

Unlocking the Fertilizer Potential of Waste-Derived Biochar

Wolfram Buss,* Anna Bogush, Konstantin Ignatyev, and Ondřej Mašek

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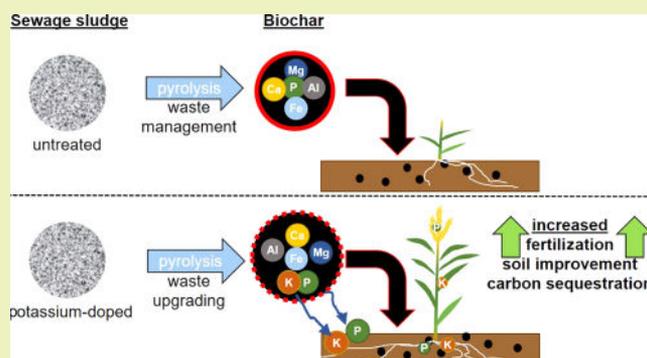
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ABSTRACT: Mankind is facing a phosphorus (P) crisis. P recycling from anthropogenic waste is critical to close the P loop. Pyrolysis could be the ideal treatment for materials, such as sewage sludge (SS), producing a safe, nutrient-rich biochar product while sequestering the inherent carbon (C). However, pyrolyzed sewage sludge typically contains low levels of potassium (K) and plant available P, making the material rather unsuitable for use as fertilizer. Here, a novel treatment was investigated to produce an optimized P and K biochar fertilizer. We doped sewage sludge with a low-cost mineral (2 and 5% potassium acetate) and pyrolyzed it at 700 °C. The percentage water extractable of the total P content in biochar increased by 237 times with 5% K addition compared to the undoped biochar. After six water extractions, all of the K and 16% of P were obtained. Further optimization is feasible through adjustments of the biochar pH or doping the feedstock with other forms of K. Using X-ray absorption near-edge spectroscopy (XANES) and synchrotron X-ray fluorescence (XRF) mapping, we identified highly soluble potassium hydrogen phosphate up to 200–300 μm below the biochar surface. This simple and cost-effective modification enables the use of sewage sludge as safe biochar fertilizer with tailored P availability that also supplies K, improves soil properties, and sequesters C.

KEYWORDS: biochar, pyrolysis, sewage sludge, fertilizer, phosphorus, potassium, XANES



INTRODUCTION

Phosphorus (P) is an essential plant nutrient and a vital planetary resource on which global agriculture and food production depend. While nitrogen (N) can be biologically or artificially fixed from air with the help of energy, this is not an option for P, and existing P resources are limited (estimated to last for 170–890 years).^{1–3} This makes recycling of P from crop residues and anthropogenic wastes to agricultural land an essential sustainability strategy that will help ensure continuing access to P for food security, and as a positive side effect reduce negative effects of P leaching into water bodies (algae blooms and eutrophication). Changes in legislation, e.g., in Germany, Switzerland, and the EU, have increased the incentive and urgency of finding alternative ways of P recycling from waste materials, such as sewage sludge (SS).^{4–6}

Sewage sludge, a waste product from sewage treatment in wastewater treatment plants, has agronomically relevant levels of N (1.5–4.0%), P (0.3–1.2%), and organic carbon,⁷ and therefore, it has been used as a soil amendment on agricultural soils for centuries to maintain productivity.⁸ However, application of untreated sewage sludge poses health risks due to pathogens and organic and inorganic contaminants.⁷ Emerging contaminants, such as pharmaceuticals, personal care products, flame retardants, and microplastics, have further increased the concern about the agricultural use of sewage sludge.^{9–11} Processes, such as

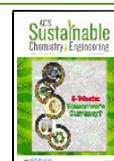
composting and anaerobic digestion, can address these associated health concerns at best only partially.^{11–13}

Thermal treatments, incineration and pyrolysis, are alternative processing methods. Pyrolysis is a thermochemical conversion process under oxygen-limited conditions. It sterilizes the material and reduces the concentration of organic toxins (e.g., polycyclic aromatic hydrocarbons (PAHs), polychlorinated biphenyls (PCBs), and dioxins), pharmaceuticals (e.g., hormones), and other microconstituents (plastics, surfactants).^{12–17} It also immobilizes potentially toxic elements (PTE) present in the feedstock, such as copper (Cu), cadmium (Cd), and chromium (Cr), reducing the risk of leaching and plant uptake.^{18,19} In contrast to incineration, pyrolysis retains most of the carbon present in sewage sludge and even converts it into a recalcitrant form. Therefore, if pyrolysis vapors and liquids are used to fuel the process or renewable energies, pyrolysis can be considered the most sustainable treatment for sewage sludge from the perspective of atmospheric carbon removal and storage, and environmental impacts.

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Sewage sludge biochar, pyrolyzed sewage sludge that is applied to soil, typically has levels of potassium (K) of 0.2–1.7%,²⁰ rather low for use as an agricultural fertilizer and can even sorb K from soil.²¹ Although most sewage sludge biochars and incineration ashes contain agriculturally relevant levels of P (1.0–20%),^{20,22} little P is mobile with the P availability being lower than in the source material.^{23–25} Slow-P-release fertilizer might be desired in certain applications, yet, in modern agriculture, immediate and, most importantly, predictable P release is needed. Since the P availability in biochar is highly variable,²² current biochars are unsuitable as P-fertilizer and modification and optimization of biochar are required.

