

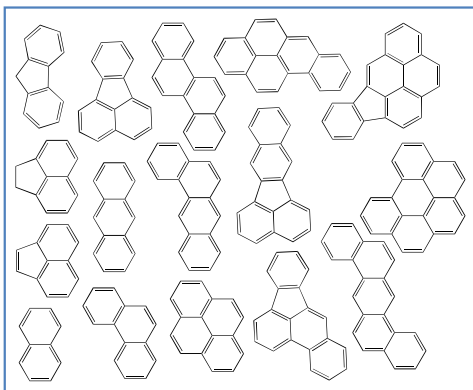
Phase equilibrium application example:

# Passive Samplers

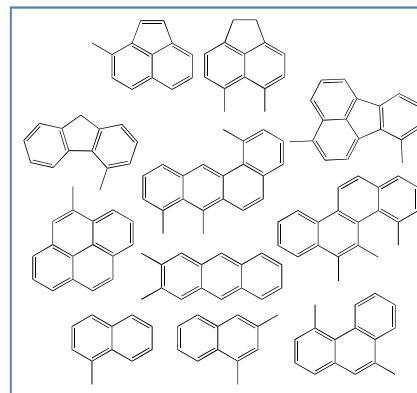
# Hydrophobic organic contaminants (HOCs)

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- **Low water solubility ( $\log K_{ow} \geq \sim 4$ )**
- **Major classes of HOCs of concern**
  - Polycyclic aromatic hydrocarbons (PAHs)
    - Generated by incomplete combustion
    - Found in urban areas (vehicles, etc.), thermal power plants, coal gasification plants, petroleum-related facilities, etc.
    - Main focus has been on parent-PAHs, but alkylated-PAHs can be also of interest (similar behavior and comparable toxicity)
    - Alkylated-PAHs dominant in petroleum (>90% of total)



Parent-PAHs



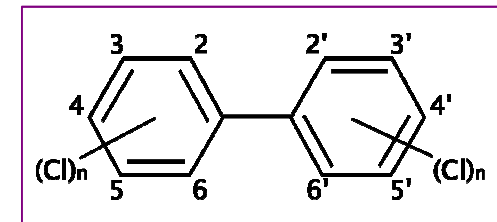
Alkylated-PAHs

# HOCs (cont'd)

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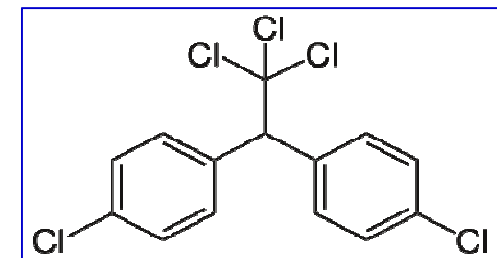
- Polychlorinated biphenyls (PCBs)

- Used as insulator, plasticizer, flame retardants, etc.
- Banned in the 1970s
- But still of concern (highly persistent, endocrine disrupting)



- Chlorinated pesticides

- DDT and its metabolites
- Methoxychlor, dieldrin, chlordane, toxaphene, ...
- Highly persistent



