Influence of Highway Traffic on Contamination of Roadside Soil with Heavy Metals

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Abstract

This study is one of the first works which examined the assessment of heavy metal contamination of pavement-side soils in Algeria. It deals with the section of National Highway 3 (RN3), which crosses the wilaya of Batna. In the environment of sampling sites there is no industry or dangerous activity on the environment, the heavy metals addressed in this study are (Pb, Cu, Cr, Fe, Ni, Zn), their origin being road traffic. The objectives of this study were to: (1) Determine the concentrations of heavy metals in road dust; (2) Identify the sources of different heavy metals in soils and road dust; (3) Exploring the extent of heavy metal pollution in neighbouring soils. To this end, 33 samples were collected, including 03 road dust and 30 soil samples over different distances from 1m to 80m. The samples were analyzed by FRX. Results indicated that concentrations in road dust were higher than in soil. The distribution of heavy metal concentrations in dust is Fe>Pb>Zn>Cu>Cr>Ni, and the distribution in the ground is Fe>Pb>Cu>Zn>Cr>Ni in the direction of Biskra and in the opposite direction and decreases away from the road, while the distribution in the central solid ground is Fe>Cu>Cr>Pb>Zn>Ni. Climatic conditions such as wind, rainfall, temperature, humidity and the nature of the terrain were also significantly related to their enrichment in these roadside soils. The enrichment factor (EF) and the geo-accumulation index (Igeo) were calculated, as well as all elements with a (EF) that ranges from moderate to high to extremely contaminated, reflecting the high anthropogenic load of these metals in the study area and the results of the Igeo accumulation indices confirm the results obtained for the enrichment factor (EF).

Keywords: Pollution; Contamination; Heavy Metals; Traffic; Road; Rainfall; Water; Runoff; Soil; Dust; Environment; Wind; Pavement.

1. Introduction

Heavy metal soil pollution in rural areas near major highways is a major issue in roads with varying loads of traffic in rural areas. Traffic in major Algerian roads in rural areas is one of the sources of roadside soil pollution, especially with regard to heavy metal concentrations and their impact on the immediate environment of the road. For this purpose, it is necessary to identify, assess and quantify the concentrations of heavy metals in road dust and in the adjacent soil arable layer. Traffic and industrial emissions, are considered important factors in metal pollution in road...
dust [1-3]. With the phasing out of leaded gasoline in developed countries and the implementation of other control measures, traffic-related metal emissions had significantly decreased[4, 5]. Whereas traffic-related emissions dominate metal pollution in road dust [6, 7]. This can contribute to the enrichment of the Cr, Cu, Fe, Ni, Pb and Zn elements [8, 9]. Nevertheless, the chemical, resulting from the mixture of heavy metals linked to road traffic, varies considerably from study to study [9, 10]; probably due to the difference in metal concentration in brake pads and tire materials, as well as the difference in road traffic density [1, 8]. Several studies have indicated that the contamination of soil and several types of roadside vegetation species by significant concentrations of heavy metals is due to the volume of road traffic [11, 12].

In addition, the results could be used appropriately to monitor the aerial deposition of these roadside metals. Pagotto et al. (2001) [13] assessed the impact of certain physical infrastructure parameters (safety slides, pavement type, pavement slope) on the extent of heavy metal contamination, as well as the composition of the material near the emission source (road dust), according to Fakayode and Olu-Owolabi (2003) [14]; roads are known as the second most common source of urban pollution. Other studies have focused on the contamination of dust from roads and roadside soils in urban areas by heavy metals [15, 16]. They showed that levels of heavy metal concentration such as Pb, Cd, Cu, Cr, Ni, Zn, Fe and Co far exceeded the limits set for clean soils and their source came mainly from industrial derivatives, combined with traffic sources. Studies have examined the influence of industrial activities and waste incineration on the accumulation and extent of heavy metal pollution in the complex system of surface soils and road dust [17, 18]. They showed that the concentrations of these metals in road dust were generally higher than those in soils. Various works have examined the spatial distribution of heavy metals in dust and surface soils next to roads and their origins [19, 20]. They showed that the increase in anthropogenic trace metal elements in the surface environment can most likely be attributed to urbanization, rapid industrialization and increased vehicle emissions into the atmosphere. They revealed that the distribution of heavy metals in soil samples is affected by wind direction.

