Simultaneous amorphous silica and phosphorus recovery from rice husk poultry litter ash†

Laura Fiameni, Ahmad Assi, Ario Fahimi, Bruno Valentim, Karen Moreira, Georgeta Predeanu, Valeria Slăvescu, Bogdan Ş. Vasile, Adrian I. Nicoara, Laura Borgese, Gaia Boniardi, Andrea Turolla, Roberto Canziani and Elza Bontempi

The livestock sector is one of the most important sectors of the agricultural economy due to an increase in the demand for animal protein. This increase generates serious waste disposal concerns and has negative environmental consequences. Furthermore, the food production chain needs phosphorus (P), which is listed as a critical raw material due to its high demand and limited availability in Europe. Manure contains large amounts of P and other elements that may be recycled, in the frame of circular economy and “zero waste” principles, and reused as a by-product for fertilizer production and other applications. This paper focuses on the extraction and recovery of amorphous silica from rice husk poultry litter ash. Two different extraction procedures are proposed and compared, and the obtained silica is characterized. This work shows that amorphous silica can be recovered as an almost pure material rendering the residual ash free of P. It also addresses the possibility of more specific phosphorus extraction procedures via acid leaching.

1. Introduction

Biomass waste is gaining support as a possible alternative energy source. In the Circular Economy Action Plan adopted on 11 March 2020, the European Union (EU) Commission promotes access to alternative feedstock sources such as sustainable biomass. The aim is to reduce Greenhouse Gas (GHG) emissions and landfilled wastes such as animal manure. In the future, biomass will play an important role in renewable and sustainable energy development. However, the increase in biomass consumption must correspond with suitable possibilities to recover waste deriving from them.

In the last few decades, several recycled materials from livestock farming were considered and proposed for reuse, mainly as fertilizers. In particular, the reuse of poultry litter (PL) shows several advantages since (a) compared to other by-products, it has a high nitrogen and phosphorus (P) content, (b) compared to other wastes it is relatively dry and consequently more adequate for energy recovery and (c) it is easier to transport. Different treatments are proposed to reuse PL, starting from composting and anaerobic digestion to direct combustion. A recently published paper shows that (among the proposed management strategies) Life Cycle Assessment (LCA) demonstrates that the thermal conversion of PL is the most environmentally friendly method and has a low environmental impact. Thermal treatment also has the advantage to destroy pathogen agents and organic pollutants that may be present in PL.

The waste from thermal treatments, poultry litter ash (PLA), is rich in P and has a high recycling potential. This makes PLA extremely interesting as a source for P recovery. Indeed, PLA has been proposed by various researchers as an alternative phosphorus source.

P used in fertilizers and animal feed is currently obtained from phosphate rocks, which are finite and located in only a few places on Earth. Therefore, the recovery and recycling of this critical raw material should be taken into consideration to have sustainable management of P. The main P-rich waste streams are sewage sludge, animal manure, slaughterhouse waste (e.g. meat and bone meal), and food waste. The mentioned waste materials are well described in Ohtake and Tsuneda (2019) and presents the latest research advances, innovations, and applications of P recovery and recycling.
It is also important to highlight that the depletion of natural resources requires the continuous improvement of recovery strategies, with an emphasis on the recovery of all the waste components, even if derived from incineration processes. The aim was not only to replace critical raw materials (such as P) with valorised wastes (i.e. secondary materials) but also with the ambition of simultaneously recovering and recycling the waste derived from the critical raw material extraction to subsequently be used in other applications. In this context, creative approaches are mandatory as a driving force not only to find suitable and sustainable ways to proceed in the recovery of critical raw materials but also to propose new and high-value applications for the recovered materials that are defined eco-materials.

Eco-materials enhance environmental improvement and respect the sustainability pillars while maintaining accountable performances.

This work can be classified as a study that addresses possible ways to manage and enhance specific biomass ash, namely rice husk poultry litter ash (RHPLA). As we have previously studied, this waste is mainly composed of K, P, Ca, S, and Si, where the amount of amorphous component is higher than 60%. Obviously, in such a complex waste, containing several phases and different elements (also in low quantities), it is hard to define the exact amorphous composition. But we highlight that literature reports that the combustion of rice husk (RH), a constituent of RHPLA, produces essentially amorphous silica. Silica recovered from RH has gained attention over the years due to its various applications and the virtuous example of the circular economy that it represents. Fillers, additives, desiccants, adsorbents, catalysts, mesoporous materials, nanocomposites, ceramics, biomedical sector, and heavy metal stabilization of incineration waste, are some of the fields in which value-added silica from RH could found an application.

On this basis, the objective of this work is to propose an eco-material extraction (amorphous silica), starting from RHPLA, that can be made in combination with P recovery. RHPLA, and more in general PLA, is a type of waste that obtained visibility for reuse in a very recent period, and for this reason, literature and studies are minimal, especially about characterisation and recovery. Table 1 shows the P and Si content in different biomass ash reported in the literature. To the best of authors’ knowledge, this is the first paper that presents the opportunity of a combined recovery strategy of amorphous silica and P for RHPLA.

