

DESIGN AND IMPLEMENTATION OF EISB SYSTEMS FOR CHLORINATED COMPOUND REMEDICATION

APRIL 28, 2021

LESSONS LEARNED

“We know a thing or two because we’ve seen a thing or two.”

– *Farmer’s Insurance Company*



PRESENTATION OUTLINE

When to Consider EISB

Bioremediation 101

Basic EISB Systems

The Design Process

Other Considerations

DNAPL Source Areas



Why Bioremediation ?

Usually highly cost-effective

Effective for most common GW contaminants

Applicable to source areas and plumes

Uses simple and safe substrates and nutrients

Minimal above-ground profile

Low O&M requirements and operational risk

Flexible and sustainable (low carbon footprint)

Compatible with natural attenuation

Why NOT Bioremediation?

Contaminants not biodegradable or inhibitory compounds present

Need to treat vadose zone soils

Natural conditions not supportive

Clean-up time is a major driver

Secondary GW quality is an issue

Other options are more cost-effective



MICROBIOLOGY 101

MICRO- BIOLOGY 101

Microorganisms (bacteria) are everywhere. There are $10^5 - 10^7$ bacteria in every gram of soil.

Like all organisms, bacteria need to eat and breathe. They eat electron donors and breathe electron acceptors.

Bacteria need low levels of other nutrients like nitrogen, phosphate, and trace minerals.

Bacteria also need water and shelter (mineral surfaces). Given these, they can survive in extreme environments (from -25° to 120°C and from $\text{pH} < 2$ to $\text{pH} > 12$).

MICROBIOLOGY 101

KEY DEFINITIONS

Aerobic

Environmental conditions where oxygen is present

Anaerobic

Environmental conditions where oxygen is absent

Biotic

Process mediated by bacteria

Abiotic

A purely physical or chemical process

MICROBIOLOGY 101

ELECTRON DONORS

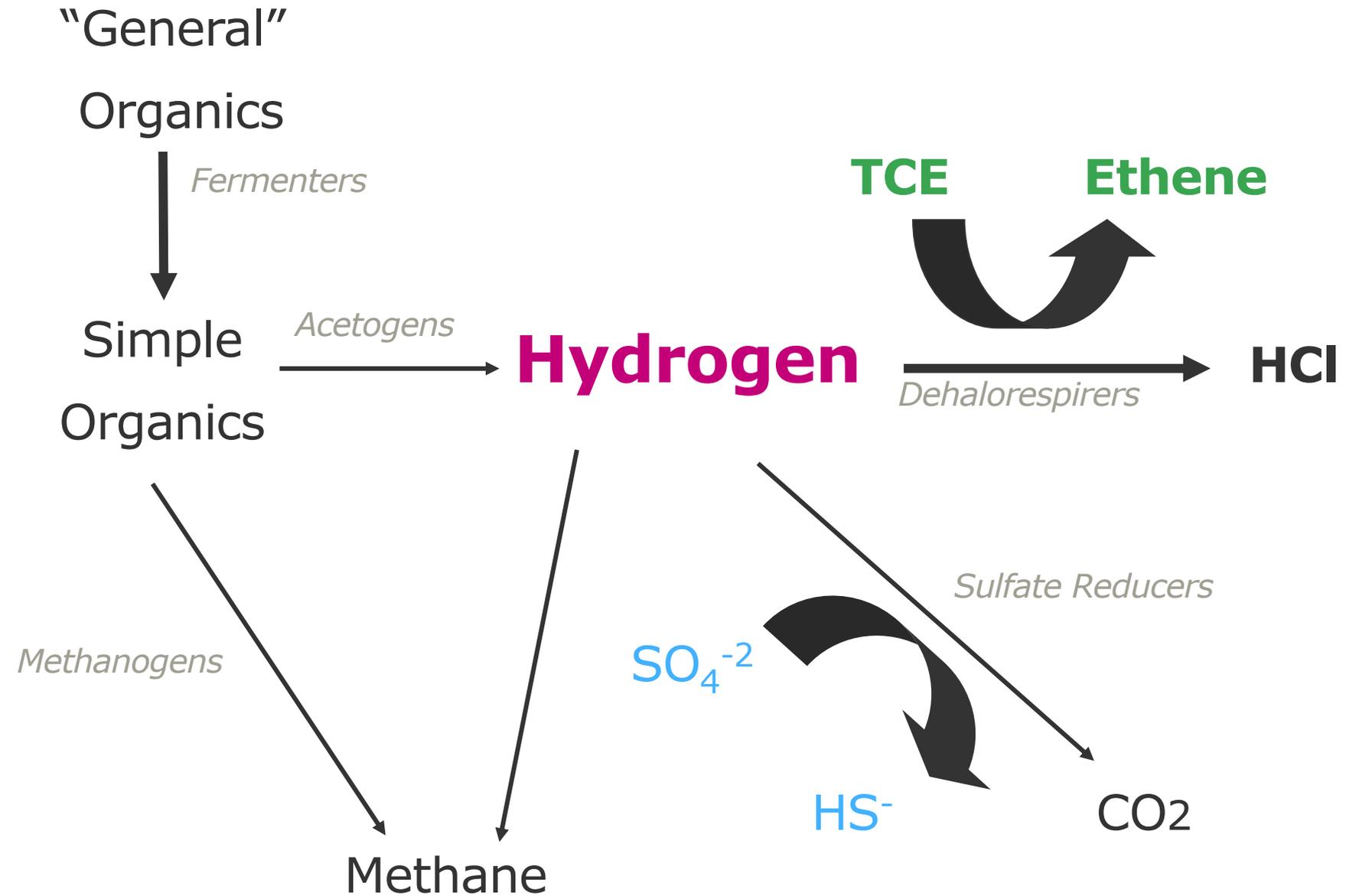
Electron Donor:
A compound
that loses
electrons during
biodegradation
(source of energy
for bacteria)

