# Using PAH Compositions in Sediment Pore Water to More Accurately Interpret "Total PAH" Concentrations

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### **Presentation Overview**

- 1 Role of pore water in evaluating and monitoring sediment PAH sites
- Evaluating pore water PAH data generated using different field sampling and laboratory procedures
- 3 Distinguishing dissolved-phase from particle-sorbed PAHs
- 4 Avoid over-estimation of dissolved-phase PAH concentrations



### Background: Why Analyze Pore Water?

### Better represents bioavailable contaminant fraction

- Assess risk to benthic receptors
- Assess groundwater discharge to surface water (along flowpath)
- Characterize sediment-pore water interaction for cap modeling/design
- Monitor remedial success

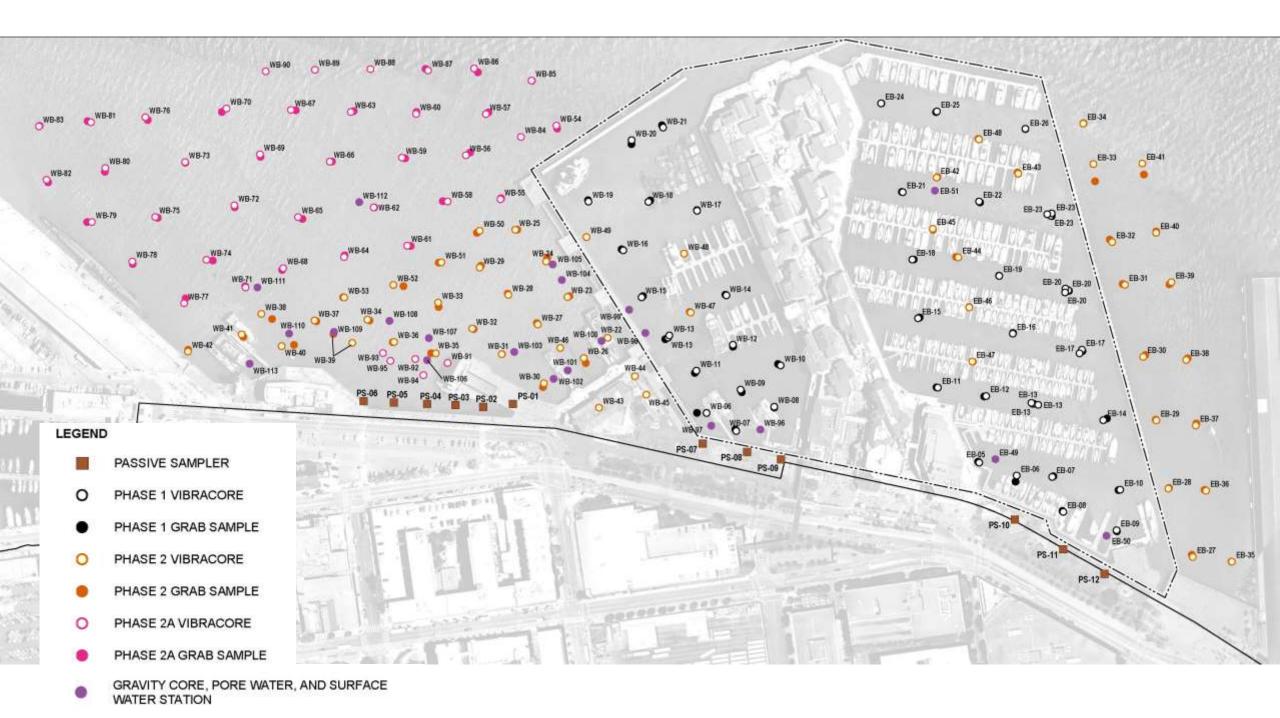
All require distinguishing entrained particle-sorbed PAHs from true dissolved-phase PAHs