Previously, it was shown that additions of Ca, Fe, and Mg to biomass before pyrolysis can decrease the P availability in biochar.²⁶ We anticipate that K-doping can do the opposite: potassium phosphate salts demonstrate very high solubilities²⁷ and our hypothesis is that pyrolysis could result in preferential formation of potassium phosphates if K is present in excess increasing P availability in biochar. We used potassium acetate for biochar doping since it is a common, low-cost salt (700–1000 USD t⁻¹)²⁸ that is used for general applications such as de-icing. The aim is to produce a more complete carbon sequestration-fertilizer product with well-balanced availability of nutrients. Sewage sludge was doped with 2% (weight %) and 5% K prior to pyrolysis at 700 °C and the nutrient and PTE availability, location of P, K, and Ca, and P speciation in biochar was studied using extractions, synchrotron X-ray fluorescence (XRF) mapping, and X-ray absorption near-edge spectroscopy (XANES).

MATERIALS AND METHODS

Feedstock Preparation and Pyrolysis. The effect of the feedstock's K content on biochar yield and properties was tested by altering the K content in sewage sludge. Sewage sludge was used because it is anthropogenic waste material with high nutrient contents that needs treatment before soil application. The sewage sludge was provided by Scottish Water in an undigested, dried, and granulated form.

As described previously,²⁹ an aqueous solution of potassium acetate was sprayed onto oven-dried sewage sludge to restore the original moisture content and add 2 and 5% (weight %) K to dry feedstock materials. The application of 2 and 5% K was based on previous experiments.

The 2 and 5% K-doped sewage sludge materials were pyrolyzed using the continuous pyrolysis unit (auger reactor) at the UK Biochar Research Centre, University of Edinburgh. The pyrolysis unit is heated by an electric split-tube furnace (inner diameter 100 mm), and the feedstock residence time in the heated zone is around 21.5 min. Details about the unit and a schematic are published in Buss et al.³⁰ Based on previous results and the availability of the 700 °C sewage sludge biochar as Edinburgh Standard Biochar³¹ (based on the same feedstock batch), a temperature of 700 °C was chosen.

The undoped sewage sludge biochar used as a reference material here was produced using the pilot scale continuous pyrolysis unit of the UKBRC (details in Mašek et al.).³² We were able to show previously that key properties of the biochars produced in our two continuous pyrolysis units are not significantly different from each other.³²

The materials are abbreviated as follows: SS, sewage sludge; SS 700, sewage sludge 700 °C; SS 700 + 2% K, sewage sludge 700 °C with 2% K; SS 700 + 5% K, sewage sludge 700 °C with 5% K doping.

Thermogravimetric Analysis (TGA). A Mettler-Toledo TGA/DSC1 was used to perform pyrolysis followed by proximate analysis for accurate yield determination of the sewage sludge biochars.

Around 70–100 mg (one granule) undoped and K-doped sewage sludge was placed in a 150 μL alumina crucible. The following

temperature profile was applied: drying at 110 °C for 10 min, 90 °C min⁻¹ dynamic phase to heat to 700 °C and a static phase of 10 min, matching the heating rate, and residence time at the highest treatment temperature (HTT) of the continuous units.³²

Subsequently, the biochar was ground up in the crucibles used for TGA pyrolysis and a standard proximate analysis protocol was used in the TGA. During proximate analysis, the biochar is heated up to 110 °C in a nitrogen atmosphere to drive-off moisture, followed by heating to 900 °C to determine the volatile matter content. The subsequent introduction of air oxidizes the sample, and the fixed carbon (FC) and ash contents were determined. The equations used to calculate the yields and FC content can be found in Buss et al.³³

Biochar Characterization. Subsamples (typically at least 20–50% of the total biochar amount) of the doped and undoped biochars were ground up with mortar and pestle. Further subsamples were taken, and the following analyses performed.

Electric Conductivity (EC) and pH. The biochars were extracted in glass vials with deionized (DI) water in a solid-to-liquid ratio of 1:20 according to Rajkovich et al.³⁴ The samples were shaken on an orbital shaker for 1.5 h at 150 rpm and subsequently analyzed with a Hach HQ40d portable meter using a Hach conductivity probe CDC 401 and a sension gel-filled pH-electrode Hach 51935-00.

Extractions and Digestions. The biochars were digested using modified dry ashing with a combination of HNO₃ and H₂O₂ and dry ashing at 500 °C for 8 h.³⁵ More details and modifications to the original method can be found in Buss et al.³⁰

Different extraction methods were applied to characterize the availability of nutrients and PTEs in biochar. Extraction with 0.01 M CaCl₂ was identified as the best extraction technique for available P in Shepherd et al.³⁶ In addition, DI water extraction and a formic acid (FA) extraction were chosen, which also showed good correlation with plant P uptake in Shepherd et al. and Wang et al.^{36,37}

The extraction protocol for CaCl₂ and water extraction was the following: 1.5 g of ground biochar was extracted with 15 mL of solution in 50 mL polypropylene centrifuge tubes. The tubes were shaken on an orbital shaker for 2 h at 150 rpm. Subsequently, the samples were filtered with Whatman No. 1 filter paper. Blanks were added and triplicate extractions conducted, respectively.

The formic acid extraction was performed according to Shepherd et al.:³⁶ 200 mg of biochar was weighted into 50 mL centrifuge tubes in triplicates, and 20 mL 2% FA was added. Reagent blanks were prepared, and the samples were shaken for 2 h at 150 rpm and filtered with Whatman No. 1.

The digests and extracts were analyzed via inductively coupled plasma optical emission spectroscopy (ICP-OES) as described in the **Elemental Analysis** section. The % available of the total elemental content was calculated.

