

Zinc speciation in organic waste drives its fate in amended soils

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1	Zinc speciation in organic waste drives its fate in						
2	amended soils						
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Recycling of organic waste (OW) as fertilizer on farmland is a widespread practice that fosters 18 19 sustainable development via resource reuse. However, the advantages of OW fertilization should 20 be weighed against the potentially negative environmental impacts due to the presence of 21 contaminants such as zinc (Zn). Current knowledge on the parameters controlling the 22 environmental fate of Zn following OW application on cultivated soils is scant. We addressed this 23 shortcoming by combining soil column experiments and Zn speciation characterization in OWs 24 and amended soils. Soil column experiments were first carried out using two contrasted soils 25 (sandy soil and sandy clay loam) that were amended with sewage sludge or poultry manure and 26 cropped with lettuce. The soil columns were irrigated with identical amounts of water twice a week 27 and the leachates collected at the column oulet were monitored and analyzed. This scheme (OW 28 application and lettuce crop cycle) was repeated for each treatment. Lettuce yields and Zn uptake 29 were assessed at the end of each cycle), The soil columns were dismantled and seven soil layers 30 were sampled and analysed at the end of the second cycle (total experiment time: 12 weeks). XAS 31 analyses were then conducted to assess Zn speciation in OW and OW-amended soils. The results 32 of this study highlighted that: (i) Zn fate in water-soil-plant compartments was similar, regardless 33 the type of soil and OW, (ii) >97.6% of the Zn input from OW accumulated in the soil surface 34 layer, (iii) Zn uptake by lettuce increased with repeated OW applications, and (iv) no radical 35 change in Zn speciation was observed at the end of the 12-week experiment and phosphate was 36 found to drive Zn speciation in both OW and amended soils (i.e. amorphous Zn-phosphate and Zn 37 sorbed on hydoxylapatite). These results suggest that Zn speciation in OW is a key determinant 38 controlling the environmental fate of this element in OW-amended soils.



42 Introduction

43 The rapid increase in the African population (by 1.3 billion people between 2017 and 2050^{1}) combined with the high urbanization rate² has led to an urban and periurban agriculture boom.^{3, 4} 44 45 In developing countries, the market garden farming intensification is being achieved via inputs of biodegradable materials such as sewage sludge, livestock manure, green waste compost, etc., as 46 47 soil fertilizers. These biodegradable materials, referred to as organic wastes (OWs), can supply a part or all of the nutrients needed for crops.^{4, 5} However, many studies have highlighted negative 48 49 impacts of agricultural OW recycling as fertilizers due to the presence of contaminants such as trace elements.^{6, 7} 50

51 A recent assessment of periurban Dakar (Senegal) agrosystems highlighted trace element contamination of OW-amended cropped soils.⁸ Zinc (Zn) accumulation in soil surface layers has 52 been explained by long-term repeated Zn-rich OW application.⁸ Short-term vegetable crop cycles 53 54 led to sewage sludge (SS) or poultry manure (PM) application up to 9 times per year over a 24 to 55 40 year period. The annual Zn input in periurban Dakar agrosystems ranged from 1369 to 5818 g.ha⁻¹year⁻¹, which is alarmingly higher than the 454-539 g.ha⁻¹year⁻¹ range in European countries 56 and China.⁹⁻¹¹ Surprisingly, Zn contamination was similar in the two main periurban Dakar 57 agrosystems (Pikine and Rufisque) despite the fact that the properties of the soils in these two 58 59 regions differ markedly. One is an Arenosol (sandy soil, dune material) and the other a Fluvisol (sandy clay loam and moderately calcareous soil).¹² However, an analytical approach⁸, involving 60 61 a comparison of Zn contents in long-term cultivated plots versus adjacent uncultivated plots, did 62 not enable (i) accurate monitoring of the Zn fate following OW spreading in the three water-soilplant system compartments, (ii) determination of the factors that could explain the similar fate of 63 Zn in contrasted soil types. 64

66 X-ray absorption spectroscopy (XAS) is one of the most widely known structural techniques 67 for direct determination of speciation of trace elements present in complex matrices and it has been successfully applied to describe Zn speciation in different OWs¹³⁻¹⁸ or amended soils¹⁹⁻²². 68 Kirpichtchikova et al.²⁰ and Formentini et al.²¹ studied Zn speciation in soils affected by long-term 69 70 application of sewage water and pig slurry, respectively. In both studies, Zn speciation in soils was 71 dominated by its associations with secondary mineral phases (clay minerals and Fe oxyhydroxides). Zn speciation in OW was assessed by Formentini et al.²¹, revealing that nano-ZnS 72 accounted for 100% of the speciation in the pig slurry. Yamamoto et al.²² recently investigated Zn 73 74 speciation in pig manure compost and in a soil amended with this compost for 23 years. They showed that oxidized species present in the compost, i.e. Zn-phosphate and Zn bound to OM, had 75 76 persisted and accumulated in the amended soil. Although these studies agreed on the fact that Zn accumulated in the soil following OW spreading, their results were not consistent with respect to 77 78 Zn speciation in OWs and in OW-amended soils.

79 The present study combined soil column experiments and Zn speciation analyses to characterize 80 the fate of Zn in periurban Dakar agrosystems. Soil column experiments were first carried out using 81 two contrasted soils that prevail in these agrosystems (i.e. an Arenosol from Pikine and a Fluvisol 82 from Rufisque). Sewage sludge (SS) or poultry manure (PM) were applied at doses consistent with 83 local agricultural practices and the soil columns were cropped with lettuce (Lactuca sativa L). 84 Laboratory studies were conducted to thoroughly control water and solute fluxes and to obtain an 85 accurate measurement of the mass balance of Zn distribution in the water-soil-plant system. XAS 86 analyses were then used to assess Zn speciation in OW and OW-amended soils from the soil column experiments.¹³⁻²² 87

This combined approach enabled us to: (i) monitor Zn fate in the three compartments of the water-soil-plant system, and (ii) identify parameters controlling the fate of Zn following OW application on contrasted types of soil.