### **Confounding Factors**

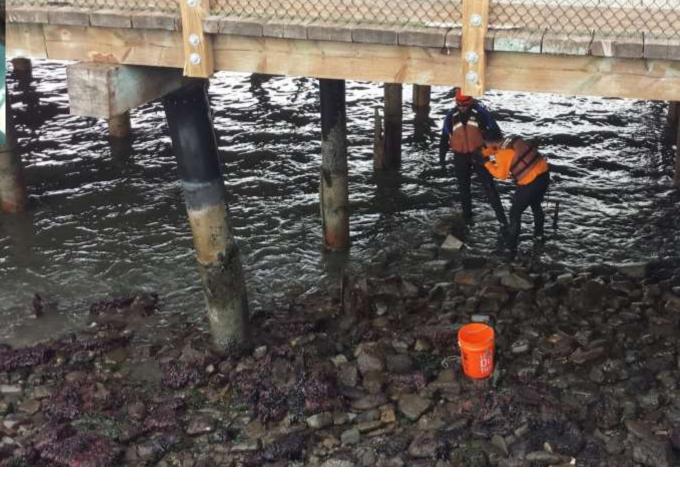
- Sediment pore water equilibrium vs. steady state conditions
  - Evaluate influence from deeper underlying contaminated sediment
- Sampling methods advantages & limitations
  - Passive sampling, depends on equilibrium assumptions, requires concentration adjustment
  - Ex situ extraction via centrifugation assures co-located sediment, limited volume
  - In situ low-flow sampling with piezometers real-time assurance of successful sample
- Particle inclusion: biasing "dissolved-phase" concentrations
  - Particle "flushing" or filtering can help, but may not eliminate sediment from pore water analysis

