

Trace Elements Distribution in Red Soils under Semiarid Mediterranean Environment

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Abstract

This study states the potential trace elements (TE's) content of red soils located at the centre region of Spain, characterized by low rainfall and slight acidity over prolonged weathering periods. For this purpose, three soil profiles from a catena were described, sampled and analyzed. The most notable characteristics are the low organic matter content and the predominantly acidic pH. Illite and kaolinite are the predominant clay minerals. The fertility of the soils is sufficient to provide most of the nutrients required, with very suitable potassium levels. The geochemical characters of this soil are: only few elements remain almost invariable across the profiles and over time, however the majority of them were directly linked with the clay content. These soils are characterized by relatively low levels of some trace elements such as Sr ($64.35 \text{ mg}\cdot\text{kg}^{-1}$), Ba ($303.67 \text{ mg}\cdot\text{kg}^{-1}$) and Sc ($13.14 \text{ mg}\cdot\text{kg}^{-1}$); high levels of other trace elements such as V ($103.92 \text{ mg}\cdot\text{kg}^{-1}$), Cr ($79.9 \text{ mg}\cdot\text{kg}^{-1}$), Cu ($15.18 \text{ mg}\cdot\text{kg}^{-1}$), Hf ($10.26 \text{ mg}\cdot\text{kg}^{-1}$), Ni ($38 \text{ mg}\cdot\text{kg}^{-1}$) and Zr ($337 \text{ mg}\cdot\text{kg}^{-1}$); while the levels for rare earth elements (REE's) such as La ($48.36 \text{ mg}\cdot\text{kg}^{-1}$), Ce ($95.07 \text{ mg}\cdot\text{kg}^{-1}$), Th ($13.33 \text{ mg}\cdot\text{kg}^{-1}$) and Nd ($42.65 \text{ mg}\cdot\text{kg}^{-1}$) are significantly high. The distribution of mayor and trace elements was directly related to weathering processes, parent material and anthropogenic activities.

Keywords: Trace Elements, Castilla-La Mancha Region, Red Soils, Soil Geochemistry

1. Introduction

Red Mediterranean soils are common in the semiarid regions of Spain, but are also broadly distributed worldwide. The name "red soil" refers to the characteristic colour found in the B horizon. Their hue is redder than 5 YR, and their chroma is over 5 [1]. Bech *et al.* [2] identified the following specific properties of Mediterranean red soils: their chroma exceeds 3.5, they possess a B₁ horizon and they are independent of their parent material.

Their characteristics can be associated with four circumstances: high anthropogenic influence, specific climate, orographic regions higher than 300 m in altitude, and presence of Saharian fine dust from the desert [1].

The common red colour of these soils has been attrib-

uted to hematite [3-5]. However, these soils are formed by a large number of different kinds of soil with different parent materials. This fact increases the geochemical variety attributable to location. Different anthropogenic activities are important as well.

In the region of Castilla-La Mancha, the existing red Mediterranean soils constitute 16.7% and represent the second most abundant soil type. They are exceeded in abundance only by the soils originating from Tertiary sediments (Miocene) [6]. These soils, formed over Ordovician-Silurian rocks, originate from the erosion and weathering of the old Hercinic socket. They are classified as Luvisols, with rhodic or chromic character, owing to the presence of iron oxides [7]. The main materials found over the Hercinic socket result from the weather-

ing and erosion of quartzite and materials from the schist-greywacke complex [8].

The main economic use of these soils in the region is the production of premium-quality wines. These soils represent 27.2% of the total regional area dedicated to the cultivation of the wine grape [6,9].

Several studies have been carried out on red soils from Mediterranean regions and areas of influence because of their different origins, parent material and characteristics [10-15]. Indeed, recently published studies have addressed the area under consideration [16,17]. However the soils from the region of Ciudad Real have been poorly studied, particularly in relation to their content of trace and major elements. This specific area of the region is very important because it includes a large percentage of the total production of wine grape [18].

Trace elements are defined as those elements that are present in a rock at concentrations below 0.1% or 1000 mg·kg⁻¹ [19]. The distribution and content of these chemical elements in soils depend on several factors: nature of the parent material, weathering processes and human activity [20]. Trace elements occur naturally in soils. Some of these elements are essential nutrients for plant growth as well as for human and animal health. However, at elevated levels some trace elements become potentially toxic, therefore acting as soil sinks for these compounds [21]. To give an idea of naturally occurring concentrations, **Table 1** shows the mean content of trace elements for the Castilla-La Mancha region [22] and in the world [23].

Marques *et al.* [24] noted that trace elements that accumulate during weathering are either pentavalent (Nb⁵⁺), tetravalent (Ti⁴⁺, Zr⁴⁺ and Th⁴⁺) or trivalent (Sc³⁺, V³⁺, Cr³⁺, Ga³⁺, Y³⁺, La³⁺, Ce³⁺) in soils that have acidic pH and are well-aerated. However, elements that are monovalent (Rb⁺) or divalent (Mn²⁺, Co²⁺, Ni²⁺, Cu²⁺, Zn²⁺, Sr²⁺, Ba²⁺ and Pb²⁺) are depleted at acidic pH.

Tetravalent cations crystallise into minerals that are very resistant to weathering and thus persist and accumulate (such as zircon, ZrSiO₄). Zr is probably the most immobile of all trace elements [25], even less so than Ti. Nb can replace Zr and Ti in clay structure [24,26]. Ti and Zr (Al and Si as major elements) are classed as residual elements after long weathering periods [27]. However, several other experiments provide evidence for the mobility of Zr [28].

In this context, threshold levels are not yet defined by law in Castilla-La Mancha. Because of the high level of industrial and agricultural activities, anthropogenic input of trace elements into the natural environment in the region represents a potential hazard [17]. The aforementioned trace elements can be absorbed by plants [23,29] and fungi [30] in different amounts and can accumulate

Table 1. Mean concentrations for trace elements in Castilla-La Mancha region [22] and in the world [23].

Trace element	Concentrations (mg·kg ⁻¹)	
	Castilla - La Mancha	World
As	7.4	
Ba	390	527
Ce	57.7	55
Co	5.8	10
Cr	54.8	67
Cs	7.7	
Cu	10.3	24
Ga	11.1	17
Hf	4.9	
La	23.6	31
Mo	0.9	
Nb	8	16
Nd	21.6	
Ni	16.9	24
Pb	19.3	29
Rb	86.2	73
Sc	23.6	9
Sn	4	
Sr	380	190
Ta	2.2	
Th	9.6	8
U	3.8	4
V	49.9	100
W	3.2	
Y	17.9	18
Zn	35.7	67
Zr	167	307

in different structures (roots, leaves, fruits). They can then pass into the human food chain from these plants and fungi.

The goals of this study are to evaluate potential problems in wine production associated with these soils and

to formulate recommendations for improved management. Accordingly, the study sought to provide pedological data from a typical red soil catena of the region and to study the associated geochemical properties in order to describe the distribution and content of trace elements in relation to other geochemical composition. The results of the study would then serve to evaluate potential risk to human health through agricultural production.

