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1 Technosol rehabilitation strategies drive soil

² physico-chemical properties and fauna diversity on a

3 former coking plant area

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7 Abstract

8 Anthropic activities such as mining, resource extraction or transformation profoundly modify 9 ecosystems and may lead to Technosol formation. Post-industrial areas are examples of potentially 10 degraded lands, due to soil contamination with metals or organic pollutants, as well as soil structure 11 damage. Soil biodiversity being closely involved in many soil functions such as organic matter decomposition, formation and stabilization of soil structure, the recovery of degraded soil functions 12 13 partly depends on soil fauna colonization. However, the relationship between Technosol abiotic 14 parameters and soil fauna colonization is still to be disentangled. In an attempt to fill this gap, we 15 studied a former coking plant area in north-eastern France, composed of Technosols resulting from 16 coking plant embankments or thermally-treated industrial soils and compared them with two local 17 soils considered as references. We hypothesized that the taxonomic and trophic diversity of 18 Technosol-dwelling invertebrates would be more diverse and abundant in the soils with the higher soil physico-chemical quality (i.e higher fertility and lower pollutions levels). To test this hypothesis, we 19 20 studied four Technosols that were settled following two different rehabilitation strategies within the 21 same post-industrial area and we compared them with two local soils of reference using nested ANOVAs as well as multivariate analyses of soil abiotic parameters and soil fauna community 22 23 indices, both within the soil and on its surface.

The results on physico-chemical analyses showed that the substrates used for Technosol rehabilitation 24 were impoverished in clay content when compared with local soils of reference (4.1 to 7.8-fold) and 25 26 enriched in sandy materials. The use of coking plant by-products for Technosol implementation have led to poor chemical quality, with low nutrient content but high organic carbon content (41 to 51 %) 27 resulting from the use of coal and correlated with the higher lead concentration in the area. The use of 28 29 thermally-treated industrial soil has led to more fertile Technosols with significantly lower lead 30 content. Meso and macrofauna densities did not vary significantly between the Technosols and the 31 local reference soils. Both Technosol-dwelling fauna trophic and taxonomic community compositions 32 were impacted by the rehabilitation strategy. Few to no earthworms were found in Technosols (mean up to 16 ind.m⁻²) compared to the local soils of reference (mean of 118.4 and 201.6 ind.m⁻²). 33 Conversely, Technosols resulting from coking plant embankments and thermally-treated industrial 34 soils were dominated by epigeic soil fauna with an intense activity of soil surface macro-saprophages 35 and micro-arthropods, as well as Formicidae. Our results suggest that the use of technogenic elements 36 37 in the rehabilitation of post-industrial area led to the development of specific soil invertebrate 38 communities, different from the reference. The gap between the high level of activity of epigeic organisms that we found on Technosols and the low trophic diversity of the litter and soil-dwelling 39 communities suggest that the soil fauna community on a former coking plant is driven rather by soil 40 41 physico-chemical properties than by colonization constraints.

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43 Keywords: Anthropogenic soil, Lumbricina, Mesofauna, Metacommunity, Soil fertility

44 **1. Introduction**

The human footprint on ecosystems is continually increasing (Kareiva et al., 2007; Vitousek et al., 1997). Over recent decades, a growing environmental concern has emerged while degradations of ecosystems by human activity have been evidenced by numerous studies (Isbell et al., 2015; Millennium Ecosystem Assessment, 2005; Moreno-Mateos et al., 2017). Soil systems are particularly affected by human activities, as testified to by urban expansion, agriculture intensification, or resource

extraction and transformation such as mining, coal or steel industries. In Europe, the ceasing of 50 industrial activities has left behind degraded ecosystems, including brownfield, mining and resource-51 52 transformation areas (Frouz et al., 2001; Mudrák et al., 2010; Šourková et al., 2005; Wong and Bradshaw, 2003). Degradation can include contamination by metals or polycyclic aromatic 53 hydrocarbons (PAH), or the overall removal of vegetation and arable soil, such as in post-mining sites. 54 Technosols, which are highly anthropized soils, containing at least 20 % of artefacts within the first 55 56 100 cm, or have a cemented or indurated layer below 10 cm depth, or a continuous geomembrane or have technic hard material starting above 5 cm depth (IUSS Working Group WRB, 2015). They 57 58 usually result directly from past industrial activities (Morel et al., 2015), but may also be a consequence of the rehabilitation strategy of a polluted site. In the past, the rehabilitation goal was 59 sometimes limited to aesthetic properties, or allowing convenient succession of the area (heaps, 60 burrowing, leveling), resulting in Spolic Technosol, with human-transported soil material (IUSS 61 62 Working Group WRB, 2015). Nowadays the aim of rehabilitation is focused more on sanitary and ecological matters and might be used to improve the ecosystem services provided by a given area 63 64 (Beroigui et al., 2020; Séré et al., 2008; Sheoran et al., 2010; Sopper, 1992). Constructed Technosols have thus emerged as one way to achieve such rehabilitation. Technogenic inputs into the soil might 65 result from a former activity (industrial wastes), the decline of the activity (dismantling of the 66 67 buildings, embankments) or from the soil rehabilitation strategy itself (input of exogenous substrates). This phenomena, called "Technosolisation" (El Khalil et al., 2013), may change the soil fauna 68 community composition and limit its colonization (Eijsackers, 2010) and may change the soil 69 70 microbial and physicochemical properties (Piotrowska-Długosz and Charzyński, 2015). Yet, soil fauna is known to play key roles in soil functioning, such as carbon and nutrient cycling, organic matter 71 72 decomposition, soil structure formation and stabilization (Orgiazzi et al., 2016). Thus, rapid Technosol 73 colonization by fauna in the first years after rehabilitation could enhance the recovery of degraded soil 74 functions and provide insight into the quality of the rehabilitation strategy (Pruvost et al., 2020; 75 Uzarowicz et al., 2020).

While studies have been conducted of the dynamics of highly anthropized soil fauna colonization in a 76 77 given rehabilitation strategy, usually revegetation or spontaneous succession of open-mining sites 78 (Frouz et al., 2006; Frouz and Nováková, 2005; Wanner and Dunger, 2002), the ways by which the 79 rehabilitation strategies shape the aboveground-belowground relationships on Technosol are still to be disentangled (Kardol and Wardle, 2010). Taxonomical and functional approaches of Technosol-80 81 dwelling fauna colonizers have recently been receiving more attention (Burrow, 2018; Pey, 2010; 82 Vergnes et al., 2017), yet the conclusions remain limited (Villenave et al., 2018) and further research 83 could provide useful clues as to how to optimize forthcoming rehabilitation strategies and increase 84 their sustainability. In an attempt to fill this gap, we studied the soil fauna metacommunity of a former 85 coking plant area, composed of several Technosols in north-eastern France. Since the dismantling of the coking plant in 1985 in this area, different strategies have been set up such as embankments with 86 coking plant by-products, a depollution process of excavated soil to reduce PAH concentration level 87 and in situ Technosol construction (Séré et al., 2008), as well as organic matter inputs, resulting in the 88 89 presence of different type of Technosols in a single geographical area. This mosaic of Technosols may 90 lead to different patterns of soil fauna colonization and thus to different spatiotemporal soil function 91 recoveries. The goals of our study were to establish (i) the influence of the soil substrates used in the 92 different rehabilitations on the soil physical and chemical fertility parameters, (ii) the recovery of the 93 soil taxonomical and trophic invertebrate communities in Technosols of the former coking plant area 94 in comparison with local soils of reference. A final goal was to establish if there is a link between the 95 quality of the Technosol physico-chemical properties (i.e. fertility, nutrient content, pollution) and the 96 level of biodiversity. We hypothesized that the physico-chemical quality of the Technosol (i) would 97 depend on the nature of substrates used for the rehabilitation, and (ii) would be lower in Technosol 98 constituted by coking plant embankments than in depolluted soils, and in soils of reference. Finally, 99 we hypothesized that Technosols with a better physico-chemical quality would exhibit a more diverse 100 taxonomic and trophic soil fauna community in comparison with the others. To test our hypothesis, we 101 assessed the soils' abiotic parameters and the soil fauna taxonomical and trophic community composition of Technosols resulting from two different rehabilitation strategies. We added to the 102 study two local soils of reference, resulting from natural pedogenetic processes and located in the 103

same area to compare both their fertility parameters and fauna community composition to those ofTechnosols.

