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Characterizing multiple water-rock interactions in the critical zone through Sr-isotope tracing of surface and groundwater.

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Abstract

Here, we report on Sr isotopes, from shallow and deep groundwater from catchments located on granite and schist around the world. This extensive approach of Sr isotope tracing, initiated in France on areas impacted by intensive agricultural activities, was enlarged to Africa (granite-gneiss and schists 2200-700 Ma of the Congo Basin; Archaean granitoid/gneisses and sedimentary greenstone 3700-2500 Ma of the Orange River); India and Nepal (Archean granites 2500 Ma and Palaeoproterozoic granodiorite and schists 3100-1600 Ma for India; Himalaya metamorphic, silicate metasediments and gneisses 630 to 490 Ma); North America (3500 to 2500 Ma silicate rocks in the Mackenzie basin; 1000 to 70 Ma silicate rocks in the Fraser Basin in Canada and 1760 to 1430 Ma anorthosite and granite in the Laramie aquifer in Wyoming); South America in French Guiana (Archean gneiss 3400 to 2700 Ma and granite-gneiss rocks 2300 to 1900 Ma); Australia (65 Ma arenaceous and argillaceous rocks; and considering both surface and groundwater). In this extensive approach, the Sr and Mg contents are well correlated and both are partly related to agricultural and weathering inputs. The relationship between Sr- isotope and Mg/Sr ratios allows definition of the relative impact on surface and groundwater of processes occurring in the Critical Zone, mainly rain, agricultural practices and water-rock interactions.

Keywords: strontium isotope, critical zone, weathering, regolith

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1 – Introduction

The Critical Zone (CZ) is the lithosphere-atmosphere boundary where complex physical, chemical and biological processes occur and control the transfer and storage of water and chemical elements. This is the place of life-sustaining resources and where rocks release nutrients. Because it is the place where we are living, this is a fragile zone and nowadays a perturbed natural ecosystem. Surface water that drains such environments can interact with groundwater. Either surface-water bodies gain water or solutes from ground-water systems or the surface-water is a source of ground-water recharge, often causing changes in ground-water quality. Surface water and shallow groundwater are particularly vulnerable to pollution, as in many countries water resources are increasingly threatened by growing demand, wasteful use and contamination, while deeper ground resources are more protected from contamination. As climate change impacts are increasingly observed, existing challenges within science and policy are being exacerbated by the rate of change within the physical system (Ostrom, 2007). Understanding the diversity of water-rock interactions is a key issue for ensuring the CZ functioning and the geochemical characterization of different hydrological compartments (weathered rock, the weathered-fissured zone, and fractured bedrock; Larsson, 1984) involved in the water circulations. This understanding will help to predict the CZ evolution in response to environmental modifications due to tectonic, climatic or anthropogenic forcing (Brantley et al., 2007; Banwart et al., 2011).

In crystalline basement rocks; which represent more than 50% of the earth's terrestrial surface (Amiotte Suchet et al., 2003), weathering leads to the formation of a cover of an unconsolidated part named alterite or saprolite, usually representing a thickness of several tens of metres. Considering the catchment scale, the base of the alterite cover can be considered as sub-horizontal, parallel to the paleosurface contemporaneous with weathering (Fig. 1). A fissured zone develops beneath the alterite, representing a thickness of several

hundred metres. The fissuring results from rock shattering under the influence of stress generated by the swelling of certain minerals during the early stages of weathering (Wyns et al., 2004). Moreover, weathering causes the dissolution and flush of soluble cations, depending on the nature and stability of the minerals, which in turn generates mineralogical phase changes, leading in particular to the formation of clayey minerals at the expense of alumina silicates. Thus, although alterite has an interstitial porosity, their clayey mineral content means that their permeability is generally low, especially in fine-grained rocks. Conversely, the global permeability of the fissured zone is generally larger than that of the alterite, even if the permeability decreases statistically with depth in close connection to fissure density.

Water resources in hardrocks commonly involve different hydrogeological compartments such as overlying sediments, weathered rock, the weathered-fissured zone, and fractured bedrock (Larsson, 1984). Aims of the present study are to characterize the strontium isotopic ratios in watersheds or continental basins worldwide draining crystalline basement, to investigate the functioning of the different compartments of the CZ in terms of weathering processes and water pathways.

2 – Samples and methods

First, fields representing different catchments in France are used as summarized in the paper by Négrel and Pauwels (2004) and illustrated in Fig. 2: shallow and deep groundwater from granite and schist of the Armorican Massif in Brittany, northwestern France (Arguenon, Coët-Dan, Pleine-Fougeres, Négrel and Pauwels (2004)); surface and shallow groundwaters from a small watershed underlain by granite or granite-gneiss in the Margeride Mountains, southern Massif Central in France (Négrel, 1999); stream- and spring-waters from the Vosges massif draining granite (NE France; Tricca et al., 1999; Probst et al., 2000; Aubert et al., 2002;

Schaffhauser et al., 2014); water flowing from the Morvan draining granite (SE Paris Basin; Négrel et al., 1988; Seimbille et al., 1989); waters flowing in the upstream part of the Hérault watershed draining granite and mica schist (S France; Petelet, 1998); and water flowing on mica schist from the Cantal (eastern Massif Central; Négrel, 1997). Other sets of samples were used: one corresponding to the Seine River draining granite (Roy et al., 1999); waters flowing in the upstream part of the Garonne River (Semhi et al., 2000); waters flowing upstream in the Pyrénées (Oliva et al., 2004); and waters flowing in the upstream part of the Moselle River (Brenot et al., 2008). These different catchments in France all drain silicate bedrocks (300-450 Ma).

We have extended to larger basins draining silicate rocks and located worldwide as illustrated in Fig. 3 for some of them: India (3a), South America (3b), Africa (3c), North America and Canada (3d), as summarized below for all considered sites:

- In India with two locations and Himalaya (Nepal). The first location in India is the Subarnarekha River Basin, located in northeastern India, rising in southern Bihar state (Négrel et al., 2007a), the river drains Precambrian metasediments, mainly mica schists, and Archean granites aged 2500 Ma. The second location in India is the Maheshwaram hydrosystem (35 km south of Hyderabad in the Ranga Reddy District, State of Andhra Pradesh). This watershed is 53 km² in surface area, purely endorheic and groundwater interact with Palaeoproterozoic granodiorite and schists 3100-1600 Ma (Négrel et al., 2010). In Nepal with the Himalayan rivers (the Bothe Kosi River; Harris et al., 1998) draining sillimanite-grade migmatites, gneisses and leucogranites in the upper section and kyanite–garnet metapelitic sediments in the lower sections, mid-Palaeozoic to Eocene age (450 – 50 Ma) and the Ganges headwaters (Bickle et al., 2005) that cross the major lithotectonic units of the

Himalaya (metamorphic units of the Himalayas and the silicate metasediments and gneisses of High Himalayan Crystalline Series 630 – 490 Ma in age).

- In South America with the French Guiana where the Maroni Basin (Négre and Lachassagne, 2000) and groundwater over the whole country were investigated (Négre and Petelet-Giraud, 2010) where weathered rocks are Archean gneiss aged 3400-2700 Ma and granite-gneiss rocks aged 2300-1900 Ma.
- In Africa with the Congo Basin with the Oubangui River (Négre and Dupré, 1995a), the Congo River (Négre et al., 1993; Négre and Dupré, 1995b) and rivers draining the western part of the Congo Basin (Négre and Dupré, 1995a). These rivers drain mainly granite-gneiss and schists, 2200-700 Ma in age. Also, in South Africa with the Orange River Basin (de Villiers et al., 2000) using the Vaal River catchment area that is underlain by granitoid/gneisses and volcano-sedimentary greenstone of the Archean Kaapvaal Basement (3700-2500 Ma) belt and below the Vaal confluence where the Orange River traverses the Namaqualand Basement Complex, lower Karoo sequences and Kalahari sands.
- In North America with the Mackenzie and Fraser Basin in Canada and the Laramie aquifer (Wyoming USA, Frost and Toner, 2004). The Mackenzie River (Slave Province, Millot et al., 2002) where rivers drain 213,000 km² of old crystalline basement from 3500 to 2500 Ma with silicate rocks predominating (potassium-rich rocks, gneisses) and the Fraser River (British Columbia, Cameron et al., 1995) draining 238,000km² and crossing through a number of geological belts (Upper Proterozoic to tertiary -1000 to 70 Ma - metamorphic and I- or S-type granitic rocks, silicate sedimentary rocks). The aquifer in the Laramie basin (Wyoming) crosses Precambrian igneous and metamorphic rocks (1760 Ma Horse Creek Anorthosite

Complex with anorthosite, monzonite, and granite, 1430 Ma Sherman Granite Batholith).

- Central Australia with groundwater from Tertiary sedimentary aquifers of the arid Ti-Tree Basin (Harrington and Herczeg, 2003). The basin is composed of a suite of arenaceous and argillaceous Cenozoic rocks (65 Ma) derived through weathering of the Proterozoic igneous and metamorphic rocks also including granite, gneiss, schist and quartzite (2500 to 540 Ma).

3 – Additional results in this study

All data compiled/reported in this paper have been already published except those from the Maheshwaram catchment in India. In this small endorheic watershed, representative of South Indian conditions in terms of geology (hard rock), land-use, overpumping of the hard-rock aquifer (over 700 bored open-end wells in use, surface waters are ephemeral and occur only during the monsoon period. Moreover, the chemical composition of groundwater is strongly impacted by evaporation linked to irrigation return flow (Perrin et al., 2011). Samples in the Maheshwaram hydrosystem (Fig. 4) were collected and analyzed in two sampling periods 2006 and 2008 (Négre et al., 2010; Pauwels et al., 2015). The 2006 and 2008 samples were analyzed for Mg contents by ICP-AES and Sr content by ICP-MS at BRGM. The 2006 samples were analyzed for Sr isotopes at National Geophysical Research Institute NGRI, Hyderabad (A. Dayal, personal communication) whereas the 2006 samples were analyzed at Laboratoire d'HYdrologie et de GÉochimie de Strasbourg LHyGeS. The Sr was separated using the classical procedure purification using ion exchange resins and the isotopic ratios were determined by thermal-ionisation mass spectrometer with an analytical uncertainty of around 20ppm ($2\sigma_{\text{mean}}$). The $^{87}\text{Sr}/^{86}\text{Sr}$ value of the NBS 987 standard was routinely measured in both labs. For more details, see Prunier et al., 2015.