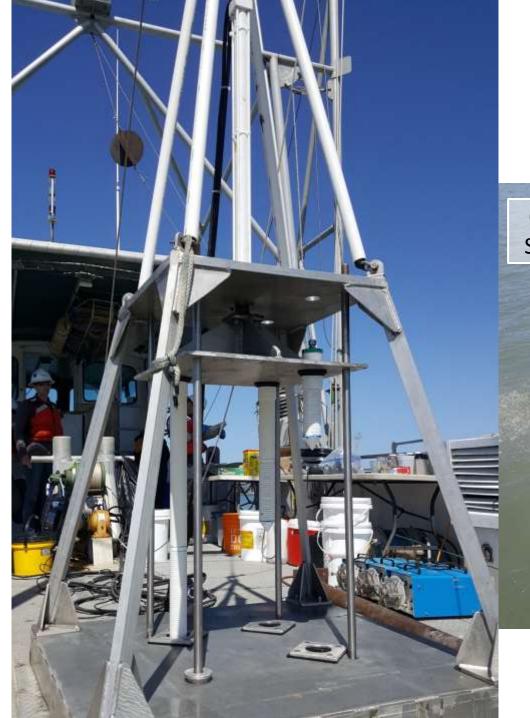




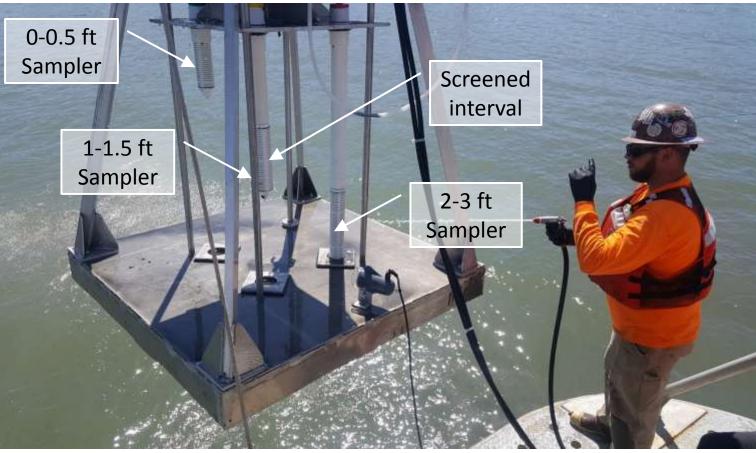


# **Passive Samplers**





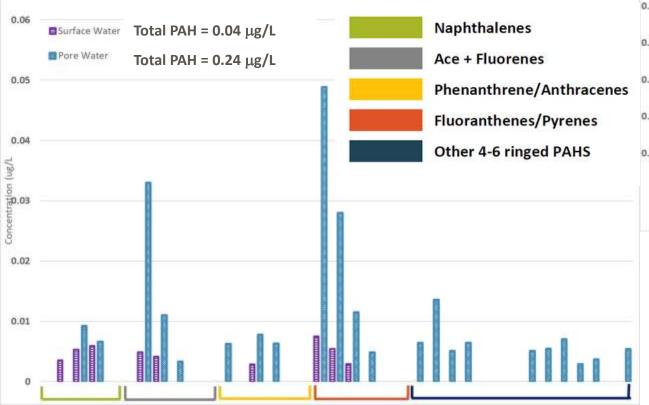
### Pore Water Sampler



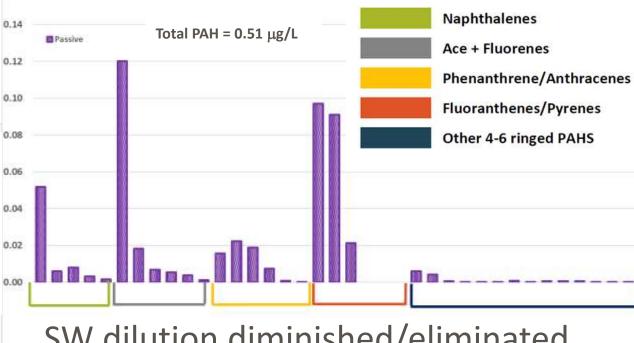
## Comparing Surface Water Measurements using Direct in-situ Surface/Pore Water vs. Passive Sampler Methods

Note dilution in SW PW contains sediment?

### WB-98 - In Situ Surface Water and 0-0.5 ft Pore Water



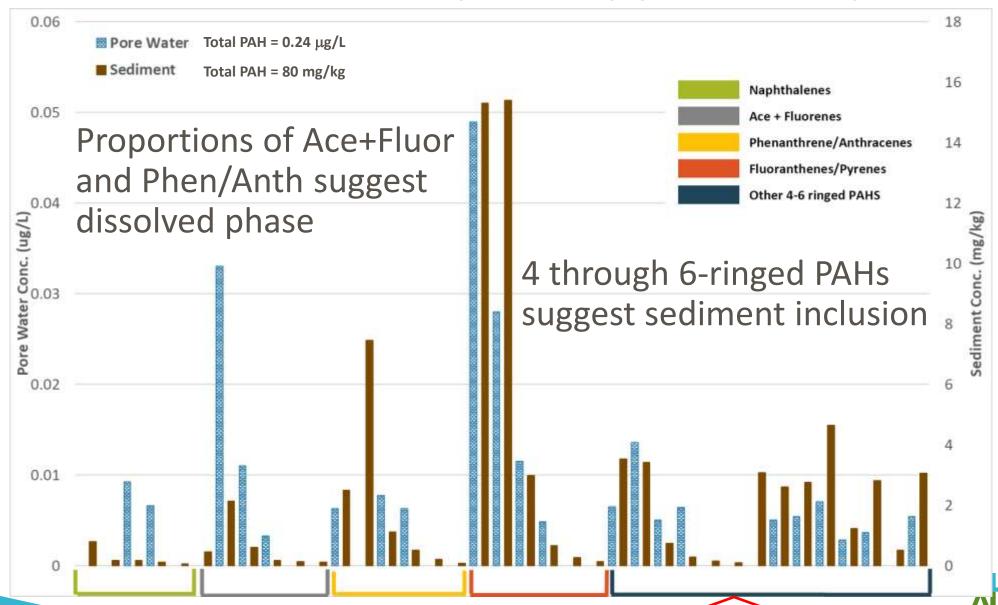
PS-01 – Passive sampler deployed at surface