106 2. Material and methods

107 2.1. Site and soils studied

108 The study was carried out at a former coking plant site (32 ha), in north-eastern France 109 (49°12'49.87"N, 05°59'43.10"E, 225 – 250 m above sea level) of the group of scientific interest on soil study (GISFI) experimental station at Homécourt (www.gisfi.univ-lorraine.fr). The mean annual 110 rainfall in the area is 720 mm and the mean annual temperature is 11°C. Two coking plants succeeded 111 112 each other between 1923 and 1980, the latter being dismantled in 1985. Their activities led to soil contamination with heterogeneous levels of heavy metals and polycyclic aromatic hydrocarbons 113 114 (PAH) (Biache et al., 2013). The site studied is now composed of a mosaic of ecological units, such as meadow, grassland, woodland, spontaneously developing or artificially implanted on different 115 116 Technosols.

On the former coking plant site, six sub-sites were studied and considered as follows: four Technosols
were chosen based on their historical background concerning the rehabilitation method they received.
Simultaneously to the four Technosols, a woodland (named WOOD) and a meadow (named MEAD)
located in the same area close to the former industrial

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site were selected as soils of reference, hereafter called "REF". Sampling sub-site localizations arepresented in Fig. 1.

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The first type of substrate that composed the first two Technosols studied (COAL and EMB, Fig.1) was embankments with coal tailing and remnants from the buildings dismantled in 1985, hereafter called "CPE". These embankments were created between 1986 and 1989 to level the soil, with no addition of organic matter nor any depollution (LECES, 2001). The absence of cemented or indurated layer starting in the first ten centimeters of the profil and the proportion of artefact exceeding 20% within the first 100 cm of the profile allow to classify the soils as Technosol. The first Technosol (named COAL) is a Spolic Technosol composed of more than 50 cm of coal embankments on which a herbaceous stratum has developed; the second, (named EMB) is a Spolic Technosol composed of a mixture of embankments more than 1.3 meters deep, with a spontaneously developed forest composed of *Populus tremula, Robinia pseudoacacia* and *Betulus pendula* (pioneer plants of the former coking plant studied).

136 The following two Technosols studied (TDT and CONS, Fig.1) share a common rehabilitation strategy, hereafter called "ITS". Implemented in 2003, this was conducted to reduce the soil PAH 137 concentration level as required by French policy. The main substrate of the two Technosols was a 138 thermally-treated industrial soil. Both soils contained more than 20% of artefact within the first 100 139 140 cm of the profile and neither exhibited a cemented or indurated layer within the first ten centimeters thus were defined as Technosols (IUSS Working Group WRB, 2015; Séré et al., 2008). A Technosol 141 (named TDT) settled in 2003, which was composed of 30 cm of this substrate, without any further 142 material addition. The last Technosol studied (named CONS) was constructed in 2008 and is 143 composed of 100 cm of a mixture of paper mill sludge and thermally-treated substrate covered by ten 144 145 centimeters of green-waste compost. This strategy was conducted to recover potential economic value 146 and to allow an improvement of the ecosystem services provided by a part of the area, using pedoengineering processes (see Séré et al. (2008) for more details). 147

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149 2.2. Technosols and soils of reference abiotic parameter characterization

All the samples were collected between the 25th and the 30th of April, 2018. At each of the 6 sub-sites, five samples were taken (except for COAL sub-site for which N=4) in order to characterize abiotic parameters (i.e. soil chemical and physical analyses) to quantify the quality of the four Technosols and two soils of reference. A cross-shaped pattern was chosen for the sampling strategy, with the center of the cross chosen randomly and the other four points were ten meters distant from the center and at least 5 m away from the edges to avoid any edge effect (Fig. 2). 157 A total of 21 abiotic variables representing generally-studied soil fertility parameters and contamination levels were measured. At each sampling point (N=5, except for COAL N=4), four 500 158 g soil samples, taken from the first 0-20 cm soil layer, were pooled into one composite sample (Fig. 2 159 160 squares "1"). Prior to analyses, soils were air-dried for 72 h then sieved at 2 mm. In order to study 161 comparable materials, we used two successive soil homogenization techniques: first reducing the size of the overall sample to 500 g using an open-bin riffle splitter (Schumacher et al., 1990), then using a 162 rotary splitter coupled with a vibratory feeder with a V-shaped channel (Laborette 24, Fritsch), thereby 163 providing two homogeneous samples. One was kept in the laboratory for chemical analyses and the 164 165 other was sent for particle size distribution analysis (NF X 31-107, Laboratoire d'Analyses des Sols,

166 www6.hautsdefrance.inra.fr/las).

Soil pH was measured on 5 mg of 2 mm-sieved soil in 25 mL distilled water (v/v) (ISO 10390) with
three replicates for each sample.

169 For the following analysis, soil samples were ground to less than 250 µm using a mixer mill (MM 400, 170 Retsch). Organic carbon and total nitrogen were measured using an elementary analyzer (vario 171 MICRO, Elementar) after soil sample combustion. Decarbonation was performed prior to analysis for organic carbon by adding 2 µL of chloridric acid (4 mol L⁻¹) until gassing ceased (ISO 10694). Total 172 and exchangeable cation exchange capacity (CEC) were measured using the cobalthyhexamine 173 174 method (ISO 23470). Available P was estimated by measuring soluble P in sodium bicarbonate (Olsen method, ISO 11263). Measurements of total and DTPA-extractable soil elements were made using 175 inductively coupled plasma with optical emission spectroscopy (ICP-OES) (NF ISO 11466 and NF 176 177 EN 13651 respectively).

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Finally, one soil corer (5 cm) was sampled at 5 cm depth in order to measure the bulk density (Fig. 2
square "2"). Soil bulk density (Bd) was calculated using the ratio of the soil dried mass at 105 °C (P_s)
to the volume of the sampled corer (NF EN ISO 11272):

182
$$Bd = \frac{Soil \, dry \, weight}{Corer \, volume}$$

183 Water holding capacity (WHC) was assessed using a porous membrane press according to the standard184 NF EN ISO 11274.

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186 *2.3. Soil fauna community analysis*

187 2.3.1 Soil fauna sampling and identification

At the time of the abiotic sampling and at close proximity (10 cm to 45 cm), three types of invertebrate communities were studied using combination of soil sampling and pitfall traps to assess the density and activity of both macrofauna and mesofauna, as previously performed on industrial areas (Hedde et al., 2018).

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Firstly, the litter and soil-dwelling macrofauna (organisms > 2 mm) was sampled using the normalized TSBF method adapted to temperate regions (ISO 23611-5). One soil monolith of 625 cm² was extracted to a depth of 30 cm, including any litter layers at a given sub-site (Fig. 2, square "3"). All organisms were hand sorted and immediately stored in 70 % (v/v) ethanol solution.

Litter and soil-dwelling micro-arthropods (0.1 to 2 mm) were sampled within the first 10 cm of each soil using successively 5 cm deep and 10 cm diameter corers (Fig. 2, circle "4"). The corers were placed in a high-temperature gradient extractor (Macfadyen, 1961) to extract micro-arthropods then stored in a 70 % (v/v) ethanol solution.

Finally, in order to study soil surface-dwelling invertebrates, we placed three pitfall traps 1 meter apart and 45 cm from each abiotic parameter sampling point and soil-dwelling invertebrate sampling points for a total of 15 pitfall traps (12 pitfall traps for COAL) carried out on each sub-site (Fig. 2, circles "5"). The traps are composed of a plastic cup, filled with glycol and soap with a 100:1 ratio (v/v). The 87 traps were collected after one week and the animals were transferred and stored in 70 % (v/v) ethanol before identification.

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All soil fauna identification was done using a stereomicroscope (MZ FLIII, Leica) at a magnification of x8 for macrofauna and x45 for micro-arthropods. Macro-invertebrates were identified at taxonomical levels of the class, order, sub-order or family depending on the group (see Table 2 for 211 litter and soil-dwelling macrofauna sampled in litter and soil monoliths; Table 3 for soil surface 212 dwellers sampled by pitfall traps method). Then, in order to study the functional profile of the soil 213 fauna communities across the 6 soils studied, we assigned a trophic type to each zoological group as 214 previously done in many studies (Hedde et al., 2018; Salmon et al., 2006). The trophic assignments are 215 shown in Tables 2 and 3.

Finally, soil micro-arthropods were identified as: collembolas (saprophages), oribatid mites (saprophages) and other mites (mostly represented by mesostigmates; predators). Distinction between Oribatida and Mesostigmata mites were performed based on the presence of both anal and genital plates on the abdomen of Oribatida mites (Balogh and Balogh, 1992; Karg, 1993).