2. Experimental

2.1. Materials and methods

2.1.1. Sampling. RHPLA was sampled at “Campoaves, S.A.” (Figueira da Foz, Portugal) from an incineration plant designed for the combustion of PL. The combustion unit consists of a chain grate stoker with an over-fire airflow which regulates the temperature for combustion at ~1000 °C (the temperature mostly depends on the amount of moisture in the environment). The fuel is composed of rice husk poultry litter (the rice husk is provided from rice fields which are located in the surrounding area close to the plant) and wood chips on a 70 : 30 ratio. The generated ash consists of bottom ash (BA) (that accumulates on the grate, falls into the BA wet tank and is deposited in an open-air storage area close to the plant), economizer fly ash (ECO), and fly ash captured by a multicyclone system (MCYC).

The conditions of the boiler are variable and change from month to month. On average, the plant generates 156 tons of ash per month. The quantity of ash samples correlates with the specific conditions implemented on the sampling day. After 5 h, the samples were collected in thick plastic bags. Approximately 327 kg of BA was collected at the water tank exit, 5 kg of ECO was collected from a duct connected to the economizer with an exit point adjoining the pit where the BA sample was collected, and 10 kg of MCYC was collected from the multicyclone hopper.

The samples were dried for 24 h in aluminium trays inside an oven operating at 50 °C (as described by Fahimi et al. (2020) and Andó (2020)). After drying, the samples were manually divided into representative fractions through the coning method and then further divided into representative 100 g sub-samples using a riffle splitter.

The ashes were then sent to Chemistry for Technologies Laboratory (Univ. of Brescia), where they were grounded, and finally sieved to obtain four different fractions: <300 μm, 300–500 μm, 500–1400 μm, >1400 μm. The sieving was manually carried out with a set of ASTM International standard sieves: 50 mesh (300 μm), 35 mesh (500 μm), and 14 mesh (1400 μm).

| Table 1 | Phosphorus (P₂O₅) and silicon (SiO₂) content in biomass ash. * Mean value, ** data converted with conversion factors P/P₂O₅ = 0.436, Si/SiO₂ = 0.468 |
|---|---|---|---|---|
| | P₂O₅% | Bottom ash | SiO₂% | Bottom ash |
| Biomass ash | Fly ash | Fly ash | Bottom ash | Bottom ash | Ref. |
| | — | 12.16** | — | — | 29 |
| | — | 22.00*, ** | — | — | 5 |
| | — | 19.38** | — | — | 9 |
| Animal by-products (meat bone meal ash) | 22.71 | — | — | — | 30 |
| | 22.45**, | 25.96*, ** | — | — | 5 |
| Pig manure ash | 32.50 | — | — | 1.80 | 30 |
| Wood ash | 0.69 ** | 3.21 ** | 7.69 ** | 12.39 ** | 31 |
2.1.2. Characterisation techniques

*X-ray diffraction (XRD) analysis.* For the structural characterisation of the raw ashes and the amorphous silica fraction obtained from the raw ashes, X-ray diffraction (XRD) analysis was conducted. XRD measurements were performed by a PANalytical X’Pert PRO diffractometer (PANalytical, Malvern, UK), equipped with a Cu Kα anode and operating at a voltage of 40 kV and a current of 40 mA. For the phase identification, Philips X’Pert software was used (associated with the crystallography open database (COD)) as described by Assi et al. (2019).14

*Scanning electron microscopy combined with energy-dispersive X-ray spectrometry (SEM-EDXS).* Morphological and dimensional characterization of the ash and silica powder samples were performed through scanning electron microscopy (SEM) (LEO EVO 40, Carl Zeiss AG, Milan, Italy), coupled with an EDXS (energy dispersive X-ray spectroscopy, Oxford instruments, Wiesbaden, Germany) probe for elemental analysis and semi-quantitative chemical characterization. The material topography was analysed using the secondary electron (SE) mode.

*Total reflection X-ray fluorescence (TXRF) analysis.* The TXRF technique was used for elemental analysis in a solution. A stock solution of 1 g L\(^{-1}\) Ga in nitric acid (Ga-inductively coupled plasma (ICP) Standard Solution, Fluka, Sigma Aldrich) was used as the internal standard. Approximately 0.010 g of 100 mg L\(^{-1}\) of Ga solution prepared from the stock solution was added to the samples and properly diluted with MQ water to obtain a final concentration of 1 mg L\(^{-1}\) Ga in 1 mL specimens. The specimens were then homogenized using a vortex shaker for 1 min at 2500 rpm. For each specimen of the digested and leachate solution, a 10 μL drop was deposited on three different plexiglass reflectors and dried on a hot plate at 50 °C under a laminar hood. Each reflector was irradiated for 600 s of live time using an S2 Picofox system (Bruker AXS Microanalysis GmbH, Berlin, Germany) equipped with a Mo tube (operating at 50 kV and 750 mA) and a silicon drift detector (SDD). An instrumental software with a routine deconvolution based on mono-element profiles was applied to evaluate the peak areas and analyse the TXRF spectra. Since lighter elements such as P and S can be underestimated by the TXRF analysis under the reported experimental conditions, a dedicated calibration curve was developed for S and P to correct for matrix effects following the approach reported by Borgese et al. (2018).35

