

Ecological Risk Assessment of Trace Elements Accumulated in Stormwater Ponds within Industrial Areas

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1 Ecological risk assessment of trace elements accumulated in stormwater ponds within
2 industrial areas

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9 quotients

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13 gratefully acknowledged for carefully reviewing the manuscript and for the assistance with the figures.

14

15

16 **Abstract**

17 Stormwater ponds can provide flood protection and efficiently treat stormwater using sedimentation.
18 As the ponds also host aquatic biota and attract wildlife there is a growing concern that the sediment
19 bound pollutants negatively affect aquatic organisms and the surrounding ecosystem. In this study we
20 used three methods to assess the accumulation and the potential ecological risk of 13 different heavy
21 metals and metalloids (e.g. trace elements) including both elements that are frequently monitored and
22 some which are rarely monitored in sediment from 5 stormwater ponds located within catchments
23 with predominately industrial activities. Ecological risk for organisms in the older ponds was observed
24 for both commonly- (e.g. Cd, Cu, Zn) and seldom- (e.g. Ag, Sb) monitored trace elements. The 3
25 methods ranked the degree of contamination similarly. We show that methods usually used for
26 sediment quality assessment in aquatic ecosystems can also be used for screening the potential risk of
27 other trace elements in stormwater ponds and may consequently be useful in stormwater monitoring
28 and management. Our study also highlights the importance of establishing background conditions
29 when conducting ecological risk assessment of sediment in stormwater ponds.

30 **Introduction**

31 Urban stormwater runoff may cause flooding and pollution in downstream watercourses if neither the
32 flow is reduced nor the pollutants are removed (Baekken 1994, Maltby et al 1995; Blecken et al. 2012,
33 Sharley et al. 2016). Today, stormwater is often retained and treated in wetlands or stormwater ponds
34 (also called stormwater detention ponds or urban ponds), either alone or in combination with other
35 techniques (Marsalek et al. 2005; Blecken et al. 2017; Jefferson et al. 2017; Sharley et al. 2017; Crane
36 2019). The type and quantity of pollutants in stormwater runoff which accumulates in a specific
37 wetland or stormwater pond depends on the catchment area, climatic factors, land use and
38 percentage impervious surfaces (Färm and Waara 2005; Casey et al. 2006; Frost et al. 2015; Sjøberg et
39 al. 2016; Blecken et al. 2017; Sharley et al. 2017; Crane 2019).

40 Recently, concern has been raised over the ecological risks to aquatic life in wetlands and stormwater
41 ponds and for the surrounding wildlife caused by the accumulated sediment. For example, Sharley et
42 al. (2017) sampled 98 wetlands in the Melbourne region and found that catchments with >10 %
43 industrial land use were at a greater risk of containing contaminants at values which exceeded
44 ecological guideline values and waste disposal guidelines. Additionally, a study in Minnesota by Crane
45 (2019) found that industrial areas in Minnesota were more contaminating than residential and
46 commercial areas, leading to statistically significant increases in zinc and some organic micropollutant

47 content in the studied stormwater ponds. This suggests that stormwater ponds in catchments with
48 industrial activity may accumulate high levels of trace elements which could pose a significant risk to
49 the ecosystem.

50 In studies of trace element accumulation in stormwater ponds, only a few selected heavy metals (e.g.
51 Cd, Cu, Ni, Pb, Zn) are typically monitored (Färm and Waara 2005; Casey et al. 2007; Egemose et al.
52 2015; Blecken et al. 2017; Sun et al. 2019). However, there are many other potentially ecologically
53 harmful trace element contaminants. Herein, 13 different trace elements representing both
54 commonly- and seldom- monitored trace elements were selected, and their accumulation in five
55 ponds in catchments with industrial activity were studied. In addition, P and S were analysed to
56 determine whether their concentrations correlated with those of the trace elements, and therefore if
57 they could provide overall contamination values. The patterns and relationships between trace
58 elements with P, S and sediment characteristics were analysed using Principal Component Analysis and
59 Classification. Furthermore, three methods which are commonly used for assessing sediment quality
60 in lakes and watercourses but rarely used for stormwater ponds and wetlands were used to assess the
61 degree of contamination and ecological risk. The accumulation of elements in the ponds compared to
62 background levels was assessed using Müller's geoaccumulation index, I_{geo} (Müller 1969), while
63 ecological risk was assessed using Håkanson's Potential Ecological Risk Index (RI) (Håkanson 1980) and
64 by the Risk Quotient Methodology (RQ-method) (European Chemicals Bureau 2003) which nowadays,
65 under REACH, is called Risk Characterisation Ratios (RCRs) (ECHA 2016) and therefore hereafter
66 referred to as the RCR-method. To our knowledge, I_{geo} and RI have not been used to assess risk in
67 stormwater pond sediment. The RCR-method is partly used as sediment guidelines, which in this study
68 serve as PNEC values for calculating RCRs, are frequently compared to measured values when assessing
69 the quality of stormwater sediment (Färm et al. 2003; Andersson et al. 2004; Färm and Waara 2005;
70 Jang et al. 2010; Blecken et al. 2012; Allen et al. 2017; Sharley et al. 2017; Crane 2019). However, RCRs
71 have not been calculated nor evaluated in previous studies. By combining the results from each
72 method, the ponds at the highest risk and the trace elements of highest concern were identified and
73 provided essential guidance for our continued contaminant monitoring.

74 **Materials and Methods**

75 *Study site*

76 The sampled stormwater ponds are in the municipality of Ängelholm in southern Sweden. The
77 characteristics of the ponds and their catchment is presented in Table 1; and their design and sediment
78 thicknesses are presented in Fig. 1. The ponds were designated Pond A and B (DUM 20 and DUM3,

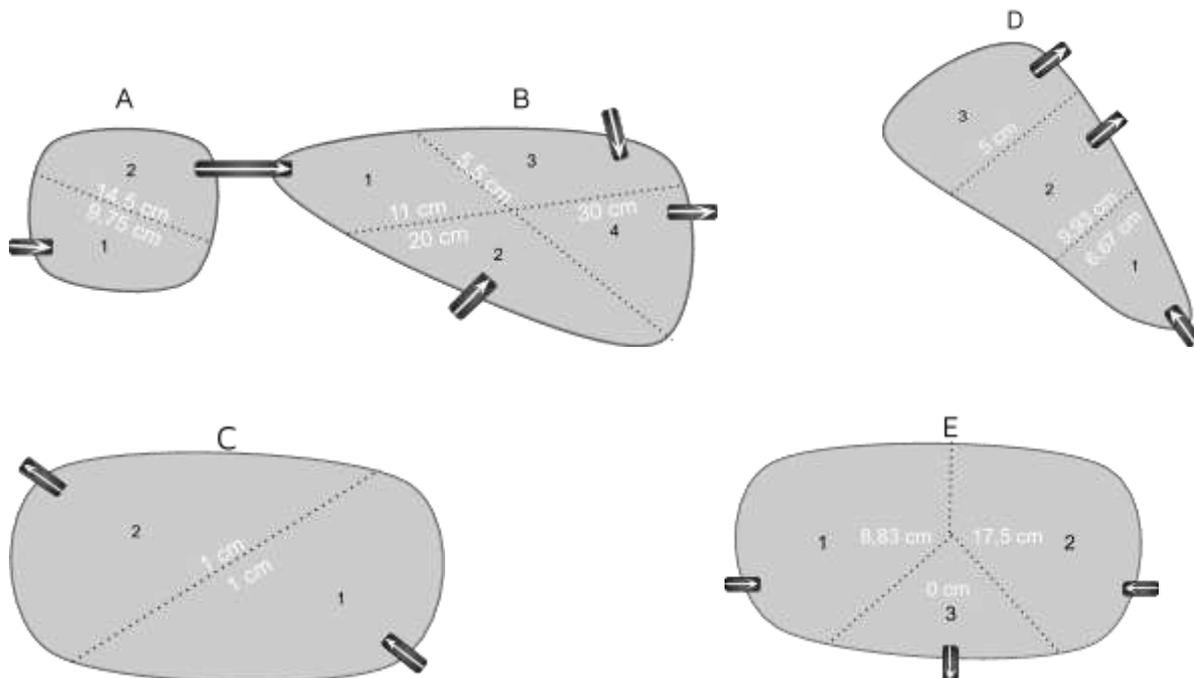
79 Åkerslund 26:3), pond C (DUM 23, Rebbelberga 14:6), pond D (DUM 8, Rebbelberga 19:2) and pond E
 80 (DUM 19, Munka-Ljungby 52:20) and they are located with catchments with mainly industrial activity.
 81 The industrial areas have been developed at different paces from when the respective ponds were
 82 built and consequently the percentage impervious area has been changing over time. The industrial
 83 activity is diverse, and it consists of workshops, shops and storage of building materials including
 84 metals, storage of vehicles for building and construction work, haulage companies, a bus company,
 85 car testing facilities, the major recycling plant in Ängelholm and the power plants of Ängelholm and
 86 Munka Ljungby. Pond A-D are located less than 1 km apart and pond E is located about 5 km east of
 87 the other ponds in the village of Munka Ljungby. The stormwater ponds were designed to prevent
 88 flooding in downstream areas.

89 **Table 1** Characteristics of the investigated ponds. Data obtained from the Municipality of Ängelholm

Pond	Area (m ²) ¹	Catchment area (ha) ²	% Impervious surface ¹	The ratio (in percentage) of pond area to impervious area	Age 2019 (years)
A (DUM 20)	300	6.5	50	0.9	6
B (DUM 3)	950	72.0	50	0.3	17
C (DUM 23)	2000	14.5	12 ¹	1.1	2
D (DUM 8)	1300	44.3	30	1.0	18
E (DUM 19)	500	24.8	70 ²	0.3	2

90 ¹ the impervious surface will increase when the area is fully developed

91 ² the pond was constructed after the development of the industrial area



94 **Fig 1** Design of the ponds and the measured thickness of sediment

95 *Sediment sampling and analysis*

96 Sediments were first sampled in April 2019. Each pond was split into as many sections as there were
97 inlets and outlets (Fig. 1), which enabled the identification of differences in sediment characteristics
98 and pollutant levels within a single pond. Sediment depth and sample collection was conducted using
99 the method described by Blecken et al. (2017). Briefly, sediment samples were taken with a sediment
100 retriever. The depth of the accumulated sediment layer (i.e. only that which was above the clay layer)
101 was measured with a ruler while the extracted core was still in the retriever. After the sediment sample
102 was emptied into a clean bucket, the clay layer was carefully removed, and the remaining sample was
103 mixed thoroughly with the other 3-6 subsamples collected from the same pond section. The number
104 of subsamples taken from each section was dependent on the section area.

105 In August 2019, surface soil around the ponds A+B, C and E and well above possible waterlines was
106 collected to determine background element concentrations in soil which in this area is dominated by
107 kaolinite clay mineral. Subsurface soil was collected randomly using a small clean spoon and then
108 mixed thoroughly before analysis. At this time, a sample from E3 was also taken as it was not possible
109 to retrieve a sample with the core sampler during the first sampling because this part of the pond is
110 covered with macadam. Additionally, although no accumulation of sediment was evident above the
111 macadam, it was possible to collect the material deposited around and under the macadam layer and
112 around the macrophyte roots using a gloved hand in order to avoid sampling the underlying clay layer.

113 The particle size distribution of the samples was determined according to ISO 11277:2009 "Soil quality
114 — Determination of particle size distribution in mineral soil material — Method by sieving and
115 sedimentation". The determination of dry matter was conducted according to Swedish Standard (SS)
116 28113 "Determination of dry matter and ignition residue in water, sludge and sediment". The
117 elemental composition of the samples was analysed with inductively coupled plasma sector field mass
118 spectroscopy (ICP-SFMS). In all cases, samples were digested in concentrated acid. For the elements
119 As, Ba, Cd, Co, Cr, Cu, Hg, Ni, P, Pb, S, V and Zn, samples were heated and mixed in concentrated HNO₃;
120 whereas for Ag and Sb, samples were treated with *aqua regia* (3:1 HCl:HNO₃). General guidelines of
121 the method are found in SS EN ISO 17294-1 "Water quality — Application of inductively coupled plasma
122 mass spectrometry (ICP-MS) — Part 1: General guidelines." Analysis was conducted according to ISO
123 17294-2:2016 "Water quality — Application of inductively coupled plasma mass spectrometry (ICP-
124 MS) — Part 2: Determination of selected elements including uranium isotopes (modified)" and USA
125 EPA 200.8 "Determination of Trace Elements in Waters and Wastes by Inductively Coupled Plasma-
126 Mass Spectrometry". All analyses were carried out by a certified laboratory.

127 The division of trace elements into commonly-monitored (i.e. Cd, Cr, Cu, Hg, Ni, Pb, Zn) and seldom-
 128 monitored (Ag, As, Ba, Co, Sb, V) was related to the frequency the elements were quantified in
 129 literature studies such as those presented in Table 5.

130 *Data analysis*

131 Calculations and statistical analyses were conducted using Statistica 13.5.0.17, TIBCO Software Inc.
 132 Multivariate analysis was conducted using the module Principal Component Analysis and Classification.
 133 Computational details are described in the software in the document "Principal Components &
 134 Classification Analysis – Computational Details".

135 *Determination of the contamination degree using the geoaccumulation index (I_{geo})*

136 The single element geoaccumulation index, I_{geo} , developed by Müller (1969) is generally used for
 137 comparing current metal levels in sediment to pre-industrial levels (Zhu et al. 2013; Manoy and Padhy
 138 2014; Duodo et al. 2016; Wang et al. 2016; Wang et al. 2018; Li et al. 2020). It is calculated using
 139 Equation 1:

140
$$I_{geo} = \log_2 \left(\frac{C_i}{1.5 \times B_i} \right) \quad \text{(Equation 1)}$$

141 where c_i is the measured concentration of the examined metal in the sediment, and B_i is the
 142 geochemical background concentration. Here B_i was calculated as the average value obtained from
 143 three soil samples taken around ponds A+B, C and E. An adjustment factor of 1.5 was used to account
 144 for possible variations in background values as well as minor anthropogenic influences. Müller (1969)
 145 used a graded scale of classification (Table 2):

146 **Table 2** Classification of I_{geo} (Müller 1969)

Class	Description	I_{geo} value
0	Unpolluted	$I_{geo} \leq 0$
1	Slightly polluted	$0 < I_{geo} \leq 1$
2	Moderately polluted	$1 < I_{geo} \leq 2$
3	Moderately/severely polluted	$2 < I_{geo} \leq 3$
4	Severely polluted	$3 < I_{geo} \leq 4$
5	Severely/Extremely polluted	$4 < I_{geo} \leq 5$
6	Extremely polluted	$I_{geo} > 5$

147 The sum of the average I_{geo} values for all trace elements determined in each pond were calculated and
 148 used to generate a risk ranking for each respective pond. These values were then compared to values
 149 obtained from two other risk ranking methods described below. Negative values (i.e. unpolluted; class
 150 0) were set to zero.