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221 2.3.2 Taxonomic and trophic fauna community indices calculation

222 Ecological indices describing edaphic fauna communities were calculated for each Technosol and soil 223 of reference: litter and soil-dwelling taxonomic richness for macrofauna and micro-arthropods (mean of zoological groups sampled at each sub-site), total macrofauna and micro-arthropods densities per 224 225 m² (mean of the densities sampled at each sub-site). The number of ants and snails were not taken into account in the calculations of the litter and soil-dwelling communities because of their high 226 aggregation capacity and the number of empty gastropod shells (Petersen and Luxton, 1982). The 227 228 trophic indices of litter and soil dwelling community were also calculated: the abundance and 229 proportion of macro-saprophages, or macro-geophages or macro-predators, and trophic richness 230 (number of trophic groups in the community).

The Shannon index (H) was calculated for soil and litter-dwelling macrofauna in each of the fourTechnosols and two soils of reference studied as follows:

233
$$H = -\sum_{i=1}^{S} p_i \log_2 p_i, \text{ where } p_i = \frac{x_i}{\sum_{i=1}^{S} x_i}$$

where p_i = proportion of organisms belonging to the zoological group "*i*", S = total number of zoological groups and x_i = number of organisms belonging to the zoological group "*i*".

For the surface soil-dwelling organisms (collected by pitfall traps), we considered that the 3 pitfalls 237 installed at a given sampling point were dependent on each other because of their proximity, but were 238 239 independent from the other sampling points of a given sub-site. Consequently, abundances of 240 invertebrates sampled in the 3 dependent pitfall traps were gathered, and we considered N=5 (=15/3) for one given site (N=4 for COAL). From this sampling method, we calculated 6 indices that described 241 the soil surface-dwelling community for each of the Technosols and soils of reference: the 242 243 taxonomical richness, the number of macro-organisms and micro-arthropods, the number of macro-244 saprophages and the proportion of macro-saprophages and macro-zoophages in pitfall traps.

245

246 2.4. Statistical analyses

Mean comparison of Technosols and soils of reference physical and chemical parameters and invertebrate communities were performed, with the soil substrate as a random effect and the sub-site as a fixed effect, using nested ANOVA after ensuring that homoscedasticity and normality of the residuals of the linear model were met. Post-hoc tests were performed for multiple mean comparison using the TukeyHSD test and using Bonferroni correction in order to reduce the risk of falsely rejecting the null hypothesis.

In order to assess the heterogeneity of our data sets, two principal component analyses (PCA) were performed using the dudi.pca function of the ade4 package in R software. The first PCA was done on abiotic parameters and the second PCA was carried out on fauna communities' taxonomical and trophic indices calculated for the three types of communities.

In order to assess the relationships (co-variation) between abiotic and biotic variables, the coinertia function of the ade4 package was used to perform a co-inertia analysis between abiotic and biotic PCAs. For the two PCAs, variables were normalized in order to ensure equal row weights for the coinertia. The statistical significance of the co-structure between the two data sets was tested by the Monte Carlo test (RV score) with 1000 permutations with the randtest function of the ade4 package. All statistical analyses were performed using RStudio software (version 1.1.383, 2017).

264 *3*. **Results**

265 *3.1 Technosol physico-chemical characteristics according to the rehabilitation strategy*

A PCA was conducted on the 21 measured abiotic variables (Fig. 3). The first two axes represent 266 267 60.4% of the total variation, with the first and second axes representing 33.7% and 26.1% of the 268 variation respectively. Mean values and standard deviations of the abiotic parameters studied at each sub-sites are presented in Table 1 with the results of the corresponding nested ANOVAs. The first axis 269 270 of the PCA is negatively correlated with clay content, WHC, potassium and phosphorus concentration and positively correlated with C:N ratio, sand content, total and DTPA extractable lead. The second 271 272 axis of the PCA is negatively correlated with pH values and Mg concentrations and is positively 273 correlated with total nitrogen concentration and exchangeable Mg. The first axis of the PCA separates 274 the characteristics of the REF soils of reference sub-sites from the ones of the CPE rehabilitation strategy while the second axis mostly separates the Technosols of the ITS strategy from the REF and 275 276 CPE. Although the PCA exhibits three clusters of points for the abiotic parameters, only 5 out of the 21 parameters differed significantly between rehabilitation strategies and soils of reference (Table 1). 277 Organic carbon content was significantly higher in the Technosols of the CPE rehabilitation strategy 278 with respectively 506.83 and 409.85 g kg⁻¹ in COAL and EMB (p-value = 0.016), which is one order 279 of magnitude greater than in TDT sub-site. Significantly different clay and sand contents between the 280 281 Technosols and soils of reference resulted from the different substrates used for rehabilitation. Clay content was higher in the REF soils of reference, with respectively 495 and 507 g kg⁻¹ in sub-sites 282 283 MEAD and WOOD (p-value = 0.001), while the two rehabilitation strategies showed a significantly higher sand content than the soils of reference, with a maximum of 646 and 756 g kg⁻¹ respectively in 284 285 COAL and TDT (p-value =0.013). Lead concentration was significantly higher in the Technosols resulting from CPE rehabilitation strategy, with mean contents of 271.2 and 282.3 mg kg⁻¹ respectively 286 found in Technosol COAL and EMB (p-value = 0.018). Finally, the total potassium content of the 287 soils was significantly higher in the REF soils of reference with a mean of 6739 and 6516 mg kg⁻¹ in 288 sub-sites MEAD and WOOD respectively (p-value = 0.007), which is two to five times higher than the 289 mean concentration found in CPE and ITS. All soils were neutral to slightly alkaline with pH ranging 290

from 7.5 for MEAD and COAL to 8.4 for TDT, the observed differences differing statistically if we 291 accept a threshold alpha of 0.051. While the other parameters did not differ significantly between 292 293 rehabilitation strategies, some of them exhibit high variability between soils sampled. The lowest C:N ratio was 26 and was found in MEAD reference soil, while it reached a mean of 84 in Technosol 294 COAL. Nitrogen content varied from 1.3 to 8.4 g kg⁻¹ in Technosols TDT and EMB respectively. 295 Metals showed a wide variability between soils, even within a given rehabilitation strategy. Zinc 296 297 especially varied from 284 in COAL to 825 in EMB Technosol. A very low bulk density was 298 measured in the Technosol COAL with a mean value of 0.69.

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300 *3.2 Soil fauna communities on the studied area*

Overall, 50,236 organisms were sampled in the study. The density per m² or the number of organisms within the 3 traps of each sub-site and for each soil zoological group collected are shown in Table 2 for litter-soil dwelling macrofauna and micro-arthropods (sampled with TSBF and corer methods), and in Table 3 for soil surface dwelling organisms (sampled with pitfall traps). The indices calculated for the three types of soil fauna communities are shown in Table 4.

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307 *3.2.1 Litter and soil-dwelling communities*

308 Quantitatively, a similar soil fauna diversity was found for the two rehabilitation strategies and for the 309 soils of reference. The density of macro-organisms in the litter and soil-dwelling community found in the study area varied between 230.4 \pm 209.7 and 483.2 \pm 203.1 individuals m⁻² in TDT and MEAD 310 respectively and the density of micro-arthropods varied between 21862 ± 6661 and 105232 ± 77889 311 individuals m⁻² in WOOD and COAL respectively and no statistical differences were found between 312 the rehabilitation strategies (Table 2). The maximum mean of Isopoda, Diplopoda, larvae of Diptera 313 314 and Lepidoptera and Lumbricina were found in REF (WOOD and MEAD). The maximum mean of Chilopoda, Protoura, Mesostigmata and Oribatida were found in Technosols COAL or EMB. Finally, 315 316 the maximum mean number of Araneae, Collembola, and Coleoptera were found in Technosols TDT

or CONS. Yet, out of the 19 taxonomical groups found in the studied soils, only the total number of 317 Lumbricina varied significantly with the nature of the substrate used in soil rehabilitation strategy (p-318 value = 0.034). The number of earthworms varied from 0 individuals m^{-2} in the EMB Technosol to 319 201.6 ± 165 individuals m⁻² in the MEAD soil of reference. Significantly more earthworms were found 320 in the soils of reference than in the Technosols following CPE and ITS rehabilitation strategies. 321 Epigeic earthworms were found in one Technosol resulting from each rehabilitation strategy. Only 3.2 322 \pm 7.2 individuals m⁻² were found in COAL Technosol and none were found in the three other 323 Technosols. Finally, only 9.6 ± 8.8 endogeic earthworms were found per square meter in the CONS 324 325 Technosol, while none were found in the three other Technosols.

