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## **Pedo-geochemical background and sediment contamination of metal(loid)s in the old mining-district of Salsigne (Orbiel valley, France)**

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### **Abstract**

15 The mining district of Salsigne in the Orbiel valley (Aude, France) was at one time the first gold mine in Europe and the first arsenic mine in the world. However, no scientific studies have evaluated the magnitude of its environmental impact. In this study, the pedo-geochemical background (PGB) was determined for 14 metal(loid) elements, including As. It appears that the PGB values for As and Sb are  
20 relatively high with  $44 \pm 12$  and  $0.9 \pm 1.2$  mg.kg<sup>-1</sup>, respectively, because of the geological particularities of this area. In a second step, these PGB values (normalized with Ti concentrations) were used as local references to determine enrichment factors (EFs) of bed river sediments for the Orbiel River and two of its major tributaries (Gresillou and Russec rivers) collected between November, 2018 and July, 2020. Results showed that riverine sediments are contaminated by past mining activity and/or current storage  
25 areas. If we except the major elements (Fe, Ti and at a lesser extent Mn), we observed that As, Cu, Sb, Pb present the highest concentrations relative to the remaining elements (Cd, Co, V, Ni and Cr). In the case of As, EFs can reach 74 in the Orbiel River, 1000 in the Gresillou River and 27 in the Russec River. These calculations were also performed for sediments transported by the extreme flood of October 14, 2018, that killed 15 people and potentially remobilized contamination in the valley. We

30 observed that the As concentrations of suspended samples from Grésillou and Russec rivers have reached 870 mg.kg<sup>-1</sup>.

Finally, the As concentrations measured in the river sediments of this valley are of the same order of magnitude than those published in the literature for environments strongly impacted by mining or mineral processing activities.

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**Keywords**

Arsenic, metal(loid)s, Orbiel valley, environmental contamination, geochemical background, soil, mines

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## 1. Introduction

Mining and ore processing activities during the so-called modern period (19<sup>th</sup> and 20<sup>th</sup> centuries) have strongly impacted the natural environments by releasing metals, metalloids as well as other products used in industrial processes (e.g., cyanide, polyacrylamide) everywhere in the world. These released compounds have a strong impact on the Critical Zone, but also on human health through food (solid and liquid) ingestion, absorption through skin, and air inhalation (Plumlee and Ziegler, 2007; Landrigan et al., 2018; Li et al., 2021). Together with the manufacturing industry, mining activities have played (and are playing) a major role in the dispersion of metals and metalloids through environment compartments, i.e. surface and ground waters, soils/sediments, atmosphere and biosphere (Purves, 1985; Nriagu et Pacyna, 1988). Face to these legacies of the past and before to carry out rehabilitation, it is necessary to establish rigorous contamination diagnostics, based on the specific study of each environmental compartment. The first compartment to be considered is the source reservoir, which corresponds to soils and sediments that have witnessed primary contamination from mining activities (Salomons and Förstner, 1984; Resongles et al., 2014). In order to assess the level of contamination of soils and sediments, it is necessary - first of all - to know the geochemical background of the area of interest in order to use it as a local reference.

Our work fits in this general context and more particularly in the Orbiel valley (South France) context, where was located the Salsigne mining district corresponding at one time to the largest gold mine in Europe and the largest arsenic mine in the world (Pujol, 2014). Following the closure of the mine in 2004, some rehabilitation has been undertaken. Nevertheless, recent works published by Khaska et al. (2015, 2018) have shown that As concentrations in surface (Orbiel River) and ground waters were high (36-40  $\mu\text{g.L}^{-1}$ ) downstream the remediated zone whereas the upstream concentrations were in the 2-6  $\mu\text{g.L}^{-1}$  range. During flood, the As concentrations measured in the Orbiel River may be significantly higher ( $\sim 120 \mu\text{g.L}^{-1}$  during the flood of May 22, 2012). These authors have explained

these high concentrations by the contribution of anthropogenic arsenic (Ca-arsenate). If these publications have revealed a strong contribution of old mining sites on the waters of the valley, no scientific studies have been published to give an overview of the contamination of soils or sediments in this valley. However, in October 2018, the potential contamination of arsenic and other metal(loid)s in this valley resurfaced with force following the extreme flash flood of the Orbiel River. Dramatic consequences have occurred on local populations (loss of 15 human lives, destroyed infrastructures), together with remobilization and dispersion of contaminants, which may have some strong environmental impacts (Girardeau, 2019; Lebouc et al., 2019). Even if this extreme hydroclimatic event has recently awakened the attention of authorities and local populations, this area has been poorly studied in the past as previously highlighted, and especially regarding the environmental consequences of the mining activities and the potential risks involved for human health. Indeed, because of its physical and chemical properties, As can induce serious health problems in humans (Duker et al., 2005; Bulka et al., 2016; Saint-Jacques et al., 2018).

The main objectives of this study were then: i) to define the pedogeochemical background (PGB) on the basis of a rigorous methodology; and ii) to use this prerequisite estimated PGB to assess the level of sediment contamination of the Orbiel River and of its major tributaries. Different soils, which were sampled from various geological formations located uphill of the mining district, in pristine environments, were analyzed. The PGB was then determined for 14 metal(oid)s, and compared to data available at national and European scales. Sediments were collected in various rivers in this mining district. Their metal(loid)s contents were determined and discussed according to the PGB determined in the first part.

This work constitutes an indispensable and fundamental preliminary study to be able to address in the near future the problems of exposure and risks for the populations living in these contaminated environments.

## 2. Material and methods

### 2.1 Site description

#### 2.1.1 Geological context

The Orbiel valley is located on the southern slope of the Montagne Noire, the south-western terminus of the Massif Central (France) (Figure 1). It consists first of a central nucleus of granites and gneiss, surrounded by a halo of micaschists (Tollon, 1970; Berger et al., 1993; Cassard et al., 1993). These micaschists form a metasedimentary cover varying in age from Cambro-Ordovician to Devonian (Cassard et al., 1993). Further south, the southern slope of the Montagne Noire is composed predominantly of Palaeozoic carbonate formations, strongly folded and dislocated, and thrust sheet dating from the Hercynian orogenesis (Cassard et al., 1993). Finally, a Tertiary cover in total discordance with the Hercynian base can be observed in the south. This Tertiary cover presents in its northern part mainly limestone and in its southern part fluvial sedimentary deposits (“Carcassonne ‘s molasse”).

The studied zone contains the Salsigne district, a major Au deposit that belong to the Variscan gold province. Mineralization processes (i.e., hydrothermal) occurred during late orogenic processes between 330 and 290 Ma and have affected late Proterozoic to early Cambrian schists, early Cambrian sedimentary rocks, Devonian limestones and sandstone (Bouchot et al., 2005). Indeed, many metalliferous bodies have been identified in the area, as evidenced by past mining activities (and particularly the second Age of Iron, the Roman period, and the modern period of the 19<sup>th</sup> and 20<sup>th</sup> centuries) (Fabre et al., 2016). These bodies can be classified in filonian and stratiform bodies, according to Tollon (1970). The “metalliferous district of Salsigne”, which is rich in arsenic, gold, lead and bismuth, is located in the southern slope of the Montagne Noire. The most represented mineral phases in the deposits are sulphide, such as pyrite ( $\text{FeS}_2$ ), arsenopyrite ( $\text{FeAsS}$ ), pyrrhotite ( $\text{Fe}_{1-x}\text{S}$  where  $0 < x < 0.2$ ), chalcopyrite ( $\text{CuFeS}_2$ ) and bismuthinite ( $\text{Bi}_2\text{S}_3$ ); and to a lesser extent galena ( $\text{PbS}$ ),

115 sphalerite ((Zn,Fe)S), scheelite (CaWO<sub>4</sub>) and wolframite ((Fe,Mn,Mg)WO<sub>4</sub>) (Marcoux and Lescuyer, 1994; Demange et al., 1986).

### 2.1.2 *Historic context of mining exploitation and waste storage*

During the modern mining period, the mining district of Salsigne included various concessions:  
120 Malabau, Pujol, Lastours, La Caunette, Salsigne, Villanière, and Villardonnell (Geoderis, 2012). These concessions are reported in Figure 1 with the main sites that have been exploited for extraction and/or treatment in the past and the main current waste deposits. It has been estimated that nearly 12 million t of ore would have been extracted to produce around 120 t of gold, and 450,000 t of arsenic during the period 1873-2004 (Girard, 2011). About 15,000 t of copper were also produced, as well as silver (~150  
125 t) and bismuth (~800 t).

During the last period of mining activity (1992-1997), the SEPS company (Société d'Exploitation et de Pyrométallurgie de Salsigne) has also reprocessed foreign waste from the area mine using the powerful water jacket furnace, inherited from the pyrometallurgical process of ore treatment. This company has also played a major role in the dispersion of contaminants in the environment. Nowadays,  
130 the main waste storage areas are located at Montredon (4 million t of waste rock containing 10 to 12% arsenic), Artus (7 million t of waste rock), and La Combe du Saut. The Montredon storage area contains mainly waste from the cyanuration of the ore, but also flotation residues, lime arsenates, residues from the Malabau mine, and sewage sludge; La Combe du Saut mainly contains waste from the demolition of the plant and buildings (furnace crucibles, bricks, drums, etc.), from the SEPS  
135 company and pyrometallurgic slag (Desaulty et al., 2016). A lime arsenate storage area is also located at La Combe du Saut, close to the Orbiel bedriver. It comes from the water treatment plant that recoils percolation water from the Montredon and Artus sites. This water treatment plant extracts most of the arsenic from these waters by precipitating lime arsenites (Ca<sub>3</sub>(AsO<sub>4</sub>)) before their released in the Orbiel

River. Unlike Montredon, the storage of the Artus is neither sealed in the background nor covered by  
140 an impermeable system. The seepage water thus passes through the 10 Million t of tailings by  
progressively loading with arsenic, cyanides and other chemical elements.