Element concentrations present in the acid leaching solution were converted into extracted mass using the formula:

\[
\text{Extracted mass (mg kg}^{-1}\text{)} = (C \times DF \times V) \times m^{-1}
\]

in which: \(C\) = concentration of the analyte by TXRF analysis (mg L\(^{-1}\)); \(DF\) = dilution factor; \(V\) = volume of extractant (mL); \(m\) = mass of ash weighed for leaching (g)

*X-ray fluorescence (XRF) analysis.* For qualitative and quantitative analysis of the elements, an X-ray Sequential Fluorescence Spectrometer Thermo Scientific ARL PERFORM’X equipped with an X-ray tube with Rh anode and Be window of 30 μm was used. The entire surface of the sample was analysed in a dry He flow.

**Phosphorus quantification via UV-Vis chemical method.** The method was developed following the Romanian standardized method.36 RHPLA raw samples were prepared at a grain size of less than 0.2 mm and the amount was 0.1 g with a 0.0001 g measured accuracy. Then, RHPLA samples were solubilized with a mixture of concentrated acids 2 : 1 H\(_2\)SO\(_4\) : HNO\(_3\). To prevent the colorimetry operation, the silicic acid that produces the opaque solution was removed by filtration. The analysis was carried out by adding ammonium molybdate and H\(_2\)SO\(_4\) 6 N solutions. The P concentration in the filtrate was encapsulated in phosphomolybdate [(NH\(_4\)\(_6\))Mo\(_7\)O\(_{24}\)-4H\(_2\)O] and the absorbance of the resulting blue solution was measured at 700 nm wavelength by a suitable optical instrument. The measurements were done in 2 cm thick cuvettes. As a reference, a blank solution of the reagents, without the ash, passed through all the analytical steps was used. The spectrophotometer was set to perform three readings and the average was taken. On the calibration curve with a 0.993 regression coefficient, the P content corresponding to the extinction obtained on the spectrophotometer was read.

The preparation of the reagents was carried out just before the analysis, performed with a JASCO V570 UV-VIS spectrophotometer at UPB - CPMTE laboratory. The P content in ash was calculated using the formula:

\[
\% \text{Phosphorus (P)} = e \times m^{-1} \times 100
\]

in which: \(e\) = P content, mg [read on the calibration curve], \(m\) = mass of the ash sample in the 50 mL colorimetric solution, mg.

2.2. Silica and phosphorus recovery

2.2.1. Procedure A. Procedure A was conducted using the method proposed by Bosio et al. (2014)37 and Kamath et al. (1998).38 50 g of each ash sample was placed in a beaker and 416 mL of 1 mol L\(^{-1}\) NaOH solution was added. Constant stirring was maintained at 100 °C for 1 h. The solutions from samples ECO and MCYC containing sodium silicate [reaction 1], still warm, were filtered under slight suction through Whatman filter paper grade 589/2 white ribbon filter ashless (pore size 4–12 μm). However, the filtration of the BA sample was slow and difficult, and it was first filtered through a Whatman filter paper grade 589/1 black ribbon ashless (pore size 12–25 μm) before passing to the finer porosity. Sedimentation, which led to phase separation, was useful for better percolation. The BA sample leachate was clear but dark brown in colour, while the other solutions were clear and colourless.

\[
\text{SiO}_2(s) + 2 \text{NaOH(aq)} \rightarrow \text{Na}_2\text{SiO}_3(aq) + \text{H}_2\text{O(l)} \quad (\text{reaction 1})
\]

After filtration, 60 mL of 1 mol L\(^{-1}\) H\(_2\)SO\(_4\) solution was slowly added, at room temperature and constant stirring, to each 200 mL of leachate. Chemical conditions of the reaction governed the kinetic, the mechanism of hydrolysis, and the condensation processes.39 Therefore, the gel formation was controlled by pH measurement to reach the best endpoint of the gelling reaction. The acid solution was further added dropwise until a pH of 9.8 was reached, and then the titration was stopped. A volume...
ranging between 20 and 40 mL, depending on the ash leachate, was added after the first \( \text{H}_2\text{SO}_4 \) addition. The formation of silica gel from the sodium silicate solution was performed in accordance with [reaction 2].