No statistical differences were found for the Shannon index, the zoological richness and the trophicrichness (Table 4).

The litter and soil-dwelling community exhibited trophic differences across the studied sub-sites. The proportion of macro-zoophages was higher in CPE sub-sites than in ITS and REF, with respectively 66 \pm 17 and 57 \pm 18% of zoophages in the COAL and EMB Technosols (p-value = 0.048). In the MEAD and WOOD soils of reference, both the density and the proportion of macro-geophages were higher than in the Technosols resulting from the two rehabilitation strategies (p-value = 0.034 and 0.017 respectively).

334 *3.2.3 Soil surface-dwelling invertebrates*

Within the studied area, the number of macro-organisms collected on the soil surface by pitfall traps 335 method varied significantly between the different rehabilitations' strategies (p-value = 0.029). The 336 highest number of soil surface macro-organisms in the coking plant area was found in ITS 337 rehabilitation. No statistical differences were found for the mean number of micro-arthropods sampled 338 in pitfall traps. Quantitatively, only the total number of Hymenoptera varied significantly between the 339 340 rehabilitation strategies (p-value = 0.040). The highest number of Hymenoptera were sampled in the 341 two Technosols resulting from the ITS rehabilitation and most organisms belonged to the Formicidae 342 family (Table 3).

The trophic composition of the soil surface-dwelling invertebrate community varied across the 343 344 different rehabilitation strategies (Table 4). The proportion of saprophages sampled at the surface of 345 the Technosols from the CPE rehabilitation strategy was higher (p-value = 0.048) than the proportion of saprophages sampled at the surface of the Technosols that were subject to the ITS rehabilitation 346 strategy. The lowest proportion of soil surface-dwelling zoophages were sampled at the Technosol of 347 ITS rehabilitation 0.040). 348 the strategy (p-value =

349 3.3 Relationship between soil fauna communities' indices and soil physico-chemical characteristics

350 A co-inertia analysis was conducted in order to test the co-structure between the abiotic parameters 351 and the soil fauna community indices (Fig. 4) for the four Technosols and two soils of reference. The 352 first two axes of the co-inertia represented 87.2% of the total co-inertia and a significant co-variation 353 between the abiotic and biotic data sets was revealed by the Monte Carlo test (RV = 0.58, p-value = 354 0.001). For a given rehabilitation strategy (CPE, ITS) or for the soils of reference), the centroids of the 355 two data sets (abiotic and biotic) were close to each other, indicating a relationship between the soil 356 abiotic parameters and the fauna indices (Fig. 3a). The first axis of the co-inertia analysis explained 357 51.2% of the total inertia and discriminated the two soils of reference from the four Technosols (Fig. 3b). For the litter and soil dwelling community, the soils with the highest fertility parameters, 358 corresponding to the two soils of reference, exhibited the highest soil fauna trophic and zoological 359 360 richness, higher Shannon indexes and the highest densities of macro-saprophages, the lowest density of micro-arthropods and the higher proportion of geophages in the community. The second axis of the 361 co-inertia analysis explained 36.0% of the total inertia and mostly discriminated between the 362 Technosols resulting from the two different rehabilitation strategies. Technosols COAL and EMB 363 364 (CPE strategy) were characterized by soil contamination variables such as soil total Zn, Pb and Cd 365 concentration, DTPA exchangeable Zn and Pb concentration, and, organic matter related variables, such as organic carbon content, and total nitrogen content. These variables co-varied positively with 366 367 the proportion of predators in the soil and at the soil surface and with the proportion of saprophages of 368 the community sampled at the soil surface. The Technosols TDT and CONS that resulted from ITS rehabilitation strategy were characterized by high soil pH values and lower metal concentrations and 369 370 metal availability than the CPE Technosols. These variables co-varied with the highest soil surface fauna activity, as attested to by the number of both macro-organisms and macro-arthropods sampled in 371 372 pitfall traps.

374 *4*. **Discussion**

375 *4.1 Technosol physico-chemical characteristics according to the rehabilitation strategy*

376 Our results on the physico-chemical characteristics of the soils from the former coking plant of 377 Homécourt support our hypothesis of a higher soil fertility in ITS than from the CPE rehabilitation 378 strategy. Technosols from the area have a lower physico-chemical quality that the REF soils of 379 reference. Our results showed that the rehabilitation methods have in fact, led to pronounced abiotic 380 parameter heterogeneity at the site scale, because of the difference between the studied Technosols. As 381 expected, the rehabilitation of the former coking plant area has resulted in sandy Technosols with significantly less clay content than the soils of reference. Indeed, Technosols of industrials areas often 382 383 exhibit a high proportion of coarse elements resulting in low fine earth contents (Burghardt, 1993) 384 associated with the nature of the industrial residues, wastes, or other technogenic substrates used (Burghardt, 1994). The high pH value that we measured in Technosols is in accordance with pH 385 values of former industrial or urban areas, which are usually neutral to slightly alkaline (Galvín et al., 386 387 2012; Joimel et al., 2016; Morel et al., 2005) as the result of the high carbonate content found in 388 building blocks (Burghardt, 1994).

Both CPE and ITS rehabilitation strategies reduced the total potassium content in the Technosols in comparison with the soils of reference, but this was not the case for the exchangeable K that we measured. According to Vincent et al. (2018), exchangeable potassium might be an important limiting factor for plant growth on derelict soils of industrial areas, but the exchangeable K found in the studied Technosols and the soils of reference is within range of the content expected to be found in the Lorraine region (French monitoring network RMQS).

The highest proportion of soil organic carbon in CPE in comparison with the other Technosols studied and soils of reference, with respectively 40 and 50 %, found in EMB and COAL Technosols, might result from the charcoal and other coal by-products used as embankments for the leveling of the area (Biache et al., 2013). The proportion of organic carbon contents above 12% that we measured in the CONS Technosol, most likely resulted from the compost used during the initial settlement of the 400 Technosol (Cortet et al., 2013). Organic carbon is often used as an indicator of soil fertility, as it contributes to the nutrient pool, formation of stable aggregates, water retention or cation exchange 401 402 capacity (Fardeau, 2014; Golchin et al., 1994), especially in nutrient-poor systems (Tiessen et al., 403 1994). Yet, in our study, organic carbon is not correlated with other fertility parameters such as clay content, water holding capacity or cation exchange capacity, but covaried positively with metal 404 405 contamination. The co-variation of organic carbon with Technosol metals that we found on the 406 Technosols resulting from CPE rehabilitation might be explained by the presence of coal by-products 407 in the soil (Colombini et al., 2020) that are known to have a high sorption capacity with metals (Zevenbergen et al., 1999). The presence of polycyclic aromatic hydrocarbons as a source organic 408 409 pollutants that have a high C content and are usually found in soils close to coking plants in addition to 410 metals (Biache et al., 2013; Liu et al., 2013; Rachwał et al., 2015; Yuan et al., 2014) might be another 411 explanation for this correlation. Within a given Technosol, various sources of organic carbon might 412 exist and can be exogenous, such as coal-products, fly ashes or compost input or neo-formed such as 413 leaf litter, living and decaying roots or fauna feces (Colombini et al., 2020). Thus, the proportion of 414 each different C source is to be considered and organic carbon content per se does not necessarily inform on Technosol fertility. The use of coking plant by-products in the CPE rehabilitation strategy 415 have led to the lowest soil chemical quality with total lead and zinc respectively 10 and 7 times higher 416 417 in EMB Technosol than in the geochemical background of the region (Darmendrail et al., 2000). 418 Nevertheless, these concentrations remain relatively low compared with soils affected by zinc smelter or iron industry deposits. For example, Gillet and Ponge (2002) recorded 34800 mg kg⁻¹ and 5840 mg 419 kg⁻¹ of Zn and Pb respectively in soils surrounding an active zinc smelter, while Huot et al. (2013) 420 measured 24800 and 21100 mg kg⁻¹ DW of Zn and Pb respectively in a Technosol of a former settling 421 422 pond in the Lorraine region. Finally, a Zn concentration 2.5 times higher in the WOOD Cambisol than 423 the mean of the geochemical background (Darmendrail et al., 2000) was found, which might result from coal fly ashes aerial deposition during the coking plant's activity (Nelson Beyer et al., 1985), as 424 425 confirmed by micromorphological observations (Colombini et al., 2020).