91 Materials and Methods

92 Organic waste and soil characteristics

93 Arenosol (Pikine; sandy soil, dune material) and Fluvisol (Rufisque; sandy clay loam and moderately calcareous soil)¹² were selected for this study as they are the main soils in the periurban 94 95 Dakar region, under the prevailing Sahelian tropical climatic conditions. The Arenosol texture was 96 dominated by the sand fraction (96.2–97.1%). Silt and clay accounted for 15.3–15.7% and 24.9– 97 28.8% of the Fluvisol content, respectively (Table 1, SI 1). The water holding capacity, total 98 limestone, pH and organic carbon were much lower in the Arenosol than in the Fluvisol. Quartz, 99 critobalite, sanidine and muscovite mainly accounted for the mineralogy of the Arenosol (SI 2). 100 This contrasted with the Fluvisol mineralogy, in which several clay minerals (kaolinite, 101 montmorillonite, palygorskite and kaolinite), calcite and quartz were detected.

102 Sewage sludge (SS) and poultry manure (PM), are widely used as fertilizers in market gardens 103 in periurban Dakar agrosystems. SS was sampled at the Pikine municipal wastewater treatment 104 plant, and PM was sampled in a henhouse at Rufisque (SI 1). Both SS and PM had been stockpiled 105 for long periods prior to being spread on the soil columns (>3 months) which is consistent with of 106 market gardeners' practices. The ash content of SS (34%) was lower than that for PM (49%), 107 whereas P concentrations were close for SS and PM (17.3 and 16.8 mg.kg⁻¹, respectively). The Zn 108 concentration was higher in SS than in PM (1003 and 320 mg.kg⁻¹, respectively). The Zn 109 concentration in SS was lower than the maximum permissible limits for land application in Europe (2500–4000 mg.kg⁻¹ DM), but higher than the most restrictive threshold values set by national 110 legislation (Sweden: 800 mg.kg⁻¹; Netherlands: 300 mg.kg⁻¹).²³ The Zn concentration in PM was 111 112 in very close agreement with Zn concentrations inventoried in United States, Canada, different European (Switzerland and Austria) and Asian (China and Japan) countries.²⁴ 113

- 116 Table 1: Soil and organic waste characteristics. Concentrations are expressed on a dry matter (DM)
- 117 basis. The analytical methods are described in SI 1 and SI 2.

-		Arenosol		Fluvisol			
		Surface layer (0-20 cm)	Deep layer (20-40 cm)	Surface layer (0-20 cm)	Deep layer (20-40 cm)	Sewage sludge	Poulty manure
Clay	%	2.9	2.3	24.9	28.8	-	-
Silt	%	1.0	0.6	15.3	15.7	-	-
Sand	%	96.2	97.1	59.9	55.5	-	-
Water-Holding Capacity	%	20.0	16.0	36.0	40.0	-	-
Total Limestone (CaCO ₃)	%	6.0	2.5	24.8	23.9	-	-
pH (water)	-	7.7	7.2	8.6	8.8	6.2	8.3
pH (KCl)	-	7.6	7.0	7.3	7.7	-	-
Organic Carbon	%	0.52	0.11	1.24	0.51	35.4	26.7
Total Nitrogen	%	0.05	0.009	0.11	0.034	5	2.44
C:N Ratio	-	10.6	12.2	11.7	15	7.1	10.9
Ash (at 500°C)	%	-	-	-	-	34.0	49.0
Total P	g.Kg ⁻¹	0.23	0.20	0.66	0.24	16.8	17.3
Total Zn	mg.Kg ⁻¹	23 ± 2	5.1 ± 1.1	28.8 ± 1.3	20.7 ± 2.9	1003 ± 40	320 ± 30
Mineralogy		Quartz, cristobalite, sanidine, muscovite		Quartz, calcite, montmorillonite, palygorskite, kaolinite		-	-

120 Soil column experiment

121 The soil columns consisted of polyethylene cylinders (50 cm h, 33 cm dia.) that were reconstituted to reproduce in situ conditions. Two soil layers (0-20 cm and 20-40 cm) were 122 123 sampled in the Arenosol and Fluvisol. Soils were air dried, decumpled and then reconstituted while 124 maintaining the soil bulk densities measured in situ (1.65 kg.dm⁻³ for the Arenosol and 1.4 kg.dm⁻ ³ for the Fluvisol). The weight of Arenosol was 56 kg.column⁻¹ (47 kg.column⁻¹ for Fluvisol). SS 125 126 and PM were applied at the surface of the soil columns at 60 T.ha⁻¹ (0.51 kg.column⁻¹). Control soil columns without OW application (0 T.ha⁻¹) were also monitored. Three replicates per 127 128 treatment (a treatment is defined as a type of soil amended with a type of OW) were performed, 129 representing a total of 18 soil columns.

Lettuces (*Lactuca sativa* L.) were cultivated 1 week after OW application. During the crop cycle (5 weeks), soil columns received 4 L of water twice a week, corresponding to 46.8 mm or 0.3 pore volume (PV) for the Arenosol and 0.25 PV for the Fluvisol. Leached soil solutions were sampled at the bottom of the soil columns weekly and analysed for pH, Zn concentration and dissolved organic carbon (DOC). Lettuce leaves were harvested at the end of the crop cycle. Dry biomass was weighed to determine yields and Zn uptake.