### 2.1.3 *Hydrological context*

The Orbiel valley has a typically continental climate with both Mediterranean and oceanic  
145 influences, with relatively high annual rainfall (900 mm), a relatively mild average temperature (13 °C)  
and frequent winds (>300 days/yr). The source of the Orbiel River is in the Montagne Noire at 900 m  
in elevation close to the city of Labruguière (Tarn, France). About 40 km long, it flows in a southern  
direction to reach the Aude River in the Trèbes town, east of Carcassonne (Figure 2). The Orbiel River  
has a watershed of approximately 250 km<sup>2</sup> and three major tributaries (Russec, Rieutord and Clamoux).  
150 Another tributary is the intermittent Gresillou River (Figure 2). At Bouilhonnac, the Orbiel River has  
an average annual flow of 2.67 m<sup>3</sup>.s<sup>-1</sup> (<http://hydro.eaufrance.fr/>). Due to the Mediterranean influence,  
this catchment is regularly affected by high and extreme flash flood events during which maximum  
flows can reach several hundreds of m<sup>3</sup>.s<sup>-1</sup> (e.g. 450 m<sup>3</sup>.s<sup>-1</sup> during the flood of 14-15 October 2018;  
Lebouc et al., 2019). The Orbiel River flowing from north to south crosses the former mining district of  
155 Salsigne. The Russec River drains: (i) the former concessions of Villardonel and Malabau, particularly  
La Messette site; (ii) the western area of the former open-pit mine of Salsigne drained by the Gourg  
Peyris River, a left bank tributary of the Russec River; and (iii) and the western flank of the Montredon  
storage site. The Gresillou River drains the waste Ramèle and Nartau deposits.

Apart from the former mining activity and its legacies (abandoned mines, waste storage areas, etc.),  
160 there are no other significant activities in this watershed. From an economic point of view the two main  
activities are tourism and viticulture. However, this tourist activity is limited (mainly hiker) with very  
little impact on the natural environment and the vineyard is relatively small with little influence on our  
area of study. The urbanized places are the villages that we can meet along the Orbiel River, in the



section we studied (Miraval-Cabardès, Mas Cabardès, Les Ilhes, Lastours). These villages have a  
165 population varying between 50 and 190 inhabitants. There is no urbanized area on the Gresillou and  
Russec rivers.

## 2.2 Soils sampling strategy

Because the Orbiel valley presents a strong variability in terms of lithology (Figure 1), we collected  
170 a representative soil sample set for each major geological formation outcropping uphill of the mining  
district, in relatively pristine zones. To this end, old and recent satellites pictures, ancient pre-industrial  
mine maps (Fabre et al., 2016) and maps of tailing location (Figure 2) were used to select pristine  
sampling sites. Forest soils were privileged as sampling sites, as the presence of trees ensure that soils  
were not, or almost not, disturbed, at least for the lifetime of the existent trees (Horckmans et al., 2005).  
175 Some sampling were performed in moorlands with the presence of shrubs, when forests were absent or  
unreachable. Totally, seven sites, labelled in accordance with the label given in the geological map for  
their parental rock, were sampled uphill of the mining district (Spd, Spe, Spf, Spg, Sph, Spi, Srk) for  
the pedo-geochemical background determination, and two sites were sampled within the district (k1  
and k2b) for elemental contents comparison (Figure 1).

180 The sampling methodology for soils in each selected site was derived from the one used in the  
French RMQS program (Jolivet et al., 2016). For each site, the methodology was based on a composite  
sampling at different depth coupled with a soil profile description. Thus, a soil pit was dug in order to  
describe the soil profile morphology and estimate succession and depths of main horizons in order to  
define the composite sampling depths. Then, composite samples were formed, following a square grid  
185 pattern of (10 m × 10 m) divided into 16 sub areas. For each horizon, or every 20 cm when the horizon  
was too thick, 16 sub-samples were taken randomly in each sub area (homogeneously distributed  
within the square), and bulked together. The sampling was conducted until the substratum or the  
saprolite (C horizon), which was sampled as well, was reached. Samples were taken using a soil auger

(Edelman Ø 7 cm), which was cleaned with demineralized water between each composite sampling  
190 depth, kept in hermetic plastic bags, which were immediately labelled. Back to the laboratory, they  
were oven dried for a week at 50 °C, sieved at 2 mm using a nylon sieve, and crushed in an agate  
mortar prior to mineralization. The soils morpho-genetic description (from field study) was reported in  
the supplementary material (SM-1) for each geological formation and sampling site.

### 195 2.3 Sediment sampling

All the sediments were collected during several sampling campaigns: 11/12/2018, 12/14/2018,  
01/17/2019, 04/16-17/2019, and 07/21/2020. Sediments were sampled directly in or the closest to the  
riverbed, and stored in clean, new, 50 mL polycarbonate tubes. Back to the laboratory, they were oven  
dried for a week at 50 °C and crushed in an agate mortar prior to mineralization.

200 Sediments were sampled in 12 locations along the three main streams in the valley: the Orbiel River,  
and **two of its** tributaries, the Grésillou and the Russec rivers (Figure 2). The Orbiel River was sampled  
at Miraval-Cabardès (called MIR), in the upstream portion, and going downstream at Mas-Cabardès  
(MAS), at Lastours (LAS) before the confluence with the Grésillou River, at Moulin d'Artigues (ART),  
at Pont de Limousis (LIM), and finally at Gué de Lassac (GUE); the Grésillou River was sampled  
205 upstream (NAU) and downstream the storage area of Nartau (NAD); the Russec River was sampled at  
Raissac (RAI), Salitis (SLT) and St-Angel (ANG); all these three sites are located downstream close to  
the confluence with Orbiel River.

In this work, we took into account two other types of solid samples: (i) samples collected in  
November and December 2018, corresponding to precipitates (labelled GUE-2 and GUE-3) formed  
210 within an excavation created in the plain bordering the Orbiel River, at the station Gué de Lassac by the  
flash flood, on October 14-15, 2018; and (ii) sediments, named LAG-1, ANG-1, ANG-2, ANG-3 and  
ANG-4, for which we are sure that they were deposited during the extreme flash flood, on October 14-  
15, 2018. During this flood, no water sampling (and suspended sediment) could be collected due to the

violence of the event. However, when the average speed of the river decreases, deposits of sediments  
215 occurred on flooded surfaces. The suspended particulate matter considered in this work are sediments  
that have been deposited on certain surfaces and that we were able to recover after the flood. Sediments  
denoted LAG-1 correspond to Grésillou suspended sediments deposited in the schoolyard of Lastours,  
sampled in April 2019. From the Russec River, sediments ANG-1 and ANG-3 were sampled at the  
same location, in November 2018 and January 2019, respectively, and corresponds to suspended  
220 sediments deposited as mud on a field. These samples were collected near the confluence between  
Russec and Orbiel rivers, and could be considered as samples representing a mixture of suspended  
sediments from both rivers. ANG-2 and ANG-4 are suspended sediments from the Russec River  
deposited in an empty wine tank, and in the warehouse storage of the farm at the Wine Castle Saint  
Angel, respectively.

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#### 2.4 Determination of elemental concentrations

Elemental concentrations in soil samples were determined after acid digestion of 50 mg of dried soil  
or sediment. This procedure was performed using a Milestone® Ultrawave microwave at the HSM  
laboratory (HydroSciences Montpellier, France) in a class 10,000 clean room. Firstly, 1 mL HF (40%,  
230 Merck Suprapur), 3 mL HNO<sub>3</sub> (67-69%, Analytika Analpure), and 3 mL HCl (30%, Merck Suprapur),  
were added together in closed Teflon vials containing 50 mg of sample. Then, the acid digestion was  
performed in the microwave, where samples were heated for 5 min at 100 °C, then 5 min at 150 °C,  
and finally 5 min at 200 °C, at a pressure starting at 40 bar (N<sub>2</sub>) and increasing to 80-100 bar. Blank  
tests indicated that the level of contamination induced by the acid digestion procedure was negligible, <  
235 2% of the less concentrated element. Samples were then let to evaporate to dryness on a hotplate and  
then diluted with 3 mL 1% HNO<sub>3</sub> and put in an ultrasound bath for 20 min. Finally, they were heated  
on a hotplate at 100 °C for 10 min. A certified reference material, “Montana II Soil” (NIST-SRM  
2711a, Mackey et al., 2010), underwent the same procedure.

Samples were then analyzed using ICP-MS (iCAP Q, Thermo Scientific- Kinetic Energy  
240 Discrimination mode using He) at the AETE-ISO platform (OSU OREME/Université de Montpellier,  
France). Metal(loid) concentrations were determined with external calibration using (Be, Sc, Ge, Rh,  
Ir) as internal standards to correct potential sensitivity drifts. Limits of quantification were: (i) lower  
than 1  $\mu\text{g.kg}^{-1}$  for Cd, Co, Cr, Ni, Pb, Sb, U and V; (ii) between 1 and 3  $\mu\text{g.kg}^{-1}$  for As, Mn, Ti and Zn;  
(iii) 35  $\mu\text{g.kg}^{-1}$  for Cu; and (iv) 121  $\mu\text{g.kg}^{-1}$  for Fe. Uncertainties associated to the measurement were  
245 inferior to 3%. The agreement of certified sample analysis with certified concentrations was better than  
85% for Cr, Pb and Ti, and 95% for As, Cd, Co, Cu, Fe, Mn, Ni, Sb, U, V, Zn.