Repeated Water Extraction (Edinburgh Biochar Toolkit D). As described in Buss et al.,³⁸ 5 g of biochar and sewage sludge were extracted with DI water and 0.05 M 2-ocatanol (nonionic surfactant to mitigate any short-term hydrophobic effects of biochar). The experiment was performed in glass bottles, which were shaken at 150 rpm for 1.5 h. After filtering, the solution was analyzed for P via ICP-OES. The biochar was extracted again with the same DI water-to-surfactant ratio. The experiment was repeated six times in triplicates (extraction steps 3, 4, 5, and 6 with 2 and 5% K-amended biochars only performed in duplicates).

The accumulated percentage elemental extractability was given as averages with standard deviation (error propagation was applied).

Elemental Analysis. All extracts and digests were analyzed by ICP-OES (Varian Vista Pro) for Al, Ca, Mg, K, P and Fe (concentrations of As, Cd, Co, Cr, Cu, Hg, Mn, Ni, Pb, and Zn in the **Supporting Information**) using calibrations from 0.01 to 25 ppm. Detailed descriptions of the ICP-OES procedure, and data processing can be found in Buss et al.³⁰

Mineralogical Composition of Sewage Sludge Samples. Biochar samples were ground up and sieved to a fraction size below 0.063 mm. X-ray diffraction (XRD) was used to characterize the crystalline phases present in the sewage sludge samples. A Bruker D8 advance XRD instrument with a Cu anode was used at 40 mA and 40

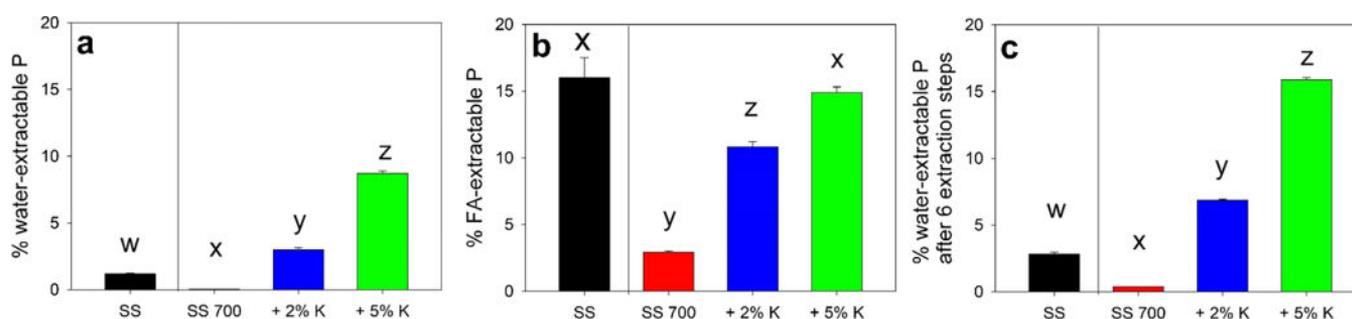


Figure 1. (a) Water- and (b) formic acid (FA)-extractable P content as % of the total content in sewage sludge (SS) and sewage sludge biochar. Panel (c) shows the P concentrations from biochar extracted in six extraction steps (Edinburgh Toolkit D). Biochar was produced at 700 °C without addition (SS 700) and with 2 and 5% K addition prior to pyrolysis. Mean and standard deviation of three replicates are shown. Letters show significant differences among the treatments ($p < 0.05$, one-way ANOVA).

Table 1. Properties of Sewage Sludge (SS) and Sewage Sludge Biochar Produced at 700 °C without Addition (SS 700) and with 2 and 5% K Addition Prior to Pyrolysis^a

		SS	SS 700	+2% K	+5% K
yield					
biochar yield	% feedstock		37.62 ± 0.37a	39.40 ± 0.09b	39.88 ± 0.22b
ash-free biochar yield	% feedstock		12.17 ± 0.21a	13.40 ± 0.04b	13.59 ± 0.07b
ash-free biochar yield	% ash-free feedstock		16.33 ± 0.31a	18.11 ± 0.06b	18.44 ± 0.12b
FC yield	% feedstock		6.44 ± 0.16a	7.66 ± 0.08b	7.10 ± 0.12c
FC yield	% ash-free feedstock		8.64 ± 0.19a	10.34 ± 0.10b	9.63 ± 0.14c
proximate analysis					
volatile matter	% ash-free	89.4 ± 0.4 ^b	47.1 ± 2.1a	42.9 ± 0.5b	47.7 ± 1.1a
FC	% ash-free	10.6 ± 0.3 ^b	52.9 ± 2.1a	57.1 ± 0.5b	52.3 ± 1.1a
FC	% dry	17.1 ± 0.6 ^b	17.1 ± 0.6a	19.4 ± 0.2b	17.8 ± 0.4a
ash	% dry	27.7 ± 0.5 ^b	67.6 ± 0.3a	66.0 ± 0.1b	65.9 ± 0.1a
pH		6.36 ± 0.10	9.60 ± 0.33	10.46 ± 0.12	10.77 ± 0.11
EC	μS cm ⁻¹	368 ± 11	162.6 ± 17.8	1134 ± 158	2575 ± 455
K concentration					
total	mg kg ⁻¹	1157 ± 42	3277 ± 35	23 316 ± 284	37 845 ± 320
CaCl ₂	mg kg ⁻¹	239 ± 3	23 ± 1.6	7135 ± 746	13 580 ± 240
water	mg kg ⁻¹	206 ± 5	32 ± 0.5	6984 ± 352	13 471 ± 80
water-6 days	mg kg ⁻¹	621 ± 15	140 ± 1.1	26 059 ± 2823	41 161 ± 3066
P concentration					
total	mg kg ⁻¹	10 984 ± 354	20 309 ± 734	18 965 ± 594	19 555 ± 265
CaCl ₂	mg kg ⁻¹	166 ± 2.3	1.1 ± 0.3	<0.26	340 ± 19
water	mg kg ⁻¹	129 ± 7.0	9.6 ± 0.5	564 ± 21	1700 ± 35
water-6 days	mg kg ⁻¹	311 ± 7.8	75 ± 1.2	1300 ± 10	3106 ± 17
formic acid (FA)	mg kg ⁻¹	1756 ± 151	592 ± 11	2055 ± 29	2920 ± 71