They concluded that concentrations of these metals in soils are higher on the surface but decrease in low-lying areas and that the highest level of metals was found in eastern parts of roads due to prevailing wind. Also, levels of contamination of these metals decrease as one moves away from the side of the road. Authors have revealed that automobile traffic is the main source of pollution from road dust and adjacent soils with a contribution of more than 50%. In addition, unregulated incineration and hazardous waste dumps along the road were responsible for these contaminations [21, 22]. Heavy metal contamination on human health was the concern of other researchers such as Wu et al. and Al-Shidi et al. (2020) [23, 24]. They found that the health risks associated with heavy metal accumulation were higher in densely populated areas, high-traffic areas and industrial areas. They also showed that tire wear and exhausting diesel engines have a higher potential threat to human health as they generate high amounts of high-grade heavy metals, and revealed that gasoline and diesel emissions contributed significantly to the presence of Cr and Ni during braking, and tire wear generated large quantities of Cu and Zn [23]. In order to assess the risks of environmental contamination by heavy metals, they concluded that the environmental risks specific to each source and the critical sources of heavy metals have changed with the seasons changes, suggesting that different strategies should be adopted according to the seasons. The aims of this study were to: (1) Determine the concentrations of heavy metals in road dust; (2) Identify the critical sources of different heavy metals in soils and road dust; (3) Exploring the extent of heavy metal pollution in neighboring soils.

2. Materials and Methods

The flowchart of the research and characterisation methodology is presented in Figure 1.

2.1. Study Area

The study area is an agricultural area far from all industrial activities, located in the commune of Oued-Echaaba wilaya of Batna in north-east Algeria, on a section of the national road three called (RN03), linking the Algerian north to the extreme south of Algeria, from the wilaya of Skikda to the wilaya of Tamanrasset with a traffic of 29521 vehicles per day. The section in question is a duplication of 7m width on either side, the stretch of road is built in embankment with full central land of 5.5m width. The type of pavement is a classic pavement, the remediation of the road is provided by earth ditches in triangular shape on both sides of the roadway, with a depth of 1.5m, a width 3m to the mirror. The city has a semi-arid climate with hot summers, cold winters and annual rainfall of around 346 mm, humidity is 53% and the maximum wind speed is 4.3m/s.

2.2. Samples and Sampling

The location of the sampling area in map mode is shown in Figure 2, soil sampling is done using a stainless transplant (cleaned after each sample with distilled water) at the level of 36 points arranged on either side of the roadway and 3 points in the TPC see Figure 3. These samples resulted in the preparation of 15 composite and homogeneous soil samples.
All samples were placed in airtight polyethylene bags, labelled and brought back to the laboratory. Prior to the analysis, all samples were put to the oven at 105°C for 24 hours for drying, crushed with porcelain mortar, then sifted to 2mm and finally ground in a type disc grinder (DM200). Road dust samples were collected about three weeks after a thunderstorm, and the weather was sunny during sampling, clean plastic brushes and brushes were used to collect road dust samples [32, 33]; six points were taken on the pavement's bearing layer on an area of 7 m² on both sides of
the roadway. A sampling was taken at the roundabout, during the preparation, small pebbles, cigarette butts, hair and other plant debris were removed. The samples were carefully mixed to obtain compound samples. The samples were placed in airtight self-sealing polyethylene bags labelled and brought back to the laboratory for drying at 105°C, and then shifted into a sieve with a mesh of 0.5mm. Soil and road dust samples were kept at 4°C until they were analyzed. Figure 4 presented the various steps of sample preparation.

2.3. Granulometric Analysis

The granulometric analysis of the samples was carried out using a laser granulometer presented in Figure 5; (Partica LA-960 HORIBA Laser Scattering Particle Size Distribution Analyzer).

The results presented in Figure 6 shown a rather coarse nature for soil samples than for road dust.

2.4. Preparing Pellets

The pellets are obtained by compressing the powder using a manual press. A boric acid binder can be added to give the pellet a good mechanical solidity. The samples are then ready to be analyzed, the pellet with a circular shape of 25mm in diameter, 5mm thick, and a weight equal to 8g of powder. Samples are analyzed by a wavelength dispersion
x fluorescence spectrometer (WDXRF) (ZSX Primus IV regaku fluorescence spectrometer). The Figure 7 shown the preparation of the pellet for analysis.