Strontium content varies greatly in groundwater samples between 2.3 and 33.1 $\mu\text{mol/l}$. The range is between 2.3 and 33.10 $\mu\text{mol/l}$ in 2006 and between 4.6 and 17.1 $\mu\text{mol/l}$ in 2008. The observed range partly agrees with that found in groundwater in the Subarnarekha River basin with Sr contents from less than 3.4 up to 100.5 $\mu\text{mol/l}$ (Négrelet et al., 2007a). Strontium isotopes display a wide variation in the groundwater from 0.71606 (lower values both in 2006 and 2008) to 0.72843 observed in 2008 (0.72823 as higher value in 2006). The $^{87}\text{Sr}/^{86}\text{Sr}$ ratios observed in groundwater agrees with those of waters draining old cratons (Guyana Shield, Edmond et al., 1995; French Guiana, Négrelet and Lachassagne, 2000; Lena River Basin in Russia, Huh and Edmond, 1999; Ganga-Brahmaputra-Indus River system in India, Krishnaswamiet al., 992; Pande et al., 1994; Singh et al., 1998; Dowling et al., 2003; Himalayan rivers, Galy et al., 1999).

4 - Discussion

4.1. The Sr isotope data from shallow and deep groundwater in silicate over France

Strontium isotopes are useful tracers of water-rock interaction (Blum et al., 1994; Brantley et al., 1998; Négrelet et al., 2001) and of mixing of water from different origins (Roy et al., 1999; Négrelet et al., 2001; Mc Nutt et al., 1990). Part of the Sr mass balance in a crystalline environment is controlled by chemical weathering of the most commonly Sr-bearing phases from the silicate host rock, either primary phases (apatite, plagioclase, K-feldspar, biotite and muscovite) or their alteration products, e.g. epidote and clay minerals (Blum et al., 1994; Brantley et al., 1998; Mc Nutt et al., 1987; Aubert et al., 2002; Pett-Ridge et al., 2009), inputs from rainwater and anthropogenic disturbance. The latter, through agricultural activity, induces an input of Sr that may be important (Négrelet, 1999; Roy et al., 1999) depending on the range of Sr contents in fertilizers and carbonate additives (3–1500 $\mu\text{g/g}$) with a Sr-isotope ratio from 0.70794 to 0.7095. This range is reflected in manure from livestock, such as hogs and poultry (Sr range from 2 to 4 $\mu\text{g/g}$, $^{87}\text{Sr}/^{86}\text{Sr}$ ratio from 0.7092 to 0.71092)

Starting from data of groundwater collected in three granitic and Brioverian ‘schist’ aquifers of the Armorican Massif, the observed Sr content is well correlated with Mg and both are partly related to agricultural practices and water rock interaction (Fig. 5, Négrel and Pauwels, 2004). Combined with additional data from surface and groundwater of catchments draining silicate bedrocks (300-450 Ma) like the Hérault, Seine, Moselle, Garonne, Morvan, Margeride, Cantal, Pyrenees and Vosges (from the literature, Fig. 5) revealed a unique trend between Sr and Mg expressed by the relationship $Mg (\mu\text{mol/l}) = A \times Sr (\mu\text{mol/l})$; with A varying between 100 and 600 (Fig. 5). As demonstrated before (Négrel and Pauwels, 2004), Mg is closely linked with Sr (and Cl) and therefore the variability in the Cl vs. Mg relationship interpreted as variations in the agricultural practices from a non-polluted status (e.g. the water rock interaction), enhanced the use of Mg/Sr ratios. The use of this ratio rather than absolute concentrations alone, avoids variations due to dilution or concentration effects and the normalization to the Sr content transforms mixing curves into straight lines (Faure, 1986).

The relationship between Sr-isotope and Mg/Sr ratios (Fig. 6) allows better definition of the different endmembers, mainly rain, agricultural practices and water-rock interaction. Actually, the 3 endmembers rain inputs, fertilizer and manure present very close characteristics with low $^{87}\text{Sr}/^{86}\text{Sr}$ and low Mg/Sr ratios (Négrel, 1999; Négrel et al., 2007b). Two binary mixing trends connect fertilizer and manure and the isotope composition of the endmembers reflecting the water rock interaction in the different compartments. This implies the occurrence of (i) high $^{87}\text{Sr}/^{86}\text{Sr}$ and low Mg/Sr ratios, and (ii) low $^{87}\text{Sr}/^{86}\text{Sr}$ and high Mg/Sr ratios endmembers, which could be related to different water rock interaction conditions. Water originating from the upper compartment of the aquifer in weathered rock (alterite) is one of the water rock interaction endmembers. The interaction of water with alterite thus led to higher $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in the water because of the weathering of residual

minerals (Bullen et al., 1997; Négrel, 1999). It is worth noting that the Vosges waters may represent partly such a compartment with regard to the $^{87}\text{Sr}/^{86}\text{Sr}$ and Mg/Sr ratios. Water from the lower compartment of the aquifer, mainly comprising fissured fresh rock, constitutes the second water rock interaction endmember. Interaction in the fissured part implies that the Sr-isotope characteristics of waters are more related to the weathering of whole rock (e.g. a lower $^{87}\text{Sr}/^{86}\text{Sr}$ ratio: Négrel, 1999; Négrel and Pauwels, 2004).

All ground- and surface waters for catchments draining silicate bedrocks are scattered between at least three geochemical signatures with some particular points plotting close to the different mixing lines. This is the case for the Garonne, Seine, Pyrénées and to a lesser degree for the Morvan, Cantal and Vosges (Négrel et al., 1988; Roy et al., 1999; Probst et al., 2000; Aubert et al., 2002; Pierret et al., 2014), and Margeride field (Négrel, 1999), which are in an intermediate location along the mixing line between fertilizer and manure and the endmember corresponding to alterite. The intermediate location suggests influences from either agricultural practices (Négrel, 1999; Roy et al., 1999) or rain, the latter being less probable. Regarding Brittany, high Mg/Sr samples from the Coet-Dan catchment plot along the mixing line between fertilizer and manure and the endmember corresponding to the fissured part of the aquifer, comprising fresh rock whereas high Mg/Sr samples from the Plaine-Fougères and Arguenon catchments plot along and/or near the mixing line between the two WRI endmembers. These locations reflect the relative lack of anthropogenic contribution to Sr content compared to water-rock interactions and, in the case of Arguenon, the occurrence of waters corresponding to a mixing between upper and lower compartments of the aquifer. The Plaine-Fougères field defines a linear trend towards the pollution endmember, with the low $^{87}\text{Sr}/^{86}\text{Sr}$ and Mg/Sr ratios of this field in agreement with the characteristics of the Hérault upstream waters (Petelet, 1998), thus reflecting considerable anthropogenic input in the upper part of the Hérault catchment as suggested by Petelet (1998). The Vosges (Probst et al., 2000;

Aubert et al., 2002; Pierret et al., 2014; Schaffhauser et al., 2014) partly agree with the alterite endmember but some waters are marked by the endmember corresponding to the fissured part of the aquifer as they plot near the mixing line between the upper and lower parts of the aquifer.

4.2. Extensive approach using Sr isotope in surface and groundwater in the critical zone

4.2.1 The focus on the Maheshwaram hydrosystem

The fluctuations of the Sr contents in the endorheic Maheshwaram hydrosystem are correlated with those of Cl by the equation $Cl = 0.14 \times Sr - 36.06$ ($N = 39$, $R^2 = 0.67$). This relationship suggests that the source of chloride is also a potential source of Sr that can be related to agricultural practices (Atal et al., 2010) as the bedrock is only comprised of silicates (e.g., granite).

The Sr-isotope studies of water have shown that variations in Sr isotopes are primarily caused by mixing of waters of various origins with different $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and Sr contents such as mixing between different water pools, each of them possibly reflecting water-rock interaction with different rock types or different stage of water rock interaction in the same bedrock. A comparison of the $^{87}\text{Sr}/^{86}\text{Sr}$ vs. Cl and Sr contents shows that all the water samples from the Maheshwaram hydrosystem are largely scattered with regard to both Sr isotopes and Sr concentrations. Comparing the Sr isotope ratios and the chloride contents argues in favour of three endmembers as samples are scattered in three directions (Fig. 7a). Rainwater inputs display low Cl contents in the Maheshwaram hydrosystem, around $113 \mu\text{mol/l}$ (Négrel et al., 2011) and anthropogenic inputs show the highest Cl contents, up to $7042 \mu\text{mol/l}$, and the natural background (e.g. the water rock interaction) has a low Cl content, near that of rainwater. The most dilute endmember in Fig. 7a could be rainwater inputs for which characteristics are usually a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio ranging between 0.714 to 0.720 (Galy et al., 1999; Négrel et al., 2007a), and Cl contents around 113 to $563 \mu\text{mol/l}$. With a mean weighted

chloride content of 25 $\mu\text{mol/l}$ in rainwater and the concentration factor related to the quantity of precipitation and the evaporation rate, a maximum for rain-derived Cl of 225 to 563 $\mu\text{mol/l}$ was determined (Négrel et al., 2011), in agreement with the range observed for this endmember. The most mineralised endmember has a broader range of Sr isotope ratios from 0.71609 to 0.72769 and Cl contents in the range 7296-8028 $\mu\text{mol/l}$. This second endmember is believed to correspond to anthropogenic inputs, mainly agricultural in origin (livestock farming, fertilizer spreading and partly related to domestic sewage and in situ sanitation). The third endmember is characterised by a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio ranging between 0.72796 and 0.72843 and a Cl content around 563 - 1408 $\mu\text{mol/l}$. This range of Cl contents agrees with the rain-derived Cl as stated above and corresponds to water rock interaction.