427 4.2 Taxonomical and trophic composition of Technosol fauna communities according to the
428 rehabilitation strategy

429 On the studied former coking plant, the densities of litter and soil-dwelling organisms (sampled in 430 monoliths) were relatively low, regardless of the rehabilitation strategy. The total macrofauna and 431 micro-arthropods densities were lower in the woodland soil of reference than in comparable temperate 432 deciduous forests (Deleporte and Tillier, 1999; Schaeffer and Shauermann, 1990). However, litter and 433 soil-dwelling macrofauna densities sampled in soils from the coking plant were similar to those 434 already observed in highly anthropized soils with natural succession (Huot et al., 2018; Nahmani and 435 Lavelle, 2002), or post-mining sites rehabilitated with an alder plantation (Frouz et al., 2007a).

The number of earthworms found for the meadow (MEAD) soil of reference in our study was comparable with the density of earthworms found in other meadows in temperate regions (Curry, 1989), while it was relatively low in the woodland (WOOD) soil of reference of the studied former coking plant compared to the density in other woodlands, where earthworm density can reach more than 600 ind m⁻² in temperate regions (Cardinael et al., 2019; Lavelle and Spain, 2001).

441 We found a high proportion of macro-zoophages (mostly Chilopoda) in the litter and soil-dwelling 442 community and a high proportion of soil surface zoophages and saprophages (Araneae, Isopoda and Diplopoda respectively) in CPE Technosols resulting from the embankments of coking plant by-443 444 products. Similar edaphic community composition has been reported in Technosols composed of 445 dumping from settling ponds with a similar spontaneously developed vegetation (Huot et al., 2018). 446 This soil fauna community composition is similar to the soil fauna community found in the moder 447 form of humus, but with lower densities of both micro-arthropods and macro-saprophages such as 448 Diplopoda and Isopoda (Salmon et al., 2006).

In comparison, the community of the two ITS Technosols was mostly composed of a high number of organisms belonging to a few taxonomic and trophic groups as revealed by the number of macroorganisms and collembola sampled in pitfalls traps and by the low Shannon index found in these two Technosols. Such community compositions indicate an early stage of colonization of reclaimed degraded area by the soil fauna (Frouz et al., 2001; Hedde et al., 2018) and occurred in the most recent rehabilitation strategy of the studied site. Micro-arthropods are usually efficient colonizers of Technosols during their early evolution stages (Frouz et al., 2001; Joimel et al., 2018) and might benefit from a low top-down regulation according to the small proportion of epigeic zoophages that we found in the pitfall traps on the TDT and CONS Technosols. When earthworms have not yet colonized reclaimed post-industrial soils, one might well expect a lower bottom-up regulation of the mesofauna community, resulting from the feeding activity of earthworms (Ponge, 2003), which might lead to high mesofauna taxonomic diversity in industrial soils (Joimel et al., 2017).

461

462 4.3 Relationship between soil fauna community indices and soil physico-chemical characteristics

463 Our results showed that only the soils of reference, with higher physical (clay content) and chemical (lower Zn and higher K contents) quality, supported an earthworm's population comparable to similar 464 ecosystems (Cluzeau et al., 2009; Curry, 1989). The sandy texture or the high metal concentration 465 466 might explain the low density or the absence of anecic and endogeic earthworms in the Technosols, as described in many other moderately to highly-polluted soils (Bradshaw, 1983; Gillet and Ponge, 2002; 467 468 Nahmani and Rossi, 2003). While Eijsackers (2010), showed that earthworms are capable of colonizing a wide range of habitats related to different rehabilitation strategies, earthworm populations 469 470 are often reported to be low if not absent from sandy soils resulting from industrial activities 471 (Emmerling and Paulsch, 2001).

ITS Technosols resulting from the depollution of the industrial substrate presented a gap between high 472 number of organisms in pitfall traps and a low soil trophic richness, thus evidencing the fact that the 473 rehabilitation promoted the recovery of the soil surface community rather than the edaphic one, in 474 475 terms of both taxonomical and trophic diversity. While the addition of organic matter has been reported to enhance the colonization of degraded soils by earthworms (Emmerling and Paulsch, 2001), 476 477 this was not the case for the CONS Technosol that received green waste compost. In this same 478 Technosol, Hedde et al. (2018) found more earthworms in the early stage of colonization indicating 479 that the soil substrate did not fit their long term establishment. The increasing abundance of 480 Formicidae that we found in the soil surface community in comparison with their study might 481 highlight the development of a top-down regulation of earthworm populations by ants (Passera and

482 Aron, 2005). On the other hand, the populations of Diptera and Coleoptera larvae seems to have 483 established more durably on CONS. Investigating their interactions with Technosol substrates after 484 rehabilitation would be of interest to specify interactions between soil fauna and Technosols functions. 485 For example, Bibionidae larvae are known to be involved in the development of the upper soil layer in 486 the latter stage of a post-mining chronosequence (Frouz et al., 2001).

On CPE Technosols resulting from the embankments of coking plant by-products, the development of 487 488 a moder humus form fauna community is in agreement with latter stage of succession found in highly 489 anthropized soils such as Technosols resulting from industrial settling pond activity (Huot et al., 490 2018). This result indicates that, additionally to the substrate used in rehabilitation, the development of the vegetation and more precisely the litter layers might be a driver of the soil fauna community on 491 Technosols and would be in agreement with studies conducted on post-mining sites (Frouz et al., 492 493 2007b; Mudrák et al., 2010). Recently, Colombini et al., (2020) showed that most of soil fauna 494 saprophage activity on the CPE Technosols resulting from the embankments of coking plant by-495 products, occurred in the neo-formed organic layers on the soil surface, with footprints of enchytreids, collembolas, isopods and diplopods feces. Action of saprophagous mites was revealed by feces in 496 497 decaying roots and woody fragments in the neo-formed organic layer and in roots within the 498 technogenic layer (Colombini et al., 2020), thus indicating that mites are less sensitive to the nature of 499 the embankments than previously- cited organisms.

500 4.4 Guidance for rehabilitation of post-industrial sites

501 The low number of earthworms and consequently of geophages that we found in the Technosols 502 resulting from the two rehabilitation strategies in the studied site, highlights the importance of the 503 nature of the substrate to be used in post-industrial area rehabilitation scenario. Technosol 504 rehabilitation should focus on fast recovery of soil functions, promoting ecosystem services and 505 biological diversity. As suggested by our results, Technosols' physico-chemical properties that were 506 highly interrelated with the quality of the substrates used in a rehabilitation scenario may limit the 507 colonization of edaphic fauna, such as endogeic and anecic earthworms that play an important role in 508 the formation of soil structure (Amezketa, 1999; Blouin et al., 2013; Foster, 1991). Earthworms, as 509 ecosystem engineers, play an important role in nutrient cycling, soil structure formation and stabilization (Angst et al., 2017, 2019; Pey et al., 2014; Vidal et al., 2019) and can limit the erosion of 510 511 polluted soils (Blouin et al., 2013; Scullion and Malik, 2000). The re-establishment of local earthworm populations should be one of the goals of post-industrial area, and can be achieved with the addition of 512 513 organic carbon at the initial stage of the rehabilitation. Studies show that direct input of organic matter 514 through organic waste materials or indirect input through neo-formed litter resulting from afforestation 515 can improve the community and activity of earthworms on open-cast coal mining sites (Emmerling 516 and Paulsch, 2001; Frouz et al., 2006). A low number of earthworms may alter the incorporation of 517 soil surface organic matter, such as tree litter. Indeed, soil surface organic matter accumulation has 518 been evidenced by many authors in different anthropized ecosystems such as meadow or deciduous forest with high metal concentrations (Gillet and Ponge, 2002; Nahmani and Rossi, 2003) and might 519 explain the soil fauna community resembling the moder humus form that we found in the COAL and 520 EMB Technosols. This community composition might be a general pattern of intermediate stages of 521 vegetation for Technosols with extreme soil physico-chemical characteristics, since it has been 522 523 reported in many studies related to metal -contaminated soils (Grelle, 2000; Huot et al., 2018).

The overall ability of soil surface saprophages to colonize Technosols regardless of their physicochemical characteristics, when organic matter is added, could represent a promising way to enhance the recovery of soil processes such as nutrient cycling and bioturbation. It can be achieved by a coupled inoculation of local soil surface saprophages and organic matter, as has previously been done in post-mining sites (Frouz et al., 2007b).

Yet, in order to obtain a better insight into the relationships between the fertility parameters and the soil fauna communities of the Technosols, the nature of the soil organic carbon should be taken into account to discriminate between both the sources and the role of technogenic and neo-formed organic matter and their interaction with the soil saprophages.