151 *Ecological risk assessment using the potential ecological risk index (RI)*

152 The RI method was developed by Håkanson (1980) for assessing the potential ecological risks in
153 sediment in freshwater ecosystems in Sweden. This was determined from the quantity of 7 elements
154 (As, Cd, Cr, Cu, Hg, Pb and Zn) and one persistent organic pollutant (PCB); and continues to be used for
155 ecological risk assessments of heavy metals in sediments (Manoj and Padhy 2014; Duodu et al. 2016;
156 Jiao et al. 2017; Wang et al. 2018; Zhuang et al. 2018; Wei et al. 2019; Li et al. 2020). The RI method
157 considers the hazard of metals to humans and ecosystems from two aspects - the abundance principle
158 and the release effect - where the potential toxicity of a metal is inversely proportional to its
159 abundance. The index includes the risk factor, Er^i , for a given substance (Equation 2), the sum of which
160 being the Potential Ecological Risk Index (RI) (Equation 4; Håkanson 1980):

161
$$Er^i = Tr^i \times C_f^i \quad \text{(Equation 2)}$$

162
$$C_f^i = \frac{C_o^i}{C_n^i} \quad \text{(Equation 3)}$$

163
$$RI = \sum_i^n Er^i = \sum_i^n Tr^i \times C_f^i \quad \text{(Equation 4)}$$

164 where Tr^i is the toxic response factor of substance i , C_f^i is the contamination factor of substance i ,
165 C_o^i is the concentration of substance i in the sediment, and C_n^i = concentration of substance i in the
166 background.

167 Here, C_f^i was calculated as the average value obtained from 3 soil samples taken around ponds A+B, C
168 and E. The toxic response factors (Tr) for each element were obtained from previously published
169 studies (Table 3). For a single substance, the risk factor value (Er) and the sum of risk factors (RI) were
170 classified according to (Håkanson 1980, Table 4).

171

172 **Table 3** Toxic response factors used for calculating the Potential Risk Index (Håkanson 1980)

Element	Tr	Source
Ag	17.5	Aksu et al. 1998
As	10	Håkanson 1980
Ba	2	Yang et al. 2015
Cd	30	Håkanson 1980
Co	5	Zhang et al. 2017
Cr	2	Håkanson 1980
Cu	5	Håkanson 1980
Ni	5	Zhang et al. 2017
Pb	5	Håkanson 1980
Sb	7	Wang et al. 2018
V	2	Zhu et al. 2013
Zn	1	Håkanson 1980

173

174 **Table 4** Classification of Potential Ecological Risk Index (Håkanson 1980)

Level of potential ecological risk	Er^i value	Level of ecological risk	RI value
Low	$Er^i < 40$	Low	RI < 150
Moderate	$40 \leq Er^i < 80$	Moderate	$150 \leq RI < 300$
Considerable	$80 \leq Er^i < 160$	Considerable	$300 \leq RI < 600$
High	$160 \leq Er^i < 320$		
Very high	$Er^i \geq 320$	Very high	RI ≥ 600

175

176 *Ecological risk assessment using the risk characterisation ratio method (RCR-method)*

177 The derivation of RCRs (the term Risk Quotients is also commonly used) for assessing ecological risk is
 178 used in many fields, such as for evaluating the risk of chemicals within the EU (ECHA 2016), the risk of
 179 emerging pollutants in landfill leachate (Nika et al. 2020), or the potential risk of pharmaceuticals in
 180 surface waters (Zhou et al. 2019). The RCR is derived by dividing the Predicted or Measured
 181 Environmental Concentration (PEC or MEC, respectively) by the Predicted No Effect Concentration
 182 (PNEC), where a RCR ≥ 1 indicates a risk of deleterious effects to the ecosystem by the pollutant in
 183 question. The PNEC is usually derived using toxicity data for sediment dwelling organisms, but where
 184 this data is lacking other methods such as the Equilibrium Partitioning Model which use toxicity data
 185 for pelagic species can be used (CIS 2011). PNEC values are often generated when Environmental
 186 Quality Standards are set, such as those under the Environmental Quality Standard Directive of the EU
 187 (Directive 2013/39/EU). In this study, we use data from when EQS were set for metals and metalloids
 188 in water, soil and sediment in the Netherlands (Crommentuijn et al. 2000); as equivalent data are still
 189 lacking for most sediment contaminants in Swedish and European legislation. Models and data from

190 the Netherlands have also been used for developing guideline values for contaminated soil in Sweden
191 (Swedish Environmental Protection Agency 2009). An RCR was consequently calculated using Equation
192 5:

$$193 \quad RCR_i = \frac{MEC_i}{NC_i} \quad (\text{Equation 5})$$

194 Where RCR_i is the risk characterisation ratio for element i , MEC_i is the measured concentration of
195 element i in the sample, and NC_i is the negligible concentration of element i . Values for NC_i were
196 calculated using the modified Eq-P method; and take into account mixture effects. Elements at
197 concentrations lower than or equal to their NC is not expected to cause negative long-term effects in
198 the ecosystem (Crommentuijn et al. 2000). NC s were available for all trace elements except Ag, for
199 which the Danish Default Guideline Value (DGV) of 1.5 mg per kg of dry weight (mg/kg DW) for
200 sediments was used as PNEC value (Ministry of Environment and Food of Denmark. 2017).

201 The sum of average RCR values for all trace elements in each pond were calculated to enable risk
202 ranking with I_{geo} and RI.

203 *Remediation requirements*

204 The samples were also classified using the generic guideline values for contaminated soil developed
205 by the Swedish Environmental Protection Agency (2009) using the guideline values in the updated list
206 from 2016 (Swedish Environmental Protection Agency 2016). Generic guideline values were available
207 for all trace elements except Ag (Supplementary Data Table S1). Two different generic guideline values
208 are derived depending upon expected land use, sensitive land use (KM) and less sensitive land use
209 (MKM). KM values are used for example if the land is going to be used for housing while MKM values
210 are used if the land use is intended for industries.

211 **Results and discussion**

212 *Concentrations of elements in sediment*

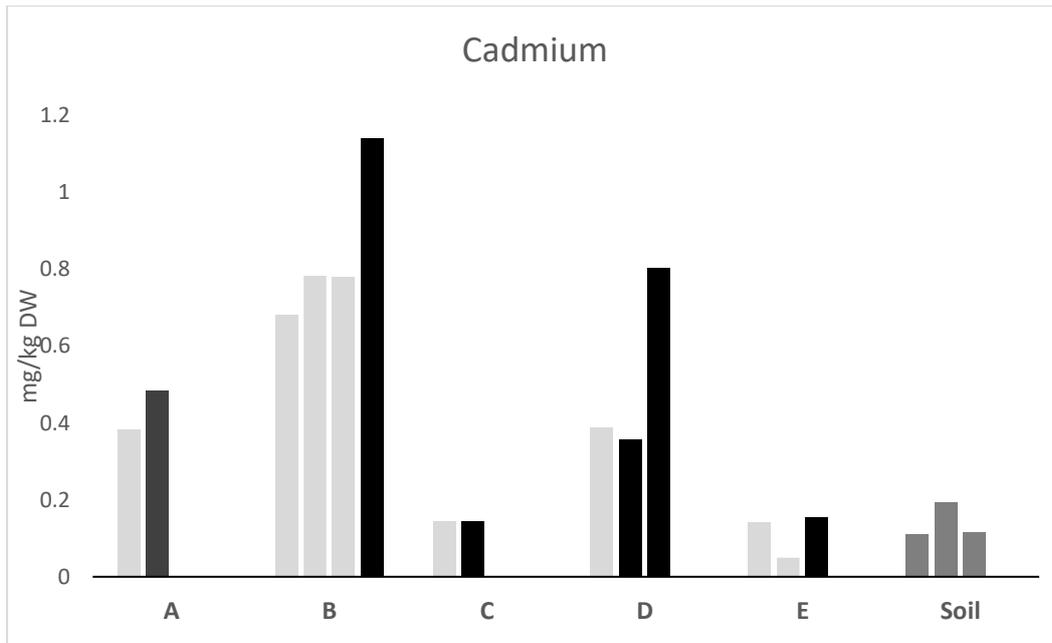
213 The concentrations for elements at all sampling sites and the average values for each element in each
214 pond are presented in Supplementary Data Table S2. The measured concentrations of four commonly-
215 monitored (Cd, Cu, Pb and Zn) and four seldom-measured (Ag, As, Ba and Sb) trace elements in
216 stormwater ponds are presented in Fig. 2 and Fig. 3, respectively. For all elements, the highest
217 concentrations were recorded in the older ponds (A, B and D) while the concentrations in recently
218 constructed ponds (C and E) were generally low or close to the concentration found in the surrounding

219 soil. The mean element concentrations were ranked as follows: P > S > Zn > Ba > Cu > V > Pb > Cr > Ni
220 > Co > As > Sb > Cd > Ag. Measured concentrations for Hg never exceeded the detection limit (0.5
221 mg/kg DW), while Ag was below the detection limit (0.7 mg/kg DW) in sediments from pond C and E.

222 The element concentrations were generally similar or higher at the outlets compared to the inlets of
223 ponds. For pond A, samples from near the outlet had higher concentrations for all elements compared
224 to the inlet except for S. In pond B, all elements were found in higher concentrations near the outlet,
225 except for Co, Cr and V which were in a higher concentration around inlet B2. In pond D, all elements
226 were found in higher concentrations at outlet D3 compared to the inlet, but only Cd, Cu, Pb, S and Zn
227 were higher in concentration at outlet D2 compared to the inlet. In pond E, element concentrations,
228 except for Ag, Co, Cu, S and Sb, were generally higher at the outlet than at the inlets. Inlet E2 presented
229 the highest concentration of Co and Cu, while inlet E1 had the highest concentrations of Ag, S and Sb.
230 Finally, in pond C, a different trend was observed where Ag was below the detection limit and all other
231 element concentrations – except for those of Ba, Cr and V - were higher at the inlet than the outlet.

232 Our measured trace element concentrations were then compared with data from other studies (Table
233 5). The concentrations of commonly-monitored trace elements (i.e. Cd, Cr, Cu, Ni, Pb, Zn) were
234 detected within the same order of magnitude as those observed in several previous international and
235 national studies (Jang et al. 2010; Blecken et al. 2012; 2017; Istenič et al. 2012; Egemose et al. 2015;
236 Frost et al. 2015; Sharley et al. 2017; Crane 2019; Sun et al. 2019), but lower maximum concentration
237 values were obtained here compared to those found in ponds within industrial areas (Sharley et al.
238 2017). Blecken et al. (2012) also detected six-fold higher Cr concentrations in sediment close to
239 stormwater outlets, while Jang et al. (2010) observed four-fold higher maximum values in road
240 residues. The concentration values for seldom-monitored metals (i.e. Ag, As, Ba, Co, Sb, V) were also
241 within the same order of magnitude as previously reported values (Färm et al. 2003; Jang et al. 2010;
242 Frost et al. 2015; Sharley et al. 2017; Crane 2019). Furthermore, higher maximum concentration values
243 for these elements were found in stormwater ponds and tunnels within industrial areas (Färm et al.
244 2003; Sharley et al. 2017). Jang et al. (2010) generally observed lower maximum values in sediment
245 from stormwater ponds but the maximum concentrations of Ba was higher.

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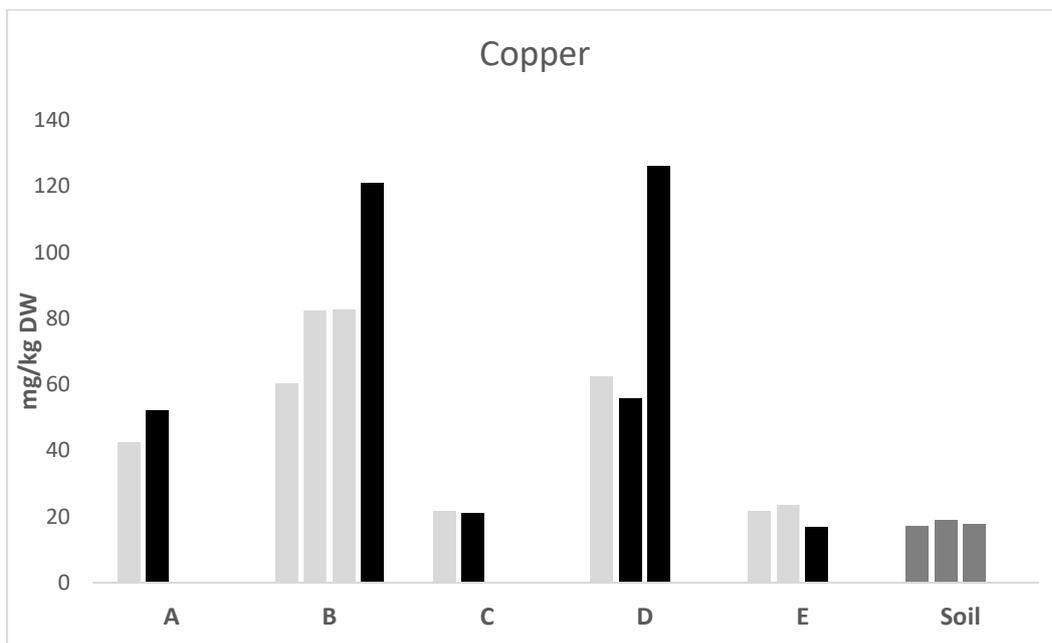