A comparison of the $^{87}\text{Sr}/^{86}\text{Sr}$ vs. Sr contents shows that all the water samples from the Maheshwaram hydrosystem are largely scattered with regard to both Sr isotopes and Sr contents. It is worth noting that (1) the large variation in the Sr contents corresponds to the highest $^{87}\text{Sr}/^{86}\text{Sr}$ ratios and (2) the relatively restricted range of the Sr contents in the lowermost values corresponds to large variations in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios. Illustrated in a diagram of $^{87}\text{Sr}/^{86}\text{Sr}$ ratios vs. $1/\text{Sr}$ (Fig. 7b), the endmembers defined in Fig. 7a cannot be fully used. Two endmembers can be considered, meaning the endmember corresponding to anthropogenic inputs with high Sr contents and variable $^{87}\text{Sr}/^{86}\text{Sr}$ ratio and the endmember corresponding to water rock interaction with a high $^{87}\text{Sr}/^{86}\text{Sr}$ ratio and large Sr content variations. The endmember previously related to rain inputs cannot be considered here as the Sr contents in the groundwater are too large compared to the values for the rain inputs (Sr content around $1\mu\text{g/l}$, Négrel et al., 2007a). This implies the existence of a second endmember corresponding to water-rock interaction with a restricted range in Sr content and a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio close to 0.715 (Fig. 7b). Such variations in the $^{87}\text{Sr}/^{86}\text{Sr}$ vs. Sr contents in groundwater

issued from a similar granitic catchment, and related to changes in water-rock interactions, have been observed by Négrel et al. (2000).

4.2.2 The extensive approach by enlarging to worldwide basins and catchments

An extensive approach was done by enlarging to Africa (granite-gneiss and schists 2200-700 Ma of the Congo Basin; Négrel et al. 1993; Archaean granitoid/gneisses and sedimentary greenstone 3700-2500 Ma of the Orange River; de Villiers et al., 2000); India and Nepal (Archean granites 2500 Ma and Palaeoproterozoic granodiorite and schists 3100-1600 Ma, Négrel et al., 2007a and this study for India; Himalaya metamorphic, silicate metasediments and gneisses 630 to 490 Ma; Harris et al., 1998; Bickle et al., 2005); North America (3500 to 2500 Ma silicate rocks in the Mackenzie basin; Millot et al., 2002 and 1000 to 70 Ma silicate rocks in the Fraser Basin in Canada; Cameron et al., 1995 and 1760 to 1430 Ma anorthosite and granite in the Laramie aquifer in Wyoming; Frost and Toner, 2004); South America in French Guiana (Archean gneiss 3400 to 2700 Ma and granite-gneiss rocks 2300 to 1900 Ma; Négrel and Petelet-Giraud, 2010); Australia (65 Ma arenaceous and argillaceous rocks; and considering both surface and groundwater; Harrington and Herczeg, 2003).

In a way identical to observations made at the scale of France, the Sr content is well correlated with Mg as illustrated in Fig. 8, when enlarging to worldwide basins and catchments. The highest values are observed in India in the Maheshwaram watershed (this study) where more than 700 boreholes are used for agricultural irrigation in this endorheic watershed of 53 km² surface area. The second place in India (the Subarnarekha River Basin), also with high agricultural activities, has high Sr-Mg contents (Négrel et al., 2007a), however with a lower range than that observed in the Maheshwaram watershed. Also, in South Africa with the Orange River Basin the high contents in these elements are related, according to de Villiers et al. (2000), to agricultural practices. The Ti-Tree Basin in Central Australia also has groundwater with high Sr-Mr contents. As this is a very arid part of the Australian country

with a ratio of mean annual rainfall over potential evaporation of less than 0.1 (Harrington and Herczeg, 2003), the groundwater resource is intensively used for a small but lucrative horticulture industry, thus inducing the high Sr-Mg contents. Contrary to the zones where the high Sr-Mg contents are related to larger agricultural practices, the lowest values are observed in French Guiana and Africa (Congo and Congo occidental basin, Oubangui) where rain inputs and water rock interactions are predominant processes. The rest of the studied sites are scattered between low and high Sr-Mg contents as representing a mix between rain inputs, water-rock interaction and anthropogenic inputs.

The weathering processes studied here all focus on old silicate environments, with ages up to 3100 Ma, such weathered silicates therefore yield clearly larger ranges in the Sr-isotope ratios as illustrated in Fig.9 when compared to the Mg/Sr ratio. The lowest $^{87}\text{Sr}/^{86}\text{Sr}$ value being observed in the Congo Basin (Kouilou river, Négrel and Dupré, 1995a, b), in French Guiana (Négrel and Lachassagne, 2000; Négrel and Petelet-Giraud, 2010) and in the Fraser River basin (Cameron et al., 1995). The higher $^{87}\text{Sr}/^{86}\text{Sr}$ value (up to 0.765) being observed in Australia (Harrington and Herczeg, 2003) as other high values are observed in in the Ganga headwaters (0.7526 and 0.7489, Bickle et al., 2005), in the Fraser river Basin (0.7513, Cameron et al., 1995) and in groundwater from Finland draining tills developed on granites (0.7508, Négrel et al., 2005). The Sr-isotopes tracing the origin of dissolved compounds define and identify the circulation pathways of waters and compare to the former plot with the French sites, the additional data define a large scatter for the Sr- isotope ratios.

More detail in Fig. 9, the scattering of the data sets from *India* is between alterite and rain or agricultural practices for Maheshwaram site and between the three endmembers (alterite, underlying weathered-fissured and fractured bedrock and rain or agricultural practices) for the Subarnarheka site reflecting the larger influence of deep alteration processes in the water in this area (Négrel et al., 2007a). For the Himalayan rivers, the Bothe Kosi River

(Harris et al., 1998) has scattered Sr-isotope ratios ranging between 0.7187 and 0.7409 and a low Mg/Sr (145 – 208) that reflects a mixing between the alterite compartment and agricultural practices (also reflected by the range of Sr – Mg contents). The Ganges headwaters (Bickle et al., 2005) are scattered between the three endmembers, one sample with high Sr – Mg content displays the lowest $^{87}\text{Sr}/^{86}\text{Sr}$ ratio reflecting a large contribution of the anthropogenic endmember. The other samples reflect the influence of the alterite endmember with increasing $^{87}\text{Sr}/^{86}\text{Sr}$ ratio (up to 0.7526) combined with Mg/Sr ratio in the range 120 – 220. One sample appears to be a mix between the alterite compartment and underlying weathered-fissured and fractured bedrock as the sample plots along the mixing line between these two endmembers with a Mg/Sr ratio of 470 and a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.7489.

In *Africa*, the scattering of the surface water data in the Congo Basin lies between the three endmembers: rain inputs as previously evidenced by Négrel et al. (1993, 1995a, b) but also the alterite compartment and underlying weathered-fissured and fractured bedrock. The weak influence of agricultural practices is confirmed by the low Sr-Mg values (Fig. 8). For the Oubangui basin (Négrel and Dupré, 1995a), the alterite compartment has a strong influence on the groundwater while the surface waters are scattered between the three endmembers with a stronger influence of underlying weathered-fissured and fractured bedrock. For the Congo Basin (Fig. 2c), both the yearly survey at Brazzaville (Négrel and Dupré, 1995b) and the investigation at the basin scale (Négrel et al., 1993) show a strong influence of the rain inputs and underlying weathered-fissured and fractured bedrock reflecting the groundwater influence with an increase of the Mg/Sr ratio during low water stage and slight decrease of the $^{87}\text{Sr}/^{86}\text{Sr}$ ratio. Rivers draining the western part of the Congo Basin (Négrel and Dupré, 1995a) show a strong influence of the alterite and underlying weathered-fissured and fractured bedrock as most of the samples plot along the mixing line between these two endmembers. In South Africa, the Orange River Basin is marked by a

strong influence of agricultural practices as confirmed by the relatively high Sr-Mg values but also the alterite compartment and a weak influence of underlying weathered-fissured and fractured bedrock with a Mg/Sr ratio remaining around 350.

In *French Guiana*, the scatter of the data is between the three endmembers. In the surface waters collected during low water stage the data clearly reflect the influence of the alterite for some samples (Négrel and Lachassagne, 2000), the influence of rain inputs for others (no water rock interaction as shown by Négrel and Petelet-Giraud, 2010) and the influence of underlying weathered-fissured and fractured bedrock for the rest. Regarding the groundwater, there is less influence of the alterite compartment and the data being scattered between the rain inputs and the underlying weathered-fissured and fractured bedrock as demonstrated by Négrel and Petelet-Giraud (2010).

In *North America*, regarding the Mackenzie River basin in Canada, scatter of the data is between the three endmembers reflecting the prominent influence of the alterite and rain inputs and less influence of underlying weathered-fissured and fractured bedrock. The role of anthropogenic endmembers can be discarded as illustrated by the low Sr-Mg values (Fig. 8). Most of the samples from the Fraser Basin in Canada plot between the anthropogenic inputs and the alterite compartment with less influence of the latter. It is worth noting the location of one sample along the mixing line between the alterite compartment and the underlying weathered-fissured and fractured bedrock. In the Laramie aquifer (Wyoming USA), one sample plots close to the weathered-fissured and fractured bedrock whereas the second one is marked by the three endmembers, reflecting a more prominent role of the alterite and less from anthropogenic inputs.

In *Australia* (Harrington and Herczeg, 2003), the groundwaters are scattered along the mixing line between the alterite compartment and the anthropogenic inputs reflecting a more

prominent role of the weathering processes in the alterite and the lack of any influence of deeper interaction of the weathered-fissured and fractured bedrock.

Groundwaters from *Finland* are close to the field defined in the Mackenzie River basin in Canada reflecting again the prominent influence of the alterite and rain inputs and less influence of underlying weathered-fissured and fractured bedrock and anthropogenic endmembers (as illustrated by the low Sr-Mg values in Fig. 8).

4.2.3 Variation in the alterite endmember: a consequence of bedrock age?

Through the use of different databanks of ground- and surface waters collected worldwide, we have illustrated large variations in Sr-isotope ratios with variations in the Mg/Sr ratios (Fig. 5, 8 and 9). Evidence of such co-variation has not previously been reported for isotopic ratios of Sr at this scale, and this result highlights the impact of the degree of silicate weathering on strontium isotopes due to different water-rock interaction processes.

In silicate rocks, a pattern is generally observed of increasing $^{87}\text{Sr}/^{86}\text{Sr}$ in the order plagioclase, hornblende, K-feldspar, and biotite, in close connection with the Rb/Sr ratio of each mineral. Meaning that minerals having a high Rb/Sr will have greater $^{87}\text{Sr}/^{86}\text{Sr}$ than minerals having low Rb/Sr because the ^{87}Rb decays to ^{87}Sr with a half-life of 48.8 billion years. The preferential alteration of strontium-rich, rubidium-poor minerals during weathering causes the increase of the Rb/Sr ratio in weathered rock compared to the Rb/Sr ratio of the fresh rock (Dasch, 1969) and thus the increase in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in the bulk alterite can be related with weathering removal over time of the remaining nonradiogenic plagioclase, apatite and hornblende (Pett-Ridge et al., 2009). This relative enrichment of rubidium over strontium is evident in most weathering profiles (Dasch, 1969) and continuous interaction in the alterite thus led to high Sr-isotope ratios in waters because of the weathering of residual minerals. However, in addition to the role of the Rb/Sr ratio, the age of rocks, in connection with decay,

promotes high ratios in minerals having high Rb/Sr. This means that for alterite developed on silicate rocks, older the rocks are, the more radiogenic the residual minerals are.

