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The effect of a fireworks event on the amount and elemental concentration of deposited dust collected in the city of Debrecen, Hungary --Manuscript Draft--

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Abstract:	<p>Many social celebrations in urban areas are followed by fireworks show. The organic and inorganic pollutants emitted during detonations are expected to affect the ambient air quality of these celebration sites. The environmental aspects of fireworks events are usually investigated by analyzing the concentration and composition of airborne particulate matter, while there is limited information regarding the effect of fireworks on the elemental concentration of deposited dust. In this study foliage dust samples were collected in the city of Debrecen (Hungary) before and after the fireworks show, organized on the 20th of August for the celebration of a historical event. Leaf samples (<i>Tilia tomentosa</i>) were collected around the location of the area of festivities. The sampling sites were further divided into 5 areas: city center (center), Southeast (SE), Southwest (SW), Northeast (NE) and Northwest (NW). We found that the amount of deposited dust particles increased significantly after the fireworks show compared to the background; we also found significant differences in the amount of dust deposition between the different locations of the city. A statistically higher level of Ca, Mg and Sr was detected in samples collected after the display compared to those collected during the previous days, while the concentration of other studied elements were not statistically different from the background level. Our study confirmed previous findings, that the relatively high altitude of detonations allows chemicals to disperse in the fine and ultrafine aerosol fractions, thus, the emitted pollutants by fireworks shows do not increase the level of elements as markedly in deposited dust as in the inhalable fraction.</p>

AIRQ-D-14-00057R1

Dear Professor Chung,

We would like to submit our revised manuscript (*The effect of a fireworks event on the amount and elemental concentration of deposited dust collected in the city of Debrecen, Hungary*) to the journal of Air Quality, Atmosphere and Health.

The manuscript is corrected according to the minor revision.

Please find attached the description of the changes indicated in *italics*.

We are grateful for your editorial work.

Yours sincerely,

Edina Baranyai and the co-authors

Rev #1:

1. In References, the numbers from 1 to 32 should be deleted (see References of a typical paper in AQAH).

Authors: Thank you for the comment, the numbers are deleted from the list of references.

2. We suggest that authors should include a couple of papers in References from AQAH journal.

Authors: We included the proposed references to the manuscript.

1 **The effect of a fireworks event on the amount and elemental concentration of deposited**
2 **dust collected in the city of Debrecen, Hungary**

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16 **Abstract**

17 Many social celebrations in urban areas are followed by fireworks show. The organic and
18 inorganic pollutants emitted during detonations are expected to affect the ambient air quality
19 of these celebration sites. The environmental aspects of fireworks events are usually
20 investigated by analyzing the concentration and composition of airborne particulate matter,
21 while there is limited information regarding the effect of fireworks on the elemental
22 concentration of deposited dust. In this study foliage dust samples were collected in the city of
23 Debrecen (Hungary) before and after the fireworks show, organized on the 20th of August for
24 the celebration of a historical event. Leaf samples (*Tilia tomentosa*) were collected around the
25 location of the area of festivities. The sampling sites were further divided into 5 areas: city
26 center (center), Southeast (SE), Southwest (SW), Northeast (NE) and Northwest (NW). We
27 found that the amount of deposited dust particles increased significantly after the fireworks
28 show compared to the background; we also found significant differences in the amount of
29 dust deposition between the different locations of the city. A statistically higher level of Ca,
30 Mg and Sr was detected in samples collected after the display compared to those collected
31 during the previous days, while the concentration of other studied elements were not

32 statistically different from the background level. Our study confirmed previous findings, that
33 the relatively high altitude of detonations allows chemicals to disperse in the fine and ultrafine
34 aerosol fractions, thus, the emitted pollutants by fireworks shows do not increase the level of
35 elements as markedly in deposited dust as in the inhalable fraction.

36 Keywords: fireworks show, urban dust pollution, elemental analysis, ICP-OES, MP-AES

37 **1. Introduction**

38 Explosive pyrotechnic devices are widely used for celebrating specific events causing an
39 unusual environmental effect on the ambient air quality (Vecchi et al, 2008). They are most
40 frequently used in the already polluted urban areas (Kulshrestha et al., 2004), emitting
41 additional amount of metal particles, gases and various organic compounds which can cause a
42 temporary decrease in air quality (Ravindra et al., 2003). The elemental concentrations of the
43 aerosol particles that reach the atmosphere during fireworks events were studied using
44 different analytical methods (Dutcher et al., 1999; Liu et al., 1997; Drewnick et al, 2005). It
45 was demonstrated that the particulate matter generated by these shows contain Sr, K, V, Ti,
46 Ba, Cu, Pb, Mg, Al, S, Mn and Zn as major components (Perry, 1999). It was found by
47 Kulshrestha et al. (2004) that the concentration of Ba, K, Al and Sr significantly increased
48 during the fireworks of Diwali festival (India) compared to the background values registered
49 on the previous days. This finding was confirmed by Sarkar et al. (2010). Most of these
50 metallic salts serve as color-generating components in pyrotechnic devices and have adverse
51 health effects if inhaled (Murty, 2000; Hirai et al., 2000). Camilleri and Vella (2010) showed
52 that most of the metals contributed by fireworks are in the inhalable dust - finding strong
53 correlations between airborne particulate matter (PM10) and Al, Ba, Cu, Sr and Sb
54 concentrations in airborne dust - hence leading to a strong concern of health risks.

55 The environmental aspects of fireworks shows are usually studied by analyzing the
56 concentration and composition of airborne particulate matter (PM10 and PM2.5) collected
57 with conventional sampling methods (Guttikunda and Kopakka, 2014; Rogula-Kozłowska et
58 al., 2014; Wang et al., 2013; Tsai et al., 2012; Shen et al, 2009). There is no information in
59 the literature on the deposited dust in urban areas from fireworks, in most of the studies the
60 fine and/or inhalable fractions were investigated due to health concerns. However, deposited
61 dust may contain toxic substances affecting plants, the quality of soil and groundwater
62 especially in urban and rural areas (Pelig-Ba et al., 2001; Goossens, 2007).

63 Trees reflect the cumulative effects of environmental pollution from both the soil and the
64 atmosphere (Madejón et al., 2004). Therefore, they have been used as bioindicators in several
65 studies (Celike et al., 2005; Baycu et al., 2006). Mostly plant tissues are analyzed to monitor
66 the accumulation of trace elements confirming their availability in the soil (Baker et al., 2000;
67 Markert et al., 2003) and atmosphere since they can capture and uptake trace elements as well
68 as concentrate them, making small amounts detectable. However, trees can also be used as
69 dust traps in environmental assessments based on urban dust analysis (Margitai and Braun,
70 2005) since deposited contaminants can be trapped by the surface of leaves thus representing
71 the environmental load. Although Simon et al. (2011) concluded that the elemental
72 concentration of foliage dust did not differ remarkably between species; the amount of
73 captured deposited dust highly depends on the surface of leaves, stomata size and density of
74 the selected species. Accordingly, these factors affect their possible use in air quality
75 monitoring investigations (Abbruzzese et al., 2009).

76 Since leaf sampling is widely used in pollution studies due to the inherent variability in
77 crowns (Luyssaert et al., 2002), our aim was to investigate the applicability of this dust
78 sampling for the analysis of deposited dust load and trace element emission resulted by a
79 fireworks event.

80 **2. Materials and methods**

81 *2.1 Sampling procedure*

82 Leaf samples were collected in the city of Debrecen the days before and after the fireworks
83 show on 20th of August (2011), which commemorated a specific historical event in Hungary.
84 Debrecen is the second largest city in Hungary with nearly 225.000 inhabitants. It is the
85 regional center of the Northern Great Plain region and situated 220 km east of the capital,
86 Budapest.

87 The fireworks event was organized simultaneously in two locations in the city center. We
88 divided the sampling sites into five sub-areas as follows: city center (center), Southeast (SE),
89 Southwest (SW), Northeast (NE) and Northwest (NW) (Fig. 1). Silver linden (*Tilia*
90 *tomentosa*) leaves were chosen since the morphology, canopy structure and the dust
91 preserving epicuticular wax on the leaf surface make linden trees ideal for biomonitoring
92 studies (Braun et al., 2007). These trees are widespread in and around the city of Debrecen
93 and can be easily distinguished from other species. We collected the control samples two days
94 before the festivities (18th and 19th of August, 2011) from the individual selected species, the

95 trees were marked and their coordinates were registered. Sample collection from the same 41
1 96 individuals was repeated two days after the fireworks show (21th and 22nd of August, 2011).
2
3 97 The sampling procedure is based on literature values (Moreno et al., 2010). The pyrotechnic
4
5 98 event was performed during stable meteorological conditions and the weather was also dry in
6
7 99 the time of the two sampling periods. The prevailing wind of Debrecen is directed NW-SE
8
9 100 (Lóki et al., 1993). The total leaf surface of each sampled tree represented 10-12 dm².
10
11 101 Samples were collected in paper bags and they were stored at +4°C for analysis.
12

13 102 *2.2 Sample preparation*

15 103 The surface area of leaves was determined by a flat scanner. The foliar dust particles were
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17 104 washed down from leaves by deionized water obtained from a Millipore Synergy Ultrapure
18
19 105 Water System. Leaves collected from the same individuals were placed into 500 ml plastic
20
21 106 container and 250 ml of deionized water was added, then samples were shaken in an
22
23 107 ultrasonic bath (Elma Transsonic, 460/H) for 10 min. The dust containing suspension was
24
25 108 filtered through a 150 µm sieve and the procedure was repeated with 50 ml deionized water.
26
27 109 This process produced 300 ml dust containing suspension which was transferred into a
28
29 110 microwave digestion system (MLS 1200 mega) where its volume was reduced to 20-30 ml.
30
31 111 Then the suspension sample was transferred into 50 ml glass beaker without loss and the rest
32
33 112 of water was evaporated at 105°C in a drying cabinet. The beakers were reweighed to
34
35 113 determine the dry weight of dust on an analytical balance (Precisa 240A). Samples were
36
37 114 prepared prior to analysis in the same vessels by acid digestion using the mixture of 5 ml 65%
38
39 115 (m/m) nitric acid (reagent grade, Merck) and 2 ml 30% (m/m) hydrogen-peroxide (reagent
40
41 116 grade, Scharlau) on an electric hotplate at 80 °C for 4 h. Only those particles were chemically
42
43 117 decomposed, which are not vaporized at temperatures up to 105°C. Digested samples were
44
45 118 diluted up to 10 ml using 1% (m/m) nitric acid (Simon et al. 2011).
46

47 120 *2.3 Elemental analysis of dust samples*

48 121 The determination of Al, As, Ba, B, Ca, Cu, Fe, K, Mg, Mn, Na, P, S and Zn was carried out
49
50 122 by inductively coupled plasma optical emission spectrometry (ICP-OES, IRIS Intrepid II
51
52 123 XSP), while the concentration of Li and Sr was determined by microwave plasma atomic
53
54 124 emission spectrometry (MP-AES 4100, Agilent Technologies). The certificated material used
55
56 125 was ERM-CZ120. The recovery for all elements was under 5%.
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126 Six-point calibration solution series were diluted from multi-element calibration stock
 127 solution of 1000 mg L⁻¹ (Merck ICP multi-element standard solution IV).

128

129 *2.4 Statistical analysis*

130 Data were log transformed prior to analyses. The distribution of data was tested with Shapiro–
 131 Wilk test. Principal Component Analysis (PCA) was used to display the effect of sampling
 132 period (before and after the fireworks event) as well as the sampling sites (center, SE, SW,
 133 NE, NW) on the concentration of the selected elements in the foliage dust. The homogeneity
 134 of variances was tested by Levene’s test. The effect of sampling locations on the amount of
 135 deposited dust on the surface of leaves and the elemental concentration of deposited dust was
 136 studied by General Linear Model (GLM). Tukey’s Multiple Comparison test was used to
 137 explore the significant differences. The t-test was used to compare the amount and elemental
 138 concentration of deposited dust before and after the fireworks show.

139

140 **3. Results**

141 *3.1 Amount of dust deposited on the surface of leaves*

142 According to the t-test there was a significant difference ($t_{80} = - 2.178$; $p = 0.032$) between the
 143 amount of dust deposited on the surface of the collected leaf samples before and after the
 144 fireworks show (Fig. 2). The foliage dust significantly increased based on the GLM results
 145 after the fireworks event in all studied areas except the Southwestern (SW) region of the city,
 146 where the amount of dust was significantly lower after the event. Sampling period and
 147 sampling area had a significant effect (period: $F = 5.274$, $p = 0.025$; area: $F = 5.665$,
 148 $p = 0.001$) on the amount of foliage dust while the interaction effect was not significant
 149 ($F = 1.939$; $p = 0.113$). The highest dust enlargement after the fireworks show occurred in the
 150 Northwestern region (NW) of the city. Considering the amount of deposited dust the
 151 Southwestern area differed significantly from the Southeastern ($p < 0.05$), Northeastern
 152 ($p < 0.05$) and Northwestern ($p < 0.001$) areas of the city both before and after the fireworks
 153 show, but did not differ significantly from the center.

154 *3.2 Elemental concentration of foliage dust*

155 The concentration of the measured elements in the foliage dust in terms of sampling time and
 156 area is indicated in Table 1. The GLM analysis showed significant differences between the

157 sampling period ($F = 4.794$, $p < 0.001$) and among sampling area ($F = 2.636$, $p = 0.002$),
 158 based on the elemental concentration of deposited dust, while their interaction effect was not
 159 significant ($F = 1.279$; $p = 0.112$).

160 *3.2.1 Elemental concentration of foliage dust before the fireworks show*

161 Significant differences occurred in the elemental concentration of foliage dust considering the
 162 sampling areas before the fireworks event. The concentration of Cu was the lowest ($p <$
 163 0.001) in the Northeastern area of the city, while the concentration of Ba was the highest
 164 ($p < 0.001$) in the Northwestern area compared to the other studied locations. The centrum
 165 area differed significantly from the Southeastern area for Al, K and Na ($p < 0.05$) and from
 166 the Northeastern are for Ca ($p < 0.05$) and Mg ($p < 0.001$). The highest Mg concentration was
 167 measured in the Northeastern area, showing a significant difference compared to the others.
 168 The Fe concentration was significantly lower ($p < 0.05$) in the Southwestern and Northeastern
 169 area of the city compared to the other studied locations. The Southeastern area showed a
 170 statistical difference in the Zn ($p < 0.001$) and S ($p < 0.05$) concentrations compared to the
 171 Northeastern area.

172 *3.2.2 Elemental concentration of foliage dust after the fireworks show*

173 According to the t-test the level of Ca ($t_{80} = -3.701$, $p < 0.001$), Mg ($t_{80} = -2.104$, $p = 0.039$)
 174 and Sr ($t_{80} = -2.292$, $p = 0.025$) increased significantly in foliage dust after the fireworks
 175 event, while the concentration of As ($t_{80} = 4.872$, $p < 0.001$) statistically decreased.

176 The GLM results proved that the concentration of Ca, Mg and Sr statistically increased in the
 177 centrum, Southwestern, Northeastern and Northwestern sampling areas, respectively
 178 ($p < 0.005$). A higher level of Al, Ba, B, Cu, Mn, Li, Na and P was also observed in the
 179 centrum, Southwestern, Northeastern and Northwestern sampling areas, although the
 180 difference was not significant. None of the studied elements showed significant concentration
 181 increase in the Southeastern sampling area after the fireworks event.

182 According to PCA there is a clear separation of elements before and after the fireworks show
 183 as well as for the studied areas except for the center. The first component (PC1) contributed
 184 32.9% while the second (PC2) contributed 26.1% to the total variance. As indicated in Fig. 3,
 185 positive correlation was found for Zn, Ba and Al with PC1, while K, B, Li and Sr correlated
 186 negatively with the same axis. Positive correlation occurred for Cu, Fe, Mn, Mg and Ca, while
 187 negative correlation was found for Sr and As with PC2.

188 4. Discussion

1
2
3 189 In the current study, the effect of a fireworks event is investigated by measuring the amount
4
5 190 and elemental concentration of foliage dust deposited on the surface of silver linden tree
6
7 191 leaves in the city of Debrecen. Earlier assessments of the effects of compounds emitted during
8
9 192 fireworks events are quite controversial. It was stated by Perry (1999) that no significant
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11 193 health concern is involved in pyrotechnic displays. There are studies in the literature which
12
13 194 challenge this statement and prove the negative effect of fireworks on the ambient air quality
14
15 195 (Ravindra et al., 2003; Camilleri and Vella, 2010). These shows may cause adverse health and
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17 196 environmental problems. Sarkar et al. (2010) investigated the amount of fine, respirable
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19 197 particles (PM₁₀) during a major fireworks festival in India and concluded, that the level of
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21 198 24h PM₁₀ was extremely high. Moreno et al. (2010) analyzed the background level of
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23 199 ultrafine particles during a fireworks pollution episode in Girona (Spain) and observed a
24
25 200 highly increased level of 24h PM_{2.5}. However, no information is found in the literature about
26
27 201 the relation of fireworks displays and the amount of deposited dust. In our study the overall
28
29 202 amount of foliage dust was significantly higher after the fireworks compared to the
30
31 203 background amount collected previously. Considering the sampling areas, the amount of
32
33 204 deposited dust statistically increased after the event in all studied areas except the
34
35 205 Southwestern part of the city. This phenomenon may be the consequence of the particular
36
37 206 climate and relief of Debrecen. The prevailing wind of Debrecen is directed NW-SE and in
38
39 207 the northwestern part of the city a woodland is located protecting that part from strong wind
40
41 208 blows (Lóki et al., 1993). On the contrary, the Southwestern region of the city is an open field
42
43 209 district with the lack of continuous tree borders and with wind tunnel effect of high buildings.
44
45 210 Probably this is the reason for the decreased dust amount in the Southwestern sampling area.

46
47 211
48 212 During fireworks occasions the concentration of elements such as Sr, Mg, K, Ba and Cu is
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50 213 reported to be remarkably higher in airborne dust (Wang et al., 2007; Moreno et al., 2010)
51
52 214 indicating the presence of highly toxic metals in fireworks devices. In the current study the
53
54 215 level of Ca, Mg and Sr increased statistically. These elements are used as color generating
55
56 216 agents in pyrotechnic devices. We found no significant increase in the concentrations of Al,
57
58 217 Ba, B, Cu, Mn, Li, Na elements after the fireworks event, and their concentrations after the
59
60 218 fireworks show are far below than reported by other research groups (Ravindra et al., 2003).
61
62 219 These results are in contrast to our previous hypothesis according to which we would expect
63
64 220 high emission of these toxic elements after the fireworks event. Yet this finding is very
65

221 similar to what Vecchi et al. (2008) concluded. They found significantly higher elemental
222 concentration in PM10 fraction but they detected no increase in the coarse fraction of dust
223 particles: the level of Sr, Mg, K, Ba and Cu were below or comparable after the fireworks
224 event in the coarse fraction of collected aerosol particles during a fireworks episode in Milan.
225 They concluded, that the ambient aerosol after the fireworks event was preferably confined in
226 the fine fraction (Vecchi et al. 2008). Crespo et al. (2012) gained very similar findings. They
227 discussed that the firework related metals were concentrated in the submicrometric region
228 (>80%) of the collected dust particles. Our study proves these statements since only Ca, Mg
229 and Sr were in significantly higher concentrations; other differences were not detected in toxic
230 metal concentrations in deposited dust after the fireworks show. Therefore, we further
231 conclude, that pyrotechnically derived aerosol particles containing inorganic pollutants are in
232 the fine fraction, while the heavy fraction of dust particles depositing after the show did not
233 pose environmental risk with respect to toxic elements.

234
235 The background elemental concentration of foliage dust showed differences between the
236 studied areas. The southeastern part of the city is the most contaminated considering the
237 measured elements, where Al, Cu, Mn, Na, P and Zn is in the highest concentration.
238 Mingorance and Oliva (2006) concluded, that the accumulation of toxic elements in leaf
239 samples depends on the urbanization levels. The Southeastern part of Debrecen represents the
240 highest level of urbanization, where the Debrecen Airport and also an industrial area are
241 located with heavy traffic load, which explain the statistically higher amount of pollutants in
242 foliage dust. The Southwestern and centrum areas proved to be the less contaminated with the
243 measured elements based on the foliage dust analysis. The centrum area is in the city center,
244 where a long pedestrian street is located with nearly no traffic load. The Southwestern part of
245 Debrecen is characterized by a green belt with garden houses and small open-space parks,
246 making this area less affected by human activities. The K concentration was the highest in
247 these two sampling areas, which is in good agreement with Simon et al. (2011) previous
248 findings. They analyzed foliage dust and leaf samples in and around the city of Vienna
249 (Austria) along an urbanization gradient and measured a significantly higher K level in the
250 rural sampling area, which represented an undisturbed part of the city with decreased human
251 impact. Beddows et al. (2004) stated that potassium is a crustal component and as a major
252 trace element plays an important role in plant grow hence it occurs frequently in rural areas. It
253 was also found by Beddows et al. (2004) that 87% of coarse particles of rural background
254 atmospheric aerosol collected in the UK contained potassium. The origin of the higher K

255 concentration in the Southwestern region may be from the biomass burning as there is a
 256 power plant located near that part of the city.

257 **5. Conclusions**

258 The fireworks show in Debrecen resulted in a higher amount of dust particles deposited on the
 259 surface of tree leaves. Our findings suggest that the relatively high altitude of detonations
 260 allow the chemicals to disperse in the fine and ultrafine fractions. Thus, emitted pollutants by
 261 firework displays did not elevate the concentration levels of elements in deposited dust since
 262 the background concentration of these inorganic components in heavy dust is already high in
 263 polluted cities like Debrecen. Pyrotechnic detonations therefore cause environmental risk by
 264 the elevated amount of dust deposition but do not contribute significantly to the background
 265 concentration of toxic metals in foliage dust.

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371 **Figure legends**

372 **Figure 1.** Locations of the 41 sampling sites and the fireworks events indicated on the city
 373 map of Debrecen.

374 **Figure 2.** The amount of foliage dust before and after the fireworks show in terms of the
 375 sampling sites given in $\mu\text{g cm}^{-2}$, (mean \pm SD). Notations: open column - before fireworks,
 376 hatched column - after fireworks. SE – Southeast, SW – Southwest, NE – Northeast, NW –
 377 Northwest.

378 **Figure 3.** Principal component biplot of the measured elements in the foliage dust of the
 379 sampling sites before and after the fireworks show. Notations: center - city center, SE -
 380 Southeast, SW - Southwest, NE - Northeast, NW - Northwest. Numbers after the notations
 381 indicate samples collected before (1) and after (2) the fireworks event.

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Table 1. Concentration of the measured elements given in mg kg⁻¹ (mean ± SD) in foliage dust of sampling sites before and after the fireworks show. Notations: CENTER - city center, SE – Southeast, SW – Southwest, NE - Northeast, NW - Northwest.

elements (mg kg ⁻¹)	CENTER		SE		SW		NE		NW	
	before	after	before	after	before	after	before	after	before	after
Al	4664±1291	5023±703	6753±1649	6018±1294	5250±1851	5941±805	4890±905	5700±1211	5955±1112	6244±1463
As	41±181	5±2	66±16	10±7	40±24	8±2	16±7	10±8	18±9	19±3
Ba	124±53	126±53	133±41	114±37	77±16	89±13	92±15	102±12	234±164	262±225
B	650±897	924±1822	819±793	752±805	510±423	750±736	963±516	1251±744	290±305	420±199
Ca	23870±5908	27594±6668	27808±6694	25310±6683	24874±5282	34357±5643	31179±9899	40483±4968	23419±7123	37417±6638
Cu	134±97	151±46	174±125	126±99	115±73	100±54	88±12	88±41	168±56	159±74
Fe	11024±2478	12561±2966	11281±2275	11316±2957	7543±1923	9217±1877	7605±1524	9823±2586	12101±4220	11203±4348
K	37334±8892	34704±12778	31408±12185	22159±13539	36482±11436	40638±16038	32215±17175	29003±18054	24581±9681	26908±11944
Li	14±9	13±12	20±8	18±8	19±16	28±33	61±106	64±116	14.2±2.5	15±7
Mg	5236±1433	5870±1694	6604±1626	5734±1756	6140±1143	7124±1400	9206±5898	9845±5323	4634±783	7692±727
Mn	333±80	364±73	458±131	426±99	375±126	470±133	404±26	497±68	421±99	404±91
Na	5906±4439	5535±5021	20304±22431	17144±18594	4654±4159	6583±8908	3504±1625	3718±1573	7243±1473	8230±8948
P	17657±4033	15805±3853	22105±5117	15079±3186	17382±9690	19507±4454	14433±2212	16970±4856	16933±3301	21469±6313
S	11809±3687	10616±3812	17450±3567	9417±2060	11815±5913	10366±2690	9089±1081	9827±1372	10959±3756	14265±2588
Sr	156±50	201±63	225±57	194±33	202±72	224±63	298±155	325±167	139±46	213±52
Zn	390±110	334±59	646±315	346±159	327±189	288±49	241±70	277±64	430±138	425±151