2. Materials and Methods

2.1. Site Description

The sampling area was located near the Almagro-Carrión

de Calatrava road (3 km from Almagro). The area is in the centre of the province of Ciudad Real (**Figure 1**), around 12 km southeast of the capital (UTM coordinates: 38°55'44"N; 3°43'30"W). Samples were carefully taken from each profile and horizon with a small shovel (**Figure 2**). The soil samples were taken from 3 profiles (toposequence) exposed by a machine along the slope of the hill called "Cerro Molino" (**Figure 3**). The selected soil profiles were identified and described morphologically according to FAO Guidelines [7].

Profile 1 (P1) was taken near the 680 m summit (**Figure 2**). The site is used as a temporary sheep pasture. Profile 3 (P3) was taken near the piedmont of the same hill, about 500 m down the slope at an altitude of 650 m.

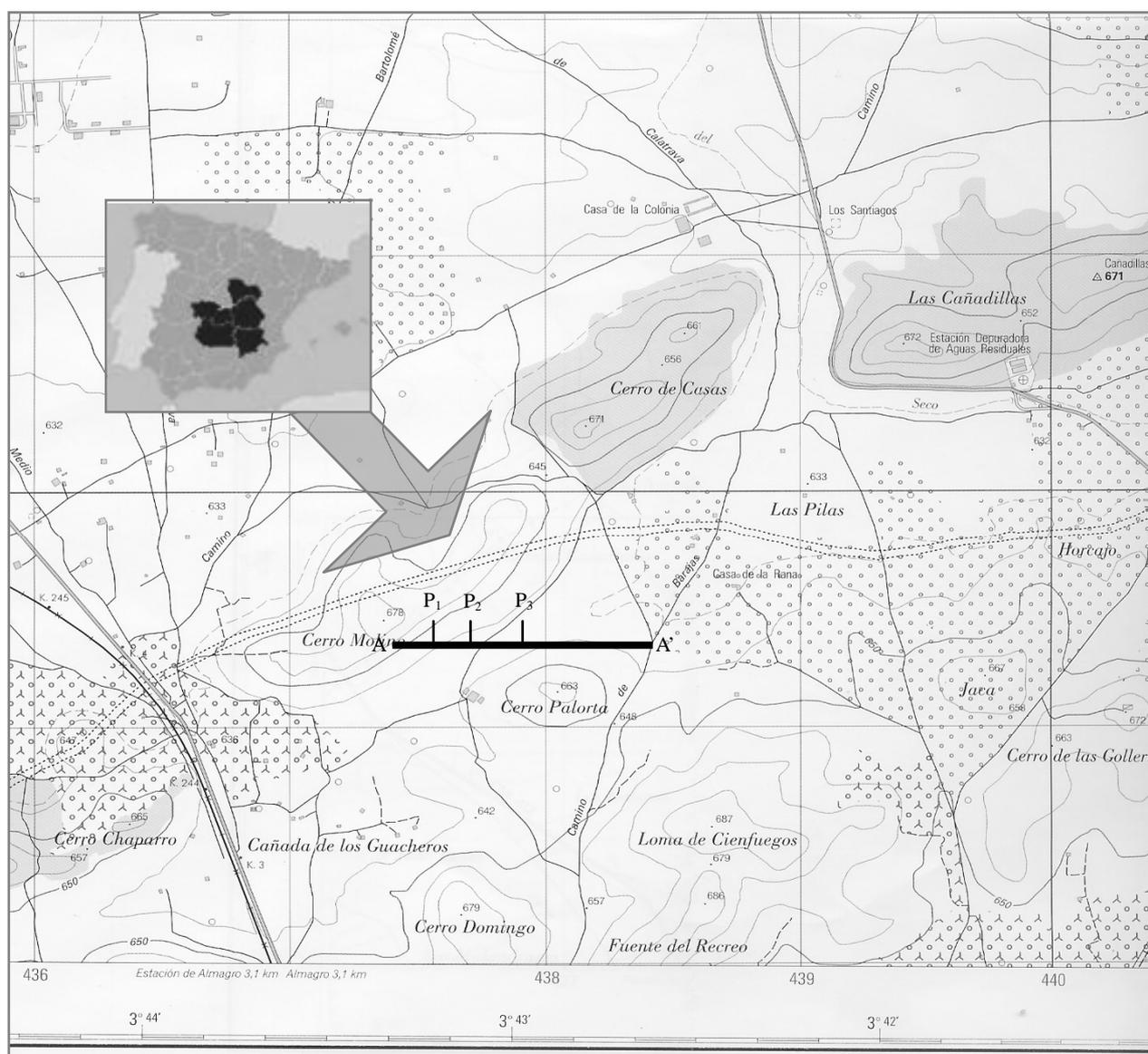


Figure 1. Geographical location of the profiles. From I.G.N. E.1:25000, Sheet 785-I "Base de Almagro" (UTM coordinates: 38°55'44"N; 3°43'30"W).

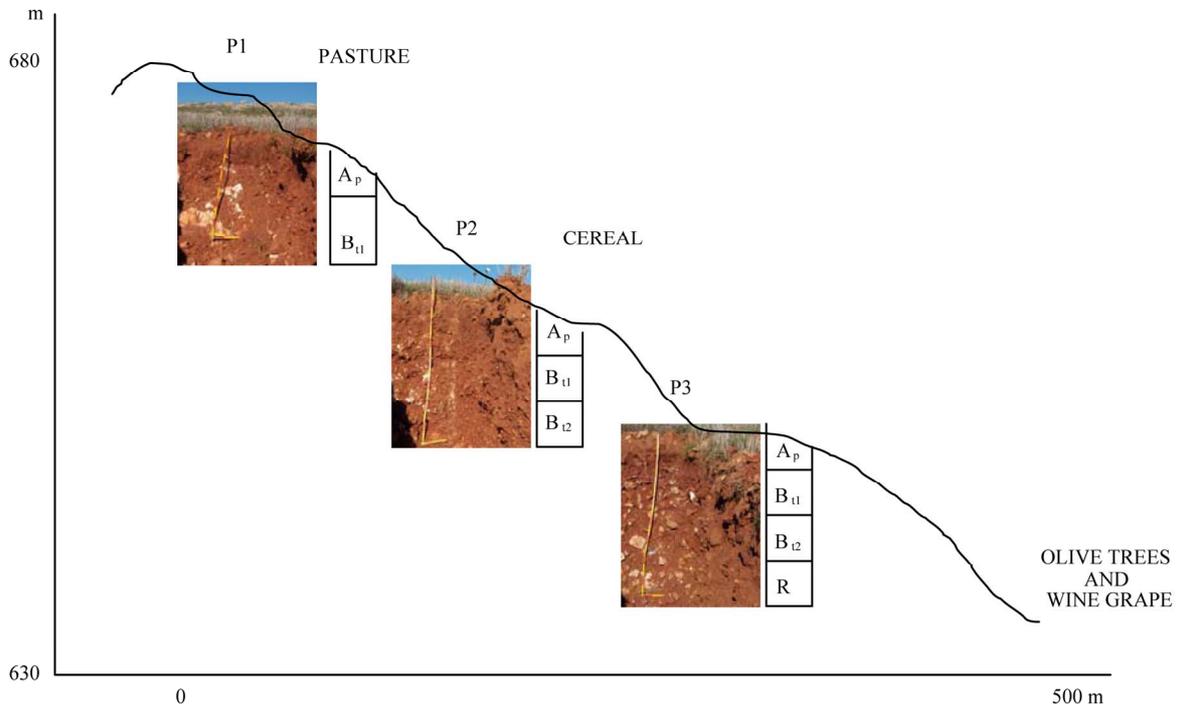


Figure 2. Location of profiles P1, P2 and P3 along the slope of Cerro Molino and their agricultural uses.