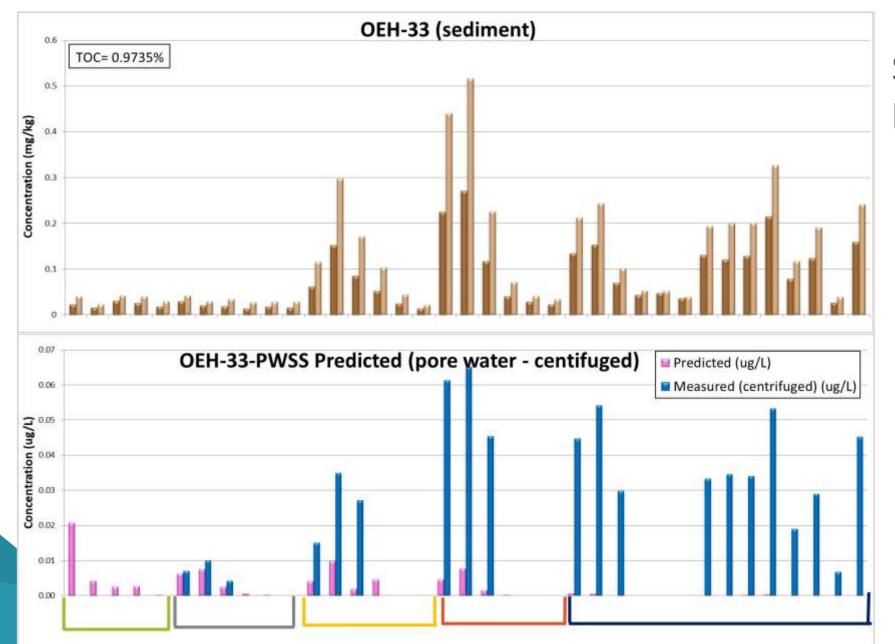
SW dilution diminished/eliminated

Passive sampler at surface and 0-0.5 ft pore water compositions very consistent

### Sediment vs. Pore Water (0-0.5 ft) (Sta WB-98)



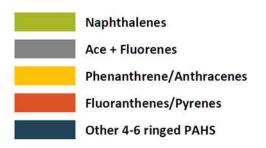
### Pore Water Collected by Centrifugation (Sta OEH-33)



# Sediment inclusion limits data usability

#### **Total PAH**

Sediment = 2.7 mg/kg Pore water = 0.51  $\mu$ g/L





### Why is Dissolved Phase vs Entrained Sediment so Important?

### Risk Assessment

- Dissolved-phase is more bioavailable fraction
- Better suited for comparison with aqueous screening concentrations
- Groundwater discharge to surface water
  - Dissolved phase is mobile, entrained sediment is not (under low flow conditions)
- Sediment pore water interaction for capping design
  - Advection/diffusion modeling assumes dissolved phase
- Post-Remediation Monitoring
  - Performance criteria based on dissolved phase



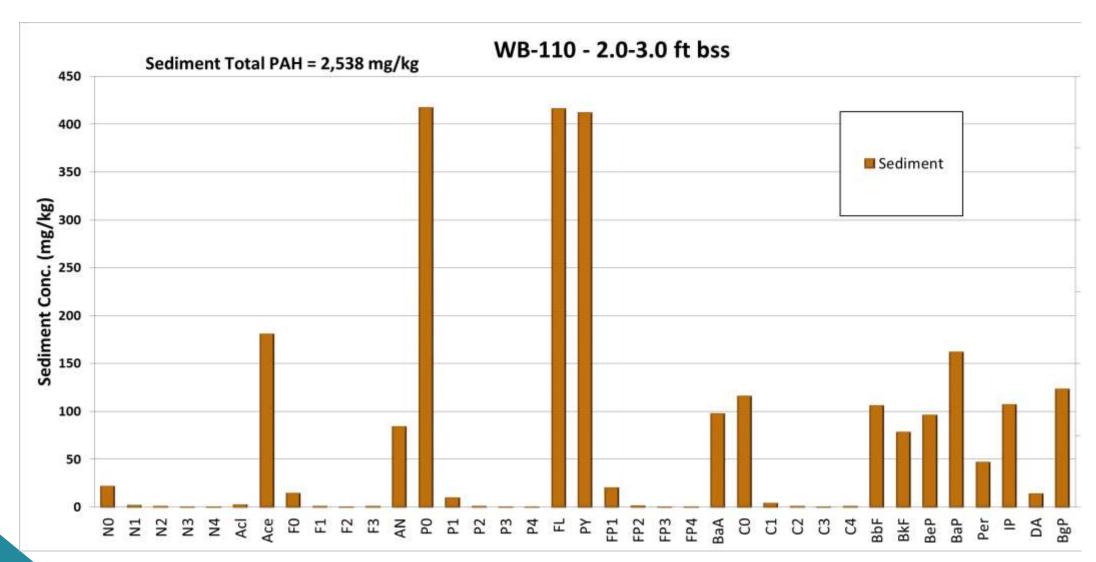
### Approach: Distinguishing Dissolved-Phase PAHs

