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Abstract

The range of metals used for industrial purposes - electrical engineering, solar panels, batteries - has increased substantially over the last twenty years. Some of these emerging metals are the subject of geopolitical conflict and are considered critical as their unique properties make them irreplaceable. Many of these elements are poorly studied and their biogeochemical cycles still raise many questions. Aim of this study is to analyse the soil-to-plant transfer of some of these chemical elements and to shed light on their uptake pathways.

For this purpose, the geological site of Jas Roux (France) was chosen as this alpine site is naturally rich in critical and potentially toxic elements such as As, Sb, Ba and Tl, but nevertheless is host to a high diversity of plants. Elemental concentrations were analysed in the topsoil and in 12 selected alpine plant species sampled in situ. Statistical tools were used to detect species dependent characteristics in elemental uptake.

Our analyses revealed accumulation of rare earth elements by *Saxifraga paniculata*, selective oxyanion absorption by *Hippocrepis comosa*, accumulation of Tl by *Biscutella laevigata* and *Galium corrudifolium* and an exclusion strategy in *Juniperus communis*. These findings advance our understanding of the environmental behaviour of critical metals and metalloids such as V, As, Y, Sb, Ce, Ba and Tl and might bare valuable information for phytoremediation applications.

Uptake patterns of critical metals in alpine plant species growing in an unimpaired natural site

Till Fehlauer¹, Blanche Collin¹, Bernard Angeletti¹, Catherine Santaella², Cedric Dentant³, Perrine Chaurand¹, Clement Levard¹, Cedric Gonneau, Daniel Borschneck¹, Jérôme Rose¹

1: Aix Marseille Univ, CNRS, IRD, INRAE, Coll France, CEREGE, Aix-en-Provence, France

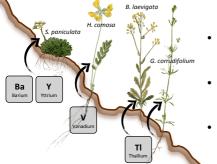
2: Aix Marseille Univ, CEA, CNRS, BIAM, LEMIRE, Laboratory of Microbial Ecology of the Rhizosphere, ECCOREV FR 3098, F-13108, St-Paul-lez-Durance, France

3: Ecrins National Park, Domaine de Charance, Gap, France

Corresponding Author:

till.fehlauer@cerege.fr

Graphinial Applicant species exposed to a cocktail of critical metals



Diverse species-related strategies

- Saxifraga paniculata
 - selective cation absorption
 - possible link between Y, Ba and Ca uptake
- Hippocrepis comosa
 - selective oxyanion absorption
 - possible link between V and Mo, Fe uptake
- Galium corrudifolium & Biscutella laevigata - Tl accumulation

Highlights:

- Critical metals: a need to understand their biogeochemical cycles
- Field-study of a plant community exposed to a natural cocktail of metals(loids)
- Multivariate analysis of elemental concentrations revealed species related uptake
- S. paniculata and H. comosa accumulating a different set of critical metals
- Observation of possible links in the soil/plant transfer of Mo/V/Fe and Y/Ca

1 1 Introduction

2 All over the world, emerging technologies require an increasingly diverse spectrum of chemical 3 elements with highly specific (photo-optical, catalytic, magnetic, etc.) properties (Celis et al., 4 2020). Metals that have practically never been used before are now indispensable for many industrial sectors and are thus the subject of soaring global demand. Due to their economic 5 importance and their supply risk, many of these metals are now classified as "critical" for high 6 7 tech applications (European Commission, 2020), examples being Sb in lead-acid batteries and 8 semiconductors, As in cell phones and solar cells and rare earth elements (REEs) in sensors, 9 displays and electric motors (Fortier et al., 2018; Mudd et al., 2018).

10 Despite their increased use, the biogeochemical cycles of many of these critical metals are not 11 vet fully understood and predicting either their environmental behaviour or their ecotoxicity is 12 consequently impossible (Espejo et al., 2018). Aim of this study is to analyse the soil-to-plant 13 transfer of some of these metals and to shed light on their uptake pathways. Our findings might 14 bare valuable information for phytoremediation and promote plant-based remediation 15 techniques of metal-contaminated sites. Phytoremediation techniques are relying on the 16 capacity of certain plant species to limit the uptake of the toxic elements or to accumulate them 17 in designated compartments (e.g. phytostabilisation, phytoextraction) (Favas et al., 2014). 18 Exclusion and accumulation are the two main strategies of plants to maintain homeostasis in 19 metal-rich habitats. Both involve a series of adaptions such as modified membrane transport, 20 chelation and complexation, or rhizosphere interactions for example (Furini, 2012).

The ability of plants to accumulate metals in their aerial parts has received considerable attention and many combinations of specific plant species tolerating or accumulating a particular set of elements have been discovered (Baker et al., 2010; Pollard et al., 2014). However, most studies have focused on "traditional" metals such as Zn, Ni, Cd and Pb, whereas the uptake pathways and metabolic impact of many elements, particularly emerging critical metals, still raise many questions (Macnair, 2003; van der Ent et al., 2013).

Elemental uptake is influenced by chemical, physical, and biological factors. Plant species, root system, root exudates, edaphic interactions, and transport mechanisms all play important roles (Tangahu et al., 2011; Trakal et al., 2015; Yadav, 2010). Despite highly specialised proteins, including transporters, pumps and channels, elemental selectivity remains challenging for plants and each uptake mechanism is likely to let pass a range of ions due to chemical similarities between the different ions (Tangahu et al., 2011). The huge number of factors and complex interactions are impossible to reproduce in the laboratory, thus, to understand the

- uptake of poorly studied elements, such as critical metals, performing in-situ studies in a natural
 environment is advantageous.
- 36 The geological site of Jas Roux, located in the Ecrins National Park (Hautes-Alpes, France), 37 was chosen for the study. The distinctive reddish colour of the Jas Roux outcrops ('Roux' 38 means red) originates from a sulphosalt-forming hydrothermal mineralisation (Johan and 39 Mantienne, 2000). These sulphosalts are one of the reasons for the characteristic metal 40 concentrations of Jas Roux as they are rich in elements with a strong affinity for S, such as Pb, 41 Sn, Cu, Tl and As (Dittrich et al., 2009; Pierrot and Roger, 2018). Many of the elements found 42 in Jas Roux are used in high-tech applications and are considered to be critical for leading 43 technologies (Dittrich et al., 2009; Graedel et al., 2015). This site is thus a good proxy of chronic 44 contamination by a cocktail of metals and metalloids. As the site has never been disturbed by 45 any human activities and is located in the heart of the Ecrins National Parc, it offers ideal 46 conditions to study an established ecosystem with a plant community that has not undergone 47 human selection. The plant community can be seen as natural response to the local mineralogy 48 and may contain candidates for phytoremediation.
- 49 To identify such plant candidates and to study the environmental behaviour of critical metals,
- 50 we were guided by the following three questions:
- 51 To what extent are critical metals absorbed by plants?
- 52 Are there any species-related differences in elemental uptake?
- 53 Do correlations exist between critical metals and other elements in different plant parts?
- 54 The approach of this study was to conduct a large-scale screening of 12 plant species to identify
- 55 individual uptake patterns using chemical analysis, soil element extraction and statistical tools.
- 56 The large number of species analysed provided insights into the actual elemental
- 57 phytoavailability of a wide range of critical metals.

58 2 Material and Methods

59 2.1 Sampling Site

The geological site of Jas Roux, located in the Ecrins National Park in the district of La 60 61 Chapelle-en-Valgaudemar (Hautes-Alpes, France), is situated at 2170 m a.s.l. (44°48' 45" N, 6° 19' 18" E; WGS84) on the right-hand side of the Chabournéou stream (Johan and Mantienne, 62 63 2000). Jas Roux is a hydrothermal stratiform deposit in dolomite limestone and has been 64 characterised as a sulphosalt site (Boulliard et al. 2010). The hydrothermal system was possibly generated by volcanic activity involving relatively low temperatures (around 100 °C) and high 65 66 pressure (Johan and Mantienne, 2000). The particular Tl-rich composition of Jas Roux has attracted many mineralogists and geologists, who have identified Zn, Sb, As, Tl, Ag, Hg, Pb 67 68 sulphides and sulphosalts including many new sulphosalt mineral species (Boulliard et al., 69 2010; Guillemin et al., 1970; Mantienne, 1974; Topa et al. 2017, 2014).

The study zone is located below one of the reddish coloured outcrops identified by Mantienne (Figure SM 1) (Mantienne, 1974). The metal-rich mineralisation is best developed in this outcrop and forms a 3.8 m thick barite-rich zone (Johan and Mantienne, 2000). The study zone measured 20 m x 5 m with a slope of approximately 45° (100%) (Figure SM 1).

- 74 Soil and plant samples were collected during two field trips, one in September 2017 and one in 75 June 2018. The steep slope, variable soil depth and the presence of large rocks impeded a 76 systematic sampling approach. Nevertheless, the samples were collected as evenly distributed 77 as possible. The soil samples were collected from 10 sampling points in the study zone and 78 consisted of 1 kg of topsoil each (< 15 cm). Among the 65 plant species identified in the study 79 zone, 12 were selected, taking into account occurrence, spatial distribution and family 80 affiliation (Table 1). For improved readability we attributed acronyms to each of the 12 species 81 throughout the paper (Table 1). Depending on their abundance, between 2 and 7 replicates of 82 each species were sampled, giving a total of 55 individuals (Table 1). All the samples were 83 stored at 4 °C in the dark.
- 84

85 **2.2 Sample preparation**

Roots and aerial parts were separated at the collar and cleaned with 18.2 M Ω ·cm ultrapure water. The plant roots were additionally brushed and vortexed in ultrapure water to remove adhering soil particles. The leaves of *Sax* were also brushed as they were partially covered with a calcite crust. Plant samples were dried at 40 °C for 48 h. Soil samples were dried at 40 °C for 72 h and sieved at 2mm before analysis.

91 All the soil and plant samples were ground at 20hz in a Retsch® Mixer Mill MM 400 for 2 min 92 to obtain a fine powder. Each sample (100mg) was solubilised in acid using a Milestone 93 UltraWAVE® microwave to complete mineralisation. The acids used for the digestion of the 94 plant samples were 4mL of HNO₃ and 2mL of H₂O₂. For the soil samples, 3mL of HNO₃, 3mL 95 of HCl and 0.5 mL of HF were used to measure the total metal concentrations. Soil pH and 96 conductivity were determined in a suspension of 1g of sieved soil (< 2mm) and 5mL of 97 ultrapure water according to the French standard X31-117 (NF X31-117, 1994). After 5 min of 98 agitation at 80 rpm and 2 h of rest, measurements were taken using a Hanna HI-2550 bench 99 meter. To estimate the mobile and phytoavailable elemental fractions in the soil, CaCl₂ 100 (0.001 mol/L) and NH₄NO₃ (1mol/L) extractions have been conducted, according to standards 101 ISO 21268-2 and ISO 19730 (ISO 19730:2008, 2009; NF X31-607-2, 2004). For the CaCl₂ 102 extraction, a liquid-solid ratio of 10:1 (20mL of ultrapure water and 2g of soil) was applied. 103 After 24 h of agitation at 10 rpm, the samples were centrifuged at 2500xG for 20 min, filtered 104 at 0.45 µm and analysed in ICP-MS. For the NH₄NO₃ extraction, 4g of soil were mixed with 105 10mL of ultrapure water, agitated for 2 h at 30 rpm, centrifuged at 1000xG for 15 min, filtered 106 at 0.45 µm and analysed in ICP-MS. 107 Soil mineralogy was analysed by X-ray diffraction using the X'Pert Pro MPD (XRD) equipped 108 with a Cobalt x-ray tube (= 1.79A) running at 40 kV and 40 mA.

109 A mixture of three bulk samples of topsoil was sent to an outside laboratory (CIRAD – US49 110 Laboratoire d'analyse de Montpellier, France) for the determination of soil parameters. The soil 111 type was determined using 5 granulometric fractions according to the French standard X31-107 112 (*NF X31-107*, 2003), organic matter by combustion analysis (Dumas), labile P content using a 113 colorimetric method (Olsen) and cation exchange capacity by quantifying the dosing adsorption 114 agent [Co(NH₃)₆]Cl₃ with inductively coupled plasma atomic emission spectroscopy according 115 to the French standard X31-130 (*NF X31-130*, 1999).

116

117 **2.3 Chemical analyses**

The samples were analysed by inductively coupled plasma mass spectrometry (ICP-MS, Perkin Elmer NexIon 300X). For the elements Mg, Al, P, K, Ca and S, the results of inductively coupled plasma optical emission spectroscopy (ICP-OES) were occasionally used to reach higher accuracy (Perkin Elmer 4300 DV). The calibration curves were established using multisolution standards CCS4, CCS5, CCS6 from Inorganic Ventures (New Jersey USA) diluted in 2% HNO₃ solution. The internal standards were Rh103 for ICP-MS and Y for ICP-OES. Oak leaves (V464) provided by the French National Institute for Agriculture Food and Environment (INRAE) were used as the external reference for plant samples. For the soil, a rendzina sample
(ISA 979) provided by Wageningen Evaluating Programs for Analytical Laboratories
(WEPAL) was used. EnviroMAT drinking water EP-L-3 and subterranean water ES-H-2 (SCP

128 Sciences) were used as internal references and for quality control.

129 ICP analyses aimed to quantify critical elements (As, V, Y, Sb, Ba, Ce, Tl) and major soil 130 elements and nutrients (Mg, Al, P, S, K, Ca, Mn, Fe, Ni, Cu, Zn, Mo) present in Jas Roux. The 131 critical elements were chosen based on the study on the EU's list of critical raw materials and a review of critical minerals and will be called critical metals throughout the manuscript 132 133 (European Commission, 2020; Hayes and McCullough, 2018). The detection limit for each 134 element was determined using handling blanks analysed at each ICP analysis (Table SM 1). 135 The coefficients of variances for all certified elements were calculated for the internal and 136 external standards (Table SM 1).

137 The translocation factor (TF) was calculated to identify the transfer strategies of the different 138 plant species. The TF is obtained by dividing the elemental concentrations in the aerial parts by 139 the concentrations in the roots (Radziemska et al., 2017). The TF was calculated for each plant 140 sample and averaged according to the respective plant species. A heatmap with adjusted 141 colormap breaks was used to illustrate the TF.

142

143 **2.4 Statistical analysis**

144 All statistical analyses were performed with the open-source software R studio. The complete 145 dataset for the plants contains 110 observations (aerial parts + roots) and 19 variables (chemical 146 elements). In the first step, a principal component analysis (PCA) was conducted on the aerial 147 parts and the roots separately to capture the essence of the datasets and to identify the features 148 that explain most of the variation in chemical composition. For this purpose, the packages 149 'FactomineR' and 'factoextra' were used (Kassambara, 2017a; Kassambara and Mundt, 2020; 150 Lê et al., 2008). Missing values were implemented using the package 'missMDA', which 151 reduced their influence on the PCA results to a minimum (Josse and Husson, 2016).

