












Spatial Modeling of Cement Dust Pollution on Soil Properties in Western Iraq

Md Munir Hayet Khan¹, Saif Saad Mansoor², M. M. Yacoub³, Alaa H. AbdUlameer⁴,
Salem M. A. Al-Ani², Ammar Hatem Kamel^{2,5}, Jumaa Mohammed Ahmad², Haitham Abdulmohsin Afan²,
Faidhalrahman Khaleel^{6*}

¹ Faculty of Engineering & Quantity Surveying (FEQS), INTI International University (INTI-IU), Nilai 71800, Malaysia

² Upper Euphrates Center for Sustainable Development Research, University of Anbar, Ramadi 31001, Iraq

³ College of Applied Science, University of Anbar, Heet 31007, Iraq

⁴ Building and Construction Techniques Engineering Department, College of Engineering and Technologies, Al-Mustaqbal University, Babylon 51001, Iraq

⁵ Dams and Water Resources Engineering Department, College of Engineering, University of Anbar, Anbar 31001, Iraq

⁶ Computer Science Department, College of Sciences, Al-Maarif University, Ramadi 31001, Iraq

Corresponding Author Email: faidhalrahman.mohammed@uoa.edu.iq

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ABSTRACT

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This research examines the effect of cement dust pollution on soil properties in Iraq's arid western region, particularly in the context of the area surrounding the Kubaisa Cement Plant. Spatial modeling methods were used by researchers to collect and analyze 32 soil samples at two different distances from the plant to assess chemical and physical changes to soil properties. Major findings showed higher concentrations of heavy metals like lead and lithium, higher alkalinity levels of soil, and higher particulate matter and CO₂ concentrations close to the source of the pollution. Even with present pollutants, ecological and environmental indices revealed low levels of contamination and ecological risk, on the whole. The findings indicate the continuing effects of industrial emissions on soil integrity and establish the necessity for focused measures to avert ecological and public health risks and help protect the environment.

1. INTRODUCTION

Considering its correlation with human existence, the environmental consequences are significant in our everyday existence. Experts have closely observed industrial operations and their detrimental effects on the surroundings to tackle the issues arising from these activities and promote environmental consciousness in developing localized industrial projects.

The cement industry is an unavoidable polluter of the environment, as it emits polluting gases such as nitrogen oxides, sulfur oxides, carbon dioxide, and carbon monoxide, in addition to the particles carried with combustion gases in the form of small-diameter dust. These pollutants significantly pollute the surrounding surroundings where these factories use black oil, one of the heaviest fuel types due to its high sulfur content, as fuel [1].

Heavy elements are environmental pollutants that spread throughout the surroundings (air, water, and soil). These parts directly and indirectly contribute to human and animal health by influencing the growth of plants that sustain living organisms. Heavy element pollution is important because its products cannot decompose, unlike other chemical pollutants [2].

Kubaisa Cement Factory was chosen because it is located within the central unit in the northern Badia desert. It is also within the secondary area: the lower valley plains. The area's

height ranges from 297.4 m above sea level. This area's lands consist of successive limestone and gypsum deposits in their layers [3].

These elements decompose due to various environmental influences, such as chemical reactions caused by heat, humidity, sunlight, or biological factors, which often reduce their toxicity. Heavy elements enter the natural cycle and move between living and non-living components of the environment, where their bioaccumulation occurs [4]. Its molecules in the air are subjected to a dilution process due to their vertical and horizontal spread. Still, their deposition on the ground leads to soil pollution and their concentration in it. Living organisms in the aquatic surroundings transfer them to other organisms in the water and on land, ultimately reaching their highest concentrations in animals at the top of the food chain, particularly humans [5]. The concentration of heavy elements in soil depends on several factors, some natural and others resulting from human activity, such as organic matter, pH, salinity, and others [6].

Cement dust impacts tree growth metrics (their height, number of leaves, weight, and area) as the dust settles on their leaves, blocking the pores and reducing gas exchange, forming a soft layer that leads to the fall of leaves and drying of plants, and also leads to the death of parts of the trees, gradually reducing the productivity of the plant and the soil [7, 8]. Researchers have discovered that cement dust has an impact

on soil properties. Numerous studies in this field have demonstrated its effects on soil properties, such as soil electrical conductivity (EC) and pH. The study also found an increase in soil alkalinity, possibly accelerating plant weakening due to cement dust's nutrient imbalance [8, 9]. They also confirmed that when cement dust comes into contact with water, it forms a hard crust on the soil's surface. This changes the soil's chemical, physical, and biological properties because it makes it less permeable to water and better able to hold onto it. It also raises the pH level because calcium hydroxide and aluminum hydroxide are formed. Further elucidated is that incorporating cement dust into the soil diminishes its organic carbon content, thereby impeding the activity of soil microorganisms. Additionally, the nitrogen concentration in the contaminated soil differs from that of the reference soil [10]. AKokatnur and Saviramath [11] studied cement dust's impact on soil, discovering that it directly contributes to strong basicity in the soil's surface layer. The study revealed that the primary constituents of cement dust, S, Na, Al, Ca, K, and Mg, were more prevalent in the soil near the cement factory, particularly in the soil's surface layer.

1.1 Study area

Anbar province is located in the arid region of western Iraq with a 33.43 N latitude and 43.33 E longitude (Figure 1), with a 138.57 km total area. It is a critical area facing significant challenges because of climate change. The average altitude is 384.64 m, with extreme temperatures of 40°C in the summer to 9°C in the winter, with scarce precipitation, great contrasting temperatures between day and night, and relative humidity (19 - 82%). Winds are south-westerly and north-westerly, with a maximum speed of about 21 m/s. The rainy season is from September to May, with an average annual rainfall of about 115 mm. 49.5% occurs between November and January, 36.3% between February and April, and 14.8% between September and October. The average annual evaporation is about 3000 mm, which raises the dryness coefficient (evaporation/rain) by 25–35. As a result of climate change and lack of rain, estimates from some studies and international organizations indicate that the average number of dusty days will reach 300 days/year.