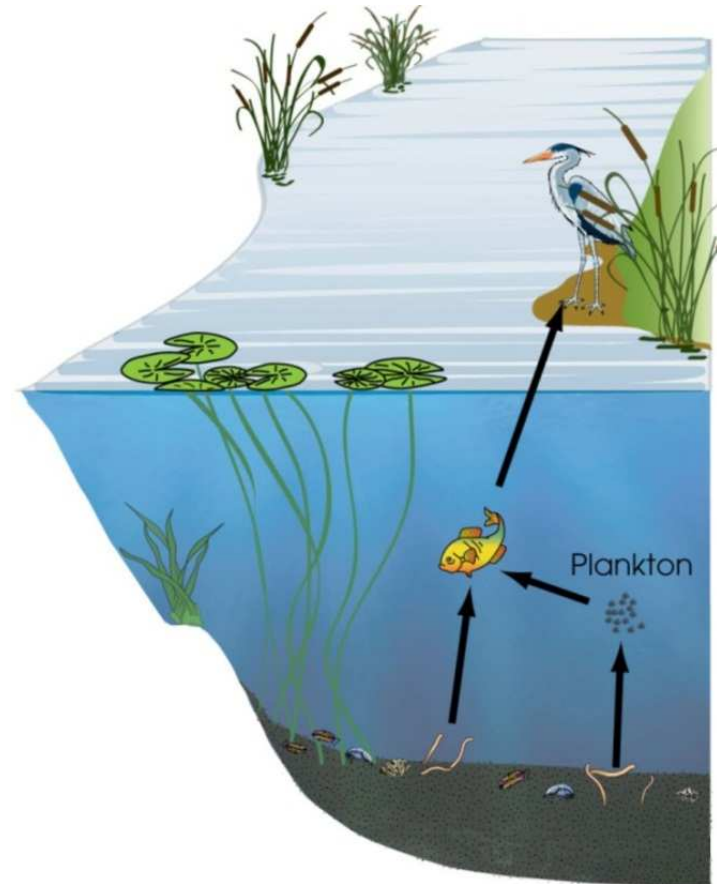
- **Significance of HOC contamination**

- PCBs & PAHs are major risk-driving contaminants at contaminated sites in the U.S. (USEPA, 2005)
  - PCBs – one of risk drivers for 58% of sediment sampling stations in the U.S. where adverse effects are probable
  - PAHs – 8%

# HOCs in sediment

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- In the presence of an external contaminant source, sediment acts as a repository of HOCs
- When the external source is eliminated, sediment acts as an HOC source
- Enters the food chain either by sediment intake by benthic organisms or release to water column



Ghosh et al. *ES&T Feature*, 2011<sup>#3</sup>

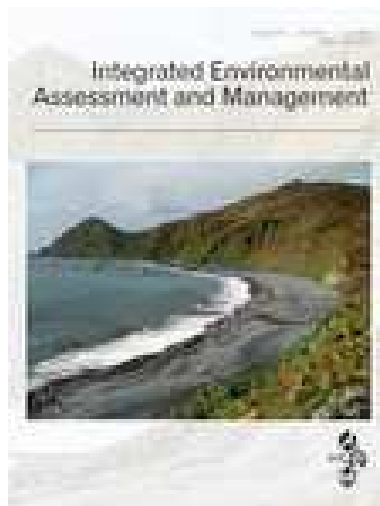
# SETAC Workshop 2012



- Society of Environmental Toxicology & Chemistry (SETAC) 2012 Workshop
- 40+ faculty-level researchers
- Passive sampling guideline series in IEAM



#4



Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 163–166  
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## Passive Sampling in Contaminated Sediment Assessment: Building Consensus to Improve Decision Making

Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 167–178  
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## Passive Sampling Methods for Contaminated Sediments: State of the Science for Organic Contaminants

Michael J Lydy,\*† Peter F Landrum,‡ Amy MP Oen,‡ Mayumi Allinson,§ Foppe Smedes,|| # Amanda D Harwood,† Huizhen Li,†† Keith A Maruya,‡‡ and Jingfu Liu,§§

Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 179–196  
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## Passive Sampling Methods for Contaminated Sediments: State of the Science for Metals

Willie JGM Peijnenburg,\*†† Peter R Teasdale,‡ Danny Reible,|| Julie Mondon, # William W Bennett,§ and Peter GC Campbell†††

Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 197–209  
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## Passive Sampling Methods for Contaminated Sediments: Scientific Rationale Supporting Use of Freely Dissolved Concentrations

Philipp Mayer,\*†§§§ Thomas F Parkerton,‡ Rachel G Adams,§ John G Cargill,|| Jay Gan, # Todd Gouin,†† Philip M Gschwend,††† Steven B Hawthorne,§§§ Paul Helm,||§§ Gesine Witt, # # Jing You,††† and Beate I Escher†††

†Department of Environmental Science, Faculty of Science and Technology, Aarhus University, Roskilde, Denmark

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## Passive Sampling Methods for Contaminated Sediments: Risk Assessment and Management

Marc S Greenberg,† Peter M Chapman,\*†† Ian J Allan,§ Kim A Anderson,|| Sabine E Apitz,‡ Chris Beegan,††† Todd S Bridges,††† Steve S Brown,§§ John G Cargill IV,||§§ Megan C McCulloch, # #†† Charles A Menzie,††† James P Shine,§§§ and Thomas F Parkerton||§§§

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Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 210–223  
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## Passive Sampling Methods for Contaminated Sediments: Practical Guidance for Selection, Calibration, and Implementation