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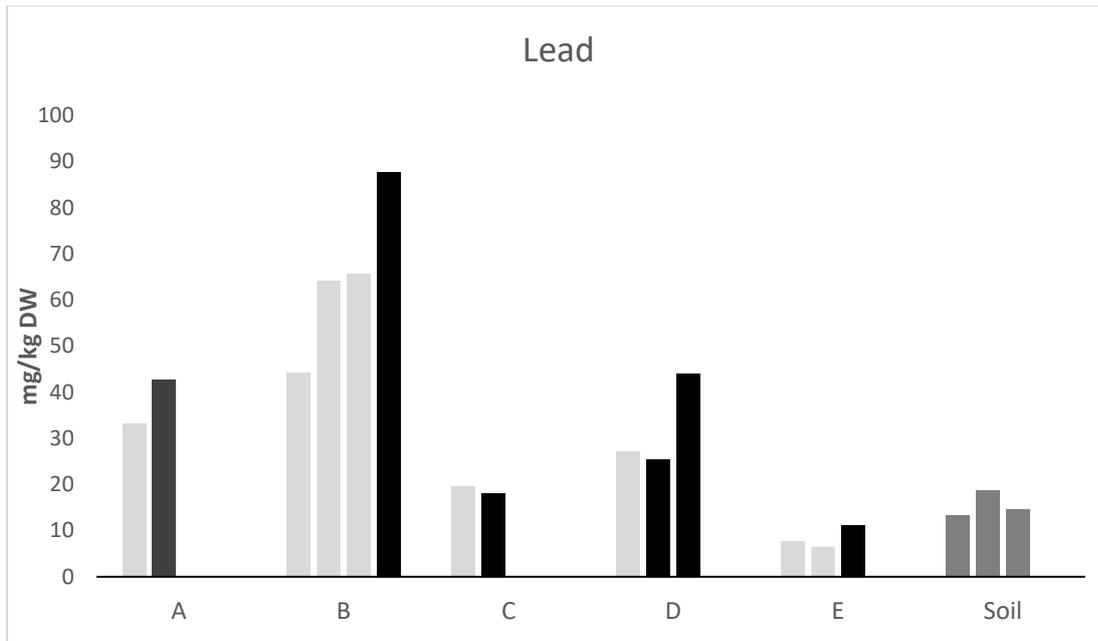
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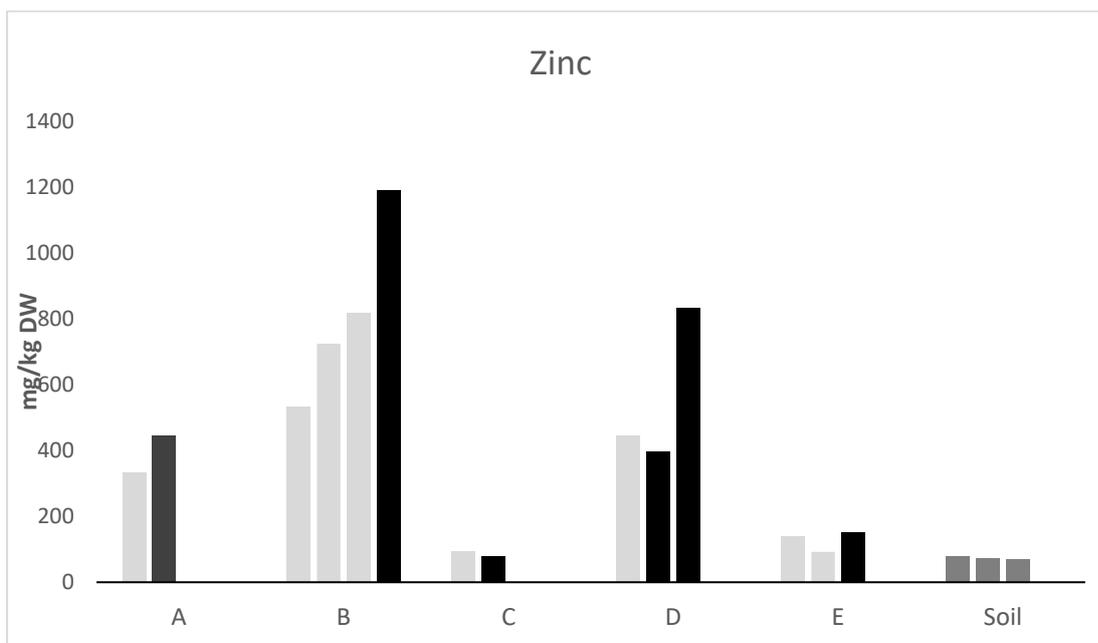
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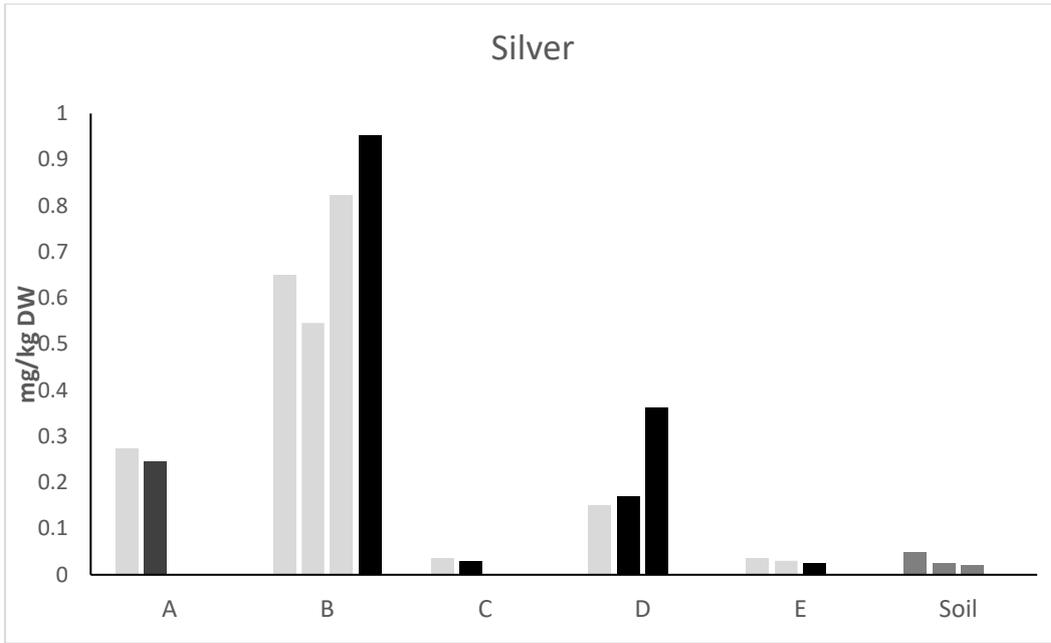


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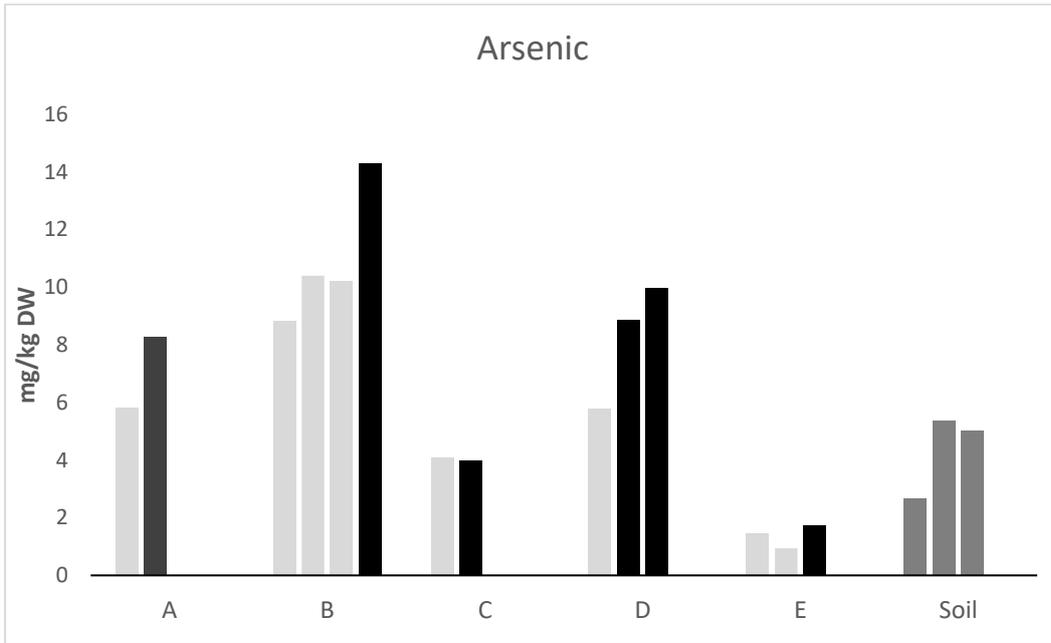
256 **Fig. 2** Concentrations of four commonly-monitored trace elements (Cd, Co, Pb, Zn) in sediments in
 257 stormwater ponds and soil from the banks of the ponds in areas with industrial activity. Light grey
 258 bars – inlets, black bars-outlets.

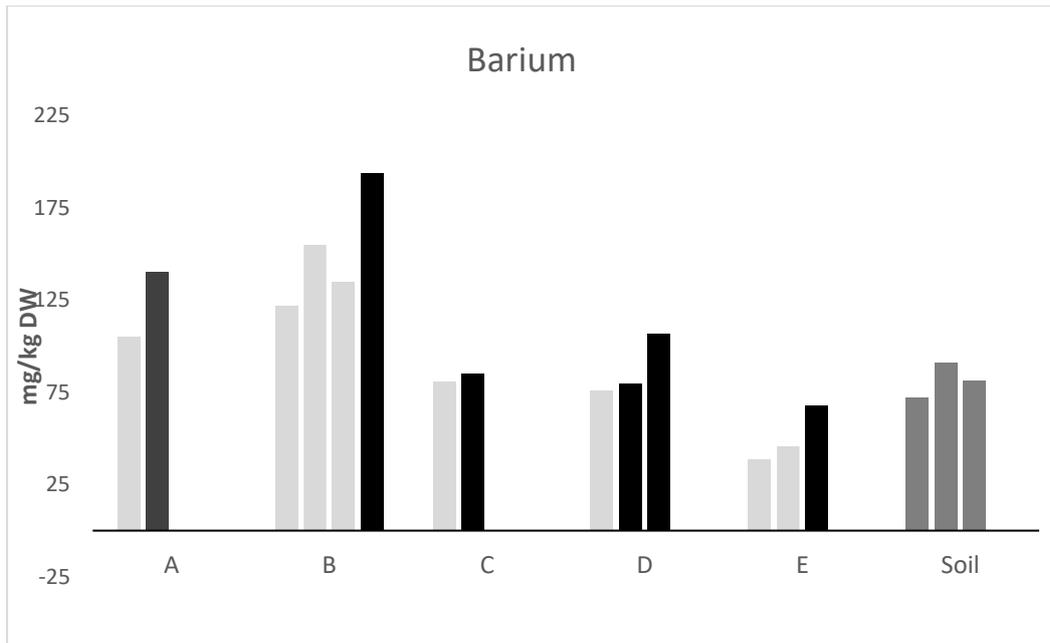
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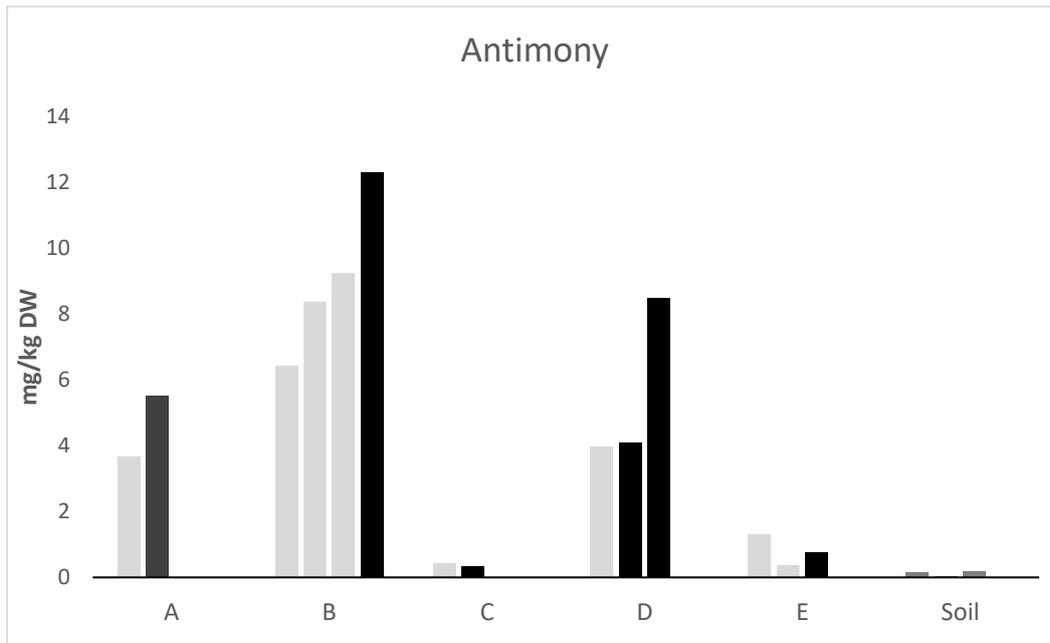


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263

264 **Fig. 3** Concentrations of four seldom monitored trace elements (Ag, As, Ba, Sb) in sediment in
 265 stormwater ponds and soil from the banks of the ponds in areas with industrial activity. Light grey bars
 266 – inlets, black bars-outlets
 267

268
269

Table 5 Concentrations of trace elements in sediments from stormwater ponds, outlets and stormwater tunnels. Data is for stormwater ponds if not stated. Min-max values are presented or if not available mean \pm standard deviation is used, n.a. = not analysed. N=the number of ponds studied