In the second step, an unsupervised clustering analysis was performed. The clustering tendency of the two datasets (aerial parts + roots) was tested using the Hopkins test from the package 'clustertend' before performing portioning clustering (K-means) (YiLan and RuTong, 2015). K-means is a clustering algorithm that returns the natural grouping of data points based on their similarity (elemental concentrations), not on their identity (plant species). The determination of the number of clusters as well as the clustering itself and the visualisation were performed using the package 'factoextra' (Kassambara, 2017b; Kassambara and Mundt, 2020). Next, a 159 PERMANOVA was performed on the complete dataset, using the package 'vegan' (Oksanen 160 et al., 2020) to examine to what degree the factors "plant species" and "plant parts" influenced 161 the elemental concentrations in the plants. Statistical significance between species was assessed 162 using a Welch one-way ANOVA with a post hoc "Games-Howell" test (Games and Howell, 163 1976), which is part of the package 'userfriendlyscience' (Peters, 2019). The paired 164 comparisons were visualised using the algorithm for a letter-based comparison embedded in 165 the package 'multcompView' (Graves et al., 2019; Piepho, 2004). For statistical analyses, a pvalue < 0.05 was used as the cutoff for significance. Statistical difference is shown by a 166 167 sequence of letters within which the absence of matches indicates significant difference. All the 168 figures were created using a colour-blind-friendly colormap (Chang, 2013).

169

170 **3 Results**

171 **3.1** Properties, parameters and elemental composition of the Jas Roux soil

172 The soil pH of the study zone was 6.8 ± 0.3 (n=10) and conductivity was 58.4 ± 12.6 µS/cm. Analysis of the soil parameters identified the soil as a poorly carbonated (0.51%) sandy loam 173 174 containing 2.08% organic matter (Table SM 2). The cation exchange capacity was 6.1 175 mmol_c/100 g (Table SM 2). The mineralogy of the topsoil was dominated by barite, calcite, 176 jarosite, kaolinite, muscovite, quartz and K-feldspar (Figure SM 2). Table 2 lists the mean 177 total concentrations in the soil (n=10) and standard deviations (SD) for the 19 selected elements 178 compared to average soil concentrations, as well as the results of the CaCl₂ and the NH₄NO₃ 179 extractions. Except for Mg, Al, P, K, Mn, Fe, Ba and Tl, the phytoavailable (NH₄NO₃) fraction 180 was inferior to 1 mg/kg DW.

181

182 **3.2 Elemental uptake of plants**

183 3.2.1 Elemental concentrations in the roots and aerial plant parts

Figure 1 shows the range of elemental concentrations in the aerial parts and the roots of all sampled individuals compared to total concentrations in the soil. With the exception of Ca, P and Mg, the soil concentrations were higher than in the plant parts. The range of concentrations of macronutrients (Mg, P, S, K, Ca) was similar in the roots and the aerial parts, whereas the concentrations of trace elements were lower in the aerial parts than in the roots (**Figure 1**). The variability was very strong among the 53 plant samples compared to variability in the soil. 191 3.2.2 PCA based examination of correlations between critical metals and other elements in192 different plant parts

- Plant responses to high metal concentrations are species dependent and can even vary from one population to another (Gonneau et al., 2014; Pošćić et al., 2013; Rajabpoor et al., 2019). In this study, data analysis was performed at species and plant part levels to identify the different strategies. In the first step, a PCA was performed to capture the essence of the datasets.
- 197 The PCA biplot (Figure 2) merges the score plot of the individuals, represented by species 198 dependant symbols, and the loading plot of the variables (elemental concentrations) represented 199 by arrows, for aerial plant parts (a) and roots (c). In the aerial parts, the two first dimensions 200 (Dim1 and Dim2) explained 61.2% of total variance (Figure 2a). A first group of positively 201 correlated variables, composed of Mg, Ca, Mn, Ni, Zn, Y, Ba and Ce concentrations, formed 202 Dim1 (33.3% of total variance) (Figure SM 3). A second group of positively correlated 203 variables, including Al, V, Fe, As, Mo and Sb concentrations, contributed to Dim2 (27.9% of 204 total variance) (Figure SM 3). The score plot shows that all Sax individuals clustered together, 205 mainly contributed to Dim 1 and were associated with high concentrations of the first group of 206 elements. A cluster of *Hip* together with some single *Glo*, *Dia* and *Gal* individuals influenced 207 Dim2 (27.9% of total variance) and was characterised by higher concentrations of the second 208 group of elements.
- 209 In roots, 59.3% of total variance was explained by the first two dimensions (Figure 2c). Three
- 210 main groups of elements with strong positive intra-group correlations, were distinguished. The
- 211 first group was composed of Al, Ca, V, Mn, Ni, Zn, Y and Ce and contributed to Dim1 (42.8%
- of total variance) (Figure SM 3). The second group was composed of P, S and Tl and mainly
- 213 contributed to Dim2 (16.5% of total variance). The third group comprised Fe, As, Sb and Ba
- and influenced both Dim 1 and Dim2 (Figure SM 3).
- All seven individuals of *Sax* and one individual of *Gal* were associated with the first group while the second group of variables was connected by individuals of *Eup* and one individual of *Gal*.
- 218 Isolated individuals of *Dac* and *Eup* were associated with the third group of variables.
- 219 Overall, the PCA shows that specific groups of chemical elements were decisive in explaining
- the variance in roots and aerial plant parts driven by the individuals of Sax and Hip and by a
- single individual each of *Dia*, *Glo*, *Gal*.
- 222 Cluster analysis confirmed the tendencies revealed by the PCA. Three clusters of individuals
- 223 were distinguished in the aerial plant parts based on their similar metal absorption patterns
- (Figure 2b). The first group contained all seven individuals of *Sax* while the second group

- contained all three individuals of *Hip* as well as a single individual each of *Glo*, *Gal* and *Dia*.
 The third group contained all other individuals showing proximity among *Pil* and *Eup*individuals.
- 228 Only two clusters were distinguished in the roots (Figure 2d). Sax and Hip individuals were
- 229 part of the same cluster whereas all individuals of *Pil*, *Bis*, *Dia*, *Ver* and *Jun* formed the second
- 230 cluster. Individuals of *Hip*, *Dac*, *Gal* and *Eup* were part of both clusters.
- 231
- 232 3.2.3 Species related differences in uptake of critical metals
- According to the PERMANOVA, the factor "plant species" explained 65.65% of the variance
- in elemental concentrations while only 4.4% was explained by the factor "plant part" (Table
 SM 3).
- A Welch one-way ANOVA test combined with a Games-Howell post hoc test (Table 3, Table
 showed that three species, *Jun*, *Sax* and *Hip*, stood out from the others.
- 238 Jun generally had lower elemental concentrations in its aerial parts than the other species.
- 239 Indeed, Jun contained the lowest mean concentrations of Mg, Al, S, K, V, Fe, Ni, Cu, Zn, As,
- 240 Y, Sb, Ce and Tl. The concentrations of all these elements were significantly lower than in at
- 241 least one other species (Table 3). However, this tendency was not observed in the roots (Table 242 4).
- In turn, *Sax* had the highest concentrations of many of the elements analysed. In the aerial parts, significant differences in the concentrations of Mg, Ca, Mn, Ni, Y and Ce were found. The concentrations of Ca, Mn, Ni and Y were significantly higher than in all the other species. In the roots, the highly concentrated elements were very similar to those in the aerial parts. The concentrations of Al, Ca, Mn, Ni, As and Ce were significantly higher than in at least three of the other species.
- The third species with characteristic elemental concentrations was *Hip*. The elements accumulated in the aerial parts of *Hip* were Al, V, Fe, As, Sb and Mo. Differences in Al, V, Fe and As were significantly higher than in the other species. Differences in the concentrations of
- 252 Sb and Mo were not significant. It should be noted that average concentrations of Mo in *Hip*
- were about 5 times higher in roots (up to 26.9 mg/kg DW) and 30 times higher in the aerial
- 254 parts (up to 53.7 mg/kg DW), than in all the other species.
- The significance between groups (species) depends on both the variability within each group and between the groups. Species with highly variable concentrations are therefore not identified as significantly differing from one another. For example, *Bis* and *Gal* had locally high concentrations of Tl in their aerial parts, up to 541 mg/kg DW in *Bis* and up to 1091 mg/kg DW

in *Gal.* Despite the fact that no individual of another species exceeded 100 mg/kg DW, the differences were not significant. The concentrations of Ba in *Sax* are another example of nonsignificant differences due to high variability. The maximum concentration measured in the aerial parts was 5007 mg/kg, while in the other species 1445 mg/kg was the maximum concentration measured.

264

265 3.2.4 Identification of candidates for phytoremediation applications

The translocation-factor (TF) is a useful tool to identify the transfer strategy of a plant. The higher the factor, the larger the share of the element translocated to the aerial parts (Macnair, 2003).

269 Figure 3 is a heatmap of the mean TFs for the plant species and chemical elements investigated 270 with a colour spectrum ranging from pale yellow to dark red. The darker the colour of a box, 271 the higher the TF. The TF was low or highly variable in the majority of the twelve plant species. 272 Bis had the highest TF for many of the critical metals analysed (Mg, V, As, Y, Sb, Ce, Tl). In 273 particular, the translocation of Tl was very high (TF=19.4±17.2). A second species with a high 274 TF for Tl was Gal (TF=2.0±0.5). Dia also translocated many critical metals to the aerial parts 275 as shown by the high TF for Mg, V, As, Ce and Tl. Hip, on the other hand, had the highest TF 276 for Mo (TF=3.8±4.4). In Sax, the TF was low for most of the critical metals (As, Y, Ce, Sb, Tl) 277 and the only elements that were preferably stored in the leaves were Mg (TF=1.3±0.2) and Ca 278 $(TF=2.1\pm0.8)$. Jun only had low TFs.

279

280 **4 Discussion**

4.1 Jas Roux - A good site to study a plant community exposed to a cocktail of metals and metalloids in a natural ecosystem

The Jas Roux outcrop has fairly been studied regarding its exceptional mineral composition (Biagioni et al., 2015; Johan and Mantienne, 2000; Topa et al., 2014). However, the surrounding ecosystem, including the soil and the plants has never been studied before.

The concentrations of Zn (602 mg/kg), As (2382 mg/kg), Mo (19.3 mg/kg), Sb (778 mg/kg), Ba (27010 mg/kg) and Tl (242 mg/kg) were far above ordinary soil concentrations (**Table 2**). Soil concentrations in European topsoil of Zn are usually around 68 mg/kg, of As, 11 mg/kg, of Mo, 0.9 mg/kg, of Sb, 1 mg/kg, of Ba, 400 mg/kg, and of Tl, 0.8 mg/kg (**Table 2**) (Salminen et al., 2005). Except for Mo, all these elements have also been identified as elements associated with the primary minerals in the Jas Roux outcrops, namely barite, quartz and pyrite, as well as

292 Zn, Sb, As, Tl, Ag, Hg and Pb sulphides and sulphosalts (Boulliard et al., 2010). In terms of

- mobility and phytoavailability Zn, As, Mo, Sb, Ba and Tl remained < 0.1% of the total 293 294 concentrations and only Ba exceeded 1 mg/kg with 16.0 mg/kg after CaCl₂ extraction and 199 295 mg/kg after NH₄NO₃ extraction. These concentrations are comparable with those measured in 296 soils close to an active barite mining region in China with up to 111 mg/kg of phytoavailable 297 Ba (Lu et al., 2019). The mobility and phytoavailability of As, Sb and Zn were low compared 298 to those in contaminated soils with comparable total concentrations (Pueyo et al., 2004; Qasim 299 et al., 2015). This low mobility indicates the dominance of stable minerals in Jas Roux. Unlike 300 other sulphosalt sites, Jas Roux has not been disturbed by mining activities which generally 301 increase metal mobility (Resongles et al., 2014; Wei et al., 2018).
- Even though many of the characteristic elements of Jas Roux are known to be toxic for plants (As, Tl etc.), plant diversity is that of a classical alpine ecosystem (Landolt, 2015; Michalet et al., 2002). It is possible that the relatively low phytoavailability of the metals helps the plants to maintain their homeostasis in the short term (**Table 2**), but long-term exposure to low doses can result in a high real exposition, to which plants need to adapt.
- Plants use different strategies to adapt to high concentrations of toxic elements in the soil. Some plants are able to exclude the metals and others to accumulate the elements in specific compartments (Tremel-Schaub and Feix, 2005). Among the accumulating metallophytes, metals are generally accumulated in the roots, but some species promote translocation to the aerial plant parts (Krämer, 2010). Different statistical approaches have been used to identify the different adaption strategies and revealed characteristic uptake-patterns for the two plant species *Sax* and *Hip* (**Figure 2b**).
- 314

315 4.2 Saxifraga paniculata - A cation accumulating species with an affinity for 316 REEs and alkaline earth metals

317 In terms of elemental concentration in roots and aerial parts Sax stood out among the 12 species 318 analysed (Figure 2). In the aerial parts, the PCA showed the individuals of Sax to be associated 319 with Mg, Ca, Mn, Ni, Zn, Y, Ba and Ce (Figure 2). Among these elements, the concentrations 320 of Ca, Mn, Ni, Y were significantly higher than in all the other species (Figure 2, Table 3). It 321 is worth noting that the majority of elements accumulated in the aerial parts of Sax are divalent 322 cations (Ba, Ni, Mn, Ca, Zn and Mg) while the three elements with the highest concentrations 323 (Mg, Ca and Ba) are alkaline earth elements. This set of accumulated elements could be caused 324 by overexpression of nonselective cation channels or increased activity of a single type of metal 325 cation transporter, as their selectivity is known to be rather low (Demidchik and Maathuis, 326 2007; Thomas et al., 2005).

The genus *Saxifraga* comprises about 400 species and some of them have already been studied regarding metal tolerance (Gornall, 1987). To give one example, *Saxifraga oppositifolia*, *Saxifraga cespitosa* and *Saxifraga azoides* growing in Spitzbergen contained higher concentrations of Pb and Zn in their aerial parts than concentrations in the soil (Jóźwik and Magierski, 1991). Further, a Cu tolerant population of *Saxifraga stellaris* growing on polluted soils was identified in southern Germany, with concentrations up to 900 mg/kg DW (Adlassnig et al., 2013; Ryszka et al., 2019; Sissolak, 1984).

334 Sax belongs to a specific taxonomic section (sect. *Ligulatae*) characterised by strong guttation 335 (Gornall, 1987). Guttation is a process to regulate root pressure and nutrient flow by excreting 336 phloem and xylem sap through openings on the leaf surface and is observable in a wide range 337 of plant species, including many Saxifragaceae (Singh, 2014, 2013). The openings are called 338 hydathodes and, in contrast to stomata, remain permanently open (Singh, 2014). The guttation 339 process is thought to help plants expel excessive nutrient salts and harmful elements (Hossain 340 et al., 2017; Neumann et al., 1997; Singh, 2013; Stahl, 1919). In some species of Saxifraga, 341 including Sax, the repeated evaporation of guttation droplets leads to the formation of a calcite 342 crust on the leaf surface (Arnott and Pautard, 1970; Schmidt, 1930; Wightman et al., 2017). 343 These deposits appear to be the result of increased Ca uptake and might be a strategy to reduce 344 the cytotoxic effects of Ca while maintaining high flow rates (Nazrul Islam and Kawasaki, 345 2015). In the present study, the accumulation of Ca by Sax is reflected in the significantly higher 346 Ca-concentrations measured in the roots and the aerial parts (Table 3, Table 4) and the high 347 translocation factor (Figure 3).

This increased Ca uptake could also explain the higher concentrations of REEs (Y, Ce) in *Sax* compared to those in the other species, as REEs are probably absorbed through the same pathway as Ca because of their similar ionic radius (Carpenter et al., 2015; Kastori et al., 2010; Rengel, 1994; van der Ent et al., 2017). Ozaki et al. (1997) observed increased REEconcentrations combined with higher Ca and Ba uptake in several REE-accumulating fern species and suspected a possible link between these elements.

