

**BRAUN Florence<sup>1,2</sup>, DELGENES Nadine<sup>1</sup>, CREUSOT Nicolas<sup>3</sup>, AIT-AISSA Selim<sup>3</sup>, LE MENACH Karyn<sup>4</sup>, BUDZINSKI H el ene<sup>4</sup>, HAMELIN J er ome<sup>1</sup>, PATUREAU Dominique<sup>1</sup>**

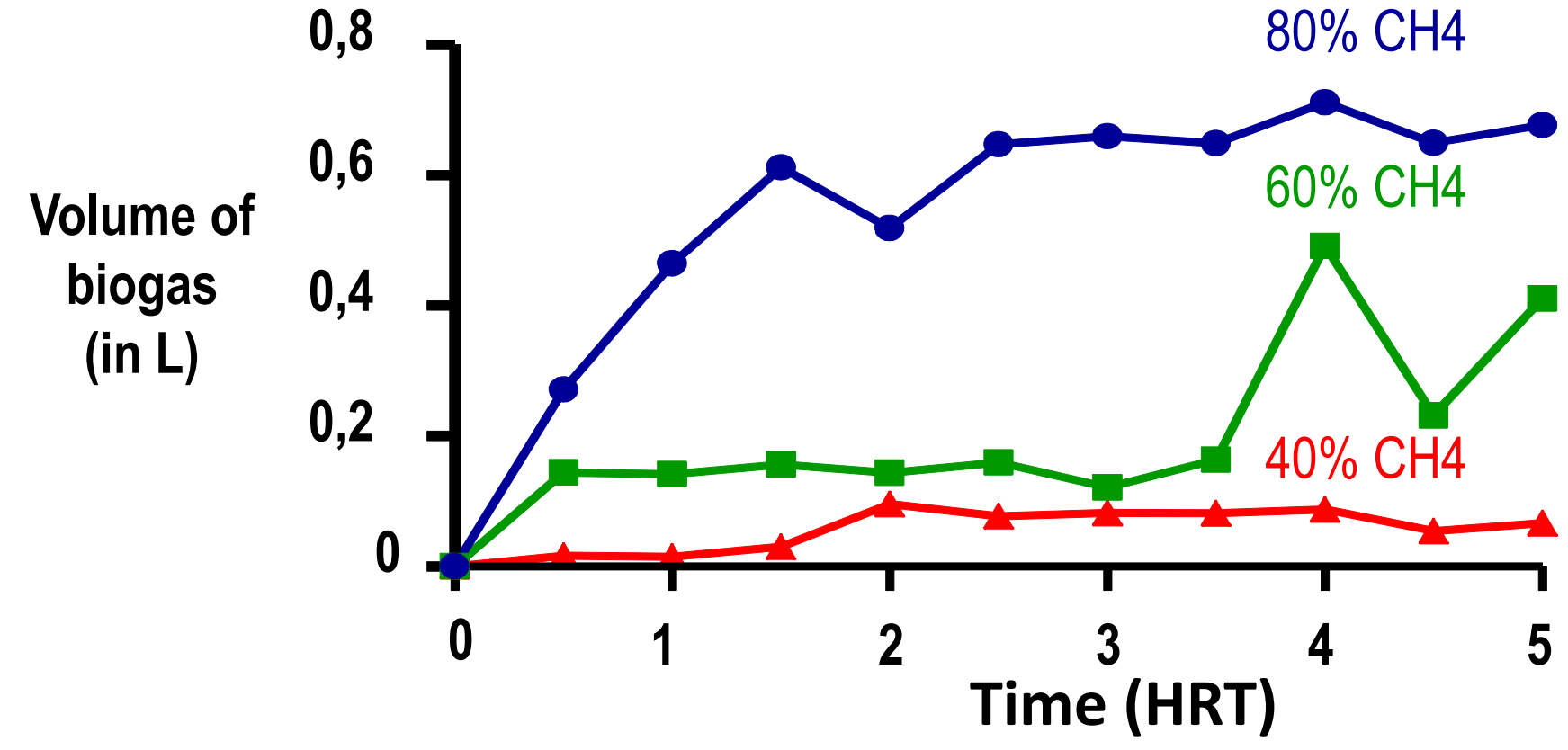
<sup>1</sup>INRA, UR050, Laboratoire de Biotechnologie de l'Environnement, Avenue des Etangs, Narbonne F-11100, France -  
<sup>2</sup>ADEME, French Environment and Energy Management Agency, 20 avenue du Gr esill -BP 90406, F-49004 Angers Cedex 01, France -  
<sup>3</sup>INERIS, Unit  Ecotoxicologie in vitro et in vivo, Parc Alata, F-60550 Verneuil-en-Halatte, France -  
<sup>4</sup>Universit  Bordeaux 1, Environnements et Pal oenvironnements O c aniques et Continentaux EPOC - UMR 5805 CNRS, Laboratoire de Physico- et Toxicochimie de l'environnement (LPTC), B timent A12, 351 cours de la Lib ration, Talence, F-33405, France.