Interactions within the alterite of silicate in France, mainly having an age of approximately 300 to 400 Ma show lower isotope ratios as proved by comparison of water samples representing this horizon when comparing to those having older age like Vosges in France and Australia for example. These interactions in alterite of different ages can be modified by the impact of agricultural inputs (Africa, Laramie aquifer in Wyoming USA, Australia, ...) or rain inputs (Himalaya, French Guiana, Mackenzie River basin in Canada, Finland...). In water in which we may state that Sr and Mg are mostly of natural sources, but due to human pressures, Sr and its isotopes are not an excellent tracer of natural processes. However, we have demonstrated that all considered sites, Sr isotopes can serve as a tracer of water-rock interaction in the context of silicate weathering.

5 – Conclusion

Strontium isotopic tracers allow the water-rock interaction and complex relationships between the different compartments of hard-rock aquifers to be more clearly defined, from small-scale catchments to the scale of continental basins, using data acquired from sites worldwide. The existence of relatively deep alteration processes, whose role in the functioning of the critical zone will have to be properly taken into account in future studies, i.e. regarding the leaching of toxic element such as fluorides (Pauwels et al., 2015) or water pathways (Chabaux et al., 2017), is evidenced by tracing Sr isotopes. The contrasting strontium-isotope signatures are linked to differences in weathered mineral phases of the bedrock regardless of the age of the geological formations. The ground- or surface waters can show either an increase or a decrease in the proportions of weathered phases, which probably are related to the functioning of the weathering profile in crystalline rock (Dasch,

1969; Négrel, 1999; Probst et al., 2000). The Sr-isotope tracing defines and identifies the relative signature of water circulation in alterite and underlying weathered-fissured and fractured bedrock. Due to its large range in abundance of the residual minerals, the alterite endmember has Sr isotope ratios that may vary as a function of the age of the bedrock. Effective water management is considered crucial for building adaptive capacity to be resilient to the impacts of climate change (Nelson et al., 2007). Thus, a greater understanding of the nature of weathering processes involved and their relationships with water circulation is seen as being at the heart of present and future water challenges (Négrel and Petelet-Giraud, 2013).

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Figure captions

Fig. 1. Conceptual model of hydrogeological properties in weathered basement rock, i.e.the hard rock aquifer concept, modified from Wyns et al. (2004).

Fig. 2. Map of the site locations of studied crystalline catchments in France (modified from Négrel and Pauwels, 2004).

Fig. 3. Map of the site locations: (a) India(Subarnarekha, Négrel et al., 2007,Maheshwaram, Négrel et al., 2010; Pauwels et al., 2015) and Himalayan rivers (Bothe Kosi River, Harris et al., 1998; Ganges headwaters, Bickle et al., 2005), (b) French Guiana (Négrel and Petelet-Giraud, 2010), Africa (Congo and Oubangui Rivers, Négrel et al., 1993; 1995a, b).

Fig. 4. Location of the study area in Andhra Pradesh (India) and simplified geological map of the Maheshwaram hydrosystem with location of sampled boreholes (Mxx for agricultural used boreholes and IFPX for scientific used boreholes).

Fig. 5. Sr vs. Mg contents (all $\mu\text{mol/l}$) in surface and groundwater interacting with silicate in France, modified from Négrel and Pauwels (2004).

Fig.6. Sr isotope ratios vs. Mg/Sr ratios modified from Négrel and Pauwels (2004) and additional data from the literature covering waters interacting with silicate in France.

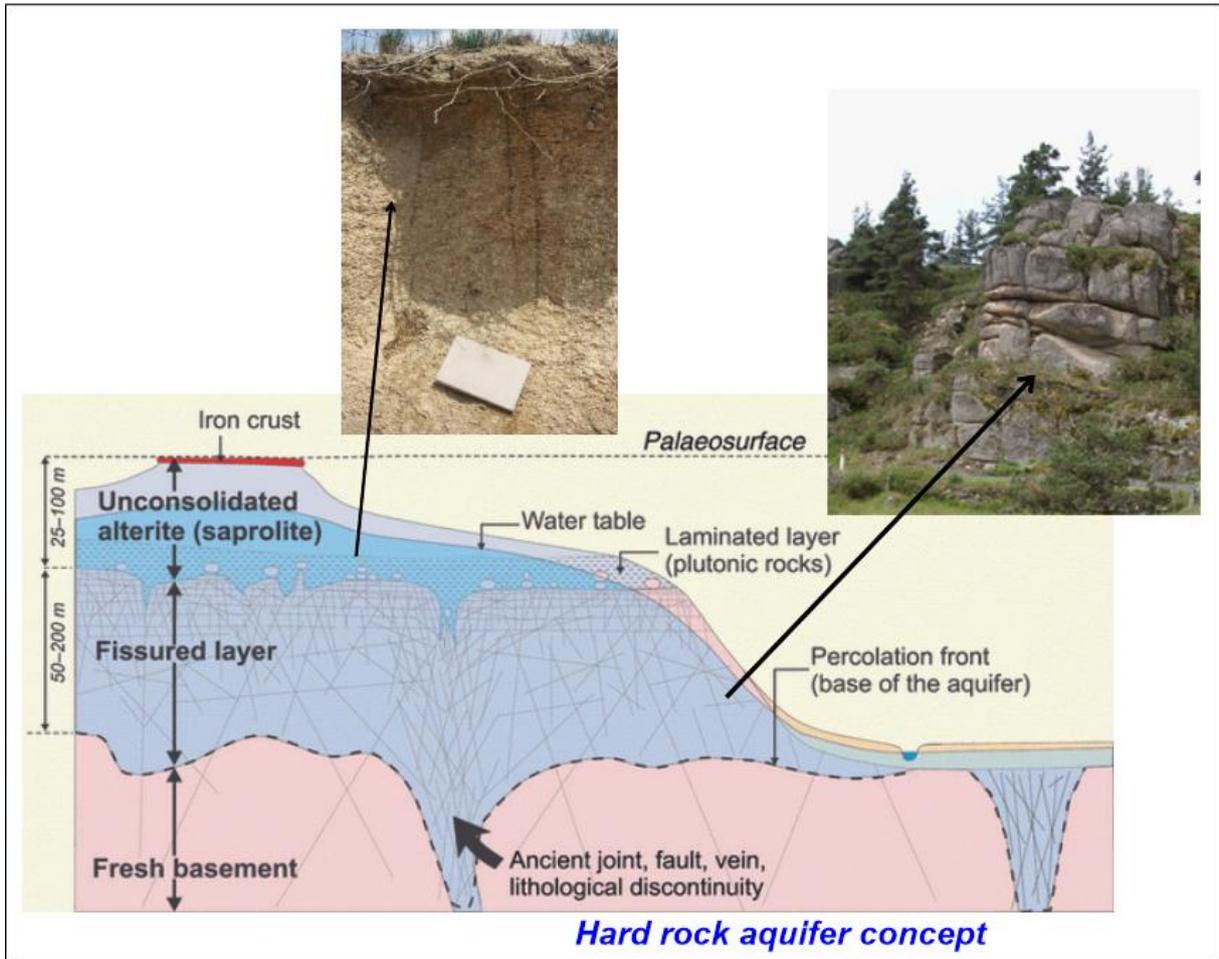
Fig.7. (a) Plot of the $^{87}\text{Sr}/^{86}\text{Sr}$ vs. Cl concentration expressed in $\mu\text{mol/l}$ for the groundwater samples collected in the Maheshwaram hydrosystem in 2006 (A. Dayal, personal communication) and 2008 (this study); (b) plot of $^{87}\text{Sr}/^{86}\text{Sr}$ vs. $1/\text{Sr}$ (Sr concentration expressed in $\mu\text{mol/l}$).

