Phase equilibrium application example:
# Passive Samplers

Hydrophobic organic contaminants (HOCs)

- **Low water solubility (log $K_{ow} \geq$ ~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)
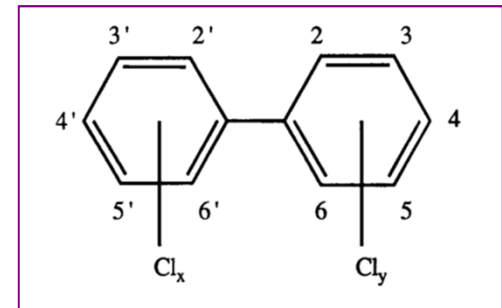   - Alkylated-PAHs dominant in petroleum (>90% of total)
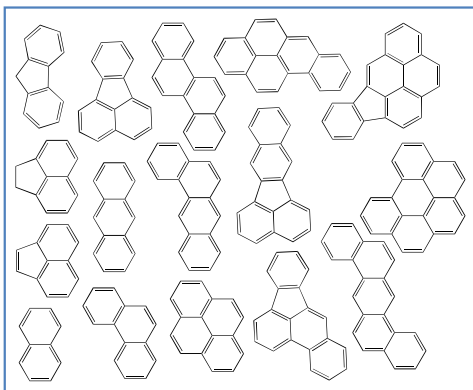
Parent-PAHs
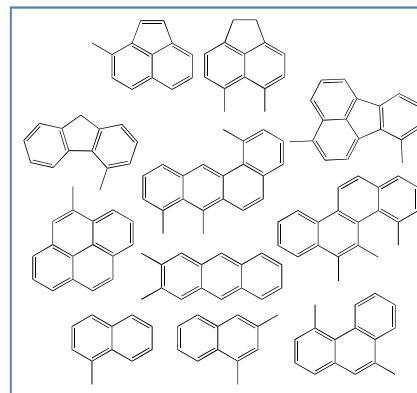
Alkylated-PAHs

HOCs (cont'd)

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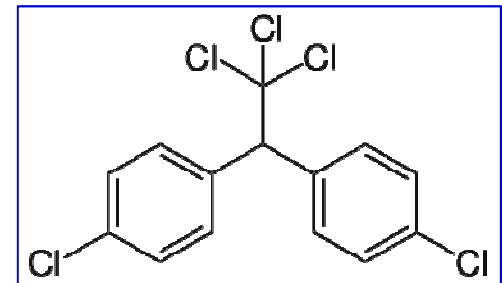
- **Significance of HOC contamination**
  - PCBs & PAHs are two major risk-driving contaminants at U.S. Superfund sites (USEPA, 2005)
    - PCBs – one of risk drivers for 58% of sediment sampling stations in the U.S. where adverse effects are probable
    - PAHs – 8%

HOCs in sediment

- 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

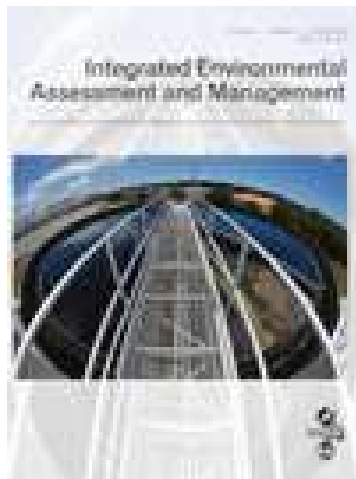- Enters the food chain either by sediment intake by benthic organisms or release to water column
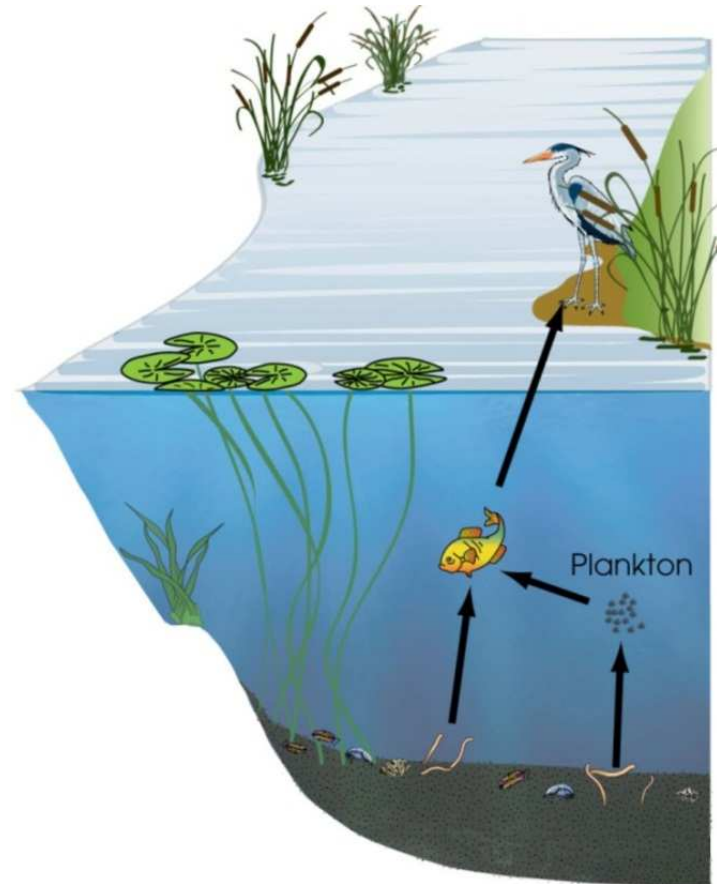
Ghosh, Luthy et al. *ES&T Feature,* 2011

