## 1 Supplementary Materials

## Suppression of lead environmental risks in cropland soil by biomass ash and its modified product

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18 **S1**:

After the modification of BA, the liquid waste of the hydrothermal reaction was
collected into a 50 ml centrifuge tube and freeze dried for 24 h by using LBJ-10C
freeze dryer (Four-Ring Science Instrument Plant Beijing Corporation, China).
Finally, the dry powder was analyzed by X-ray diffraction (XRD, D8 Advance,
Bruker AXS, Germany) over an angular range of 15 to 90 20 with Cu Kα radiation
(40kV, 40mA) to determine the crystal type of the by-products generated during the
modification procedure.

19 I. XRD Analysis of the by-products generated during the modification procedure

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28 II. XRF Analysis of BA and MA

The relative content of trace elements in BA and MA were determined by X-Ray
Fluorescence spectra (XRF, ZSX pimusII, Rigaku Corporation, Japan) with the
effective measuring range from fluorine to uranium (F to U).

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33 III. XPS Analysis of the incubation soils

34 After incubation, the mass concentration of the elements (Ca, P, O, C, and Si) in soils 35 with or without BA and MA was determined by using X-ray photoelectron 36 spectroscopy (XPS, AXIS ULTRA, Shimadzu, UK) with transmission rod through 37 ultra-high vacuum isolation valve.

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40 IV. Characterization of Soils

41 1. Soils pH was measured using a pH meter (FE20, Shanghai Mettler - Toledo
42 Instrument Co., Ltd., China) with solid / water ratio of 1:2.5 (w/w), and the mixture
43 was stirred and left for 0.5 h at room temperature (25 °C);

44 2. Organic matter (OM) was analyzed using Walkley-Black procedure (Walkley and
45 Black, 1934);

46 3. Cation exchange capacity (CEC) was determined by using ammonium acetate
47 method (USEPA, 2007);

48 4. Available phosphate (P), potassium (K), and nitrogen (N) were determined
49 according to the forest soil analysis method (State Forestry Administration, 1999);

50 **5. Total Pb contents in the soil** were determined by using inductively coupled 51 plasma optical emission spectrometer (ICP-OES, PE Optima 2100DV, Perkin-Elmer, 52 USA) after the soil sample ( $0.2\pm0.0001$  g, grounded and passed through a 0.15 mm 53 sieve) was digested with hydrochloric acid (HCl), nitric acid (HNO<sub>3</sub>), perchloric acid 54 (HClO<sub>4</sub>), and hydrofluoric acid (HF) (4:4:2:8 by volume) on a hot plate. A certified 55 soil reference material (GBW07402, CRM/RM Information center of China) was used 56 to evaluate the validity of the results.

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Walkley, A.; Black, I.A. An examination of the Degtjareff method for determining soil
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63		Solid Waste, Washington D.C. 2007.
64	3.	State Forestry Administration, P.R.C.E. Forest Soil Analysis Methods LY/T 1275-1999,
65		Standard Press of China, Beijing, China. 1999.
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77 Figure S1. XRD pattern of the soluble by-products generated during the modification

78 process/hydrothermal reaction.

Parameter	Biomass ash (BA)	Modified biomass ash (MA)
pН	10.06 <sup>b</sup>	10.24
Ca(mg/g) <sup>a</sup>	69.61	ND °
Cu(mg/kg)	52.3	30.9
Pb(mg/kg)	46.0	44.9
Zn(mg/kg)	207	132
Cr(mg/kg)	55.6	54.6

<sup>a</sup> Total content of Ca in BA was measured by using an atomic absorption spectrometer
(AAS, contrAA 700, Analytik Jena AG, Germany) with Lanthanum chloride as
deionization agent. <sup>b</sup> Mean of three replicates (n=3). <sup>c</sup> ND, no determined.

Parameter	Biomass ash (BA)	Modified biomass ash (MA)
Ca(%)	31.9	30
Si(%)	27.9	29
Fe(%)	12	10.6
K(%)	11.7	7.1
Al(%)	4.31	4.52
Mg(%)	3.13	3.14
Cl(%)	1.76	0.219
P(%)	1.65	9.96
S(%)	1.62	0.127
Ti(%)	1.16	1.06
Na(%)	0.852	2.8
Mn(%)	0.548	0.466
Ba(%)	0.342	0.336
Sr(%)	0.325	0.264

0.272

0.0506

F(%)

Treatment	O 1s	Ca 2p	C 1s	Р 2р	Si 2p
СК	57.15%	0.31%	20.71%	BDL <sup>a</sup>	21.83%
2.5% BA	56.09%	1.61%	20.46%	BDL	21.84%
5% BA	54.51%	1.72%	21.9%	BDL	21.87%
2.5% MA	58.06%	1.93%	19.36%	0.83%	19.82%
5% MA	56.27%	2.95%	18.55%	1.49%	20.74%

115 Table S3. Mass concentration of elements (Ca, P, O, C, and Si) in incubation soils116 with or without BA and MA.

117 <sup>a</sup> BDL, below detection limit.