This scheme (OW application and lettuce crop cycle) was repeated for each treatment. At the end of the second cycle (total experiment time: 12 weeks), the soil columns were dismantled and seven soil layers were sampled and analysed.

139 The experimental procedure is described in further detail in SI 3.

141 Zinc speciation in organic wastes and soils

142 X-ray absorption spectroscopy (XAS) was applied to assess the speciation of Zn in OW and the 143 surface layer (0-3 cm) of amended soils (SS-amended Fluvisol, PM-amended Fluvisol and SS-144 amended Arenosol). Zn K-edge XAS spectra were recorded at the ESRF synchrotron (Grenoble, 145 France) on the BM30B (FAME) beamline. Dried OW and soil samples were cryo-ground (Retsch 146 MM400) and pressed into pellets before analysis. Spectra were measured at liquid helium 147 temperature, to minimise beam-induced damage of the samples, in fluorescence mode with a 30-148 element solid-state Ge detector. The spectra of each sample were an average of three to seven 149 scans, depending on the Zn concentration and the signal-to-noise ratio. The acquisition of EXAFS 150 spectra for the PM-amended Arenosol was not possible due to the combination of a low Zn concentration and a very high Fe concentration. Each scan was measured on a different spot on 151 152 the sample pellet to limit beam damage. Energy calibration was performed using metallic Zn 153 reference foil recorded simultaneously with the samples. The absorption edge was defined at 9659 eV, as indicated by the zero crossing of the second derivative of the Zn-foil spectrum. A similar 154 procedure was applied to the previously acquired spectra of Zn reference compounds. 155 Normalization and data reduction were performed according to standard methods²⁵ using Athena 156 157 software.²⁶ A library of spectra from Zn reference compounds was used to identify Zn species in 158 OW and soil (SI4). These references have been synthesized or purchased and described in previous published studies: Nano-ZnS and Amorphous Zn-Phosphate¹³; Zn sorbed on Goethite and Zn 159 sorbed on Ferrihydrite ²⁷; Zn Cryptomelane ²⁸; Zn-Layered Double Hydroxide, Zn sorbed on 160 hydroxy-Al interlayered montmorillonite ²⁹; Zn sorbed on Kaolinite ³⁰; Zn sorbed on 161 Hydroxylapatite³¹; Zn-Malate, Zn-Histidine, Zn-Cysteine^{27,32}; Zn-Methionine^{15,33}; Zn-Phytate 162 ^{15, 34}; Zn-Oxalate, Crystalline Zn Phosphate and Sphalerite (reagent grade, Sigma Aldrich). 163 164 Reference material spectra and sample spectra were measured in the same conditions. Least square linear combination fitting (LCF, SI 5) was performed for each OW or amended soil spectrum over a k-range of 2.5-10.6 Å⁻¹. The residual factor of each LCF was calculated as follows: R = $\Sigma(k^3\chi(k)_{exp} - k^3\chi(k)_{fit})^2/\Sigma(k^3\chi(k)_{exp})^2$. At each step of the fitting, an additional reference spectrum was added and retained in the result if the following two conditions were true: (i) the residual factor decreased by 20% or more, and (ii) the additional reference had a contribution equal to or higher than 10% among Zn-species used for the fit. The uncertainty of this LCF method was estimated at ±15%.²⁵

172

173 Statistical Analysis

Data are presented as means of three independent soil columns and mean values were taken for statistical analyses using one-way ANOVA. Significant differences were further analysed using the Student–Newman–Keuls test to identify differences between modalities. The differences were considered significant if $p \le 0.05$.

179 **Results and Discussion**

180 Accumulation of OW-borne Zn in the soil surface layer

181 The different mass balance terms in the soil column experiments are shown in Table 2. Note that 182 the Zn mass balance ((Initial Zn content in the soil column + Zn input from OW) = (Zn leaching 183 at the soil column outlet + Zn uptake by lettuce)) was not statistically different from the final Zn content in the soil column, indicating that the different mass balance terms had been accurately 184 185 quantified and no compartments (soil, plant or leaching) were overlooked. Since Zn concentrations 186 in the OWs varied (see Table 1), Zn inputs in the amended soils were different: SS application 187 provided 3.1-fold more Zn than PM (1029 and 328 mg, respectively, for the whole experiment). 188 For the Fluvisol columns, there was no significant difference between the leached Zn

189 concentration at the outlet when comparing the control and the amended columns, indicating that 190 exogenous Zn (i.e. Zn derived from OW) was not leached from the Fluvisol column. In contrast, 191 Arenosol promoted leaching of a minor fraction of the exogenous Zn. The leached Zn at the outlet 192 of the Arenosol columns represented 0.15% of the Zn input in the SS-amended soil and 1.9% of 193 the Zn input in the PM-amended soil.

For both soils, OW application enhanced Zn uptake by lettuce as compared to the control soil. Total Zn uptake by lettuce (cumulated for the two cycles) was similar regardless of the treatment (soil+OW), ranging from 0.36% (SS-amended Fluvisol) to 0.68% (PM-amended Fluvisol) of the Zn inputs.

Soil column dismantling and analysis at the end of the two crop cycles highlighted the accumulated Zn distribution through the soil profile (Figure 1). In the Fluvisol, exogeneous Zn remained entirely in the 0–3 cm surface layer for both SS- and PM-amended soils. In the Arenosol, exogenous Zn was distributed in the 0–3 cm and 3–6 cm layers for the SS-amended soil and 0–3 202 cm for the PM-amended soil. Zn derived from OWs was therefore not transferred through the soil203 profile to deeper layers.