The accumulation of REEs is of particular interest for this study as REEs are crucial for modern technologies. Compared to the average concentration in the soil $(4.22\pm1.04 \text{ mg/kg})$ the roots of *Sax* individuals contained nearly twice as much Y $(7.37\pm3.24 \text{ mg/kg})$ (**Figure 4**). This raises the question of how *Sax* is able to absorb such a high ratio of Y (**Figure SM 4**). Y soil mineralogy in Jas Roux soil has not been studied so far. Y can be associated with several mineral phases, but one common soil mineral based on Y is Xenotime (YPO₄) (Ramos et al., 2016). Xenotime has low solubility and phosphate minerals are generally difficult for plants to 361 access (Darcy et al., 2018; Liu and Byrne, 1997; Tropper et al., 2011). However, there are some 362 exceptions - like white lupin or cassava - which can take up a fraction of soil P that is not 363 accessible to other plant species. In the case of cassava, the mechanisms allowing the increased 364 P-uptake are strong mycorrhization (Zaag et al., 1979). In white lupin, on the other hand, 365 increased root exudation is combined with a cluster root system (Braum and Helmke, 1995). In 366 the case of Sax, root exudates are more likely to play an important role, as the plant is able to 367 grow in very extreme habitats, in high altitudes and with little organic matter (Burga et al., 368 2004; Neuner et al., 1999).

369

4.3 *Hippocrepis comosa* - A oxyanion accumulating species with a particular affinity for Mo

The second plant species showing a characteristic pattern in elemental uptake was *Hip* (**Figure 2b**). In the aerial parts, all three individuals were positively correlated with the elements As, Fe, Sb, Mo, V and Al (**Figure 2**) with a particularly high TF for Mo (TF= 3.8 ± 4.4) (**Figure 3**). The fact that none of these elements accumulated in the aerial parts of *Sax* led to two easily distinguishable groups, separating the two species (**Figure 2b**).

Among the elements accumulated by *Hip* typical elemental correlations became visible. Due to their geochemical similarities, As and Sb (for example) are considered to be related elements (Herath et al., 2017). A positive correlation between these two elements has already been observed in plants and increased Sb uptake is presumed to be the result of damaged cell membrane integrity caused by As (Feng et al., 2013; Müller et al., 2013).

There is also a strong link between the uptake mechanisms of Mo and Fe (Bittner, 2014). Most Mo-enzymes also require Fe-containing redox groups and many studies reported increased Mo uptake in the presence of Fe and vice versa (Bittner, 2014; Kannan and Ramani, 1978; Zimmer and Mendel, 1999).

386 Three major hypotheses are proposed to explain the characteristic uptake pattern of *Hip*:

I) One hypothesis is a characteristic modification of the rhizosphere that would influence the
redox potential. This modification could locally increase phytoavailability of the accumulated
elements, as, except for Al, all are redox sensitive elements (As, Fe, Sb, Mo, and V).

390 II) The second hypothesis concerns the uptake pathway of the elements. It is known that As,

391 Sb, Mo, and V are usually absorbed as oxyanions by plants (Hossain et al., 2018; Kabata-

392 Pendias, 2011). Vanadate (VO_4^{3-}) and arsenate (AsO_4^{3-}) enter the plant through P transporters,

393 due to similarities in structure and charge (Chen et al., 2021; Meharg and Hartley-Whitaker,

394 2002, Han et al., 2016). Antimonate, on the other hand, is more likely to be absorbed through

- less selective anion transporters, such as those for Cl^- and NO_3^- , or an incompletely sealed or damaged Casparian strip (Feng et al., 2013; Tschan et al., 2009b, 2009a). However, little is known about Mo uptake by roots. Because of structural and behavioural similarities between molybdate (MoO₄²⁻) and sulphate (SO₄²⁻), it is plausible that sulphate transporters are involved (Bittner, 2014; Zimmer and Mendel, 1999).
- 400 Overexpression or increased activity of so many different transporters, responsible for the
 401 particular elemental concentrations found in *Hip*, is unlikely. Nevertheless, a very unselective
 402 anion transporter, able to transport phosphate, sulphate, nitrate and chlorate, could be involved.
 403 Such a transporter was identified in barley by Preuss et al. (2010), for example.
- 404 III) The third hypothesis is based on the N fixing property of *Hip*, as Mo and Fe are known to 405 play an essential role as cofactors during symbiotic N fixation (Hossain et al., 2018). For this 406 reason, legume plants like *Hip* have a higher demand for Mo than non-N fixing plants (Kabata-407 Pendias, 2011). It is plausible that *Hip* is able to promote Mo absorption in order to guarantee 408 N fixation under Mo deficient conditions. However, in Mo rich soils like Jas Roux (19.9±3.71 409 mg/kg), this property might lead to an increased uptake and thus explain the comparatively high 410 Mo concentrations in *Hip* (Figure 4). It is worth mentioning that V may also interfere during 411 N fixation by replacing Mo in the nitrogenase enzyme (Hernandez et al., 2009).
- 412 Individually, none of the three hypotheses explains all the elemental concentrations observed 413 in *Hip*. Nevertheless, a combination of the mechanisms could be a valid explanation. Al is the 414 only element whose accumulation by *Hip* remains difficult to explain, as this species is 415 generally found in Ca rich sites and is sensitive to high Al concentrations (Michalet et al., 2002). 416 Concentrations of Al > 1000 mg/kg, like those we measured in aerial parts of *Hip* (1336 mg/kg 417 DW), are rare in higher plants and are considered as Al accumulation (Kabata-Pendias, 2011). 418 The tolerance of toxic elements (Al, As) makes *Hip* an interesting candidate for remediation of 419 polluted soils as its ability to fix N thereby improving soil fertility could be an additional
- 420 advantage (Pajuelo et al 2008).
- 421

422 **4.4 Translocation factor reveals specific plant strategies**

The differentiation of *Hip* and *Sax* from the other species was mainly caused by high concentrations of a particular set of elements that made both species stand out in the PCA and the cluster analysis. However, this statistical approach failed to detect affinities for single elements, or adaption strategies associated with particularly low concentrations. The translocation factor in combination with the mean elemental concentrations, revealed further adaption strategies and species-related tendencies in elemental uptake (**Figure 3**). 429 For instance, Bis and Dia did not form a distinct group in the PCA or in the cluster analysis but 430 both had TFs > 1 for a large number of critical metals (Mg, V, Mn, As, Y, Sb, Ba, Ce, Tl). In 431 particular, the TF of *Bis* for Tl was remarkably high (TF=19.4±17.2) indicating Tl accumulation 432 (256±224 mg/kg) (Figure 4). While no explanation was found in the literature for the high TF 433 of *Dia*, *Bis* belongs to the Brassicaceae family, which includes many hyperaccumulating 434 species (Babula et al., 2008; van der Ent et al., 2013). Bis itself has already been mentioned as a Tl accumulating species (LaCoste et al., 1999; Pošćić et al., 2013). But Bis was not the only 435 436 Tl accumulating species, as *Gal*, which has never previously been associated with metalliferous 437 sites, also showed high concentrations of Tl in its aerial parts (254±410 mg/kg) and high 438 translocation (TF= 2.0 ± 0.5).

439 Another species that did not stand out in the statistical analysis was Jun. Nevertheless, Jun had 440 the lowest concentrations of the elements Mg, Al, S, K, V, Fe, Ni, Cu, Zn, As, Y, Sb, Ce and 441 Tl in its aerial parts (**Table 3**). This exclusion strategy seems to consist in limited translocation 442 to the aerial parts, as the TF was low for all elements (Figure 3). Given that Jun is the only 443 lignifying plant among the 12 species we analysed, its exclusion capacity might be due to 444 different plant-internal barriers or reduced flow rates due to its small leaf surface area. Even 445 though no studies have demonstrated metal exclusion in Juniperus species, metal concentrations 446 in other conifers (Pinales) have already been reported. For example, Mingorance et al. (2007) 447 found *Pinus pinea* growing in an industrial area was able to exclude Al, Ba, Cr, Fe, and Pb. In 448 light of the results of our study, Jun could be an interesting candidate for phytostabilisation 449 thanks to its high resistance to metal contamination, its widespread occurrence and its 450 ruggedness (Saladin, 2015).

451

452 **5** Conclusions

453 Among 12 plant species exposed to a cocktail of metals and metalloids in an entirely natural 454 environment, two species stood out: Hippocrepis comosa and Saxifraga paniculata, each of 455 which accumulated an individual set of elements. Saxifraga paniculata preferably absorbed 456 divalent cations, but also the REEs Y and Ce. This uptake pattern could be explained by the Ca 457 accumulation property of Saxifraga paniculata. The affinity to Ca might cause an increased 458 absorption of other elements with a similar ionic radius like Y and Ce, or similar ionic charge 459 like Mn and Ba. *Hippocrepis comosa*, on the other hand, preferably absorbed oxyanions, 460 perhaps due to overexpression or increased activity of a non-selective anion transporter. 461 Furthermore, *Hippocrepis comosa* is an interesting candidate for phytoremediation of polluted 462 soils, as its ability to fix N₂ improves soil fertility and could thus facilitate the establishment of 463 other species.

464 Different tolerance strategies were identified among the studied plant species. Biscutella 465 *laevigata* and *Dianthus sylvestris* favoured translocation of multiple critical metals to their 466 aerial parts while, Juniperus communis avoided translocation and used an exclusion strategy. 467 Particularly high concentrations of Tl were found in Biscutella laevigata and Galium 468 corrudifolium. While hyperaccumulation of Tl has already been reported in Biscutella 469 laevigata, it is a new discovery in Galium corrudifolium. The exclusion capacity of Juniperus 470 communis subsp. nana might represent a potential for phytostabilisation applications. Erosion 471 of a polluted soil could be prevented in the long term with no risk of transfer into the food chain 472 because Juniperus communis subsp. nana prevents elemental translocation and the shrub is not 473 consumed by grazing animals.

These results advance our understanding of soil-to-plant transfer of poorly studied but increasingly used critical metals and point to the potential of several plant species for future phytoremediation techniques.

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482

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504

490 Abbreviations:

491 ANOVA..... Analysis Of Variance 492 a.s.l. Above Sea Level 493 DW Dry Weight 494 ICP-MS Inductively Coupled Plasma Mass Spectrometry 495 ICP-OES Inductively Coupled Plasma Optical Emission Spectroscopy 496 PCA Principal Component Analysis 497 PERMANOVA..... Permutational Multivariate Analysis Of Variance 498 PGMs Platinum Group Metals 499 REEs Rare Earth Elements 500 SD Standard Deviation 501 subsp. Subspecies 502 TF Translocation Factor 503 XRD X-Ray Diffraction

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1 **1 Introduction**

2 All over the world, emerging technologies require an increasingly diverse spectrum of chemical 3 elements with highly specific (photo-optical, catalytic, magnetic, etc.) properties (Celis et al., 4 2020). Metals that have practically never been used before are now indispensable for many 5 industrial sectors and are thus the subject of soaring global demand. Due to their economic 6 importance and their supply risk, many of these metals are now classified as "critical" for high 7 tech applications (European Commission, 2020), examples being Sb in lead-acid batteries and 8 semiconductors, As in cell phones and solar cells and rare earth elements (REEs) in sensors, 9 displays and electric motors (Fortier et al., 2018; Mudd et al., 2018).

10 Their increasing use also involves increased risk of their dispersion in the environment during

11 their extraction, use, and disposal. HoweverDespite their increased use, the biogeochemical 12 cycles of many of these critical metals are not yet fully understood and predicting either their 13 environmental behaviour or their ecotoxicity is consequently impossible (Espejo et al., 2018). 14 This knowledge, however, Aim of this study is essential to choose analyse the right remediation 15 techniquesoil-to-plant transfer of some of these metals and to shed light on their uptake 16 pathways. Our findings might bare valuable information for polluted sites. In recent 17 years, phytoremediation and promote plant-based remediation techniques (phytoremediation) 18 have become a feasible and environmentally friendly solution to treat of metal-contaminated soils (Geras'kin et al., 2011; Tangahu et al., 2011; van der Ent et al., 2017).sites. 19 20 Phytoremediation techniques are relying on the capacity of certain plant species to limit the 21 uptake of the toxic elements or to accumulate them in designated compartments (e.g. 22 phytostabilisation, phytoextraction...).) (Favas et al., 2014). Exclusion and accumulation are the two main strategies of plants to maintain homeostasis in metal-rich habitats. Both involve 23 24 a series of adaptions such as modified membrane transport, chelation and complexation, or 25 rhizosphere interactions for example (Furini, 2012).

The ability of plants to accumulate metals in their aerial parts has received considerable attention and many combinations of specific plant species tolerating or accumulating a particular set of elements have been discovered (Baker et al., 2010; Pollard et al., 2014). However, most studies have focused on "traditional" metals such as Zn, Ni, Cd and Pb, whereas the uptake pathways and metabolic impact of many elements, particularly emerging critical metals, still raise many questions (Macnair, 2003; van der Ent et al., 2013).

Elemental uptake is influenced by chemical, physical, and biological factors. Plant species, root
system, root exudates, edaphic interactions, and transport mechanisms all play important roles
(Tangahu et al., 2011; Trakal et al., 2015; Yadav, 2010). Despite highly specialised proteins,

including transporters, pumps and channels, elemental selectivity remains challenging for plants and each uptake mechanism is likely to let pass a range of ions due to chemical similarities between the different ions (Tangahu et al., 2011). The huge number of factors and complex interactions are impossible to reproduce in the laboratory, thus, to understand the uptake of poorly studied elements, such as critical metals, performing in-situ studies in a natural environment is advantageous.

41 The geological site of Jas Roux, located in the Ecrins National Park (Hautes-Alpes, France), 42 was chosen for the study. The distinctive reddish colour of the Jas Roux outcrops ('Roux' 43 means red) originates from a sulphosalt-forming hydrothermal mineralisation (Johan and 44 Mantienne, 2000). These sulphosalts are one of the reasons for the characteristic metal 45 concentrations of Jas Roux as they are rich in elements with a strong affinity for sulphurS, such 46 as lead, antimony, copper, thalliumPb, Sn, Cu, Tl and arsenicAs (Dittrich et al., 2009; Pierrot 47 and Roger, 2018). Many of the elements found in Jas Roux are used in high-tech applications 48 and are considered to be critical for leading technologies (Dittrich et al., 2009; Graedel et al., 49 2015). This site is thus a good proxy of chronic contamination by a cocktail of metals and 50 metalloids. As the site has never been disturbed by any human activities and is located in the 51 heart of the Ecrins National Parc, it offers ideal conditions to study an established ecosystem 52 with a plant community that has not undergone human selection. The plant community can be 53 seen as natural response to the local mineralogy and may contain candidates for 54 phytoremediation.

- 55 To identify such plant candidates and to study the environmental behaviour of critical metals,
- 56 we were guided by the following three questions:
- 57 To what extent are critical metals absorbed by plants?
- 58 Are there any species-related differences in elemental uptake?
- 59 Do correlations exist between critical metals and other elements in different plant parts?
- 60 The approach of this study was to conduct a large-scale screening of 12 plant species to identify
- 61 individual uptake patterns using chemical analysis, soil element extraction and statistical tools.
- 62 The large number of species analysed provided insights into the actual elemental
- 63 phytoavailability of a wide range of critical metals.