Figure 1. The location of the study area

International reports indicate that Iraq is the fifth most affected country by climate change, which is dramatically affecting the environment. Concentrations of pollutants, especially those related to air pollution, have increased significantly. Cement factories are among the primary sources of pollution in the governorate, such as the Kubaisa Cement Factory, located approximately 70 km from Ramadi, the center of Anbar Governorate (Figure 1).

1.2 Heavy metals

Heavy metals are among the most dangerous environmental pollutants, as their continued emission leads to their increasing accumulation in the soil. Some elements, such as iron and copper, are essential for human health, while others are toxic to living organisms [12]. Heavy metals are characterized by their high specific gravity and include elements with a relatively high density, atomic number, or atomic mass, such as transition elements, lanthanides, and actinides. The ecosystem naturally contains these elements, but their concentrations vary greatly, and their recent increase in levels

has resulted in industrial pollution and industrial wastewater [13]. Metals are commonly associated with biological compounds containing oxygen, sulfur, and nitrogen, which can disrupt specific enzyme systems. This is particularly true for enzymes directly or indirectly involved in the production of ATP. In acute poisoning, large excesses of metal ions can disrupt membrane and mitochondrial function and generate free radicals. In most cases, this results in general weakness and malaise [14].

Air pollution sources are divided according to their type into natural sources and sources from human activity. It is sometimes difficult to separate between these two sources, as what is emitted by human activity in one area may be supported by a natural source such as wind in another location, making it difficult to determine the origin of the source. In industrialized countries, factories emit many types of air pollutants. The type varies according to the industry; some emit harmful gases, while others emit dust containing solid pollutants [15]. Both the permitted [15-19] local determinants (Iraq 2012) and the permitted local determinants (NAAQS) have been explained as shown in Table 1.

Table 1. Some permissible limits for some pollutants

Air Pollutants	Allowed Local Parameters (Iraq 2012)	Locally Allowed Queuing Specifiers (NAAQS)
Carbon monoxide (CO)	ppm/ hour 35 ppm/24-hour 9	ppm/ hour 9 ug/m ³ / hour 100
Carbon dioxide (CO ₂)	-	-
Sulfur dioxide (SO ₂)	ppm/24-hour 0.01 ppm/ hour 0.002	ppm/ hour 0.01 ug/m ³ /24 hour 0.03
Ozone (O ₃)	ppm/ hour 0.12	ppm 0.12 / hour
Hydrogen sulfide (H ₂)	ppm/ hour 5	ppm/ hour 5
Nitrogen dioxide (NO ₂)	0.1 ppm/ year ppm /24 hour 0.05	0.11 ppm/year ppm/24 hour 0.03
Total suspended minutes (T.S.P)	350 ug/m ³ / 24 hour 150 ug/m ³ / year	150 ug/m ³ / 24 hour 60 - 90 ug/m ³ /year
Lead Pb	0.5 ug/m ³ /year	2 ug/m ³ / 24 hour 1.5 ug/m ³ / 3 month

Soil pollution is the destruction of the soil's surface layer due to the deposition of complex chemical compounds or reactive synthetic materials. This leads to changes in the natural properties of environmental elements. Soil retains lead as hydrated oxides or organic complexes, with mobility determined by organic content and pH level. As the lead resistance period increases, its biological availability decreases in humans and exposed organisms [19]. Heavy metal contamination primarily impacts soil through pedogenetic processes, where parent materials weather at low levels that are seldom harmful. The buildup of heavy metals in soil layers is typically driven by human activities that disturb and hasten the natural geochemical cycle [20]. In the last three decades, the soil's buildup of these harmful substances (heavy metals) has escalated significantly, threatening both the ecosystem and human health. Pollution from heavy metals has seriously affected people, impacting the environment in multiple ways and leading to industrial unrest in several nations. Azher et al. [2] have reviewed several studies, which evaluate and analyze the numerous approaches for preparing heavy metal-free soil by adopting different soil remediation methods.

Franzaring et al. [21] stated that the critical limits for lead (Pb) in agricultural soils of some countries range from 50–300 ppm, while background concentrations of lithium in soil range from less than 1 to 200 mg/kg [22, 23], with higher concentrations in arid and saline soils [24]. Lithium generally enters the soil solution primarily through weathering sedimentary minerals [22, 25], and surface soils typically contain less lithium than the underlying layers [26]. The clay fraction of the soil contains higher concentrations of lithium than the organic soil fraction [27], with lithium found in biconcave cavities within clay minerals [28].

1.3 Research problem

The vicinity of the Kubaisa cement factory in Anbar Governorate faces significant environmental challenges resulting from increased soil pollution, which is directly threatening the surroundings and public health. This pollution requires a comprehensive study to determine its scope and different types, in addition to analyzing the environmental characteristics of the affected soil. The present study aims to use spatial modeling to assess the distribution and characteristics of pollution in the soil surrounding the factory to provide accurate data that helps develop effective strategies to reduce negative environmental impacts and improve soil quality in the region. Research has proven that cement dust impacts the soil's electrical conductivity (EC) and pH levels.

1.4 Study objectives

The study's objectives are as follows:

- 1) Detecting the most important environmental effects of cement dust on the soil adjacent to cement factories.
- 2) Identifying and diagnosing some metals and heavy elements in the soil surrounding the Kubaisa Cement Factory.
- 3) Evaluating the level of pollution in the soil surrounding the Kubaisa Cement Factory using spatial modeling techniques to determine the extent of the spread of pollutants and their type in the area.
- 4) Analyzing the environmental characteristics of the affected soil in order to understand the environmental impacts resulting from pollution on the local ecosystem and public health.
- 5) Providing an accurate scientific database contributes to developing effective strategies to reduce pollution and improve soil quality in the region.
- 6) Proposing solutions and remedial measures to reduce the negative environmental impacts resulting from industrial processes in the factory is crucial to preserving the surroundings and achieving sustainable development in the region [29].