\[
\text{Na}_2\text{SiO}_3(\text{aq}) + \text{H}_2\text{SO}_4(\text{aq}) \rightarrow \text{SiO}_2(\text{gel}) + \text{Na}_2\text{SO}_4(\text{aq}) + \text{H}_2\text{O}(l)
\]  

(reaction 2)

After 7 days of aging, the gels were completely formed, however, they were still mixed with reaction water. A centrifuge (Eppendorf, Centrifuge 5804 R) was used to separate the gel from the solution. The mixture was transferred to a 50 mL centrifuge tube and centrifuged at 1000 rcf for 10 min at 20 °C. The supernatant was separated and stored in a polyethylene tube, while the gel was dried with an electric oven at 120 °C for 7–10 h (halogen convection furnace Bonny 2172339, Kooper, Rome, Italy). The final BA silica gel was brown, while the other two gels were whitish, even after drying. For subsequent XRD analysis, the gels were ground using a mixer mill MM 400 (Retsch, Haan, Germany).

2.2.2. Procedure B. The second procedure is based on work from Santana Costa et al. (2018).

30 g of each ash was placed in a beaker with a 1 mol L\(^{-1}\) HCl solution at a solid/liquid (S/L) ratio of 1 : 10 and constantly stirred at room temperature for 2 h. After this period, the acid leachates of ECO and MCYC were filtered under slight suction through Whatman filter paper grade 589/2 (pore size 4–12 μm) and stored in polyethylene tubes for further TXRF analysis. The solid residues were washed with MQ water until a constant pH was reached, dried, and then the washed water was stored separately in polyethylene tubes. It was not possible to filter the BA acid suspension due to its fine porosity and the sample was instead centrifuged at 20 °C for 20 min at 1000 rcf with a 50 mL centrifuge tube (Eppendorf, Centrifuge 5804 R). The supernatant was separated and stored in a polyethylene tube for further TXRF analysis. The solid residue was washed with MQ water until a constant pH was reached, dried, and then, by using the procedure described previously, separated from the wash waters and stored in polyethylene tubes.

All the solid samples after acid pre-treatment were dried in an electric oven at 120 °C for 9 h (halogen convection furnace Bonny 2172339, Kooper, Rome, Italy). A weight loss of 64.1% for BA and 72.8% for MCYC was recorded.

To conform with [reaction 1], these samples were reacted with a 4 mol L\(^{-1}\) NaOH solution, in an S/L ratio 1 : 10, under constant stirring at 80 °C for 4 h. At the end of the reaction, the mix was allowed to cool and settle at room temperature for 24 h.

The separation of sodium silicate solutions from the solid residues was performed under slight suction through Whatman filter paper grade 589/2 and 589/1 (pore size 4–12 μm and 12–25 μm). The solid residues were then washed with MQ water until a constant pH was reached, dried with an electric oven (halogen convection furnace Bonny 2172339, Kooper, Rome, Italy), and stored in plastic bags. The washing waters were stored in polyethylene tubes. During the procedure, a volume loss for the sodium silicate solutions was recorded: 25% for ECO and MCYC and 40% for BA.

To comply with [reaction 2], a 5 mol L\(^{-1}\) \( \text{H}_2\text{SO}_4 \) solution was added dropwise to each 50 mL of sodium silicate solution, under constant stirring at room temperature, in order to lower the pH to 9.8 and to form the gel. After the period of aging, the silica gel separation from the reaction water was carried out in the same way as described in procedure A. A Neya centrifuge model 16-R was used, operated at 5000 rpm for 20 min at 25 °C. Subsequent drying in the oven lasted 5–6 h. The BA silica gel was brown, while the ECO and MCYC gels were whitish. For subsequent XRD analysis, the gels were ground with a mixer mill MM 400 (Retsch, Haan, Germany).

3. Results and discussion

3.1. Rice husk poultry litter ash (RHPLA) characterisation

Fig. 1 shows the XRD patterns of the raw RHPLA (BA, ECO, and MCYC) samples compared to the same samples after the acid pre-treatment step of procedure B. As shown in Fig. 1A (spectrum BA-A), BA contains quartz [\( \text{SiO}_2 \)], sylvite [\( \text{KCl} \)], and arcanite [\( \text{K}_2\text{SO}_4 \)] as identified by the main peaks at 26.62°, 28.31° and 30.77° in 2θ (2 theta) respectively. All the other crystalline phases are phosphate compounds of Na, Ca, K, and Mg namely sodium hydrogen phosphate [\( \text{NaH}_2\text{PO}_4 \)], sodium-calcium phosphate [\( \text{Na}_2\text{CaP}_2\text{O}_7 \)], calcium phosphate [\( \text{Ca}_3(\text{PO}_4)_2 \)], sodium hydrogen phosphate hydrate [\( \text{NaH}_2\text{PO}_4(\text{H}_2\text{O}) \)], and potassium magnesium orthophosphate hydrate [\( \text{K}_2\text{MgPO}_4(\text{OH})_2 \)].