Figure 3. Sampling area.

The site is used for the culture of woody crops (olive tree and wine grape). Profile 2 (P2) was taken at an intermediate point between P1 and P3. This location is traditionally used for dry farming (cereal-fallow).

Selected climatic information is given in **Figure 4**, based on data supplied by the Meteorological Station of Ciudad Real. This station is located near the sampling area. The mean temperature and rainfall throughout the year were sampled from 1970 to 2009. A Mediterranean climate modified by altitude is evident and is typical of the central high plateau of Spain. This area has low mean rainfall (410 mm/year), especially in summer when a xeric period occurs from mid-June to the end of September. The evapotranspiration is negative (average 400 mm/year). The mean temperature is 14.4°C and exhibits a wide range between winter and summer.

2.2. Analytical Procedures

All the analytical determinations were carried out according to SCS-USDA [31]. All samples were extracted and analysed in triplicate.

Specifically, soil texture was determined using the hydrometer method [32]. Soil pH was measured in H₂O and in 0.1 M KCl using a 1:2.5 soil/solution ratio. Electrical conductivity was measured in a 1:5 soil:water extract. The method of Olsen *et al.* [33] was used to estimate available P. The soil organic matter was quantified by

the Walkley and Black wet oxidation method [34]. Exchangeable cations were determined using an ammonium acetate extraction method [35]. Exchangeable Na, K, Ca and Mg were determined by atomic absorption spectrometry. Total nitrogen content was determined by the Kjeldahl method [36].

Semi-quantitative mineralogical analyses were carried out by X-ray diffraction (XRD) techniques. About 2 g of sample was hand-milled to below 53 µm in an agate mortar and used for the bulk mineralogy determination (random powder method). For the detailed study of phyllosilicates, 100 g of sample was treated to remove components that prevent complete dispersion (e.g. carbonates, sulphates, organic matter). The clay fraction (< 2 µm) particles were extracted by sedimentation techniques and analysed on “thick” glass slides by XRD according to Moore and Reynolds [37]. Samples were chemically treated [(a) ethylene glycol to detect expandable minerals and (b) dimethyl sulphoxide to differentiate chlorite and kaolinite] and thermally treated (550°C for 2 h, to study the behaviour of phyllosilicates). The samples were analysed using a CuK α radiation source (Philips-Panalytical X-PERT diffractometer) with a graphite monochromator, 40 kV and 40 mA, and a sensitivity of 2.103 cps. The ranges measured were 2 - 75° or 2 - 50° 2 θ , goniometer speed of 0.04 or 0.05 and time constant of 0.4 or 1 second for random powder or glass slides, respectively.

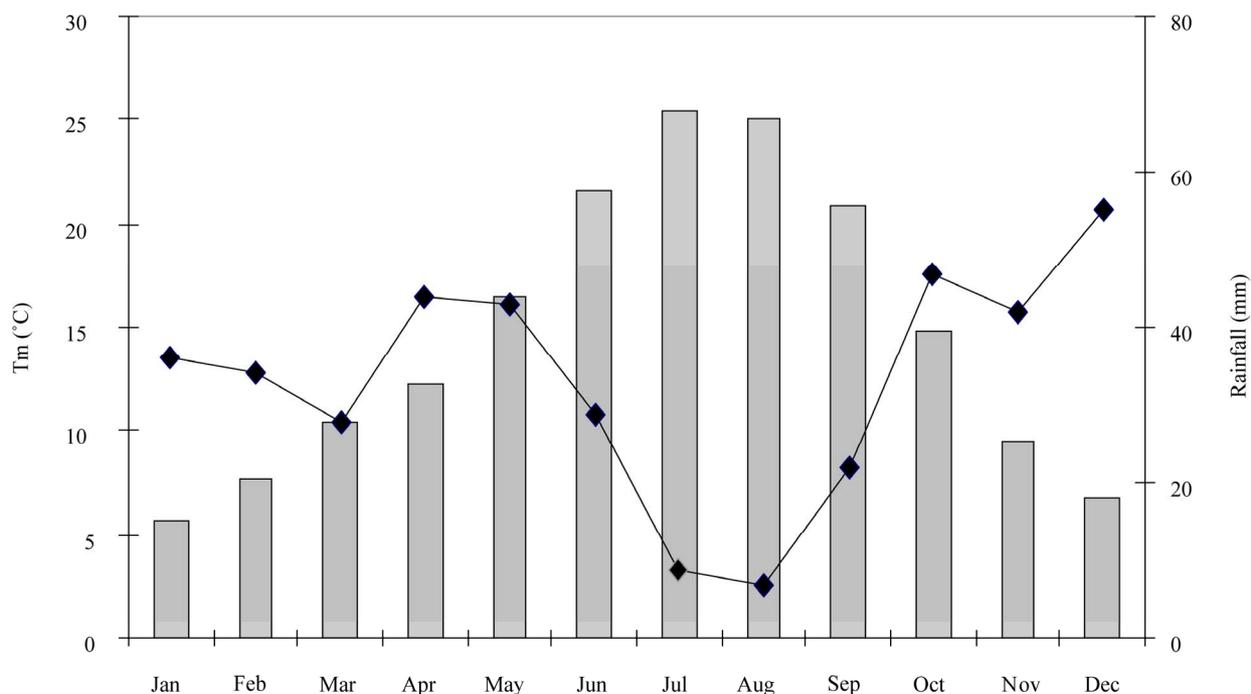


Figure 4. Mean temperatures (T_m, °C, in bars) and monthly rainfall (mm in line) throughout the year (meteorological station of Ciudad Real, Spain, 1970 - 2009).

The trace element content of whole samples was determined using an X-ray fluorescence spectrometer (PHILIPS PW 2404) in solid mode.

2.3. Statistical Analysis

The software packages XLSTAT_Pro version 7.5 (© 1995-2006 Addinsoft) and XLSTAT_3DPlot version 3.0 (©1995-2006 Addinsoft) were used for statistical and PCA analyses.