Figure 7. Preparation of the pellet for analysis: (a) The manual press; (b) The pellets; (c) The WDXRF

3. Estimated Intensity of Contamination

The intensity of soils contamination by heavy metals was assessed from two indices; the enrichment factor (EF) and the geo-accumulation index (Igeo), their principle is based in the comparison of measured values against reference values.

3.1. Enrichment Factor

The Enrichment factor, is a parameter that, allows to see whether the concentration of heavy metals obtained in soils are anthropogenic or natural in origin [26]; if EF close to 1 indicates a natural source while EF > 10 suggests an anthropogenic source. According to Yuen et al. [3] the Enrichment factor (EF) is defined by:

$$
EF = \frac{(C_i/Fe)_{Sample}}{(C_i/Fe)_{Background}}
$$

where: \((C_i/Fe)_{Sample}\) and \((C_i/Fe)_{Background}\) are the ratios of metal concentration (i), standardizer (Fe) in sample, metal concentration (i) and standardizer (Fe) in the background material, respectively. In this research, the (Fe) has been used as a standardizer because it has low variability in occurrence and is a main component of the Earth's crust [27, 28]; and soils 80m from the roadway were used as reference materials [3, 13]. The level of metal pollution can be categorized into five categories Sutherland, R. A.

- EF < 2  Deficiency to minimal enrichment;
- 2 < EF < 5  Moderate enrichment;
- 5 < EF < 20  Significant enrichment;
- 20 < EF < 40  Very high enrichment;
- EF > 40  Extremely high enrichment;

3.2. Geo-accumulation Index (Igeo)

To quantify the degree of metal contamination of dust and road side soils [28, 30]; the geo-accumulation index (Igeo) was calculated on the basis of [1]:

$$
Igeo = \log_2 \frac{C_i}{B_{1.5}}
$$

Where: Igeo: Geo-accumulation index; \log_2: basic logarithm 2; i: item considered; C: concentration measured in the sample; B: geochemical background; 1.5: is a factor used because of possible variations in background data due to lithological variations [28, 31].

In addition, Gasser and Müller (1979) [32] defined a scale of values with six classes based on the intensity of pollution. As shown below:

- Igeo < Uncontaminated.
- 0 < Igeo < 1 uncontaminated to moderately contaminated.
- 1 < Igeo < 2 moderately contaminated.
2 < Igeo < 3 moderately to strongly contaminated.
3 < Igeo < 4 strongly contaminated.
4 < Igeo < 5 strongly to extremely contaminated.
Igeo > 5 extremely contaminated.

3.3. Statistical Analysis

The results were analyzed statistically using SPSS ver 20 software to determine the different correlations that exist between the metals (Fe, Zn, Ni, Cu, Cr, Pb) studied. Pearson correlation matrixes were established for a confidence interval of 95 and the level of significance was set at P< 0.05.

4. Results and Discussions

In this study, the results show that heavy metals concentrations in road dust are higher than those found in soils, and that heavy metals in soils decrease in soils by moving away from the road axis. Tables 1, 2 and 3. The distribution of heavy metals in road dust is in the order of Pb > Zn > Cu. > Cr > Ni. In the soils (Batna to Biskra and Biskra to Batna direction) is the order of Pb > Cu > Zn > Cr > Ni. In the central solid earth, the distribution is in the order of Cu > Cr > Pb > Zn > Ni. The high levels of lead in the dust are due to traffic-related exhaust and the overuse of leaded gasoline in third world countries, notably Algeria. In addition, the concentrations of (Cu, Zn, Cr, Ni) are due to the use of spare parts, braking system pads (discs and brake pads) and tire wear whose counter-manner is very marked in the markets. For iron, there is a high surface and roadside content. Presumably there is a road-related contribution that is more or less masked by the high concentrations of this element in natural soils. It has also been observed that the transport of dust through the various channels (hydraulic, wind, dry fallout) affects directly the quality and degree of soil pollution, which is confirmed by the differences in the concentrations of heavy metals in the two directions of the road as shown in Figures 8 to 10.