**INTRODUCTION**

Urban sludge is often contaminated by organic pollutants such as polycyclic aromatic hydrocarbons (PAH). **Biodegradation under methanogenic conditions** was already reported for PAH [1]. However, the potential of degradation relied on the compounds availability, depending on both the organo-mineral composition of the sludge and on the intrinsic performances of the microbial populations. We investigated the influence of 3 microbial communities with **contrasting pollution history (PAH contaminated soil, PCB contaminated sediment, anaerobic sludge)** on **13 PAH removal**, their **distribution** in each physical compartment (aqueous, particulate) and their **endocrine disrupting activity** while physico-chemical conditions are strictly controlled.

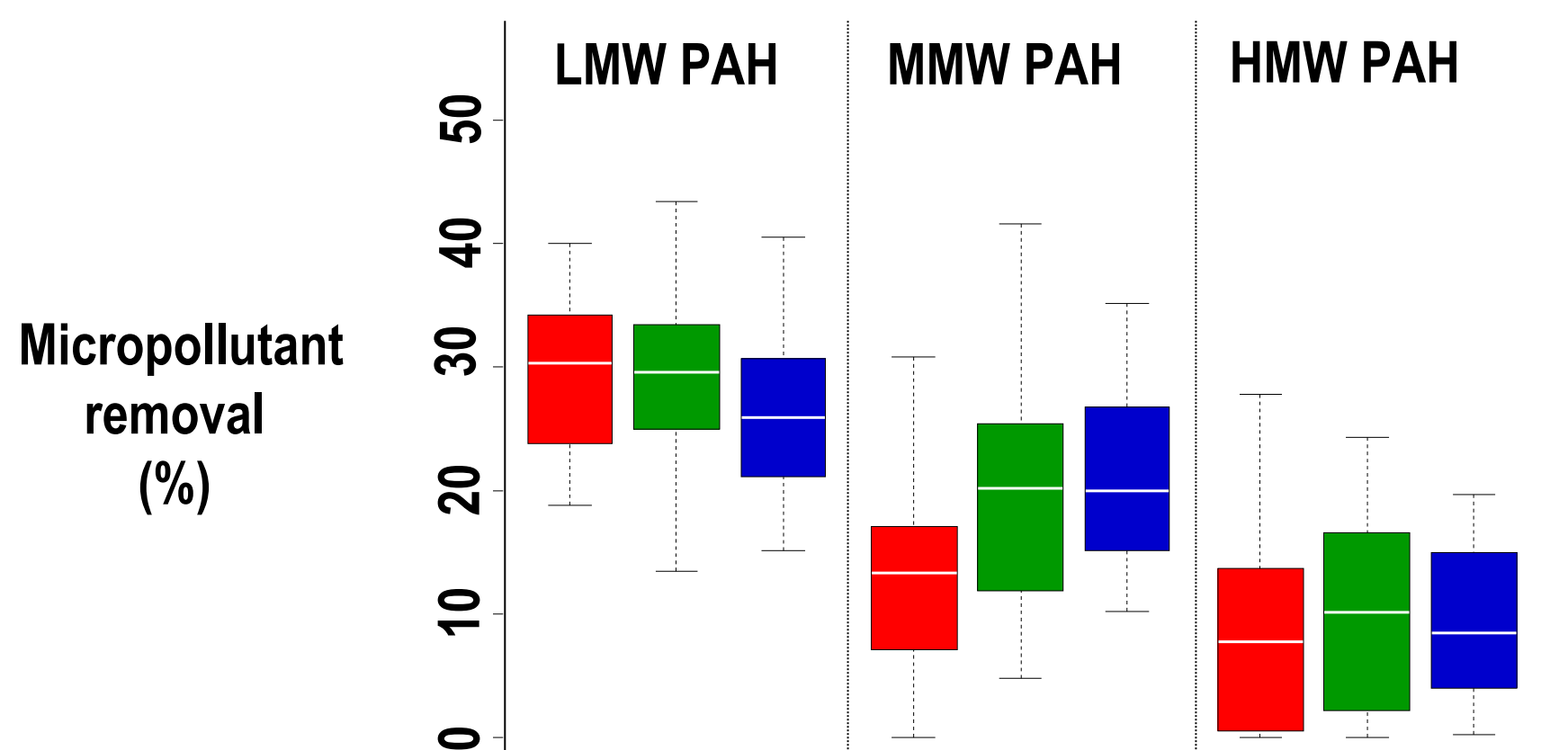
**MAIN RESULTS**

**1) Different metabolic pathways**



Biogas production over 100 days and methane proportion at steady state according to the origin of the inocula (soil, sediment, sludge) [5].

**2) Same PAH removal rates whatever the inoculum and the metabolic pathway**



The PAH removals at steady state are correlated with the molecule characteristics [5].  
 MW: molecular weight (L: low, M: medium, H: high).

**REFERENCES**

[1] Barret M, Cea-Barcia G, Guillon A, Carr re H, Patureau D. 2010. Influence of feed characteristics on the removal of micropollutants during the anaerobic digestion of contaminated sludge. Journal of Hazardous Materials 181, 241-247.  
 [2] Braun F, Hamelin J, G evaudan G, Patureau D. 2011. Development and application of an enzymatic and cell flotation treatment for the recovery of viable microbial cells from environmental matrices such as anaerobic sludge. Applied of Environmental Microbiology 77, 8487-8493  
 [3] De Perre C. 2009. Etude des interactions mati re organique dissoute - contaminants organiques dans l'environnement aquatique. PhD University Bordeaux 1.  
 [4] Kinani S, Bouchonnet S, Creusot N, Bourcier S, Balaguer P, Porcher J-M, Ait-Aissa S. 2010. Bioanalytical characterisation of multiple endocrine- and dioxin-like activities in sediments from reference and impacted small rivers. Environmental Pollution 158, 74-83  
 [5] Braun F, Hamelin J, Bonnafous A, Delgenes N, Steyer J-P, Patureau D. Similar micropollutants fate in anaerobic digesters fed with the same sludge but exhibiting different microbial populations and metabolic routes (submission in process)

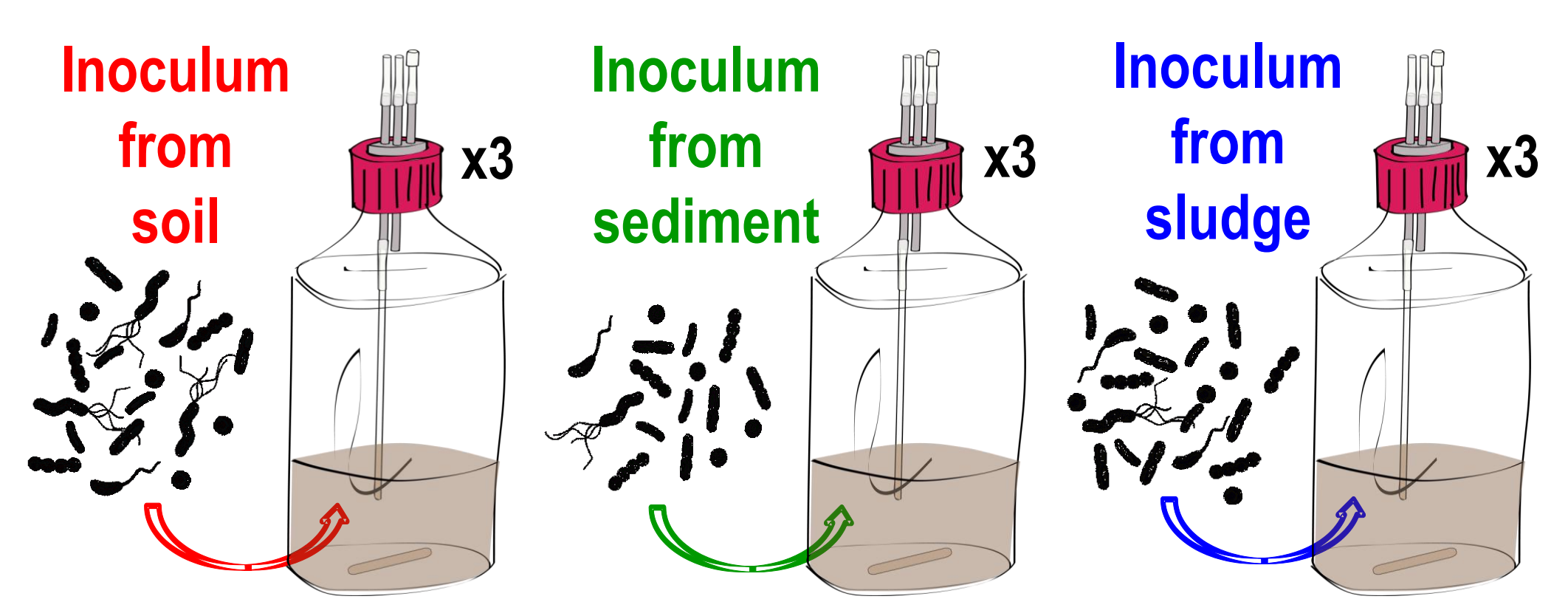
**PNRPE- Recent advances on the environmental and health effects of endocrine disrupters 10-11 D cembre 2012, Paris (France)**

**MATERIAL & METHODS**

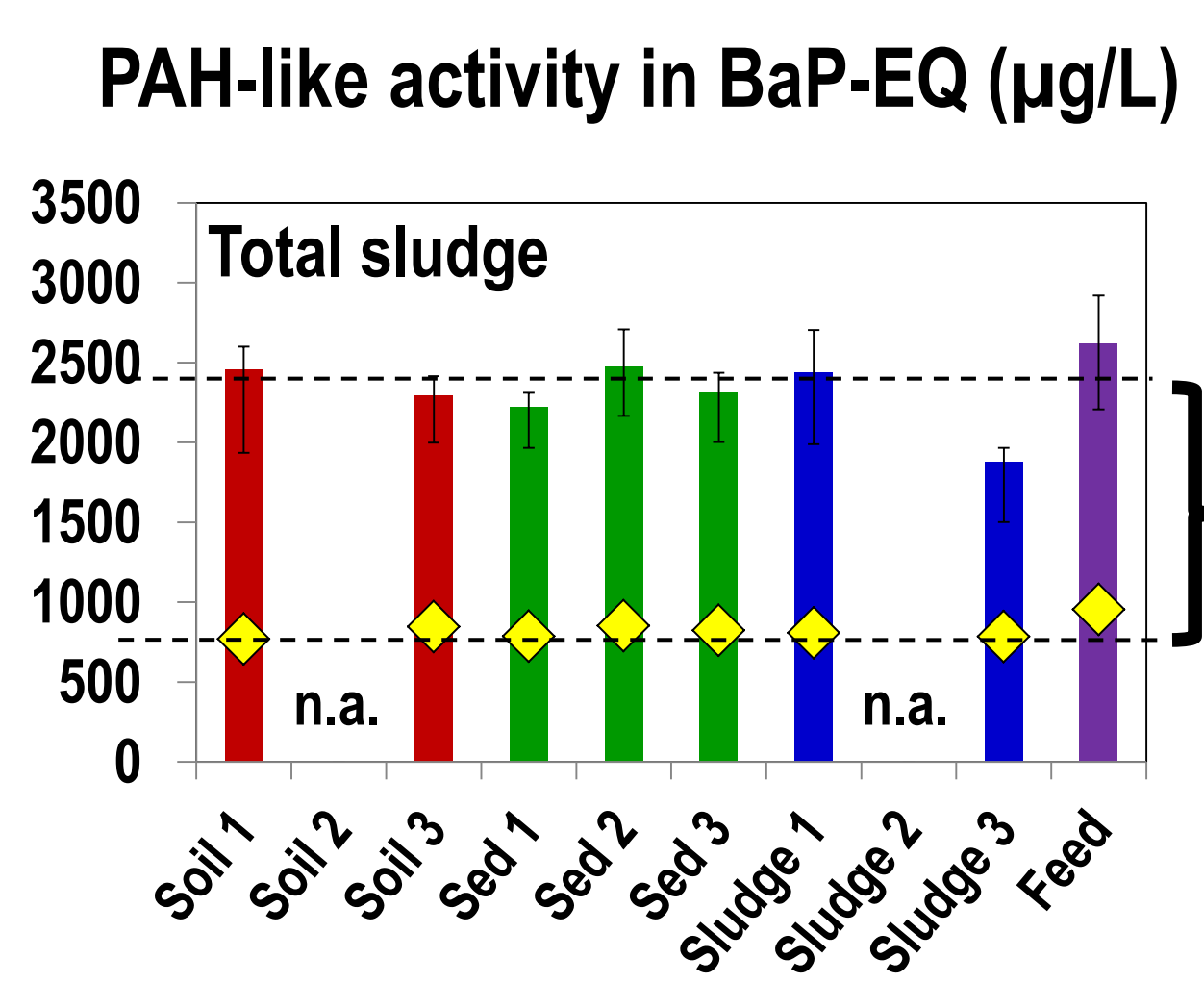
**9 anaerobic reactors** were operated at 37 C with a hydraulic retention time (HRT) of 20 days. The inoculated microorganisms were extracted from **3 ecosystems (soil, sediment, sludge)** [2]. The **sterile feed** was a sludge **spiked with 13 PAH and 7 PCB** (5 mg.kg<sup>-1</sup> DM) and **NP** (100 mg.kg<sup>-1</sup> DM).

- **Monitoring:** biogas production, biogas composition, dry matter and volatile fatty acid concentrations, chemical oxygen demand, PAH concentrations.
- **At steady state:** measure of PAH concentrations (in total and particulate compartment by ASE-HPLC and in aqueous compartment by SPME-GC-MS [3]).

Measure of PAH-like activity (*in vitro* bioassays) in total, particulate and aqueous compartments. The bioassays are based on the ability of PAH-like compounds to act through the AhR signaling pathway [4].



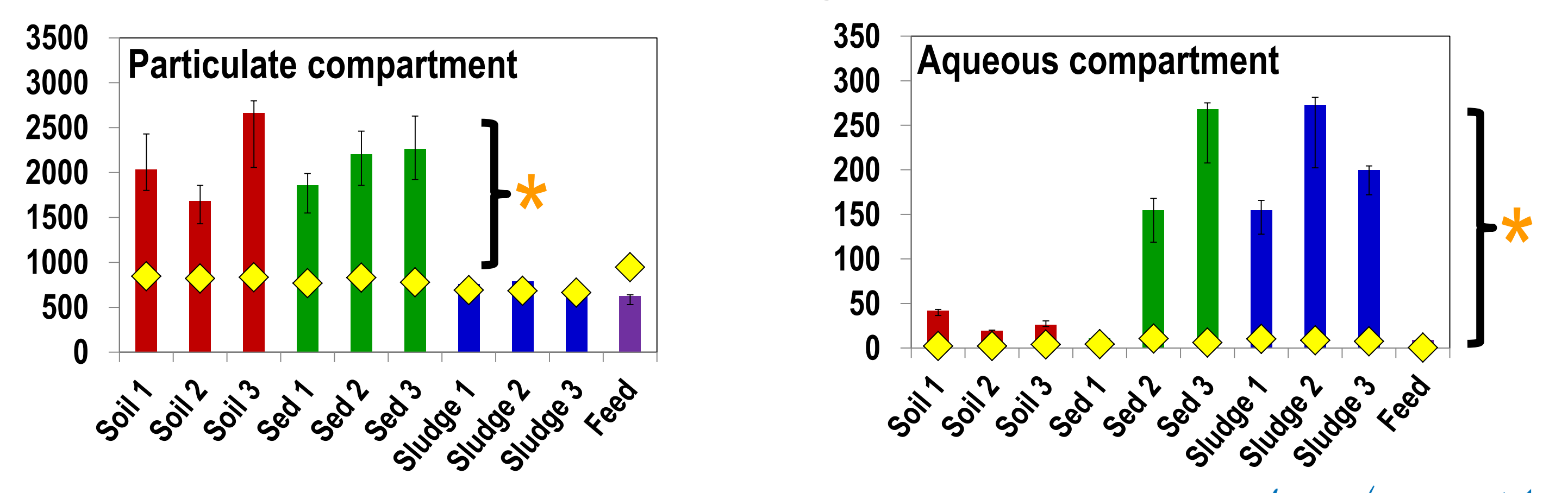
**3) In total sludge, same PAH-like activity between feed and digested sludge**



The PAH-like activities calculated by PAH chemical concentrations ( ) don't explain the totality of the biological activity measured at steady state for the 3 inocula (soil, sediment, sludge) and feed. (\* Presence of PAH by-products or other compounds that interact with AhR receptors.)

**4) PAH-like activities in the compartments linked to the metabolic pathways**

**5) PAH repartition and PAH-like activity in the particulate and aqueous compartments relied on the anaerobic digestion level**



*Take-Home Message*



The PAH removal converged to the same level, regardless of the inoculum or the metabolic pathway but were correlated to the molecule characteristics. PAH repartition and PAH-like activity in the sludge compartments depend on the digestion degree of the organic matter. Other compounds or PAH degradation by-products maintained the global PAH-like sludge activity.