Unal Ghosh,\*†† Susan Kane Driscoll,‡ Robert M Burgess,§ Michiel TO Jonker,|| Danny Reible, # Frank Gobas,††† Yongju Choi,‡† Sabine E Apitz,§§ Keith A Maruya,||§§ William R Gala, # # Munro Mortimer,††† and Chris Beegan†††

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||Southern California Coastal Water Research Project Authority, Costa Mesa, California, USA

#Chevron Energy Technology Company, San Ramon, California, USA

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†††California State Water Board-Division of Water Quality, Sacramento, California, USA

#5-#10

5

# SETAC Workshop 2012 - Consensus

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- **Consensus: absolute HOC concentration in sediment ( $C_{sed}$ ) is NOT a valid indicator of HOC risk**
- **$C_{free}$  as an alternative indicator**
  - **Freely-dissolved aqueous HOC concentration**: refers to the dissolved form of HOCs in water that is not associated with dissolved organic matter (DOM) or other species
  - Consensus: (at least) **a better indicator or HOC bioavailability than  $C_{sed}$**
  - $C_{free}$  is a direct indicator of “chemical activity” (or fugacity) in a multi-phase system
    - At equilibrium, bioaccumulation of HOCs would be proportional to chemical activity
  - Movement towards the use of  $C_{free}$  for regulation

# $C_{free}$ measurement by pore-water sampling

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- Large volume requirement
  - $C_{free}$  for HOCs is quite low  $\rightarrow$  large volume is needed to ensure detectability
- Disturbance of the equilibrium during collection and analysis
- Difficulty to separate freely-dissolved species from DOM-bound species
- Labor and cost issues



*Comparison of a water sample before (right) and after (left) the DOM removal: DOM removed by a simplified (!!) flocculation technique developed by Ghosh et al. (2000, ES&T)<sup>#11</sup>*

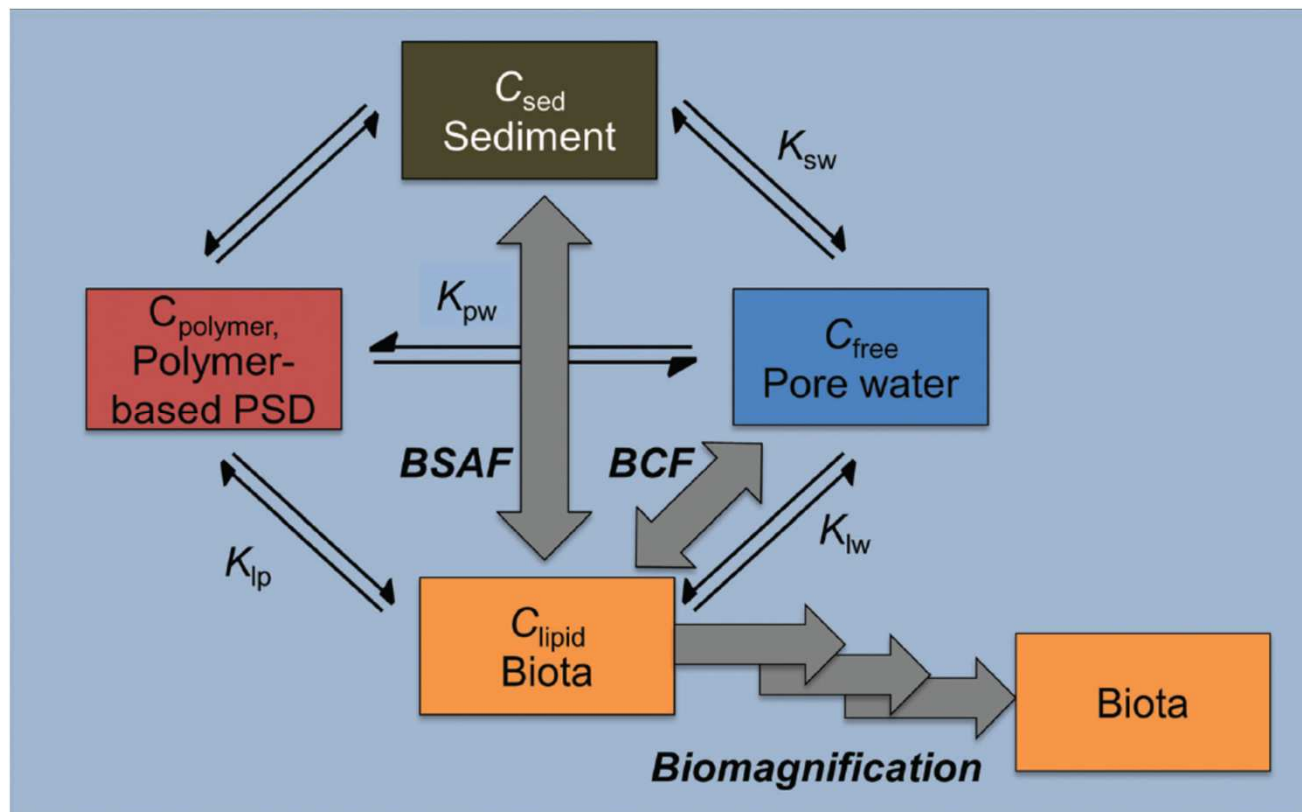
*Now, the remaining steps are: liquid-liquid extraction ( $\geq 3$  times) using hexane, moisture removal using sodium sulfate anhydrous, sample concentration under nitrogen stream, solvent exchange to cyclohexane (for PAHs), cleanup column, another sample concentration, and instrumental analysis*