| Elements | Cr   | Fe   | Zn   | Ni   | Pb   | Cu   |
|----------|------|------|------|------|------|------|
| TPC      | 197.74 | 38958.42 | 110.87 | 99.81 | 171.74 | 197.96 |
| 1 m      | 145.74 | 37699.44 | 210.49 | 90.38 | 262.72 | 234.23 |
| 5 m      | 84.74  | 46722.13 | 122.92 | 82.52 | 208.88 | 77.37  |
| 10 m     | 97.11  | 54555.78 | 51.53  | 12.95 | 41.78  | 28.71  |
| 20 m     | 92.99  | 53926.29 | N.D    | 17.02 | 91.91  | 28.71  |
| 40 m     | 84.74  | 46722.13 | 122.92 | 82.52 | 208.88 | 77.37  |
| 80 m     | 71.74  | 52597.37 | 43.04  | 8.59  | 9.75   | 2.09   |
| V max    | 197.74 | 53926.29 | 210.49 | 99.81 | 262.72 | 234.23 |
| V min    | 71.74  | 37699.44 | 43.03  | 8.59  | 9.75   | 2.09   |
| V med    | 112.75 | 47611.40 | 97.69  | 116.56 | 79.84  |
| Gap-type | 43.96  | 6932.24  | 69.97  | 41.43 | 98.47  | 96.99  |

| Elements | Cr   | Fe   | Zn   | Ni   | Pb   | Cu   |
|----------|------|------|------|------|------|------|
| TPC      | 154.48 | 41896.04 | 164.7 | 99.81 | 253.44 | 237.39 |
| 1 m      | 93.16  | 51758.05 | 118.1 | 98.23 | 248.79 | 78.32  |
| 5 m      | 92.99  | 54206.06 | 66.86 | 96.24 | 160.6  | 31.51  |
| 10 m     | 91.58  | 52177.72 | 59.63 | 19.16 | 90.05  | 18.48  |
| 20 m     | 92.99  | 52107.76 | 40.23 | 62.41 | 72.41  | 9.09   |
| 40 m     | 86.99  | 54415.89 | 48.07 | N.D  | 11.42  | 4.84   |
| 80 m     | 103.66 | 51093.58 | 97.69 | 116.56 | 79.84  |
| V max    | 154.48 | 54415.89 | 164.7 | 99.81 | 253.44 | 237.39 |
| V min    | 86.99  | 41896.04 | 48.07 | 16.23 | 11.42  | 4.84   |
| V med    | 32.91  | 4646.84  | 48.51 | 42.67 | 98.71  | 89.340 |
| Gap-type | 43.96  | 6932.24  | 69.97 | 41.43 | 98.47  | 96.99  |
Table 3. Concentration of heavy metals in road dust (D, G, A)

| Elements | Cr     | Fe    | Zn    | Ni    | Pb    | Cu    |
|----------|--------|-------|-------|-------|-------|-------|
| dust D   | 215.53 | 34062.39 | 316.54 | 103.74 | 368.65 | 276.82 |
| Dust G   | 214.16 | 26928.17 | 156.67 | 101.38 | 380.62 | 269.72 |
| dust R   | 212.11 | 25319.48 | 261.91 | 99.81  | 309.13 | 272.88 |
| V max    | 215.53 | 34062.39 | 316.54 | 103.74 | 380.62 | 276.82 |
| V min    | 212.11 | 25319.48 | 156.67 | 99.81  | 309.13 | 269.72 |
| V med    | 213.93 | 28770.01 | 245.04 | 101.64 | 352.8 | 273.14 |
| Gap- type| 1.72   | 4653.38 | 81.26  | 1.98   | 38.28  | 3.55  |

Note: D: in the direction to Biskra; G: in the direction to Batna; R: in the roundabout.