2. MATERIALS AND LABORATORY WORK

2.1 Data collection

Soil samples were taken from the vicinity of the present study's Kubaisa cement factory. After obtaining the laboratory's location from the coordinates, a preliminary site survey was conducted to determine the changes in the area. A field visit was conducted to the Kubaisa Cement Factory, and a soil sample was taken from 8 sites and two complete circular perimeters (Figure 2), which were coded (A and E) from a distance of 100 meters and 200 meters and on the perimeter of the Kubaisa Cement Factory, with three replicates. Thus, the number of samples is 32 samples. A soil sample was taken from two different depths, 0–15 cm and 15–30 cm.

Several tools were used to take samples from the soil and portable air pollutant measuring devices, including a coordinate finder (GPS), auger tool, plastic bags, CO₂ air pollution meter, and H₂S and particulate air pollution meter. Furthermore, two types of sites were selected for the model as follows:

- The sites not affected by the materials emitted from the factory exhaust are those outside the influence of the factory deposits. These sites include four paths in the directions of north (N), east (E), south (S), and west (W), with distances of 100 and 200, respectively.
- The affected locations in the factory exhaust (the southeastern part of the factory) are because the prevailing winds in the area are north-westerly, and the resulting points of intersection of the distances specified above included the direction of the factory exhaust.

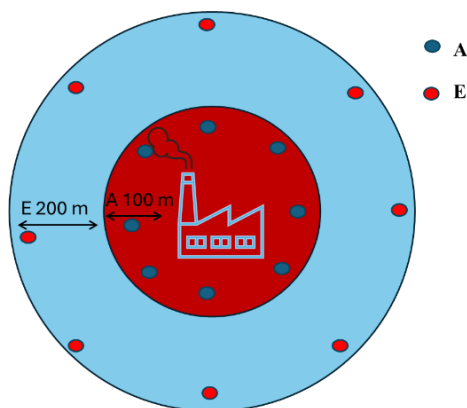


Figure 2. The locations of the selected models for the study

2.2 Laboratory work

After the data collection process, the samples were taken to a private laboratory for the necessary tests. The laboratory work was conducted at the University of Anbar in the College of Applied Sciences-Heet. First, soil samples were dried using an electric oven (The manufacturing company is Memmert) at a temperature of (105) Celsius. Then, the soil extraction process was carried out at an extraction ratio of 1:1 (water: soil), i.e., 100 ml of water was taken for every 100 g of soil for all 32 samples, and then these water extracts were kept in opaque bottles to conduct some measurements such as pH, electrical conductivity, heavy metals (calcium, potassium, and lithium), and finally the soil digestion, which were carried out utilizing the following steps:

- The sample was placed in a beaker, washed with distilled water, dried, and then put in the oven at a temperature of (105) degrees Celsius for an hour to dry.
- Weight 1 gm of the dried sample with a sensitive balance and place it in a clean beaker with a capacity of 250.
- 10 mL of aqua regia was added in a ratio of 3:1, consisting of 7.5 mL of concentrated hydrochloric acid HCL and 2.5 mL of concentrated nitric acid HNO₃.
- Heating the sample to 80 degrees Fahrenheit until the coffee vapors stop appearing, noting that the sample did not dry out.
- Cooling the beaker, add 5 mL of concentrated HCL acid with 50 mL of hot distilled water to wash the sides of the beaker from traces of the dissolved sample.
- Heating the mixture to boiling point for 2–3 minutes.
- Filtering the mixture with filter paper, place the filtrate in a 50 ml volumetric bottle and fill the bottle with distilled water to the mark.
- The samples were stored in opaque polyethylene bottles at 4°C until the atomic absorption device examined them.

2.3 Environmental index

2.3.1 Geochemical Index (I_{geo})

This index is widely used to assess the degree of pollution by comparing heavy metals obtained with the background levels initially used with the bottom sediments. It can also be applied to determine soil pollution, as shown in Eq. (1).

$$I_{geo} = \log \left(\frac{C_n}{1.5 B_n} \right) \quad (1)$$

where, C_n represents the concentration of the element in the

enriched samples. B_n represents the concentration of the background soil component or the element's initial value. The constant is 1.5 to reduce the effect of possible differences in background values that can be attributed to rock variations.

This method evaluates the degree of metal contamination in terms of seven degrees, based on the increasing values of the index, as shown in Table 2.

Table 2. Values and degree of geological accumulation coefficient

I_{geo}	Degree	Pollution Level
≤ 0	0	Unpolluted
0-1	1	Slightly polluted
1-2	2	Moderately polluted
2-3	3	Moderately severely polluted
3-4	4	Severely polluted
4-5	5	Severely extremely polluted
> 5	6	Extremely polluted

2.3.2 Contamination factor (C_F)

The level of sediment contamination with metals is expressed in terms of the contamination factor (C_F), and it is calculated as in Eq. (2).

$$C_F = \frac{C_{metal}}{C_{background}} \quad (2)$$

C_{metal} is the concentration of a particular metal in river sediments. $C_{background}$ is the background metal concentration. The C_F values that describe the pollution are shown in Table 3.

Table 3. Values and degree of contamination factor

Degree	Pollution Level
< 1	Low contamination
$1 \leq C_F < 3$	Moderate contamination
$3 \leq C_F < 6$	Considerable contamination
> 6	Very high contamination

2.3.3 Potential Ecological Risk Index (E_r^i)

The Potential Ecological Risk Index (E_r^i) was calculated using Eq. (3), as follows:

$$E_r^i = T_r^i \times C_f^i \quad (3)$$

where, T_r^i is the toxic response factor for a given heavy metal, C_f^i is the contamination factor. T_r^i is the Toxic response factor proposed by Hakanson, Pb, Cu, Ni, Mn, Zn, As, and Co have toxic response factors of 5, 5, 5, 1, 1, 10, and 5, respectively. The potential ecological risk (E_r^i) of heavy metals is classified into five levels, which are shown in Table 4.

Table 4. The levels of E_r^i

Degree	Pollution Level
< 40	Low
40 - 80	Moderate
80 - 160	Moderate to high
160 - 320	High
≥ 320	Very high

2.4 Statistical analysis

After obtaining the data, they were statistically analyzed

according to the randomized complete block design (RCBD). The means were compared using the least significant difference (L.S.D) test at a significance level of 0.05 using the Gestates program.