Referen- ce	Ag	As	Ba	Cd	Co	Cr	Cu	Hg	Ni	Pb	Sb	V	Zn
This study N=5	0.030- 0.952	0.9- 14.3	39- 194	0.05-0.80	3.5- 13.2	9.0- 45.6	20.9- 126.0	<0.5	6.4- 34.9	6.5- 87.6	0.325- 12.3	15.9- 75.3	78.6- 1190
Crane 2019 N=15	0.99 \pm 2.4	5.5 \pm 2.3	147 \pm 43	1.5 \pm 0.57	8.2 \pm 2.8	42.0 \pm 14.7	31.6 \pm 10.1	0.059 \pm 0.022	23.3 \pm 7.2	39.2 \pm 25.5	<1.2	45.3 \pm 11.7	154.3 \pm 79.7
Sun et al. 2019 N=12	n.a.	n.a.	n.a.	n.a.	n.a.	15-86	18-200	n.a.	17-110	9.9-76	n.a.	n.a.	57-850
Blecken et al. 2017 N=25	n.a.	n.a.	n.a.	<0.1-2.3	n.a.	2-81	3-109	n.a.	2-39	3-77	n.a.	n.a.	14-597
Sharley et al. 2017 N=98	<2-6	<5-51	40- 430	<1-11	2-31	12-121	6-1090	<0.1- 2.9	4-159	9-456	<5-17	14-130	12-4940
Ege- mose et al. 2015 N=37	n.a.	n.a.	n.a.	0.1-1.4	n.a.	18-62	18-62	n.a.	18-62	18-62	n.a.	n.a.	166-451
Frost et al. 2015	0.04-0.37	n.a.	64.7- 507.3	0.06-0.39	3.8- 20.3	9.6- 60.5	9.5-42.3	n.a.	6.6- 45.4	6.0-38.1	n.a.	14.3-67.5	35.5- 175.4
Blecken et al. 2012 N=32	n.a.	n.a.	n.a.	0.15-1.23	3.8- 25.0	64-307	17-263	n.a.	10-50	8-42	n.a.	n.a.	68-470
Istenič et al. 2012	n.a.	n.a.	n.a.	<0.5	n.a.	17-80	4-3293	n.a.	10-42	<2-220	n.a.	n.a.	26-1361

N=3													
Jang et al. 2010 N=22	<0.8	0.6-24.8	8.1-1019	<0.37 1st -5.3	n.a.	5.8-174.5	4.5-90.4	<0.02	5.4-40.4	5.6-196	n.a.	n.a.	5.4-711
Färm et al. 2003 N=3	0.314-1.70	10.2-20.7	n.a.	1.15-2.93	19.6-34.7	51.0-75.7	106-159	Kan finnas	32.0-50.5	61.4-98.2	n.a.	73.2-106	609-974

270 ¹ wetlands

271 ² stormwater outlets

272 ³ road residues from stormwater systems

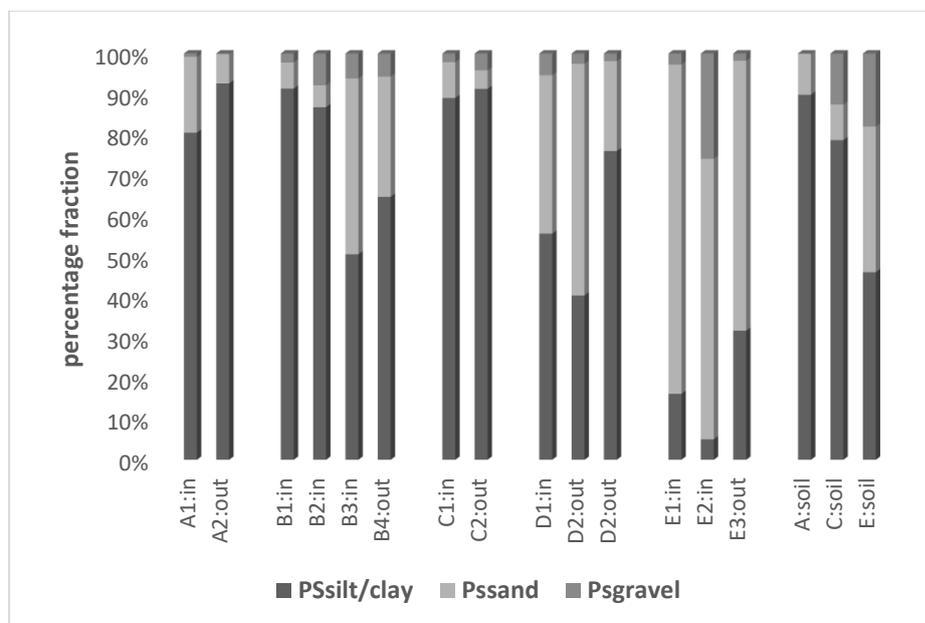
273 ⁴ sediment from stormwater tunnels

274 *Sediment depth and particle size distribution*

275 The sediment depth measurements in different ponds showed that there were no significant
 276 differences in sediment depth between the areas surrounding the inlets and outlets (Fig. 1). The largest
 277 sediment depths were observed in one of the older ponds (B). Accumulation of material had already
 278 occurred at the inlets in one of the newer ponds (E), while the accumulation was still low in the other
 279 new pond (C).

280 The particle size distribution and the sediments varied between sampling sites across individual ponds
 281 and between ponds (Fig. 4, all data shown in Supplementary Data Table S2). Most of the sampling sites
 282 contained mainly fine material (PSsilt/clay), except for pond E. During sampling in August 2019, erosion
 283 of material from the surrounding step banks into the ponds was observed, and thus the sediment may
 284 constitute material from both the catchment and the areas directly surrounding the ponds.

285 A well-designed pond should accumulate coarse material at the inlets and fine material at the outlets,
 286 but we observed mixed results. Here, the stormwater ponds were constructed to prevent flooding
 287 downstream and have several inlets (e.g. B and E) or several outlets (e.g. D), therefore the hydraulic
 288 efficiency may be low. However, previous studies also reported varied results which suggested that
 289 fine sediment may also accumulate preferentially at inlets in ponds with high hydraulic efficiency
 290 (Istenič et al. 2012; Blecken et al. 2017).



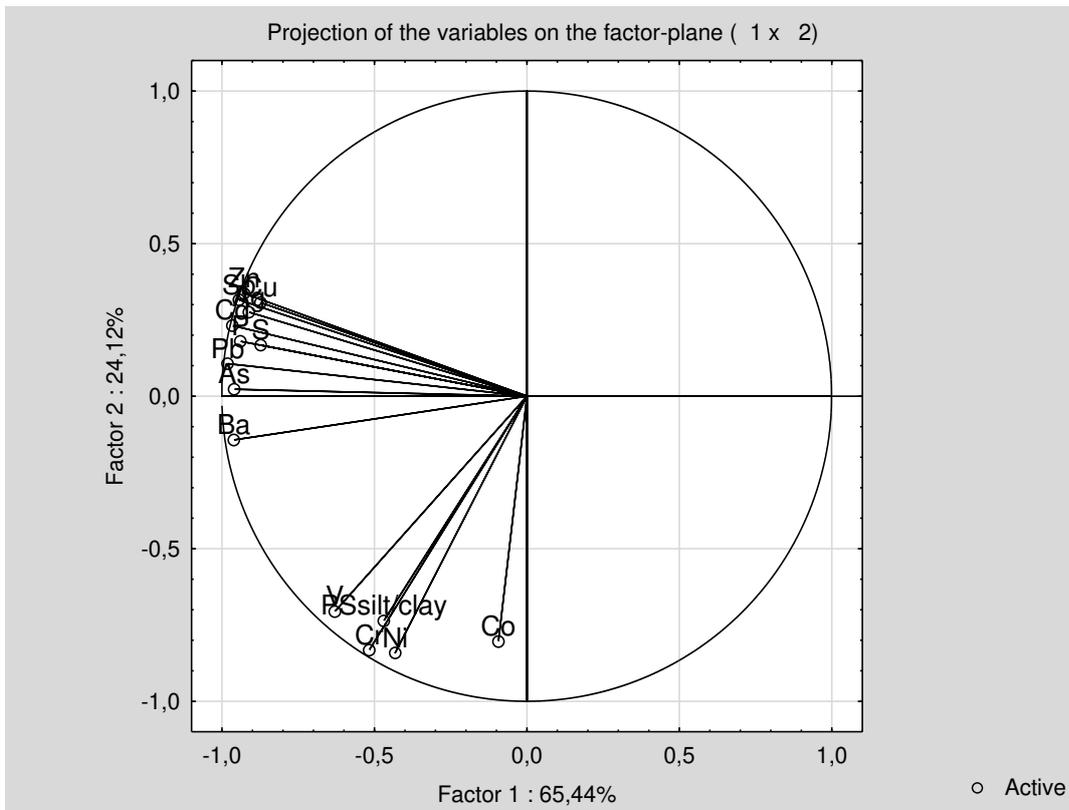
291
 292 **Fig. 4** Particle size distribution of the sediment in the different sampling sites in five stormwater ponds.
 293 Particle size (PS) for PSsilt/clay < 0.063 mm, PS sand 0.063- 2mm and PS gravel > 2mm

295 Trends in variation and the relationship between measured variables were further explored using PCA
296 and classification methods. PCA on all element concentration data and sediment characteristics was
297 not possible because the correlation matrix became singular indicating that the data could be
298 replicated as the linear combination of fewer variables. Therefore, PCA was conducted with the
299 element concentration and PSsilt/clay data (Fig. 5). In Fig. 5a the PCA for the 2 first Principle
300 Components is presented for the samples including the soil samples (cases n=17), the 12 trace
301 elements detected, P, S and the silt/clay fraction (variables n=15). Only 2 PCs had an eigenvalue above
302 1 and PC1 and PC2 describe 65.44% and 24.12% of the variation respectively. Two distinct groups of
303 variables were identified (Fig. 5a). The first group (Fig 5b), hereafter labelled Group A, consisted of
304 elements with a high negative loading (score) on PC1 include (in descending order) Pb, Cd, Ba, As, Sb,
305 P, Zn, Ag, Cu and S. They also all have a low positive loading on PC2 except for Ba. The second group,
306 hereafter labelled B (Fig. 5c) , consisted of elements and parameters with lower negative loadings on
307 PC1 and higher negative loadings on PC2 included (in descending order of PC 1) V, Cr, PSsilt/clay (i.e.
308 PS < 63 μm) and Ni. Co was included in the second group, but we remark that it showed a very low
309 negative loading on PC1.

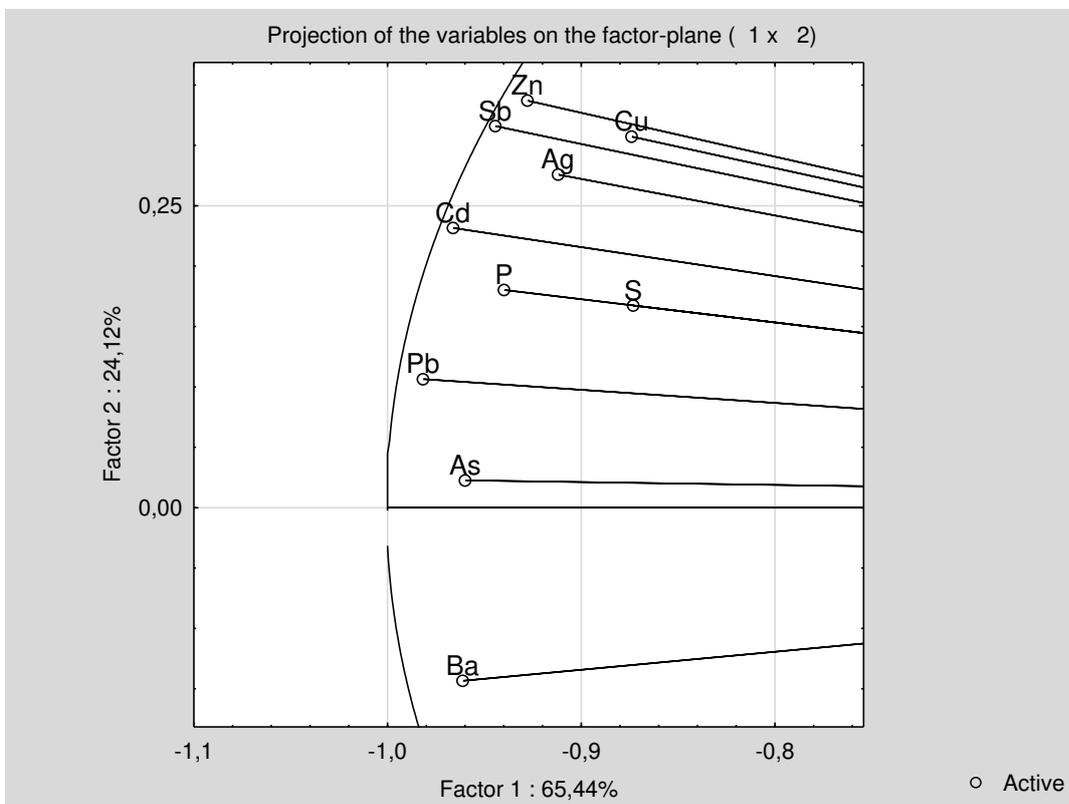
310 The projection of the cases on PC1 and PC2 showed that samples from a given pond were loosely
311 grouped (Fig. 5d). This suggested that samples taken from a single pond were more similar to each
312 other than to those from other ponds, which reflects differences in catchment characteristics. Samples
313 with the highest contamination level had the highest negative loading in PC1 (i.e. located furthest to
314 the left in Fig. 5b), which indicated that pond B was the most contaminated and pond E was the least
315 contaminated.