## 2.5 Enrichment factor (EF) and Geoaccumulation index determination

EF were determined using the following relationship:

$$250 \quad EF = \left( \frac{[X]}{[Ti]} \right)_Y / \left( \frac{[X]}{[Ti]} \right)_Z \quad (\text{Eq. 1})$$

For soil EF calculation, with respect to national or international references, Y = PGB data and Z =  
reference data. For sediment EF calculation, Y = sediment data and Z = PGB data. Ti concentration was  
chosen as reference in order to erase pedological variations and anomalies due to natural pedogenetic  
processes, external sources or anthropogenic activities. An EF close to 1 indicates an absence of  
255 contamination while a value significantly greater than 1 indicates a contribution by an anthropogenic  
source.

Considering the high lithological variability from upstream to downstream, it was difficult, if not  
impossible, to define a natural geochemical background for sediments. We could have defined this  
sediment geochemical background in the uppermost part of the watershed, but considering this  
260 variability of lithology, would it have been representative of sediments all along the stream? We believe  
that no. Therefore, using the PGB as a reference for sediments was a much more robust process for us.

Geo-accumulation index ( $I_{geo}$ ) have been calculated for sediments according to the formula proposed by Müller (1969):

$$I_{geo} = \log_2 (C_{sed} / 1.5 C_{ref}) \quad (\text{Eq. 2})$$

265 where  $C_{sed}$  is the element concentration in the sediment and  $C_{ref}$  the element concentration in the reference, PGB for us here. Müller (1969) proposed several classes to define sediment quality: 0: uncontaminated; between 0 and 1: uncontaminated to moderately contaminated; between 1 and 2: moderately contaminated; between 2 and 3: moderately to strongly contaminated; between 3 and 4: strongly contaminated; between 4 and 5: strongly to extremely contaminated; above 5: extremely  
270 contaminated.

## 2.6 Statistical analyses for soil data treatment

Two Kruskal-Wallis tests were performed using PAST 4.01® software, in order to assess significant differences between: (i) surface horizon and substratum (C horizon), and (ii) between soils sampled  
275 uphill of the mining district and those sampled within the district. The Kruskal-Wallis test is a non-parametric test, which therefore does not assume a normal distribution of data, based on medians, and used to compare different populations of small sizes (Zolfaghari et al., 2018). This test is then adapted to our preliminary study in the Orbiel valley.

## 280 3. Results

### 3.1 Morpho-genetic description, classification and metal(loid) concentrations of soils

The different soil profiles from the different lithologies are described in Supplementary Information SI-1. According to field observations and the international World Reference Base classification (WRB from FAO, 2015), among all soils studied, 3 belong to Luvisols (*SPd*, *SPe*, *SRk*), 3 can be classified as

285 Leptosols (*SPf*, *SPh*, *SPi-j*), 2 of them belong to Cambisols (*SPg*, *kI*) and the last one can be classified  
as Calcisol (*k2b*), as shown in SI-1.

Soil metal(oid) concentrations are reported in Table 1 for 14 elements (As, Cd, Co, Cr, Cu, Fe, Mn, Ni, Pb, Sb, Ti, U, V, Zn). The first Kruskal-Wallis test performed in the soil data revealed that surface horizons and substratum or C horizon did not exhibit difference in terms of elements concentrations. Consequently, data from all horizons were considered for the following presentation of the results. The second Kruskal-Wallis test - comparing uphill soils and soils sampled within the mining district area - showed significant differences for As, Cd, Mn, Pb, Sb, U, V and Zn. For this first group of elements, soils located uphill of the district exhibited lower concentrations, except for U and V (see Table 1). For U, we think the difference of concentration should be related to the lithology behind these two soils [k2 (sandstone) and k2b (dolomitic limestone)]. Sandstone are generally poor in U and carbonates exhibit generally lower U concentration than felsic rocks (Condie, 1993). On the contrary, Cr, Co, Cu, Fe, Ni and Ti did not show significant differences between soils uphill and within the mining district. This latter set of elements will be called group 2. Considering the uphill soils, we classified the elements by decreasing range of concentrations (in mg.kg<sup>-1</sup> except Fe and Ti in %): Fe (2.29 – 5.09%) > Ti (0.37 – 0.50%) > Mn (200 – 1600) > V (103.6 – 136.5) > Zn (44.8 – 153.5) > Cr (68.5 – 94.4) > As (35.6 – 75.1) > Cu (200 – 70.2) > Pb (27.0 – 61.9) > Ni (11.0 – 43.1) > Co (2.6 – 18.9) > U (2.13 – 2.91) > Sb (0.32 – 6.25) > Cd (0.15 – 0.35). Arsenic concentrations range from 88.1 to 162.6 mg.kg<sup>-1</sup> within the district, i.e. higher values than uphill of the district (35.6 to 75.1 mg.kg<sup>-1</sup>). A more detailed comparison of As concentrations in soils (location / depth) is given in supplementary information SI-2

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### 3.2 Metal(loid) concentrations in sediments

Metal(loid) concentrations in sediments are reported in Table 2 for all the sampled rivers. Considering the Orbiel River, we classified elements by decreasing order of concentration (in mg.kg<sup>-1</sup>

except Fe and Ti in %): Fe (0.68 – 6.85%) > Ti (0.05 – 0.53%) > Mn (160 – 1050) > Pb (17.3 –  
310 832.5) > As (7.7 – 515) > Zn (66.8 – 204.4) > V (14.6 – 152.7) > Cr (8.4 – 120.1) > Cu (2.6 – 99.0) >  
Ni (4.2 – 61.1) > Co (2.4 – 21.1) > Sb (0.02 – 5.11) > U (0.64 – 3.51) > Cd (0.1 – 0.98). Arsenic  
concentrations increase along the watercourse of the Orbiel River, from 8 mg.kg<sup>-1</sup> upstream, to 515  
mg.kg<sup>-1</sup> downstream (Gué de Lassac sampling point, see Figure 2). We observed that precipitates  
collected in the excavation of the floodplain bordering the Orbiel River showed very high As (1.44 –  
315 2.79%) and Fe (11.1 – 14.1%) concentrations. In the case of the Grésillou River, it comes the following  
range of concentrations (in mg.kg<sup>-1</sup> except Fe and Ti in %) for river sediments: Fe (3.96 – 9.58%) > Ti  
(0.06 – 0.28%) > As (118 – 6472) > Mn (100 – 980) > Cu (46.0 – 409.2) > Pb (20.1 – 202.3) > Zn  
(91.2 – 185.1) > V (89.5 – 136.3) > Cr (62.3 – 96.6) > Ni (30.6 – 52.2) > Sb (0.67 – 36.17) > Co (11.5  
– 21.1) > U (2.16 – 3.37) > Cd (0.21 – 1.64). In the Grésillou River, the As sediment concentrations,  
320 which can reach up to 0.6%, were much higher than in the bed Orbiel River sediments. The sample  
LAG-1, collected in the schoolyard of Lastours, showed that suspended sediments transported during  
the flash flood by the Grésillou River can have a very high As content, i.e. 873 mg.kg<sup>-1</sup>. It can also be  
noted that the sediment Co, Cu and Cd concentrations were in a high range in this river. For the Russec  
River, the following order was observed (concentrations in mg.kg<sup>-1</sup> except Fe and Ti in %): Fe (1.86 –  
325 4.85%) > Ti (0.09 – 0.26%) Mn (290 – 970) > As (143 – 631) > V (45.5 – 127.2) > Cu (26.1 – 107.4) >  
Cr (30.4 – 83.1) > Pb (28.1 – 54.1) > Ni (15.9- 50.0) > Co (6.7 – 21.0) > U (1.17 – 3.74) > Sb (1.44 –  
3.43) > Cd (0.23-0.70). We observed that ANG-1 and ANG-3 samples present a similar high As  
concentration with 554 and 631 mg.kg<sup>-1</sup>, respectively. As a reminder, these two samples were collected  
at the same location (as mud in the field) at two different times (November, 2018, and January, 2019).  
330 ANG-2 and ANG-4 present an As concentration of 470 and 143 mg.kg<sup>-1</sup>, respectively.

Globally, As appears to be one of the analysed elements present with the highest concentrations for  
the three monitored rivers. On the other hand, samples ANG-1 to ANG-3, collected at the wine castle

Saint-Angel and deposited after the flash flood of October 14-15, 2018, showed very high concentrations of As, Cu, Cr and Mn compared to other sediments collected in the Russec River bed.