Finally, the creation of ecological corridors at the site scale could improve the rate of earthworms and other saprophagic organism colonization, which usually occurs within 5 years for pioneer earthworm species (Snyder and Hendrix, 2008), but would not be sufficient if the nature of the soil substrates used does not fit their long-term requirements. The development of different trophic networks over

time could induce a top-down or bottom-up regulation, but a frequent sampling would be needed in 537 order to prove a competitive exclusion of certain taxa. Moreover, different colonization dynamics may 538 539 occur as a function of the dispersal strategy. Indeed, the presence of Diptera and Coleoptera larvae on 540 Technosols, even in earlier stages of colonization can be explained by the dispersion of flying adults, while endogeic earthworms would need suitable ecological corridors to colonize new environments 541 (Eijsackers, 2010). Thus, studies of soil fauna functional traits that improve the colonization success 542 543 might help us to improve our understanding of the actual diversity of the soil fauna community on 544 post-industrial sites. Finally, the gap between the high level of activity of epigeic organisms that we 545 found on Technosols and the low trophic diversity of the litter and soil-dwelling communities suggest 546 that the soil fauna community on a former coking plant is driven rather by soil physico-chemical 547 properties than by colonization constraints. Our results suggest that the physico-chemical and 548 biological properties of the studied Technosols are evolving. Following the dynamics of the physicochemical and biological properties of the Technosols studied in future years to decades could be help 549 to improve our understanding of their evolution pathways and lifespan. 550

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1 Figures & Captions

2

Fig. 1. Localization of the studied former coking plant and its 6 sampled sub-sites: 4 Technosols and 2
soils of reference. WOOD: woodland, MEAD: meadow, COAL: Spolic Technosol composed of coal
embankments, EMB: Spolic Technosol with multiple embankments from a dismantled coking plant,
TDT: Technosol composed of thermally-treated industrial soil, CONS: constructed Technosol
composed of a mixture of paper mill sludge, thermally-treated industrial soil and green waste compost.

8

Fig. 2. Sampling design followed for the characterization of (i) the physico-chemical quality and (ii) the
fauna communities of the 4 Technosols and 2 soils of reference (N=5, except for COAL, N=4). Numbers
on the right part of the figure indicate the sampling procedure: (1) Sampling for chemistry analyses. (2)
Sampling of a soil corer for the bulk density assessment. (3) Litter and soil-dwelling macrofauna
sampling. (4) Soil corer for the litter and soil-dwelling micro-arthropod sampling. (5) Pitfall traps for
the sampling of the soil surface-dwelling invertebrates.

15

Fig. 3. Principal Component Analysis (PCA) obtained for the abiotic parameters measured for the 4 16 Technosols and 2 soils of reference. Thicker symbols represent the barycenter for the 10 samples of the 17 REF soils of reference (WOOD and MEAD) or the ITS (Technosols COAL and EMB) and CPE 18 19 (Technosols CONS and TDT) strategy (9 samples for CPE). Pb = lead concentration; Zn = zinc20 concentration; K = potassium concentration; Mg = magnesium concentration; CEC = cation exchange capacity; WHC = water holding capacity; Ca = calcium concentration; Cd = cadmium concentration; 21 C_{org} = organic carbon content; N_{tot} = nitrogen content; P_2O_5 = available P concentration; DTPA: 22 23 Diethylene triamine penta-acetic acid extractible metals.

24

Fig. 4. Co-inertia analysis between the two PCA on abiotic parameters and soil fauna indices. Axis1 =
first projected axis, Axis2: second projected axis. A) Projection of the Technosols that followed 2

rehabilitation strategies and the soils of reference on the two first axes, grey squares represent centroids 27 for soil fauna indices used in the PCA and blue dots represent centroids for abiotic parameters. Large 28 squares and dots are centroids for the 10 points (9 points for CPE). B) Projection of the 2 soils' 29 classification (Technosols or Cambisol) on the two first axes, grey squares represent centroids for soil 30 31 fauna indices used in the PCA and blue dots represent centroids for soil abiotic parameters. Large 32 squares and dots are centroids for the 10 points (9 points for CPE). C) Scatter plot of the co-inertia, 33 abiotic parameters and fauna indices from Technosols and soils of reference are encoded, see Tables 1 to 4 for correspondences. 34



36 Fig. 1.









40 Fig. 3.



43 Fig. 4.

1 Tables & captions

2 Table 1.

3 Soil abiotic parameters of the 6 sub-sites (4 Technosols + 2 soils of reference) studied (mean \pm sd, n = 5 except for COAL n = 4). WHC: Water Holding Capacity;

4 Corg: organic carbon concentration; Ntot: nitrogen concentration; CEC: cation exchange capacity; P2O5: available P concentration; Cd: cadmium concentration;

5 Pb: lead concentration; Zn: zinc concentration; K: potassium concentration; Mg: magnesium concentration, Exch: exchangeable; DTPA: Diethylene triamine 6 penta-acetic acid extractible metals. Different letters in the same row indicate a significant difference between means tested with one-way nested ANOVA

penta-acetic acid extractible metals. Different letters in the same row indicate a significant difference between means tested with
between REF, CPE and ITS, p-value < 0.05.

Soil classification	Cam	bisol	Technosol							
Rehabilitation strategy	Soils of refe	rence (REF)	Coking plant emb	ankments (CPE)	Industrial trea	Industrial treated soil (ITS)				
Site	MEAD	WOOD	COAL	EMB	TDT	CONS	p-value			
WHC (%)	26.2 ± 1	26 ± 0.9	19.8 ± 3.5	21.2 ± 2.1	17.2 ± 1.9	22.1 ± 2.1	0.094			
Bulk density (g cm ⁻³⁾	1.26 ± 0.22	0.97 ± 0.14	0.69 ± 0.07	0.98 ± 0.20	1.15 ± 0.25	1.11 ± 0.21	0.289			
pH	7.5 ± 0.2	7.9 ± 0.1	7.5 ± 0.3	7.6 ± 1.4	8.4 ± 0.1	8.2 ± 0.1	0.051			
C _{org} (g kg ⁻¹)	100 ± 17	186 ± 36 b	507 ± 62	410 ± 140	a 46 ± 12	122 ± 72 b	0.016			
N _{tot} (g kg ⁻¹)	3.8 ± 0.4	6.2 ± 0.96	6.7 ± 1.5	8.4 ± 2.5	1.3 ± 0.2	3.6 ± 1.2	0.118			
C/N	26 ± 4	31 ± 9	84 ± 26	49 ± 9	38 ± 15	32 ± 8	0.153			
CEC (cmolc kg ⁻¹)	31 ± 2	47 ± 2	18 ± 5	28 ± 18	12 ± 4	26 ± 14	0.216			
P_2O_5 (mg kg ⁻¹)	14 ± 5	11 ± 2	8 ± 1	10 ± 5	5 ± 2	29 ± 12	0.753			
Clay (g kg ⁻¹)	495 ± 31	507±19 a	78 ± 21	112 ± 41	b 64 ± 9	120±19 b	0.001			
Sand (g kg ⁻¹)	214 ± 40	157 ± 17 b	646 ± 65	640 ± 97	a 756 ± 21	577±39 a	0.013			
Cd _{tot} (mg kg ⁻¹)	0.5 ± 0.0	0.9 ± 0.2	1.0 ± 0.4	2.9 ± 0.8	3.0 ± 1.5	1.7 ± 0.5	0.329			
Pb _{tot} (mg kg ⁻¹)	34 ± 3	72±14 b	271 ± 190	282 ± 102	a 67±13	93 ± 20 b	0.018			
Zn _{tot} (mg kg ⁻¹)	148 ± 11	297 ± 80	284 ± 70	825 ± 431	224 ± 36	383 ± 134	0.449			
K _{tot} (mg kg ⁻¹)	6739 ± 231	6516±3029 a	1696 ± 327	1287 ± 225	b 2463 ± 269	3128 ± 1415 b	0.007			
Mg _{tot} (mg kg ⁻¹)	5623 ± 1739	4778 ± 84	5335 ± 1237	3782 ± 627	6440 ± 518	5660 ± 997	0.307			
Mg _{Exch} (cmolc kg ⁻¹)	2.4 ± 0.3	0.7 ± 0.1	0.9 ± 0.4	1.5 ± 1.1	0.3 ± 0.1	1.8 ± 1.3	0.883			
K _{Exch} (cmolc kg ⁻¹)	0.7 ± 0.0	0.6 ± 0.2	0.3 ± 0.0	0.4 ± 0.1	0.4 ± 0.1	0.8 ± 0.1	0.362			
Cd _{DTPA} (mg kg ⁻¹)	0.1 ± 0.0	0.2 ± 0.0	0.2 ± 0.1	0.6 ± 0.4	0.1 ± 0.0	0.1 ± 0.0	0.444			
Pbdtpa (mg kg ⁻¹)	4.1 ± 0.3	9.6 ± 0.9	18.9 ± 15.5	34.9 ± 13.5	3.1 ± 0.4	6.6 ± 1.5	0.089			
Zn _{DTPA} (mg kg ⁻¹)	3.9 ± 0.6	18.4 ± 0.6	22.0 ± 11.0	109.4 ± 43.3	3.4 ± 0.4	12.9 ± 6.9	0.339			
Base staturation	1.02 ± 0.02	0.86 ± 0.02	1.06 ± 0.03	0.99 ± 0.05	0.99 ± 0.15	0.99 ± 0.05	0.534			