316 A Pearson's r correlation analysis was also conducted to identify correlations between the elements
317 and the sediment characteristics including PSclay/silt (i.e. PS < 63 μm), PSsand (i.e. PS 0.63 μm – 2 mm),
318 PSgravel (i.e. PS > 2 mm), PScoarse material (i.e. PSsand and PSgravel) and sediment depth (Table 6).
319 Statistically significant positive correlations ($p \leq 0.05$) were observed for PSclay/silt and (in order of
320 decreasing correlation) V, Cr Ni, S, Ba and Co; where the same elements also showed significant
321 negative correlation with similar amplitude with PSsand. No significant correlation between PSgravel
322 and the elements was observed. Additionally, (in order of decreasing correlation) P, Pb, Ba, Cd, Sb, Zn,
323 Ag and S were significantly correlated with sediment depth. Finally, a correlation analysis was also
324 conducted for the trace elements and the potential indicators of contamination P and S (Table 6).
325 Elements constituting Group A as identified in the PCA (see Fig. 5a,b) showed higher correlation

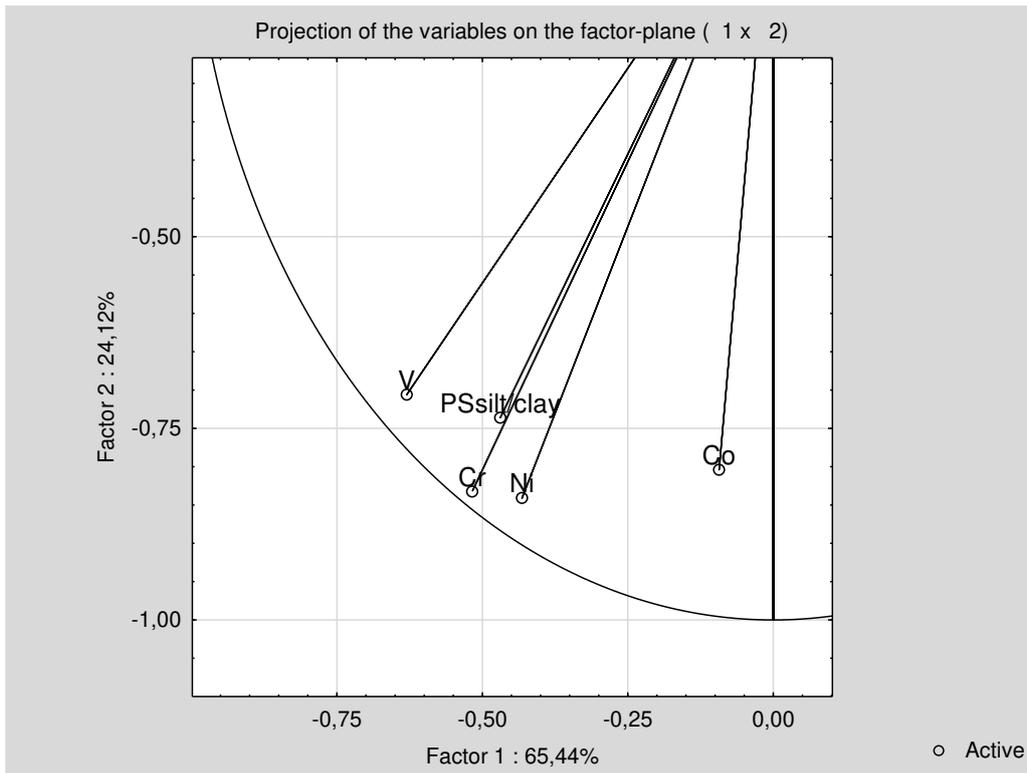
326 coefficients with P and S than those in Group B (Fig 5a,c) except for V which showed a significant
 327 correlation with S. Group A elements were more strongly correlated with P than with S.



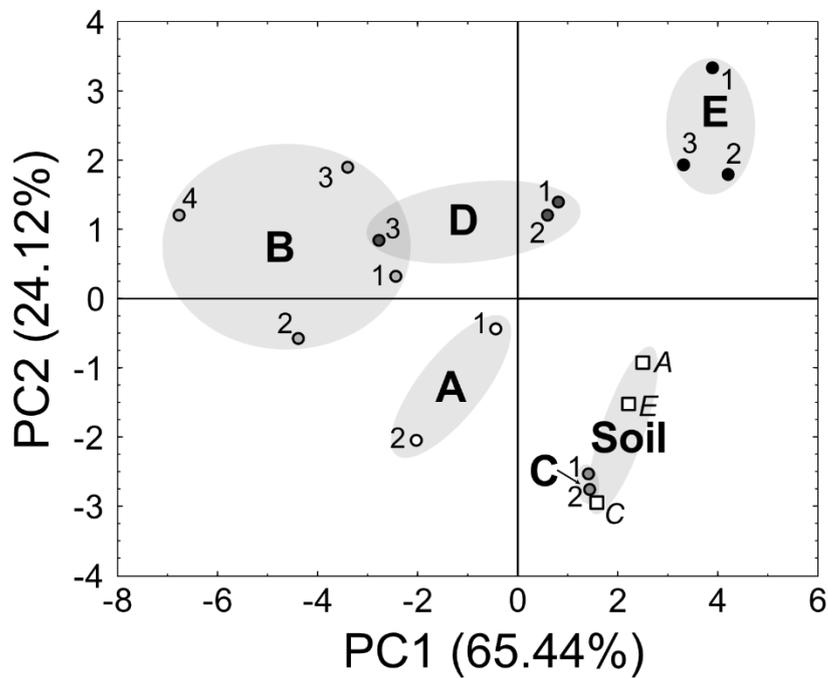
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331

332 **Fig. 5** a) Projection of the variables on PC 1 and PC2. b) Enlarged section of PCA with Group A
 333 elements c) Enlarged section of PCA with Group B elements and PSsilt/clay d) Projection of the
 334 samples on PC1 and PC2

335 **Table 6** Pearson r correlations between sediment characteristics and elements. Significant correlations
 336 are presented in in bold. Statistical significance was set to <0.05

Element mg/kg DW	PSsilt/clay ¹ (%)	PSsand ² (%)	Sediment depth (cm)	P mg/kg DW	S mg/kg DW
Ag	0.29	-0.31	0.57	0.96	0.78
As	0.48	-0.47	0.53	0.90	0.82
Ba	0.58	-0.59	0.60	0.91	0.87
Cd	0.38	-0.38	0.59	0.93	0.86
Co	0.54	-0.68	0.16	0.10	0.15
Cr	0.91	-0.92	0.16	0.46	0.59
Cu	0.28	-0.28	0.48	0.76	0.75
Ni	0.85	-0.86	0.14	0.42	0.47
P	0.38	-0.40	0.62	-----	0.84
Pb	0.44	-0.44	0.61	0.96	0.86
S	0.59	-0.58	0.54	0.84	-----
Sb	0.31	-0.31	0.59	0.92	0.84
V	0.93	-0.92	0.16	0.49	0.70
Zn	0.26	-0.26	0.58	0.89	0.80

337 ¹Particle size < 0.063mm

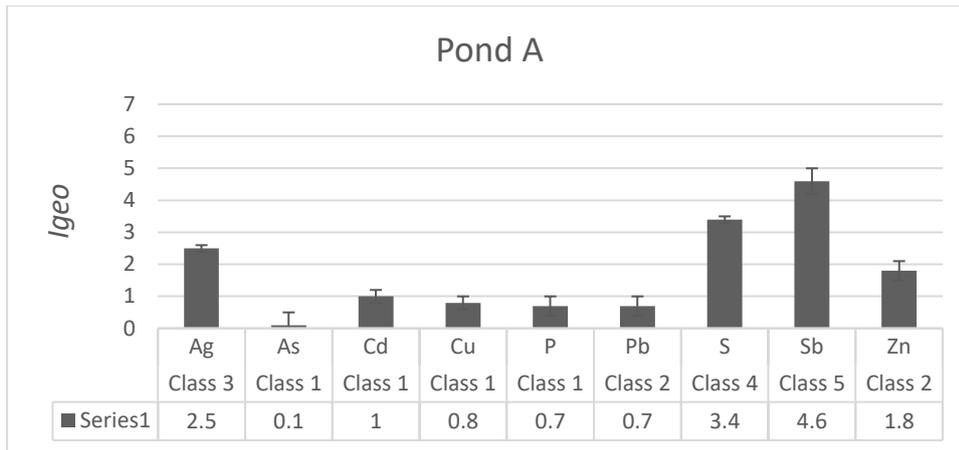
338 ²Particle size 0.063 mm – 2 mm

339

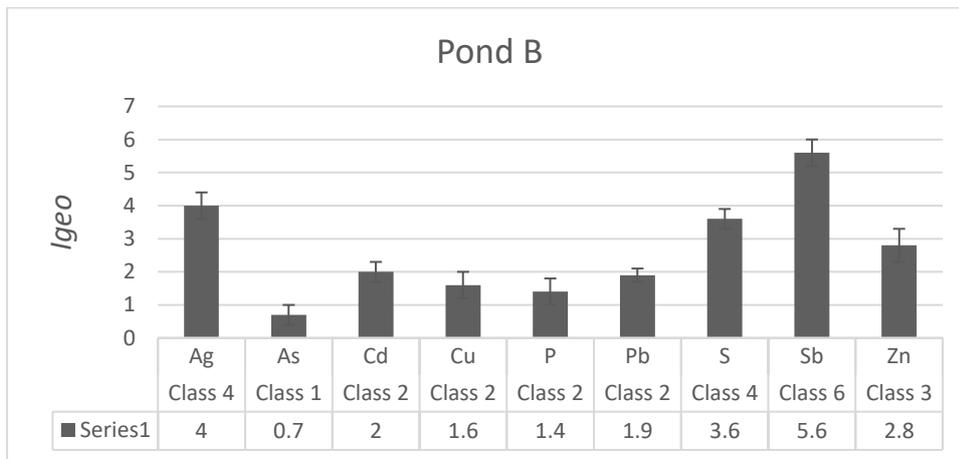
340 *Contamination degree using the geoaccumulation index I_{geo}*

341 The single element index, I_{geo} was used to assess the degree of contamination in the sediment. The
 342 mean I_{geo} values for the elements Co, Cr, Ni and V in all ponds were all negative and the sediment were
 343 therefore classified as unpolluted (Class 0) with respect to these elements. The mean I_{geo} for Ba only
 344 exceeded 0 in pond B (0.3±0.3). The I_{geo} for the elements Ag, As, Cd, Cu, P, Pb, S, Sb and Zn are shown
 345 for each pond in Fig. 6. The complete data set for I_{geo} values is shown in Supplementary Data Table S3.
 346 The highest mean I_{geo} values were obtained for Ag, S, Sb and Zn. The older ponds (A, B and D) showed
 347 higher contamination degree than the newest ponds (C and E). Overall, Pond B was the most polluted,
 348 showing the highest I_{geo} values for Ag (Class 4), S (Class 4) and Sb (Class 6) and Zn (Class 3). Ponds C and
 349 E only showed contamination by S and Sb (both ponds) and Zn (pond E only).

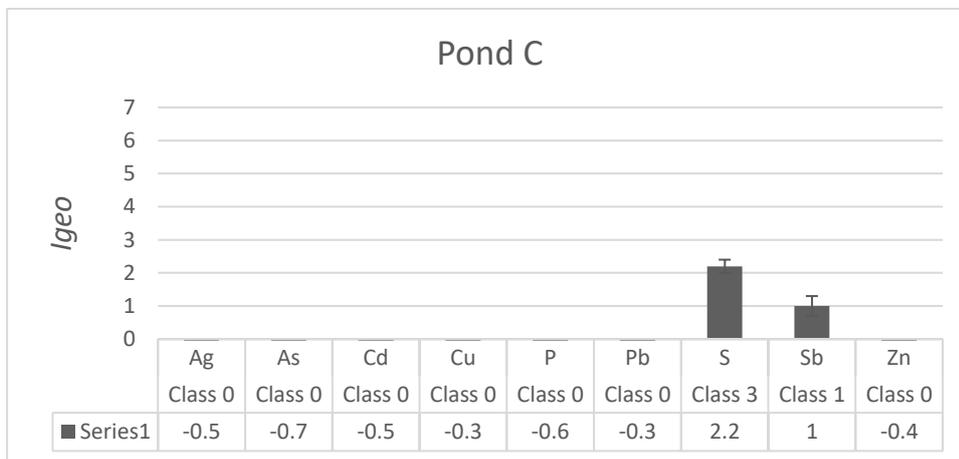
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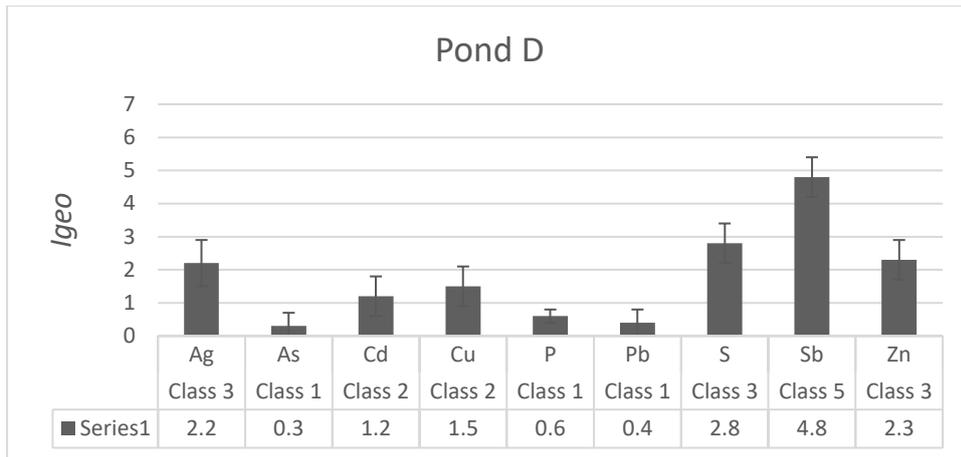


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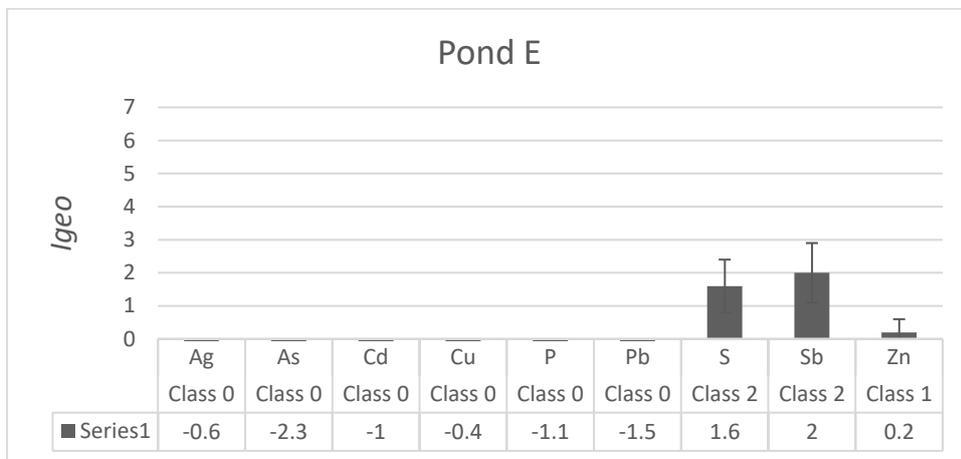


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353



354

355 **Fig. 6** Contamination degree for trace elements in the different ponds using the geoaccumulation
 356 index I_{geo} . Classification was made using mean values, and error bars present standard deviation
 357

358

359 *Ecological risk assessment using the Potential Ecological Risk Index (RI)*

360 The mean risk factor, Er , for the elements As, Ba, Co, Cr, Cu, Ni, Pb, V and Zn was below 40 which
 361 suggested they provided low potential ecological risk. The Er values for Ag, Cd and Sb are shown in
 362 Table 7, and all data are summarised in Supplementary Data Table S4. Low risk factors ($Er < 40$) were
 363 obtained for the newer ponds C and E while the older ponds A,B and D showed very high risks ($Er \geq$
 364 320) for Ag and Sb.