204 Overall, 99.6, 99.3 and 99.4% of OW-borne Zn accumulated in the upper soil layers (0-3 and 205 3-6 cm) for the SS-amended Fluvisol, PM-amended Fluvisol and SS-amended Arenosol, 206 respectively, whereas 97.6% of Zn accumulated in the 0-3 cm soil layer for the PM-amended 207 Arenosol. This exogenous Zn accumulation in the upper soil layers after OW spreading has already been observed. For instance, Formentini et al.⁶ assessed the impact of 11 years of pig slurry 208 209 spreading in a clayey Hapludox soil. They concluded that most exogenous Zn accumulated within 210 the topsoil layers, with assimilation by crops and leaching virtually negligible in the overall Zn 211 balance.

This study specifically highlighted a soil effect. The Arenosol promoted leaching of a small fraction of Zn in drained water (SI 6) and migration from 0 to 6 cm depth (Figure 1). The sandy texture of the Arenosol associated with its high hydraulic conductivity might have favored particulate or dissolved Zn movement. Conversely, higher clay, carbonate and organic matter contents, basic pH and the higher cation exchange capacity of the Fluvisol might have favored Zn accumulation in the surface layer (Figure 1).

An OW effect was also highlighted. We observed a 1 pH unit time-course decrease in the solution collected at the outlet of the SS-amended Arenosol (SI 6). This decrease in pH is common and attributed to the acidic effect of OW mineralization and the ammonium nitrification process^{27,} 135, 36. It is well established that pH is the major driver of Zn solubility in soils³⁷ and the observed decrease in pH favoured Zn leaching in the SS-amended Arenosol. However, Zn leaching at the outlet of the PM-amended Arenosol column (1.9% of the Zn input) was surprising as the pH of the soil solution was stable and slightly alkaline (pH=7.3-8) throughout the experiment (SI 6).

225 Nevertheless, Zn leaching was closely correlated with the DOC at the column outlet (SI 7). 226 Complexation with organic ligands could have mobilized a fraction of exogenous Zn.Several 227 studies concluded that colloidal transport can favour trace element transfer in soils following OW spreading. For example, Karathanasis et al.³⁸ studied Cu, Zn and Pb elution in a soil monolith. 228 They demonstrated the elution of significant soluble metal loads in association with water-229 dispersible biosolid colloids. Similarly, Yamamoto et al.²² studied the fate of P, Zn and Cu in soil 230 231 affected by a long-term application of swine manure compost. They demonstrated that continual 232 compost application enhanced the formation of a soil water dispersible colloid in which Zn, Cu 233 and P were accumulated.



- 236
- 237 Figure 1: Zn concentration (mg.kg⁻¹) in the Arenosol and Fluvisol soil layers at the end of the soil
- 238 column experiment. Dashed lines represent the analysed soil layers.

		Arenosol			Fluvisol	
	Control	SS-amended	PM-amended	Control	SS-amended	PM-amended
(1) Initial Zn content in the soil column	809 ± 67	809 ± 67	809 ± 67	1168 ± 98	1168 ± 98	1168 ± 98
(2) Zn input from OW ^a	-	1029 ± 41	328 ± 31		1029 ± 41	328 ± 31
(3) Zn leaching at the soil column outlet ^b	3.7 ± 0.3	$\begin{array}{c} 5.2\pm 0.5 \\ (0.15\%\pm 0.08\%) \end{array}$	9.9 ± 0.9 (1.9% ± 0.56 %)	12.6 ± 2.2	9.3 ± 3.5 (0% ± 0.50 %)	$\begin{array}{c} 11.9 \pm 1.9 \\ (0\% \pm 1.02\%) \end{array}$
(4) Zn uptake by lettuce ^c	0.2 ± 0.04	$\begin{array}{l} 4.6 \pm 0.7 \\ (0.42\% \pm 0.09\%) \end{array}$	1.8 ± 0.2 (0.46%±0.12%)	0.7 ± 0.1	$\begin{array}{l} 4.4 \pm 0.7 \\ (0.36\% {\pm} 0.10\%) \end{array}$	$\begin{array}{c} 2.9\pm 0.3\\ (0.68\%\pm 0.18\%)\end{array}$
(5) Final Zn content in the soil column ^d	768 ± 163	1618 ± 230	912 ± 130	1366 ± 133	2267 ± 132	1567 ± 91
(6) Zn balance: $(1) + (2) - ((3) + (4))$	$805\pm67^{NS~e}$	$1828 \pm 110^{\text{NS}}$	$1126\pm99^{\rm NS}$	$\begin{array}{c} 1154 \pm \\ 100^{NS} \end{array}$	$2183\pm143^{\text{NS}}$	$1481\pm131^{\text{NS}}$

239 Table 2: Mass balances in the soil column experiments. The Zn content is expressed in mg.

^a Zn input from $OW = Zn_{OW C1} + Zn_{OW C2}$, where $Zn_{OW C1}$ is the Zn input from SS or PM for cycle 1 and $Zn_{OW C2}$ is the Zn input from SS or PM for cycle 2.

^b Zn leaching at the soil column outlet = $\sum_{i=1}^{n} [Zn]_i \times F_i$, where $[Zn]_i$ is the Zn concentration measured in the leachate at week i, and F_i is the water flux measured at the bottom of the column at week i. n is the number of weeks (n = 12). Percentage of OW-derived Zn calculated after normalization by Zn leaching measured for the control soil column.

^c Zn uptake by lettuce = $Zn_{PC1} + Zn_{PC2}$, where Zn_{PC1} is the mass Zn uptake by lettuce for cycle 1, and Zn_{PC1} is the mass Zn uptake by lettuce for cycle 2. Percentage of OW-derived Zn derived calculated after normalization by Zn uptake measured for the control soil column.