3. Results and Discussion

3.1. Profile Description, Granulometric and Mineralogic Properties

The characteristics of the three profiles (P1, P2 and P3) in terms of depth, texture, stoniness and boundaries between studied profiles are included in **Table 2**, detailed by horizon. The main differences among the profiles concern the thickness of each horizon and the boundaries between them. The plow layer (A_p) has almost the same thickness in each case, a result consistent with the use of soil. The B_{t1} horizon shows the same behaviour, but B_{t2} increases in depth towards the foot of the hill.

The textures of the profiles tend to be rich in clay, especially at depth. This property will increase the Cation Exchange Capacity (C.E.C.) and the potential water retention [12,38]. This clayey character of soils from the sequence increases their adsorptive properties.

The B_{t1} horizon has a level of clay about 15% higher than that of the layer above. The layer thickness in all three cases is very similar, a value of about 30 cm that increases slightly at the bottom of the slope (P3). Finally, in horizon B_{t2} the clay content reaches 60% (P2). At the foothill site, the profile is much deeper (more than 140 cm depth).

In each case the stoniness is high and increases with depth. The boundaries between the horizons A_p/B_{t1} are gradual and irregular (G.I.) and gradual and undulated (G.U.). The limit is net and plane (N.P.) between B_{t1}/B_{t2} owing to the higher compactness of the B_{t2} horizon.

The mineralogy of the finer and clay fractions is represented in **Figure 5**. The high content of phyllosilicates (more than 70% in all horizons) is noteworthy. Illite is predominant, with lower amounts of Kaolinite (15 to 20%). Illite usually provides significant amounts of native potassium [38]. Smectite was not detected, and this finding is consistent with the lack of vertic character.

3.2. Chemical Characteristics and Soil Classification

The main chemical characteristics of the profiles studied

are given in **Table 2**. Trace element content is considered in the next section. As indicated in **Table 2**, the pH is slightly acidic.

Only in the case of P3 does the B_{t2} horizon possess a slightly higher pH. This property is probably the result of migration of calcium carbonate from some source material (Miocene) below or at the same level. The neutral or slightly acidic pH induces the availability of certain elements (mainly Fe, Mn and Zn), whose migration is blocked in calcareous soils [10,39,40].

The C.E.C. values of the soils under investigation are over 20 $\text{cmol}\cdot\text{kg}^{-1}$ (**Table 2**). This result is consistent with results obtained in surrounding areas [16] and in other regions in Spain [12] that have Mediterranean character and similar types of soil. The low pH - BS relation in the surface layers can be attributed to a possible effect of nitrification as a result of intensive nitrogen fertilization. Salinity problems were not detected in the profiles studied.

It is worth noting that organic matter content is low, ranging from 0.4% - 2.4%. The organic matter content is only substantial in the A_p horizon and drops significantly in B_t (to around 1% in fine earth). The observed values of organic matter content are mainly the result of the warm climate and tillage management of ploughed land.

The available P levels are given in **Table 2**. The amounts found in these profiles show that the P is exogenous in origin (fertilisers and O.M. mineralisation).

The exchange complex is not saturated (BS%), especially in the A_p horizon, owing to leaching and removal of crops (dystric character). In the B_{t1} horizon, a greater degree of saturation is observed, with values above 60% (eutric character) that increase to 100% in the B_{t2} horizon of P3. Such variation has been reported for similar soils and environments [12].

The exchange cations are dominated by calcium, particularly in the B_t horizons. The magnesium and potassium levels are normal and balanced according to levels reported by Lanyon *et al.* [41]. The levels of sodium were not significant.

The three soils under investigation are classified as Typic Rhodoxeralf [31]. In terms of the F.A.O. (2006) classification, these soils are poorly developed Luvisols (prefix Haplic for P1), highly coloured (Rhodic second suffix), with a clear and sharp contact with bedrock near the surface in P1 and P2 (Ruptic first suffix) and high stoniness in P3 (Skeletal first suffix).

The contents of nine major elements were analysed. The results are reported in **Table 3**. From the content obtained for each element, the following behaviour patterns were observed:

1) Some elements, such as Al, Fe, Ca and Mg, tend to increase with depth in relation to the content of the clay

Table 2. Soils characteristics between studied profiles (P1, P2 and P3) detailed by horizon (P is given in mg·kg⁻¹ and C.E.C., Ca²⁺, Mg²⁺, K⁺ and Na⁺ is given in cmol·kg⁻¹).

Horizon	Depth (cm)	Colour		Clay (%)	Coating	Stoniness (%)	Boundary (%)	Texture	pH		Electrical Conductivity (dS/m)	P (Olsen)	Organic matter (%)	Cations			C.E.C cmol·kg ⁻¹	BS (%)	
		Dry	Wet						H ₂ O	KCl				Ca ²⁺	Mg ²⁺	K ⁺			Na ⁺
PROFILE 1																			
A _p	0-12	5 YR 4/6	2,5 YR 4/4	21.4		30	G.I.	Sandy-Clay-Loam	6.3	5.2	0.25	2	1.4	3.5	0.8	0.6	1.9	23.4	29.05
B _{1t}	12-55	2,5 YR 5/6	2,5 YR 3/4	37.4	slim and discontinued	>50	N.P.	Clay-Sandy	6.7	5.8	0.33	0.6	0.4	10.3	1.9	0.3	2.1	21.9	66.0
PROFILE 2																			
A _p	0-15	2,5 YR 4/4	2,5 YR 3/4	20.9		30	G.I.	Sandy-Clay-Loam	6.2	5.0	0.28	3.7	1.5	4.7	0.8	0.5	2.0	22.0	36.36
B _{1t}	15-50	2,5 YR 4/6	2,5 YR 3/6	38.9	thick and dense	25	N.P.	Clay-Sandy	6.7	5.7	0.47	0.3	0.8	12.7	1.3	0.3	1.7	25.7	62.25
B ₂	50-125	10 R 4/3	2,5 YR 5/8	52.2		>50	N.I.	Clay	6.4	5.5	0.39								
PROFILE 3																			
A _p	0-16	5 YR 4/6	2,5 YR 4/4	25.8		20	G.U.	Sandy-Clay-Loam	6.2	5.6	0.17	1.7	2.4	5.9	1.2	0.6	2.0	25.2	38.49
B _{1t}	16-50	2,5 YR 4/6	2,5 YR 3/4	35.8	slim and discontinued	40	G.I.	Clay-Sandy	6.4	5.2	0.13	0.2	1.1	12.7	2.4	0.4	1.4	19.8	85.35
B ₂	50-145	2,5 YR 4/6	2,5 YR 4/8	61.4		>50	D.I.	Clay	7.3	6.4	0.2								
R	>145																		

*G.I.: Limit gradual and irregular; G,U.: Limit gradual and undulated; N.P.: Limit net and plane; N.I.: Limit net and irregular; D.I.: Limit diffuse and irregular.