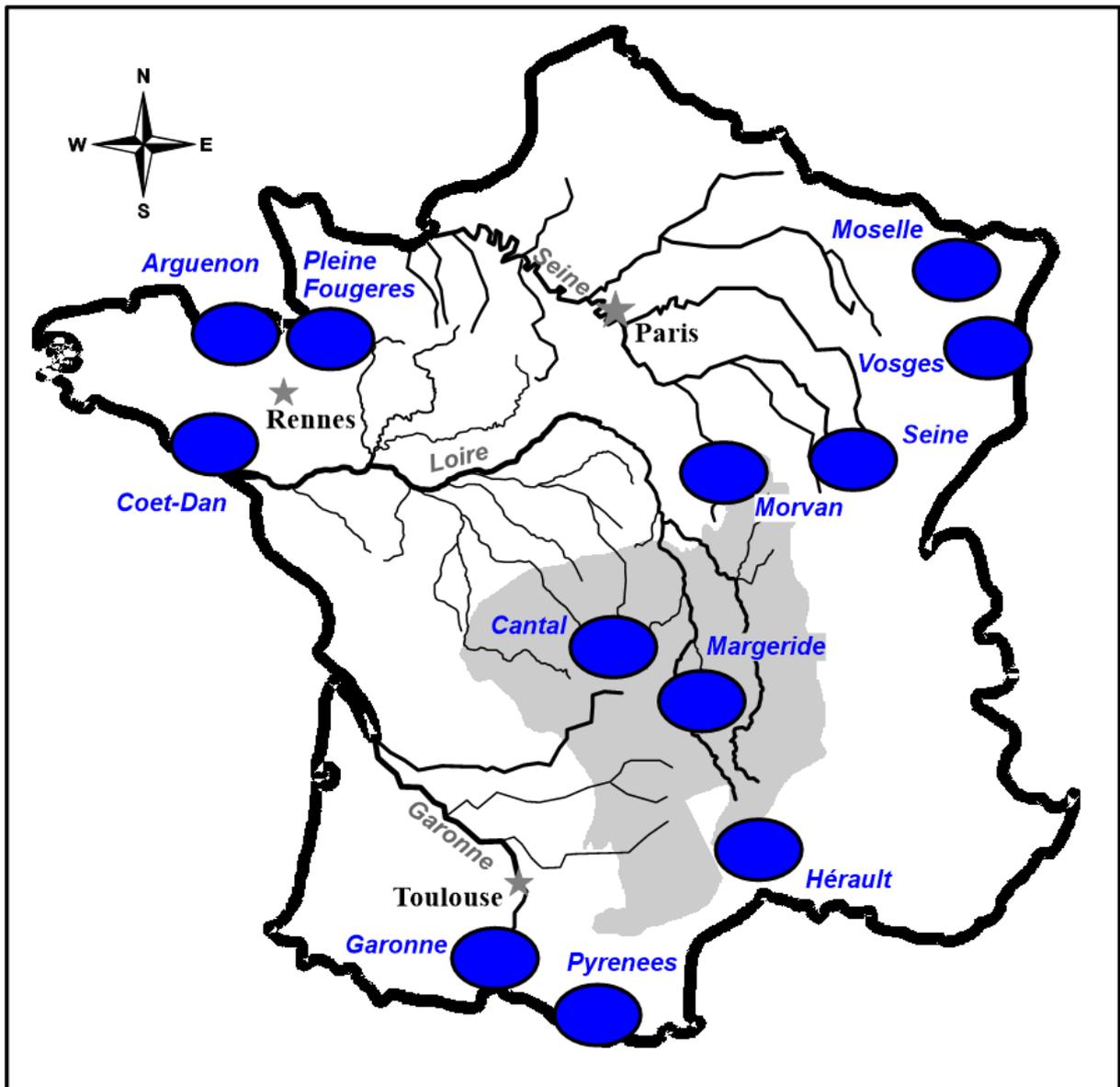
Fig. 8. Sr vs. Mg contents (all $\mu\text{mol/l}$) when enlarging to worldwide basins and catchments in Africa (Congo Basin Négrel et al. 1993; Orange River; de Villiers et al., 2000); India and Nepal (Négrel et al., 2007a and this study for India; Himalaya metamorphic, silicate metasediments and gneisses 630 to 490 Ma; Harris et al., 1998; Bickle et al., 2005); North America (3500 to 2500 Ma silicate rocks in the Mackenzie basin; Millot et al., 2002 and 1000 to 70 Ma silicate rocks in the Fraser Basin in Canada; Cameron et al., 1995 and 1760 to 1430 anorthosite and granite in the Laramie aquifer in Wyoming; Frost and Toner, 2004); South America in French Guiana (Archaen gneiss 3400 to 2700 Ma and granite-gneiss rocks 2300 to 1900 Ma; Négrel and Petelet-Giraud, 2010); Australia (65 Ma arenaceous and argillaceous rocks; and considering both surface and groundwater; Harrington and Herczeg, 2003).

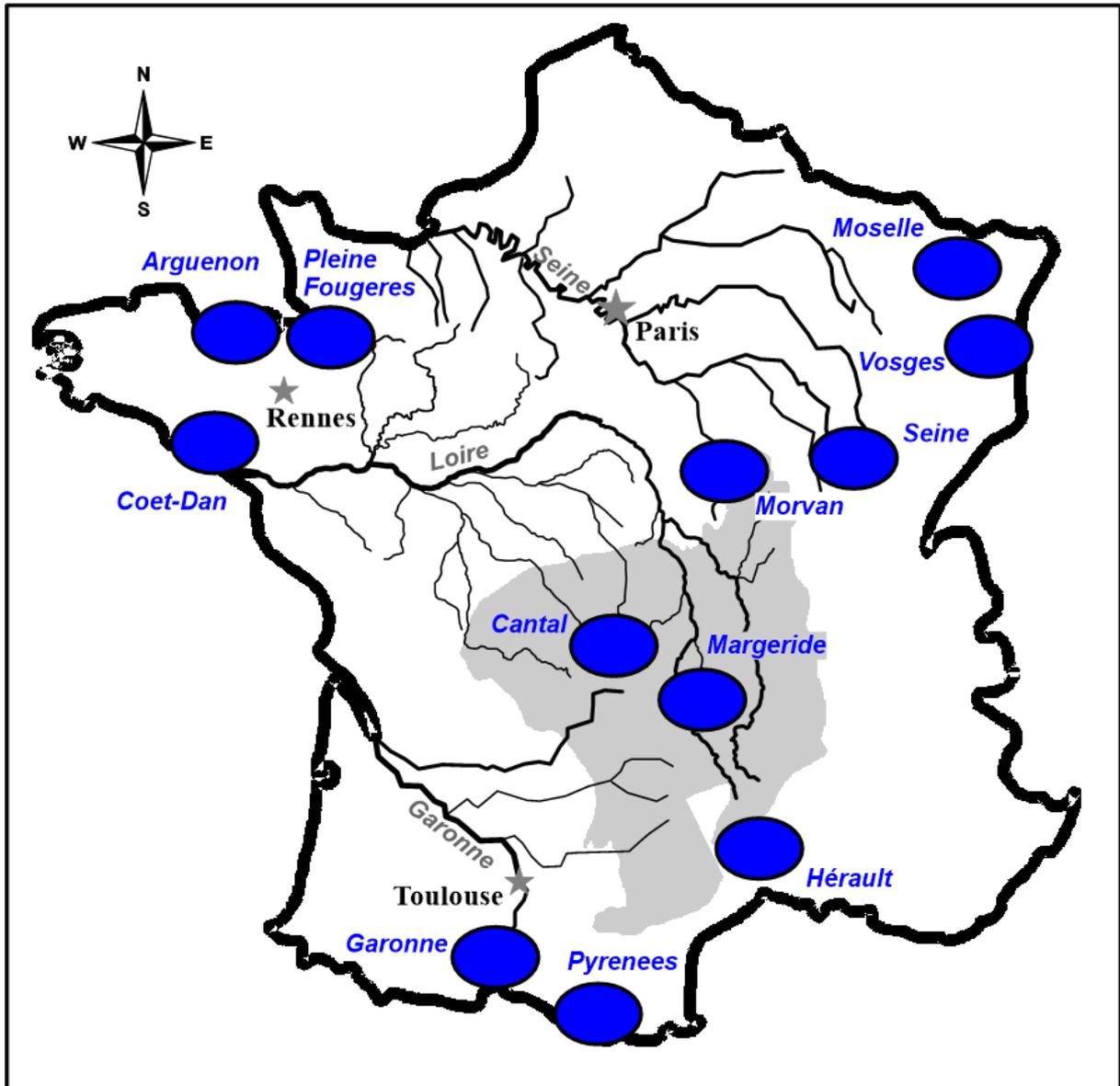
Fig. 9. Sr isotope ratios vs. Mg/Sr ratios, modified from Fig. 6, and additional data when enlarging to worldwide basins and catchments in Africa (Congo Basin Négrel et al. 1993; Orange River; de Villiers et al., 2000); India and Nepal (Négrel et al., 2007a and this study for India; Himalaya metamorphic, silicate metasediments and gneisses 630 to 490 Ma; Harris et al., 1998; Bickle et al., 2005); North America (3500 to 2500 Ma silicate rocks in the Mackenzie basin; Millot et al., 2002 and 1000 to 70 Ma silicate rocks in the Fraser Basin in Canada; Cameron et al., 1995 and 1760 to 1430 anorthosite and granite in the Laramie aquifer in Wyoming; Frost and Toner, 2004); South America in French Guiana (Archaen gneiss 3400 to 2700 Ma and granite-gneiss rocks 2300 to 1900 Ma; Négrel and Petelet-Giraud, 2010); Australia (65 Ma arenaceous and argillaceous rocks; and considering both surface and groundwater; Harrington and Herczeg, 2003).