^aFC, fixed carbon; EC, electric conductivity. In most cases, mean and standard deviation of three replicates shown. Yield and proximate analysis results are on dry basis. Different numbers indicate statistical differences among the treatments. ^bNot tested: for proximate analysis, only the biochars, not the feedstock, were compared with each other statistically, as appropriate.

kV with a NaI detector, which analyzed at 6 s/step from 2 to 65° in 0.025°/step increments. The raw data were evaluated with the software TOPAS 3.0 Rietveld. The biochar samples were spiked with 20% calcite to quantitatively measure the composition of mineral in the samples relative to a known concentration of calcite.

Synchrotron X-Ray Analysis of P-Species in Biochar. The P distribution and speciation in the dehydrated SS 700 and SS 700 + 5% K biochars were investigated by X-ray fluorescence (XRF) elemental mapping and X-ray absorption near-edge spectroscopy (XANES) at the I18 beamline at Diamond Light Source (U.K.) using a monochromator with Si(111) crystals and a helium-purged sample chamber to reduce X-ray attenuation.

X-Ray Fluorescence (XRF) Elemental Mapping. The spherical particles of biochars were cut and placed onto Kapton tape. The XRF elemental mapping was run on the dehydrated SS 700 and SS 700 + 5% K biochar particles with a resolution of 2–3 μm to investigate the P distribution and association of P with Ca and K in the samples.

X-Ray Absorption Near-Edge Spectroscopy (XANES). P K-edge XANES measurements of particular phases enriched with P and average P spectra were collected to examine the P speciation in the dehydrated SS 700 and SS 700 + 5% K biochars. The reference materials of P-containing compounds of analytical quality reagents (Sigma-Aldrich) selected for the XANES analysis were: hydroxyapatite (Ca₅(PO₄)₃(OH)), calcium phosphate (Ca₃(PO₄)₂), calcium phosphate dibasic (CaHPO₄), calcium phosphate monobasic (Ca(H₂PO₄)₂), potassium phosphate (K₃PO₄), potassium phosphate monobasic (H₂KPO₄), potassium phosphate dibasic (K₂HPO₄), iron(III) phosphate dihydrate (FePO₄·2H₂O), and aluminum phosphate (AlPO₄). The reference material spectra were recorded in transmission mode using an ionization chamber. The sample spectra were recorded in fluorescence mode using a four-element energy-dispersive fluorescence detector.

Data Analysis and Statistics. The data were tested for normality with a Shapiro–Wilk test before conducting one-way analysis of

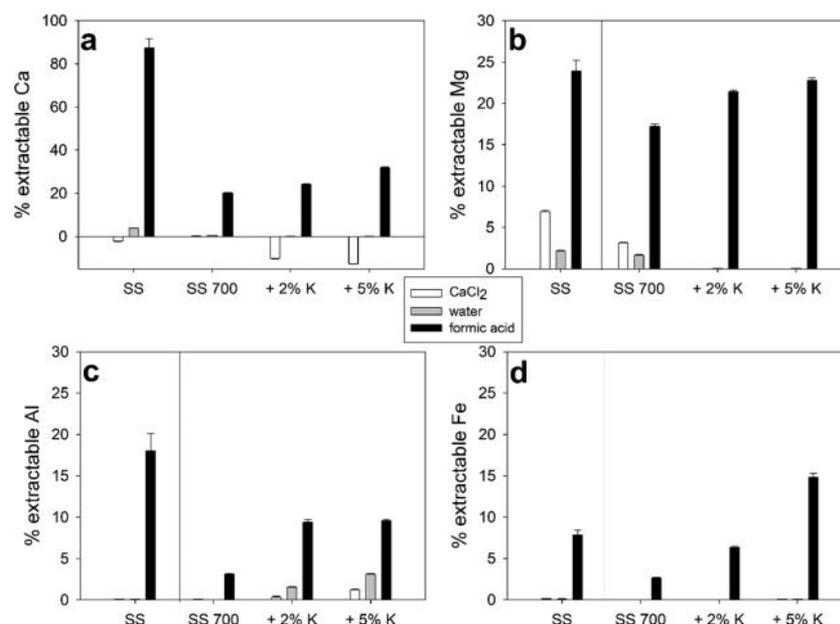


Figure 2. Percentage water-, 0.01 M CaCl₂, and formic acid extractable elements. (a) Ca, (b) Mg, (c) Al, and (d) Fe content as % of the total content in sewage sludge (SS) and sewage sludge biochar. Biochar was produced at 700 °C without addition (SS 700) and with 2 and 5% K addition prior to pyrolysis. Mean and standard deviation of three replicates are shown.

variance (ANOVA) with SigmaPlot 11 followed by Student–Newman–Keuls pairwise post hoc comparison test. The data set from single-step and repeated water extraction of P failed the Shapiro–Wilk test. Yet, one-way ANOVAs are robust against violation of the normality assumption.³⁹

The XANES spectral analysis was performed using Athena.⁴⁰ Linear combination fitting (LCF) of the spectra for P in the dehydrated SS 700 and SS 700 + 5% K biochars with those of the reference materials was performed in derivative space, to identify P species that were present in the investigated samples.