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## 4. Discussion

### 4.1 Determination of the pedo-geochemical background (PGB) of the Salsigne mining district

Equation (3) was used to calculate the PGB for a given element, as proposed by Reimann et al. (2005).

$$340 \quad PGB = Median \pm 2 \times MAD \quad (Eq. 3)$$

where *MAD* corresponds to the median absolute deviation. It has been proven to be less sensitive to extreme values than the “classical” equation (mean  $\pm 2 \times$  standard deviation), and therefore is now being used in several studies focused on the definition of a geochemical background (e.g. Esmaeili et al., 2014; Tapia et al., 2012; Rothwell and Cooke, 2015). Depending on groups 1 or 2, the median and the MAD were calculated on either all soil samples, i.e. soils located uphill and soils located within the mining district, or only on soils located uphill of the district. Thus, for group 1 elements, the PGB was only based on soils located uphill of the district. On the contrary, for group 2 elements, the PGB was based on all soil samples. Median and MAD values were calculated considering all sub-samples from each soil, i.e. each depth. Globally, the total numbers of samples for the calculations were 19 for group 1 and 23 for group 2. This method ensured that the calculated PGB included both the geological variability and the variability vs. depth due to pedogenesis. The set of estimated PGB values, considering the 14 selected elements for the Orbiel valley soils, was reported in Table 3. For As, the PGB for the Orbiel valley is estimated to  $44 \pm 12 \text{ mg.kg}^{-1}$ . It can be observed that the PGB ranges of Co, Ni, Pb, Sb and Zn were quite wide:  $13 \pm 10$ ,  $28 \pm 18$ ,  $38 \pm 21$ ,  $0.9 \pm 1.2$  and  $92 \pm 44 \text{ mg.kg}^{-1}$ , respectively. These broad ranges of Co, Ni, Pb, Sb and Zn concentrations could be due to the natural heterogeneity among the soils in the area.

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The PGB determined in the Orbiel valley are then compared to median values (baselines) obtained for soils at the scale of France (Saby et al., 2019a,b) and Europe (Salimen et al., 2005), in the framework of the RMQS (<https://www.gissol.fr/>) and FOREGS (<http://weppi.gtk.fi/publ/foregsatlas/>) programs, respectively (Table 3). It comes that these two baselines appear to be very similar, and significantly lower than the PGB calculated in this Orbiel area. Estimated median concentrations, or PGB in the case of Orbiel soils, for As, Cd, Cr, Cu, Pb, V and Zn, seem to be significantly higher in soils from the Orbiel valley than median concentrations observed in soils at the French or European scales. As and other trace metal and metalloid contents in soils and sediments in the vicinity of Au mining areas and As-rich deposits are known to be higher than those found in area with lower mineral indexes, so much so that arsenic has long been used as a geochemical indicator of the presence of mining potential by geologists. The values we find for the geochemical background of the soils of the Orbiel Valley for As, Cu, Pb and Zn (As:  $44 \pm 12$  mg.kg<sup>-1</sup>; Cu:  $38 \pm 16$ ; Pb:  $38 \pm 21$ , and Zn:  $92 \pm 44$ ) are comparable to those of other mining provinces in the published literature. Threshold concentration in As, Cu, Pb and Zn for natural geochemical background in soils in the surroundings of mines located in the Iberian Pyrite Belt in the South of Portugal (Pelica et al., 2018 and references therein) are respectively 50, 300, 55 and 140 mg.kg<sup>-1</sup> for the Neves-Corvo mine area (copper-rich deposits) and 50, 260, 30-90 and 150-250 mg.kg<sup>-1</sup> for the Aljustrel mine area (lead-zinc rich deposits). In the Oruro mining region of Bolivia (Andean tin belt), the work of Tapia et al. (2012) determined the geochemical background (in the form [minimum; median; maximum]) for As at [38; 52; 72] mg.kg<sup>-1</sup>, for Cu with [40; 52; 66] mg.kg<sup>-1</sup> and Zn with [104; 119; 147] mg.kg<sup>-1</sup>. The natural richness in As in soils located in mining environments is not always verified. Indeed, there are mining environments with low arsenic geochemical background values such as in the vicinity of the Cachoeira Au deposit in the Para state in Brazil where the deep ferralitic soils display geochemical background in As very low, with a mean concentration varying between 0.7 to 6 mg.kg<sup>-1</sup>, comparable to values for tropical forested

soils without any anthropic influence (Souza Neto et al., 2020). Soils from the old mining area of the Antiform of Martinamor in Spain display As median concentrations of 26 mg.kg<sup>-1</sup> (García-Sánchez et al., 2010). In Poland, Galuszka et al. (2018) investigated soils geochemistry in the vicinity of an historic lead mining area (Mt. Karczowka) and found concentration values for geochemical background for As of respectively 8-16 mg.kg<sup>-1</sup>. Similar geochemical background values were determined from the sediment study for the state of Morelos, Mexico with 12.5 mg.kg<sup>-1</sup> As (Barats et al., 2020). On the opposite, other studies have shown higher values for the geochemical background of As, in particular, a British Geological Survey study for a former As mine in Devon, England, determined a geochemical background in soils of 93 mg.kg<sup>-1</sup> (Palumbo-Roe and Klinck, 2007). The recently published geochemical background data of the Orbiel valley by Melleton and Girardeau (2019) have obtained “ PGB ranges between 45 and 339 mg.kg<sup>-1</sup> for As, depending on the geological formation. Unfortunately, this PGB cannot be used in the present study since it was defined using data previously selected and chosen for another use, i.e. to calculate mineralization indices from the “national mining inventory” (Lambert, 2005). Moreover, data to establish these so-called PGBs have been lacking from the point of view of sampling information (e.g., sampling date, depth, sampling protocol), analytical techniques (limits of detection/quantification) or the mathematical relationship used. Compared to our study, for which As PGB was estimated to 44 ± 12 mg.kg<sup>-1</sup>, it is obvious that such a difference would have strong impacts on the establishment of an objective diagnostic of the contamination by As. In conclusion, the establishment of the geochemical background, which can vary greatly depending on local lithology and pedogenetic processes, needs to be considered with particular attention on a case by case basis. Using geochemical background data, which did not follow the PGB determination guidelines, could be considered as a wrong way leading on the one hand to overestimate PGB data and on the other hand to under-estimate the impact of human activities in relation to mining on the global environment contamination of the Orbiel valley.

## 4.2 Enrichment factor in Salsigne 'soils compared to soil national and international references

We calculated the EF of the soils of the Orbiel valley (PGB values) with respect to the baselines obtained for the soils at the scale of France (Saby et al., 2019a,b) and Europe (Salimen et al., 2005) (see paragraph 4.1), and with respect to upper crust data from Taylor and McLennan (1985) and Rudnick and Gao (2003) (Figure 3). Exception of As and Sb, all the references used give more or less the same flat pattern of enrichment factors, suggesting that our values of PGB were very similar to both chemical composition of the upper crust and soil baselines (France, Europe). The As EF estimated in our work is 8 times higher than the European soil baseline and upper crust data from Rudnick and Gao (2003), and very high, around 30 times more, with respect to the upper crust data from Taylor and McLennan (1985). We also observed that our As PGB value is close to the value proposed by Saby et al. (2019a,b) for the baseline of soils from French territory. In the case of Sb, the EF value is around 7 times higher than the upper crust one (Taylor and McLennan, 1985). Our data set suggests a relative natural As enrichment for the preserved soils within the Orbiel valley. Finally, the data homogeneity with respect to the different references underlines the data robustness and the choice to use them as a local reference for studying sediments of the Orbiel valley (see paragraph 4.3).

## 4.3 Contamination of sediments by metal(loid)s

For sediment metal(loid)s of the Orbiel valley rivers (Orbiel and its tributaries Russec and Grésillou), EF values were calculated from normalization with respect to Ti as before and (Eq. 1) by using  $Y = \text{sediment data}$  and  $Z = \text{PGB data}$  (see Figure 4). From left to right one goes from upstream to downstream for each river. Samples deposited both by the Grésillou River in the Lastours schoolyard and by the Russec River in the Saint-Angel wine castle during this event are reported in Figure 4, in order to better assess the potential risks involved for human health after flash flood of October 14-15, 2018. It should be noted that for the Russec River, all samples were collected in its downstream part,

430 just before the confluence with the Orbiel River (see Figure 2). Therefore, we cannot compare sediment chemical composition before and after former mine extraction/storage areas for the Russec River.

For the Grésillou River (NAU, NAD and LAG sites, Figure 4), we found that As, Pb, Sb, Cd, Co, Cu and Fe have high EF values ( $2.3 < EF < 1042$ ), while Mn, Ni, Cr, U, V and Zn exhibit low to moderate EF values ( $1.5 < EF < 10.0$ ). Given the limited number of samples, it is thus difficult to  
435 determine the difference in concentrations between upstream (NAU) and downstream (NAD) points compared to the Nartau storage area. However, it clearly appears that the EF values of Grésillou sediments were higher after the Nartau deposit area than upstream (NAU). By considering the LAG-1 sample, we can introduce constraints on the chemical composition of suspended sediments transported by the Grésillou River during the flash flood on October 14-15, 2018. In that specific case, it comes  
440 that the sediment EFs were 190, 108, 33, 20 and 13 for As, Sb, Cd, Pb and Cu, respectively. For all these elements the EF was particularly high. We have reported on the Supplementary Information (SI-4) the granulometry of some of the sediment samples presented in this study. We observed that the suspended sediments from the flood of October, 2018, such as LAG-1 have the finest particle size. The high EF observed are not surprising because the finest fraction is generally more concentrated than the  
445 coarser one; this finest fraction may contain clays and Fe oxyhydroxydes, minerals known to control metals and metalloids.