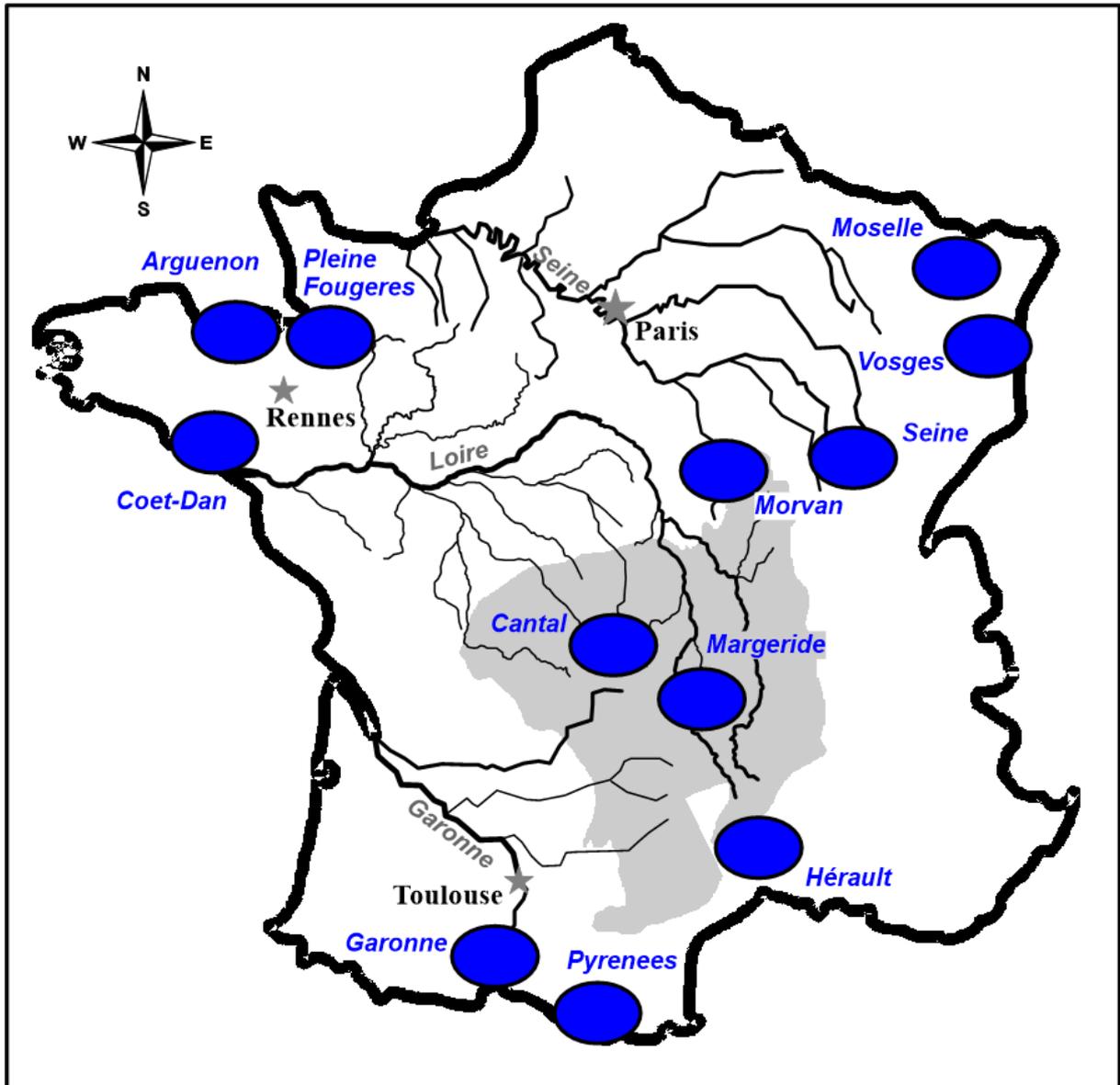
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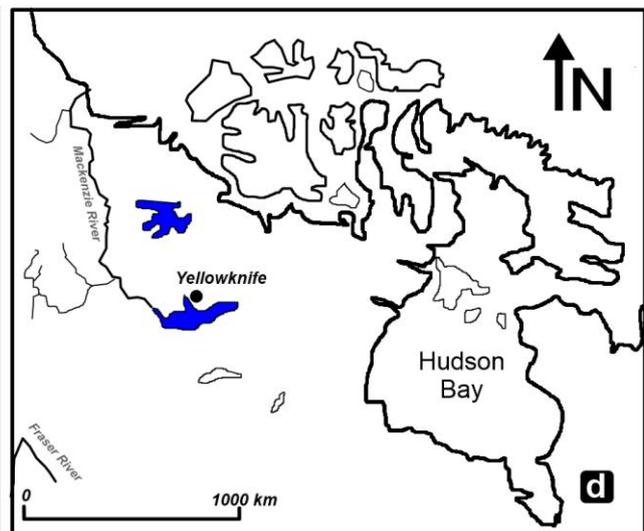
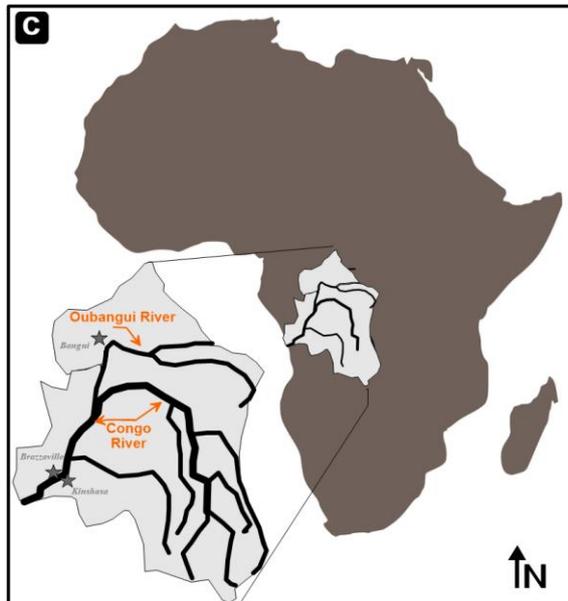
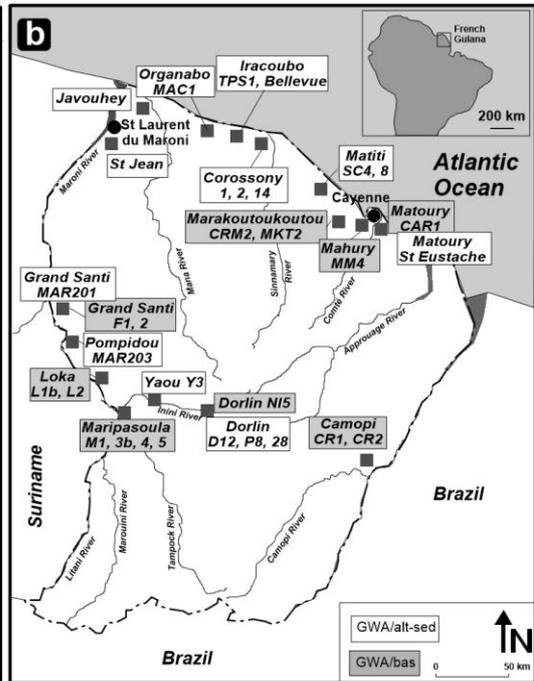
Table 1. Mg and Sr concentration expressed in $\mu\text{mol/l}$, $^{87}\text{Sr}/^{86}\text{Sr}$ ratios for the groundwater samples collected in the Maheshwaram hydrosystem in 2006 (A. Dayal, personal communication) and 2008 (this study).

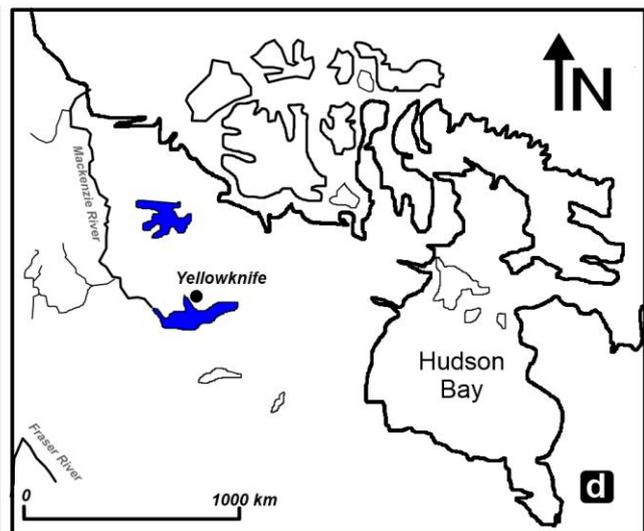
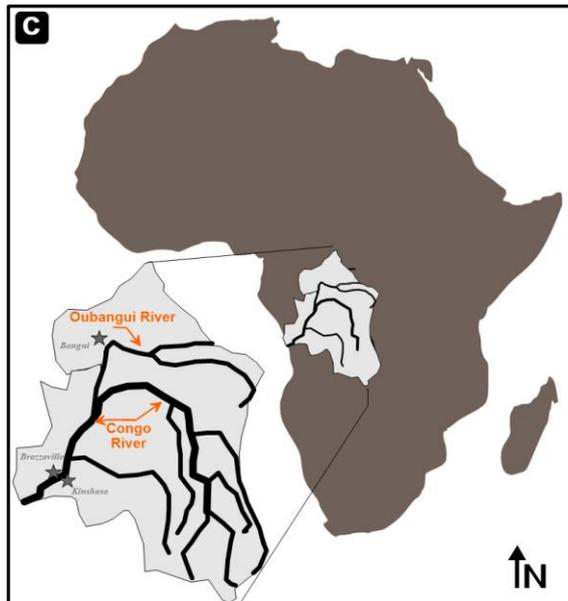
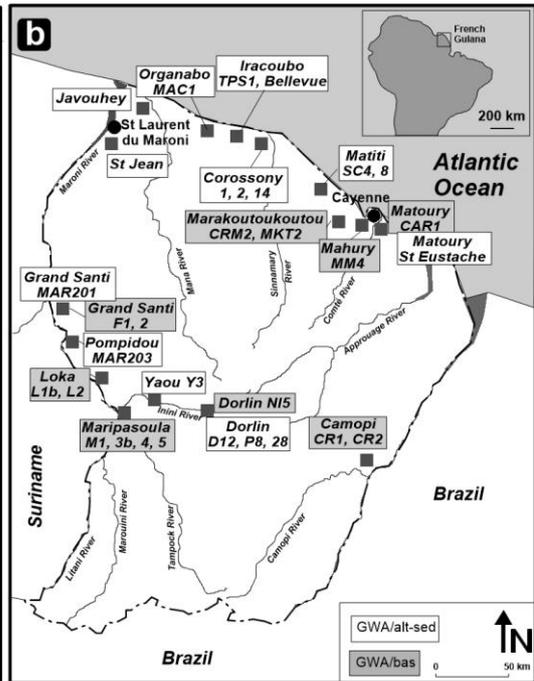