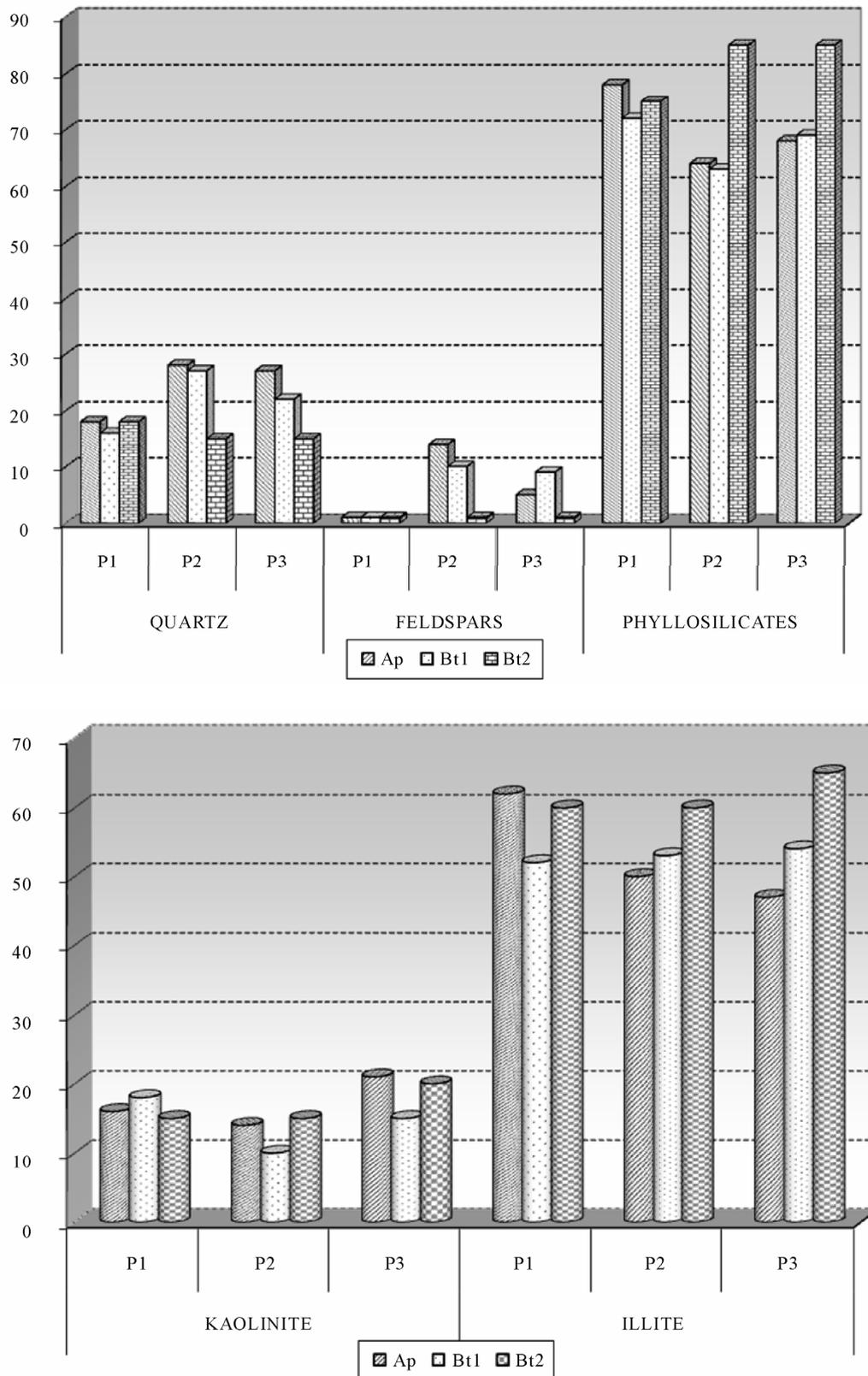


Figure 5. Mineralogy of finer and clay fractions. Top: percentage of minerals <math>< 2 \mu\text{m}</math>. Bottom: Percentages of major clay types detected (smectite was not detected).

fraction. K, Ti and Si do not vary with depth, and only a small increase is observed in the intermediate horizon B₁₁. The other elements, such as Mn and Na, become depleted with depth.

2) Slight enrichment in Mn was found at the surface in the three profiles. This result suggests an allochthonous origin for Mn.

3) Finally, variations in the levels of the major elements within a given toposequence are related to the parent material, slope gradients and vegetation cover.

3.3. Trace Element Content

The results for trace elements are reported in **Table 4**. In comparison with the values shown in **Table 1**, the content of trace elements in this particular red soil in La Mancha was within the range of background values for Castilla-La Mancha and is comparable to average values worldwide. The most obvious variations with respect to average levels are discussed in this section in order to show that trace element content can serve as a fingerprint of the studied soil catena. In general, when the concentration is less than 5 mg/kg (detection threshold) only inconsistent conclusions can be drawn. Such was the case for U, Ta, W, Sn and Mo.

Ti, Nb and Th content remained constant in all profiles and horizons. The permanence observed at the surface for Zr and Hf content is consistent with the direct linkage of these elements with sand content.

Zr minerals are linked to sand particle size and were more abundant in A_p horizons, as noted in other studies [42,43]. These elements showed higher content in the soil under investigation than in the average values reported for Castilla-La Mancha (**Table 1**).

The behaviour of trivalent elements is more complex because of their ionic radius (similar to that of Al and Fe).

This behaviour allows them to be incorporated into the structures of kaolinite, hematite and goethite, which make up most of the clay fraction. In the samples, this fraction increased with depth in the B₁₁ and B₁₂ horizons. This finding is consistent with results reported by Marques *et al.* [24] and is also the case for the values found for Sc, V, Cr and Ga.

Lanthanides, also known as rare earth elements (REEs), have an affinity to oxygen as trivalent cations. Their presence can be a strong indicator of pedological processes. We found a uniform distribution of REEs within profiles and horizons. No enrichment as a function of depth was evident. It can be concluded that the studied profiles have the same origin and that no lithological discontinuity exists.

In contrast, lithological discontinuity has been found in other soils in Castilla-La Mancha (Malpica) [44] and in other regions [45] based on examination of REE distributions. Moreover, the relatively high content of La, Ce, Nd and Th represents part of the fingerprint of these soils in comparison with others from Castilla-La Mancha that are calcareous, volcanic or igneous in origin [9,17,46]. Zn and Ni levels were relatively high (46 mg·kg⁻¹), as expected, given that the original rock was quartzite. In our case, slightly acidic soils without smectite (**Figure 2**) yielded lower Ca, Mg, Ba and overall Sr content (3 or 4 times less) compared with the average values found in soils in Castilla-La Mancha (**Table 1**). Ca is the most important element in change complex saturation. In the case described here, it almost never reached sufficiently high levels to influence the depletion of REE. Sr is more abundant in the gypsum and calcium carbonate-rich soils of Castilla-La Mancha [22]. Rb content was relatively low in comparison with levels in soils that originate over granitic rocks.

In order to enhance our understanding of the distribu-

Table 3. Major elements (in alphabetic order) in oxide form given in % from all the horizons.