3. RESULTS AND DISCUSSION

3.1 pH

Figure 3 shows how the surroundings and location affect the pH concentration of soil samples. The statistical analysis showed significant differences, as the highest pH value was

8.3 in the second surroundings (E), while the lowest was 6.9 in the first surroundings (A). The decrease in pH is attributed to the factory being out of operation for at least five years, which reduced the deposition of alkaline calcium carbonate, one of the components of cement dust in the soil. This dust contains many components, such as calcium, aluminum, and some oxides, that can raise its value. These results align with studies (38, 39) on the subject.

The statistical analysis results showed significant differences in the effect of location, as the highest value reached 8.3 at location 8 and the lowest value reached 7.1 at location 4 (Table A1, (Appendix A)).

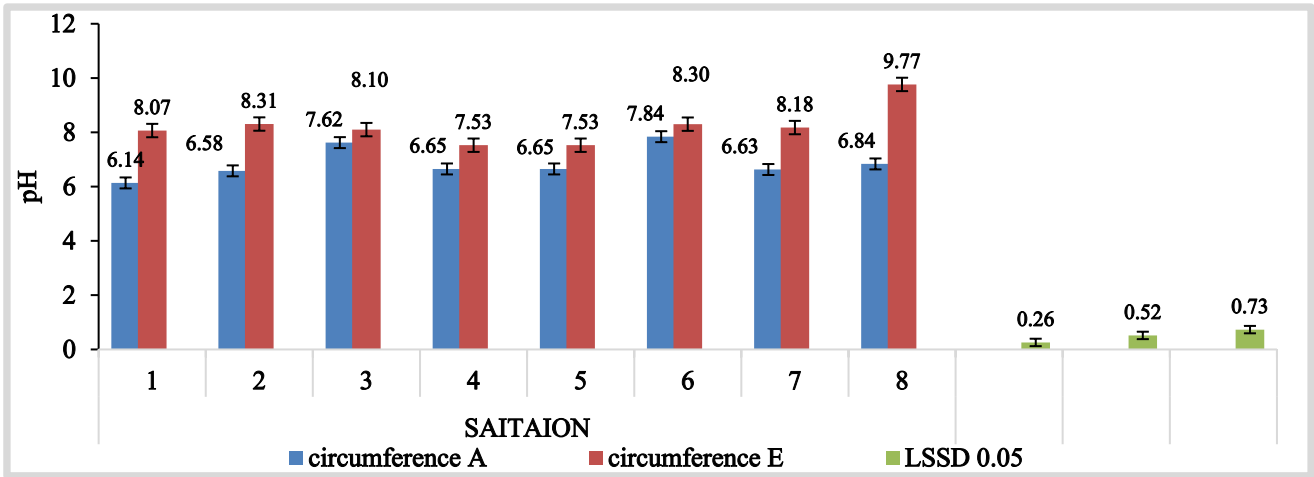


Figure 3. The effect of the surroundings and location on the pH concentration

3.2 Lithium (Li)

Statistical analysis revealed significant differences in the lithium concentration in the soil. The highest lithium concentration in surroundings E was 2.008 mg/L; in surroundings A, the lowest was 1.708 mg/L, as shown in Figure 4. The statistical analysis revealed significant differences regarding the effect of location on lithium

concentration, with the highest value being 2.375 mg/L and the lowest value being 0.560 mg/L in site 5. Since this site is directly in front of the chimney, it may be more susceptible to pollutant deposition (Table A2, (Appendix A)).

The statistical analysis results showed significant differences regarding the effect of location and surroundings on lithium concentration. The highest value was 2.250 mg/L, while the lowest was 0.455 mg/L.

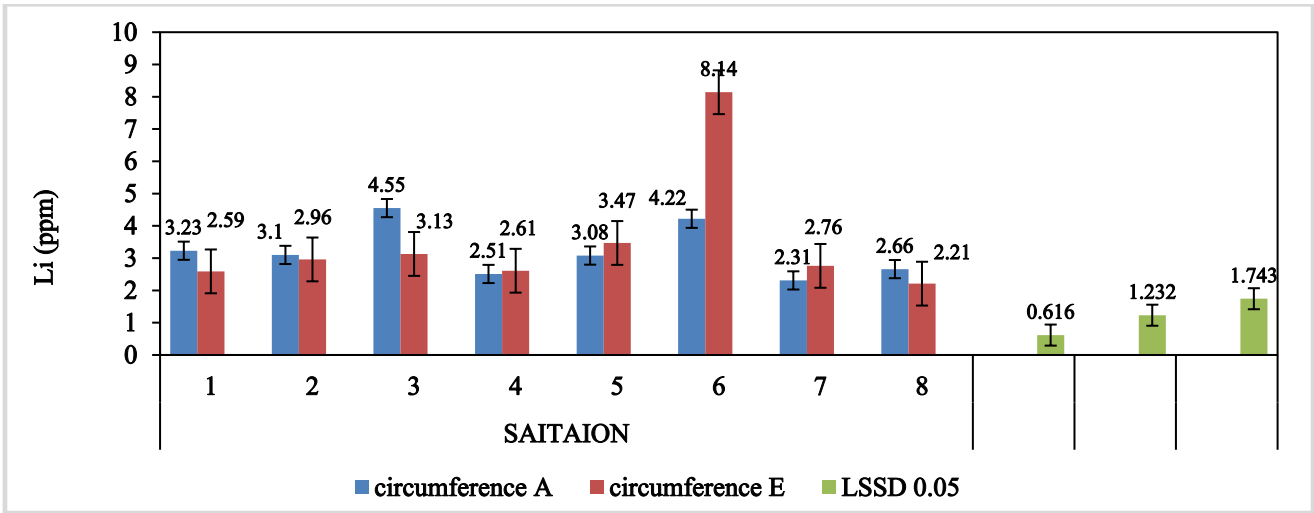


Figure 4. The effect of the surroundings and location on the lithium concentration

3.3 Lead (Pb)

Figure 5 shows the effect of dust and oxides produced by the Kubaisa Cement Plant on the lead concentration in the soil.