*(estimated total labor time of  $\sim 10$  hrs/sample, except for time required for sampling)*



## Alternative $C_{free}$ measurement: Polymeric passive sampler

- Add a polymeric passive sampler into the sediment, collect the sampler after a certain period of exposure, determine  $C_{polymer}$ , and use the value to calculate  $C_{free}$





# Polymeric passive samplers

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- Low-density polyethylene (LDPE), polyoxymethylene (POM), polydimethylsiloxane (PDMS), ...
- Linear HOC partitioning between polymer and water
- Large partitioning coefficient → can accumulate substantial amount of HOCs → remarkably enhanced sensitivity
- Inexpensive and easily available
- Easy to analyze HOC concentration (using a simple extraction technique and minimal cleanup requirements)
- Versatile applications: can fit into any scaffold



# Application example (1)

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*Measurement of depth-profile of sediment pore-water HOC concentrations in the field*

## Application example (2)

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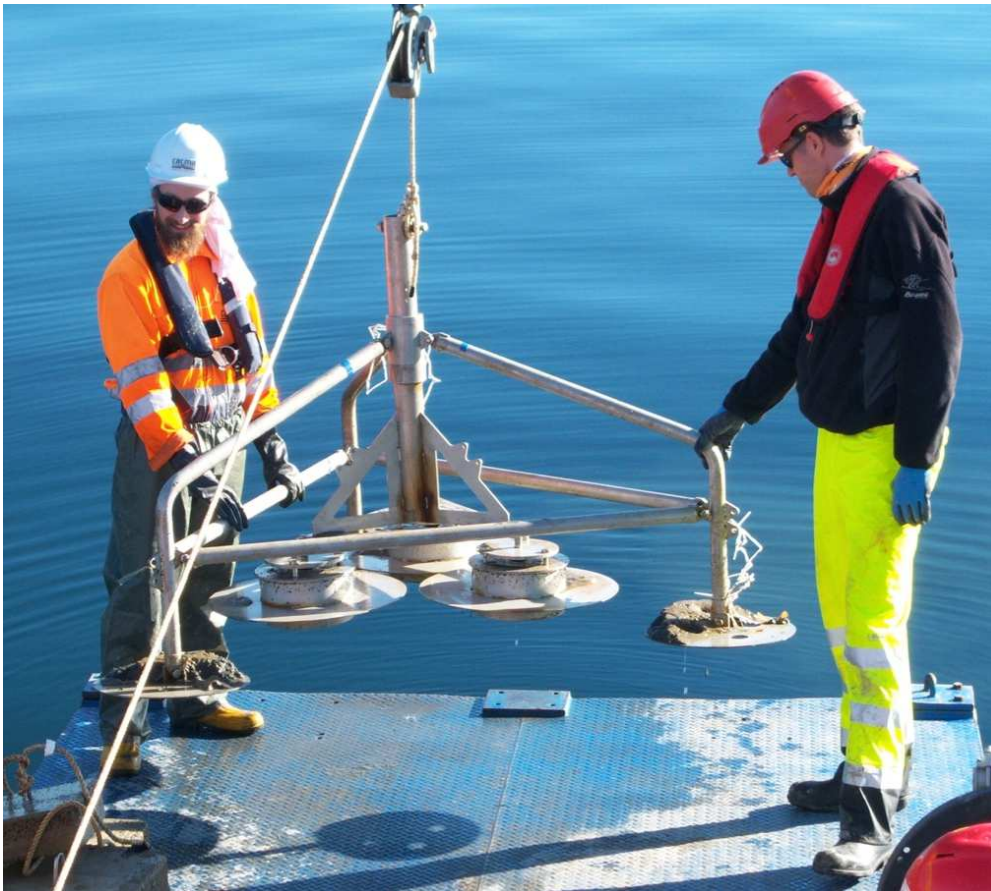