365 The RI for ponds C and E were classified as low while the classifications for RI for ponds A, B and D
 366 ranged from considerable ecological risk (Class 3) to very high ecological risk (Class 4). The highest RI
 367 was obtained in pond B.

368 **Table 7** Mean risk factor, Er , values for Ag, Cd and Sb with standard deviation; the RI for the sum of
 369 risk factors and the contribution, in percentage, of Ag, Cd and Sb to the RI in sediment from the five
 370 stormwater ponds

Pond	Er -Ag ¹	Er -Cd ¹	Er -Sb ¹	RI ²	% Ag of RI	% Cd of RI	% Sb of RI	% Ag+Cd+Sb of RI
A	147±11 Class 3	93±16 Class 3	267±75 Class 4	593 Class 3	25	16	45	86
B	421±102 Class 5	182±44 Class 4	528±143 Class 5	1248 Class 4	34	15	42	91
C	18±2 Class 1	31±0 Class 1	22±4 Class 1	134 Class 1	14	23	16	53
D	129±66 Class 3	111±54 Class 3	321±150 Class 5	646 Class 4	20	17	50	86
E	17±3 Class 1	25±12 Class 1	48±27 Class 2	117 Class 1	16	20	39	75

371 ¹ The risk of single element was classified as according to Håkanson (1980) and here, for convenience,
 372 it has been given a numeric classification: Class 1 Low potential ecological risk $Er < 40$; Class 2 Moderate
 373 potential ecological risk $40 \leq Er < 80$, Class 3 Considerable ecological risk $80 \leq Er < 160$; Class 4 High
 374 potential ecological risk $160 \leq Er < 320$ and Class 5 Very high ecological risk $Er > 320$.

375 ² The RI, the sum of the risk factors, was classified according to Håkanson (1980) and here, for
 376 convenience, it has been given a numeric classification: Class 1 Low ecological risk $RI < 150$; Class 2
 377 Moderate ecological risk $150 \leq RI < 300$; Class 3 Considerable ecological risk $300 \leq RI < 600$; Class 4 Very
 378 high ecological risk $RI \geq 600$.

379

380 *Ecological risk assessment using the RCR-method*

381 Elements with mean RCRs exceeding 1 in at least one sample are summarised in Table 8, while all data
 382 is presented in Supplementary Data Table S5. Mean RCRs in soil were < 1 for all elements except for
 383 Co and V. In pond sediment mean RCRs for Ag, As, Cd, Cr, Pb were always ≤ 1 . For Ba, two samples (B2
 384 and B4) had a $RCR \geq 1$ (1.0 and 1.2, respectively). For Cd and Ni, only samples B4 and C1 had a $RCR \geq 1$
 385 (both 1.0). For Co and V all ponds except pond E had a mean RCRs of $2 > RCR \geq 1$. Finally, mean RCRs
 386 for Cu, Sb and Zn exceeded 2 in numerous samples, where the highest mean RCRs for Zn (5.6), Sb (2.8)
 387 and Cu (2.4) - and therefore the highest Sum-RCR - was obtained in pond B. The lowest sum-RCR values
 388 were obtained for the newest ponds C and E. Indeed, the sum-RCR for the surrounding soil exceeded
 389 that of pond E.

390

391 **Table 8** Mean Risk Characterisation Ratio (RCR) with standard deviation for elements measured in soil
 392 and sediments in five stormwater ponds. Negligible concentrations (NC, in mg/kg DW sediment) was
 393 collected from Crommentuijn et al. (2000). Mean RCRs in bold text ≥ 1 , ponds with a at least 1 sample
 394 with a RCR ≥ 1 in italics.

Pond	Ba (157) ¹	Cd (1.1)	Co (9.1)	Cu (36)	Ni (35)	Sb (3.2)	V (42)	Zn (145)	Sum- RCR ²
A	<i>0.8±0.2</i>	0.4±0.1	1.2±0.2	1.3±0.2	0.7±0.1	1.4±0.4	1.6±0.3	2.7±0.5	11.1±2.1
B	1.0±0.2	<i>0.8±0.2</i>	1.1±0.2	2.4±0.7	0.7±0.2	2.8±0.8	1.3±0.2	5.6±1.9	17.3±4.2
C	0.5±0.0	0.1±0.0	1.4±0.0	0.6±0.0	1.0±0.0	0.1±0.0	1.3±0.1	0.6±0.1	6.3±0.1
D	0.6±0.1	0.5±0.2	1.0±0.2	2.3±1.1	0.5±0.1	1.7±0.8	1.0±0.2	3.8±1.6	12.3±4.5
E	0.3±0.1	0.1±0.1	<i>0.8±0.5</i>	0.6±0.1	0.3±0.1	0.3±0.1	0.5±0.2	<i>0.9±0.2</i>	3.9±0.4
soil	0.5±0.1	0.1±0.0	1.4±0.3	0.5±0.0	0.8±0.2	0.0±0.0	1.1±0.1	0.5±0.0	5.6±0.8

395 ¹NC in parenthesis

396 ²sum mean RCR for all elements analysed except P and S

397

398 *Remediation requirements*

399 The complete data set is shown in Supplementary Data Table S2. The mean concentration of each
 400 element in background soil was below the respective generic guideline values for both less sensitive
 401 land use (MKM) and sensitive land use (KM). The KM value for Co was, however, exceeded in the
 402 surrounding soil from pond C. The generic guideline value for less sensitive land use (MKM) was never
 403 exceeded in pond sediments except for Zn where all samples from pond B exceeded the MKM value
 404 as did the sample D3 from pond D. Based upon means the KM value was exceeded in pond A for Zn,
 405 for pond B for As, Cd, Cu, Pb and Zn (i.e. exceeds MKM) and for pond D for Cu and Zn (i.e. Zn exceeds
 406 MKM). No sediment concentrations exceeded KM values in pond C and E. This suggest that some
 407 elements - particularly Zn - will limit the re-use of the sediments from the older ponds and they may
 408 require costly and laborious clean-up procedures (Zubala et al. 2018) when excavated in the future.

409 *Priority ranking of ponds and elements*

410 The ponds were then ranked according to their I_{geo} , RI and RCR values (Table 9). The rankings were
 411 similar for all methods except I_{geo} was higher in pond E than in pond C. Overall, Pond B was the most
 412 contaminated. The degree of contamination was summarised as B >>> D \approx A > C \approx E. The degree of
 413 contamination in ponds C and E was low such that their RCR values matched those of the surrounding
 414 soil. This confirmed that the younger ponds (C and E) were less contaminated than the older ponds. In
 415 theory, the annual accumulation of trace element should be the same if it is not influenced by other
 416 factors. In this case, it was assumed that low rainfall during 2018, 60% less than normal (Swedish
 417 Meteorological and Hydrological Institute 2021), contributed to a low accumulation. Additionally, the

418 percentage impervious surface of pond C was low (12%) as the pond was constructed before the
419 industrial area has been fully developed.

420 The 3 ranking methods used were originally developed for sediment quality assessment in lakes and
421 watercourses (Müller 1969; Håkanson 1980; Crommentuijn et al. 2000). To our knowledge, this study
422 represented the first use of I_{geo} and RI in the risk assessments of stormwater ponds sediment. The RCR-
423 method is rarely used but sediment guidelines are frequently compared to measured values when
424 assessing the quality of stormwater sediment (Färm et al. 2003; Andersson et al. 2004; Färm and Waara
425 2005; Jang et al. 2010; Blecken et al. 2012; Sharley et al. 2017; Crane 2019). It is encouraging that I_{geo} ,
426 for which there is no need to find neither Tr nor PNEC values, provided a similar ranking as the RI and
427 the RCR-method. This suggested that the I_{geo} method is suitable for evaluating the risk caused by the
428 accumulation of emerging contaminants in stormwater ponds.

429 Although the different ranking methods indicated somewhat different elements of concern (see Table
430 9), their scales for classification were different and P and S were not included in the RI and Sum-RCR
431 values. P and S are not considered toxic trace elements, but they are used in this study to investigate
432 if they are correlated with the accumulated trace elements. Both elements belong to the Group A
433 elements identified in the PCA (see Fig. 5a,b) and they were significantly correlated with the trace
434 elements with a high I_{geo} (Fig. 6). P and S are rarely analysed in sediment in stormwater ponds, but the
435 P level measured here was within the same range (358-4130 mg/kg DW) as reported by Istenič et al.
436 2012 (90-1810 mg/kg DW). The levels for S found in this study (347-3170 mg/kg DW) are within the
437 lower range of the interval found by Blecken et al. in 2012 (358-16000 mg/kg DW). Finally, while both
438 P and S showed high and significant correlation coefficients with trace elements with a high I_{geo} , P
439 showed very low accumulation compared to S. This indicated that S was generally a better indicator of
440 contamination than P, but additional studies are necessary to explore if S content can be a general
441 indicator of pollution in stormwater ponds.

442 It was assumed that the RCR values for Co, Ni and V (> 1) were questionable since I_{geo} indicated no
443 accumulation of these elements above the background reference soil (Table 9). The NC value used as
444 PNEC was derived for risk assessments made in the Netherlands, which accounted for higher local
445 background concentrations than what we observed in own reference soil (except Co and V;
446 Crommentuijn et al. 2000). There are currently no available environmental quality standard (EQS) for
447 sediment in lakes and watercourses in Sweden for most of the trace elements analysed, except for Cd
448 (EQS=2.3 mg/kg DW, NC = 1.1 mg/kg DW), Cu (EQS=36 mg/kg DW, NC = 37 mg/kg DW) and Pb
449 (EQS=130 mg/kg DW, NC = 132 m/kg DW) (HVMFS2019:25). However, replacing the NCs used in this

450 study with the national EQSs only marginally changed the RCRs for Cu and Pb, although the average
451 RCR for Cd fell under 1. As both the I_{geo} and Er values indicated that Cd was an element of concern in
452 pond B, Cd was kept as an element of concern overall. Denmark has set a national guideline for V to
453 23.6 mg/kg DW using the added risk approach (Ministry of Environment and Food of Denmark. 2017),
454 which can be added to the background concentration. In our case, the concentration in a sample must
455 have exceeded 71.7 mg/kg DW (i.e. $48.1 + 23.6 = 71.7$) to affect the ecosystem negatively, and this level
456 was only exceeded in sample A2 (Supplementary Data Table S2).

457 The NCs and the national EQSs for Cd, Cu and Pb were calculated under the assumption that the total
458 organic carbon (TOC) was 5%, and that at a higher TOC the trace elements would bind to the organic
459 matter and become biologically unavailable (CIS 2011). Organic C was not measured in this study, but
460 we assume that organic C content would vary depending upon land use and amount of vegetation in
461 the ponds.

462 Ag is considered an element of concern despite low RCR levels (i.e. $RCR < 1$) because the I_{geo} value
463 indicated extensive accumulation and the Er value was high. Cu was also classified as an element of
464 concern even though the RCR was also below 1 because the I_{geo} and Er value indicated a risk and the
465 KM guideline was exceeded in pond B and D. As and Pb are interesting cases as the I_{geo} indicated
466 accumulation in pond A, B and D and the RCRs were below 1, but the KM values for Pb (KM=50 mg/kg
467 DW; NC = 132 mg/kg DW) and As (KM=10 mg/kg DW; NC =31 mg/kg DW) were exceeded in pond B.
468 This was attributed to the setting of the generic guideline values for contaminated soil for both
469 ecological and human risk. The RCR for Ba was above 1 only in pond B, while I_{geo} indicated only slight
470 pollution (Class 1) and the Er value indicated low risk in the same pond. Consequently, Ba may not be
471 classified as an element of concern. The high RCRs in pond A, B and D for Sb and Zn agreed directly
472 with the I_{geo} and Er classification and were thereby classed as elements of high concern. Sb and Zn also
473 show high correlation in the PCA (Fig. 5b) and may therefore have a common source of origin. In
474 conclusion, the use and comparison of three risk assessment methods confirmed that both commonly
475 monitored trace elements (Cd, Cu, Zn) and seldom monitored trace elements (Ag, Sb) may cause an
476 ecological risk in the ponds.

477 For Ag (Färm et al. 2003; Frost et al. 2015; Sharley et al. 2017; Crane 2019) and Sb (Sharley et al. 2017;
478 Crane 2019) we have found only a few records of measurements in stormwater ponds with catchments
479 featuring industrial land yet the current data showed that the monitoring these elements is paramount
480 as they showed very high accumulation compared to the background. Ag is currently used in many
481 nanomaterials in textiles, medical products, food containers, cosmetics, paints and nano-

482 functionalised plastics (McGillicuddy et al. 2017). Meanwhile, Sb is used in catalysts for production of
483 polyethylene terephthalate (PET), and in flame retardants for various materials such as paper, plastic,
484 paints and textiles (Wang et al. 2018); and brake linings (Wang et al. 2018, Müller et al. 2020). It has
485 recently been found in high concentration in plastic waste fractions (Viczek et al. 2020); and shown to
486 be mobilised from microplastic in coastal estuarine sediments when extracted with fluids that simulate
487 the digestive system of sediment dwelling invertebrates (James and Turner 2020). Significantly, Pond
488 D is located near to a municipal recycling plant. As the industrial areas are less than 5 km apart, they
489 may also experience similar sources of local and long-distance transport of airborne pollutants. Clearly,
490 further studies are needed to determine whether the high accumulation of Ag and Sb in the studied
491 stormwater ponds is a local phenomenon or not.

492 In this study, the total amount of certain elements was measured and risk-assessed. There are several
493 studies that indicate that heavy metals preferentially accumulate in the roots of macrophytes and are
494 therefore unlikely to become bioavailable via plant uptake and translocation (Istenič et al. 2012).
495 Benthic organisms have also been shown to accumulate heavy metals in stormwater ponds such as Cd,
496 Cr, Ni and Cu (Stephansen et al. 2014), but they might not reach toxic levels as was demonstrated by
497 Andersson et al. (2004) and Casey et al. (2006). Recently, Sun et al. (2019) studied the biodiversity in
498 roadside ponds and found that most of the taxa displayed in the ordination diagram were negatively
499 correlated with the pollution levels in the water column and sediments, while a positive correlation
500 with the pond size and the number of neighbouring ponds was observed. Whether the trace elements
501 studied here will cause toxicity and thereby reduce the biodiversity in catchments with industrial
502 activity remains to be explored.