For the Russec River (Figure 4), we can classify analyzed elements into three groups: (i) As, Cd, Cu, and Sb with high enrichment compared to the geochemical background,  $1.5 < EF < 27.5$ , the highest EF values being for As (9 – 27.5); (ii) Pb with moderate EF values,  $1.8 < EF < 6.8$ ; and (iii) Co, Zn, Fe,  
450 Mn, U, Ni, Cr, and V with low EF values or no enrichment,  $0.9 < EF < 3.3$ . Although we did not have sediments from the upstream part of this river, we can nevertheless observe that the sediments from the downstream part are strongly impacted by anthropogenic activities and most likely the former mines or mine waste storage areas located in the Russec River watershed with respect to As, Cd, Sb and Cu. It is

important to notify that the suspended sediments deposited in the confluence area between Orbiel and  
455 Russec rivers (ANG-1 and ANG-3), after the flood of October, 2018, are marked by high EF for these  
elements (As, Cd, Sb, and Cu) since these EFs are between 5 (Cu) and 20 (As). Again, like for the  
suspended sediment from the Gresillou River, suspended sediment from the Russec river exhibit a finer  
granulometry which can explain the high EFs of these specific samples.

For the Orbiel River, we found that some elements show a regular increase in the EF values from  
460 upstream to downstream. This is the case for As, Cr, Cu, Fe, Ni, Sb and V, which have EF values close  
to 1, i.e. no enrichment, in the upstream part, and which can reach 74, 4.1, 6, 4.5, 4.8, 11 and 3.5,  
respectively, downstream at the Gué de Lassic location (GUE). This statement revealed a strong  
progressive enrichment of sediments in these elements relative to the PGB going downstream. The  
Orbiel River, flowing from north to south, crosses the old mining zone (former mines and current  
465 storage zones) and its sediments were gradually enriched in As, Cr, Cu, Fe, Ni, Sb and V. The EF  
values decreased at the Water Treatment Station (WTS) and then increased again between WTS and  
the Gué de Lassic sites. In the case of Co, Mn and Zn, the EF values were low, close to 2, and constant  
along the river. However, there were two peculiarities: (i) the EF values were high for Zn and Pb at  
Miraval-Cabardès, the most upstream point of the Orbiel River; and (ii) the systematic increased in Mn  
470 and Co EF values between Lastours (LAS) and Pont de Limousis (LIM), at Moulin d'Artigues (ART)  
sites. For Pb and U, the EF values were constant and low, up to the GUE site, where a significant  
increase occurred. For Cd, high EF values, around 5, were calculated, more or less constant from  
upstream to downstream, with an increase at the GUE site. If we return specifically to the case of As,  
the EF calculated for the Orbiel and Russec rivers sediments (up to 100) appear to be in the high range  
475 of values reported in the literature while the EF calculated for the Gresillou River sediments (up to  
1000) are much higher ( $1 < EF < 285$ , Au Giant Mine in Yellowknife, USA, Cheney et al., 2020;  $1 <  
EF < 133$ , Oruro mine, Bolivia, Tapia et al., 2012).

The sediment contamination, i.e. excess of element with respect to geochemical background, could be explained by the two following ways: either by transportation of primary bearing minerals (for example sulphides) or secondary minerals such as iron arsenates (scorodite), characterized at the Nartau site (Lucia Perez Serrano, pers. com.), which were extracted by mechanical weathering and carried from upstream to downstream; or by a chemical transfer from the dissolved phase (e.g., < 0.20  $\mu\text{m}$ ) to the particulate phase (e.g., > 0.20  $\mu\text{m}$ ) via sorption process (Sigg et al., 2014). We know that the dissolved As charge of the Orbiel River water is also increasing from upstream to downstream (Khaska et al., 2015) by the input of water rich in As, during stable hydrological periods or during extreme events (flash floods as in October, 2018) relatively frequent in this region of Southern France (Raynaud et al., 2015). By studying As concentrations and  $^{87}\text{Sr}/^{86}\text{Sr}$  ratio in surface and ground water, Khaska et al. (2015) have showed that most of the As transported by the Orbiel River originate from leaching of Ca-arsenate stored on site. These Ca-arsenate were produced by the WTS to remove As from contaminated waters, As being co-precipitated with whitewash as  $\text{Ca}_3(\text{AsO}_4)_2$ .

Numerous studies have shown the major role of Fe minerals (particularly hydrous iron oxides or iron sulfate minerals), which control the arsenic fate in the environments impacted by mining activity (e.g., Casiot et al., 2003; Sanchez-Espana et al., 2005). In this study, the Fe-As relationship (see Supplementary Information SI-3) also suggested a control of arsenic by Fe minerals in the Orbiel valley sediments, at least for the most concentrated samples. As mentioned before, a preliminary study conducted on the waste storage area of Nartau identified the presence of scorodite, a hydrated iron arsenate ( $\text{FeAsO}_4 \cdot 2\text{H}_2\text{O}$ ) in abundant quantities. As this area is absolutely not isolated, these wastes can be mobilized towards the Gresillou and the Orbiel rivers.

The calculation of the geoaccumulation index (Müller, 1969) is another method for assessing the degree of metal and metalloids contamination of sediments in the Orbiel Valley. According to this index (see section 2.5), some samples may be extremely contaminated with respect to Cr (Water Treatment Station, WTS), Pb (WTS) and Sb (downstream Grésillou). Others sediments are strongly to extremely

contaminated with respect to As (WTS, Gué de Lassac, suspended sediments from the Grésillou and Orbiel/Russec), Cd (Gue de Lassac), Cr (WTS), Cu (downstream Gresillou), Pb (WTS) and Sb (downstream Gresillou). Considering this index, it appears that most of the sediments from the Orbiel River and its two tributaries, Russec and Gresillou, are moderately to extremely contaminated in As. Finally, this index gives results consistent with those of the enrichment factors.

From our set of data, it was possible to compare the measured As concentrations in the sediments of the Orbiel Valley with those from other environments, such as rivers, deltas, lakes, in preserved area [1-14] or on the contrary, in strongly impacted by human activities due to mining or mineral processing [15-22] (Figure 5). This comparison, which is not exhaustive, can nevertheless give us some lessons:

- i)* In the case of preserved environments, As concentrations range from a few  $\text{mg.kg}^{-1}$  to a few dozen  $\text{mg.kg}^{-1}$  (typically  $< 50 \text{ mg.kg}^{-1}$ ). Moreover, estimates of As average suspended sediment concentrations for major world rivers have been published, yielding 36 (+/- 27)  $\text{mg.kg}^{-1}$  (Viers et al., 2009), 5  $\text{mg.kg}^{-1}$  (Martin and Meybeck, 1976) or 14  $\text{mg.kg}^{-1}$  (Savenko, 2006).
- ii)* Impacted sediment concentrations can reach several thousands of  $\text{mg.kg}^{-1}$ .
- iii)* Finally, this figure shows that sediment concentrations collected in the Orbiel valley, after the Salsigne mining district area, can reach the highest values measured elsewhere in strongly impacted zones.

## 5. Conclusion

This first study conducted on the soils and sediments of the Orbiel valley enabled us to obtain a few important results:

- 530 i) we first determined pedo-geochemical background (PGB) concentrations for As and also other metal(loid) elements. It appears that compared to soil baselines at national or European level, pristine soils of the Orbiel valley present a marked natural enrichment for As and Sb, with concentrations of  $44 \pm 12 \text{ mg.kg}^{-1}$  and  $0.9 \pm 1.2 \text{ mg.kg}^{-1}$ , respectively. The other elements belong to the magnitude order of soil baselines at national or European level.
- 535 ii) The PGB was then used as a local reference to assess the level of contamination of sediments collected along the Orbiel River and two of its major tributaries. We showed the contamination of riverine sediments by past mining activity and/or current storage areas of the former mining district of Salsigne. Indeed, if some peculiarities may appear from one river to another, it appears that the sediments of these rivers are strongly impacted for As, Cu, Sb and Pb after the district mining area. At Gué de Lassac station, at the outlet of the mining district, the enrichment factors can reach 74, 15, 11 and 6 for As, Pb, Sb and Cu, respectively.
- 540 iii) Finally, by collecting sediments that have been transported during and deposited after the flash flood of October 14-15, 2018, we can estimate the chemical composition of the suspended sediments transported by the rivers of the valley. In particular, the concentrations of As in suspended samples from the Grésillou River have reached  $870 \text{ mg.kg}^{-1}$ . In the case of the Russec and Orbiel rivers, suspended sediment concentrations belong to the range 143
- 545 –  $631 \text{ mg.kg}^{-1}$ .

Finally, following these new insights brought by this preliminary study, further studies are currently in progress in the Orbiel valley, based on a multidisciplinary approach in order to better take into account the different environmental compartments and to better understand elemental transfers occurring between water, soil, vegetation and atmosphere. The major issue will concern the population health by proposing adapted mitigation solutions in a context of social and political tensions facing the environmental and health challenges of this Orbiel valley.