SETAC Workshop 2012

- Society of Environmental Toxicology & Chemistry (SETAC) 2012 Workshop

- 40+ faculty-level researchers

- Passive sampling guideline series in IEAM

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

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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§§

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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††

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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

Integrated Environmental Assessment and Management — Volume 10, Number 2—pp. 224–236
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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‖‖‖‖
†USEPA Office of Superfund Remediation & Technology Innovation, Edison, New Jersey

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

Upal 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‡‡‡
†Department of Chemical, Biochemical, and Environmental Engineering, University of Maryland Baltimore County, Baltimore, Maryland, USA
‡Exponent, Maynard, Massachusetts, USA
§USEPA, Office of Research and Development, Narragansett, Rhode Island, USA
‖Institute for Risk Assessment Sciences, Utrecht University, Utrecht, the Netherlands
#Department of Civil and Environmental Engineering, Texas Tech University, Lubbock, Texas, USA
††School of Resource and Environmental Management, Simon Fraser University, Burnaby, British Columbia, Canada
‡‡Department of Civil and Environmental Engineering, Stanford University, Stanford, California, USA
§§SEA Environmental Decisions, The Ford, Little Hadham, Hertfordshire, United Kingdom
‖‖Southern California Coastal Water Research Project Authority, Costa Mesa, California, USA
##Chevron Energy Technology Company, San Ramon, California, USA
†††National Research Centre for Environmental Toxicology, The University of Queensland, Brisbane, Australia
‡‡‡California State Water Board-Division of Water Quality, Sacramento, California, USA

Integrated Environmental Assessment and Management

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SETAC Workshop 2012 - Consensus

- **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

- Large volume requirement
  - C_{free} for HOCs is quite low → 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 <u>simplified</u> (!!) flocculation technique developed by Ghosh et al. (2000, ES&T)