The statistical analysis revealed significant differences, with the highest value reaching 3.48 mg/L in the E surroundings and the lowest value reaching 3.21 mg/L in the A environment. This is likely due to fuel burning and airborne transmission to

distant areas. On the other hand, lead is less mobile. Lead in the soil is mainly associated with clay minerals, manganese oxides, iron and aluminum hydroxides, and organic matter. Adding phosphate to soil contaminated with lead results in a decrease in the soluble part of it, a significant reduction in the part associated with oxides and carbonates, and an increase in the part related to organic matter. In soils severely contaminated with lead, adding phosphate leads to the formation of lead pyromorphite metal Pbs (POCI). The solubility of lead in the soil can be significantly reduced when lime is added.

The high pH of lime-treated soil may precipitate lead in the

form of hydroxide, phosphate, or carbonate and encourage the formation of organic lead complexes characterized by their stability. Increasing the soil's acidity may improve the lead's ability to move. Still, this process is usually slower than its accumulation in layers rich in organic matter. The statistical analysis results (Table A3, (Appendix A)) showed significant differences in the site's effect on the lead concentration, as the highest value reached 6.18 mg/L at the sixth site, and the lowest value was 2.31 mg/L at the seventh site. The statistical analysis showed significant differences in lead concentration between the site and the environment, with the highest value reaching 8.14 mg/L and the lowest value being 0.455 mg/L.

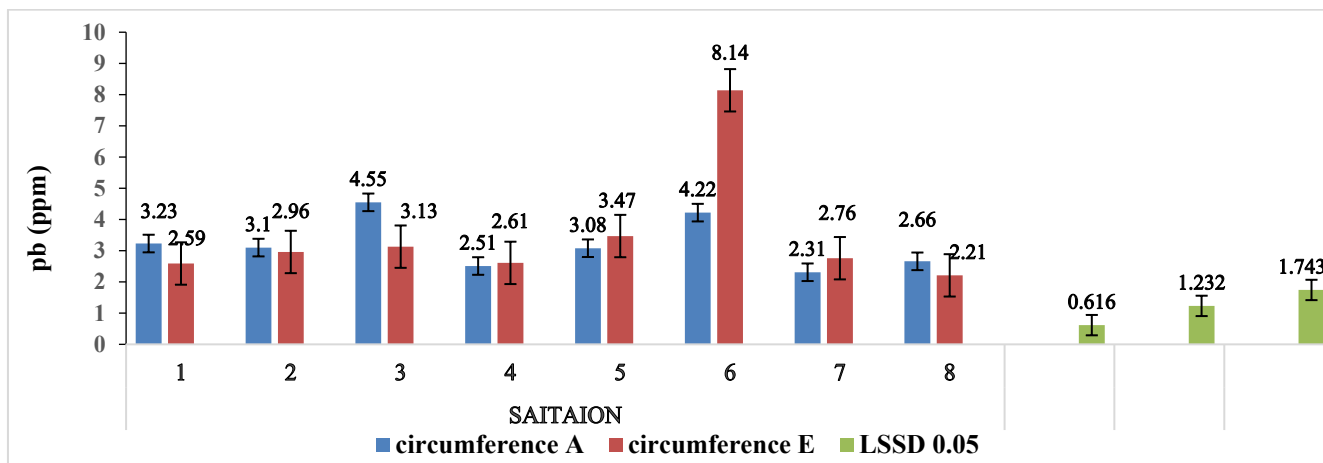


Figure 5. The effect of the surroundings and location on the lead concentration

When comparing the confidence intervals for the values of the first circumference A, which ranged between (2.21 - 3.13) mg/L, and circumference E, which ranged between (2.49 - 3.78) mg/L, with the permissible limits for lead in Table 1, it becomes clear that the confidence intervals were higher than the permissible limits, which are 0.5 ug/m³/year according to Allowed local parameters (Iraq 2012), while they reached 2 ug/m³/24 hour, 1.5 ug/m³/3 month, which reflects the adverse effect of the soil lead content.

3.4 Calcium (Ca)

Figure 6 depicts the calcium content in the soil. The statistical analysis results indicated that the surroundings had no significant effect, and no significant differences appeared. The maximum calcium concentration recorded was 34.30

mg/L in surroundings E. In contrast, the minimum was 33.84 mg/L in surroundings A. Dust contains several components, including calcium, aluminum, and some oxides, which may elevate the calcium concentration.

The statistical analysis showed significant differences in the effect of location on calcium concentration. The highest calcium concentration was 38.83 mg/L at site 8, while the lowest was 27.38 mg/L at site 5. The statistical analysis results also showed significant differences regarding the effect of location and surroundings on calcium concentration. At site 2, the highest calcium concentration was 41.15 mg/L; at site 5, the lowest was 15.90 mg/L. Table A4 (Appendix A) shows that site five is located behind the chimney, away from the influence of pollutants, while site two is situated towards the chimney and the east wind.