^d Final Zn content in the soil column = $\sum_{j=1}^{m} [Zn]_j \times MSl_j$, where $[Zn]_j$ is the Zn concentration measured in soil layer j, and MSl_j is the mass of soil layer j. m is the number of soil layers (m=7).

^e NS= no significant difference between the final Zn content in the soil column (5) and the Zn balance (6) at the 5% level.

251 Zinc uptake by lettuce increases with repeated organic wastes application

For both soils, PM or SS application significantly increased the lettuce biomass in comparison to the unfertilized control soil (Figure 2a). No significant differences in yields were found between the first and second lettuce crop cycle within a same treatment (except for the Fluvisol control, for which a decrease in dry biomass was observed). This highlights the fertilization value of OW, which represents a potential alternative to inorganic fertilizers for crop production.³⁹

Significantly higher Zn uptake by lettuce was measured in the OW-amended soils than in the control soils (Figure 2b). Moreover, we observed a significant increase in Zn uptake by lettuce between the first and second OW application for the SS-amended Arenosol and PM-amended Fluvisol. This was also observed for the other two treatments (PM-amended Arenosol and SSamended Fluvisol), but these findings were not statistically significant.

If this trend, i.e. constant crop yield combined with increased Zn uptake, were to be confirmed by long-term experiments with repeated OW application and successive crop cycles, OW fertilization could represent an opportunity to improve the Zn content of staple food crops. Such agronomic biofortification⁴⁰ could help tackle Zn deficiency in human populations which is a cause of illness and death in developing countries.⁴¹ The advantages of using OW to increase the plant Zn concentration should, however, be assessed together with the potential uptake of toxic trace elements such as Cd or Pb.





Figure 2 (a) Mean harvested lettuce dry biomass per soil column (g) and (b) mean Zn uptake by
lettuces per soil column (mg). Different letters represents significant differences between
treatments at the 5% level. PM: Poultry manure; SS: Sewage sludge

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276 Phosphate drives Zn speciation in organic wastes and amended soils

The quantitative LCF results (expressed as a percentage of each Zn species) obtained for both OW and OW-amended soils (0-3 cm soil layer) collected at the end of the experiment are shown in Figure 3. The experimental data were modelled with two Zn references per sample and total contributions ranged from 89 to 111% (SI 5, SI8). Zn species contributions were normalized to 100% for easier comparison. The R-factor of the LCFs ranged from 0.029 to 0.063.





Figure 3: Zn speciation determined from the linear combination fitting of the Zn K-edge extended
X-ray absorption fine-structure spectroscopy data (PM = poultry manure, SS = sewage sludge).

287 2

Zn speciation in OWs

Zinc phosphate (organic, amorphous or crystalline) accounted for 100% of Zn speciation in SS and PM, which is in agreement with the high affinity of Zn for phosphate compounds.⁴² Amorphous Zn-phosphate was found to be the main Zn species in SS (85%) and PM (92%). The second Zn species was another form of Zn-phosphate: organic Zn-phosphate (15% in SS) or crystalline Zn-phosphate (8% in PM). Hopeite (crystalline Zn-phosphate) has already been documented in previous studies as a major Zn phase in OWs.^{15, 16, 18, 22, 43} In the present study, the amorphous Zn phosphate reference closely matched the Zn speciation

of PM and SS. These findings are in accordance with those of a recent survey of Zn speciation in

a large series of OW sampled in full-scale plants¹³ where it was reported for the first time that the 296 297 main Zn species in compost or stockpiled OW was amorphous Zn-phosphate (40–100%). In this survey, Le Bars et al.¹³ also demonstrated that nano-sized Zn sulfide (nano-ZnS, with a crystallite 298 299 size of 3 ± 0.4 nm) was a major Zn species in liquid agricultural OW and SS. This contrasts with 300 our result as zinc phosphate accounted for 100% of Zn speciation in the studied SS. Nevertheless, 301 these discrepancies could be explained by the chemical instability of nano-ZnS which become a minor species after OW composting or stockpiling.^{13, 16} As the SS and PM in the present study had 302 303 been stockpiled for a long period (>3 months), they were exposed to atmospheric conditions, which 304 favors nano-ZnS oxidative-dissolution and could explain why nano-ZnS were no longer observed 305 and were replaced by Zn-phosphate species.

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- 307

Zn speciation in amended soils

308 The best LCFs for both SS-amended and PM-amended Fluvisol were achieved by combining 309 the same references at almost the same proportions. Amorphous Zn-phosphate was the main 310 species, i.e. 84% for PM-amended Fluvisol and 90% for SS-amended Fluvisol. Zn sorbed on 311 hydroxy-Al interlayered montmorillonite (HIM) was the secondary Zn species, i.e. 16% for PM-312 amended Fluvisol and 10% for SS-amended Fluvisol. The latter mineral reference was supported 313 by the XRD clay mineral findings in the Fluvisol (SI 2). For SS-amended Fluvisol, the Zn 314 speciation pattern was in very close agreement with the Zn distribution in the 0–3 cm soil layer: 315 the proportion of exogenous Zn derived from SS represented 90% of the total soil Zn concentration 316 while the proportion inherited from the pedo-geochemical background represented 10% (Figure 317 1). This agreement was less satisfactory for PM-amended Fluvisol: Zn speciation was 318 characterized as 84% Amorphous Zn phosphate + 16% Zn sorbed on HIM, whereas the proportion 319 of exogenous Zn derived from PM represented 66% of the total soil Zn concentration while the 320 proportion of endogenous Zn represented 34% (Figure 1). In this case, the poor quality of the 321 EXAFS spectra (SI 5), as well as the uncertainty of the LCF method ($\pm 15\%$), could explain this 322 discrepancy. The SS-amended Arenosol contained 87% of Zn sorbed on hydoxylapatite and 13% 323 of Zn-phytate, an organic Zn-phosphate already detected in the SS. Zn sorbed on hydroxylapatite 324 was not detected in SS, in which amorphous Zn-phosphate was found to be the main Zn species. 325 However, the featureless shape of the Zn sorbed on the hydoxylapatite EXAFS spectrum suggested 326 that this phase was a Zn-sorption surface complex or a poorly crystalline Zn-phosphate precipitate.³¹ This change of speciation (amorphous Zn-phosphate to Zn sorbed on hydoxylapatite) 327 328 was not fully understood but could be explained by the acidification of the soil solution (-1 pH 329 unit, SI 6) due to organic matter (OM) mineralization. In brief, no radical change in Zn speciation 330 was observed at the end of the 12-week experiment (two OW applications and two lettuce crop 331 cycles) and phosphate was the main driver of Zn speciation in both OWs and amended soils.