The presence of a broad band in all the XRD patterns BA-B (Fig. 1A), ECO-B (Fig. 1B), and MCYC-B (Fig. 1C) between 15 and 35° in 2θ, demonstrates that an amorphous phase is also present in these samples.

The broad X-ray diffraction halo in the XRD pattern is typical for amorphous compounds. Consequently, even if this type of bottom ash (BA) was characterized for the first time by the XRD technique in this paper, the presence of an amorphous phase can be assumed seeing as it has already been reported for fly ash with the same origin by Fahimi et al. (2020). This amorphous component in all the samples may consist of both silicon oxide, widely found and studied for ash originating from rice husk combustion, and P phases, as confirmed by the Raman and XRD analyses in ECO and MCYC samples made by Fahimi et al. (2020) also per Stammeyer et al. (2018).

The spectrum BA-B in Fig. 1A shows that after the acid leaching, the initial complex matrix loses its soluble components, maintaining only two crystalline phases: quartz [\( \text{SiO}_2 \)] and sylvite [\( \text{KCl} \)]. The compound KCl is present in a different crystalline phase from the raw BA sample seeing as it shows the main peak at 34.88° in 2θ which was previously absent. Also, the differences between the crystalline phases in BA-A and BA-B (Fig. 1A) suggest that phosphate soluble compounds are leached by HCl. The same can be assumed for the ECO and MCYC samples.

Spectrum ECO-A (Fig. 1B) and spectrum MCYC-A (Fig. 1C) show that ECO and MCYC, similar to BA, contain quartz [\( \text{SiO}_2 \)], sylvite [\( \text{KCl} \)], and potassium sulphate (or arcanite) [\( \text{K}_2\text{SO}_4 \)] as identified by the same main peaks found in BA-A [26.6°, 28.3° and 30.8° in 2θ]. All the raw samples contain the same crystalline phosphate phase: sodium hydrogen phosphate [\( \text{NaH}_2\text{PO}_4 \)]. Sodium calcium

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phosphate \([\text{Na}_2\text{CaP}_2\text{O}_7]\) was observed in MCYC as well as in the BA sample, while hydroxyapatite \([\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2]\) was found only in ECO (Fig. 1B ECO-A). Hydroxyapatite and sylvite were previously found in similar wastes and some of the detected crystalline phases are those previously found by Fahimi et al. (2020). Lastly, Periclase \([\text{MgO}]\) was found in both fly ash samples (spectra ECO-A and MCYC-A in Fig. 1B and C). After the acid pre-treatment, spectra ECO-B and MCYC-B (Fig. 1B and C) show the presence of an amorphous phase and remaining crystalline phases quartz/silicon oxide \([\text{SiO}_2]\) and bassanite \([\text{Ca}_2(\text{SO}_4)_2\text{H}_2\text{O}]\). A residual amount of potassium chloride \([\text{KCl}]\) was detected for MCYC-B (Fig. 1C) in a phase very similar to that found in BA-B (Fig. 1A). Similar to BA, after the acid treatment, phosphatic compounds are absent in ECO and MCYC.

The crystalline phase identification is corresponding to the XRF data reported in Table 2 for each raw RHPLA and its corresponding ash after acid leaching with procedure B.

The raw ashes formed from the combustion of three main fuels: rice husk, poultry droppings, and wood chips that have high concentrations of silicon \([\text{SiO}_2]\), phosphorus \([\text{P}_2\text{O}_5]\), and calcium \([\text{CaO}]\) respectively. However, as shown in Table 2, the main constituent of these samples is potassium \([\text{K}_2\text{O}]\). Additionally, \(\text{SO}_3\), \(\text{MgO}\), \(\text{Na}_2\text{O}\), and \(\text{Cl}\) are present in lower concentrations. Other metals, such as \(\text{Al}_2\text{O}_3\), \(\text{MnO}\), \(\text{Fe}_2\text{O}_3\), \(\text{NiO}\), \(\text{CuO}\), and \(\text{ZnO}\) are present in negligible amounts. These results confirm the importance of recovering both P and silicon \([\text{Si}]\) from this waste typology.

XRF analysis of ash samples after acid pre-treatment reported in Table 2, also corroborates the previous hypothesis that P-compounds are leached by HCl, while Si remains in the

Fig. 1 XRD patterns obtained for the samples BA (A), ECO (B) and MCYC (C): raw RHPLA in black (BA-A, ECO-A, MCYC-A) and raw RHPLA after the acid leaching with HCl 1 mol L\(^{-1}\) for procedure B in grey (BA-B, ECO-B, MCYC-B).
matrix and is purified for subsequent recovery. As shown in Table 2, during the leaching process, BA lost P2O5, K2O and CaO, while maintaining SiO2, SO3, and Cl. In ECO evident leaching of K2O, P2O5, and other minor components occurred. However, SiO2, SO3, and CaO remain in high percentages in the sample. For MCYC, the P-leaching was not so effective, and the amounts of P2O5, SiO2, and SO3 in the RHPLA residues remain quite high (P2O5 at 21.41%). The leaching of K2O occurred. According to the results expressed as w/w% it can be observed that the major content of SiO2 after the acid pre-treatment is detected in BA. After acid leaching with HCl 1 mol L−1 as described in procedure B, BA seems more purified compared to ECO and MCYC because most of the components are contained in minor concentrations. However, the BA complex matrix, rich in organic matter and hydrocarbon compounds, is not the best starting material for silica recovery, due to its composition and dark colour, as seen during the experimental.