*Measurement of aqueous HOC concentration in the lab*



## Application example (3)

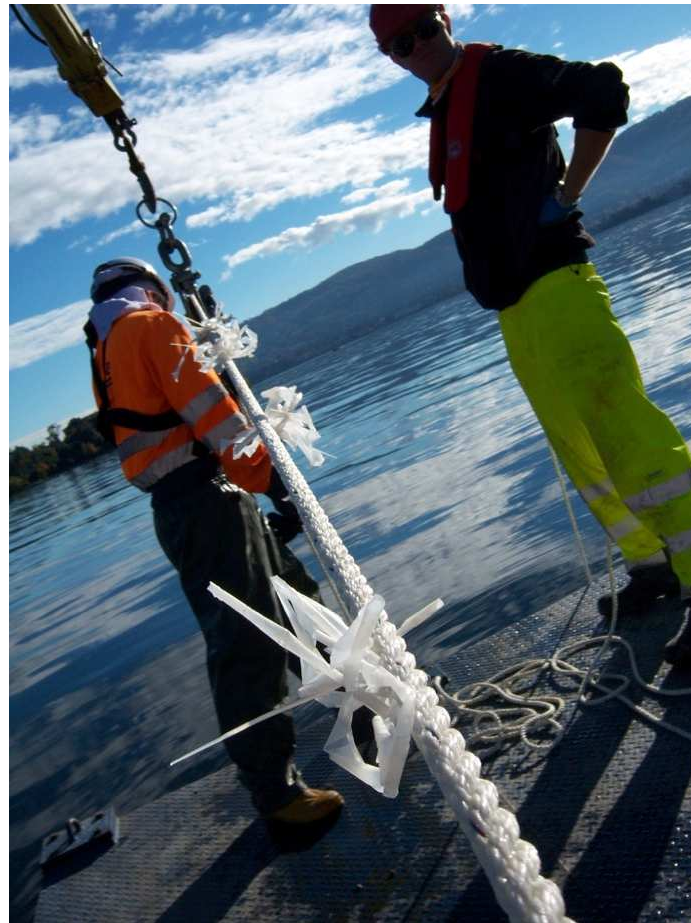
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*Sediment-water HOC flux measurement  
(courtesy: Diana Lin, Stanford Univ.)*

# Application example (4)

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*Depth-profile of HOC  $C_{free}$  in the water column  
(courtesy: Diana Lin, Stanford Univ.)*

# Sampling approaches (1)

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- Equilibrium sampling
  - Exposure the passive sampler to the environment for a duration that is sufficient to achieve equilibrium for HOC partitioning
  - Applicable for laboratory deployment (or field deployment for relatively small/less hydrophobic contaminants)
  - Basic concept

$$C_{polymer,eq} = K_{polymer-water} \cdot C_{free}$$

$$\Rightarrow C_{free} = C_{polymer,eq} / K_{polymer-water}$$

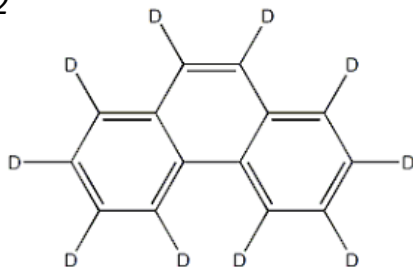
- Ways to ensure that the exposure duration is sufficient for equilibrium
  - Collect passive samplers at different exposure durations  
[equilibrium if  $C_{polymer} \neq f(time)$ ]
  - Deploy passive samplers with different thicknesses  
[equilibrium if  $C_{polymer} \neq f(thickness)$ ]