64 2 Material and Methods

65 2.1 Sampling Site

The geological site of Jas Roux, located in the Ecrins National Park in the district of La 66 67 Chapelle-en-Valgaudemar (Hautes-Alpes, France), is situated at 2170 m a.s.l. (44°48' 45" N, 6° 19' 18" E; WGS84) on the right-hand side of the Chabournéou stream (Johan and Mantienne, 68 69 2000). Jas Roux is a hydrothermal stratiform deposit in dolomite limestone and has been 70 characterised as a sulphosalt site (Boulliard et al. 2010). The hydrothermal system was possibly 71 generated by volcanic activity involving relatively low temperatures (around 100 °C) and high 72 pressure (Johan and Mantienne, 2000). The particular thalliumTl-rich composition of Jas Roux 73 has attracted many mineralogists and geologists, who have identified Zn, Sb, As, Tl, Ag, Hg, 74 Pb sulphides and sulphosalts including many new sulphosalt mineral species (Boulliard et al., 75 2010; Guillemin et al., 1970; Mantienne, 1974; Topa et al. 2017, 2014). 76 The study zone is located below one of the reddish coloured outcrops identified by Mantienne 77 (Figure SM 1) (Mantienne, 1974). The metal-rich mineralisation is best developed in this 78 outcrop and forms a 3.8 m thick barite-rich zone (Johan and Mantienne, 2000). The study zone 79 measured 20 m x 5 m with a slope of approximately 45° (100%) (Figure SM 1). 80 Soil and plant samples were collected during two field trips, one in September 2017 and one in 81 June 2018. The steep slope, variable soil depth and the presence of large rocks impeded a 82 systematic sampling approach. Nevertheless, the samples were collected as evenly distributed 83 as possible. The soil samples were collected from 10-evenly distributed sampling points in the 84 study zone and consisted of 1 kg of topsoil each (< 15 cm). Among the 65 plant species 85 identified in the study zone, 12 were selected, each belonging to a different plant family (Table 1).taking into account occurrence, spatial distribution and family affiliation (Table 1). For 86 87 improved readability we attributed acronyms to each of the 12 species throughout the paper 88 (Table 1). Depending on their abundance, between 2 and 7 replicates of each species were

sampled, giving a total of 55 individuals-<u>(Table 1)</u>. All the samples were stored at 4 °C in the
dark.

91

92 **2.2 Sample preparation**

Roots and aerial parts were separated at the collar and cleaned with 18.2 MΩ·cm ultrapure water. The plant roots were additionally brushed and vortexed in ultrapure water to remove adhering soil particles. The leaves of *Saxifraga paniculataSax* were also brushed as they were partially covered with a calcite crust. Plant samples were dried at 40 °C for 48 h. Soil samples were dried at 40 °C for 72 h and sieved at 2mm before analysis.

98 All the soil and plant samples were ground at 20hz in a Retsch® Mixer Mill MM 400 for 2 min 99 to obtain a fine powder. Each sample (100mg) was solubilised in acid using a Milestone 100 UltraWAVE® microwave to complete mineralisation. The acids used for the digestion of the 101 plant samples were 4mL of HNO₃ and 2mL of H₂O₂. For the soil samples, 3mL of HNO₃, 3mL 102 of HCl and 0.5 mL of HF were used to measure the total metal concentrations. Soil pH and 103 conductivity were determined in a suspension of 1g of sieved soil (< 2mm) and 5mL of 104 ultrapure water according to the French standard X31-117 (NF X31-117, 1994). After 5 min of 105 agitation at 80 rpm and 2 h of rest, measurements were taken using a Hanna HI-2550 bench 106 meter. To estimate the mobile and phytoavailable elemental fractions in the soil, CaCl₂ 107 (0.001 mol/L) and NH₄NO₃ (1mol/L) extractions have been conducted, according to standards 108 ISO 21268-2 and ISO 19730 (ISO 19730:2008, 2009; NF X31-607-2, 2004). For the CaCl₂ 109 extraction, a liquid-solid ratio of 10:1 (20mL of ultrapure water and 2g of soil) was applied. 110 After 24 h of agitation at 10 rpm, the samples were centrifuged at 2500xG for 20 min, filtered at 45µm0.45 µm and analysed in ICP-MS. For the NH₄NO₃ extraction, 4g of soil were mixed 111 112 with 10mL of ultrapure water, agitated for 2 h at 30 rpm, centrifuged at 1000xG for 15 min,

- filtered at $0.45 \,\mu\text{m}$ and analysed in ICP-MS.
- 114 Soil mineralogy was analysed by X-ray diffraction using the X'Pert Pro MPD (XRD) equipped
- 115 with a Cobalt x-ray tube (= 1.79A) running at 40 kV and 40 mA.

116 A mixture of three bulk samples of topsoil was sent to an outside laboratory (CIRAD – US49 117 Laboratoire d'analyse de Montpellier, France) for the determination of soil parameters. The soil 118 type was determined using 5 granulometric fractions according to the French standard X31-107 119 (*NF X31-107*, 2003), organic matter by combustion analysis (Dumas), labile <u>phosphorousP</u> 120 content using a colorimetric method (Olsen) and cation exchange capacity by quantifying the 121 dosing adsorption agent [Co(NH₃)₆]Cl₃ with inductively coupled plasma atomic emission 122 spectroscopy according to the French standard X31-130 (*NF X31-130*, 1999).

123

124 **2.3 Chemical analyses**

The samples were analysed by inductively coupled plasma mass spectrometry (ICP-MS, Perkin Elmer NexIon 300X). For the elements Mg, Al, P, K, Ca and S, the results of inductively coupled plasma optical emission spectroscopy (ICP-OES) were occasionally used to reach higher accuracy (Perkin Elmer 4300 DV). The calibration curves were established using multisolution standards CCS4, CCS5, CCS6 from Inorganic Ventures (New Jersey USA) diluted in 2% HNO₃ solution. The internal standards were Rh103 for ICP-MS and Y for ICP-OES. Oak leaves (V464) provided by the French National Institute for Agriculture Food and Environment (INRAE) were used as the external reference for plant samples. For the soil, a rendzina sample
(ISA 979) provided by Wageningen Evaluating Programs for Analytical Laboratories
(WEPAL) was used. EnviroMAT drinking water EP-L-3 and subterranean water ES-H-2 (SCP)

- 135 Sciences) were used as internal references and for quality control.
- I36 ICP analyses aimed to quantify critical metalselements (As, V, Y, Sb, Ba, Ce, Tl) and major
- 137 soil elements and nutrients (Mg, Al, P, S, K, Ca, Mn, Fe, Ni, Cu, Zn, Mo) present in Jas Roux.
- 138 <u>The critical elements were chosen based on the study on the EU's list of critical raw materials</u>
- and a review of critical minerals and will be called critical metals throughout the manuscript
- 140 (European Commission, 2020; Hayes and McCullough, 2018). The detection limit for each
- 141 element was determined using handling blanks analysed at each ICP analysis. (Table SM 1).
- 142 The coefficients of variances for all certified elements were calculated for the internal and
- 143 external standards (**Table SM 1**).

The translocation factor (TF) was calculated to identify the transfer strategies of the different plant species. The TF is obtained by dividing the elemental concentrations in the aerial parts by the concentrations in the roots (Radziemska et al., 2017). The TF was calculated for each plant sample and averaged according to the respective plant species. A heatmap with adjusted colormap breaks was used to illustrate the TF.

149

150 **2.4 Statistical analysis**

151 All statistical analyses were performed with the open-source software R studio. The complete 152 dataset for the plants contains 110 observations (aerial parts + roots) and 19 variables (chemical 153 elements). In the first step, a principal component analysis (PCA) was conducted on the aerial 154 parts and the roots separately to capture the essence of the datasets and to identify the features 155 that explain most of the variation in chemical composition. For this purpose, the packages 156 'FactomineR' and 'factoextra' were used (Kassambara, 2017a; Kassambara and Mundt, 2020; 157 Lê et al., 2008). Missing values were implemented using the package 'missMDA', which 158 reduced their influence on the PCA results to a minimum (Josse and Husson, 2016).

In the second step, an unsupervised clustering analysis was performed. The clustering tendency of the two datasets (aerial parts + roots) was tested using the Hopkins test from the package 'clustertend' before performing portioning clustering (K-means) (YiLan and RuTong, 2015). K-means is a clustering algorithm that returns the natural grouping of data points based on their similarity (elemental concentrations), not on their identity (plant species). The determination of the number of clusters as well as the clustering itself and the visualisation were performed using the package 'factoextra' (Kassambara, 2017b; Kassambara and Mundt, 2020). Next, a

PERMANOVA was performed on the complete dataset, using the package 'vegan' (Oksanen 166 167 et al., 2020) to examine to what degree the factors "plant species" and "plant parts" influenced 168 the elemental concentrations in the plants. Statistical significance between species was assessed 169 using a Welch one-way ANOVA with a post hoc "Games-Howell" test (Games and Howell, 170 1976), which is part of the package 'userfriendlyscience' (Peters, 2019). The paired 171 comparisons were visualised using the algorithm for a letter-based comparison embedded in 172 the package 'multcompView' (Graves et al., 2019; Piepho, 2004). For statistical analyses, a p-173 value < 0.05 was used as the cutoff for significance. Statistical difference is shown by a 174 sequence of letters within which the absence of matches indicates significant difference. All the 175 figures were created using a colour-blind-friendly colormap (Chang, 2013).

176

177 **3 Results**

178 **3.1** Properties, parameters and elemental composition of the Jas Roux soil

179 The soil pH of the study zone was 6.8 ± 0.3 (n=10) and conductivity was 58.4 ± 12.6 µS/cm. 180 Analysis of the soil parameters identified the soil as a poorly carbonated (0.51%) sandy loam 181 containing 2.08% organic matter (Table SM 2). The cation exchange capacity was 6.1 182 mmol_c/100 g (Table SM 2). The mineralogy of the topsoil was dominated by barite, calcite, 183 jarosite, kaolinite, muscovite, quartz and K-feldspar (Figure SM 2). Table 2 lists the mean 184 total concentrations in the soil (n=10) and standard deviations (SD) for the 19 selected elements 185 compared to average soil concentrations, as well as the results of the CaCl₂ and the NH₄NO₃ 186 extractions. Except for Mg, Al, P, K, Mn, Fe, Ba and Tl, the phytoavailable (NH₄NO₃) fraction 187 was inferior to 1 mg/kg DW.

188

189 **3.2 Elemental uptake of plants**

190 3.2.1 Elemental concentrations in the roots and aerial plant parts

Figure 1 shows the range of elemental concentrations in the aerial parts and the roots of all sampled individuals compared to total concentrations in the soil. With the exception of Ca, P and Mg, the soil concentrations were higher than in the plant parts. The range of concentrations of macronutrients (Mg, P, S, K, Ca) was similar in the roots and the aerial parts, whereas the concentrations of trace elements were lower in the aerial parts than in the roots (**Figure 1**). The variability was very strong among the 53 plant samples compared to variability in the soil.

- 198 3.2.2 PCA based examination of correlations between critical metals and other elements in199 different plant parts
- Plant responses to high metal concentrations are species dependent and can even vary from one
 population to another (Gonneau et al., 2014; Pošćić et al., 2013; Rajabpoor et al., 2019). In this
 study, data analysis was performed at species and plant part levels to identify the different
 strategies. In the first step, a PCA was performed to capture the essence of the datasets.
- 204 The PCA biplot (Figure 2) merges the score plot of the individuals, represented by species 205 dependant symbols, and the loading plot of the variables (elemental concentrations) represented 206 by arrows, for aerial plant parts (a) and roots (c). In the aerial parts, the two first dimensions 207 (Dim1 and Dim2) explained 61.2% of total variance (Figure 2a). A first group of positively 208 correlated variables, composed of Mg, Ca, Mn, Ni, Zn, Y, Ba and Ce concentrations, formed 209 Dim1 (33.3% of total variance) (Figure SM 3). A second group of positively correlated 210 variables, including Al, V, Fe, As, Mo and Sb concentrations, contributed to Dim2 (27.9% of 211 total variance) (Figure SM 3). The score plot shows that all Sax individuals clustered together, 212 mainly contributed to Dim 1 and were associated with high concentrations of the first group of 213 elements. A cluster of *Hip* together with some single *Glo*, *Dia* and *Gal* individuals influenced 214 Dim2 (27.9% of total variance) and was characterised by higher concentrations of the second 215 group of elements.
- 216 In roots, 59.3% of total variance was explained by the first two dimensions (**Figure 2c**). Three
- 217 main groups of elements with strong positive intra-group correlations, were distinguished. The
- 218 first group was composed of Al, Ca, V, Mn, Ni, Zn, Y and Ce and contributed to Dim1 (42.8%
- of total variance) (Figure SM 3). The second group was composed of P, S and Tl and mainly
- 220 contributed to Dim2 (16.5% of total variance). The third group comprised Fe, As, Sb and Ba
- and influenced both Dim 1 and Dim2 (Figure SM 3).
- All seven individuals of *Sax* and one individual of *Gal* were associated with the first group while the second group of variables was connected by individuals of *Eup* and one individual of *Gal*.
- Isolated individuals of *Dac* and *Eup* were associated with the third group of variables.
- 226 Overall, the PCA shows that specific groups of chemical elements were decisive in explaining
- the variance in roots and aerial plant parts driven by the individuals of Sax and Hip and by a
- single individual each of *Dia*, *Glo*, *Gal*.
- 229 Cluster analysis confirmed the tendencies revealed by the PCA. Three clusters of individuals
- 230 were distinguished in the aerial plant parts based on their similar metal absorption patterns
- 231 (Figure 2b). The first group contained all seven individuals of *Sax* while the second group

- contained all three individuals of *Hip* as well as a single individual each of *Glo*, *Gal* and *Dia*.
 The third group contained all other individuals showing proximity among *Pil* and *Eup*individuals.
- 235 Only two clusters were distinguished in the roots (Figure 2d). Sax and Hip individuals were
- part of the same cluster whereas all individuals of *Pil*, *Bis*, *Dia*, *Ver* and *Jun* formed the second
- 237 cluster. Individuals of *Hip*, *Dac*, *Gal* and *Eup* were part of both clusters.
- 238
- 239 3.2.3 Species related differences in uptake of critical metals
- According to the PERMANOVA, the factor "plant species" explained 65.65% of the variance
- in elemental concentrations while only 4.4% was explained by the factor "plant part" (Table
 SM 3).
- A Welch one-way ANOVA test combined with a Games-Howell post hoc test (**Table 3, Table**
- 4) showed that three species, *Jun*, *Sax* and *Hip*, stood out from the others.
- 245 *Juniperus communis (Jun)* generally had lower elemental concentrations in its aerial parts than
- the other species. Indeed, Jun contained the lowest mean concentrations of Mg, Al, S, K, V, Fe,
- 247 Ni, Cu, Zn, As, Y, Sb, Ce and Tl. The concentrations of all these elements were significantly
- lower than in at least one other species (Table 3). However, this tendency was not observed inthe roots (Table 4).
- In turn, *Saxifraga paniculata* (*Sax*) had the highest concentrations of many of the elements analysed. In the aerial parts, significant differences in the concentrations of Mg, Ca, Mn, Ni, Y and Ce were found. The concentrations of Ca, Mn, Ni and Y were significantly higher than in all the other species. In the roots, the highly concentrated elements were very similar to those in the aerial parts. The concentrations of Al, Ca, Mn, Ni, As and Ce were significantly higher than in at least three of the other species.
- 256 The third species with characteristic elemental concentrations was *Hippocrepis comosa* (*Hip*).
- 257 The elements accumulated in the aerial parts of *Hip* were Al, V, Fe, As, Sb and Mo. Differences
- in Al, V, Fe and As were significantly higher than in the other species. Differences in the concentrations of Sb and Mo were not significant. It should be noted that average concentrations
- of Mo in *Hip* were about 5 times higher in roots (up to 26.9 mg/kg DW) and 30 times higher in
- the aerial parts (up to 53.7 mg/kg DW), than in all the other species.
- The significance between groups (species) depends on both the variability within each group and between the groups. Species with highly variable concentrations are therefore not identified as significantly differing from one another. For example, *Bis* and *Gal* had locally high concentrations of Tl in their aerial parts, up to 541 mg/kg DW in *Bis* and up to 1091 mg/kg DW

in *Gal.* Despite the fact that no individual of another species exceeded 100 mg/kg DW, the differences were not significant. The concentrations of Ba in *Sax* are another example of nonsignificant differences due to high variability. The maximum concentration measured in the aerial parts was 5007 mg/kg, while in the other species 1445 mg/kg was the maximum concentration measured.