503

504 **Table 9** Priority ranking of contamination in the ponds and identified elements of concern based upon
 505 I_{geo} , RI and RCR

Pond	Rank Sum- I_{geo}^1	Rank RI ²	Rank Sum- RCR ³	Rank ⁴	I_{geo}^5	mean- $Er > 40^6$	mean-RCR $\geq 1^7$
Ranking of ponds					Elements of concern		
A	3 (11.8)	3 (593)	3 (11.1)	3	Ag, S, Sb, Zn	Ag, Cd, Sb	Cu, Sb, V, Zn
B	1 (18.5)	1 (1248)	1 (17.3)	1	Ag, Cd, Cu, P, Pb, S, Sb, Zn	Ag, Cd, Sb	Ba, Cd ⁸ , Co, Cu, Sb, V, Zn
C	5 (1.0)	4 (134)	4 (6.3)	4	S	None	Co, Ni, V,
D	2 (12.6)	2 (646)	2 (12.3)	2	Ag, Cd, Cu, S, Sb, Zn	Ag, Cd, Sb	Co, Cu, Sb, V, Zn
E	4 (2.1)	5 (117)	5 (3.9)	5	S, Sb	Sb	none
soil	n.a. ⁹	n.a.	5.6	n.a.	n.a.	n.a.	Co, V
List elements of concern					Ag, Cd, Cu, (Pb), Sb, Zn		

506 ¹ Sum I_{geo} in parenthesis, when calculating sum I_{geo} negative I_{geo} values have been set to 0. I_{geo} was
 507 calculated for all elements except Hg which was below detection limit in all samples. Sum I_{geo} has been
 508 calculated to enable comparison in the risk ranking of the ponds.

509 ² Sum Mean Potential Ecological Risk Index in parenthesis, RI was calculate including all elements
 510 except Hg, P and S, see also Table 7.

511 ³ Sum Mean RCR in parenthesis, Sum RCR was calculated for all elements except Hg, P and S, see also
 512 Table 8

513 ⁴ Based upon sum of the 3 ranking methods

514 ⁵ Mean I_{geo} values classified in Class 2 or higher are included, values in Class 3 or higher in bold

515 ⁶ Mean Er values higher than Class 2 are included, values in bold are Class 3 or higher, see Table 7

516 ⁷ mean RCR >2 in bold

517 ⁸ for one sampling site (B4) the RCR is above 1

518 ⁹ n.a. - not applicable, soil is used in calculation of I_{geo} and RI

519

520

521 Conclusions

522 There is an urgent need to update the list of trace elements that are routinely monitored in sediment
 523 in stormwater ponds when evaluating ecological risk, especially in ponds with catchments in areas of
 524 industrial land use. The ponds in this study were mainly constructed to protect downstream areas from
 525 flooding during high flows and therefore have low hydraulic efficiency. Accordingly, some trace
 526 elements that bind to finer particles that did not sediment in the ponds might therefore flow into the
 527 river downstream. Currently, there is no information about background concentrations nor the degree
 528 of pollution of Ag and Sb in the downstream catchment nor in other catchments in the region of study.
 529 Therefore, the development of a program for monitoring numerous trace elements – significantly
 530 more than what is currently monitored - is essential. Identifying the sources of the high accumulation
 531 of Ag and Sb found in this study is another priority. Finally, we showed that the use and comparison of
 532 different assessment methods including multivariate analyses when evaluating ecological risk of

533 stormwater pond sediment is valuable as it highlighted different aspects of risk. For example,
534 information from the geoaccumulation index combined with the multivariate analyses clearly
535 demonstrated that Co, Ni and V were not accumulating and therefore their risk as identified by the
536 RCR-method was overestimated. On the other hand, for Ag, RCRs are below 1 also in the most
537 contaminated pond B while the geoaccumulation index and RI indicates high accumulation and high
538 risk, triggering the need for additional studies.

539 **Declarations**

540 Ethics approval and consent to participate: Not applicable

541 Consent for publication: Not applicable

542 Availability of data and materials: All data generated or analysed during this study are included in this
543 published article and its supplementary information files.

544 Competing interests: The authors declare that they have no competing interests.

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546 Authors' contributions: Sylvia Waara: conceptualization, methodology, investigation, analysis and
547 writing, Frida Johansson: writing, investigation and analysis. Both authors read and approved the
548 final manuscript.

549

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Figures

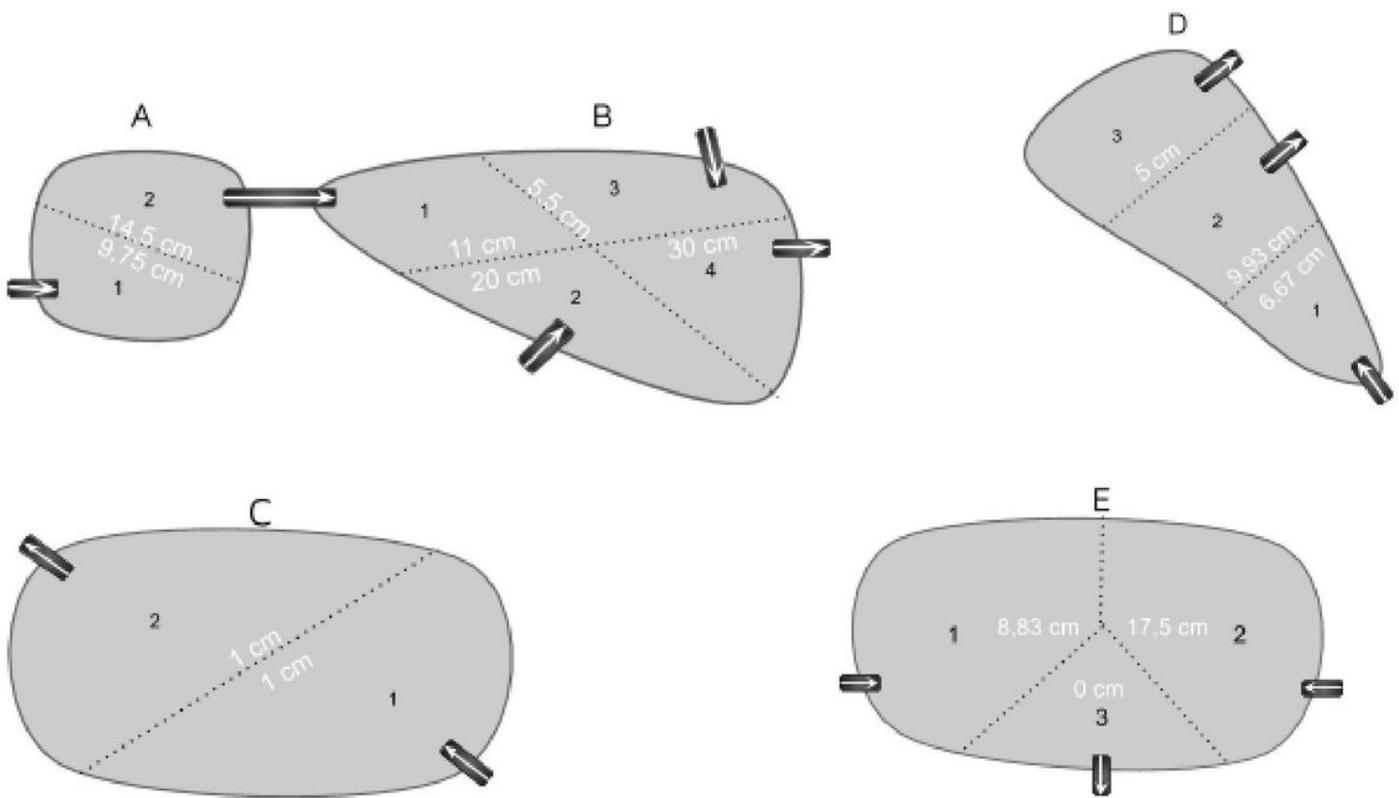


Figure 1

Design of the ponds and the measured thickness of sediment

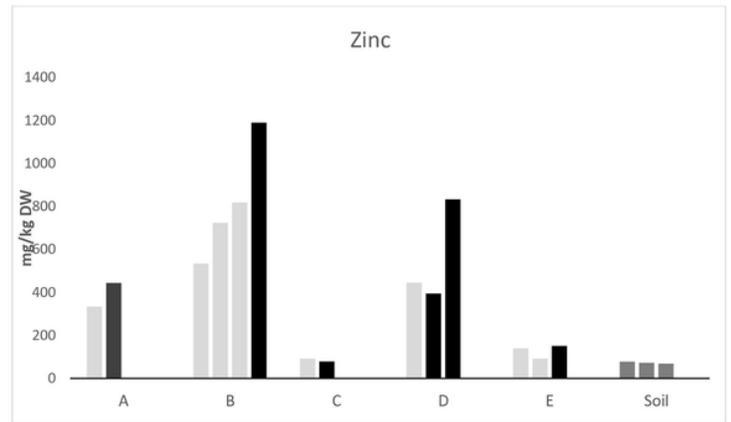
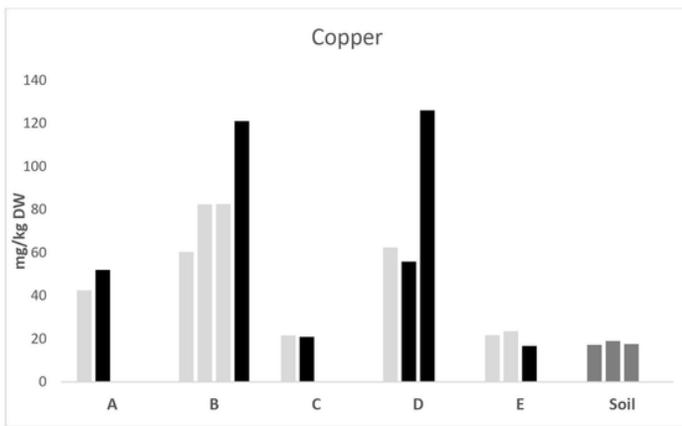
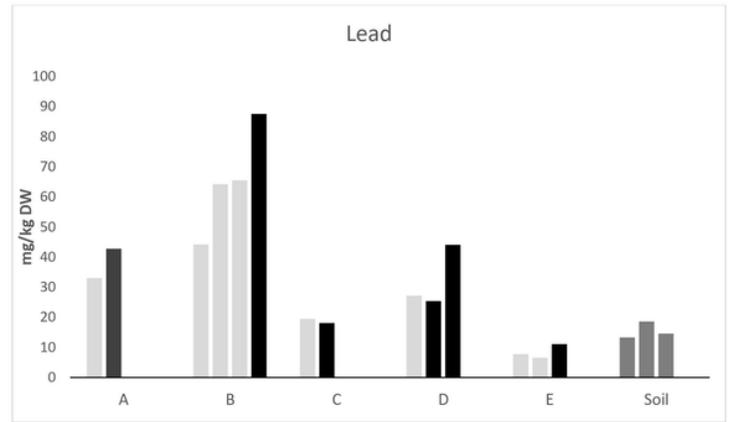
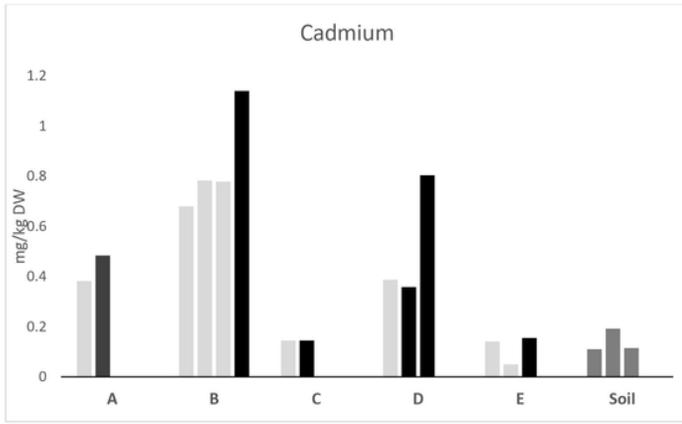


Figure 2

Concentrations of four commonly-monitored trace elements (Cd, Co, Pb, Zn) in sediments in stormwater ponds and soil from the banks of the ponds in areas with industrial activity. Light grey bars – inlets, black bars-outlets.

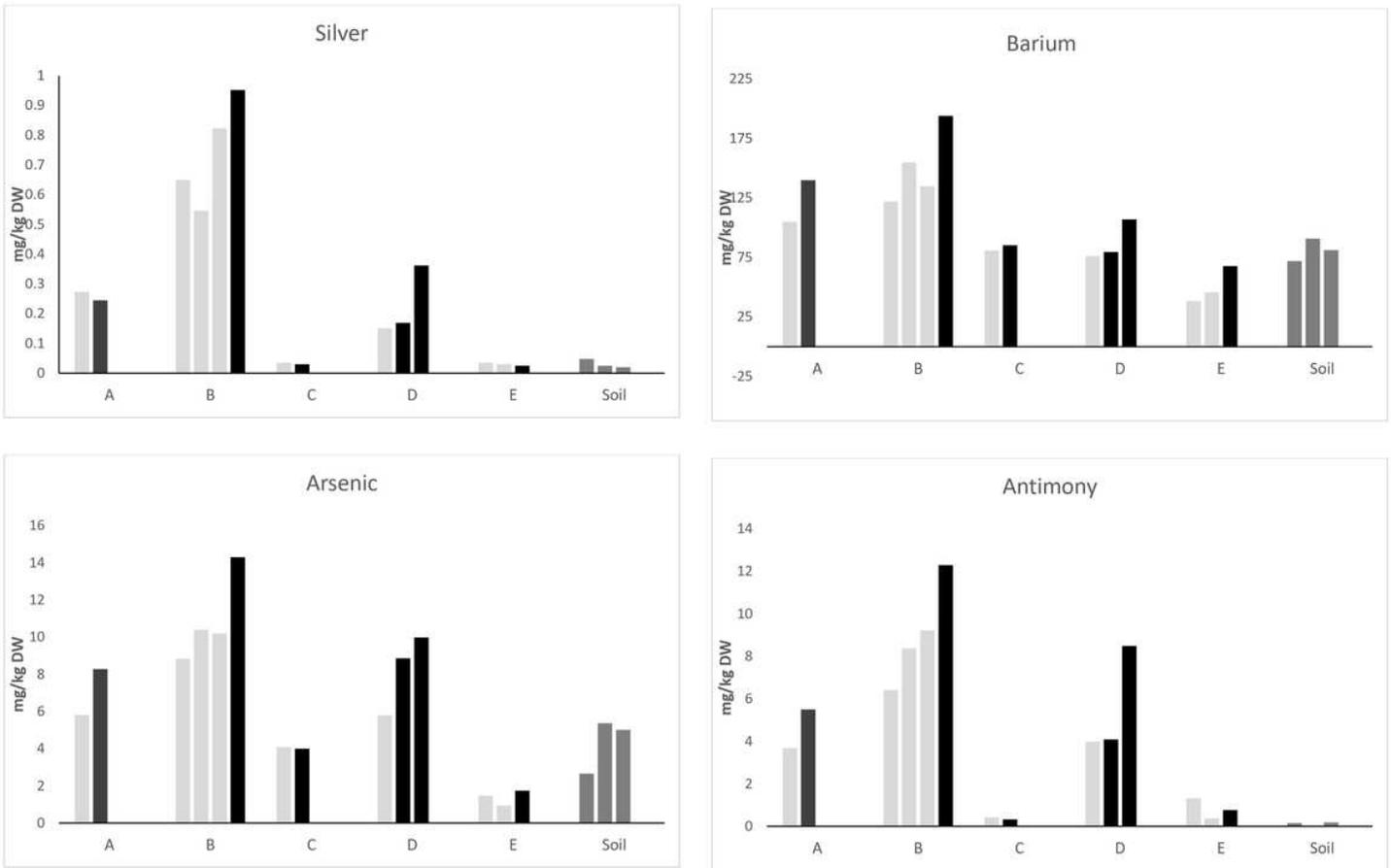


Figure 3

Concentrations of four seldom monitored trace elements (Ag, As, Ba, Sb) in sediment in stormwater ponds and soil from the banks of the ponds in areas with industrial activity. Light grey bars – inlets, black bars-outlets