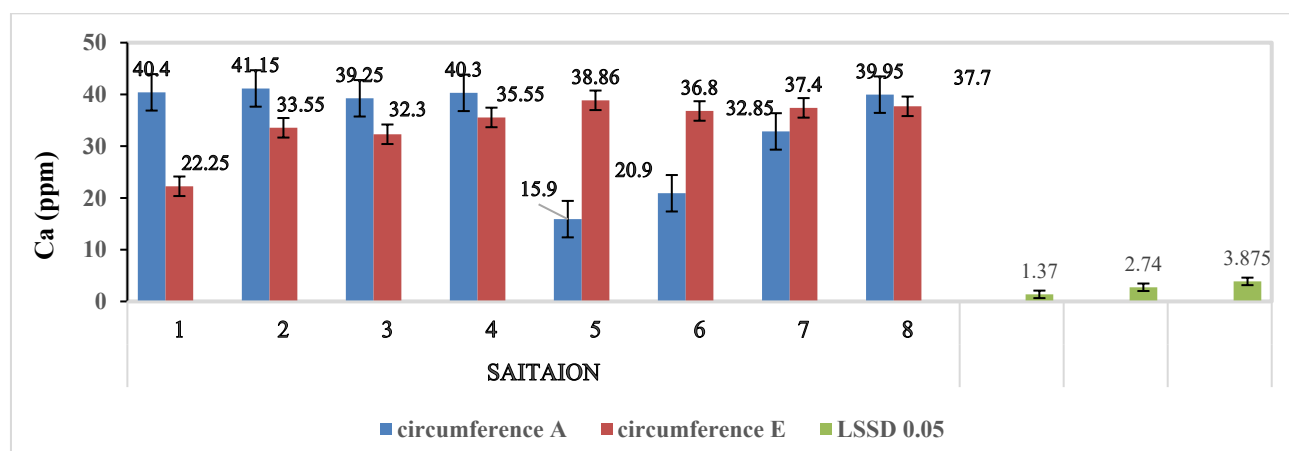


Figure 6. The effect of the surroundings and location on the calcium concentrations

3.5 Potassium (K)

Figure 7 shows the concentration of potassium in the soil. The statistical analysis results showed that the surroundings have a significant effect, as the highest value reached 0.2287 mg/L at surroundings E, and the lowest value was 0.2206 at surroundings A.

The statistical analysis results showed significant differences in the effect of location on calcium concentration, with the highest value reaching 0.2775 mg/L at site eight and the lowest value being 0.1500 at site 6. The statistical analysis results showed significant differences in the effect of the location and surroundings on calcium concentration. The highest value reached 0.3200 mg/L at site two and surroundings A, and the lowest was 0.0400 mg/L at site 6. This is because site 2 faces the chimney and the easterly wind, whereas site 5 faces the back of the chimney and is shielded from the effects of pollutants, as illustrated in Table A5 (Appendix A).

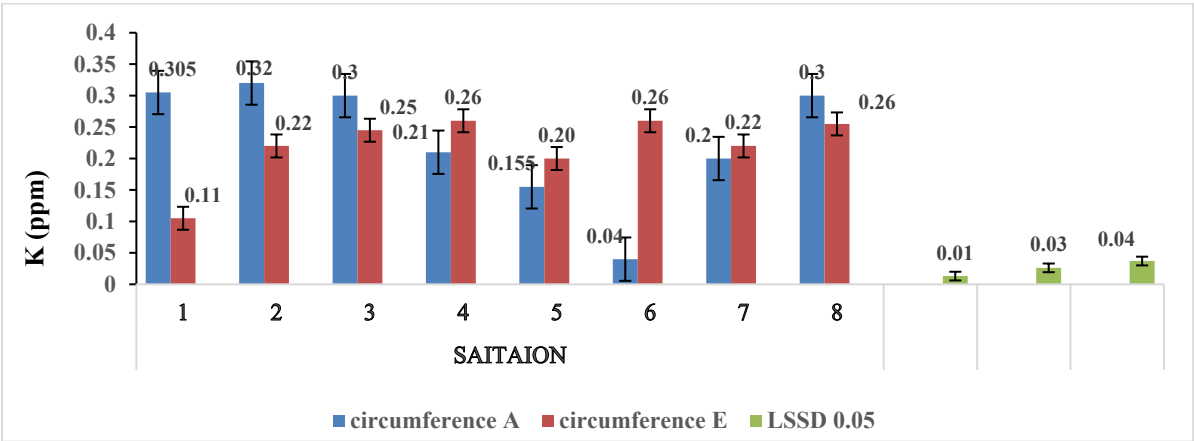


Figure 7. The effect of the surroundings and location on the potassium concentrations

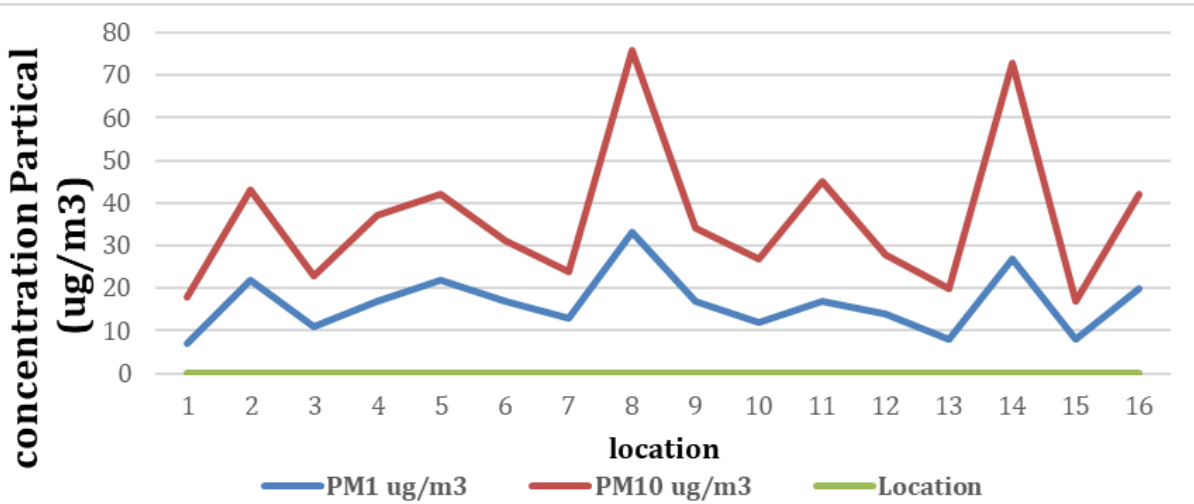


Figure 8. The effect of location and surroundings on the concentration of suspended particles of PM₁ and PM₁₀ µg/m³ sizes

3.7 CO₂ concentration

Figure 9 and Table 6 show the relationship between the surroundings and carbon dioxide gas concentration. The results showed an increase in carbon dioxide gas concentration at the first site, surroundings A, site 10, and surroundings 9.