10 Table 2.

11 Density (mean number of individuals per square meter ± standard deviation) of zoological groups for litter-soil dwelling macrofauna and micro-arthropods

12 communities sampled in the 4 Technosols and 2 soils of reference (n = 5 except for COAL n = 4). Trophic types were determined based on Salmon et al.,

13 (2006). Zoo: zoophage, mi: microphage, geo: geophage, phy: phytophage, sapro: saprophage. PCA code refers to the code used in both PCA and co-inertia

14 analyses for the fauna indices. Different letters in the same row indicate a significant difference between means tested with one-way nested ANOVA, between

15 REF (MEAD and WOOD), CPE (COAL and EMB) and ITS (TDT and CONS), p-value < 0.05.

Soil classification			Can	nbisol	Technosol				
Rehabilitation strategy			Soils of reference (REF)		Coking plant emba	ankments (CPE)	Industrial treate	Industrial treated soil (ITS)	
Taxonomic level	Zoological group	Trophic type	MEAD	WOOD	COAL	ЕМВ	TDT	CONS	p- value
Class	Isopoda	Sapro	6.4 ± 8.8	44.8 ± 26.3	9.6 ± 21.5	6.4 ± 14.3	0 ± 0	0 ± 0	0.375
Class	Chilopoda	Zoo	16 ± 16	92.8 ± 68.3	118.4 ± 50.1	92.8 ± 59.2	12.8 ± 28.6	0 ± 0	0.128
Class	Diplopoda	Sapro	0 ± 0	54.4 ± 36.8	12.8 ± 17.5	44.8 ± 30.8	51.2 ± 88.7	12.8 ± 13.4	0.987
Class	Collembola	Sapro/mi	11237 ± 3603	8383 ± 2433	9555 ± 2683	6115 ± 2368	23263 ± 8998	6778 ± 4739	0.619
Class	Protoura	Sapro/mi	0 ± 0	51.0 ± 114.0	1401.4 ± 1348.3	1070.2 ± 1620.5	1248.5 ± 2511.8	25.5 ± 57.0	0.307
Order	Araneae	Zoo	32 ± 32	16 ± 16	41.6 ± 31.2	44.8 ± 34.7	35.2 ± 61.3	54.4 ± 26.8	0.442
Order	Mesostigmata	Zoo	2930.2 ± 2190	1732.6 ± 545	5478.2 ± 3661.5	7083.4 ± 6686.7	2930.2 ± 1164.2	1503.3 ± 520.6	0.106
Sub-order	Oribatida	Sapro	17963 ± 7667	11746 ± 4735	90199 ± 58942	26958 ± 16035	62961 ± 11811	36411 ± 37765	0.381
Order	Diptera larvae	Sapro	28.8 ± 39.8	12.8 ± 13.4	22.4 ± 18.2	6.4 ± 8.8	9.6 ± 14.3	22.4 ± 26.8	0.831
Order	Lepidoptera larvae	Phy	12.8 ± 20.9	6.4 ± 8.8	6.4 ± 8.8	0 ± 0	0 ± 0	0 ± 0	0.228
Sub-order	Homoptera	Phy	0 ± 0	0 ± 0	3.2 ± 7.2	3.2 ± 7.2	0 ± 0	3.2 ± 7.2	0.443
Order	Total Coleoptera	Zoo/phy	185.6 ± 107.1	51.2 ± 28.6	64 ± 54.3	67.2 ± 36.5	76.8 ± 61.3	294.4 ± 99.7	0.577
Order	Coleoptera larvae	Zoo/phy	96 ± 69.7	16 ± 22.6	6.4 ± 14.3	25.6 ± 33.2	3.2 ± 7.2	64 ± 134.3	0.669
Family	Curculionidae larvae	Phy	6.4 ± 8.8	6.4 ± 8.8	0 ± 0	3.2 ± 7.2	6.4 ± 14.3	44.8 ± 41.4	0.391
Family	Elateridae larvae	Phy	9.6 ± 14.3	0 ± 0	0 ± 0	16 ± 19.6	19.2 ± 26.3	83.2 ± 111.7	0.305
Family	Carabidae	Zoo	25.6 ± 21.5	0 ± 0	35.2 ± 34.7	0 ± 0	35.2 ± 44.4	60.8 ± 51.1	0.319
Family	Staphylinidae	Zoo	28.8 ± 26.3	12.8 ± 17.5	9.6 ± 14.3	16.0 ± 22.6	6.4 ± 8.8	9.6 ± 14.3	0.394
Family	Staphylinidae larvae	Zoo	0 ± 0	16 ± 19.6	3.2 ± 7.2	3.2 ± 7.2	3.2 ± 7.2	12.8 ± 13.4	0.784
Sub-order	Total Lumbricina	Sapro/geo	201.6 ± 165	118.4 ± 47.5 a	16 ± 16	0±0 b	6.4 ± 14.3	9.6±8.8 b	0.034
Earthworm	Epigeic	Sapro	70.4 ± 55	28.8 ± 13.4	12.8 ± 17.5	0 ± 0	6.4 ± 14.3	0 ± 0	NT
ecological	Anecic	Geo/sapro	9.6 ± 14.3	28.8 ± 34.7	3.2 ± 7.2	0 ± 0	0 ± 0	0 ± 0	NT
categories	Endogeic	Geo	121.6 ± 106.5	60.8 ± 20.9	0 ± 0	0 ± 0	0 ± 0	9.6 ± 8.8	NT

18 Table 3.

19 Activity of surface soil dwelling invertebrates (mean number of individuals collected in 3 traps ± standard deviation) and zoological richness (mean number of

20 zoological groups \pm standard deviation) (n = 5 after gathering 3 pitfalls per sampling square). Trophic types were determined based on Salmon et al., (2006).

21 Zoo: zoophage mi: microphage, phy: phytophage, sapro: saprophage. Different letters in the same row indicate a significant difference between means tested

22 with one-way nested ANOVA, between REF, CPE and ITS, p-value < 0.05. No test (NT) was performed when less than 5 individuals of a given group were

collected across the 6 sub-sites.