271

272 3.2.4 Identification of candidates for phytoremediation applications

The translocation-factor (TF) is a useful tool to identify the transfer strategy of a plant. The higher the factor, the larger the share of the element translocated to the aerial parts (Macnair, 2003).

276 Figure 3 is a heatmap of the mean TFs for the plant species and chemical elements investigated 277 with a colour spectrum ranging from pale yellow to dark red. The darker the colour of a box, 278 the higher the TF. The TF was low or highly variable in the majority of the twelve plant species. 279 Bis had the highest TF for many of the critical metals analysed (Mg, V, As, Y, Sb, Ce, Tl). In 280 particular, the translocation of Tl was very high (TF=19.4±17.2). A second species with a high 281 TF for Tl was Gal (TF=2.0±0.5). Dia also translocated many critical metals to the aerial parts 282 as shown by the high TF for Mg, V, As, Ce and Tl. Hip, on the other hand, had the highest TF 283 for Mo (TF=3.8±4.4). In Sax, the TF was low for most of the critical metals (As, Y, Ce, Sb, Tl) 284 and the only elements that were preferably stored in the leaves were Mg (TF=1.3±0.2) and Ca 285 $(TF=2.1\pm0.8)$. Jun only had low TFs.

286

287 **4 Discussion**

4.1 Jas Roux - A good site to study a plant community exposed to a cocktail of metals and metalloids in a natural ecosystem

The Jas Roux outcrop has fairly been studied regarding its exceptional mineral composition (Biagioni et al., 2015; Johan and Mantienne, 2000; Topa et al., 2014). However, the surrounding ecosystem, including the soil and the plants has never been studied before.

The concentrations of Zn (602 mg/kg), As (2382 mg/kg), Mo (19.3 mg/kg), Sb (778 mg/kg), Ba (27010 mg/kg) and Tl (242 mg/kg) were far above ordinary soil concentrations (**Table 2**). Soil concentrations in European topsoil of Zn are usually around 68 mg/kg, of As, 11 mg/kg, of Mo, 0.9 mg/kg, of Sb, 1 mg/kg, of Ba, 400 mg/kg, and of Tl, 0.8 mg/kg (**Table 2**) (Salminen et al., 2005). Except for Mo, all these elements have also been identified as elements associated with the primary minerals in the Jas Roux outcrops, namely barite, quartz and pyrite, as well as

299 Zn, Sb, As, Tl, Ag, Hg and Pb sulphides and sulphosalts (Boulliard et al., 2010). In terms of

mobility and phytoavailability Zn, As, Mo, Sb, Ba and Tl remained < 0.1% of the total 300 301 concentrations and only Ba exceeded 1 mg/kg with 16.0 mg/kg after CaCl₂ extraction and 199 302 mg/kg after NH₄NO₃ extraction. These concentrations are comparable with those measured in 303 soils close to an active barite mining region in China with up to 111 mg/kg of phytoavailable 304 Ba (Lu et al., 2019). The mobility and phytoavailability of As, Sb and Zn were low compared 305 to those in contaminated soils with comparable total concentrations (Pueyo et al., 2004; Qasim 306 et al., 2015). This low mobility is due toindicates the fact that dominance of stable minerals in Jas Roux-is a natural ecosystem that. Unlike other sulphosalt sites, Jas Roux has not been 307 308 disturbed by mining activities that which generally increase metal mobility- (Resongles et al., 309 2014; Wei et al., 2018).

310 The particularity of Jas Roux is its location in the heart of the Ecrins National Parc and the fact

311 that, unlike other sulphosalt sites, it has never been disturbed by mining activities (Ilmen et al.,

312 2016; Klemm et al., 2004). The plant community in Jas Roux has had millennia to adapt to an

- 313 environment with no human intervention.
- Even though many of the characteristic elements of Jas Roux are known to be toxic for plants (As, Tl etc.), plant diversity is that of a classical alpine ecosystem (Landolt, 2015; Michalet et al., 2002). It is possible that the relatively low phytoavailability of the metals helps the plants to maintain their homeostasis in the short term (**Table 2**), but long-term exposure to low doses can result in a high real exposition, to which plants need to adapt.
- Plants use different strategies to adapt to high concentrations of toxic elements in the soil. Some plants are able to exclude the metals and others to accumulate the elements in specific compartments (Tremel-Schaub and Feix, 2005). Among the accumulating metallophytes, metals are generally accumulated in the roots, but some species promote translocation to the aerial plant parts (Krämer, 2010). Different statistical approaches have been used to identify the different adaption strategies and revealed characteristic uptake-patterns for the two plant species *Saxifraga paniculata* and *Hippocrepis comosaSax* and *Hip* (Figure 2b).
- 326

327 4.2 Saxifraga paniculata - A cation accumulating species with an affinity for 328 REEs and alkaline earth metals

In terms of elemental concentration in roots and aerial parts *Saxifraga paniculataSax* stood out among the 12 species analysed (**Figure 2**). In the aerial parts, the PCA showed the individuals of *Sax* to be associated with Mg, Ca, Mn, Ni, Zn, Y, Ba and Ce (**Figure 2**). Among these elements, the concentrations of Ca, Mn, Ni, Y were significantly higher than in all the other species (**Figure 2, Table 3**). It is worth noting that the majority of elements accumulated in the aerial parts of *Saxifraga paniculataSax* are divalent cations (Ba, Ni, Mn, Ca, Zn and Mg) while
the three elements with the highest concentrations (Mg, Ca and Ba) are alkaline earth elements.
This set of accumulated elements could be caused by overexpression of nonselective cation
channels or increased activity of a single type of metal cation transporter, as their selectivity is
known to be rather low (Demidchik and Maathuis, 2007; Thomas et al., 2005).

The genus *Saxifraga* comprises about 400 species and some of them have already been studied regarding metal tolerance (Gornall, 1987). To give one example, *Saxifraga oppositifolia*, *Saxifraga cespitosa* and *Saxifraga azoides* growing in Spitzbergen contained higher concentrations of Pb and Zn in their aerial parts than concentrations in the soil (Jóźwik and Magierski, 1991). Further, a Cu tolerant population of *Saxifraga stellaris* growing on polluted soils was identified in southern Germany, with concentrations up to 900 mg/kg DW (Adlassnig et al., 2013; Ryszka et al., 2019; Sissolak, 1984).

346 Saxifraga paniculata, the species analysed in this study, Sax belongs to a specific taxonomic 347 section (sect. Ligulatae) characterised by strong guttation (Gornall, 1987). Guttation is a 348 process to regulate root pressure and nutrient flow by excreting phloem and xylem sap through 349 openings on the leaf surface and is observable in a wide range of plant species, including many 350 Saxifragaceae (Singh, 2014, 2013). The openings are called hydathodes and, in contrast to 351 stomata, remain permanently open (Singh, 2014). The guttation process is thought to help plants 352 expel excessive nutrient salts and harmful elements (Hossain et al., 2017; Neumann et al., 1997; 353 Singh, 2013; Stahl, 1919). In some species of Saxifraga, including Saxifraga paniculataSax, 354 the repeated evaporation of guttation droplets leads to the formation of a calcite crust-of calcium 355 carbonate on the leaf surface (Arnott and Pautard, 1970; Schmidt, 1930; Wightman et al., 2017). 356 These deposits appear to be the result of increased Ca uptake and might be a strategy to reduce 357 the cytotoxic effects of Ca while maintaining high flow rates (Nazrul Islam and Kawasaki, 358 2015). In the present study, the accumulation of Ca by Saxifraga paniculataSax is reflected in 359 the significantly higher Ca-concentrations measured in the roots and the aerial parts (Table 3, 360 Table 4) and the high translocation factor (Figure 3).

This increased Ca uptake could also explain the higher concentrations of REEs (Y, Ce) in *Saxifraga paniculataSax* compared to those in the other species, as REEs are probably absorbed through the same pathway as Ca because of their similar ionic radius (Carpenter et al., 2015; Kastori et al., 2010; Rengel, 1994; van der Ent et al., 2017). Ozaki et al. (1997) observed increased REE-concentrations combined with higher Ca and Ba uptake in several REEaccumulating fern species and suspected a possible link between these elements. 367 The accumulation of REEs is of particular interest for this study as REEs are crucial for modern 368 technologies. Compared to the average concentration in the soil $(4.22\pm1.04 \text{ mg/kg})$ the roots of 369 Saxifraga paniculataSax individuals contained nearly twice as much Y (7.37±3.24 mg/kg) 370 (Figure 4). This raises the question of how *Saxifraga paniculataSax* is able to absorb such a 371 high ratio of Y (Figure SM 4). The most Y soil mineralogy in Jas Roux soil has not been studied 372 so far. Y can be associated with several mineral phases, but one common soil mineral based on 373 Y is Xenotime (YPO₄) which (Ramos et al., 2016). Xenotime has low solubility (Liu and Byrne, 1997; Ramos et al., 2016; Tropper et al., 2011). However, and phosphate chemistry in iron-rich 374 375 environments like Jas Roux is highly complex and the formation of mixed amorphous phases is possible. Phosphate minerals are generally difficult for plants to access (Darcy et al., 2018).; 376 377 Liu and Byrne, 1997; Tropper et al., 2011). However, there are some exceptions - like white 378 lupin or cassava - which can take up a fraction of soil P that is not accessible to other plant 379 species. In the case of cassava, the mechanisms allowing the increased P-uptake are strong 380 mycorrhization (Zaag et al., 1979). In white lupin, on the other hand, increased root exudation 381 is combined with a cluster root system (Braum and Helmke, 1995). In the case of Saxifraga 382 *paniculataSax*, root exudates are more likely to play an important role, as the plant is able to 383 grow in very extreme habitats, in high altitudes and with little organic matter (Burga et al., 384 2004; Neuner et al., 1999).

385

4.3 *Hippocrepis comosa* - A oxyanion accumulating species with a particular affinity for Mo

The second plant species showing a characteristic pattern in elemental uptake was *Hippocrepis comosa<u>Hip</u>* (Figure 2b). In the aerial parts, all three individuals were positively correlated with the elements As, Fe, Sb, Mo, V and Al (Figure 2) with a particularly high TF for Mo (TF=3.8±4.4) (Figure 3). The fact that none of these elements accumulated in the aerial parts of *Saxifraga paniculataSax* led to two easily distinguishable groups, separating the two species (Figure 2b).

- Among the elements accumulated by *Hippocrepis comosa<u>Hip</u>* typical elemental correlations became visible. Due to their geochemical similarities, As and Sb (for example) are considered to be related elements (Herath et al., 2017). A positive correlation between these two elements has already been observed in plants and increased Sb uptake is presumed to be the result of damaged cell membrane integrity caused by As (Feng et al., 2013; Müller et al., 2013). There is also a strong link between the uptake mechanisms of Mo and Fe (Bittner, 2014). Most
- 400 Mo-enzymes also require Fe-containing redox groups and many studies reported increased Mo

401 uptake in the presence of Fe and vice versa (Bittner, 2014; Kannan and Ramani, 1978; Zimmer

402 and Mendel, 1999).

Three major hypotheses are proposed to explain the characteristic uptake pattern of *Hippocrepis comosa:<u>Hip:</u>*

I) One hypothesis is a characteristic modification of the rhizosphere that would influence the
redox potential. This modification could locally increase phytoavailability of the accumulated
elements, as, except for Al, all are redox sensitive elements (As, Fe, Sb, Mo, and V).

- 408 II) The second hypothesis concerns the uptake pathway of the elements. It is known that As, 409 Sb, Mo, and V are usually absorbed as oxyanions by plants (Hossain et al., 2018; Kabata-Pendias, 2011). Vanadate (H_2VO_4)VO₄³⁻) and arsenate ($H_2AsO_4AsO_4^{3-}$) enter the plant through 410 411 phosphorousP transporters, due to similarities in structure and charge (Chen et al., 2021; 412 Meharg and Hartley-Whitaker, 2002)., Han et al., 2016). Antimonate, on the other hand, is more 413 likely to be absorbed through less selective anion transporters, such as those for Cl⁻ and NO₃⁻, 414 or an incompletely sealed or damaged Casparian strip (Feng et al., 2013; Tschan et al., 2009b, 415 2009a). However, little is known about molybdenumMo uptake by roots. Because of structural 416 and behavioural similarities between molybdate ($HMoO_4MoO_4^2$) and sulphate ($HSO_4SO_4^2$), it 417 is plausible that sulphate transporters are involved (Bittner, 2014; Zimmer and Mendel, 1999). 418 Overexpression or increased activity of so many different transporters, responsible for the 419 particular elemental concentrations found in *Hippocrepis comosaHip*, is unlikely. Nevertheless, 420 a very unselective anion transporter, able to transport phosphate, sulphate, nitrate and chlorate, 421 could be involved. Such a transporter was identified in barley by Preuss et al. (2010), for 422 example.
- 423 III) The third hypothesis is based on the *nitrogenN* fixing property of *Hippocrepis comosaHip*, 424 as Mo and Fe are known to play an essential role as cofactors during symbiotic nitrogenN 425 fixation (Hossain et al., 2018). For this reason, legume plants like *Hippocrepis comosaHip* have 426 a higher demand for Mo than non-nitrogenN fixing plants (Kabata-Pendias, 2011). It is 427 plausible that *Hippocrepis comosaHip* is able to promote Mo absorption in order to guarantee 428 nitrogenN fixation under Mo deficient conditions. However, in Mo rich soils like Jas Roux 429 (19.9±3.71 mg/kg), this property might lead to an increased uptake and thus explain the 430 comparatively high Mo concentrations in *Hippocrepis comosaHip* (Figure 4). It is worth 431 mentioning that vanadiumV may also interfere during nitrogenN fixation by replacing Mo in
- 432 the nitrogenase enzyme (Hernandez et al., 2009).
- Individually, none of the three hypotheses explains all the elemental concentrations observed
 in *Hippocrepis comosa.Hip.* Nevertheless, a combination of the mechanisms could be a valid

435 explanation. <u>Aluminium</u>Al is the only element whose accumulation by <u>*Hippocrepis comosa*Hip</u> 436 remains difficult to explain, as this species is generally found in Ca rich sites and is sensitive to 437 high Al concentrations (Michalet et al., 2002). Concentrations of Al > 1000 mg/kg, like those 438 we measured in aerial parts of *Hippocrepis comosaHip* (1336 mg/kg DW), are rare in higher 439 plants and are considered as Al accumulation (Kabata-Pendias, 2011). The tolerance of toxic 440 elements (Al, As) makes *Hippocrepis comosaHip* an interesting candidate for remediation of 441 polluted soils as its ability to fix nitrogenN thereby improving soil fertility could be an 442 additional advantage (Pajuelo et al 2008).

