:patrick.mcnamara@mu.edu)  
phone: (414) 288-2188

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## S1. Triclosan structure and physical-chemical properties

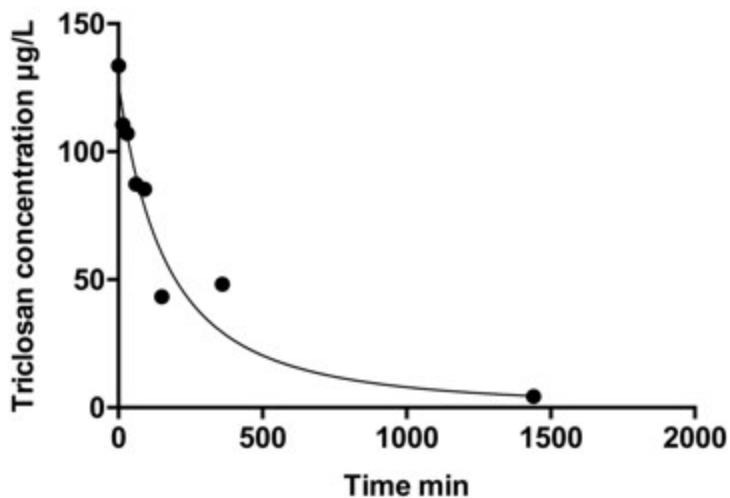
The triclosan molecule, shown in Figure S1, is a trichlorinated binuclear aromatic that is classified as a pesticide and antimicrobial drug<sup>1</sup>. Triclosan has a pKa value of 7.9. The log K<sub>ow</sub> value of 4.76 suggests that the compound is hydrophobic. The vapor pressure of 4.65E-06 mm Hg indicates it is characterized by low volatility.



**Figure S1-** Triclosan chemical structure

## S2. Kinetics studies

To determine the adsorption equilibrium, 600°C HCl-biochar was added at a concentration of 0.4 g/L to deionized water spiked with triclosan at an initial concentration of approximately 130 µg/L. Samples were taken over time and were quantified using the LC-MS. The results are shown in Figure S2. The equilibrium time was determined as 24 hours.



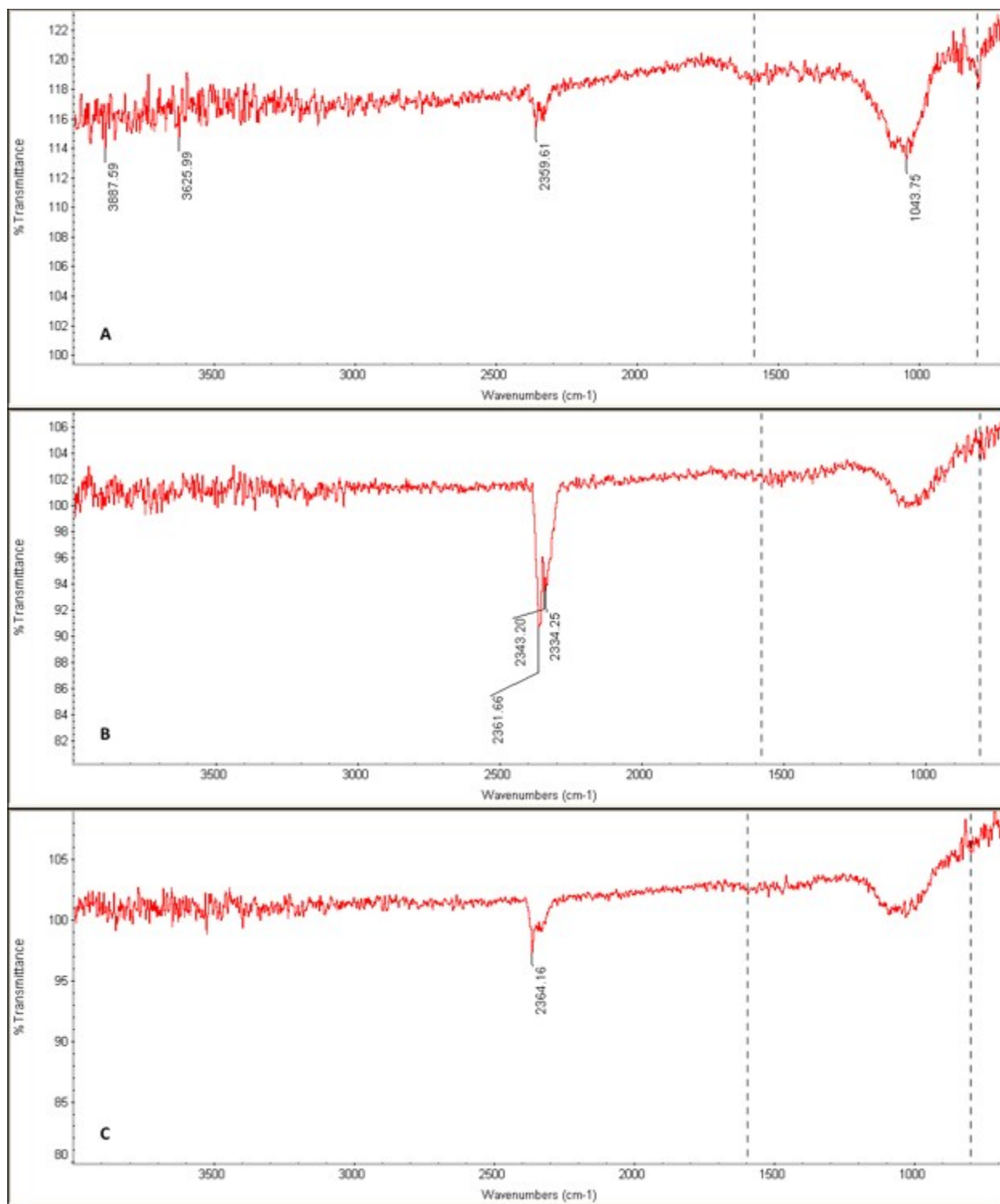
**Figure S2-** Triclosan adsorption kinetics curve