RESULTS AND DISCUSSION

Elevated Phosphorus (P) and Potassium (K) Availability. The extractable P content in untreated sewage sludge was low, only 1.5 and 1.2% of the total P was extracted with 0.01 M CaCl₂ and water, respectively; formic acid (FA) extracted 16% of the total P (Figure 1 and Table 1). After pyrolysis at 700 °C, the P availability was even lower, CaCl₂ extracted <0.01% of the total P, water 0.04%, and FA 2.9%. In Zhang et al.,²² no P was water extractable in sewage sludge biochar produced >400 °C supporting our results.,

Doping of sewage sludge with K successfully increased the water extractable of the total P content in the one-step extractions from 0.04% in the undoped biochar to 3.0 and 8.7% with 2 and 5% K addition, respectively, an increase by 81 and 237 times (Figure 1). In addition, the treatment increased 0.01 M CaCl₂ and FA extractable P (Table 1). Repeated water extractions (Edinburgh Tool Kit D), simulating P release over time, continued to extract P (Figure S1–Sn). At the end of sixth extraction steps, 7 and 16% of the total P were extracted in the 2 and 5% K-amended biochar, respectively, compared to only 0.4 and 3% in the SS 700 and untreated sewage sludge (Figure 1). In the 5% K-amended biochar, the same amount of P was extracted with FA in one extraction step as with water in six extraction steps.

The total K concentration in sewage sludge was around 10 times lower than the concentration of P, which is not unusual for sewage sludge samples.^{21,41} In the undoped sewage sludge

biochar, water extracted only low amounts of K (Table 1). The addition of K prior to pyrolysis increased the extractable fraction of K (CaCl₂ and water) to 30–50% of the total K content in a single extraction step (Table 1). Repeated water extraction resulted in 100% recovery of the total K content in the two amended biochars (Table 1 and Figure S1–Sn). High amounts of K immediately available to plants can cause localized toxic effects and low nutrient-use efficiency due to K leaching through soil, an issue observed for biochar and combustion ashes.^{23,42} Therefore, initial availability of one-third to one-half of K but eventually full accessibility of K in the enhanced biochar could be a good balance but needs to be confirmed in plant trials. Overall, the results are very promising for K-enhanced sewage sludge biochar to be used as P and K fertilizers.

K-Doping Changes P Speciation in Biochar. Extractability of P-Binding Partners. Elements such as Ca, Mg, Al, and Fe bind to P to form minerals of low solubility²⁷ that limit P mobility and plant uptake in soil and biochar.^{43–45} Sorption of P to Al, Ca, Mg, and Fe is also the main mechanism responsible for P retention by biochar.⁴⁶ Changes in extractability of P-binding partners in sewage sludge after pyrolysis and with K-doping can be evidence for a change in P speciation.

The extractability of Ca, Mg, Al, and Fe in undoped sewage sludge biochar was lower than in the untreated sewage sludge (in all three extractants; Figure 2). The P in biochar feedstocks from manure and plant residues is predominantly present in organic forms, e.g., in DNA and phytate,^{47,48} while sewage sludge is mostly composed of inorganic, ortho-P.⁴⁴ Pyrolysis converts a proportion of P in both types of material into P-containing minerals of low water solubility (Ca, Mg, Al, and Fe phosphates),^{25,44,45,47,48} reducing the extractability of both, P and P-binding partners.

The hypothesis is that with K addition, instead of forming salts of low water solubility, during pyrolysis P binds preferentially to K to form salts of high water solubility, such

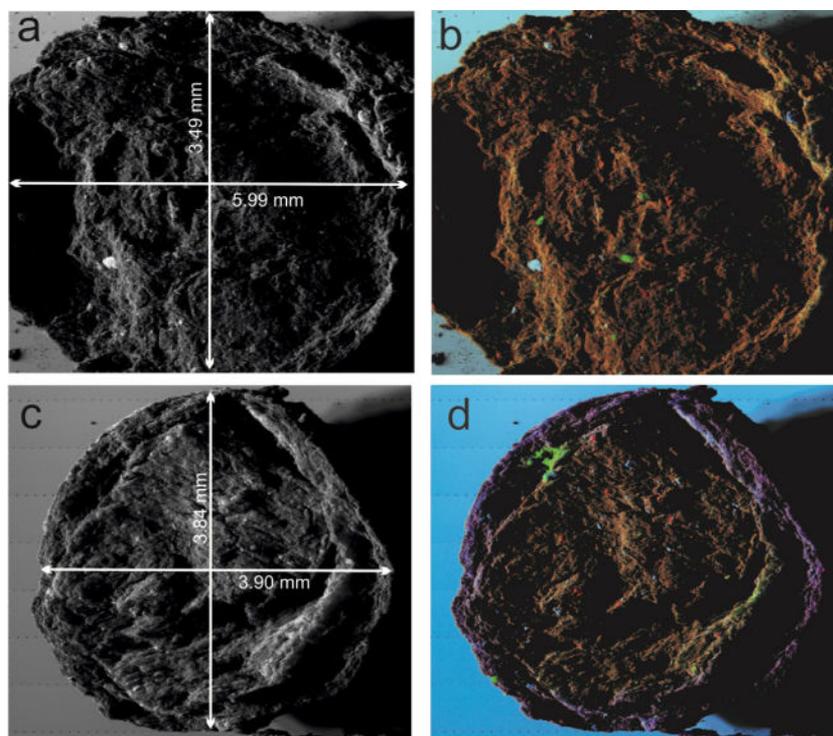


Figure 3. Picture (a–d) are images of the internal view of the biochar spherical particles (a: SS 700; c: SS 700 + 5% K) and the synchrotron-XRF maps (b: SS 700; d: SS 700 + 5% K) with P—red, Ca—green, K—blue.