# Sampling approaches (2)

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- Non-equilibrium sampling
  - Equilibration time estimated for typical HOCs (e.g., PCBs, PAHs) in the field: months to years to decades
  - Equilibrium sampling not practically viable in most cases of field applications
- Current approach for non-equilibrium passive sampling
  - Use of “performance reference compounds (PRCs)”
    - PRCs: analytically non-interfering chemicals that are embedded in the passive sampler prior to environmental exposure (Ghosh, ..., Choi et al., *IEAM*, 2014)
  - ex) contaminant: phenanthrene → PRC: d10-phenanthrene

#12

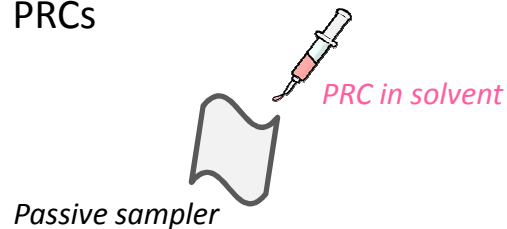


*d10-phenanthrene structure. All hydrogens are substituted with deuterons.*



# Sampling approaches (2) – cont'd

- 1) Impregnate passive sampler with PRCs



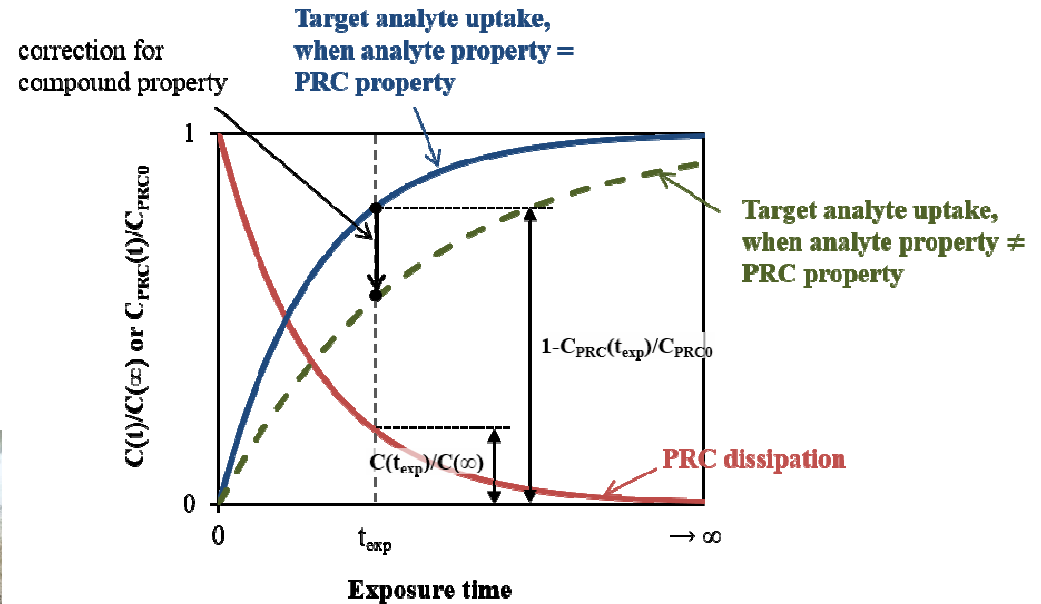
- 2) Expose the sampler to the target media



- 3) After exposure, analyze both the target contaminant and PRC concentration in the sampler  $[C(t_{exp}), C_{PRC}(t_{exp})]$



- 4) Calculate the fraction of the PRC released out of sampler  $[1 - C_{PRC}(t_{exp})/C_{PRC0}]$



- 5) Calculate the fraction of the target contaminant accumulated in the sampler compared to the equilibrium value  $[C(t_{exp})/C(\infty)]$
- 6) Calculate target contaminant concentration at equilibrium  $[C(\infty)]$
- 7) Obtain  $C_{free} [= C(\infty)/K_{polymer-water}]$

# Addt'l note for non-eq. sampling

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- PRC method reasonably works for most situations, but has been criticized by some researchers
- Basic assumption of the PRC method: “isotropic exchange kinetics”
  - The rate of PRC dissipation from the sampler is expressed in exactly the same way as the rate of target contaminant accumulation by the sampler
- Experiments show that the isotropic exchange kinetics assumption may not hold
  - Because of the resistance of a PRC for dissipation from the sampler
  - More significantly, because of the resistance of a target contaminant for desorption from the sediment