4.1. Enrichment Factor (EF)

These results suggest that for road dust: Cu is extremely enriched; Pb has very high enrichment and Zn, Ni and Cr have moderate to significant enrichment, while for soils (in both directions Batna-Biskra, Biskra-Batna and in the TPC) (Tables 4 to 6): the Pb has a very high enrichment for distances (1m; 5m) and significant to moderate for distances (10m, 20m, 40m) and minimum at 80m. For the Enrichment Cu is extremely high in (TPC, 1m, 5m) and significant to moderate to (10m, 20m, 40m) and minimal at 80m; for the neither enrichment and significant in (TPC, 1m, 5m, 10m) and moderate in (20m, 40m) and minimal at 80m (Figures 11 to 13). All elements have an FE that ranges from (moderate to strong) to (extremely contaminated), reflecting the high anthropogenic load of these metals in the study area. However, differences can be attributed to the different approaches used in methods of calculating enrichment factors.
Table 4. Enrichment Factor Values (EF) in the soil (direction to Biskra)

| Distance | EF (Pb) | EF (Cr) | EF (Zn) | EF (Ni) | EF (Cu) | EF (Fe) |
|----------|---------|---------|---------|---------|---------|---------|
| TPC      | 17.62   | 2.76    | 2.57    | 11.62   | 94.91   | 0.74    |
| 1 m      | 26.95   | 2.03    | 4.89    | 10.53   | 112.30  | 0.72    |
| 5 m      | 21.43   | 1.19    | 2.86    | 9.61    | 37.09   | 0.89    |
| 10 m     | 9.43    | 1.29    | N.D     | 1.98    | 13.76   | 1.02    |
| 20 m     | 4.3     | 1.35    | 1.19    | 1.51    | 5.83    | 1.04    |
| 40 m     | 2.99    | 1.38    | 1.1     | 2.11    | 3.04    | 0.93    |
| 80 m     | 1       | 1       | 1       | 1       | 1       | 1       |

Note: N.D: not defined

Table 5. Enrichment Factor Values (EF) in the soil (direction to Batna)

| Distance | EF (Pb) | EF (Cr) | EF (Zn) | EF (Ni) | EF (Cu) | EF (Fe) |
|----------|---------|---------|---------|---------|---------|---------|
| 1 m      | 22.2    | 2.93    | 3.43    | 14.23   | 49.11   | 0.77    |
| 5 m      | 21.79   | 1.77    | 2.46    | 14.00   | 16.20   | 0.95    |
| 10 m     | 14.07   | 1.75    | 1.39    | 9.87    | 6.52    | 0.99    |
| 20 m     | 7.9     | 1.74    | 1.24    | 2.73    | 3.82    | 0.96    |
| 40 m     | 6.34    | 1.65    | 0.83    | 2.31    | 1.88    | 0.96    |
| 80 m     | 1       | 1       | 1       | 1       | 1       | 1       |

Table 6. Enrichment Factor Values (EF) in Road Dust

| Elements  | EF (Pb) | EF (Cr) | EF (Zn) | EF (Ni) | EF (Cu) | EF (Fe) |
|-----------|---------|---------|---------|---------|---------|---------|
| Dust D    | 37.82   | 4.09    | 7.36    | 14.79   | 132.72  | 0.62    |
| Dust G    | 39.05   | 4.06    | 3.64    | 14.45   | 129.31  | 0.49    |
| Dust R    | 31.71   | 4.02    | 6.08    | 14.23   | 130.83  | 0.46    |

Figure 11. Enrichment factors (EF) in the soil (Direction to Biskra)

Figure 12. Enrichment factors (EF) in the soil (Direction to Batna)

Figure 13. Enrichment factors (EF) in road dust
The values of the calculated geo-accumulation indices (Igeo) are presented in Tables 7 to 9. The results of the index (Igeo) in road dust show different levels of contamination depending on the element:

For Pb and Cu, Igeo ranges from (high to extremely contaminated) to (extremely contaminated); for Cr and Zn, Igeo varies between (moderately contaminated to highly contaminated), and for Ni Igéo is highly contaminated. The results of the index (Igeo) in soils (Batna-Biskra direction, Biskra-Batna, TPC) indicate different levels of contamination, depending on the element and distance: For Pb and Cu, Igeo in (TPC, 1m, 5m) varies between (highly contaminated) to (highly contaminated) and for distances of 10m, 20m, 40m to (moderately to highly contaminated) and at 80m the Igeo varies between (untaminated) to (not to moderately contaminated). For Cr and Zn, Igéo varies from (untaminated) to (not moderately contaminated) to (moderately contaminated) at all points. For the Ni, the Igeo for (TPC, 1m, 5m, 10m) varies between (moderately to highly contaminated) to (highly contaminated) and for distances (20m, 40m, 80m) varies from (untaminated) to (not to moderately contaminated) (Figures 14 to 16).