as potassium dihydrogen phosphate (KH_2PO_4), potassium hydrogen phosphate (K_2HPO_4), and potassium phosphate (K_3PO_4).²⁷ Ca, Mg, Al, and Fe that typically bind to P in biochar instead form other, more soluble compounds, increasing their extractability in the K-amended biochar compared to the unamended biochar. This is indeed the case for most samples giving evidence supporting our hypothesis (Figure 2). Yet, the concentration of Ca and Mg in the water and CaCl_2 extracts was lower in the K-doped biochars compared to the undoped SS 700 (net decrease of Ca in Figure 3 is a result of subtracting Ca concentration in the blank from the analyte).

Two different factors need to be considered that determine the concentration of P and Ca, Mg, Al, and Fe in the analyte: first, the initial solubility of the biochar inherent minerals and, second, the solution conditions and composition as once dissolved P can precipitate over time, e.g., with Ca, which then is not quantified by the detector. Formation and precipitation of significant quantities of calcium phosphate are only possible when high phosphate concentrations are present in the solution⁴⁹ as is the case in the 5% K-doped biochar. The solution pH in water and CaCl_2 extracts were close or slightly lower than the pH displayed in Table 1 (pH \sim 10 for the amended biochars), which are suitable conditions for calcium phosphate but not for aluminum and iron phosphate precipitation. Subsequently, with ample P present in the solution and a suitable pH for Ca and Mg–P precipitation, Ca and Mg concentrations in the water and CaCl_2 extracts of the K-amended biochars were lower than in the undoped biochar (less P present for precipitation) and sewage sludge sample (lower pH unsuitable for precipitation).

In the 5% K-doped biochar, FA and water extracted the same amount of P in one and six extraction steps, respectively (Table 1). In contrast, FA extracted nearly 10 times more P

than the water extractions in the SS 700 biochar as also shown in Zwetsloot et al.⁵⁰ This indicates that the P in the doped sample is extracted regardless of the solution pH and type of extractant, while specific conditions are necessary to extract the P from SS 700. This is further proof that the P species have changed with K addition.

With pyrolysis, the extractability of Ca, Mg, Al, and Fe decreased due to the conversion of organic to mineral P that forms minerals with these P-binding partners. While K addition prior to pyrolysis increased the water solubility of P-binding partners in biochar, it also increased Ca and Mg precipitation (as phosphates) when extracted at the native biochar pH, hence limiting Ca, Mg, and likely P availability. According to Barrow, in soils, calcium phosphate precipitation is not a factor that significantly contributes to P immobilization.⁴⁹ Still, in certain soils (excess availability of Ca and Mg), a lower biochar pH (or biochar-soil solution pH) could further increase the immediate P availability. This is an important aspect to consider in the design of biochar for specific soils and applications requiring controlled P delivery.

P-Speciation Analyzed by XRD, Sy-XRF, and XANES. We used Sy-XRF mapping to identify the location of Ca, K, and P in biochar to track any changes with K-doping, Ca–P being the most common P-containing mineral at high pyrolysis temperatures.⁴⁵ The brown background of the interior of SS 700 biochar (Figure 3b) shows an even distribution of P and Ca, which likely associate with each other (red—P and green—Ca results in brown). In the K-amended biochar (Figure 3d), a purple ring enveloping the biochar center is visible up to 200–300 μm from the biochar surface, clearly showing that K and P associate with each other (red—P and blue—K results in purple). The center of the biochar particle is still colored brown demonstrating that there are still significant amounts of Ca–P present in the center. The existence of the ring of K is

likely explained by K not having fully penetrated into the biochar center during initial doping. A more equal distribution of K during doping, e.g., through longer soaking in the K-rich solution could further increase the K–P association and, henceforth, P availability. This hypothesis will be tested in follow-up studies. The results demonstrate not only the feasibility of increasing the P availability but also that further optimization of K distribution for maximum P availability is viable.

We could neither identify distinct differences between the XRD spectra of the biochars (Figure S2–Sn) nor any particular P crystalline phases. This indicates that most of the P phases were amorphous or present in small concentrations of many different minerals, below the level of detection. This was confirmed in Qian and Jiang, who did not identify any P crystalline phases in sewage sludge biochar produced by pyrolysis at 400–700 °C using XRD; only stanfieldite ($\text{Ca}_4\text{Mg}_5(\text{PO}_4)_6$) was identified in biochar produced at 800 °C.⁴⁴

Therefore, we analyzed SS 700 and the 5% K-doped biochar with P K-edge XANES in five replicates enabling a higher resolution. The results (Figure 4) of three replicates of SS 700 (SS 700(1)–SS 700(3)) demonstrate that the spectra mainly associated with aluminum phosphate (AlPO_4) with a dominant narrow peak at around 2152.5 eV (the P absorption edge)