### S3. LC-MS operation

Filtered water samples were mixed with methanol (50/50) in 2 mL amber vials. A binary gradient of Milli-Q water and 100% HPLC-grade methanol was used as the eluent. The method described by Ross et al. (2016) was modified and applied. The gradient began at 80% methanol, raised to 100% methanol at 8 minutes, ramped down to 80% methanol from 8 to 9 minutes and remained at 80% methanol to 13 minutes to allow column re-equilibration. The eluent flow rate was 0.4 mL/min. Sample injections of 20  $\mu$ L were passed through a Phenomenex<sup>®</sup> (Torrance, CA, USA) Luna 3u C18 reverse-phase column (150 $\times$ 3 mm, 100 $\text{\AA}$  pore size). Triclosan was detected in mass-spectrometry with negative electrospray ionization (ESI<sup>-</sup>), at a mass-to-charge ( $m/z$ ) ratio of 287.

#### S4. FT-IR spectra of HCl, NaOH, and Milli-Q water treated biochar

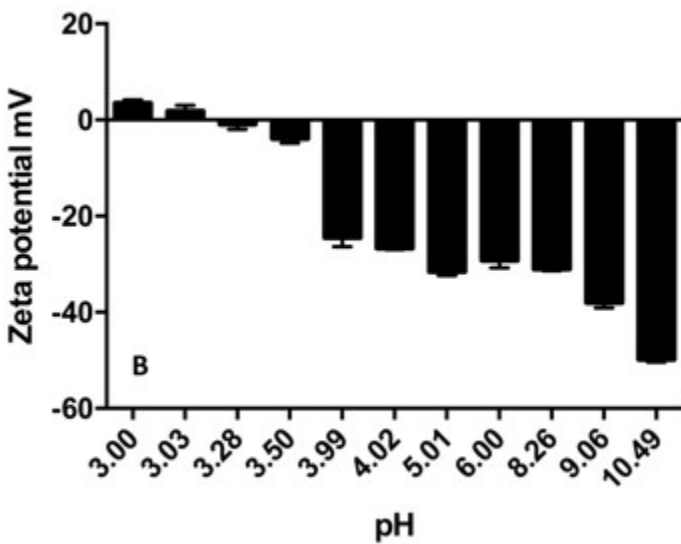
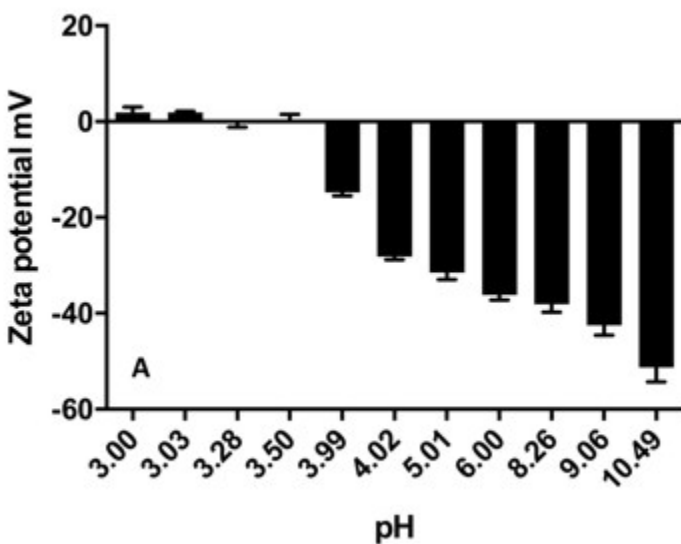
Biochar, pyrolyzed at 600°C and pretreated with HCl, NaOH or Milli-Q water, was ground to fine powder (<10 µm). The scanning of each sample was 32. Variability observed in the spectra may be due to the amorphous nature of biosolids-derived biochar.