332

333 Previous studies and new findings: the benefits of assessing the source speciation to predict 334 the environmental fate

335 To our knowledge, no previous studies have assessed the fate of OW-borne amorphous Zn-336 phosphate species in the soil after OW spreading. More importantly, our results differed substantially from those of previous studies using XAS techniques.¹⁹⁻²² Mamindy-Pajany et al.¹⁹ 337 338 described Zn speciation in a soil mixed with SS and spiked with a Zn nitrate solution. Their results 339 (58% Zn sorbed to Fe oxyhydroxides and 42% Zn bound to OM) highlighted the soil retention 340 mechanism with respect to Zn rather than the fate of exogenous OW-derived Zn. In two studies by Kirpichtchikova et al.²⁰ and Formentini et al.²¹,Zn speciation in soils affected by long-term 341 342 sewage water and pig slurry application, was dominated by its associations with secondary mineral phases (clay minerals and Fe oxyhydroxides). Surprisingly, nano-ZnS accounted for 100% of the Zn speciation in the pig slurry²¹, but nano-ZnS was not detected in the amended soil. The authors assumed that the nano-scale of the ZnS favoured oxidation of this species in the soil, in turn releasing Zn that was subsequently retained by the soil matrix. We think that an identical mechanism could explain the Zn speciation in the soil amended by sewage water²⁰, since the conditions in liquid OW are often reducing hence leading to the formation of nano-ZnS¹³, which might have oxidized and dissolved after sewage water spreading.

Yamamoto et al.²² investigated Zn speciation in pig manure compost and in a soil amended with this compost for 23 years. Their results highlighted that oxidized species present in the compost, i.e. Zn-phosphate and Zn bound to OM, had persisted and accumulated in the amended soil. Our results confirmed that Zn present in OW stockpiled for a long period (>3 months) did not undergo a considerable change in speciation after OW application and was mainly accumulated in the soil surface horizon, despite the highly contrasting properties of both studied soils.

After application of fresh OW with a high Zn-sulfide content, the final Zn speciation would be soil dependent as the sulfide would oxidize and release Zn, which in turn would be sorbed/precipitated on the soil matrix.^{13, 21} Conversely, if the OW had been stockpiled/composted with sufficient phosphate present, then OW-borne Zn-phosphate species accumulate and persist in the amended soil.

The results of the present study and the aforementioned studies jointly point towards the same conclusion. Speciation of trace elements in the source, i.e. OW, is the main driver that controls the Zn fate in the soil-plant-water compartments following OW application. Neglecting trace element speciation in the contamination source (OW in this case) hampers gaining further insight into the environmental fate of contaminants in amended soils.

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376

377 Supporting Information

378 Soil, plant, leached water and organic waste sampling procedure, analytical methods and 379 experimental design; X-ray diffraction; Zn K-edge EXAFS spectra and radial distribution function 380 of reference compounds; linear combination fitting results; variations in pH, DOC and Zn in the 381 drained water at the soil column outlets; Comparison of EXAFS spectra and radial distribution 382 function of reference compounds.

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- 507
- 508

509 Supporting Information

510	Zinc speciation in organic waste drives its fate in
511	amended soils
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- 529 number of tables 1

530 SI 1 Soil, plant, leached water and organic waste sampling procedure and analytical methods

531

Soils

532 Soils were sampled in the 0-20 and 20-40 cm layers from two sites located in suburban 533 agricultural areas (Pikine and Rufisque market gardening areas located on an Arenosol and a 534 Fluvisol, respectively) of Dakar (Senegal). Only steel or plastic tools were used for sampling in 535 order to avoid trace element contamination of soils. After air-drying and declumping the 536 aggregates, soils samples were sieved through a 2 mm mesh sieve.

537 After organic matter (OM) dissolution and carbonate removal, soil particle size analyses (sand, 538 silt and clay) were conducted using the combined pipette sieving and sedimentation method (NF 539 X 31-107). The total limestone or carbonate content in soil was estimated by the volumetric method 540 (ISO 10693). Total nitrogen (ISO 13878) and total carbon (ISO 10694) were determined by dry 541 combustion with an elemental NC 2100 Soil Analyzer (Thermo Electron Corp.), and organic 542 carbon was determined according to the difference with mineral carbon as calculated from the carbonate content (ISO 10694). Soil and OW pH were determined according to the potentiometric 543 544 method in the supernatant of the soil-water and OW-water suspension with 1:2 and 1:5 ratios, 545 respectively (ISO 10390). For total P and Zn analyses, sieved soil sub-samples were ground to 100 546 um particle size in zirconium material before dissolution. After 2 h calcination at 500°C, total 547 dissolution was performed by acid digestion using a mixture of hydrofluoric (HF), nitric (HNO3) 548 and perchloric (HClO4) acids (ISO 14869-1). Total P and Zn concentrations were then determined 549 using inductively coupled plasma mass spectrometry (ICP-MS). Furthermore, two reference 550 standard certified soil samples, i.e. CRM 7001 (light sandy soil) and CRM 7004 (loam), were used 551 to assess the accuracy of the results according to the ISO 9001 certification of the laboratory. The 552 variation coefficient for both of these checks must be less than 5%.