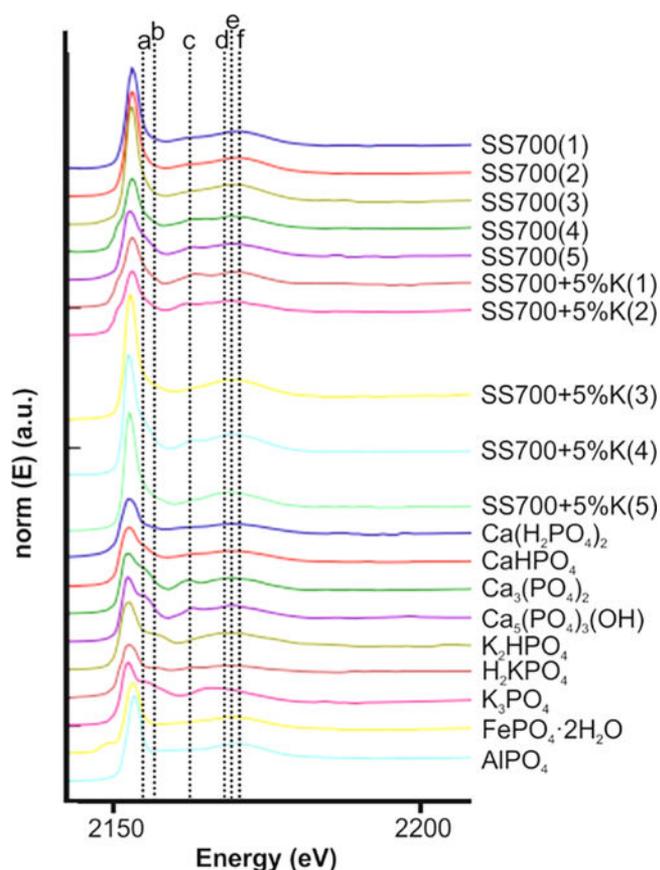


Figure 4. Phosphorus K-edge XANES spectra of sewage sludge biochars with 5% K addition prior to pyrolysis (SS 700 + 5% K) and without (SS 700) produced at 700 °C in five replicates ((1)–(5)). In addition, reference materials of P-containing compounds likely present in sewage sludge (biochar) are shown. The dotted lines and letters are references indicating energy levels that characterize unique spectral features for different P species.

followed by a much broader peak at reference line (f). In contrast, the spectra SS 700(4) and SS 700(5) mainly associated with the hydroxyapatite ($\text{Ca}_5(\text{PO}_4)_3(\text{OH})$) spectrum that has a dominant, much broader peak at around 2152.5 eV with a pronounced shoulder at reference line (a) followed by two broad peaks at around lines (c) and (e) (Figure 4). Results from linear combination fitting (LCF) of the P K-edge XANES spectra for sewage sludge biochars confirm the occurrence of AlPO_4 and $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$ (Table S1–Sn). Both compounds were also identified as key P-containing compounds in Sun et al.⁵¹ Hydroxyapatite and aluminum phosphate are both poorly soluble P compounds with solubility products of 3.04×10^{-59} for $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$ and 6.3×10^{-19} for AlPO_4 explaining the low solubility of P in SS 700 (Figure 1).⁵²

The LCF results indicate that hydroxyapatite and aluminum phosphate are still present in the 5% K-doped biochar; however, in addition K_2HPO_4 , highly water soluble (168/100 g H_2O^{27}), appeared in the modified biochar (Table S1–Sn). The K_2HPO_4 spectrum has a dominant narrow peak at 2152.5 eV with a broad post-edge shoulder at line (b) followed by a much broader peak at line (d) (Figure 4). The LCF of the spectra for P in the 5% K-doped biochar (replicates 3–5) shows residues and a poor correlation fit with the tested reference materials indicating the presence of other P-bearing phases that were not covered by used reference materials.

In summary, our results demonstrate that with the addition of K before pyrolysis P-minerals of very high solubility, such as potassium hydrogen phosphates, were formed confirming our hypothesis that P availability can be modified and customized with simple and cost-effective mineral additions.

Carbon Sequestration Potential of Doped Biochar.

Besides agronomically relevant levels of nutrients, sewage sludge biochar contains stable carbon and when applied to soil can sequester atmospheric carbon dioxide for centuries.⁵³ Therefore, we tested the biochar yield and the yield of fixed carbon (FC) as influenced by K addition, FC being a proxy for what remains in soil after ~100 years.^{54–56}

The biochar yield of sewage sludge (in % of dry feedstock) was 37.6%, which increased to 39.4 and 39.9% with the addition of 2 and 5% K, respectively (Table 1). The FC yield per feedstock and per ash-free feedstock input both increased by around 20 and 10% with 2 and 5% K, respectively (in relative terms, Table 1) (see more details on calculation and interpretation in Buss et al.³³). Therefore, 2% K-doping maximizes the carbon sequestration potential of sewage sludge biochar.

Previously, we demonstrated that 2% K-doping boosted the carbon sequestration potential of *Miscanthus* biochar by 45% due to catalytic effects of alkali (and earth alkaline) metals, which increase biochar formation and yield.²⁹ The sewage sludge in our current study had a five times higher concentration of inherent alkali and earth alkaline metals (1.5% K, Ca and Mg; Table S1–Sn) than the *Miscanthus* biomass (0.3% K, Ca and Mg; unpublished data). Therefore, the addition of 2% K had a smaller effect on overall alkali and earth alkaline metal content in sewage sludge compared to the content in *Miscanthus*, likely resulting in less catalytic effects and biochar yield increases.