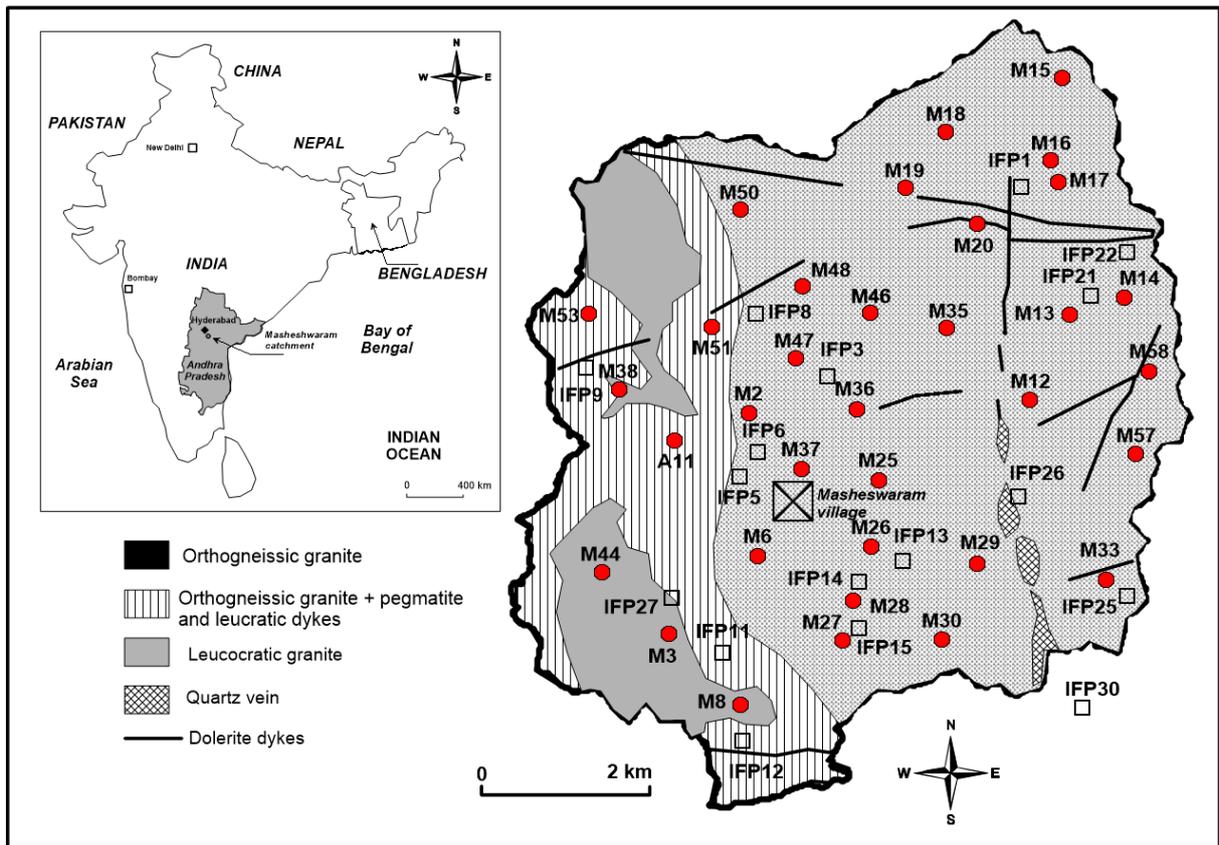


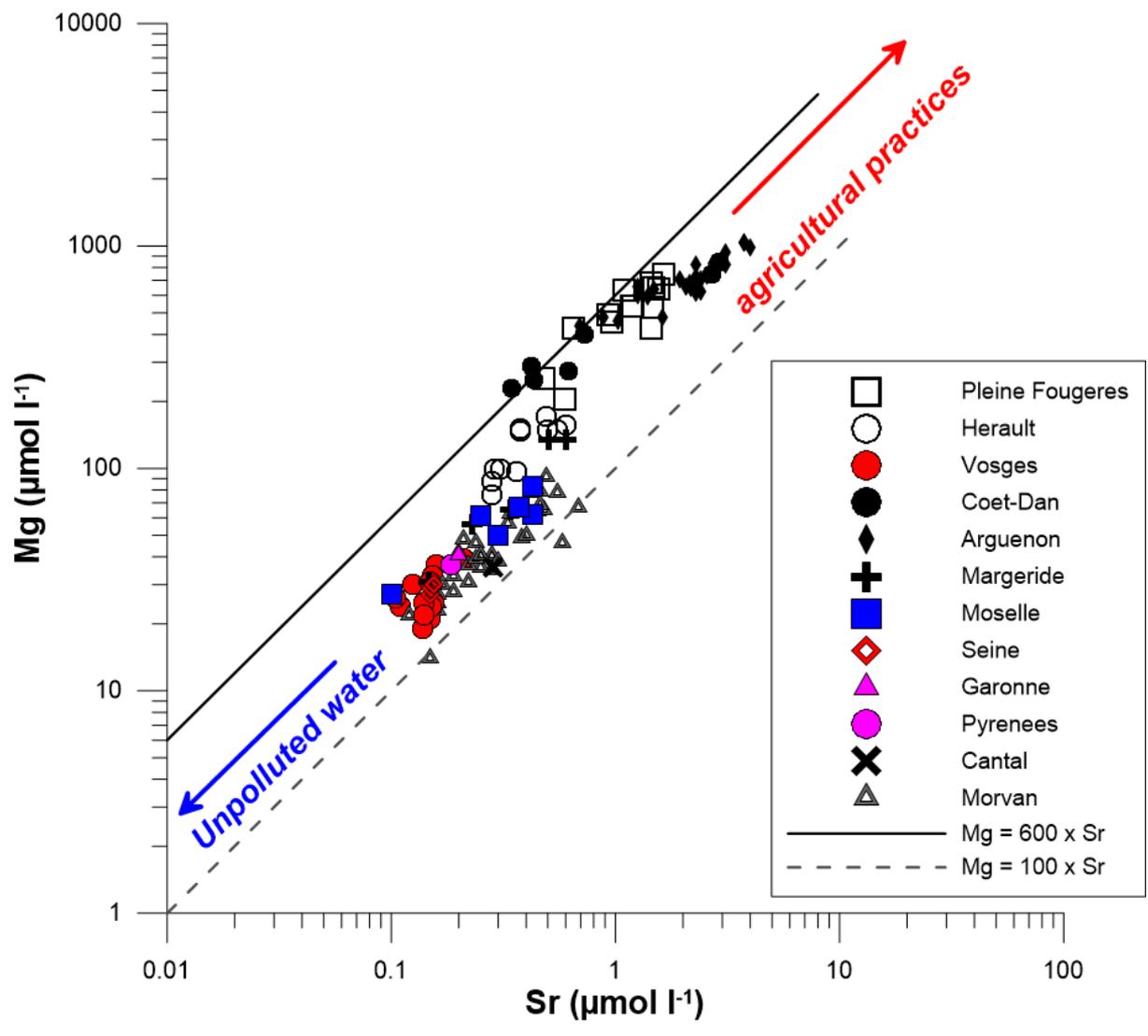


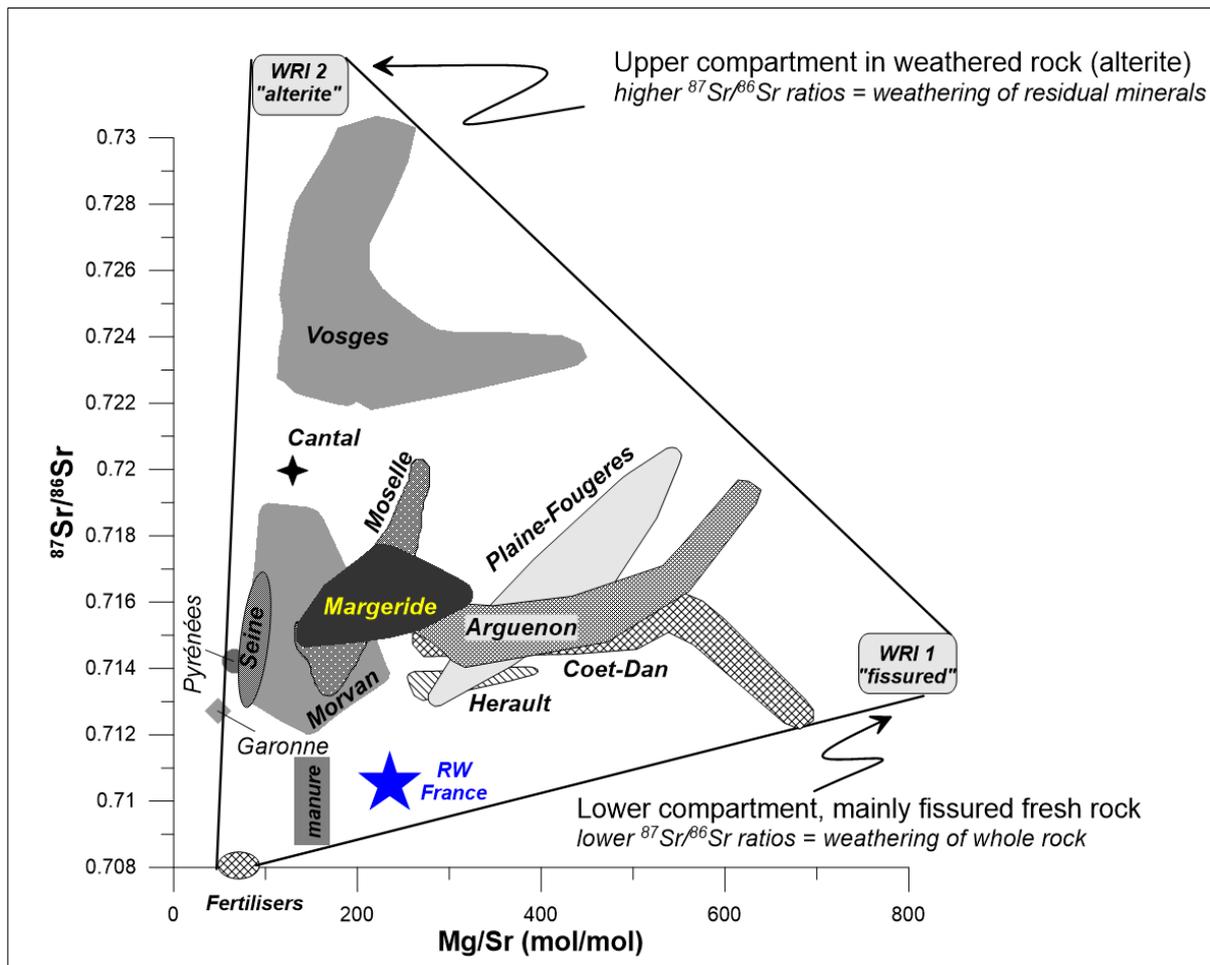


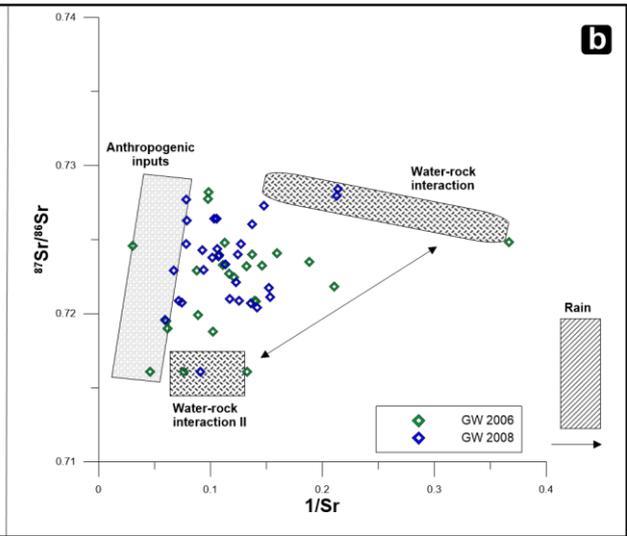
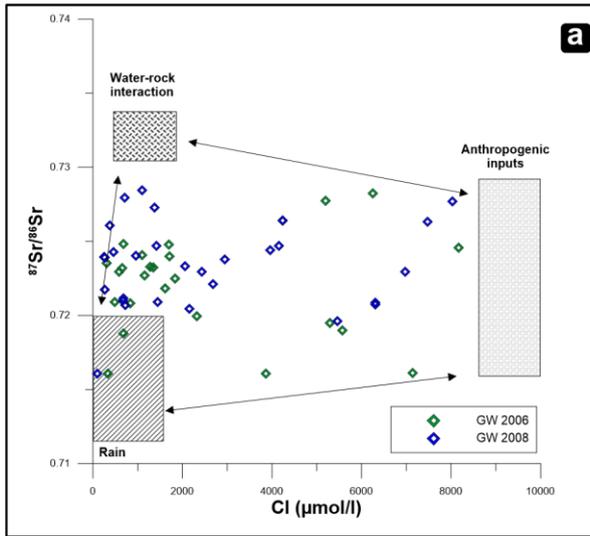


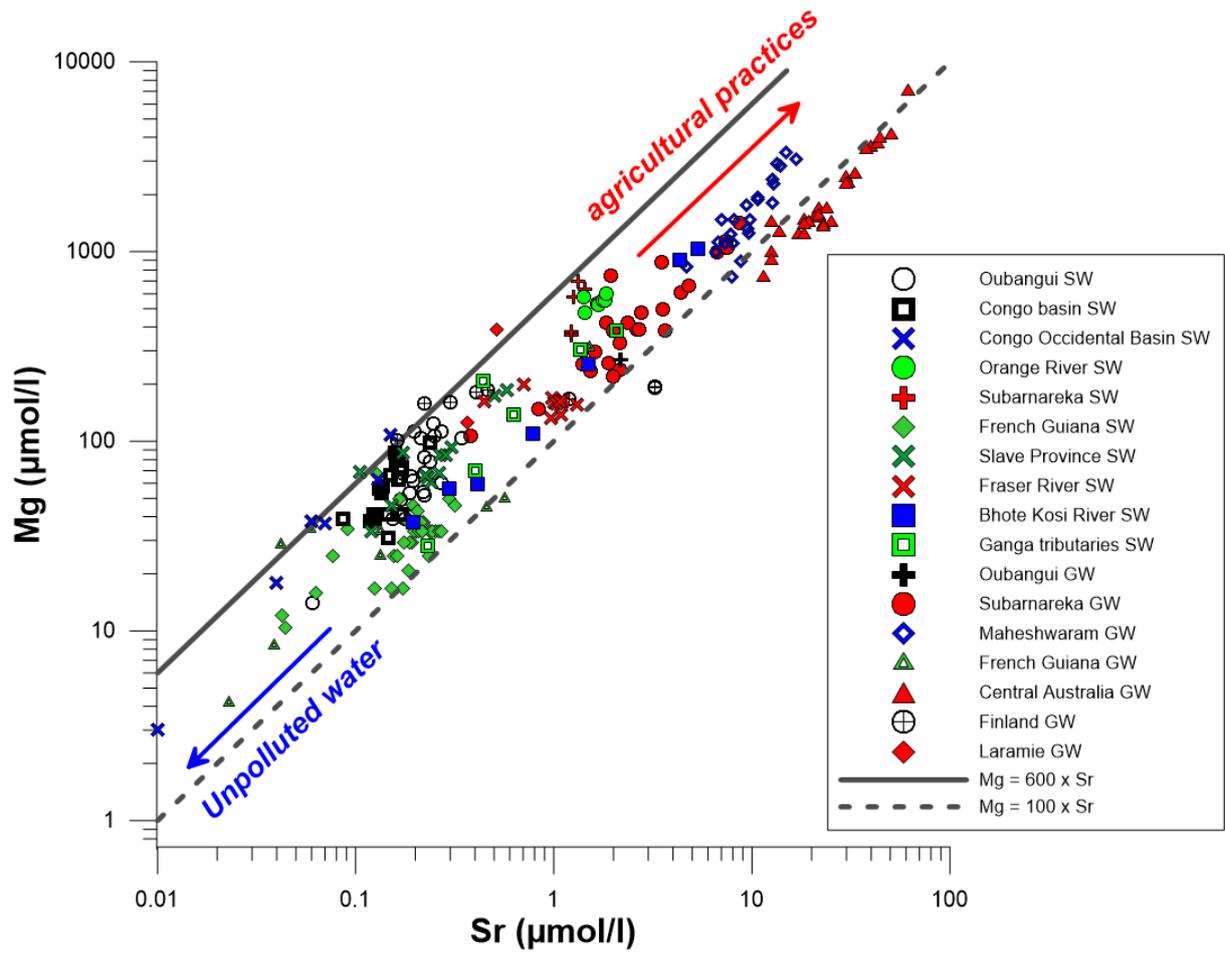


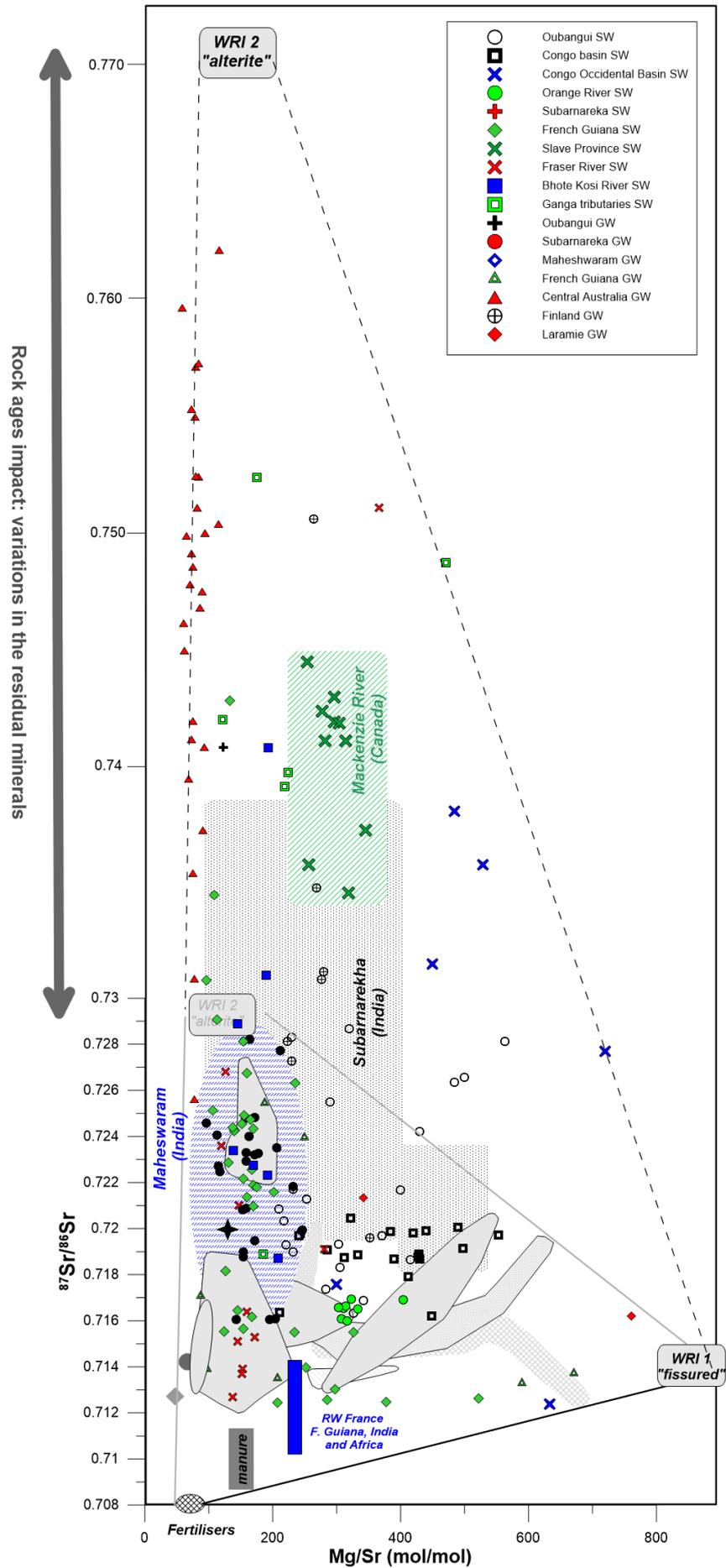












Sample	Mg μmol	Sr μmol	$^{87}\text{Sr}/^{86}\text{Sr}$		Sample	Mg μmol	Sr μmol	$^{87}\text{Sr}/^{86}\text{Sr}$
2006	$\mu\text{mol/L}$	$\mu\text{mol/L}$			2008	$\mu\text{mol/L}$	$\mu\text{mol/L}$	
IFP 12	1747	10.1	0.728226		M2	2404	12.7	0.726301
A11	1145	7.1	0.72083		M3	888	8.8	0.723328
HP3	4664	21.8	0.716094		M6	1105	8.0	0.724011
M03	1041	8.6	0.722701		M8	739	8.0	0.720892
M06	1496	9.0	0.723293		M12	1101	7.4	0.720694
M08	740	6.3	0.724064		M30	999	6.6	0.721741
M12	1583	9.8	0.718774		M35	1877	10.7	0.722948
M13	2674	13.1	0.71606		M48	1241	7.9	0.724700
M15	3337	33.0	0.724582		M50	1114	6.8	0.727282
M17	2964	16.5	0.719486		M51	2272	12.8	0.727688
M18	2628	16.3	0.718987		M53	1935	10.8	0.724285
M20	1155	4.7	0.72183		M57	3065	16.8	0.719595
M25	1192	7.2	0.720891		IFP1/1	1474	7.1	0.720433
M26	1895	11.4	0.722921		IFP1/1a	1481	8.1	0.722106
M27	1362	7.6	0.723186		IFP1/7	2910	13.5	0.720743
M30	1146	5.3	0.723503		IFP1/7a	2821	13.9	0.720857
M33	1130	7.5	0.71606		IFP1/7b	3335	14.8	0.722931
M35	1272	6.8	0.723252		IFP5	1478	9.8	0.723768
M38B	1244	7.3	0.723981		IFP6	1747	9.4	0.724389
M44	1460	8.9	0.724783		IFP6	1813	12.8	0.724689
M46	1014	8.3	0.722471		IFP8	1316	9.5	0.726401
M48	490	2.7	0.724826		IFP8	1248	9.7	0.726405
M51	2265	10.2	0.727728		IFP9	836	4.7	0.728431
M58	2895	11.2	0.719925		IFP9	785	4.7	0.727958
					IFP13	1228	9.3	0.723954
					IFP14	1060	7.3	0.726056
					IFP15	1093	8.5	0.721001
					IFP25	1572	11.0	0.716063

Table 1