553 At the end of the second cycle, the soil columns were dismantled and seven soil layers were 554 sampled as follows: 0-3, 3-6, 6-10, 10-15, 15-20, 20-30 and 30-40 cm. Soils were air-dried, sieved 555 through a 2 mm mesh sieve, and ground to 100 µm using a planetary grinder made of zirconium 556 oxide to avoid trace element contamination. Zn contents were determined by ICP-MS after 2 h 557 calcination of the soil samples at 500°C, followed by total dissolution using a mixture of 558 hydrofluoric (HF), nitric acid (HNO₃) and perchloric acid (HClO₄) (ISO 14869-1). Furthermore, 559 two reference standard certified soil samples: CRM 7001 (light sandy soil) and CRM 7004 (loam) 560 were used to assess the accuracy of the results according to the ISO 9001 certification of the 561 laboratory. Deviations from the certified values of these controls were less than 5%.

562

Leached water

pH, dissolved organic carbon (DOC) and Zn were analysed in the leached water. For this 563 564 purpose, weekly leachate samples collected at 40 cm depth (i.e. at the bottom of the soil columns) 565 were filtered using grade GF/D Whatman microfiber filter paper (pore size 2.7 µm) and subdivided 566 in three subsamples. pH was determined in the first leachate subsamples immediately after 567 sampling. The second leachate subsamples were conditioned and acidified with 2.5% ultrapure 568 nitric acid (v/v) in 10 mL scintillation flasks (previously washed with a 10% nitric acid bath and 569 rinsed with ultrapure water), and stored at 4°C. The Zn concentration in acidified leachate 570 subsamples was determined using ICP-MS. The third leachate subsamples conditioned in 20 mL 571 brown amber glass bottles (prewashed in a 10% NaOH bath and rinsed with ultrapure water), were 572 treated with 0.1% sodium azide (NaN₃) and stored at 4°C until use. DOC in azide-treated leachate 573 subsamples was determined using a Total Organic Carbon (TOC) Analyzer (Shimadzu, TOC-L) 574 in NPOC mode. Samples were bubbled for 90 s with 1.5% HCl to eliminate carbonates before oven injection (680°C). The DOC measurement quality was checked with the certified reference
sample MISSIPPI-03 and the variation coefficient was less than 10%.

577

Plants

578 Freshly harvested leaves of mature lettuce plant samples were rinsed several times with distilled 579 water, cut into small pieces, and oven-dried at 60°C to constant mass. Then these samples were 580 ground into powder separately through a grinder made of zirconium oxide (bowls and balls) in 581 order to avoid trace element contamination. Total digestion of ground lettuce samples consisted of 582 introducing 1 g of each ground lettuce sample into a platinum crucible which was subsequently 583 placed in a muffle furnace for 2 h calcination at 500°C. After cooling, the ash was moistened with 584 a few drops of ultrapure water, and 2 mL of 6N HCl was added. Dry evaporation was then carried 585 out on a warming plate. Hydrofluoric acid was added to the ash residue and then evaporated again 586 on a warming plate. The silica contained in the ash thus volatilized as SiF6. The ash residue was 587 taken up in 2 mL of 6N HCl and filtered through Whatman No.42 filter paper into a 50 mL vial. 588 The filter containing the residue was placed in the same crucible and then calcined at 500°C. 589 Finally, the crucible was rinsed with 1 mL of 6N HCl, and then with ultrapure water (18.2 MΩ.cm) 590 from a Millipore Milli-Q system, and transferred to the same vial. The vial was gauged and then 591 homogenized by manual stirring. Furthermore, three strawberry plant samples (Fragaria vesca) 592 with internal references were also analyzed for verification and validation of the results. Deviations 593 from the certified values of these controls were always less than 5%. All reagents used were Merck 594 analytical grade (AR). The Zn concentration in digestates was determined by inductively coupled 595 plasma mass spectrometry (ICP-MS, Varian Ultramass).

06 Organic wastes

597 Sewage sludge (SS) was sampled at the Pikine municipal wastewater treatment plant. This 598 lagoon-type plant is run on the basis of biological processes with free culture in basins and 599 produces 1905 t of activated sludge per year. SS are sun dried in a refining tank.

600 Poultry manure (PM) from laying hens was sampled in a henhouse at Rufisque. The manure is 601 mixed with litter (sawdust or rice husking waste) used as a henhouse substrate.

Both SS and PM had been stockpiled for long periods prior to being spread on the soil columns

603 (>3 months), which is consistent with market gardeners' practices.

Organic waste was dried at 40°C then ground with a grinder made of zirconium oxide (bowls and balls) in order to avoid trace element contamination, and then ground and sieved through a 1 mm mesh sieve. The characteristics of these OW soils were obtained using standard soil analysis

607 methods.

608

610 SI 2 X-ray diffraction

611 X-ray diffractograms of the soils were obtained with a Philips PW 3710 X-ray diffractometer

612 using CoK α radiation at 40 kV and 40 mA (a counting time of 12 s per 0.02° step was used for the





616 Figure SI-2: X-ray diffractograms for (a) Arenosol and (b) Fluvisol

617 SI 3 Experimental design

The soil columns were composed of cylindrical polyethylene columns (50 cm h; 33 cm dia.). The bottom of each column was covered by a nylon membrane (Nitex, 50 μ m mesh) to prevent any soil particle release. The bottom was perforated with five holes (1 cm dia.) to collect the drained water. At the outlet, each column is equipped with a funnel and a polyethylene bottle to collect the leachate. Two soil layers (0-20 and 20-40 cm) were sampled for the Arenosol and the Fluvisol. Soils were air dried, decumpled and then assembled while maintaining the soil bulk

624 densities measured *in situ* (2.65 kg.dm^{-3}).