The quantification of P by XRF and chemical method is reported (for comparison) in Fig. 2. Also, Fig. S.1 (ESI†) reports the description of the UV-Vis method that was adapted for phosphorus quantification: it was considered suitable for RHPLA since it is a material with high P content and the method is based on colorimetric tests to obtain the extinction values on the calibration curve. XRF results reveal, in the three raw RHPLA, a P content between 4.9 and 7.1%, while UV-Vis data between 3.9 and 7.9%.

In Fig. 3 an image for each RHPLA sample (scale bar 2 mm), BA (A), ECO (B), MCYC (C), is provided. For points P.1 (Fig. 3B) and P.2 (Fig. 3C) respectively located in the ECO and MCYC samples, a higher magnification analysis was carried out as reported in Fig. 3D and E. The chemical composition measured with EDXS of points P.1, P.2, P.3, P.4 is shown in the spectra below. Phosphospheres were identified in ECO (Fig. 3B P.1) and MCYC (Fig. 3C P.2) and are distinguishable due to their (spherical) shape and semi-smooth surface which was covered with micrometric particles of other elements. Although the phosphosphere shown in Fig. 3E is very “clean” compared to the phosphosphere in Fig. 3D, their respective compositions reported in spectra P.1 and P.2 are relatively similar.

The main constituent element for these two points, besides O and K, is P. Other spheres like these were detected during the analysis and, as can be seen from Fig. 3, their dimensions vary from less than 100 to 300 μm. Similar to findings made by Fahimi et al. (2020), the two fly ash samples (ECO and MCYC) consist of micrometric and supermicrometric (<10 μm and >10 μm respectively) P-rich morphotypes. Valentim et al. (2016) reported that Si-, Ca-, K- and Mg-phosphospheres (P-rich glassy spheres) are a type of amorphous material that can frequently be observed in biomass fly ash. Points P.3 and P.4 (Fig. 3D and spectra P.3 and P.4) show high concentrations of Si and O, and their surfaces are covered mainly with K but also with Na, Mg, P, S, Cl, and Ca. The presence of an amorphous P-phase (P.1 and P.2) and an amorphous Si-phase (P.3 and P.4) are reconfirmed by the SEM-EDXS analysis.

Table 2 XRF chemical composition of raw RHPLA and raw RHPLA after acid leaching with HCl 1 mol L−1 in procedure B. nd: not detected. Others contains: TiO2, Cr2O3, Rb2O, BaO, SrO, Br, V2O5 and SeO2.

<table>
<thead>
<tr>
<th></th>
<th>BA Raw RHPLA</th>
<th>Leaching HCl 1 mol L−1</th>
<th>ECO Raw RHPLA</th>
<th>Leaching HCl 1 mol L−1</th>
<th>MCYC Raw RHPLA</th>
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<td>0.97</td>
<td>3.93</td>
</tr>
<tr>
<td>NiO</td>
<td>0.01</td>
<td>0.02</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
</tr>
<tr>
<td>CuO</td>
<td>0.11</td>
<td>0.49</td>
<td>0.07</td>
<td>0.26</td>
<td>0.08</td>
<td>0.24</td>
</tr>
<tr>
<td>ZnO</td>
<td>0.30</td>
<td>0.25</td>
<td>0.43</td>
<td>0.13</td>
<td>0.36</td>
<td>0.42</td>
</tr>
<tr>
<td>Others</td>
<td>0.17</td>
<td>0.53</td>
<td>0.09</td>
<td>0.27</td>
<td>0.11</td>
<td>0.38</td>
</tr>
</tbody>
</table>

Fig. 2 Comparison of XRF and UV-Vis results for P quantification in raw RHPLA.
3.2. Silica characterisation

As previously mentioned, the XRF analysis in Table 2 shows a significant reduction or total leaching of P₂O₅ and alkali metals, while SiO₂ remains in the matrix. The acid pre-treatment of procedure B therefore allows the lowering of salt contents in the final dried silica, as shown by the XRD analysis in Fig. 4. Fig. 4A shows that, for silica recovered with procedure A, thernadite and sodium sulphate [Na₂SO₄] with potassium sodium sulphate phases were identified. It should be noted that there are several crystalline phases for procedure A, while for procedure B (Fig. 4B), the final product is contaminated only by thernadite and sodium sulphate with the same chemical nature.
The presence of these compounds is justified by [reaction 2] where Na₂SO₄ is a reaction product together with SiO₂. During the aging of the gel, a certain amount of Na₂SO₄, soluble in the reaction solution, may be trapped in the hygroscopic silica network, and then recrystallized during drying.