# Some relevant publications

#13



Article  
pubs.acs.org/est

## Polyethylene–Water Partitioning Coefficients for Parent- and Alkylated-Polycyclic Aromatic Hydrocarbons and Polychlorinated Biphenyls

Yongju Choi<sup>1</sup>, Yeo-Myoung Cho, and Richard G. Luthy\*

<sup>1</sup>Department of Civil and Environmental Engineering, Stanford University, Stanford, California 94305-4020, United States

Journal of Hazardous Materials 318 (2016) 579–586



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journal homepage: [www.elsevier.com/locate/jhazmat](http://www.elsevier.com/locate/jhazmat)



## Non-equilibrium passive sampling of hydrophobic organic contaminants in sediment pore-water: PCB exchange kinetics

Yongju Choi<sup>2,\*</sup>, Yanwen Wu<sup>1</sup>, Richard G. Luthy<sup>1</sup>, Seju Kang<sup>2</sup>

<sup>1</sup> Department of Civil and Environmental Engineering, Seoul National University, Seoul 0826, Republic of Korea  
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#9

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## Passive Sampling Methods for Contaminated Sediments: Practical Guidance for Selection, Calibration, and Implementation

Upal Ghosh,<sup>\*,†</sup> Susan Kane Driscoll,<sup>‡</sup> Robert M Burgess,<sup>§</sup> Michiel TO Jonker,<sup>||</sup> Danny Reible,<sup>#</sup> Frank Gobas,<sup>††</sup> Yongju Choi,<sup>‡‡</sup> Sabine E Apitz,<sup>§§</sup> Keith A Maruya,<sup>|||</sup> William R Gala,<sup>##</sup> Munro Mortimer,<sup>†††</sup> and Chris Beegan<sup>†††</sup>

<sup>\*</sup>Department of Chemical, Biochemical, and Environmental Engineering, University of Maryland Baltimore County, Baltimore, Maryland, USA  
<sup>†</sup>Exponent, Maynard, Massachusetts, USA  
<sup>‡</sup>USEPA, Office of Research and Development, Narragansett, Rhode Island, USA  
<sup>§</sup>Institute for Risk Assessment Sciences, Utrecht University, Utrecht, the Netherlands  
<sup>||</sup>Department of Civil and Environmental Engineering, Texas Tech University, Lubbock, Texas, USA  
<sup>††</sup>School of Resource and Environmental Management, Simon Fraser University, Burnaby, British Columbia, Canada  
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<sup>##</sup>Chevron Energy Technology Company, San Ramon, California, USA  
<sup>†††</sup>National Research Centre for Environmental Toxicology, The University of Queensland, Brisbane, Australia  
<sup>††††</sup>California State Water Board-Division of Water Quality, Sacramento, California, USA

#14



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One This: Environ. Sci. Technol. 2016, 52, 3574–3582



#15

## Advancing the Use of Passive Sampling in Risk Assessment and Management of Sediments Contaminated with Hydrophobic Organic Chemicals: Results of an International Ex Situ Passive Sampling Interlaboratory Comparison

Michiel T. O. Jonker,<sup>\*,†</sup> Stephan A. van der Heijden,<sup>‡</sup> Dave Adelman,<sup>‡</sup> Jennifer N. Apell,<sup>§</sup> Robert M. Burgess,<sup>||</sup> Yongju Choi,<sup>††</sup> Loretta A. Fernandez,<sup>‡‡</sup> Geanna M. Flavetta,<sup>‡‡</sup> Upal Ghosh,<sup>‡‡</sup> Philip M. Gschwend,<sup>‡‡</sup> Sarah E. Hale,<sup>‡‡</sup> Mehregan Jalalizadeh,<sup>‡‡</sup> Mohammed Khairy,<sup>‡‡</sup> Mark A. Lampi,<sup>‡‡</sup> Wenjian Lao,<sup>‡‡</sup> Rainer Lohmann,<sup>‡‡</sup> Michael J. Lydy,<sup>‡‡</sup> Keith A. Maruya,<sup>‡‡</sup> Samuel A. Nuttle,<sup>‡‡</sup> Amy M. P. Oen,<sup>‡‡</sup> Magdalena I. Rakowska,<sup>‡‡</sup> Danny Reible,<sup>‡‡</sup> Tatsiana P. Rusina,<sup>‡‡</sup> Foppe Smedes,<sup>‡‡</sup> and Yanwen Wu<sup>‡‡</sup>

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