625

626 Figure SI 3: Soil column

Sewage sludge (SS) and poultry manure (PM) were applied at 60 t DM/ha (60 T.ha⁻¹). Control soil columns without OW application (0 T.ha⁻¹) were also included. Predried OWs were uniformly spread on the soil surface and immediately incorporated into the surface soil layer (0–2 cm) by hand before sowing. About 25 lettuce (*Lactuca sativa L.*, Eden variety) seeds were sown in the soil column subsurface and covered with a thin soil layer (about 2 mm depth). After germination, at the 3 to 4 leaf stage, the seedlings were thinned in order to leave a dozen vigorous lettuce plants
per column. Lettuces (*Lactuca sativa* L.) were cultivated 1 week after OW application. Each
lettuce crop cycle lasted 5 weeks.

635 The pore volume (PV) of each soil column was calculated using the equation reported by Feder
636 et al. (2015)³⁶:

- 637 [Pore volume] = Porosity \times Volume of the soil column.
- 638 with Porosity = $1 \frac{AD}{RD}$

where AD (kg.dm⁻³) corresponds to the apparent density or bulk density measured for each soil *in situ*, and RD corresponds to the real particle density (expressed in kg.dm⁻³) estimated at 2.65 kg
.dm⁻³ for both soils.

Each column received 8 L of water weekly, corresponding to 0.6 PV for the Arenosol columns and 0.5 PV for the Fluvisol columns. Water (4 L) was applied on fixed days (Monday and Thursday). 24 h after each water supply, the leachates drained and collected at the column outlets were sampled. On a weekly basis, the two leachate samples from the same soil column were combined and stored before analysis.

At the end of the lettuce crop cycle, only the above-ground parts (leaves) of the plants were harvested. The plants were cut, then washed three times successively with tap water, and rinsed twice with distilled and ultra-pure water. The lettuces were dried in an oven at 60°C to constant weight and the dry biomass was weighed to determine the lettuce yields.

651 After the first lettuce harvest, the whole experiment (OW application, lettuce crop, water 652 irrigation and sampling) was repeated a second time.

Three replicates were performed for each treatment (a treatment is defined as a soil and an OW)

654 representing a total of 18 soil columns.





Figure SI 4a: Zn K-edge extended X-ray absorption fine-structure spectroscopy spectra of allreference compounds.



660 Figure SI 4b: Radial distribution function (window Kaiser-Bessel, dk=2) of all reference661 compounds.





Figure SI 5: Experimental (black) and modeled (dotted, red) Zn K-edge extended X-ray absorption
fine-structure spectroscopy spectra and radial distribution function (window Kaiser-Bessel, dk=2)
for each sample of OW and soil + OW. SS= sewage sludge, PM=poultry manure

668 Table SI 5: Detailed linear combination fitting results. SS = sewage sludge, PM = poultry manure

	Amorphous Zn- phosphate	Zn- phytate	Cristalline Zn- phosphate	Zn sorbed on hydroxylapatite	Zn sorbed on hydroxy-Al interlayered montmorillonite	R- factor*
Arenosol + SS		14%		97%		0.050
Fluvisol + PM	75%				14%	0.063
Fluvisol + SS	91%				10%	0.029
РМ	96 %		10%			0.030
SS	86 %	15 %				0.037

*R-factor = $\Sigma (k^3 \chi(k)_{exp} - k^3 \chi(k)_{fit})^2 / \Sigma (k^3 \chi(k)_{exp})^2$

670



672 SI 6 variations in pH, DOC and Zn in the drained water at the soil column outlets.



Figure SI 6: Variations in pH, DOC and Zn concentrations in drained water at the outlet of the (a)
Arenosol columns and (b) Fluvisol columns. Arrows represent the OW applications. PM: Poultry
manure; SS: Sewage sludge

677 SI 7 Zn concentration as a function of the dissolved organic carbon concentration in drained

678 water at the soil column outlets.

679



Figure SI 7: Zn concentration as a function of dissolved organic carbon in drained water at the column outlets of the Arenosol (a) and Fluvisol (b). R corresponds to the Pearson linear correlation coefficient; NS = not significant; Significance level: * = p < 0.05, and *** = p < 0.0001.

686 **<u>SI 8 X-ray absorption spectroscopy – comparison of reference compounds.</u>**

687 The EXAFS spectra of Zn sorbed on hydroxyapatite, amorphous Zn phosphate and Zn sorbed on Ferrihydite look very similar, although some minor features can be observed. But, these distinct 688 689 features are much easier to detect on the radial distribution function. The radial distribution 690 function obtained from Fourier transforming the reduced EXAFS data consists of different peaks 691 resulting from Zn-backscatter interactions. The peaks between 2 and 4 Å indicate the presence of atoms in the next-nearest shells around Zn. For amorphous Zn phosphate, Zn sorbed on 692 693 hydroxylapatite and Zn sorbed on Ferrihydrite, the radial distances of these peaks are different (see 694 arrows), thus indicating a modification of the local Zn environment for these three references.



Figure SI 8: Zn K-edge extended X-ray absorption fine-structure spectroscopy spectra and radial
distribution function (window Kaiser-Bessel, dk=2) for amorphous Zn phosphate, Zn sorbed on
hydroxylapatite and Zn sorbed on Ferrihydrite.

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