443

444 **4.4 Translocation factor reveals specific plant strategies**

The differentiation of *Hippocrepis comosa* and *Saxifraga paniculata<u>Hip</u> and <i>Sax*</u> from the other species was mainly caused by high concentrations of a particular set of elements that made both species stand out in the PCA and the cluster analysis. However, this statistical approach failed to detect affinities for single elements, or adaption strategies associated with particularly low concentrations. The translocation factor in combination with the mean elemental <u>concentrations, revealed further adaption strategies and species-related tendencies in elemental</u> <u>uptake (Figure 3).</u>

452 Biological factors are a useful tool to identify adaption strategies. Together with the mean

453 elemental concentrations, the translocation factor indeed revealed further species related

454 tendencies in elemental uptake (Figure 3).

455 For instance, *Biscutella laevigataBis* and *Dianthus sylvestrisDia* did not form a distinct group 456 in the PCA or in the cluster analysis but both had TFs > 1 for a large number of critical metals 457 (Mg, V, Mn, As, Y, Sb, Ba, Ce, Tl). In particular, the TF of *Biscutella laevigataBis* for Tl was 458 remarkably high (TF=19.4±17.2) indicating Tl accumulation (256±224 mg/kg) (Figure 4). 459 While no explanation was found in the literature for the high TF of *Dianthus sylvestris*, 460 Biscutella laevigataDia, Bis belongs to the Brassicaceae family, which includes many 461 hyperaccumulating species (Babula et al., 2008; van der Ent et al., 2013). Biscutella laevigataBis itself has already been mentioned as a Tl accumulating species (LaCoste et al., 462 463 1999; Pošćić et al., 2013). But *Biscutella laevigataBis* was not the only Tl accumulating species, 464 as Galium corrudifoliumGal, which has never previously been associated with metalliferous 465 sites, also showed high concentrations of Tl in its aerial parts (254±410 mg/kg) and high 466 translocation (TF= 2.0 ± 0.5).

- Another species that did not stand out in the statistical analysis was *Juniperus communis subsp.*
- 468 *nana.<u>Jun.</u>* Nevertheless, *Juniperus communis subsp. nana<u>Jun</u>* had the lowest concentrations of

- 469 the elements Mg, Al, S, K, V, Fe, Ni, Cu, Zn, As, Y, Sb, Ce and Tl in its aerial parts (Table 3). 470 This exclusion strategy seems to consist in limited translocation to the aerial parts, as the TF 471 was low for all elements (Figure 3). Given that *Juniperus communis subsp. nanaJun* is the only 472 lignifying plant among the 12 species we analysed, its exclusion capacity might be due to 473 different plant-internal barriers or reduced flow rates due to its small leaf surface area. Even 474 though no studies have demonstrated metal exclusion in Juniperus species, metal concentrations 475 in other conifers (Pinales) have already been reported. For example, Mingorance et al. (2007) 476 found Pinus pinea growing in an industrial area was able to exclude Al, Ba, Cr, Fe, and Pb. In 477 light of the results of our study, Juniperus communis subsp. nana.Jun could be an interesting 478 candidate for phytostabilisation thanks to its high resistance to metal contamination, its 479 widespread occurrence and its ruggedness (Saladin, 2015).
- 480

481 **5** Conclusions

Jas Roux is a unique site to understand the soil-to-<u>Among 12</u> plant transfer of critical metals in an ecosystem<u>species</u> exposed to a cocktail of metals and metalloids in an entirely natural environment that has never been affected by human activities. The statistical approach used in the present study revealed species related uptake patterns in the aerial plant parts. The, two species that stood out were: *Hippocrepis comosa* and *Saxifraga paniculata*, each of which accumulated an individual set of elements.

- 488 Different hypotheses for these particular uptake patterns are proposed. Saxifraga paniculata 489 preferably absorbed divalent cations, but also the REEs Y and Ce. The ionic radius of both 490 elements resembles that of calcium, which was strongly accumulated This uptake pattern could 491 be explained by the Ca accumulation property of Saxifraga paniculata. The affinity to Ca might 492 cause an increased absorption of other elements with a similar ionic radius like Y and Ce, or 493 similar ionic charge like Mn and Ba. Hippocrepis comosa, on the other hand, preferably 494 absorbed oxyanions, perhaps due to overexpression or increased activity of a non-selective 495 anion transporter. In turn, this anion affinity is probably linked to the nitrogen fixation activity 496 of Hippocrepis comosa, which increases its demand for the anionic cofactor Mo (Kabata-Pendias, 2011; Zimmer and Mendel, 1999). Furthermore, Hippocrepis comosa is an interesting 497 498 candidate for phytoremediation of polluted soils, as its ability to fix N₂ improves soil fertility 499 and could thus facilitate the establishment of other species.
- 500 Further, different Different tolerance strategies were identified among the 12studied plant 501 species. On the one hand, Biscutella laevigata and Dianthus sylvestris favoured translocation 502 of multiple critical metals to their aerial parts while, on the other hand, Juniperus communis 503 avoided translocation and used an exclusion strategy. Particularly high concentrations of Tl 504 were found in *Biscutella laevigata* and *Galium corrudifolium*. While hyperaccumulation of Tl 505 has already been reported in Biscutella laevigata, it is a new discovery in Galium corrudifolium. 506 Likewise, the The exclusion capacity of Juniperus communis subsp. nana has never previously 507 been documented and might represent a potential for phytostabilisation applications. Erosion 508 of a polluted soil could be prevented in the long term with no risk of transfer into the food chain 509 because Juniperus communis subsp. nana prevents elemental translocation and the shrub is not 510 consumed by grazing animals.
- 511 To conclude, several promising species among the 12 candidates were identified in the course
- 512 of this study. These results advance our understanding of soil-to-plant transfer of poorly studied
- 513 but increasingly used critical metals and point to the potential of several plant species for future
- 514 phytoremediation and phytoextraction techniques.

- 515
 516
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 - 527

528 Abbreviations:

- 529 ANOVA..... Analysis Of Variance
- 530 a.s.l. Above Sea Level
- 531 DW Dry Weight
- 532 ICP-MS Inductively Coupled Plasma Mass Spectrometry
- 533 ICP-OES Inductively Coupled Plasma Optical Emission Spectroscopy
- 534 PCA Principal Component Analysis
- 535 PERMANOVA..... Permutational Multivariate Analysis Of Variance
- 536 PGMs Platinum Group Metals
- 537 REEs Rare Earth Elements
- 538 SD Standard Deviation
- 539 subsp. Subspecies
- 540 TF Translocation Factor
- 541 XRD X-Ray Diffraction
- 542

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Response to reviewers

We would like to thank both reviewers for their efforts reviewing our revised manuscript. Thanks to their comments we were able to improve consistency and straightforwardness of the paper.

Reviewer #1: The authors have addressed most issues raised by the reviewers. I miss a deeper discussion on the control of mineralogy on the metal bioavailability for these plants, and comparison of metal phytoextraction with the same plants but in cleaner environment. We fully agree that it would be valuable to perform more soil characterization to have a better knowledge on soil mineralogy. However, the majority of the observed variability in metal concentrations could be explained by the plant species. Therefore, we focused our discussion on species related particularities. We identified interesting couples of plant/critical element in this study, for which we will perform further studies, including advanced mineralogy characterization and a plantation of Jas Roux species on an artificially doped soil. Despite that, I consider that the manuscript could be published after minor changes below.

Some minor comments:

I found the Graphical Abstract low informative

We revised the graphical abstract to include additional information and hope the new version is an improvement.

Line 19: avoid the use of ellipsis

We replaced the ellipsis by e.g. in L17: "(e.g. phytostabilisation, phytoextraction)"

Line 60: be consistent with section numbering, Introduction should be the point 1 Thank you for this comment, the oversight has been corrected accordingly

Line 77: the difficulties commented by the authors in the response to reviewers letter should be included in the manuscript.

We agree with the reviewers' comment and added an additional sentence L75-78: "The steep slope, variable soil depth and the presence of large rocks impeded a systematic sampling approach. Nevertheless, the samples were collected as evenly distributed as possible. The soil samples were collected from 10 sampling points in the study zone and consisted of 1 kg of topsoil each (< 15 cm)."

Line 102: 0.45µm ??

The mistake has been corrected for CaCl2 and NH4NO3 extractions

Line 283: which soils? European, worldwide? I am afraid that these values range notably according to background geology. I guess that the authors refer to European average values Table 2 indicates average European topsoil concentrations according to Salminen et al., 2005. The information has been added to the text according to the reviewers' proposition L288-290: "Soil concentrations in European topsoil of Zn are usually around 68 mg/kg, of As, 11 mg/kg, of Mo, 0.9 mg/kg, of Sb, 1 mg/kg, of Ba, 400 mg/kg, and of Tl, 0.8 mg/kg (Table 2)"

Lines 293-295: also, the mineralogy, with predominance of stable minerals, plays an important role in the low metal mobility

We agree with the reviewer that this information was missing in this paragraph. We modified the paragraph accordingly. L299-301: This low mobility indicates the dominance of stable

minerals in Jas Roux. Unlike other sulphosalt sites, Jas Roux has not been disturbed by mining activities which generally increase metal mobility (Resongles et al., 2014; Wei et al., 2018).

Lines 357-361: other minerals can host significant concentrations of Y, so I find this part too speculative without a deeper mineralogical study of the substrate by SEM-EDS or similar techniques.

We agree with the reviewers' comment and reduced the speculative part of this paragraph. L357-361: "Y soil mineralogy in Jas Roux soil has not been studied so far. Y can be associated with several mineral phases, but one common soil mineral based on Y is Xenotime (YPO4) (Ramos et al., 2016). Xenotime has low solubility and phosphate minerals are generally difficult for plants to access (Darcy et al., 2018; Liu and Byrne, 1997; Tropper et al., 2011)."

Line 399: this is not sulphate but bisulphate. Please clear it up

We changed the chemical formula in L392 + L398 to the unprotonated forms to be more consistent.

Line 413: please be consistent, use the chemical symbol or the name We changed all element names to the chemical symbol.

Line 422: accumulators??

This sentence seems right as it is. The subject of the sentence is "concentrations". L416-417: "Concentrations of Al > 1000 mg/kg, like those we measured in aerial parts of Hippocrepis comosa (1336 mg/kg DW), are rare in higher plants and are considered as Al accumulation (Kabata-Pendias, 2011)."

Line 437: As is a metalloid not a metal. Neither is critical

Indeed, this is an important comment. We used the term "critical metals" instead of "critical elements" as it is a standing term in literature and includes metalloids like As and Sb (https://doi.org/10.1016/j.resconrec.2019.104669). We added a sentence to explain our choice of critical elements and justify the presence of As which is considered critical in many studies summarized by Hayes and McCullough - 2018. L130-133: "The critical elements were chosen based on the study on the EU's list of critical raw materials and a review of critical minerals and will be called critical metals throughout the manuscript (European Commission, 2020; Hayes and McCullough, 2018)."

Reviewer #3: Metal accumulation in plants have been widely studied in the past few decades. As stated by Fehlauer et al., however, the case of rare metals accumulation in plants is limited. Totally, the work present here is very important and interesting, but I think the manuscript should be revised again according to the following suggestions or comments. We thank reviewer #3 for the kind words. Thanks to the comments and suggestions we were able to further improve our manuscript.

Line 10: "Their increasing use also involves increased risk of their dispersion", what is the main focus of your study? You introduced rare element application in industry, its risks to humans and how to remove them by phytoremediation. I recommend the authors to highlight the most important goal of the study throughout the manuscript.

Indeed, the main goal of the study was mentioned too late in the introduction. Thanks to the reviewers' comments we modified our second paragraph to improve the straightforwardness of the introduction. L12-15: "Aim of this study is to analyse the soil-to-plant transfer of some

of these metals and to shed light on their uptake pathways. Our findings might bare valuable information for phytoremediation and promote plant-based remediation techniques of metal-contaminated sites."

Furthermore, the main goal of the study is specified again in the introduction, the abstract and the conclusion.

L32-35: "The huge number of factors and complex interactions are impossible to reproduce in the laboratory, thus, to understand the uptake of poorly studied elements, such as critical metals, performing in-situ studies in a natural environment is advantageous."

L474-476: "These results advance our understanding of soil-to-plant transfer of poorly studied but increasingly used critical metals and point to the potential of several plant species for future phytoremediation techniques."

Line 43: "lead, antimony, copper, thallium and arsenic". You used the abbreviated names of some heavy metals before.

We increased consistency by using exclusively the atomic symbol instead of the names.

Line 79: "12 were selected", If two or more plants belonging to a certain species, how did you make the choice?

We chose the sampled species according to occurrence, spatial distribution and family. The family was different for each of the 12 species, except for *Dactylis glomerata* and *Festuca laevigata* that both belong to the family of Poaceae. We modified the sentence to improve clarity. L78-80: "Among the 65 plant species identified in the study zone, 12 were selected, taking into account occurrence, spatial distribution and family affiliation (Table 1)."

Lines 201-225: Pls add the figure captions of Hip, Glo, Dia, etc., following the figure 2. The acronyms of each species are introduced in Table 1. Additionally, we added the information to the caption of each figure as proposed by the reviewer.

Figure 2: Another question, did the PCA show statistical significance?

No, a PCA does not allow to draw any conclusions regarding statistical significance. It is more of a visualization of the dataset. The clustering however gives statistical information about clusters within the dataset. Statistical differences between species regarding the elemental concentrations have been tested using a Games Howel test and are illustrated in Table 3 and 4.

Section 4.4: Pls reference to previous article <u>http://dx.doi.org/10.1016/j.jhazmat.2016.08.079</u> We added the reference to the citation in L 392-394: "Vanadate (VO43-) and arsenate (AsO43-) enter the plant through phosphorous transporters, due to similarities in structure and charge (Chen et al., 2021; Meharg and Hartley-Whitaker, 2002, Han et al., 2016)."

Lines 227-258: Jun, Sax, Hip, etc., pls add the information of the abbreviated names in the first use. See Q4 and check the mistakes throughout the manuscript. You can use the full names and give the abbreviated information in the first use and use the abbreviated names in other parts.

We followed the reviewers' advice and replaced all full names by the acronyms. We kept the full names only in the conclusion and the abstract as these parts need to be understandable without reading the full paper. In order to introduce the acronyms, we added a sentence in material and methods referring to Table 1. L80-81: "For improved readability we attributed acronyms to each of the 12 species throughout the paper (Table 1)". Additionally, all acronyms are explained in the figure captions.

Lines 260-273:TF is one of the indexes to isolate the potential plants for phytoremediation. Why didn't you focus on other indexes, e.g., BF?

We also calculated the bioconcentration factor (C_{roots}/C_{soil}) and the migration ratio $(C_{leavves}/C_{soil})$ using the total soil concentrations. Both factors can give additional information about elemental uptake by plants. In our case however the translocation factor $(C_{leavves}/C_{roots})$ was sufficient to illustrate the main results.

Line 433: Biological factors? You only showed TF in this study.