3.6 Particle diameters

Figure 8 and Table 5 show the relationship between the number of respiratory particulates PM₁₀ (diameters more significant than 10) and particulates with diameters less than 1-micron PM₁. The percentage of fine particles smaller than 1-micron increases significantly as one move away from the plant. This is because fine aerobic particulates settle more slowly.

Table 5. The effect of location and surroundings on the concentration of suspended particles of sizes PM₁ and PM₁₀ µg/m³

E8	A8	E7	A7	E6	A6	E5	A5	E4	A4	E3	A3	E2	A2	E1	A1	Location
42	76	17	24	73	31	20	42	28	37	45	23	27	43	34	18	PM10 µg/m ³
20	33	8	13	27	17	8	22	14	17	7	11	121	22	17	7	PM1 µg/m ³

Table 6. The effect of location and surroundings on CO₂ concentration µg/m³

E8	A8	E7	A7	E6	A6	E5	A5	E4	A4	E3	A3	E2	A2	E1	A1	Location
48	51	46	46	41	41	42	44	41	44	45	47	43	50	44	45	Concentratio
0	0	0	2	5	3	1	3	8	3	7	3	0	2	7	9	n CO2 µg/m ³

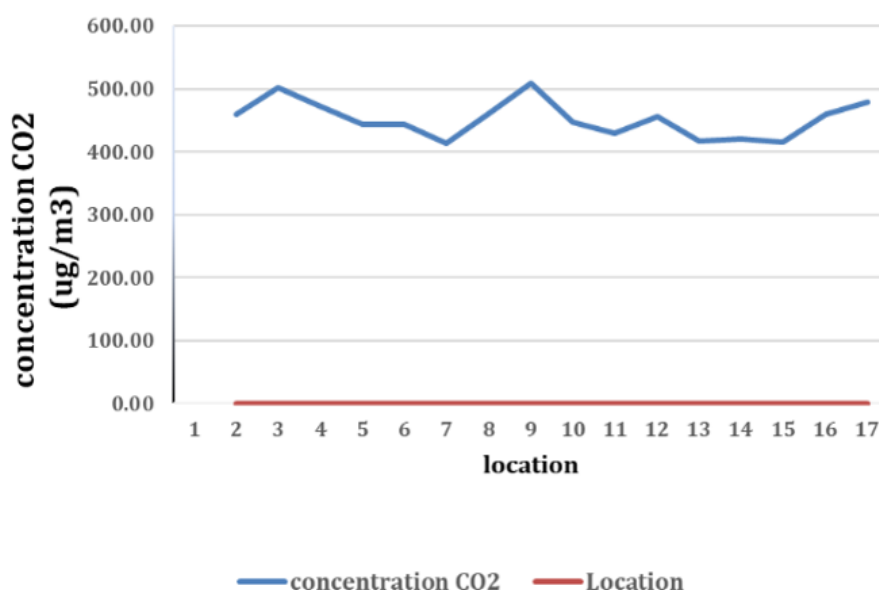


Figure 9. The effect of location and surroundings on CO₂ concentration $\mu\text{g}/\text{m}^3$

3.8 Geochemical Index (I_{geo})

Table 7 shows that the pollution load index (I_{geo}) for all sites within the two perimeters is less than zero. This could be attributed to elevated calcium concentrations, or it aligns with the findings presented in Table 6, suggesting that pollutants from human activity, such as industrial activities or vehicles, are affecting the soil. The anthropogenic source here contributed to soil pollution due to emitted gases and materials resulting from fuel combustion in the power plant, vehicle pollutants, and any other human activity that may be present in the study area. This indicates that the terrestrial source's contribution to supplying the soil with heavy elements was small. Further analysis is shown in Table A6 (Appendix A).

3.9 Contamination Factor (C_F)

Table 8 displays the values of (C_F), and the results demonstrate that the values were less than 1, signifying a low pollution level. This is attributed to the impact of humans and their activities, precisely the anthropogenic effect (Anthropogenic), which stems from fuel combustion from furnaces and power plants and pollutants from small industrial workshops and car plumbing workshops dispersed on public roads. Additionally, vehicle fuel combustion and the gases released from their exhausts in soil samples near traffic roads contain high lead levels. The increase in traffic exacerbates this pollution, leading to higher pollution levels in the air and soil. Further analysis is shown in Table A7 (Appendix A).

Table 7. The effect of location and surroundings on the Geochemical Index (I_{geo})

Surroundings	Location								Average
	1	2	3	4	5	6	7	8	
A	-3.226	-3.293	-2.816	-3.580	-3.324	2.840	3.725	-3.512	-3.290
E	-3.539	-3.373	3.278	-3.527	-3.117	1.896	3.443	-3.764	-3.242
Average	-3.383	-3.333	3.047	-3.554	-3.220	2.368	-3.584	-3.638	-3.266
		Surroundings			Location			Location & Surroundings	
LSD _{0.05}		0.2331			0.4662			0.6593	

Table 8. The effect of location and surroundings on the contamination factor

Surroundings	Location								Average
	1	2	3	4	5	6	7	8	
A	0.1614	0.1550	0.2274	0.1256	0.1539	0.2111	0.1154	0.1332	0.1604
E	0.1293	0.1479	0.1562	0.1306	0.1736	0.4070	0.1380	0.1104	0.1741
Average	0.1453	0.1515	0.1918	0.1281	0.1638	0.3091	0.1267	0.1218	0.1673
		Surroundings			Location			Location & Surroundings	
LSD _{0.05}		0.03080			0.06161			0.08713	

3.10 Potential Ecological Risk Index (E_r^i)

Table 9 shows the results of the statistical analysis of the potential environmental hazard index (E_r^i) values for heavy metals. The results demonstrate the influence of location and the surrounding environment on the potential environmental hazard values (E_r^i). The heavy element lead was classified as a low potential environmental hazard, and no data exceeded this

level. On the other hand, the results show that dust particles surrounding the cement plant generally contain significant amounts of lead and that these values are higher than the background values. The potential environmental hazard index and the geological accumulation index show that lead in sediments are the primary sources of environmental hazards. At the same time, it is not difficult to observe that the environment surrounding the cement plant has been severely

polluted due to mining activities. This soil contamination seriously threatens the safety of workers surrounding the plant, causing all kinds of health and safety problems. These results are consistent with what was reported by Liu et al. [28] when our results are compared with the potential environmental

hazard values for heavy metals in reclaimed soil by Cheng et al. [30], which are shown in Table 10. We find that the risk values for lead fall within the range of 0–27, indicating that lead represents an environmental hazard at the ecological level. Further analysis is shown in Table A8 (Appendix A).