	P1 A _p	P1 B ₁₁	P2 A _p	P2 B ₁₁	P2 B ₁₂	P3 A _p	P3 B ₁₁	P3 B ₁₂	Mean
Al ₂ O ₃	14.42	19.01	14.89	19.30	24.82	13.42	20.93	23.99	18.85
CaO	0.28	0.46	0.33	0.48	0.71	0.35	0.60	0.80	0.50
Fe ₂ O ₃	4.08	5.49	4.11	5.32	8.02	3.64	5.82	7.73	5.53
K ₂ O	2.51	2.73	2.59	2.91	2.77	2.17	2.47	2.16	2.54
MgO	0.82	1.14	0.87	1.14	1.41	0.69	1.05	1.30	1.05
MnO	0.07	0.05	0.08	0.06	0.04	0.06	0.06	0.03	0.06
Na ₂ O	0.22	0.18	0.25	0.19	0.11	0.24	0.14	0.08	0.17
SiO ₂	71.97	64.09	71.19	63.59	51.48	73.60	60.98	51.77	63.58
TiO ₂	0.87	0.89	0.84	0.94	0.82	0.76	0.84	0.78	0.84

Table 4. Trace element content for horizons belonging to the three profiles studied. Average content in mg·kg⁻¹.

	P1 A _p	P1 B ₁₁	P2 A _p	P2 B ₁₁	P2 B ₁₂	P3 A _p	P3 B ₁₁	P3 B ₁₂	Average Concentration
As	12.1	13	11.6	14.1	17.3	10.6	13.2	16	13.48 ± 2.10
Ba	304.4	304.3	316.6	352.7	304.3	258.7	295.7	292.7	303.67 ± 24.44
Ce	97.4	103.9	103.3	98.7	96.9	88.7	87.7	84	95.07 ± 6.93
Co	9.1	11.4	10.1	12.6	11	9.4	12.5	12.4	11.06 ± 1.31
Cr	68.5	70.2	64.8	78.6	102.3	60	86.4	108.4	79.90 ± 16.60
Cs	4	8.7	6.8	7.3	12.3	2.2	6.6	11.6	7.43 ± 3.22
Cu	13.5	15.4	15.3	16.4	17.5	11.1	14	18.3	15.18 ± 2.16
Ga	9.3	13.3	10	13.1	20.5	8.9	14.3	21.3	13.83 ± 4.48
Hf	12.7	11.6	13.5	9.8	8	10.8	7.8	7.9	10.26 ± 2.10
La	42.9	50.7	46.8	53.2	56.5	40.4	48.6	47.8	48.36 ± 4.88
Mo	0.4	0.2	0.6	0.5	0.7	0.4	0.3	0.9	0.50 ± 0.21
Nb	15	16.5	15.8	17.1	17.5	14.4	16.2	17.8	16.28 ± 1.11
Nd	40.1	46.3	43.2	42.2	47.6	40.4	41	40.4	42.65 ± 2.68
Ni	26.3	32.9	28.3	34.6	49.6	27.5	43.1	61.7	38.00 ± 11.73
Pb	19.9	18.5	20.6	19.6	21.3	21.7	21	20.1	20.33 ± 0.96
Rb	61.9	77.9	67.1	81.7	95.7	55.9	73.7	89.4	75.41 ± 12.69
Sc	9.6	13.5	9.9	12.9	18.9	8.4	14.3	17.6	13.14 ± 3.53
Sn	0.7	0.9	0.2	1.5	1.8	0.6	1.1	1	0.97 ± 0.47
Sr	57.6	67.4	61.1	68.8	74.6	56.2	60.5	68.6	64.35 ± 6.01
Ta	1.6	1.7	0.9	1.1	2.5	1.6	1.1	1.6	1.51 ± 0.46
Th	13.6	14.2	13.5	14.4	13.6	12	12.4	13	13.33 ± 0.77
U	3	3	3.2	3.6	2.8	3.3	2.5	2.8	3.02 ± 0.31
V	76.7	101.9	79.7	100.3	147.8	70.6	109.6	144.8	103.92 ± 27.57
W	1.4	1.2	1.9	2.8	1.1	2	1.8	2.7	1.86 ± 0.59
Y	21.9	21.5	22.6	23.4	20.4	20.5	20	18.9	21.15 ± 1.37
Zn	36.3	45.5	39.7	46.9	63.2	32.5	46.4	60.7	46.40 ± 10.17
Zr	434.5	346.9	444.5	373.2	234	368.8	286.4	212.1	337.55 ± 80.81

tion of trace elements (**Table 4**), a Principal Components Analysis (PCA) was performed on the data (**Figure 6**). Each horizon was considered to be a vector with 26 variables (trace element content). Of the seven principal components given by the program, F1 and F2 explained 76.89% of the total variance. Thus, the best representation was that in which the X axis corresponded with F1

(56, 74%) and the Y axis corresponded with F2 (20, 15).

Several relationships between trace elements can be inferred from the correlation matrix (**Table 5**). Zr was highly correlated with Hf (0.92), and both accumulate in surface horizons. However, the elements that are depleted correlate with each other [for example Sr correlates with Rb (0.96), As (0.9), Zn (0.89), Cu (0.88), V

Table 5. Correlation matrix. (Highlighted in black significant values; $\alpha = 0.050$ (bilateral probe)).