We improved the sentence according to the reviewers' comment. L426-428: "The translocation factor in combination with the mean elemental concentrations, revealed further adaption strategies and species-related tendencies in elemental uptake (Figure 3)."

Conclusions: Its too long, pls cut down this part. Conclusion is not a summary for each part but a concise conclusion of your findings.

We fully agree with the reviewers' comment we cut down the conclusion significantly (291 words instead of 352 words). We hope that this improved overall readability and precision.

	Acronym	n
Brassicaceae	Bis	4
Poaceae	Dac	3
Caryophyllaceae	Dia	3
Euphorbiaceae	Eup	7
Poaceae	Fes	7
Rubiaceae	Gal	6
Globulariaceae	Glo	3
Fabaceae	Hip	3
Cupressaceae	Jun	3
Asteraceae	Pil	7
Saxifragaceae	Sax	7
Plantaginaceae	Ver	2
	Poaceae Caryophyllaceae Euphorbiaceae Poaceae Rubiaceae Globulariaceae Fabaceae Cupressaceae Asteraceae Saxifragaceae	PoaceaeDacCaryophyllaceaeDiaEuphorbiaceaeEupPoaceaeFesRubiaceaeGalGlobulariaceaeGloFabaceaeHipCupressaceaeJunAsteraceaePilSaxifragaceaeSax

Table 1: Selected plant species, n = number of replicates

Table 2: Average total, mobile $(0.001 \text{ mol/L CaCl}_2)$ and phytoavailable $(1 \text{ mol/L NH}_4\text{NO}_3)$ concentrations and standard deviations (SD) in mg/kg DW of topsoil from Jas Roux (n=10), as well as average concentrations in European topsoil (n≥840) from Salminen et al. 2005. The concentrations were measured by ICP-MS except for total concentrations of Al, K, Ca and S that where measured by ICP-OES.

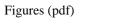
				· · ·						2									
-	Mg	Al	Р	S	К	Ca	v	Mn	Fe	Ni	Cu	Zn	As	Y	Мо	Sb	Ва	Ce	TI
Total	1679	12172	678	11258	21153	3890	110	3401	58930	59.9	44.6	602	2382	4.22	19.3	778	27182	6.61	242
SD	457	1467	112	4892	2134	3651	11.1	802	11219	8.12	3.96	161	975	1.04	3.71	287	6157	2.61	145
CaCl ₂	55.53	1.65	0.595	NA	79.8	NA	0.004	1.62	5.95	0.049	0.043	0.272	0.649	0.003	0.016	0.446	16.0	0.006	0.778
SD	19.6	0.986	0.350	NA	26.0	NA	0.002	1.18	4.85	0.027	0.013	0.146	0.560	0.002	0.012	0.219	6.22	0.003	0.720
NH₄NO ₃	106	0.237	1.52	NA	112	NA	0.003	2.61	0.718	0.048	0.121	0.350	0.103	0.006	0.006	0.113	199	0.001	7.60
SD	34.9	0.176	0.794	NA	27.9	NA	0.001	3.25	0.610	0.024	0.044	0.166	0.096	0.003	0.004	0.054	93.3	0.001	6.78
Europea	n topso	oil					68.1			37.3	17.3	68.1	11.6	22.7	0.943	1.04	400	52.2	0.821
SD							47.0			136	19.0	141	20.1	15.7	1.31	2.04	213	31.3	1.02

indicat	es sigi	nificant di	fference	(NA=no	ot availab	le). Excep	t for Mg,	P, K, Ca	and S,	which w	ere mea	sured in	ICP-O	ES, all co	oncentrat	tions we	re measu	red by l	ICP-MS.	
	U	Mg	Al	P	S	ĸ	Ca	v	Mn	Fe	Ni	Cu	Zn	As	Y	Мо	Sb	Ba	Ce	Tİ
Biscutel	la laevig	ata																		
n=4	Mean	3781 ^{cde}	522 ^{abc}	1954 abcd	6022 abcd	14251 ^d	23583 ^{abc}	0.828 abc	<u>71.3</u> a	589 ^{abc}	1.68 ^a	2.59 ^a	173 ^{ab}	15.9 abcd	0.396 abc	0.951 ^{ab}	5.80 ^{ab}	568 ^a	0.637 abc	256 a
11-4	SD	881	385	698	1937	1091	9538	0.669	51.5	406	0.317	0.458	129	9.61	0.275	0.176	3.24	299	0.579	224
Dactylis	glomera	ita				_				_			_							
n=3	Mean	1838 ^{abcd}	189 ^a	471 ^{ab}	1375 ^{cb}	7315 ^{abcd}	4911 ^{ab}	0.320 ^a	230 ^{ab}	324 ^{ab}	1.84 ^a	3.47 ^{ab}	37.9 ^{ab}	9.34 ^{ab}	0.267 ^{abc}	1.77 ^a	3.29 ^{ab}	<u>193</u> a	0.263 ^{ab}	5.85 ^a
11-5	SD	247	72.2	123	112	2630	1566	0.119	84.6	168	0.395	0.421	27.3	5.79	0.073	0.300	1.92	124	0.118	4.07
Dianthu					-															
n=3	Mean	2672 ^{abcde}	659 ^{abc}	526 ^b	1243 ^{abc}	7520 ^{abcd}	25214 ^{abc}	1.10 ^{abc}	219 ^{ab}	797 ^{abc}	2.19 ^a	4.69 ^{ab}	64.3 ^{ab}	20.3 abcd	0.445 abc	0.982 ^{ab}	6.14 ^{ab}	948 ^a	0.887 abcd	6.68 ^a
11-5	SD	931	686	85.7	231	1938	6796	1.18	91.7	870	0.949	2.10	10.7	24.3	0.420	0.385	6.86	439	0.902	3.17
Euphorb	ia cypar	issias			_					_	_	_			_			_		
n=7	Mean	2995 ^{ce}	405 ^{ab}	1818 ^{cd}	2401 ^{bd}	16302 ^{cde}	12840 ^{bc}	0.667 ^{ab}	77.5 ^a	517 ^{ab}	1.73 ^a	5.07 ^b	94.9 ^b	14.2 ^{ab}	0.230 ^{ab}	0.594 ^{ab}	4.28 ^{ab}	642 ^a	0.440 ^{ab}	26.9 ^a
11-7	SD	858	433	504	716	5342	4526	0.727	31.8	500	0.689	1.38	44.3	12.2	0.224	0.213	3.51	469	0.461	31.5
Festuca	violaced	1					•													
n=7	Mean	917 ^b	377 ^{ab}	613 ^{ab}	1558 ^{abc}	8885 ^{bc}	<u>3715</u> a	0.616 ^{ab}	80.5 ^a	506 ^{ab}	2.04 ^a	2.60 ^a	29.0 ^{ab}	13.4 ^{ab}	0.320 ^{ab}	1.87 ^a	3.53 ^{ab}	310 ^a	0.500 ^{ab}	6.87 ^a
	SD	379	327	285	549	1943	1817	0.557	49.2	504	2.34	0.556	14.8	11.2	0.274	0.786	3.18	279	0.477	10.5
Galium	-			1			1					1								
n=6	Mean	1903 ^{adc}	550 abc	810 abcd	1939 ^{abcd}	10407 ^{abcd}	15129 ^c	0.941 ^{abc}	149 ^{ab}	753 ^{abc}	3.98 ^a	5.29 ^{ab}	79.0 ^{ab}	21.4 ^{abcd}	0.532 abc	1.27 ^{ab}	6.63 ^{ab}	627 ^a	0.732 abc	254 ^a
-	SD	220	464	422	660	4019	4398	0.740	100	570	6.73	2.58	39.1	14.4	0.643	0.567	4.80	363	0.541	410
Globula	-		1	i	i		i					i		1	1	1				
n=3	Mean	1734 ^{abcd}	1111 ^{abc}	734 ^{abcd}	2598 ^d	5378 ^{ab}	13248 ^{bc}	1.80 abc	235 ^{ab}	1249 ^{abc}	3.33 ^a	5.50 ^{ab}	36.9 ^{ab}	37.7 ^{abcd}	0.749 abc	0.859 ab	10.8 ^{ab}	838 ^a	1.57 ^{abcd}	5.56 ^a
-	SD	337	794	298	123	1184	2191	1.38	207	1054	1.25	0.990	5.90	35.6	0.590	0.117	10.1	298	1.25	2.42
Hippocre																				
n=3	Mean	1313 ^{abd}	1336 ^C	903 ^a	1622 ^c	7089 ^{abc}	14237 ^c	<u>2.05</u> ^c	241 ^b	1686 ^C	8.89 ^a	7.22 ^{ab}	39.4 ^b	55.9 ^{cd}	0.954 ^c	<u>30.8</u> ab	<u>16.1</u> ab	1029 ^a	1.72 ^{cd}	18.1 ^a
	SD	196	118	52.0	143	1124	1883	0.190	25.8	274	5.24	2.28	4.57	10.4	0.151	19.9	4.05	354	0.168	14.6
Juniperu	is comm	unis ssp. nan			;															
n=3	Mean	789 ^{ab}	<u>178</u> a	1010 ^{ac}	<u>885</u> a	<u>5272</u> a	10831 ^{bc}	0.292 a	102 ^a	<u>203</u> a	<u>1.49</u> a	2.31 ^a	<u>14.1</u> a	2.61 ac	<u>0.127</u> a	0.514 ^{ab}	0.888 ^a	249 ^a	<u>0.227</u> ^a	2.18 ^a
	SD	321	79.3	34.5	69.2	390	582	0.127	25.4	71.3	0.039	0.169	2.81	1.59	0.042	0.128	0.497	96.8	0.094	0.345
Pilosella																				
n=7	Mean	1952 ^{acd}	563 ^b	1391 ^d	1852 ^{cd}	<u>22525</u> e	9919 ^{bc}	0.893 ^b	144 ^{ab}	575 ^{bc}	4.39 ^a	<u>11.7</u> ab	49.5 ^b	16.2 ^{bd}	0.442 ^{bc}	0.873 ^{ab}	4.56 ^b	492 ^a	0.709 ^b	6.50 ^a
	SD	184	130	184	447	2935	1933	0.210	69.7	146	1.81	5.99	14.6	7.41	0.169	0.559	1.73	166	0.182	3.16
Saxifrag			1	1		1	1			1		1	1	1 .	1	1 .	1		1 . 1	
n=7	Mean	4056 e	604 ^{ab}	773 ^{ab}	995 ^{ab}	5914 ^{ab}	<u>52243</u> d	0.423 ^{ab}	1282 ^c	318 ^{ab}	<u>36.6</u> b	3.12 ^{ab}	636 ab	7.63 ^{abcd}	<u>3.31</u> d	<u>0.445</u> b	2.41 ^{ab}	2323 a	<u>2.67</u> d	28.6 ^a
	SD	695	275	256	128	1931	10053	0.407	356	219	11.6	0.535	405	3.81	1.18	0.362	1.53	1359	0.974	18.5
Veronico	-		1			1				1				1	1	1				
n=2	Mean	1309 ^{NA}	711 ^{NA}	534 ^{NA}	1195 ^{NA}	6816 ^{NA}	6569 ^{NA}	1.16 ^{NA}	194 ^{NA}	876 ^{NA}	3.31 ^{NA}	10.1 ^{NA}	83.9 ^{NA}	22.5 ^{NA}	0.535 ^{NA}	1.50 ^{NA}	6.95 ^{NA}	625 ^{NA}	1.07 ^{NA}	10.2 ^{NA}
	SD	171	109	85.1	265	525	674	0.236	82.4	130	0.023	4.18	24.5	11.0	0.087	0.090	2.65	68.5	0.300	6.37

Table 3: Average concentrations (mg/kg DW) and standard deviation (SD) in the aerial parts of each plant species analysed. The <u>highest</u> and <u>lowest</u> concentrations of each element are highlighted. The statistical difference between the species (α =0.05), is shown for each element by a sequence of letters within which the absence of matches indicates significant difference (NA=not available). Except for Mg, P, K, Ca and S, which were measured in ICP-OES, all concentrations were measured by ICP-MS.