Now, the remaining steps are: liquid-liquid extraction (≥ 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 ~ 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}
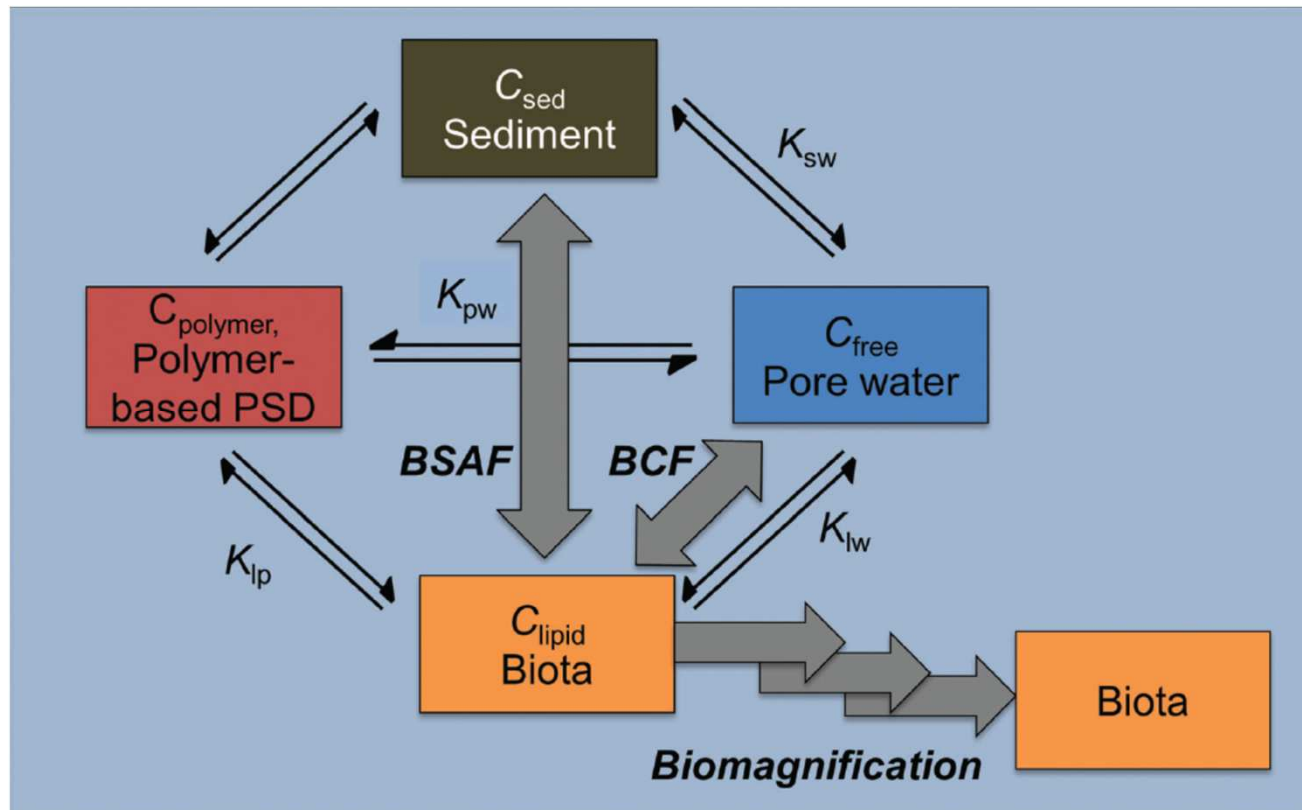
Lydy et al., IEAM, 2014

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Polymeric passive samplers

- 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
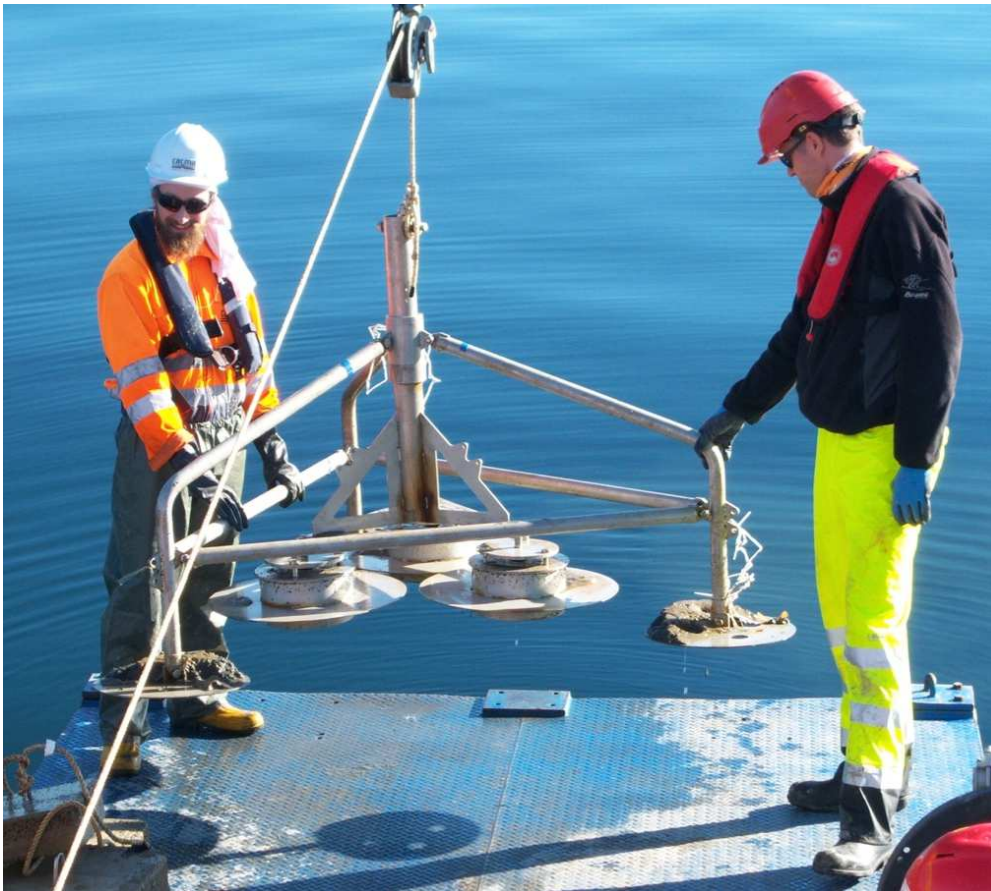
Application example (1)