In Buss et al., blending of pine biomass with wood ash, rich in alkali and earth alkaline metals, increased the FC yield (ash-free feedstock) significantly and the highest wood ash addition had the strongest effect.³³ In our current study, 2% K

performed better in terms of FC yield than 5% K-doping (Table 1), demonstrating that there is an optimum K addition for maximum FC yield. This is a significant finding and highlights the need to investigate different feedstocks separately and tailor the biochar-mineral formulation individually to maximize the carbon sequestration potential.

Sewage Sludge Biochar Fertilizer in Agriculture. The P content in the biochar under investigation (2%) is low for biochar produced from sewage sludge, which can have P contents of 6–20%, more P than present in refined super-P fertilizer.^{18,22,41} In our study, water and FA both extracted 16% of the total P in six and one extraction step, respectively. Assuming an average sewage sludge biochar P content of 10%^{18,22,41} and a P availability of 16%, in total 1.6% of the biochar, would likely be immediately plant-available P. At a P application rate of 40 kg ha⁻¹, ~2.5 t ha⁻¹ of biochar would satisfy the typical P demand of winter wheat.⁵⁷ In addition to the immediately available P, this amendment adds extra P to the soil that can be used in subsequent growing cycles.

Application of 2.5 t ha⁻¹ of a sewage sludge biochar with the same composition as the one in our study would also supply ~95 kg K ha⁻¹ (demand for winter wheat ~133 kg K ha⁻¹).⁵⁷ More K could be added to the sewage sludge to make even more P available and to provide enough K for specific crops, depending on the application. In addition, such biochar fertilizer (at 2.5 t ha⁻¹) sequesters 445 kg ha⁻¹ of carbon likely stable for >100 years (fixed carbon dry basis in Table 1), supply micronutrients, such as Ca, Mg, B, Mn, and Mo (Table S1–S_n), and increase the pH of acidic soil (pH 10.77; Table 1). The effect of sewage sludge biochar on soil pH was demonstrated in Buss et al.⁵⁸ Therefore, K-amended biochar can be a great multifunctional, environmentally sustainable fertilizer.

An often raised concern regarding the application of sewage sludge in agriculture is its levels of potentially toxic elements (PTEs),^{14,59} which are enriched in biochar due to decomposition and volatilization of a proportion of the organic matter during pyrolysis, while the PTEs remain.³⁰ Yet, the availability of PTEs drastically decreased after sewage sludge pyrolysis, in both, undoped and doped biochar (Table S2), e.g., the water-extractable contents of Cu, Ni, and Zn were all reduced by 82–93% after pyrolysis of 5% K-amended sewage sludge. Therefore, as concluded by other authors, in terms of PTE toxicity, sewage sludge biochar application in agriculture is likely to be much safer than the still widely practiced application of unpyrolyzed sewage sludge.^{18,42,60} Importantly, pyrolysis destroys a whole suite of organic contaminants present in sewage sludge resulting in markedly increased safety.^{12–17}

To further optimize (sewage sludge) biochar as a fertilizer, refinement of the method presented here is possible, e.g., through a change in biochar pH, feedstock K/P ratio, concentration, and form of K minerals and K distribution in the feedstock (soaking time). This new class of fertilizer contributes to nutrient recycling and reduced nutrient leaching, reduces the dependence on fertilizer imports (as intended as part of the new fertilizer regulation in the EU⁶), and helps to comply with new circular economy legislation, e.g., the German and Swiss legislation on P recycling from wastewater.^{4,5}

CONCLUSIONS

Sewage sludge-based biochar fertilizers offer a number of advantages compared to synthetic fertilizers, namely, they are produced from sustainable resources, improve soil properties, and store carbon. However, to be able to replace synthetic fertilizers in conventional farming systems, it is essential for biochar-based fertilizers to have a predictable and consistent release of nutrients. In this study, potassium acetate addition in low concentrations (5%) before pyrolysis increased the water-extractable P content in biochar by several orders of magnitude over the undoped biochar. In addition, it significantly increased the 2% formic acid and 0.01 M CaCl₂ extractable P and the P extracted after six water extraction steps. The results from this study open up a new, simple way to modify and, hence, control the P availability in biochar using a low-cost K additive, yielding a new class of PK fertilizer. Our new multifaceted approach links waste management, soil management, and food production in a virtuous cycle fully aligned with the circular economy and zero-waste concepts.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acssuschemeng.0c04336>.

Results for repeated water extraction for P and K, XRD spectra of the biochars and tables showing results from P K-edge XANES and elemental analyses of the biochars (PDF)

AUTHOR INFORMATION

Corresponding Author

Wolfram Buss – UK Biochar Research Centre, School of GeoSciences, University of Edinburgh, EH9 3FF Edinburgh, U.K.; Research School of Biology, Australian National University, 2601 Canberra, Australia; Conversion Technologies of Biobased Resources, University of Hohenheim, 70599 Stuttgart, Germany; orcid.org/0000-0002-9653-0895; Email: wolfram.buss@anu.edu.au

Authors

Anna Bogush – Centre for Agroecology, Water and Resilience, Coventry University, CV8 3LG Coventry, U.K.

Konstantin Ignatyev – Diamond Light Source, Diamond House, Harwell Science and Innovation Campus, OX11 0DE Didcot, U.K.

Ondřej Mašek – UK Biochar Research Centre, School of GeoSciences, University of Edinburgh, EH9 3FF Edinburgh, U.K.; orcid.org/0000-0003-0713-766X

Complete contact information is available at: <https://pubs.acs.org/doi/10.1021/acssuschemeng.0c04336>

Notes

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■ ABBREVIATIONS

FC, fixed carbon; PTE, potentially toxic element; LCF, linear combination fitting

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