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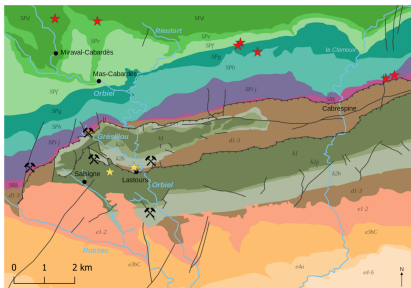
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## Legend

Cenozoic

### Aquitain Basin

- e4-6 - Carcassonne molasse (upper Ypresian - Bartonian)
- e4-a - Ventenac lacustrine limestones (upper Ypresian)
- e3bC - Alveolinids-bearing limestones (lower Ypresian)
- e1-2 - Conglomerates, sand, silt, lacustrine limestones (Paleocene)

Paleozoic

### Minervois nappe

- d1-3 - Indian bead limestones, black limestones, gray dolomites, white limestones, flinty limestones (lower Devonian)
- k2b - Archaocyathids-bearing dolomitic limestones (lower Cambrian)
- k2a - Alternating sandstones and archaeocyathids- and trilobites-bearing limestones (lower Cambrian)
- k1 - Marcory sandstones (lower Cambrian)

### Axial zone

#### Roc-Suzadou group

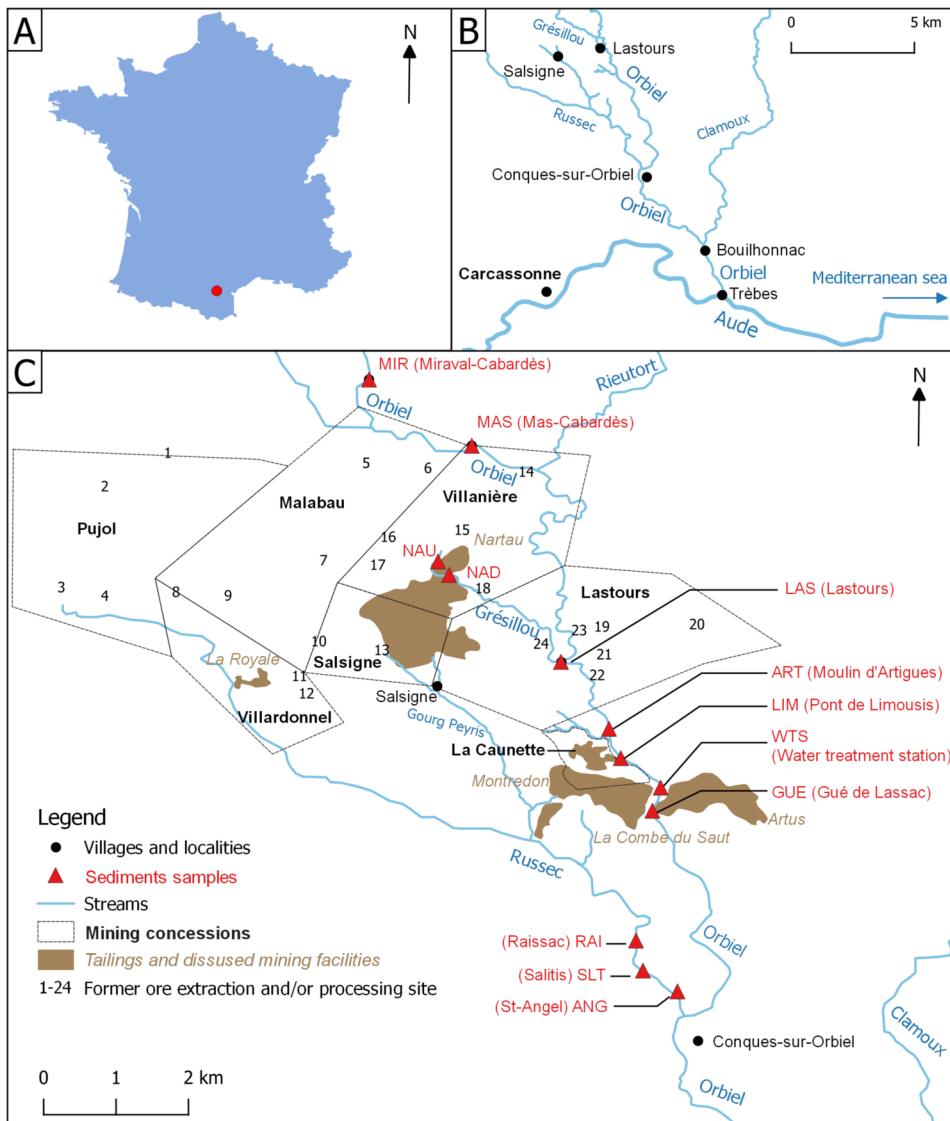
- SRk - Conglomerates, sandstones, quartzites (upper Ordovician - Silurian)

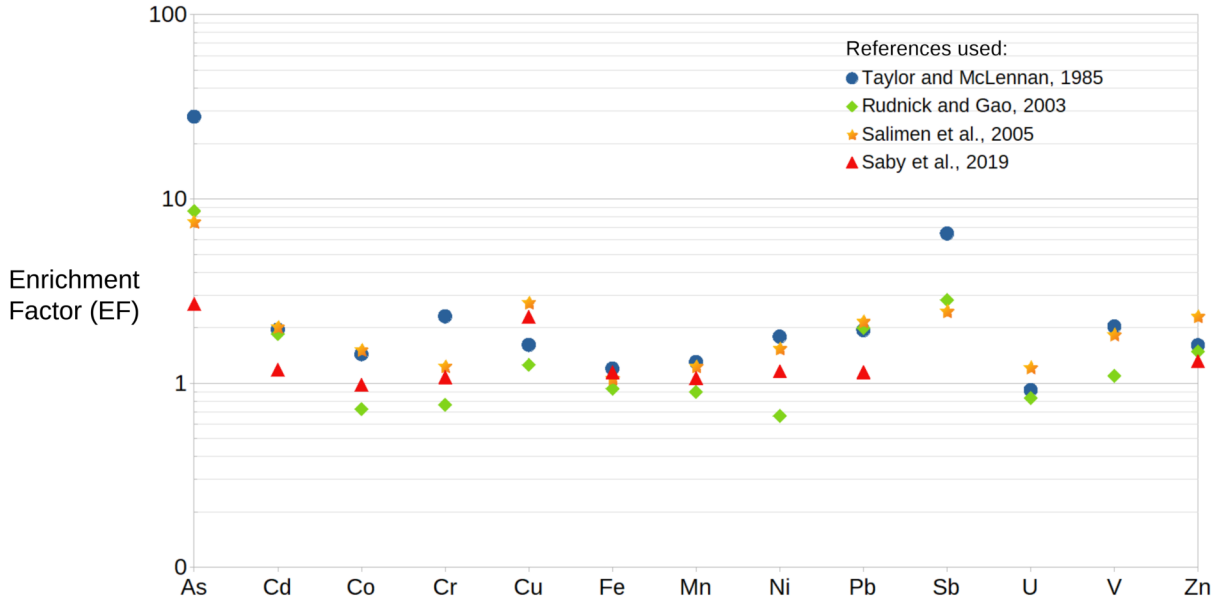
#### Saint-Pons - Cabardès group

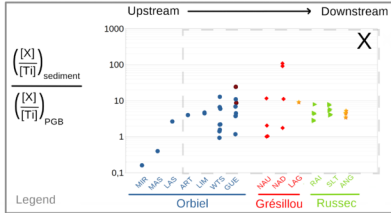
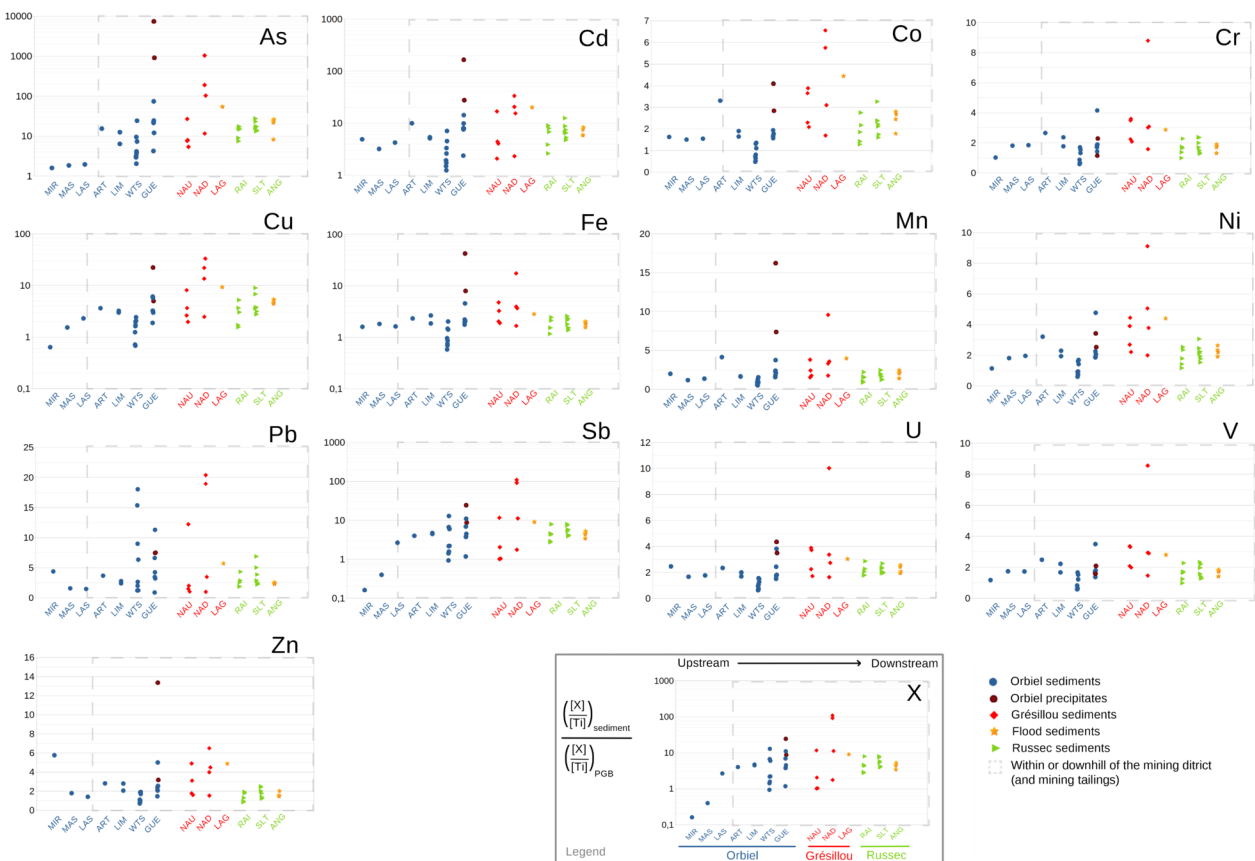
- SPI-j - Black schists (Cambrian - Ordovician)
- SPh - Clastic sandstones (Cambrian - Ordovician)
- SPg - Black schists and clastic sandstones, banded schists (Cambrian - Ordovician)
- SPf - Biotite-bearing schists and quartzites (Cambrian - Ordovician)
- SPe - Pelitic sandstones formation of Mas-Cabardès (Cambrian - Ordovician)
- SPd - Metapelitic formation of Miraval-Cabardès (Cambrian - Ordovician)