Table 7. Values of geo-accumulation indices (Igeo) in the soil (Direction to Biskra)

| Distance | Igéo (Pb) | Igéo (Cr) | Igéo (Zn) | Igéo (Ni) | Igéo (Cu) | Igéo (Fe) |
|----------|-----------|-----------|-----------|-----------|-----------|-----------|
| TPC      | 3.55      | 0.88      | 0.78      | 2.95      | 5.98      | -1.02     |
| 1 m      | 4.17      | 0.44      | 1.70      | 2.81      | 6.22      | -1.06     |
| 5 m      | 3.83      | -0.34     | 0.93      | 2.68      | 4.63      | -0.76     |
| 10 m     | 2.65      | -0.21     | N.D       | 0.40      | 3.2       | -0.55     |
| 20 m     | 1.51      | -0.15     | -0.32     | 0.01      | 1.96      | -0.53     |
| 40 m     | 0.99      | -0.11     | -0.45     | 0.49      | 1.02      | -0.69     |
| 80 m     | -0.58     | -0.58     | -0.58     | -0.58     | -0.58     | -0.58     |

Table 8. Values of geo-accumulation indices (Igeo) in the soil (Direction to Batna)

| Distance | Igéo (Pb) | Igéo (Cr) | Igéo (Zn) | Igéo (Ni) | Igéo (Cu) | Igéo (Fe) |
|----------|-----------|-----------|-----------|-----------|-----------|-----------|
| 1 m      | 3.89      | 0.97      | 1.19      | 3.24      | 5.03      | -0.96     |
| 5 m      | 3.86      | 0.24      | 0.71      | 3.22      | 3.43      | -0.65     |
| 10 m     | 3.23      | 0.22      | -0.11     | 2.72      | 2.12      | -0.59     |
| 20 m     | 2.39      | 0.21      | -0.27     | 0.86      | 1.35      | -0.64     |
| 40 m     | 2.08      | 0.14      | -0.84     | 0.62      | 0.32      | -0.64     |
| 80 m     | -0.58     | -0.58     | -0.58     | -0.58     | -0.58     | -0.58     |

Table 9. Values of geo-accumulation indices (Igeo) in road dust

| Elements | Igéo (Pb) | Igéo (Cr) | Igéo (Zn) | Igéo (Ni) | Igéo (Cu) | Igéo (Fe) |
|----------|-----------|-----------|-----------|-----------|-----------|-----------|
| Dust D   | 4.65      | 1.45      | 2.29      | 3.30      | 6.47      | -1.26     |
| Dust G   | 4.70      | 1.44      | 1.28      | 3.27      | 6.43      | -1.6      |
| Dust R   | 4.40      | 1.42      | 2.02      | 3.24      | 6.44      | -1.69     |
4.3. Correlations between Heavy Metals

Tables 10 and 11 show Pearson’s correlation matrixes between the heavy metals studied in soil samples. Analysis of these matrixes shows very significant and positive correlations (P < 0.05) between Ni/Pb, Pb/Ni and Ni/Cu, Cu/Ni; and negative correlations between Fe/Ni, Ni/Fe and Fe/Cu, Cu/Fe, and significant and positive correlations between Zn/Ni, Ni/Zn, Zn/Pb, Pb/Zn, Zn/Cu, Cu/Zn Cu/Pb, Pb/Cu, Cu/Cr/Cu, and negative between Cr/Fe, Fe/Cr, Fe/Zn, Zn/Fe Pb/Fe, in the ground (Batna to Biskra) and in the opposite direction (Biskra to Batna) very significant correlations were observed between Zn/Cu, Cu/Zn, Ni/Pb, Pb/Ni and negative between Cr/Fe, Fe/Cr Fe/Cu, Cu/Fe and positive significant correlations between Cu/Zn, Zn/Cu, Cr/Cu, Cu/Cr, Zn/Ni, Ni/Zn, Zn/Pb, Pb/Zn and negative between Zn/Fe and Fe/Zn. The high correlations noted suggest that these parameters are governed by the same mechanism. On the other hand, the positive and significant correlations between different heavy metals reflect a common source, while the significant and negative correlations between the heavy metals studied indicate different sources.