signifi	cant di	fference	(NA=no	t availab	ole). Exc	ept for M	g, P, K, Ca	and S, w	hich wei	e measu	red in IC	P-OES	, all con	centratio	ons were	measure	d by ICl	P-MS.		
		Mg	AI	Р	S	K	Ca	v	Mn	Fe	Ni	Cu	Zn	As	Y	Мо	Sb	Ва	Ce	Tİ
Biscute	lla laevig	jata		•																
n=3	Mean	1654 ^{abc}	704 ^a	1068 ^{ab}	5100 ab	12449 ^{abc}	7720 ^{bc}	0.913 a	<u>133</u> a	1124 ^{ab}	7.37 ^a	<u>4.60</u> a	241 ^a	31.3 ^a	0.804 ab	<u>0.609</u> a	11.9 ^a	1073 ^a	1.19 ^a	23.1 ^{ab}
11-5	SD	295	555	305	1058	3469	2199	0.660	87.7	1173	4.45	3.30	210	32.7	0.717	0.229	13.3	1339	1.09	18.9
Dactylis	glomera	1		1							1									
n=3	Mean	1000 ^{ab}	2241 ^{ab}	591 ^{ab}	2991 ^{ab}	3485 ^a	13851 ^{abcd}	4.34 ^{abc}	339 ^{ab}	5449 ab	8.60 ^a	19.1 ^a	95.3 ^a	<u>210</u> ab	2.23 abc	3.58 ^{ab}	39.1 ^a	4155 a	2.98 ^{ab}	125 ^{ab}
-	SD	369	1867	146	2047	867	1476	4.35	191	6101	4.09	11.0	47.5	274	1.08	1.50	26.5	5324	2.35	43.4
Dianthu	s sylvest	1	I	!		l	1	!	!			I	I	I		l	!		l	1
n=3	Mean	1588 ^{ab}	<u>640</u> a	454 ^a	1242 ^a	3760 ^a	10156 ^{abcde}	1.02 ^{ab}	171 ^a	<u>785</u> a	<u>4.80</u> a	21.6 ^a	<u>52.4</u> a	<u>21.3</u> a	0.881 ^a	3.52 ^{ab}	7.77 ^a	1321 ^a	<u>1.08</u> a	6.89 a
	SD	189	528	132	305	904	5642	0.876	91.0	658	2.25	23.2	20.1	19.9	0.341	0.747	5.84	710	0.794	2.80
Euphorb	pia cypar	1	ı .		1 . 1	ı .	ı .	ı .		1 . 1		ı	ı	ı .				1 1		ı .
n=7		2697 ^c	2397 ^{ab}	1293 ^b	3041 ^b	8129 ^b	8735 ^{ab}	3.94 ^{bc}	273 ^a	3641 ^{ab}	7.05 ^a	13.7 ^a	169 ^a	156 ^{ab}	1.78 ^{abc}	2.14 ^b	47.6 ^a	3913 ^a	3.07 ^{ab}	<u>174</u> ab
	SD	585	746	392	710	1740	3179	1.44	121	1801	1.96	6.87	109	150	0.614	0.683	37.2	4295	1.13	156
Festuca	violaced	i	i.	i.	i.,	i	i .	i.	i	i.,		i	i	i .	. i	i .	i	i	i .	i.
n=6		1040 ^a	3446 ^{ab}	608 ^{ab}	1880 ^{ab}	4430 ^a	9012 ^{ab}	5.50 ^{abc}	335 ^a	4280 ^{ab}	9.04 ^a	17.5 ^a	84.5 ^a	141 ^{ab}	3.08 ^{bc}	2.65 ^b	35.6 ^a	2413 ^a	4.88 ^{ab}	71.3 ^{ab}
	SD	167	1746	256	813	1132	1398	3.07	173	2297	1.95	10.8	31.3	75.9	1.07	0.627	15.2	1609	2.34	63.2
Galium	corrudif	1		i		- 1-		I	I .		_	I .	I .				i .			1
n=6		1341 ^{ab}		1	2133 ^{ab}	5846 ^{ab}	16531 ^{cde}	4.04 ^{abc}	i	3171 ^{ab}		9.62 ^a	i	99.7 ^{ab}				2730 ^a	3.41 ^{ab}	i
<u>Chilin Ia</u>	SD	506	2200	126	1047	1981	3479	4.04	130	2494	3.50	2.88	75.2	81.1	1.48	0.777	29.7	2683	2.95	223
Globula	ria cordi	í				ab	ab	abc							a la abo					l
n=3		1095 ^{abc}	-		1404 ^a	3831 ^{ab}	7257 ab	3.70 ^{abc}		2728 ^{ab}				86.8 ^{ab}						31.5 ^{ab}
Hinnoor	SD	591	1263	198	108	2731	796	2.14	225	1586	0.688	9.27	29.9	57.7	0.808	0.778	17.2	290	1.57	11.9
пірросі	epis cor	1	aaca ah			aaaa ah	anan abrd	= to abc	co. ah	tere ab	to o ah			l .== ah	a Ta ahr	a a a ah			4 To ah	in c ab
n=3	Mean		3362 ab		1546 ^a	3232 ab	9595 ^{abcd}	5.48 ^{abc}		4558 ^{ab}			i	i				2143 a		47.6 ^{ab}
luningr	SD	187 unis ssp. n	1433 ana	85.0	150	1476	1772	2.68	280	2101	6.50	4.18	20.6	90.2	1.12	11.1	25.6	1554	2.02	44.0
Jumpen			:	628 ^{ab}	1005 8	3972 ^{ab}	25231 ^{ade}	3.11 ^{abc}	201 ^a	2009 ^{ab}	7.79 ^a	25.8 ^a	105 ^a	51.4 ^a	1.95 ^{abc}	2.60 ^{ab}	2013	1179 ^a	2.24 ^a	38.4 ^{ab}
n=3		1017 ^{ab} 170	1852 48	110	1635 -	3972 do 1374	3922	0.293	201 ° 50.5	2009 00	7.79 ° 1.62	25.8 ° 5.02	43.0	51.4 ° 8.25	0.481	2.60 a. 0.550	30.1 ° 11.1	1179 5	2.24 ° 0.240	i
Pilosella	SD a neleter		130	110	192	1574	5922	0.295	50.5	200	1.02	5.02	45.0	0.25	0.461	0.550	11.1	100	0.240	14.7
	-	1690 ^{bc}	1360 ^a	<u>1380</u> ab	1226 ab	24555 ^c	9481 ^{abcd}	2.38 ^{abc}	231 ^a	1478 ^a	6.01 ^a	20.0 ^a	70.0 ^a	42.2 ^a	1.55 ^{ab}	1.44 ^{ab}	11.3 ^a	1746 ^a	2.10 ^a	54.7 ^b
n=7	SD	354	964	973	462	6961	2683	1.83	101	1478 -	2.53	20.0 - 16.1	33.9	42.2 - 35.8	0.808	1.44 0.666	8.68	830	1.37	23.5
Saxifrao	a panicı	i	504	575	402	0501	2005	1.05	101	1145	2.55	10.1	55.5	55.0	0.000	0.000	0.00	050	1.57	23.5
		2048 abc	6453 b	623 ap	1888 ^{ab}	4081 ^a	26773 ^e	9.34 ^c	1252 b	4851 ^b	30.9 b	32.4 ^a	795 a	177 ^b	<u>7.37</u> ^c	2.12 ^b	41.3 ^a	3802 ^a	8.07 b	50.4 ^{ab}
n=7	SD	756	2688	171	534	1345	6062	4.42	510	1806	8.07	17.6	616	68.5	3.24	0.721	18.9	1843	3.27	30.0
Veronic	a frutica		2000		557	10.0	0002		510	1000	0.07	1.10	010	00.5	5.27	0.7 = 1	10.5	10.0	3.27	
		1002 ^{NA}	1149 ^{NA}	434 ^{NA}	<u>900</u> NA	5819 ^{NA}	7097 ^{NA}	1.81 ^{NA}	273 ^{NA}	1261 ^{NA}	8.48 ^{NA}	33.9 NA	122 NA	36.8 ^{NA}	1.27 ^{NA}	2 20 NA	12.5 ^{NA}	<u>889</u> NA	1 74 ^{NA}	44.5 ^{NA}
n=2	SD	18.4	217	124	183	1012	1470	0.596	21.7	449	6.28	27.3	83.0	22.4	0.071	0.077	5.70	205	0.504	47.4
	50																			<u> </u>

Table 4: Average concentrations (mg/kg DW) and standard deviation (SD) in the roots of each plant species analysed. The <u>highest</u> and <u>lowest</u> concentrations of each element are highlighted. The statistical difference between the species (α =0.05), is shown for each element by a sequence of letters within which the absence of matches indicates significant difference (NA=not available). Except for Mg, P, K, Ca and S, which were measured in ICP-OES, all concentrations were measured by ICP-MS.



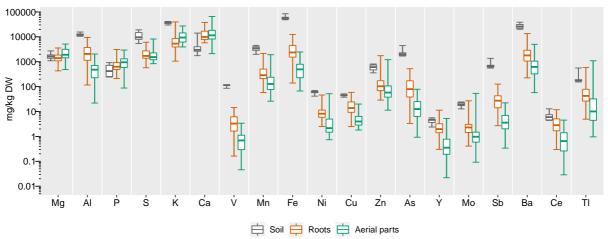


Figure 1: Boxplot with whiskers (minimum to maximum) of elemental concentrations in aerial plant parts (n=55) and roots (n=55) of all individuals of the 12 species compared to the total concentrations in the soil (n=10) in mg/kg DW. The bottom and top of the boxes represent the lower and upper quartile and the median is represented by the horizontal line inside each box.

(coloured print)

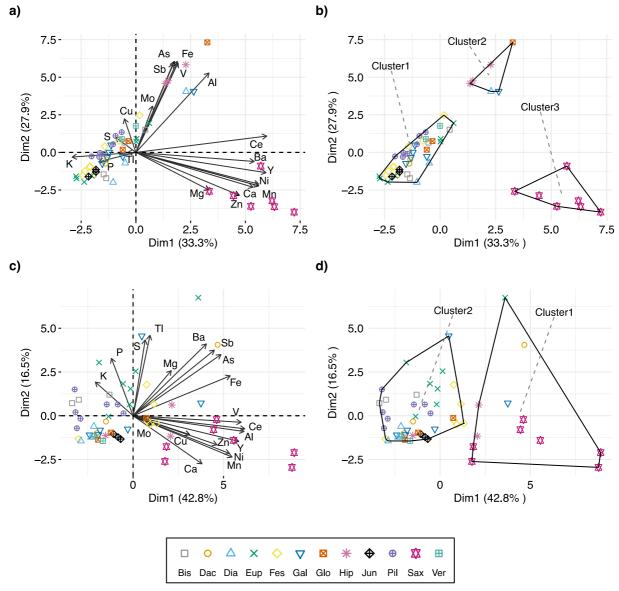


Figure 2: Biplot of the results of a primary component analysis of the elementals concentrations in aerial parts (a) and roots (c) including the individuals according to their species and results of partitioning clustering (K-means) on elemental concentrations in aerial plant parts (b) and roots (d) with a Hopkins clustering tendency of respectively 0.19 and 0.20. Coloured symbols representing the 12 plant species Bis (*Biscutella laevigata*), Dac (*Dactylis glomerata*), Dia (*Dianthus sylvestris*), Eup (*Euphorbia cyparissias*), Fes (*Festuca laevigata*), Gal (*Galium corrudifolium*), Glo (*Globularia cordifolia*), Hip (*Hippocrepis comosa*), Jun (*Juniperus communis subsp. nana*), Pil (*Pilosella peleteriana*), Sax (*Saxifraga paniculata*) and Ver (*Veronica fruticans*).

(coloured print)

		Mg	AI	Р	S	к	Ca	v	Mn	Fe	Ni	Cu	Zn	As	Y	Мо	Sb	Ва	Ce	тΙ
Bis	Mean	2.2	2.0	1.6	1.3	1.2	3.8	2.0	1.0	1.5	0.4	0.7	1.0	1.7	1.0	1.6	1.4	1.6	1.3	19.4
DIS	SD	0.8	2.2	0.4	0.7	0.3	2.3	2.0	0.8	1.3	0.3	0.5	0.3	1.9	0.8	0.3	1.2	1.3	1.1	17.2
Dac	Mean	1.9	0.1	0.8	0.6	2.1	0.4	0.1	0.8	0.1	0.2	0.2	0.4	0.1	0.1	0.6	0.1	0.1	0.1	0.0
Dac	SD	0.4	0.1	0.1	0.4	0.4	0.0	0.1	0.6	0.1	0.1	0.2	0.3	0.1	0.1	0.5	0.0	0.0	0.1	0.0
Dia	Mean	1.7	1.6	1.2	1.0	2.1	2.8	1.8	1.6	1.6	0.6	0.4	1.5	1.5	0.5	0.3	0.8	0.9	1.1	1.1
	SD	0.7	1.7	0.4	0.2	0.8	1.1	2.0	0.9	1.6	0.4	0.2	0.9	1.5	0.3	0.1	0.6	0.6	0.9	0.7
Eup	Mean	1.1	0.2	1.5	0.8	2.1	1.5	0.1	0.3	0.2	0.3	0.4	0.6	0.1	0.1	0.3	0.1	0.3	0.1	0.2
Lup	SD	0.2	0.2	0.6	0.3	0.7	0.3	0.1	0.1	0.1	0.1	0.3	0.2	0.1	0.1	0.1	0.1	0.3	0.1	0.1
Fes	Mean	0.9	0.4	1.0	0.8	2.2	0.4	0.6	0.6	0.6	0.3	0.2	0.4	0.6	0.2	0.7	0.2	0.4	0.4	0.1
105	SD	0.5	0.7	0.3	0.4	1.2	0.2	1.2	0.8	1.2	0.5	0.1	0.3	1.2	0.2	0.4	0.3	0.8	0.6	0.1
Gal	Mean	1.6	0.4	1.2	1.1	1.8	1.0	0.4	0.6	0.4	0.7	0.6	0.6	0.3	0.3	0.7	0.3	0.5	0.3	2.0
<u>001</u>	SD	0.5	0.3	0.6	0.6	0.5	0.6	0.4	0.4	0.3	1.1	0.4	0.4	0.3	0.5	0.4	0.3	0.4	0.3	0.5
Glo	Mean	2.4	0.5	1.8	1.9	2.1	1.8	0.5	0.5	0.4	0.4	0.3	0.5	0.4	0.3	0.3	0.3	0.7	0.4	0.2
010	SD	2.2	0.1	0.3	0.1	1.5	0.2	0.1	0.3	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.3	0.1	0.1
Hip	Mean	1.8	0.5	2.3	1.1	2.5	1.5	0.5	0.4	0.5	0.7	0.6	0.5	0.4	0.4	3.8	0.4	0.6	0.4	0.7
<u>mp</u>	SD	0.7	0.2	0.4	0.1	1.1	0.5	0.3	0.2	0.3	0.2	0.0	0.2	0.3	0.3	4.4	0.3	0.3	0.3	0.7
Jun	Mean	0.8	0.1	1.6	0.5	1.5	0.4	0.1	0.5	0.1	0.2	0.1	0.2	0.0	0.1	0.2	0.0	0.2	0.1	0.1
<u>7411</u>	SD	0.5	0.1	0.3	0.1	0.5	0.1	0.1	0.3	0.0	0.1	0.0	0.1	0.0	0.0	0.0	0.0	0.1	0.1	0.0
Pil	Mean	1.2	0.7	1.5	1.7	1.0	1.1	0.7	0.7	0.7	0.9	0.7	0.8	0.6	0.4	0.6	0.6	0.3	0.5	0.1
<u>r 11</u>	SD	0.2	0.5	0.9	1.0	0.2	0.3	0.5	0.4	0.4	0.2	0.3	0.2	0.4	0.3	0.4	0.3	0.2	0.3	0.1
Sax	Mean	2.2	0.1	1.3	0.6	1.6	2.1	0.1	1.2	0.1	1.2	0.1	1.0	0.1	0.5	0.3	0.1	0.8	0.4	0.7
<u>547</u>	SD	1.0	0.1	0.6	0.2	0.8	0.8	0.1	0.5	0.1	0.5	0.1	0.9	0.1	0.3	0.3	0.1	0.6	0.2	0.3
Ver	Mean	1.3	0.6	1.3	1.4	1.2	1.0	0.7	0.7	0.7	0.5	0.4	0.8	0.6	0.4	0.7	0.6	0.7	0.6	0.4
ver	SD	0.2	0.0	0.6	0.6	0.3	0.3	0.1	0.2	0.2	0.4	0.2	0.3	0.1	0.1	0.1	0.0	0.1	0.0	0.2

Figure 3: Heatmap of average translocation-factors (mean) and standard deviation (SD) for each plant species and chemical element (Concentration in the leaves / Concentration in the roots). The acronyms are standing for the plant species *Biscutella laevigata* (Bis), *Dactylis glomerata* (Dac), *Dianthus sylvestris* (Dia), *Euphorbia cyparissias* (Eup), *Festuca laevigata* (Fes), *Galium corrudifolium* (Gal), *Globularia cordifolia* (Glo), *Hippocrepis comosa* (Hip), *Juniperus communis subsp. nana* (Jun), *Pilosella peleteriana* (Pil), *Saxifraga paniculata* (Sax) and *Veronica fruticans* (Ver).

(coloured print)

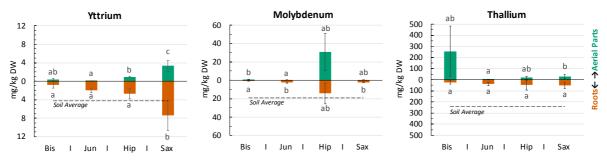


Figure 4: Average concentrations of Y, Mo and Tl in aerial plant parts and root of *Biscutella laevigata* (Bis, n=4), *Juniperus communis* (Jun, n=3), *Hippocrepis comosa* (Hip, n=3) and *Saxifraga paniculata* (Sax, n=4) compared to the average soil concentration (n=10) in mg/kg DW.

Supplementary Material

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Till Fehlauer	Writing - Original Draft, Visualization, Formal analysis, Investigation
Blanche Collin	Writing - Review & Editing, Funding acquisition, Supervision,
Conceptualization	
Bernard Angeletti	Resources, Investigation
Catherine Santaella	Writing - Review & Editing, Formal analysis
Cedric Dentant	Writing - Review & Editing, Supervision
Perrine Chaurand	Resources
Clement Levard	Resources
Clement Levard Cedric Gonneau	Resources, Investigation
Cedric Gonneau	
Cedric Gonneau	Resources, Investigation
Cedric Gonneau Daniel Borschneck	Resources, Investigation Resources, Investigation

Declaration of interests

 \boxtimes The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

□The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: