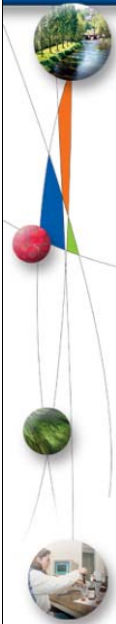


Dissipation of polar xenobiotics from pharmaceutical POCIS and suggestion of a performance reference compound

N. Mazzella, T. Debenest, J. Grange, C. Madigou, F. Delmas

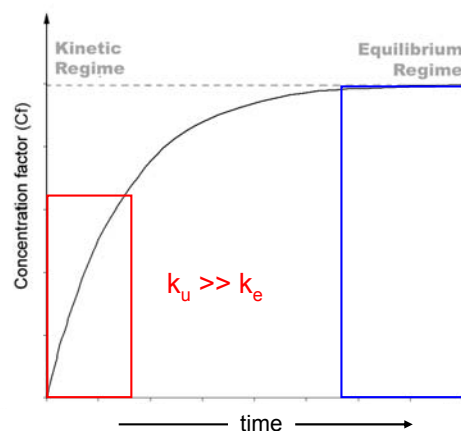


Evaluation of the water quality and its consequences on river biological organisms (WFD)

Effects of pesticides and metabolites on diatom communities

Time-weighted average concentration estimates :

- ✓ Classical grab sampling → **Representative ?**
- ✓ Automated/extensive sampling → **Time-consuming and expensive**
- ✓ Passive sampling → **Validity of quantitative results ?**



$$Cf(t) = \frac{C_{POCIS}}{C_{water}} = \frac{k_u}{k_e} (1 - e^{-k_e t})$$

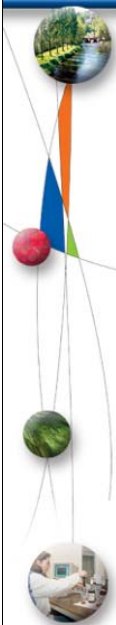
Equilibrium regime

$$Cf = \frac{k_u}{k_e} = K_{sw}$$

Linear uptake regime

$$Cf(t) = k_u \times t$$

Assumption of linear isotherms and isotropic exchanges



1. Methodologies

Analytical techniques and analyte recoveries

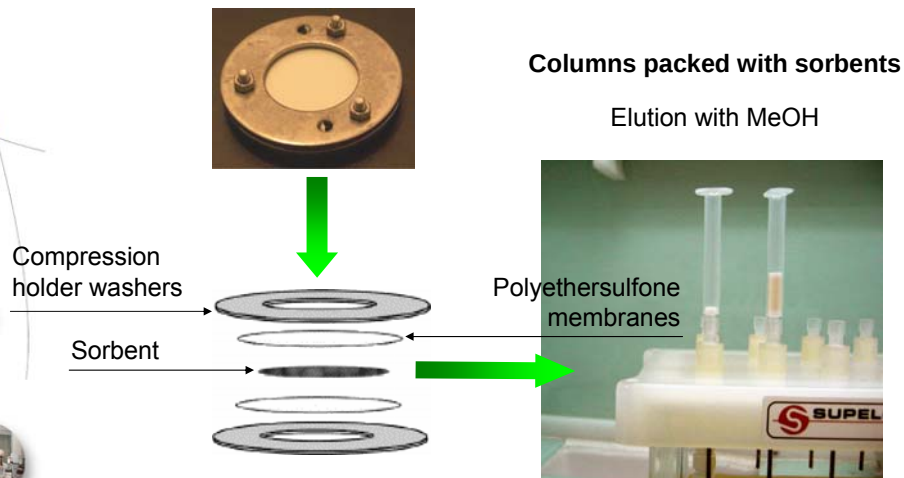
2. Calibration of pharmaceutical POCIS

Determination of k_u and R_s

3. Study of analyte elimination rates

Suggestion of a potential PRC

Analyte recoveries





Herbicides analyzed

Large polarity range : -1.7 (Nicosulfuron) $\leq \text{Log Kow} \leq 5.07$ (trifluraline)

Different classes : Triazines, Phenylureas, Sulfonylureas, Diphenyl ethers, Chloroacetanilides, Dinitroanilines, Benzonitriles, Triketones

Triazines and Phenylureas metabolites

Analytical method

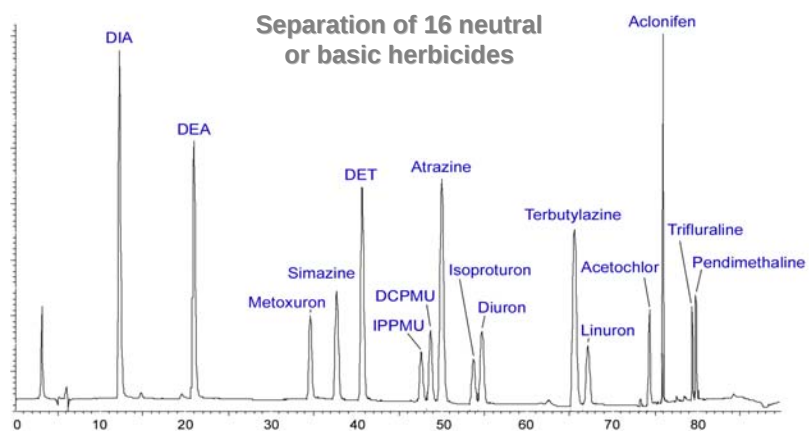
2 multiresidue methods :

HPLC-DAD technique

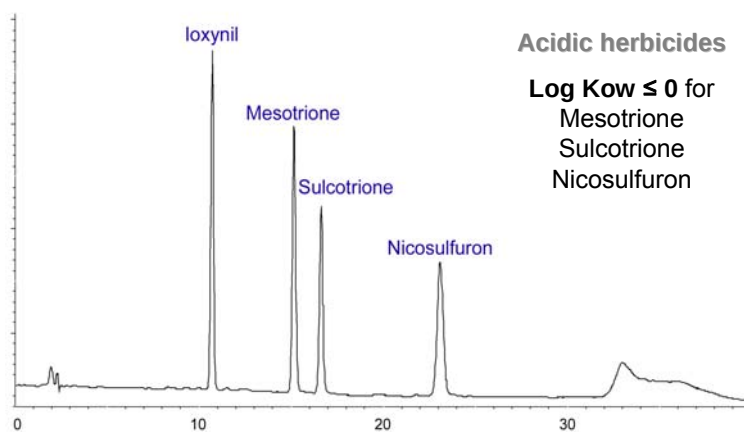
Reverse phase (ODS 2) and weakly anion-exchanger (polyamine) columns



Analysis with an ODS 2 column



Analysis with a polyamine column



Acidic herbicides

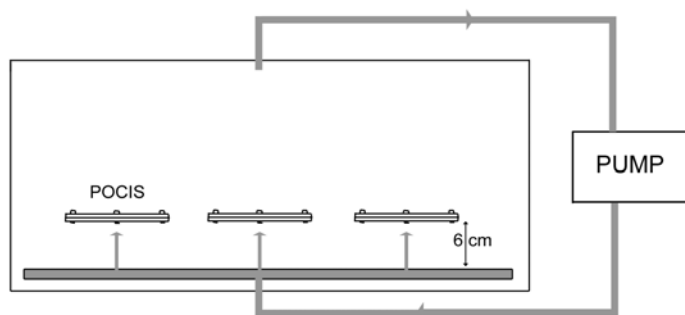
Log K_{ow} ≤ 0 for

Mesotrione

Sulcotrione

Nicosulfuron

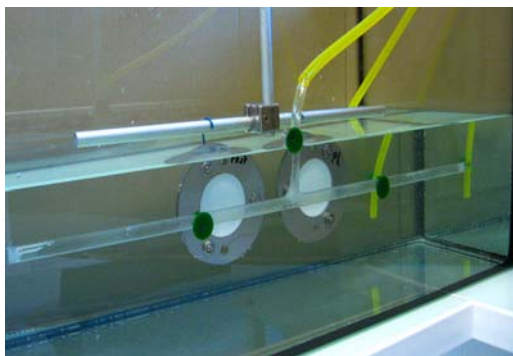
Microcosm experimental design



Constant temperature ($17 \pm 1^\circ \text{C}$) and obscurity

Large volume (80 L)

Turbulent conditions : constant flow velocity $2\text{-}3 \text{ cm.s}^{-1}$



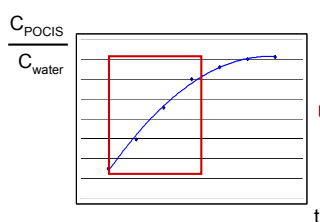
Microcosm experiment

Constant temperature and flow velocity, obscurity

Tap water

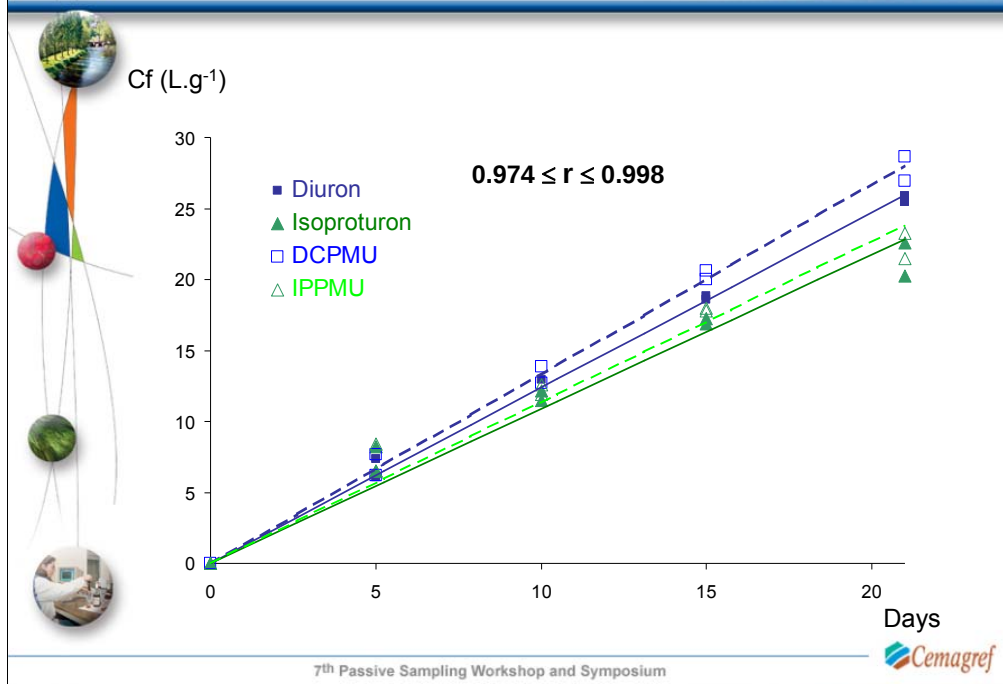
Constant concentrations over **21 days** of exposure

Duplicate analyses



1. Linearity and kinetic regimes

2. Sampling rates R_s



Sampling rates over 21 days

log K_{ow}

-1.7

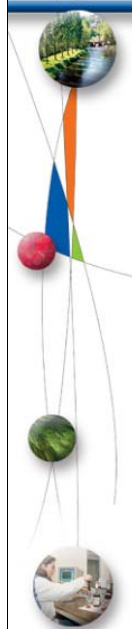


3.03

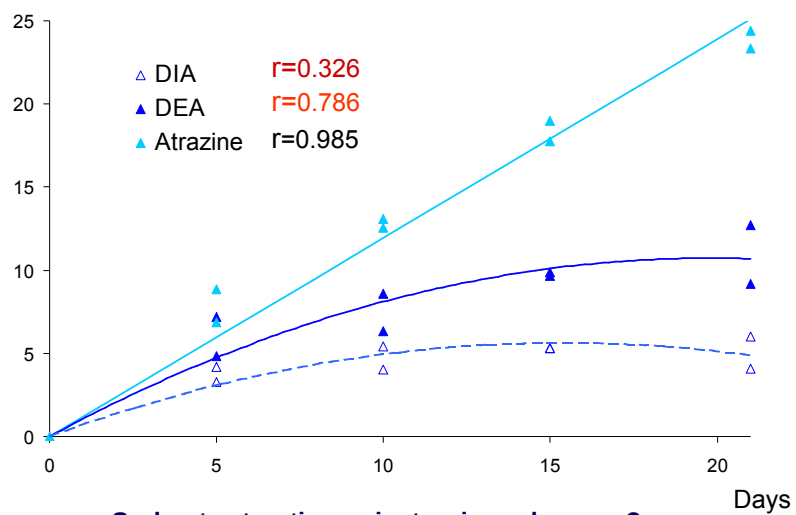
5.18

Herbicides	ku (mL g ⁻¹ d ⁻¹)	Rs (mL d ⁻¹)	% RSD	Linearity over 21 days
Nicosulfuron	219	43.9	3.5	0.910
Sulcotrione	146	29.3	23.8	0.806
Mesotrione	104	20.8	13.0	0.391
Ioxynil	884	176.8	6.2	0.971
DIA	318	63.6	16.5	0.326
DEA	608	121.5	16.5	0.786
Metoxuron	989	197.7	2.4	0.950
Simazine	1051	210.3	0.6	0.955
DET	1025	205.0	2.7	0.974
IPPMU	1135	226.9	3.4	0.981
DCPMU	1334	266.9	3.4	0.998
Atrazine	1195	239.0	3.4	0.985
Isoproturon	1088	217.6	4.7	0.977
Diuron	1236	247.3	0.1	0.998
Terbutylazine	1253	250.7	3.8	0.994
Linuron	1179	235.9	0.4	0.996
Acetochlor	1126	225.2	7.2	0.989

Pendimethalin



Cf (L.g⁻¹)



Sorbent saturation or isotropic exchanges ?

Partial conclusion

Linear uptakes of 16 herbicides for 21 days

Curvilinear uptakes of **DIA** and 3 other polar herbicides :

- Assumption of isotropic exchanges for DIA
- Estimation of the elimination rate constant
- Use of DIA as PRC ?

$$C_w = \frac{C_{\text{POCIS}} \times M_{\text{POCIS}}}{R_s \times t}$$

Rs depends on : Temperature

Turbulence

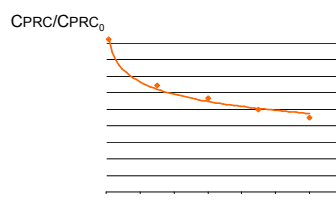
Biofouling

Performance Reference Compound

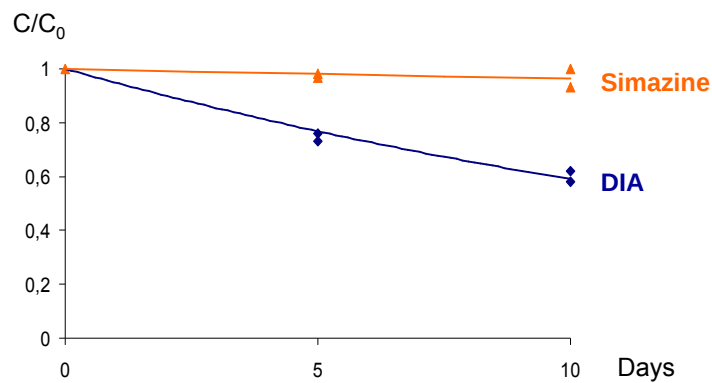
$$K_{\text{SW}} = \frac{k_u}{k_e}$$

$$k_{e\text{PRC}} = \ln(C_{\text{PRC}_0}/C_{\text{PRC}})/t$$

$$R_{s\text{ in situ}} = (k_{e\text{ in situ}}/k_{e\text{ cal}}) \times R_{s\text{ cal}}$$



Dissipation of DIA and simazine from pharmaceutical POCIS

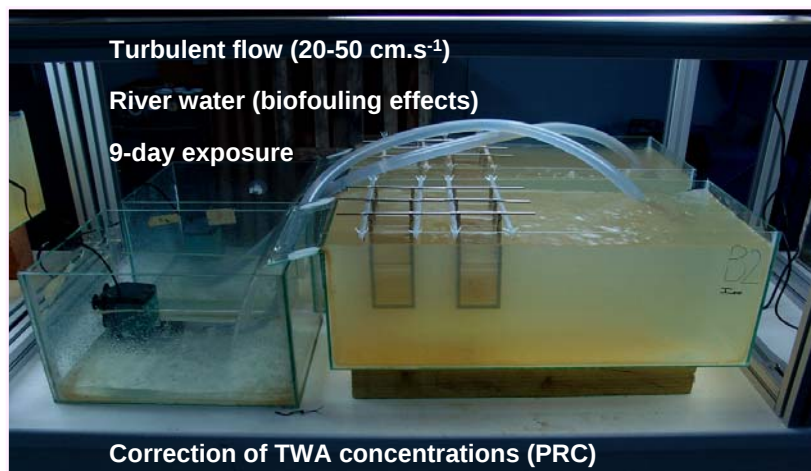


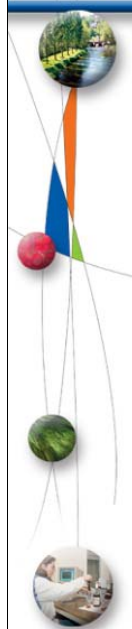
Elimination rate constant estimates :

$k_e = 0.047 \pm 0.005 \text{ d}^{-1}$ (dissipation) and $k_e = 0.08 \pm 0.02 \text{ d}^{-1}$ (uptake)

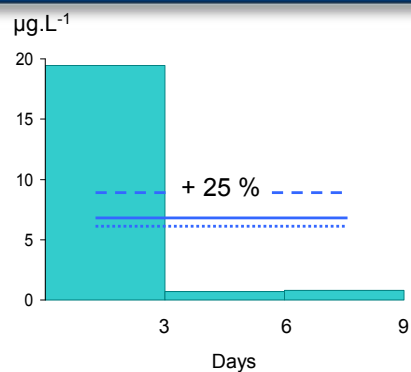
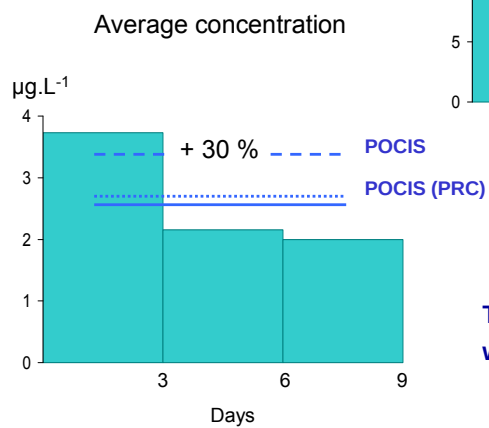
Isotropic exchanges might exist for DIA

Microcosm experiment



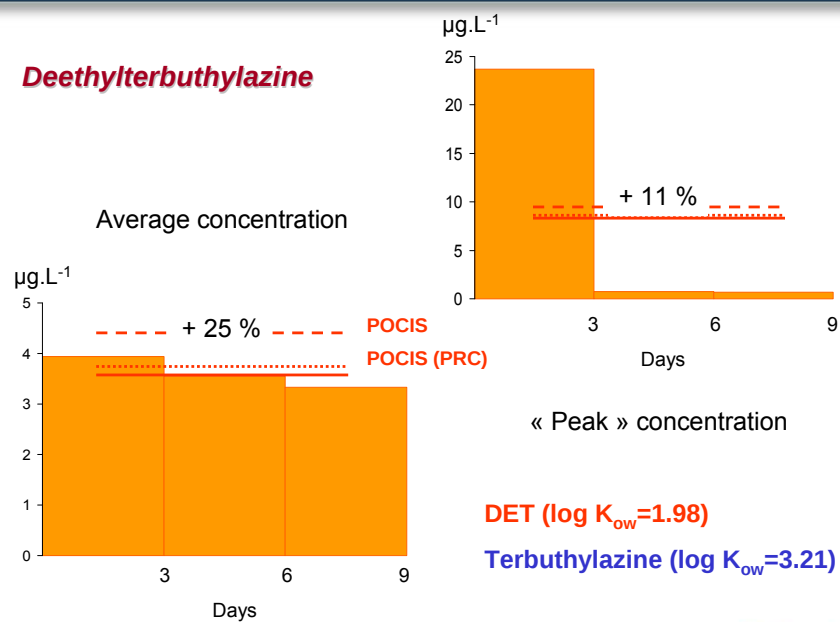


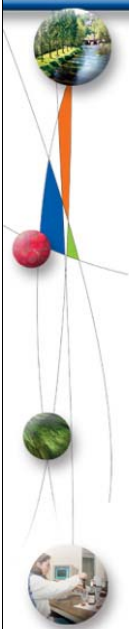
Terbutylazine



The « peak » concentration was integrated during 9 days

Deethylterbuthylazine





Isotropic exchanges for DIA (first-order monophasic kinetic)

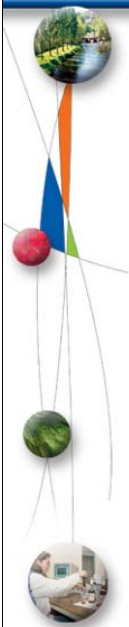
« Peak » concentrations are integrated for the 2 test chemicals

Correction of TWA concentration estimates for 9 days

**Desorption of DIA and other herbicides for 28 days
(biphasic kinetics)**

Correction of TWA concentrations for 21-28 days

***In situ* validation with one or several PRCs**



Thank you for your attention !