| Pearson Correlation | Cr     | Fe     | Zn     | Ni     | Pb     | Cu     |
|---------------------|--------|--------|--------|--------|--------|--------|
| Cr      | 1      | -0.816∗ | 0.526  | 0.733  | 0.544  | 0.839∗ |
| Fe      | -0.816∗ | 1      | -0.868∗ | -0.912∗ | -0.811∗ | -0.947∗ |
| Zn      | 0.526  | -0.868∗ | 1      | 0.836∗ | 0.862∗ | 0.859∗ |
| Ni      | 0.733  | -0.912∗ | 0.836∗ | 1      | 0.917∗ | 0.904∗ |
| Pb      | 0.544  | -0.811∗ | 0.862∗ | 0.917∗ | 1      | 0.871∗ |
| Cu      | 0.839∗ | -0.947∗ | 0.859∗ | 0.904∗ | 0.871∗ | 1      |

| Pearson Correlation | Cr     | Fe     | Zn     | Ni     | Pb     | Cu     |
|---------------------|--------|--------|--------|--------|--------|--------|
| Cr      | 1      | -0.928∗ | 0.845∗ | 0.714  | 0.778  | 0.917∗ |
| Fe      | -0.928∗ | 1      | -0.855∗ | -0.582 | -0.635 | -0.958∗ |
| Zn      | 0.845∗ | -0.855∗ | 1      | 0.881∗ | 0.890∗ | 0.954∗ |
| Ni      | 0.714  | -0.582 | 0.881∗ | 1      | 0.981∗ | 0.761  |
| Pb      | 0.778  | -0.635 | 0.890∗ | 0.981∗ | 1      | 0.774  |
| Cu      | 0.917∗ | -0.958∗ | 0.954∗ | 0.761  | 0.774  | 1      |

5. Conclusion

The objective of this study is to assess the level of heavy metal contamination of roadside soils exposed to road traffic on a major road (national road three (RN03) located in the commune of Oued-Echaab, wilaya of Batna, Algeria. The results obtained show that all the heavy metals detected, except Fe, (Zn, Ni, Cu, Cr, Pb) have a close link with road traffic. All of these elements have some higher levels than the reference values. The general sequence of metal element levels in these soils is: Fe > Zn > Cr > Cu > Ni > Pb. High concentrations of heavy metals were found in soils near the road at distances of 1 to 10 m depending on the metal and in central solid earth (TPC). These concentrations decrease exponentially as they move away from the highway. Although all elements have achieved moderate to extremely enrichment for road dust, and minimal enrichment in soils at 80m distance, to extremely high in.
soils (TPC, 1m, 5m), the results of FE are confirmed by the results of Igeo whose contamination is varied between uncontaminated at a distance of 80m, to highly contaminated in (TPC, 1m, 5m). Contamination of the soil by Cu and Pb persists up to 20m. The results of this study are of great importance for the quantitative identification of the sources of contamination. They will be used to raise awareness of the pollution of vehicles by heavy metals. The contamination of these soils is, on the one hand, a risk of poisoning throughout the food chain for the populations who use them for their crops and, on the other hand, a risk of groundwater contamination. Thus, an easement must be taken into account in order to avoid such contamination.

5.1. Future Aspects of the Research

This work will constitute an inventory and a starting point. Further research such as pollution of plants, deep soils should be carried out to reduce the health risks posed by heavy metals in road dust, plants and road sewage water.

6. Declarations

6.1. Author Contributions

Conceptualization, B.A. and B.N.; writing—original draft preparation, B.A. and B.N.; writing—review and editing, B.A. and B.N.; Sample taking and preparation sample testing by B.A, M.T and MA; The manuscript was written by all authors; results, discussions and conclusion section was completed by B.A and B.N; All authors have read and agreed to the published version of the manuscript.

6.2. Data Availability Statement

The data presented in this study are available in article.

6.3. Funding

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6.5. Conflicts of Interest

The authors declare no conflict of interest.

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