Moreover, from the XRD analysis of recovered silica from procedure B (Fig. 4B), an amorphous phase is identified from the broad halo present in all the XRD patterns from about 15 to 35° in 2 theta. These assessments suggest that procedure B was more effective and data in Table 3 validate this hypothesis. XRF analysis of the six samples of recovered amorphous silica, results provided in Table 3, indicates that the SiO₂ content in the gels from procedure A was very low compared to procedure B. Na₂O, SO₃ and K₂O contaminate amorphous silica A much more than the amorphous silica B. The absence of an acid pre-treatment in procedure A leads to P₂O₅ still being present in the final amorphous recovered product, wasting P and contaminating the silica. On the other hand, SiO₂, Na₂O and SO₃ are the significant components in amorphous silica B, and the only crystalline phase detected by XRD, namely thenardite/sodium sulphate [Na₂SO₄] (Fig. 4B), justifies the presence of sodium and sulphur in the XRF analysis of these samples. As reported in Table 3, ECO B has the highest silica content at 80.30%. Nevertheless, all the samples produced with procedure B show higher values of SiO₂% due to the low presence of crystalline phases compared to the silica from procedure A that contains a lot of salts. This means that procedure B allows the obtainment of an almost pure final product in three steps: acid leaching pre-treatment, [reaction 1], and [reaction 2].

### 3.3 Acid leachate characterisation

In procedure B for silica extraction, a pre-treatment with 1 mol L⁻¹ HCl solution was used, as proposed by Alvarez et al. (2015), to obtain better results in terms of purity of the extracted silica. As is well known for other different starting materials, such as sewage sludge ash (SSA), the most common method for P-recovery is acid leaching. The choice to use HCl as an extractant in this work was done in accord with the reference article on which the procedure B was based, following literature about SSA: Hong et al. (2005), Biswas et al. (2009), Xu et al. (2012), and the recent Semerci et al. (2020) that demonstrates the effective recovery of P with HCl.

A recent paper shows that wet chemical leaching approaches appear to be the most sustainable method for P extraction from SSA and the current research proceeds in this direction.

The main advantage of the proposed method is that the RHPLA contains low quantities of heavy metals compared to SSA. In particular, great attention to fertilizer regulations (the first area of interest for the application of recovered P) must be given regarding maximum concentrations of As, Cd, Hg, Cr(VI),...
Ni, Cu, Pb, and Zn. Regulation (EC) No 2003/2003 of the European Parliament and of the Council relating to fertilizers will be repealed with effect from 16 July 2022 by regulation (EU) 2019/1009 laying down rules on the market availability of EU fertilizing products. The limits of contaminants in fertilizers and soil improvers are reported in Table S.2 (ESI†) and compared with Fig. 5. Fig. 5 highlights the concentration of the elements of interest for the discussion, in the acid leaching solution, derived from TXRF results. The complete TXRF elemental analysis is reported in Table S.3 (ESI†). P is detected between 57 and 75 g kg\(^{-1}\), or 5.7 and 7.5%, in the three RHPLA leachate samples. Arsenic (As), cadmium (Cd), and mercury (Hg) were not detected in the samples. Hexavalent chromium (Cr\(\text{VI}\)) is fixed at 2 mg kg\(^{-1}\), but TXRF analysis provides an elemental analysis that cannot discriminate between elemental speciation. To set the precise value of hexavalent chromium in the samples, different analytical techniques are required. In any case, the highest concentration of total chromium is recorded for BA about 46 mg kg\(^{-1}\). Nickel (Ni) must respect the limit of 50 or 100 mg kg\(^{-1}\), and in MCYC it is present at more or less 44 mg kg\(^{-1}\), with a limit fixed at 300 or 600 mg kg\(^{-1}\). The same for lead (Pb), which has a limit of 120 mg kg\(^{-1}\), and it is present in BA at more or less 5 mg kg\(^{-1}\). Contrary, zinc (Zn) shows the following concentrations: about 0.7 g kg\(^{-1}\) for BA, about 2.5 g kg\(^{-1}\) for ECO, and about 1.9 g kg\(^{-1}\) for MCYC, while the limit is fixed at 0.8 or 1.5 g kg\(^{-1}\). In this specific case, the limit is only respected by BA; the use of P recovered from RHPLA to prepare fertilizers or soil improvers, through the precipitation of a solid product from the RHPLA acid leachate, shall be developed to keep this contaminant within the legal limit. So, the leaching with an acid that has a high P extraction capacity, which is not preferred in literature for SSA due to metal leaching, may be a good solution for P-recovery from RHPLA for fertilizer applications, because the main limitation is only provided by Zn concentration in the solution. The P thus extracted may undergo the same treatments already studied for SSA, and recently proposed for poultry manure, to obtain