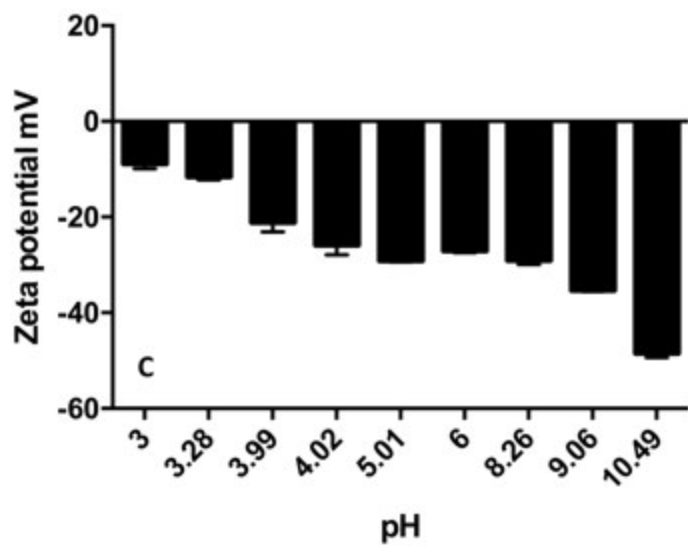


**Figure S3-** FT-IR spectra of biochar produced at 600°C, treated with A) HCl, B) NaOH, and C) Milli-Q water.

### S5. Biochar zeta potential

Biochar, pyrolyzed at 600°C and pretreated with HCl, NaOH or Milli-Q water, was ground to fine powder (<10 µm). Approximately 0.01 g of biochar powder was suspended in 40 mL of Milli-Q water. The solution pH was then adjusted with HCl or NaOH. The zeta potential was immediately measured using a Malvern Zetasizer Nano ZS (Malvern Instruments Ltd, MA, USA). Results are shown in Figure S4.





**Figure S4-** Zeta potentials of biochar produced at 600°C, pretreated with A) HCl, B) NaOH, and C) Milli-Q water.



S6. Isotherm model fitting

**Table S1**-Isotherms of HCl-biochar produced at multiple temperatures and activated carbon fitted with Linear, Langmuir and Freundlich models.

Isotherm model	Equation	Parameter	Sorbent					Activated carbon
			300°C	500°C	600°C	700°C	800°C	
Linear	$Q_e = AC_e + B$	A	0.345	3.09	2.40	4.82	8.29	30.0
		B	120	190	237	153	332	1440
		R <sup>2</sup>	0.049	0.936	0.757	0.917	0.859	0.877
Langmuir	$Q_e = \frac{Q_{max}K_a C_e}{1 + K_a C_e}$	Q <sub>max</sub>	$5.07 \times 10^3$	$9.02 \times 10^4$	$5.70 \times 10^4$	$1.44 \times 10^4$	$4.42 \times 10^3$	$1.50 \times 10^5$
		K <sub>a</sub>	0.0389	0.0113	0.0151	0.0405	0.204	0.0345
		R <sup>2</sup>	0.070	0.867	0.896	0.76	0.977	0.967
Freundlich	$Q_e = K_F C_e^{1/n}$	K <sub>F</sub>	56.5	43.2	62.0	62.9	254	554
		1/n	0.22	0.54	0.45	0.46	0.30	0.44
		R <sup>2</sup>	0.0593	0.912	0.835	0.85	0.977	0.928

S7. Treated wastewater effluent qualities

**Table S2**-Treated municipal wastewater effluent qualities

<b>pH</b>	<b>COD (mg/L)</b>	<b>TOC (mg/L)</b>	<b>Turbidity (NTU)</b>	<b>TSS (mg/L)</b>
<b>7.2</b>	BD*	70.0	2.1	4.6

\*: BD: below detection. Detection limit: 125 mg/L

## S8. References

- 1 R. U. Halden and D. H. Paull, *Environ. Sci. Technol.*, 2005, **39**, 1420–1426.
- 2 J. J. Ross, D. H. Zitomer, T. R. Miller, C. A. Weirich and P. J. McNamara, *Environ. Sci. Water Res. Technol.*, 2016.