	Sc	V	Cr	Co	Ni	Cu	Zn	Ga	As	Rb	Sr	Y	Zr	Nb	Mo	Sn	Cs	Ba	La	Ce	Hf	Ta	W	Pb	Th	U	Nd
Sc	1.00	0.99	0.95	0.67	0.90	0.83	0.99	0.98	0.96	0.96	0.86	-0.52	-0.90	0.91	0.52	0.76	0.94	0.15	0.77	-0.25	-0.80	0.53	-0.02	-0.04	0.18	-0.54	0.47
V	0.99	1.00	0.97	0.64	0.94	0.83	0.99	1.00	0.96	0.94	0.84	-0.57	-0.92	0.90	0.60	0.72	0.93	0.10	0.71	-0.32	-0.80	0.54	0.04	0.01	0.12	-0.52	0.41
Cr	0.95	0.97	1.00	0.63	0.97	0.79	0.95	0.97	0.93	0.88	0.72	-0.61	-0.90	0.85	0.68	0.67	0.85	0.08	0.59	-0.47	-0.84	0.44	0.17	0.09	0.02	-0.55	0.19
Co	0.67	0.64	0.63	1.00	0.66	0.62	0.63	0.63	0.58	0.69	0.59	-0.19	-0.61	0.79	0.18	0.57	0.60	0.43	0.64	-0.26	-0.70	-0.17	0.47	-0.28	0.20	-0.22	0.11
Ni	0.90	0.94	0.97	0.66	1.00	0.74	0.91	0.96	0.85	0.81	0.65	-0.72	-0.94	0.82	0.68	0.54	0.82	-0.06	0.47	-0.58	-0.84	0.36	0.26	0.10	-0.12	-0.56	0.11
Cu	0.83	0.83	0.79	0.62	0.74	1.00	0.89	0.83	0.87	0.92	0.88	-0.13	-0.57	0.96	0.67	0.53	0.94	0.52	0.76	0.07	-0.44	0.25	0.21	-0.32	0.53	-0.18	0.43
Zn	0.99	0.99	0.95	0.63	0.91	0.89	1.00	0.98	0.98	0.97	0.89	-0.47	-0.86	0.93	0.64	0.72	0.96	0.21	0.77	-0.22	-0.74	0.51	0.06	-0.03	0.23	-0.44	0.46
Ga	0.98	1.00	0.97	0.63	0.96	0.83	0.98	1.00	0.95	0.93	0.82	-0.61	-0.93	0.89	0.65	0.68	0.92	0.06	0.66	-0.37	-0.80	0.52	0.10	0.02	0.08	-0.50	0.36
As	0.96	0.96	0.93	0.58	0.85	0.87	0.98	0.95	1.00	0.97	0.90	-0.38	-0.81	0.91	0.62	0.81	0.92	0.27	0.79	-0.18	-0.73	0.58	0.03	-0.03	0.29	-0.37	0.46
Rb	0.96	0.94	0.88	0.69	0.81	0.92	0.97	0.93	0.97	1.00	0.96	-0.28	-0.77	0.97	0.54	0.78	0.96	0.38	0.88	-0.05	-0.68	0.46	0.06	-0.17	0.41	-0.30	0.56
Sr	0.86	0.84	0.72	0.59	0.65	0.88	0.89	0.82	0.90	0.96	1.00	-0.09	-0.63	0.92	0.47	0.77	0.92	0.45	0.92	0.16	-0.52	0.51	0.00	-0.25	0.56	-0.09	0.70
Y	-0.52	-0.57	-0.61	-0.19	-0.72	-0.13	-0.47	-0.61	-0.38	-0.28	-0.09	1.00	0.80	-0.22	-0.34	-0.14	-0.34	0.71	0.10	0.80	0.67	-0.43	0.04	-0.38	0.66	0.74	0.13
Zr	-0.90	-0.92	-0.90	-0.61	-0.94	-0.57	-0.86	-0.93	-0.81	-0.77	-0.63	0.80	1.00	-0.72	-0.48	-0.65	-0.74	0.22	-0.49	0.60	0.92	-0.53	-0.08	-0.19	0.22	0.60	-0.22
Nb	0.91	0.90	0.85	0.79	0.82	0.96	0.93	0.89	0.91	0.97	0.92	-0.22	-0.72	1.00	0.57	0.68	0.94	0.47	0.83	-0.07	-0.64	0.26	0.24	-0.28	0.44	-0.24	0.44
Mo	0.52	0.60	0.68	0.18	0.68	0.67	0.64	0.65	0.62	0.54	0.47	-0.34	-0.48	0.57	1.00	0.20	0.59	0.07	0.22	-0.34	-0.36	0.23	0.43	0.28	-0.02	-0.06	-0.03
Sn	0.76	0.72	0.67	0.57	0.54	0.53	0.72	0.68	0.81	0.78	0.77	-0.14	-0.65	0.68	0.20	1.00	0.59	0.34	0.80	-0.12	-0.75	0.54	-0.01	0.01	0.30	-0.15	0.44
Cs	0.94	0.93	0.85	0.60	0.82	0.94	0.96	0.92	0.92	0.96	0.92	-0.34	-0.74	0.94	0.59	0.59	1.00	0.29	0.80	0.00	-0.56	0.45	-0.02	-0.20	0.38	-0.39	0.59
Ba	0.15	0.10	0.08	0.43	-0.06	0.52	0.21	0.06	0.27	0.38	0.45	0.71	0.22	0.47	0.07	0.34	0.29	1.00	0.61	0.56	0.10	-0.32	0.26	-0.52	0.83	0.39	0.24
La	0.77	0.71	0.59	0.64	0.47	0.76	0.77	0.66	0.79	0.88	0.92	0.10	-0.49	0.83	0.22	0.80	0.80	0.61	1.00	0.31	-0.48	0.33	-0.10	-0.23	0.60	-0.12	0.75
Ce	-0.25	-0.32	-0.47	-0.26	-0.58	0.07	-0.22	-0.37	-0.18	-0.05	0.16	0.80	0.60	-0.07	-0.34	-0.12	0.00	0.56	0.31	1.00	0.68	-0.05	-0.43	-0.50	0.77	0.42	0.60
Hf	-0.80	-0.80	-0.84	-0.70	-0.84	-0.44	-0.74	-0.80	-0.73	-0.68	-0.52	0.67	0.92	-0.64	-0.36	-0.75	-0.56	0.10	-0.48	0.68	1.00	-0.34	-0.23	-0.30	0.28	0.50	-0.05
Ta	0.53	0.54	0.44	-0.17	0.36	0.25	0.51	0.52	0.58	0.46	0.51	-0.43	-0.53	0.26	0.23	0.54	0.45	-0.32	0.33	-0.05	-0.34	1.00	-0.55	0.15	0.05	-0.30	0.55
W	-0.02	0.04	0.17	0.47	0.26	0.21	0.06	0.10	0.03	0.06	0.00	0.04	-0.08	0.24	0.43	-0.01	-0.02	0.26	-0.10	-0.43	-0.23	-0.55	1.00	-0.01	-0.08	0.41	-0.59
Pb	-0.04	0.01	0.09	-0.28	0.10	-0.32	-0.03	0.02	-0.03	-0.17	-0.25	-0.38	-0.19	-0.28	0.28	0.01	-0.20	-0.52	-0.23	-0.50	-0.30	0.15	-0.01	1.00	-0.76	-0.19	-0.19
Th	0.18	0.12	0.02	0.20	-0.12	0.53	0.23	0.08	0.29	0.41	0.56	0.66	0.22	0.44	-0.02	0.30	0.38	0.83	0.60	0.77	0.28	0.05	-0.08	-0.76	1.00	0.38	0.52
U	-0.54	-0.52	-0.55	-0.22	-0.56	-0.18	-0.44	-0.50	-0.37	-0.30	-0.09	0.74	0.60	-0.24	-0.06	-0.15	-0.39	0.39	-0.12	0.42	0.50	-0.30	0.41	-0.19	0.38	1.00	-0.08
Nd	0.47	0.41	0.19	0.11	0.11	0.43	0.46	0.36	0.46	0.56	0.70	0.13	-0.22	0.44	-0.03	0.44	0.59	0.24	0.75	0.60	-0.05	0.55	-0.59	-0.19	0.52	-0.08	1.00

(0.83), Ga (0.81) and Cr (0.72)].

Figure 6 indicates that when elements accumulate near the surface, they appear on the left of the y axis. Examples are Zr and Hf. Elements that accumulate at depth are grouped on the right of the y axis. Elements that appear in the centre of the graph and near to the y axis are not well represented in this PCA. It is difficult to draw any conclusions about them (Y, Ce, Th, Ba, Pb).