Measurement of depth-profile of sediment pore-water HOC concentrations in the field
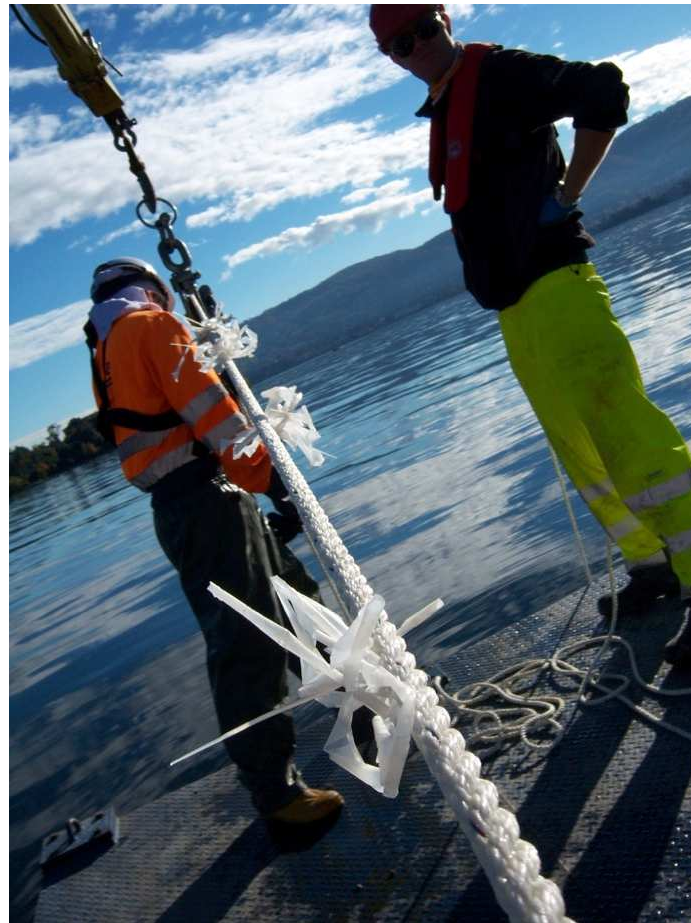
Application example (2)

Measurement of aqueous HOC concentration in the lab

Application example (3)

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

Application example (3)

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

Sampling approaches (1)

- 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

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

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

  - 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)

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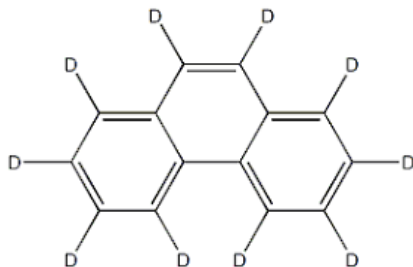
d10-phenanthrene structure. All hydrogens are substituted with deuterons.