24

Soil classification			Cam	bisol	Technosol					
Rehabilitation strategy			Soils of refere	ence (REF)	Cokinç embankme	g plant ents (CPE)	Industrial treated soil (ITS)			
Taxonomic level	Zoological group	Trophic type	MEAD	WOOD	COAL	EMB	TDT	CONS	p- value	
Class	Collembola	Sapro/mi	251.4 ± 133.8	75.4 ± 40.5	71.0 ± 20.0	61.0 ± 23.4	2246.8 ± 1359.3	220.2 ± 64.0	0.412	
Sub-class	Acari	Zoo/sa	66.0 ± 36.2	20.8 ± 13.3	69.4 ± 59.9	10.6 ± 5.0	60.8 ± 40.1	217.8 ± 62.5	0.462	
Order	Araneae	Zoo	17.0 ± 5.8	2.4 ± 1.8	20.0 ± 4.2	13.4 ± 3.4	23.0 ± 6.1	36.4 ± 4.8	0.275	
Order	Opiliones	Zoo	0 ± 0	1.2 ± 1.1	0.2 ± 0.4	1.4 ± 1.1	4.8 ± 3.6	0 ± 0	0.685	
Class	Isopoda	Sapro	6.0 ± 3.2	3.8 ± 2.2	21.0 ± 22.0	6.4 ± 4.3	3.0 ± 4.5	1.0 ±2.2	0.335	
Class	Diplopoda	Sapro	0.2 ± 0.4	2.6 ± 0.1	3.8 ± 4.1	5.2 ± 1.3	5.6 ± 5.1	7.4 ± 8.8	0.301	
Class	Gastropoda	Phy/sa	13.6 ± 6.3	2.4 ± 0.9	2.0 ± 2.0	2.0 ± 1.0	1.2 ± 1.1	0.4 ± 0.5	0.436	
Sub-order	Homoptera	Phy	4.6 ± 5.4	0.2 ± 0.4	2.8 ± 2.3	0.6 ± 0.9	5.2 ± 2.2	9.2 ± 5.7	0.241	
Sub-order	Heteroptera	Phy	0 ± 0	0.4 ± 0.5	0.2 ± 0.4	0 ± 0	1.8 ± 1.6	0 ± 0	NT	
Order	Total Hymenoptera	Phy	12.6 ± 10.9	5.6 ± 3.6	8.2 ± 8.2	14.2 ± 10.7	259.6 ± 137.8	210.2 ± 272.7	0.040	
Family	Formicidae	Phy/sa	12.4 ± 10.5	4.6 ± 2.6	7.8 ± 7.3	13.0 ± 9.1	258.8 ± 137.0	208.8 ± 271.0	NT	
Order	Lepidoptere larvae	Phy	0 ± 0	0.6 ± 0.4	0 ± 0	1.0 ± 0.7	0.2 ± 0.4	0 ± 0	NT	
Order	Diptera larvae	Sapro	1.4 ± 2.6	1.8 ± 1.9	0 ± 0	1.0 ± 0.7	0.4 ± 0.5	0.6 ± 0.9	0.290	
Order	Coleoptera	Zoo/phy	6.2 ± 2.9	14.2 ± 4.4	5.8 ± 2.6	8.2 ± 4.0	8.2 ± 2.5	22.0 ± 12.5	0.549	
Order	Coleoptera_UND	Zoo/phy	0.4 ± 0.5	0.4 ± 0.9	1.8 ± 2.0	2.0 ± 2.9	0.8 ± 0.4	9.8 ± 1.2	NT	
Order	Coleoptera_larvae	Zoo/phy	0.2 ± 0.4	1.2 ± 0.4	0.2 ± 0.4	2.0 ± 1.0	0.6 ± 0.54	1.0 ± 0.7	NT	
Family	Elateridae	Phy	0.6 ± 0.9	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0.6 ± 0.9	NT	
Family	Chrysomelidae	Phy	0 ± 0	0.2 ± 0.4	0 ± 0	0 ± 0	0.2 ± 0.4	0 ± 0	NT	
Family	Curculionidae	Phy	0 ± 0	0.4 ± 0.5	0.2 ± 0.4	0 ± 0	0.2 ± 0.4	1.8 ± 0.8	NT	
Family	Carabidae	Zoo	3.8 ± 2.6	3.8 ± 2.6	1.0 ± 1.2	0.2 ± 0.4	3.2 ± 1.5	4.0 ± 2.8	0.072	
Family	Staphylinidae	Zoo	1.2 ± 0.8	8.2 ± 3.2	2.6 ± 2.3	4.0 ± 2.2	3.2 ± 2.3	4.8 ± 2.2	0.880	

27 Table 4.

Trophic indices of soil fauna communities: Litter and soil-dwelling community (LSDC): Shannon index, zoological richness, total macrofauna and microarthropods densities, trophic richness, proportion of macro-zoophages, saprophages and geophages, densities of macro-zoophages, saprophages and geophages and densities of micro-zoophages and saprophages. Soil surface-dwelling community (SSDC): trap richness, number of soil surface macrosaprophages, proportion of soil surface saprophages and zoophages, number of macro-organisms and microarthropods sampled in the 4 Technosols and 2 soils of reference studied (mean ± standard deviation) (n = 5 except for COAL n = 4). PCA code refers to the code used in both PCA and co-inertia analyses for the fauna indices. Different letters in the same row indicate a significant difference between means tested with one-way nested ANOVA between REF, CPE and ITS, p-value < 0.05.

Soil classification	Cambisol			Technosol							
		Soils of reference (REF)		-	Coking plant embankments (CPE)		· -	Industrial treated soil (ITS)			
Indices	Fauna community	MEAD	WOOD	-	COAL	ЕМВ		TDT	CONS		p- value
Shannon	macrofauna LSDC	1.38 ± 0.45	2.26 ± 0.25		1.76 ± 0.25	1.35 ± 0.5		0.85 ± 0.68	0.64 ± 0.17		0.150
ZOO_Rich	total LSDC	7.4 ± 1.1	7.8 ± 0.83		7.6 ± 0.9	5.4 ± 1.3		5.2 ± 1.9	6.0 ± 1.0		0.279
Macro_dens (ind m ⁻²)	macrofauna LSDC	483.2 ± 203.1	403.2 ± 160.6		304 ± 143.7	268.8 ± 53.5		230.4 ± 209.7	396.8 ± 86.5		0.248
Micro dens (ind m ⁻²)	microarthropods LSDC	32130 ± 6253	21862 ± 6661		105232 ± 77889	40157 ± 214178		89155 ± 28067	44692 ± 17672		0.420
Troph_Rich	LSDC	6 ± 0.71	6.4 ± 0.89		4.75 ± 0.5	4.6 ± 0.55		4 ± 1	5.4 ± 0.55		0.135
P_Macro_PRED (%)	macrofauna LSDC	15 ± 11	33 ± 11	b	66 ± 17	57 ± 18	а	18 ± 19	29 ± 23	b	0.048
P_Macro_SAPRO (%)	macrofauna LSDC	24 ± 16	41 ± 5		21 ±12	22 ± 15		43 ± 39	12 ± 6		0.771
P_Macro_GEO (%)	macrofauna LSDC	24 ± 12	17 ± 7	а	0 ± 0	0 ± 0	b	0 ± 0	3 ± 3	b	0.017
Dens_M_Sapro (ind m ⁻²)	macrofauna LSDC	35.2 ± 36.5	112.0 ± 57.7		40.0 ± 27.7	57.6 ± 41.7		60.0 ± 81.7	35.2 ± 17.5		0.726
Dens_M_Zoo (ind m ⁻²)	macrofauna LSDC	48.0 ± 46.6	108.8 ± 71.0		160.0 ± 90.5	137.6 ± 51.3		48.0 ± 89.8	54.4 ± 26.8		0.010
Dens_M_Geo (ind m ⁻²)	macrofauna LSDC	201.6 ± 165	118.4 ± 47.5	а	16 ± 16	0 ± 0	b	9.6 ± 8.8	6.4 ± 14.3	b	0.034
Dens_Micro_Sapro (ind m ⁻²)	microarthropods LSDC	29200 ± 7266	21276 ± 7064		100710 ± 57381	33200 ± 16094		86224 ± 13858	43189 ± 39153		0.459
Dens_Micro_Zoo (ind m ⁻²)	microarthropods LSDC	2930 ± 2190	1733 ± 545		5478 ± 3662	7083 ± 6687		2930 ± 1164	1503 ± 521		0.101
TRAP_Rich	SSDC	11.6 ± 0.9	13.9 ± 1.9		11.2 ± 1.3	14.4 ± 0.9		14.6 ± 1.3	12.6 ± 0.9		0.995
TRAP_Nb_M_Sapro (ind trap ⁻¹)	SSDC	7.6 ± 4.3	8.2 ± 4.1		23.5 ± 25.4	12.6 ± 4.6		9.0 ± 7.1	9.0 ± 10.7		0.248
TRAP_P_Sapro (%)	SSDC	12.0 ± 6.6	22.2 ± 7.4	ab	33.4 ± 18.1	23.5 ±6.6	а	3.5 ± 3.6	5.0 ± 6.1	b	0.048
TRAP_P_Pred (%)	SSDC	36.7 ± 14.6	43.1 ± 9.7	а	41.9 ± 14.9	36.5 ± 8.5	а	12.6 ± 6.5	22.3 ± 9.1	b	0.040
TRAP_Nb_Micro (ind trap-1)	SSDC	317.4 ± 146.1	96.2 ± 51.7		140.4 ± 69.6	71.6 ± 21.4		2307.6 ± 1379.7	438.0 ± 123.8		0.329
TRAP_Nb_Macro (ind trap-1)	SSDC	61.6 ± 7.2	35.2 ± 10.1	b	64.0 ± 30.0	54.8 ± 19.4	b	313.0 ± 123.9	287.2 ± 267.4	а	0.029
35											

Graphical abstract