### Environment, Structure and Sampling

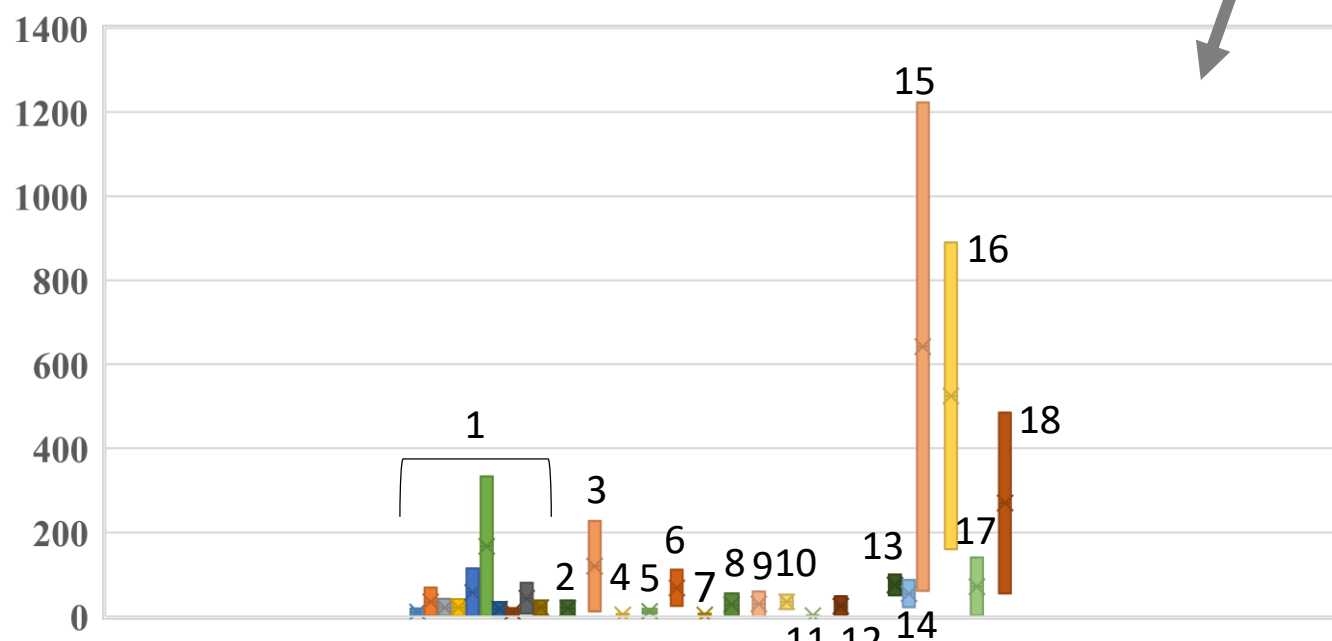
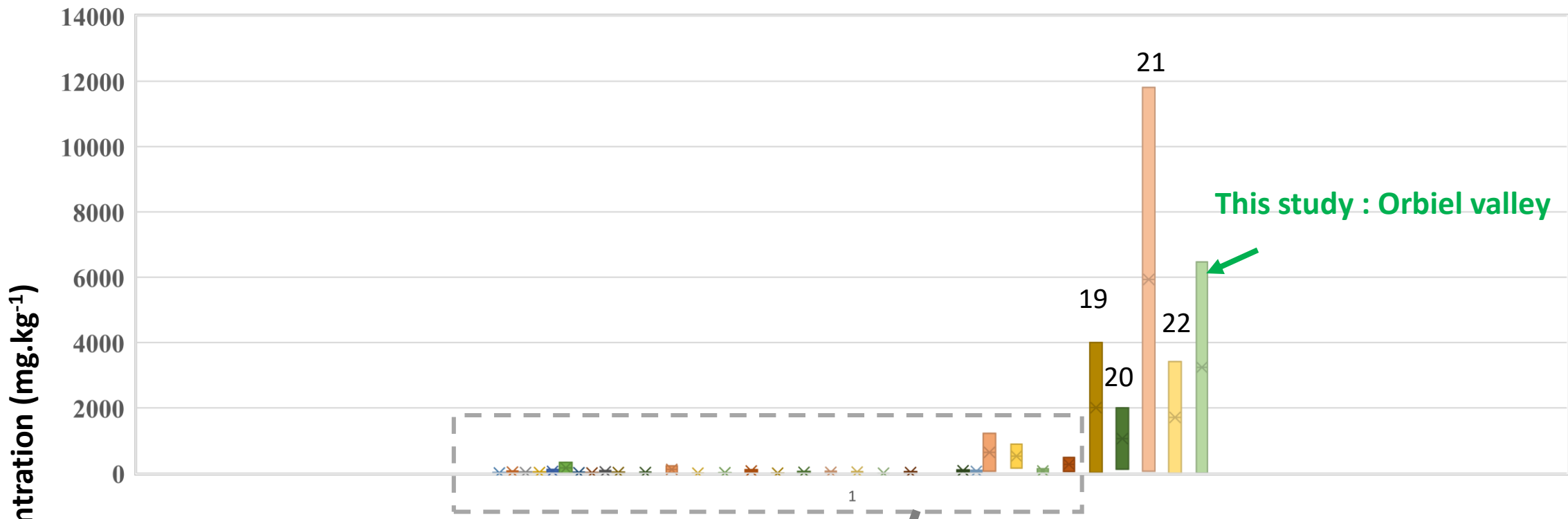
- Streams
- Faults
- Thrust faults
- Disused mines
- ★ Soils sampled (geochemical background)
- ☆ Soils sampled (for comparison)







- Orbiel sediments
- Orbiel precipitates
- Grésillou sediments
- ★ Flood sediments
- ▲ Russec sediments
- Within or downhill of the mining district (and mining tailings)



**Figure 5:** Comparison of As concentration ranges between this study (Orbiel Valley) and other environments: preserved (rivers, deltas, lakes) [1-14], or impacted by mining activities [15-22].

**Table 1 : Mean concentrations in metal(oid)s in soils sampled in the Orbiel valley**

	Samples	As*	Cd*	Co	Cr	Cu	Fe	Mn*	Ni	Pb*	Sb*	Ti	U*	V*	Zn*
<b>Uphill of the mining district (pedo-geochemical background)</b>	<b>Spd</b> (n = 5)	35.6	0.15	12.9	79.6	38.4	3.68	0.07	33.6	27.7	0.32	0.37	2.66	117.3	106.43
	<b>Spe</b> (n = 4)	42.6	0.16	18.9	94.4	48.8	5.09	0.07	43.1	31.1	0.42	0.47	2.72	136.5	153.53
	<b>Spf</b> (n = 2)	47.4	0.30	12.4	77.6	70.2	4.09	0.08	28.2	43.4	1.30	0.41	2.13	117.0	83.75
	<b>Spg</b> (n = 2)	45.0	0.35	5.2	68.5	29.2	3.93	0.03	18.3	61.9	1.64	0.38	2.16	103.6	73.32
	<b>Sph</b> (n = 2)	75.1	0.27	14.1	76.8	23.7	3.69	0.16	27.9	42.6	0.91	0.46	2.18	114.9	90.45
	<b>Spi</b> (n = 2)	47.2	0.18	4.1	89.4	33.2	4.01	0.02	17.0	46.4	6.25	0.50	2.91	131.4	67.68
	<b>Srk</b> (n = 2)	43.0	0.14	2.6	84.2	20.0	2.29	0.02	11.0	31.6	3.68	0.45	2.53	124.3	44.82
<b>Within the mining district</b>	<b>k1</b> (n = 2)	88.1	0.31	19.7	82.3	37.1	4.58	0.16	41.6	71.7	4.60	0.50	1.99	114.1	127.21
	<b>k2b</b> (n = 2)	162.6	1.96	8.3	34.8	44.8	2.81	0.16	27.0	162.8	4.60	0.20	1.70	61.2	232.98

Concentrations are given in mg.kg<sup>-1</sup> except for Fe, Mn and Ti, which are given in %. n corresponds to the number of horizons collected for each soil and used for the mean calculation. \* is displayed when there is a statistical difference (Kruskall-Wallis test, p-value : 0.05) between soils located uphill and within the mining district. For the localisation of the samples refer to Figure 1. Elements are presented in alphabetical order.