Use expanded PAH list for compositional interpretation, e.g., "PAH-34"

- 1. Plot PAH compositions in sediment
- 2. Plot PAH compositions in PW <u>predicted</u> applying Equilibrium Partitioning to sediment
- 3. Plot "co-located" measured PAH compositions in PW
- 4. Compare PAH compositions in PW vs predicted PW
  - Tease out dissolved vs particle-sorbed PAHs in PW

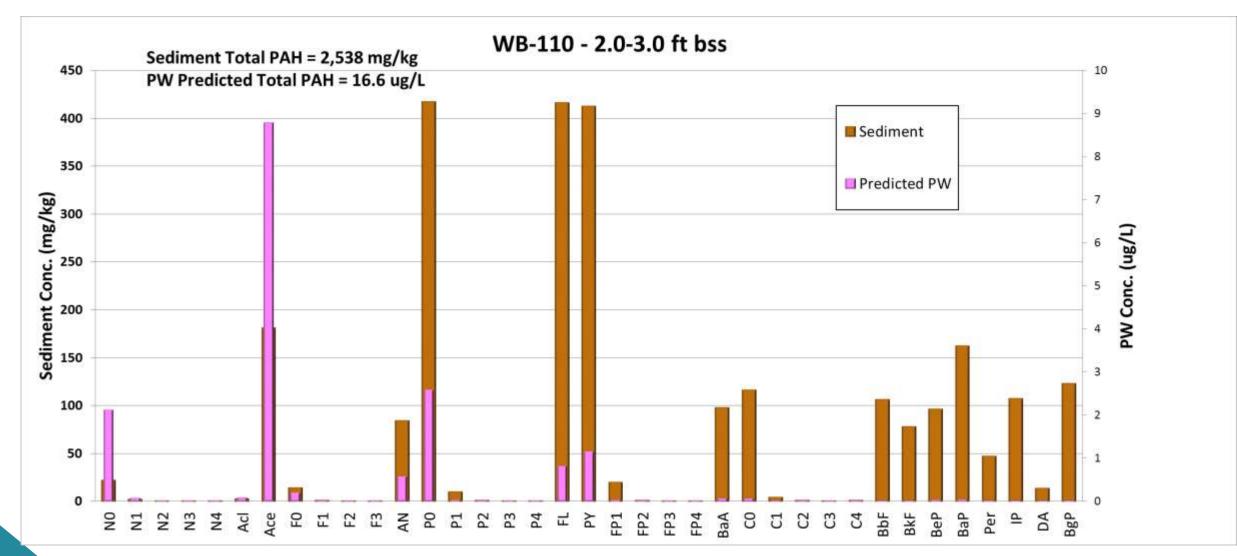


### 1) Plot PAHs in Sediment



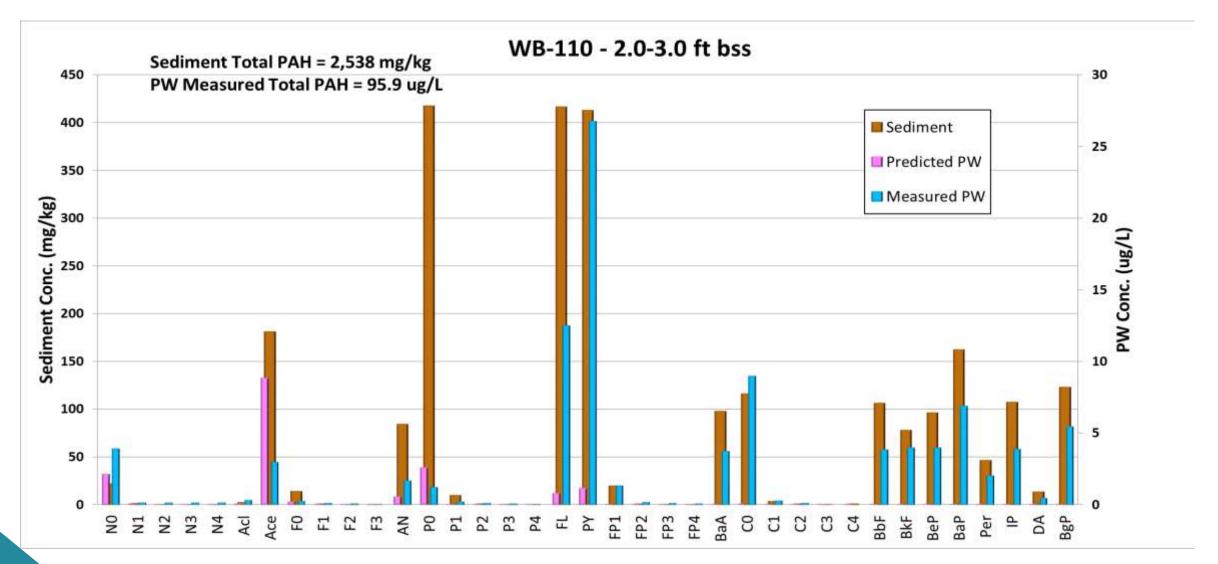


### 2) Plot PAHs Predicted in Pore Water



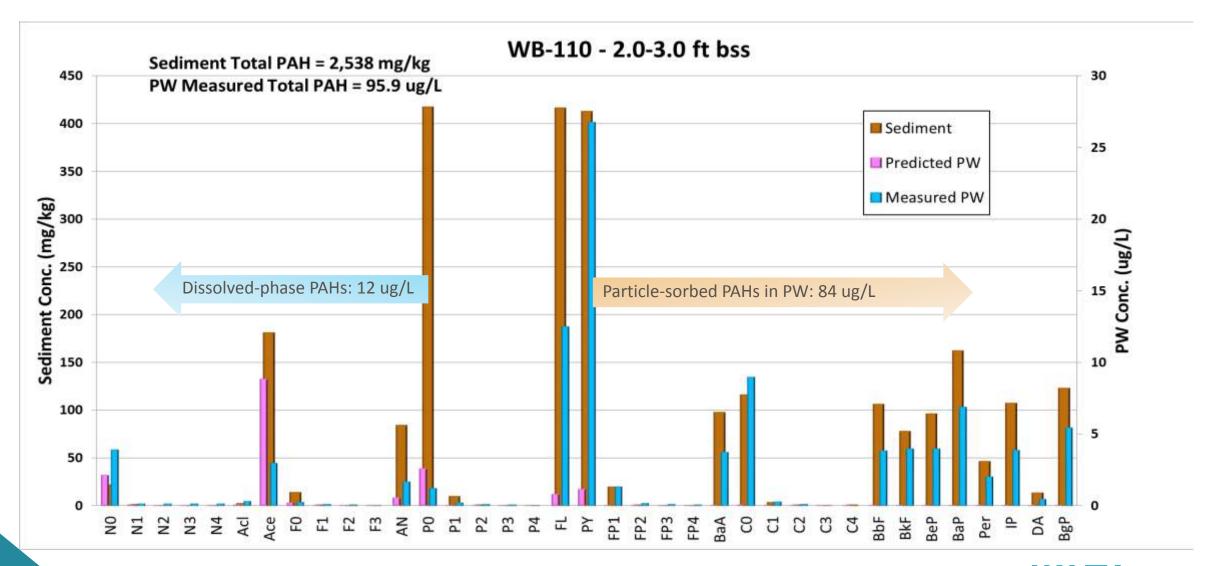


### 3) Plot PAHs Measured in "co-located" Pore Water





### 4) Compare PAH compositions in PW vs predicted PW





### Results/Lessons Learned

- Dissolved phase matters!
- Minimize the need to recognize & adjust for particle influence through careful PW sampling, processing, analysis
- Recognize particle-sorbed PAH influence in PW
- Conservative assumptions are ok, even GOOD....but at least recognize the extent of conservatism:
  - Potential cost savings (e.g., reactive vs isolation/sand capping), or
  - Credit for conservative margin of safety