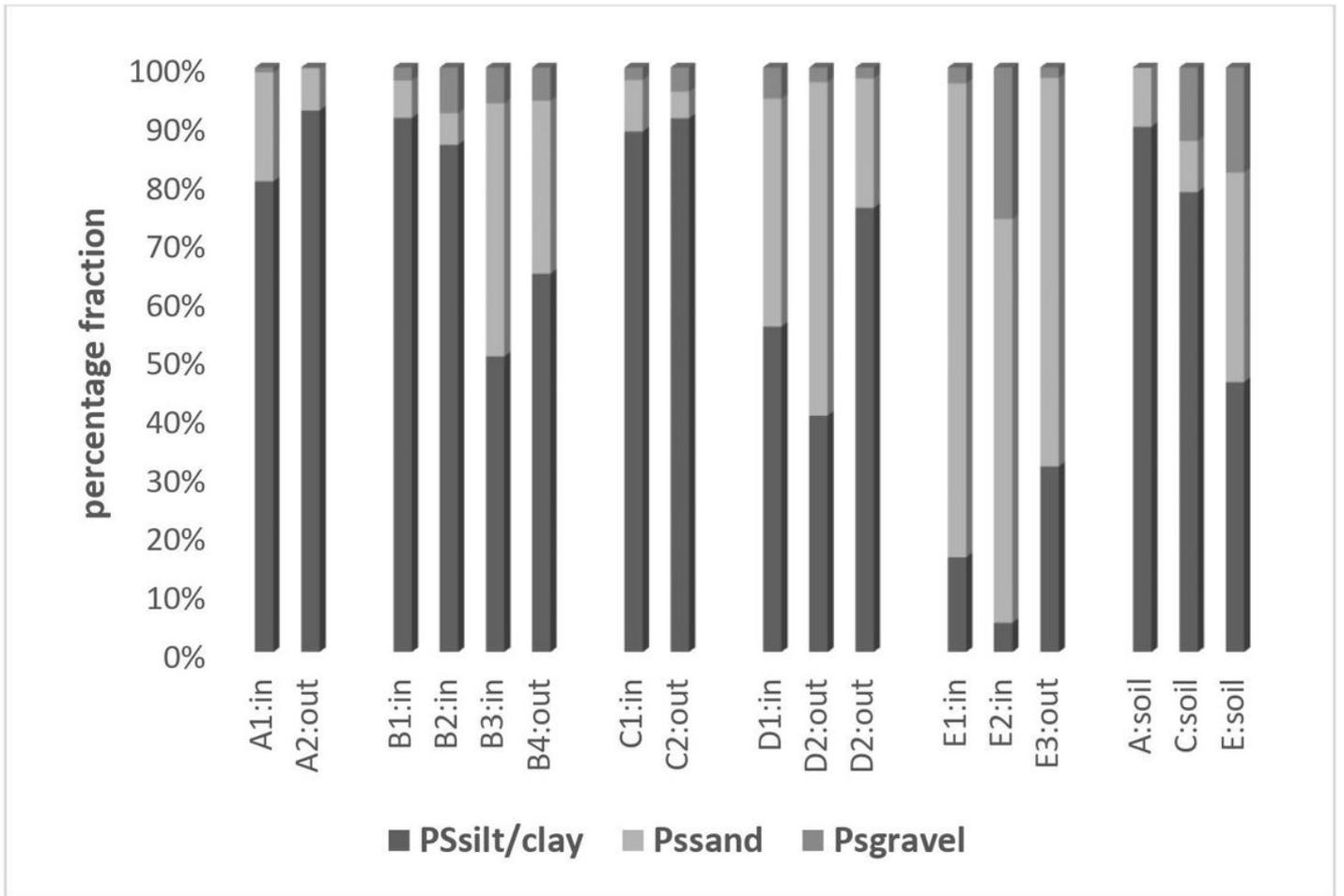


Figure 4

Particle size distribution of the sediment in the different sampling sites in five stormwater ponds. Particle size (PS) for PSsilt/clay < 0.063 mm, PS sand 0.063- 2mm and PS gravel > 2mm

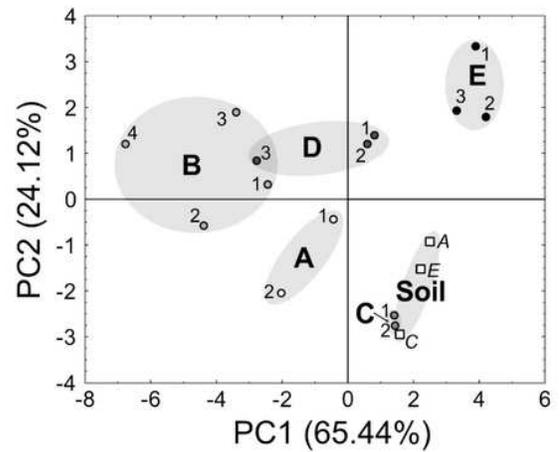
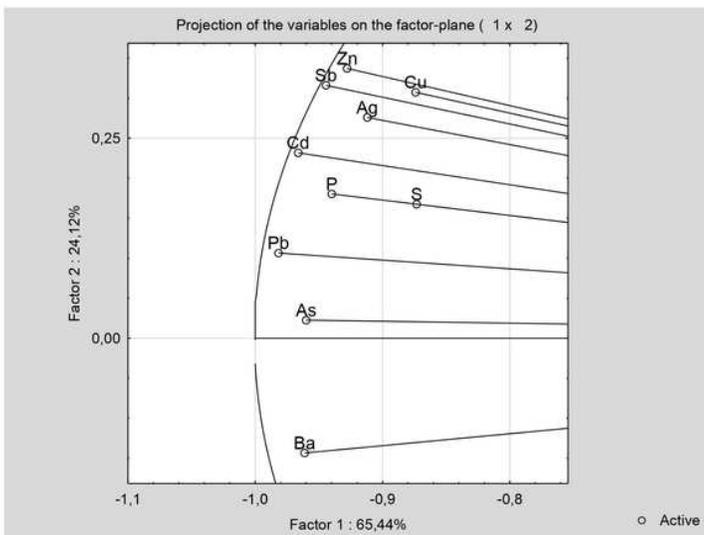
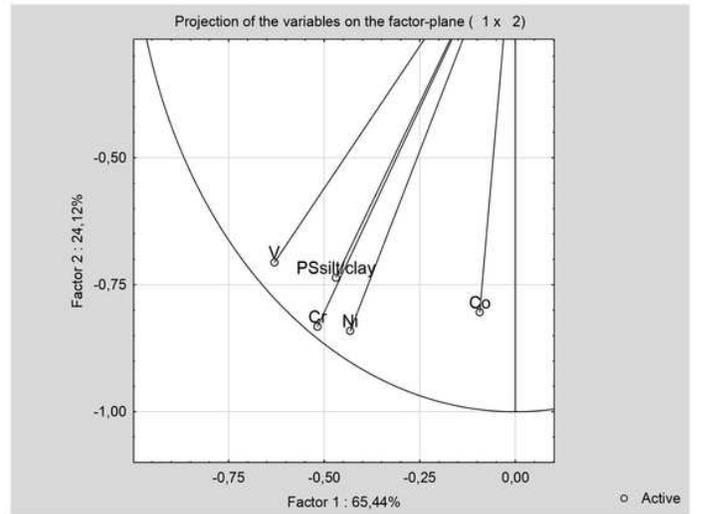
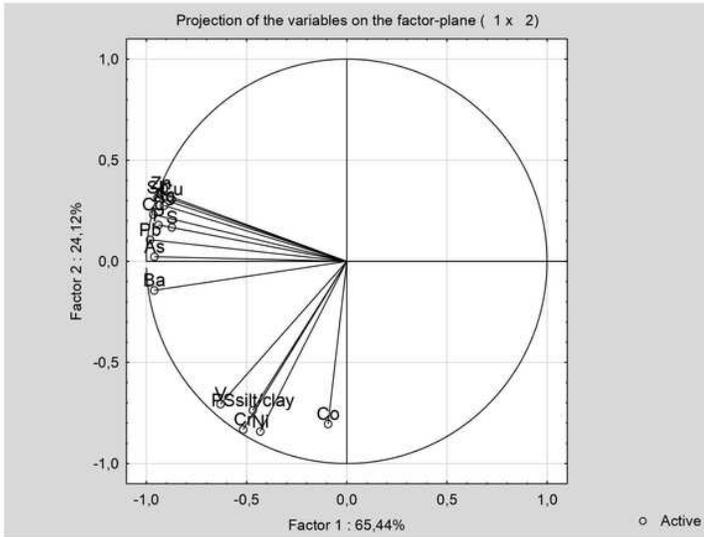


Figure 5

a) Projection of the variables on PC 1 and PC2. b) Enlarged section of PCA with Group A elements c) Enlarged section of PCA with Group B elements and PSsilt/clay d) Projection of the samples on PC1 and PC2

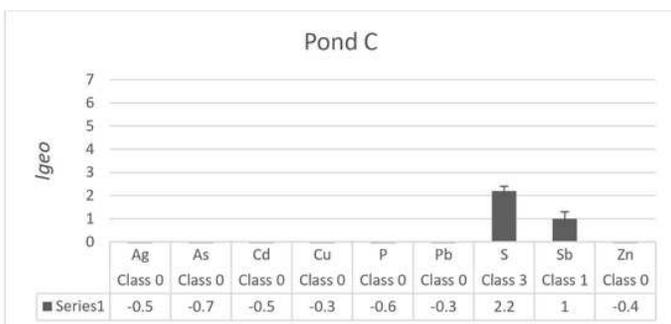
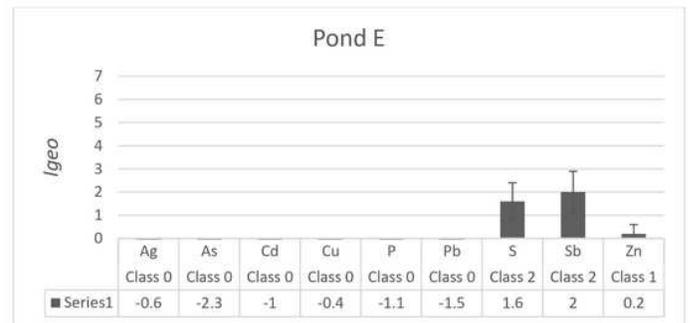
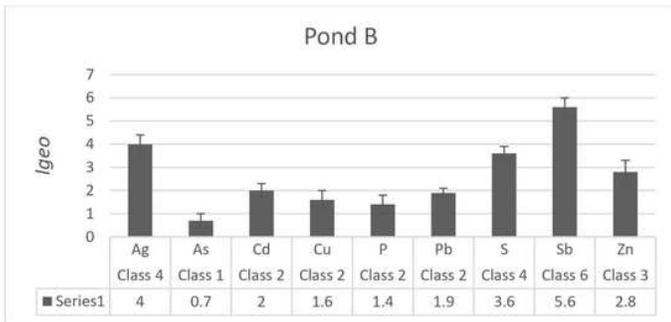
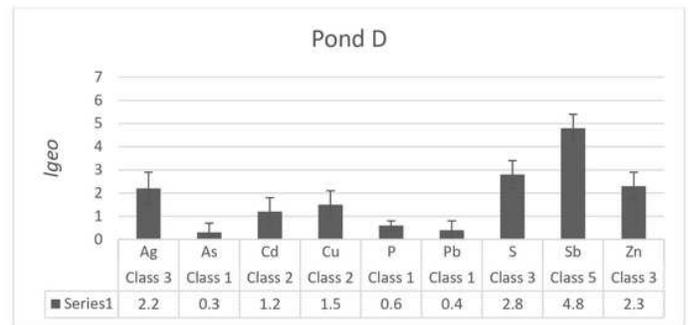
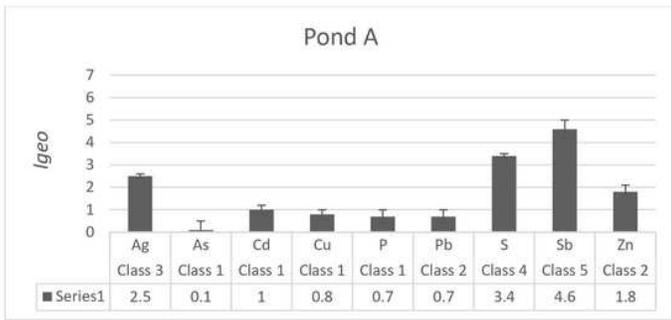


Figure 6

Contamination degree for trace elements in the different ponds using the geoaccumulation index Igeo. Classification was made using mean values, and error bars present standard deviation

Supplementary Files

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