4. Conclusions

The results of this study indicate that the distribution of trace elements in the red soil sequence studied under semiarid conditions in the province of Ciudad Real can provide specific composition of this Mediterranean red soil.

This distribution is affected mainly by the following factors:

1) The specific parent material yields residual elements which are resistant to weathering. This is the case for Ti, Zr, Hf and Th.

2) The pedogenetic process, which explains the accumulation of clay at depth, allows the existence of numerous associated elements in the studied profiles, such as Fe, Ca, Sc, V, Cr and Ga.

3) The profiles have a significant degree of evolution. The semiarid environment can explain this finding. Lithological discontinuities were not observed.

4) In general, the three profiles have similar characteristics with respect to their main geochemical and pedological properties. The most notable characteristics are the low organic matter content and the predominantly acidic pH. Illite and kaolinite are the predominant clay minerals. The fertility of the soils is sufficient to provide most of the nutrients required, with very suitable potassium levels.

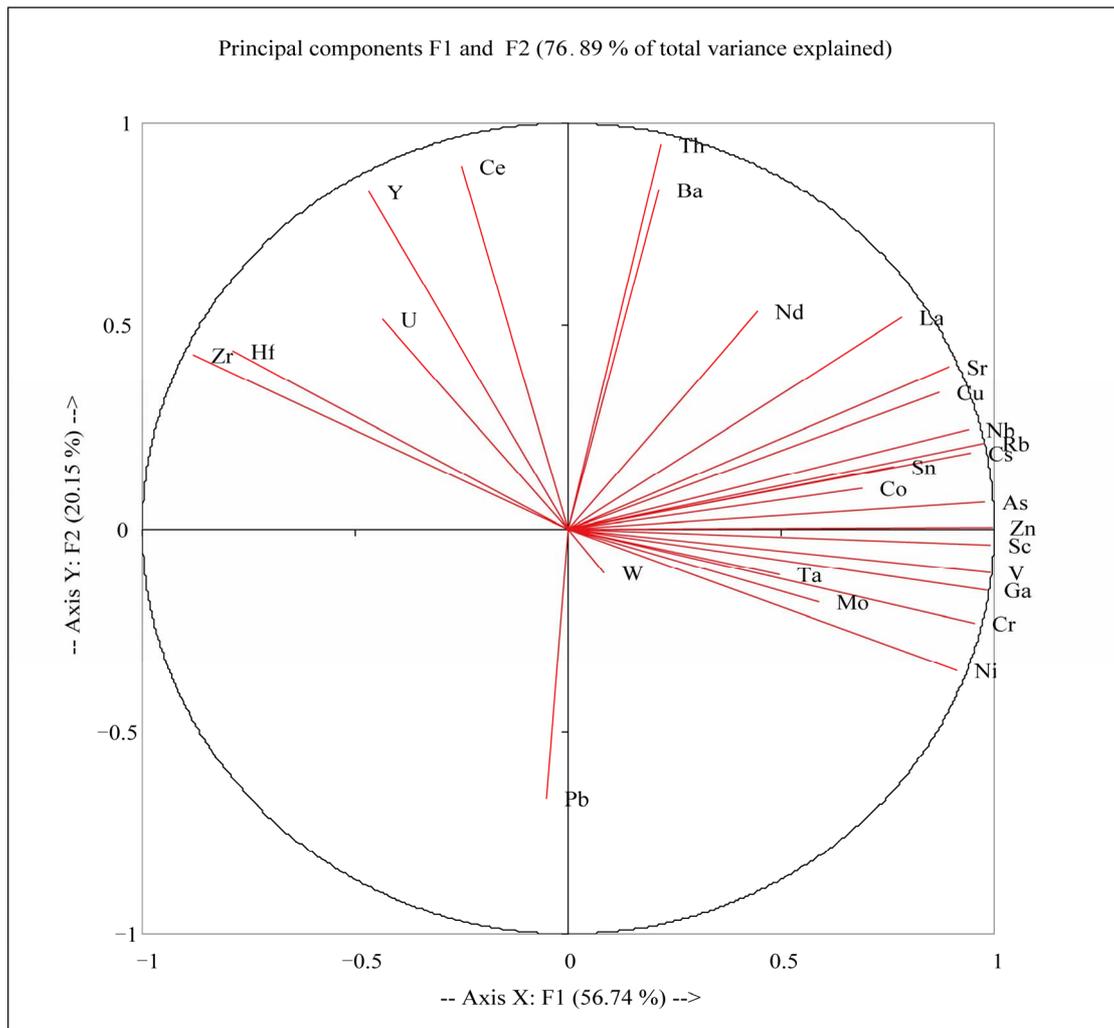


Figure 6. Variable representation of the PCA carried out on data from Table 5 with the two first principal components F1 (X axis) and F2 (Y axis).

5) Finally, the relatively low influence of human activity is indicated by the absence of surface enrichment of trace elements considered to be pollutants, namely Cu, Pb, Ni, Cr or Zn.

In general, the geochemical characteristics for this soil are:

1) Low levels of some trace elements such as Sr ($64.35 \text{ mg}\cdot\text{kg}^{-1}$), Ba ($303.67 \text{ mg}\cdot\text{kg}^{-1}$) and Sc ($13.14 \text{ mg}\cdot\text{kg}^{-1}$) and major elements like Ca and Mg.

2) High levels of some trace elements such as V ($103.92 \text{ mg}\cdot\text{kg}^{-1}$), Cr ($79.9 \text{ mg}\cdot\text{kg}^{-1}$), Cu ($15.18 \text{ mg}\cdot\text{kg}^{-1}$), Hf ($10.26 \text{ mg}\cdot\text{kg}^{-1}$), Ni ($38 \text{ mg}\cdot\text{kg}^{-1}$) and Zr ($337 \text{ mg}\cdot\text{kg}^{-1}$) and major elements like Fe.

3) High levels of REEs such as La ($48.36 \text{ mg}\cdot\text{kg}^{-1}$), Ce ($95.07 \text{ mg}\cdot\text{kg}^{-1}$), Th ($13.33 \text{ mg}\cdot\text{kg}^{-1}$) and Nd ($42.65 \text{ mg}\cdot\text{kg}^{-1}$).

In general, the results show that the texture of the soils and their stoniness can allow the development of crops as long as these crops do not have to compete for water. In particular, these soils can be used for the cultivation of high-quality grapes. To ensure the settling of the crop and as well as water-holding capacity, 1 m of soil must be left to accommodate the roots.

Finally, it is important to mention that the data obtained in this study can serve as a pattern for extended agro-edaphic areas (almost 20% from Ciudad Real province and more than 10% from Toledo province). The study data are potentially useful for evaluating potential uses of the soils in accordance with their content of trace elements. However, specific studies must be considered, depending on location, in order to eliminate other anthropogenic and allochthonous influences.

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