<table>
<thead>
<tr>
<th>Composition</th>
<th>Procedure A</th>
<th>Procedure B</th>
</tr>
</thead>
<tbody>
<tr>
<td>SiO(_2)</td>
<td>23.38</td>
<td>77.48</td>
</tr>
<tr>
<td>Na(_2)O</td>
<td>22.99</td>
<td>9.29</td>
</tr>
<tr>
<td>SO(_3)</td>
<td>35.35</td>
<td>10.67</td>
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<tr>
<td>P(_2)O(_5)</td>
<td>3.68</td>
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<tr>
<td>K(_2)O</td>
<td>11.08</td>
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<tr>
<td>Cl</td>
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<tr>
<td>CaO</td>
<td>0.59</td>
<td>0.09</td>
</tr>
<tr>
<td>MgO</td>
<td>0.54</td>
<td>0.13</td>
</tr>
<tr>
<td>Al(_2)O(_3)</td>
<td>0.52</td>
<td>0.42</td>
</tr>
<tr>
<td>Others</td>
<td>0.13</td>
<td>0.47</td>
</tr>
</tbody>
</table>

Table 3 XRF chemical composition of recovered amorphous silica using procedure A and B. nd: not detected. Others contains: Fe\(_2\)O\(_3\), ZnO, CuO, MnO and BaO

Fig. 5 TXRF analysis of acid leachate with HCl 1 mol L\(^{-1}\) for the three RHPLA under examination: BA in orange, ECO in green and MCYC in blue. Phosphorus and contaminants of interest for the preparation of fertilizers are reported. Concentration values can be compared with the limit values in Table S.2 (ESI†). Other elements concentrations for TXRF analysis are reported in Table S.3 (ESI†).
a novel product and a novel eco-material for the fertilizer industry. It is evident that the proposed procedure is suitable to promote P separation before proceeding with silica recovery and the two stages can be studied with a mutual approach as already achieved for SSA. This first attempt of double recovery (P and SiO2) from raw RHPLA could also present the opportunity for the reuse of RHPLA residual. The alkali leaching step with NaOH in the silica recovery procedure could leave a certain amount of Si-amorphous phase in the matrix: this may suggest a more in-depth study of extraction efficiency or a new study for further application of the RHPLA residual in Portland cement-based materials as already proposed for incinerated sewage sludge ash (ISSA).

4. Conclusions

To the authors’ knowledge, this is the first work that proposes a procedure to recover P and silica simultaneously from RHPLA.

Firstly, XRD and SEM-EDXS characterization analyses of poultry litter bottom ash (BA) were made. The results show that BA is composed of different crystalline phosphate compounds namely: sodium-calcium phosphate [Na2CaP2O7], calcium phosphate [Ca3(PO4)2], sodium hydrogen phosphate hydrate [NaH2PO4·H2O], and potassium magnesium orthophosphate hydrate [K(MgPO4·6H2O)]. In all the ash samples (BA, ECO and MCYC), together with quartz, syilte, and arcancite, phosphate is present as sodium hydrogen phosphate [NaH2PO4], while Hydroxyapatite emerges only in the ECO sample. Amorphous phases of both Si and P are present in ECO and MCYC and phosphospheres were identified. The XRD analysis clearly shows these amorphous phases are purified from the solubilized forms after procedure B acid leaching. EDXS analysis confirms that P, Na, Mg, S, K, and Ca are leached by the acid, while Si remains in the solid matrix. Therefore, in procedure B, Si and P were extracted separately and acid leaching was essential to obtain purified dried silica. The best result was obtained for sample ECO (about 80% SiO2 grade). It can be concluded that procedure B can be a combined process to recover firstly P and then silica in two subsequent steps. Moreover, considering this process as a flow that produces, respectively, acid leachate containing P, RHPLA residues, and amorphous silica, we can further report that: evidence that the acid leachate contains a very small quantity of contaminants (As, Cd, Hg, Ni, and Pb) compared to the sewage sludge ash (SSA) leachate, allows the conclusion that RHPLA is a good secondary source of P that can be precipitated to obtain an eco-material candidate for industrial fertilizer production such as struvite, with particular attention to Zn concentration. RHPLA residues can be used as a binder in porous materials to improve mechanical performance or can be studied for further application in the Portland cement industry. And finally, the dried silica can be tested for the stabilization of heavy metals in Municipal Solid Waste Incinerated Fly Ash (MSWI-FA). Further studies are needed to assess whether silica may be obtained together with P from SSA.

In conclusion, optimization of procedure B can be a forthcoming activity to find the best conditions for the acid leaching and the alkali steps to obtain a good yield for both P and silica (DoE).

Author contributions


Conflicts of interest

The authors report that there are no conflicts of interest to declare.

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