**Table 3 : Pedo-geochemical background in the Orbiel valley.**

Element	Unit	Pedo-geochemical background (Orbiel valley)	Baselines	
			France <sup>1</sup>	Europe <sup>2</sup>
<b>As</b>	mg.kg <sup>-1</sup>	44 ± 12	12	6
<b>Cd</b>	mg.kg <sup>-1</sup>	0.18 ± 0.06	0.11	0.09
<b>Co</b>	mg.kg <sup>-1</sup>	13 ± 10	10	9
<b>Cr</b>	mg.kg <sup>-1</sup>	84 ± 16	51	62
<b>Cu</b>	mg.kg <sup>-1</sup>	38 ± 16	12	14
<b>Fe</b>	%	3.90 ± 1.37	2.49	3.75
<b>Mn</b>	%	0.07 ± 0.03	0.05	0.06
<b>Ni</b>	mg.kg <sup>-1</sup>	28 ± 18	21	22
<b>Pb</b>	mg.kg <sup>-1</sup>	38 ± 21	23	17
<b>Sb</b>	mg.kg <sup>-1</sup>	0.9 ± 1.2	-	0.5
<b>Ti</b>	%	0.44 ± 0.09	0.42	0.57
<b>U</b>	mg.kg <sup>-1</sup>	2.6 ± 0.3	-	2.0
<b>V</b>	mg.kg <sup>-1</sup>	126 ± 16	-	63
<b>Zn</b>	mg.kg <sup>-1</sup>	92 ± 44	59	47

Mean concentrations in soils (baselines) are reported at the French and European scales are reported for comparison : <sup>1</sup> Saby et al. (2019) ; <sup>2</sup> Salimen et al. (2005).



**Table 4:** Geo-accumulation index have been calculated according to the formula proposed by Müller (1969)

	As	Cd	Co	Cr	Cu	Fe %	Mn ‰	Ni	Pb	Sb	Ti ‰	U	V	Zn
MIR-1	-1.9	-0.3	-1.9	0.3	-3.3	-1.9	-1.6	-2.2	0.3	-4.8	0.6	-1.4	-2.5	0.1
MAS-1	0.4	1.2	0.2	1.0	0.2	0.4	-0.1	0.7	1.0	-1.3	2.8	0.2	0.2	0.6
LAS-1	0.6	1.7	0.3	1.0	0.8	0.3	0.1	0.9	1.0	1.5	2.8	0.3	0.3	0.3
ART-1	2.9	2.3	0.7	1.6	0.8	0.2	1.1	0.9	1.6	1.4	2.2	0.1	0.1	0.7
LIM-1	3.0	1.7	0.3	1.6	1.0	0.7	0.1	0.8	1.6	1.9	2.6	0.2	0.3	1.0
LIM-2	2.2	1.8	0.3	1.6	1.1	0.4	0.3	0.7	1.6	2.2	2.7	0.2	0.1	0.8
WTS-1	2.2	1.5	0.1	1.6	0.8	0.5	0.5	0.7	1.6	1.5	3.8	0.4	0.1	0.9
WTS-2	1.5	1.2	-0.1	1.6	0.4	0.0	-0.1	0.4	1.6	3.6	2.7	0.0	0.0	0.6
WTS-3	2.5	1.8	-0.2	0.7	0.6	0.1	-0.5	0.4	0.7	1.2	2.8	0.1	0.1	0.6
WTS-4	1.7	0.9	0.0	2.4	0.0	0.1	-0.3	0.5	2.4	2.2	3.8	0.0	-0.1	0.6
WTS-6	2.5	1.5	-0.1	5.7	0.5	0.2	0.2	0.5	5.7	1.3	4.2	0.2	0.0	0.7
WTS-7	2.8	2.3	0.7	5.9	1.7	0.7	1.1	1.1	5.9	2.0	4.2	0.9	0.1	1.2
WTS-8	4.1	2.3	-0.1	2.9	0.7	0.5	0.2	0.5	2.9	2.5	2.7	-0.3	-0.4	0.6
WTS-9	3.8	1.4	-0.3	4.5	1.3	0.1	-0.2	0.3	4.5	3.7	3.8	0.1	-0.3	0.6
GUE-1	3.1	2.5	0.4	2.0	1.1	0.6	0.6	0.8	2.0	2.1	2.7	0.3	0.2	1.1
GUE-2 *	-0.1	4.4	-0.9	0.7	1.5	2.4	1.1	-0.9	0.7	2.1	0.2	-1.0	-2.5	1.0
GUE-3 *	-1.0	3.9	0.6	2.8	1.4	2.1	2.0	0.7	2.8	2.7	2.3	0.8	0.0	1.0
GUE-4	3.9	2.4	0.4	2.9	2.0	0.5	0.7	0.8	2.9	2.6	2.6	0.5	0.0	0.8
GUE-5	2.0	1.2	0.7	0.6	0.8	0.8	0.7	1.1	0.6	0.6	3.2	0.4	0.5	0.7
GUE-6	3.8	3.0	1.3	3.5	1.7	1.4	1.2	1.7	3.5	3.1	2.4	1.0	0.9	1.7
GUE-7	-4.4	2.6	0.2	2.2	1.0	0.4	0.7	0.5	2.2	1.7	2.6	0.1	-0.3	0.6
GUE-8	4.1	2.6	0.4	2.3	1.4	0.7	0.6	0.9	2.3	2.0	2.9	0.4	0.2	1.0
NAU-1	2.0	1.1	1.0	0.8	0.9	0.7	0.4	1.4	0.8	0.5	2.2	0.8	0.6	0.9
NAU-2	2.7	0.8	1.0	1.1	1.2	0.8	0.5	1.5	1.1	0.2	3.0	0.8	0.7	0.8
NAU-3	3.3	2.6	0.4	2.9	1.5	0.8	0.5	0.7	2.9	2.5	1.7	0.3	0.1	1.0
NAU-4	2.1	1.7	0.8	0.5	0.6	0.6	0.6	1.1	0.5	0.2	2.9	0.3	0.5	0.6
NAD-1	-4.3	2.9	0.7	1.6	4.0	0.9	0.9	1.2	1.6	2.9	2.2	0.3	0.4	1.4
NAD-2	3.6	1.2	0.8	0.8	1.3	0.8	0.9	1.3	0.8	1.3	3.3	0.6	0.4	0.9
NAD-3	-2.2	2.1	0.5	2.8	2.2	1.9	1.1	1.2	2.8	4.7	1.0	1.0	0.7	0.7
NAD-4	-3.7	3.8	1.3	3.9	2.5	0.7	0.5	1.3	3.9	5.9	1.9	0.4	0.1	0.9
LAG-1 **	4.9	3.4	1.3	2.4	2.3	0.6	-2.2	1.5	2.4	2.7	2.3	0.6	0.4	1.6
RAI-1	2.5	1.4	-0.2	1.5	0.2	-0.1	-0.7	0.1	1.5	2.1	1.9	0.0	-0.3	-0.3
RAI-2	3.2	2.1	0.6	1.4	1.5	0.2	0.3	0.7	1.4	1.7	2.3	0.2	-0.2	0.3
RAI-3	3.1	2.3	0.1	1.3	1.0	-0.2	-0.1	0.2	1.3	1.8	2.4	0.1	-0.3	-0.3
RAI-4	2.5	0.9	0.1	1.2	0.2	0.2	-0.6	0.3	1.2	1.4	2.8	0.3	-0.3	-0.3
RAI-5	2.5	1.2	-0.3	1.0	0.0	-0.5	-0.6	-0.2	1.0	1.3	2.5	0.0	-0.9	-0.7
SLT-1	2.4	1.1	-0.4	1.9	0.2	-0.3	-0.6	-0.1	1.9	1.8	1.6	-0.6	-0.6	-0.1
SLT-2	3.8	2.4	0.5	1.5	2.0	0.1	0.3	0.6	1.5	2.1	2.5	0.1	-0.3	0.4
SLT-3	3.1	1.5	0.0	1.2	0.7	-0.2	-0.4	0.2	1.2	1.7	2.5	0.1	-0.5	-0.2
SLT-4	3.1	1.7	0.1	1.4	0.9	-0.1	-0.3	0.3	1.4	1.8	2.5	0.2	-0.5	-0.2
SLT-5	2.8	1.9	0.2	1.8	0.5	0.3	-0.1	0.5	1.8	2.0	2.3	0.4	0.0	0.1
SLT-6	3.2	2.1	0.2	1.5	1.6	-0.5	-0.2	0.3	1.5	1.8	1.6	-0.4	-0.7	-0.1
SLT-7	3.5	2.1	0.5	1.4	1.3	0.0	0.2	0.6	1.4	1.8	2.6	0.3	-0.3	0.1
ANG-1 **	4.2	2.6	1.0	1.6	1.9	0.4	0.7	0.9	1.6	2.3	2.7	0.4	0.1	0.7
ANG-2 **	4.0	2.3	0.9	1.5	1.8	0.4	0.8	1.0	1.5	2.0	2.6	0.6	0.1	0.2
ANG-3 **	4.4	2.5	1.3	1.9	2.1	0.9	1.1	1.4	1.9	2.5	3.1	1.1	0.6	0.7
ANG-4 **	2.3	1.8	0.1	1.3	1.4	-0.1	-0.2	0.4	1.3	1.4	2.4	0.1	-0.4	0.0

Value	Class	
0	0	uncontaminated
0 et 1	1	uncontaminated to moderately contaminated
1 et 2	2	moderately contaminated
2 et 3	3	moderately to strongly contaminated
3 et 4	4	strongly contaminated
4 et 5	5	strongly to extremely contaminated
> 5	6	extremely contaminated