Table 9. The effect of location and surroundings on the contamination factor

Surroundings	Location								Average
	1	2	3	4	5	6	7	8	
A	0.807	0.775	1.137	0.628	0.770	1.055	0.577	0.666	0.802
E	0.647	0.740	0.781	0.653	0.868	2.035	0.690	0.552	0.871
Location	0.727	0.758	0.959	0.640	0.819	1.545	0.727	0.609	0.836
	Surroundings			Location			Location & Surroundings		
LSD _{0.05}	0.1540			0.3080			0.4356		

Table 10. Values of the Potential Ecological Risk Index

Element	Degree	Average
Lead	0.00–27.43	6.21±3.21

4. CONCLUSIONS AND RECOMMANDATIONS

Limited measurements conducted in surroundings A, 50 meters from the laboratory, and surroundings E, 200 meters from the laboratory, showed that the pH, lead, lithium, and respiratory dust particles were well above the permissible limits. The results indicate that industrial activity surrounding the laboratory has contributed significantly to increased levels of air and soil pollution, which may negatively affect public health and the environment. Besides harmful gas emissions, factories also contributed to the buildup of heavy metals like lead in the surrounding soil, increasing the risk of soil quality degradation and its unsuitability for agricultural use. Although some sites showed high levels of pollutants, more measurements and studies are needed on a larger scale to determine the long-term impact of pollution and its relationship to industrial development. Unregulated industrial development, especially in the absence of appropriate environmental measures, contributes to environmental degradation and increases the percentage of pollutants that threaten the ecological balance in the region. The impact of human activities on pollution is still more significant than the contribution of natural sources, indicating the need to take practical steps to reduce these harmful activities.

In the light of these findings. This study recommends the following:

- 1) Installing dust precipitators in proportion to the factory's production size is necessary.
- 2) Step two is taking the necessary precautions to prevent the emission of various pollutants from factory plants to assess the current situation and the potential damage these activities could cause to the environment.
- 3) It is recommended that more similar studies be conducted on various sources of pollution, such as factories and power.
- 4) Establishing new factories before conducting a comprehensive assessment of the environmental impact that may result from them is prohibited. Educating the local community and factories about the importance of preserving the surroundings and reducing pollutant emissions through workshops and training programs.
- 5) Enhancing cooperation between government agencies and academic institutions to conduct continuous research

on air and soil pollution and the impact of human activities on them.

- 6) Adopting clean and sustainable energy technologies in factories and power plants as an alternative to traditional sources to reduce pollution.
- 7) Motivating companies to use clean manufacturing technologies and techniques while reducing their reliance on fossil fuels.
- 8) Develop emergency plans to monitor and deal with pollution in pollutant leakage or environmental accidents.

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APPENDIX

Table A1. Potential of Hydrogen (PH)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	0.03	0.0032	0.0032	1	rep stratum
<.001	136.89	16.0744	16.0744	1	steram
<.001	7.23	0.8486	5.9403	7	Sation
0.002	5.78	0.6788	4.7517	7	steram.sation
		0.1174	1.7614	15	Residual
			28.5310	31	Total

Table A2. Lithium (Li)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	15.82	0.23978	0.23978	1	rep stratum
<.001	47.70	0.72300	0.72300	1	steram
<.001	96.93	1.46934	10.28537	7	Sation
<.001	42.20	0.63972	4.47802	7	steram.sation
		0.01516	0.22737	15	Residual
			15.95355	31	Total

Table A3. Lead (Pb)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	3.12	2.0859	2.0859	1	rep stratum
0.356	0.91	0.6072	0.6072	1	steram
<.001	9.13	6.1031	42.7218	7	Sation
0.014	3.80	2.5415	17.7903	7	steram.sation
		0.6684	10.0257	15	Residual
			73.2309	31	Total

Table A4. Calcium (Ca)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	6.70	22.145	22.145	1	rep stratum
0.482	0.52	1.721	1.721	1	steram
<.001	22.79	75.308	527.156	7	Sation
<.001	54.56	180.295	1262.064	7	steram.sation
		3.304	49.566	15	Residual
			1862.650	31	Total

Table A5. Potassium (K)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	1.75	0.0005281	0.0005281	1	rep stratum
0.205	1.75	0.0005281	0.0005281	1	steram
<.001	29.65	0.0089388	0.0625719	7	Sation
<.001	51.11	0.0154067	0.1078469	7	steram.sation
		0.0003015	0.0045219	15	Residual
			0.1759969	31	Total

Table A6. Geochemical index (I_{geo})

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	2.59	0.24826	0.24826	1	rep stratum
0.671	0.19	0.01793	0.01793	1	steram
<.001	7.15	0.68415	4.78908	7	Sation
0.114	2.06	0.19700	1.37899	7	steram.sation
		0.09568	1.43526	15	Residual
			7.86953	31	Total

Table A7. Contamination factor (C_F)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	3.12	0.005215	0.005215	1	rep stratum
0.356	0.91	0.001518	0.001518	1	stream
<.001	9.13	0.015258	0.106805	7	Sation
0.014	3.80	0.006354	0.044476	7	Stream station
		0.001671	0.025064	15	Residual
			0.183077	31	Total

Table A8. Potential ecological risk (E_r^i)

P-value	F-statistic	Sum of Squares	Mean Square	Degrees of Freedom	Source of Variation
	3.12	0.13037	0.13037	1	rep stratum
0.356	0.91	0.03795	0.03795	1	stream
<.001	9.13	0.38145	2.67012	7	Sation
0.014	3.80	0.15884	1.11190	7	Stream station
		0.04177	0.62660	15	Residual
			4.57693	31	Total