Figure

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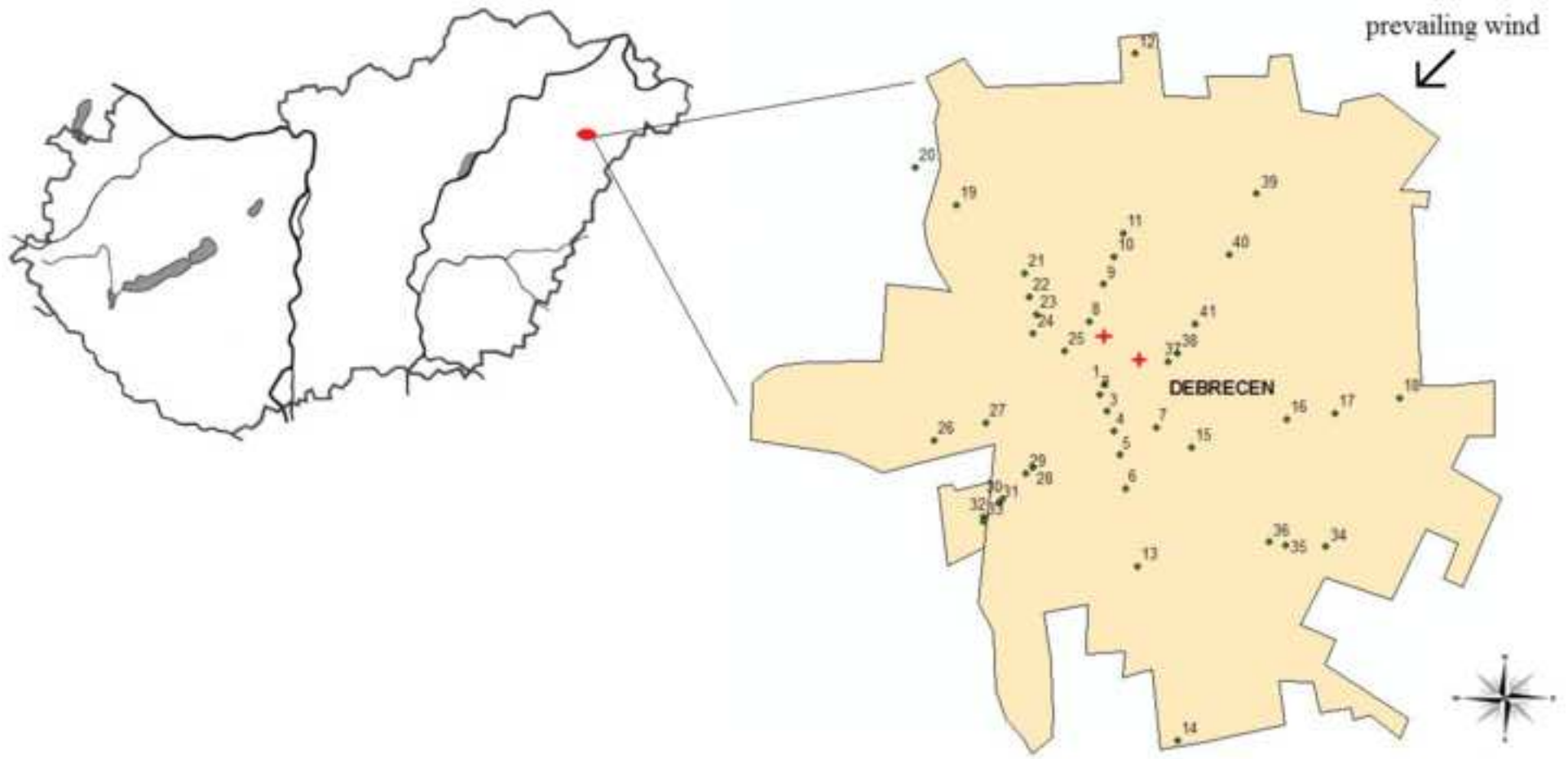


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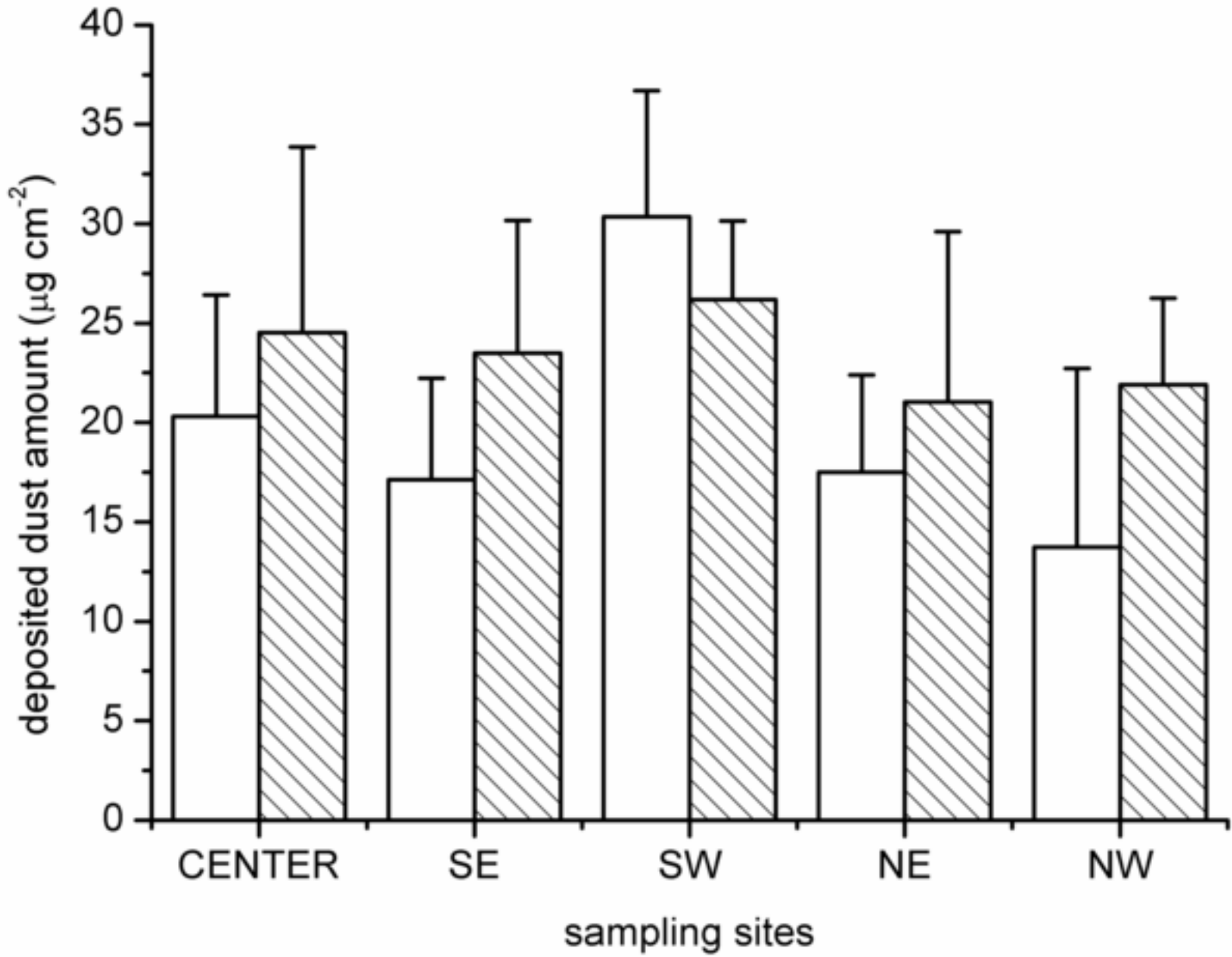


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