## Anthropogenic and lithologic Pb behaviours in acid atmospheric polluted catchment (the Strengbach case study, Vosges, France)

 N. Pycke
 Laboratoire de Géochimie, Université Paul Sabatier, OMP et

 D. Baque
 CNRS UMR 5563, 38 rue des 36 ponts, 31400 Toulouse, France

 A. Probst
 Centre de géochimie de la surface, CNRS UPR 06251, 1 rue

 Blessig, 67084 Strasbourg Cedex, France

 B. Dupre
 Laboratoire de Géochimie, Université Paul Sabatier, OMP et

 M. Loubet
 CNRS UMR 5563, 38 rue des 36 ponts, 31400 Toulouse, France

Two main reasons lead us to study the behaviour of lead in an equiped natural catchment - the Strengbach bassin in the Vosges massif, France.

(1) Like other radiogenic isotopes, Pb is an interesting tracer to provide insights on the fonctionning the natural system.

(2) Lead is also one of the most significant and dangerous polluting agent. Therefore the knowledge of its behaviour in environnmental systems is primordial. In such natural systems Pb has two origins: a natural one and an anthropogenic one. In an interesting way Pb isotopic composition allows to distinguish these two origins.

Geochemical characteristics of open field rain water, throughfall, soil solution, spring and streamwaters in the small granitic Strengbach catchment (Vosges, France) have been regulary determined for ten years (Probst *et al.*, 1995; Dambrine *et al.*, 1995) and specifically during hydrological event (Ladouche et al., 1995). Major elements, dissolved organic carbon (DOC), trace element contents were particulary analysed. The different water fluxes distinguish one another by specific geochemical features. Following these previous studies, a specific analysis of the Pb absolute content and isotopic signature was undertaken in these different water fluxes in order to identify the behaviour of this element in such an acid atmospheric polluted catchment and to distinguish the anthropogenic Pb from the substratum Pb.

The following main points were reached:

Isotopic compositions of the atmospheric anthropogenic Pb (from the rain) and of the substratum (granite) distinguish clearly one another. The water fluxes display Pb isotopic compositions in between these two extremum values ( $17.8 < {^{206}Pb}/{^{204}Pb} < 18.9$ ) (Fig. 1).

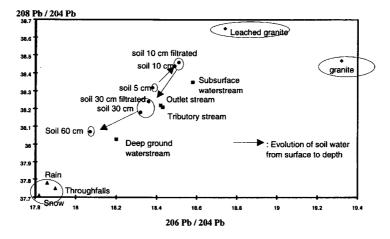


FIG. 1. (<sup>208</sup>Pb/<sup>204</sup>Pb) vs (<sup>206</sup>Pb/<sup>204</sup>Pb).

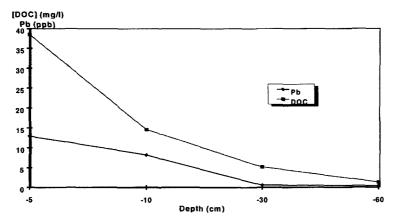


Fig. 2. Lead and DOC concentration in soil solutions (depth 5, 10, 30, 60 cm).

(2) Pb isotopic compositions evolve from surface to depth. However in a surprising way, waters draining the subsurface compartments display compositions close to those of the substratum. Waters from the deeper soil profile or saprolites (Springs) are close to Pb atmospheric signature (i.e. anthropogenic origin)(Fig. 1).

(3) In water fluxes, Pb content decreases significantly with depth and Pb content is correlated with DOC (Fig. 2)

A preferential complexation of Pb by organic compounds explains the specific behaviour of this element within the basin. Fluxes analys show that Pb is selectively removed from the basin with subsurface water fluxes.

Various hypotheses can explain the surprising Pb isotopic evolution with depth: water pathway, low complexation kinetics, disinct complexation forms of Pb in rain and soil solution.

## References

Dambrine et al., (1995) In: Forest decline and atmospheric deposition effects in the french mountains, Landmann, G.& Bonneau, M. (eds), 233-58.

Ladouche et al., DBT II, (1995) 51.

Probst et al., WASP, (1995)79, 39-59.