Sampling approaches (2) – cont'd

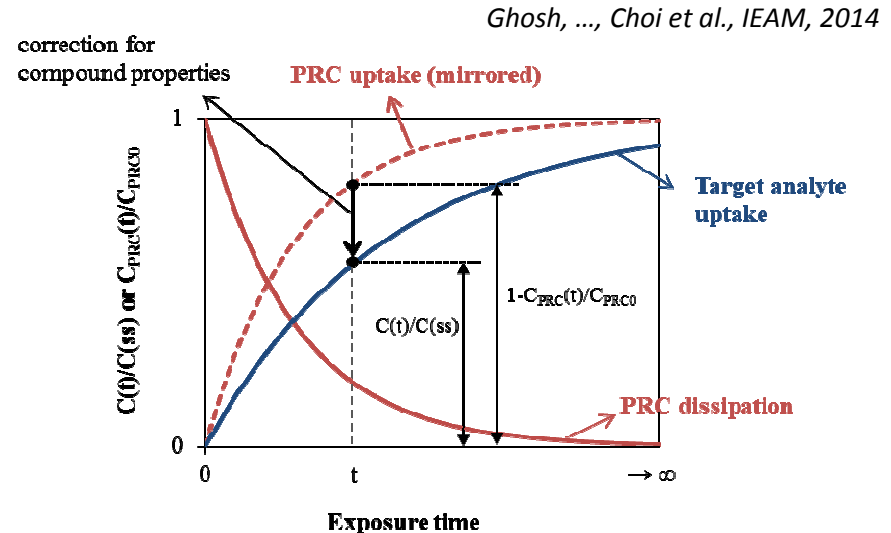1) Impregnate passive sampler with PRCs

PRC in solvent

Passive sampler
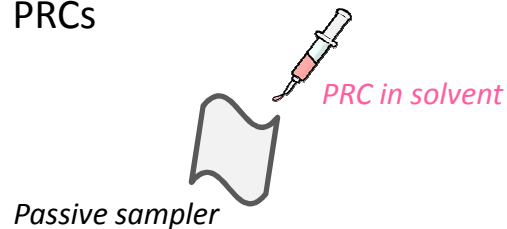
2) Expose the sampler to the target media

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

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

Ghosh, ..., Choi et al., IEAM, 2014

correction for compound properties

PRC uptake (mirrored)

$C(t)/C(ss)$ or $C_{PRC}(t)/C_{PRC0}$

Target analyte uptake

$1 - C_{PRC}(t)/C_{PRC0}$

$C(t)/C(ss)$

PRC dissipation

Exposure time

5) Calculate the fraction of the target contaminant accumulated in the sampler compared to the steady-state (equilibrium) [$C(t)/C(ss)$]

6) Calculate the steady-state (equilibrium) target contaminant concentration [$C(ss)$]

7) Obtain C_{free} [$= C(ss)/K_{polymer\text{-}water}$]

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Addt'l note for non-eq. sampling

- 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

ENVIRONMENTAL Science & Technology
Article
pubs.acs.org/est

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

Yongju Choi, Yeo-Myoung Cho, and Richard G. Luthy*

Department of Civil and Environmental Engineering, Stanford University, Stanford, California 94305-4020, United States

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journal homepage: www.elsevier.com/locate/jhazmat

ELSEVIER

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

Yongju Choi [a,*], Yanwen Wu [b], Richard G. Luthy [b], Seju Kang [a]

[a] Department of Civil and Environmental Engineering, Seoul National University, Seoul 08826, Republic of Korea
[b] Department of Civil and Environmental Engineering, Stanford University, Stanford, CA 94305-4020, United States

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†Department of Chemical, Biochemical, and Environmental Engineering, University of Maryland Baltimore County, Baltimore, Maryland, USA
‡Exponent, Maynard, Massachusetts, USA
§USEPA, Office of Research and Development, Narragansett, Rhode Island, USA
‖Institute for Risk Assessment Sciences, Utrecht University, Utrecht, the Netherlands
#Department of Civil and Environmental Engineering, Texas Tech University, Lubbock, Texas, USA
††School of Resource and Environmental Management, Simon Fraser University, Burnaby, British Columbia, Canada
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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,*† Stephan A. van der Heijden,† Dave Adelman,‡ Jennifer N. Apell,§ Robert M. Burgess,‖ Yongju Choi,‡,# Loretta A. Fernandez,▽ Geanna M. Flavetta,▽ Upal Ghosh,○ Philip M. Gschwend, Sarah E. Hale, Mehregan Jalalizadeh,○ Mohammed Khairy,‡,¶ Mark A. Lampi,▲ Wenjian Lao,○ Rainer Lohmann, Michael J. Lydy, Keith A. Maruya,○ Samuel A. Nutile, Amy M. P. Oen, Magdalena I. Rakowska,△ Danny Reible,△ Tatsiana P. Rusina, Foppe Smedes, and Yanwen Wu

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