Hydrogen is the main electron donor in most anaerobic biodegradation processes.

Hydrogen is produced by the fermentation of complex organic compounds.

Examples include natural organic matter, small organic acids, fermentable substrates, waste solvents (i.e. methanol, acetone), landfill leachate, BTEX compounds

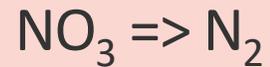
MICROBIAL
COMMUNITIES
INVOLVED IN
ANAEROBIC
REDUCTIVE
DECHLORI-
NATION



MICRO- BIOLOGY 101

CHLORINATED COMPOUNDS AS ELECTRON ACCEPTORS

Respiration



Redox Range

+0.8 to +0.2

+0.5 to +0.2

+0.3 to 0

-0.1 to -0.3

-0.2 to -0.4

-0.2 to -0.4

Bacteria

Aerobic

Nitrate reducing

Iron reducing

Sulfate reducing

Methanogenic

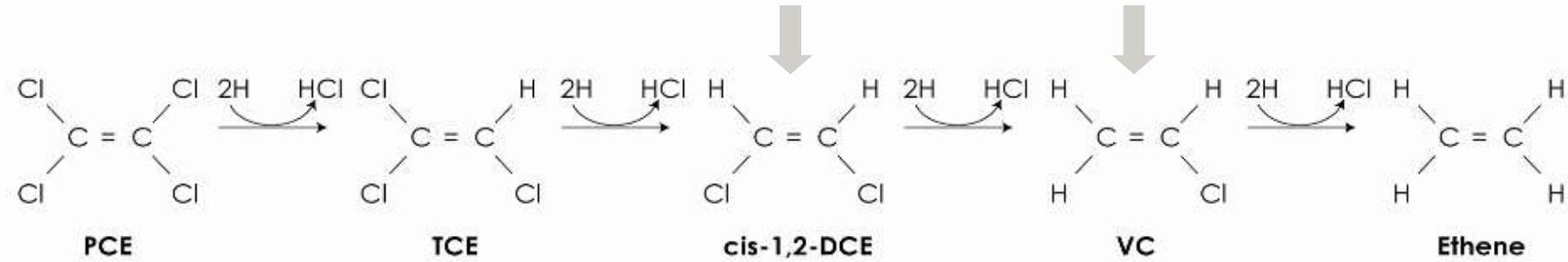
Acetogenic



MICROBIOLOGY 101

REDUCTIVE DECHLORINATION PATHWAY

Can accumulate if requisite bacteria
are absent



Dehalobacter
Dehalospirillum
Desulfitobacterium
Desulfuromonas
Dehalococcoides
Geobacter

Multiple strains within
a single group
(*Dehalococcoides*)



ESIB SYSTEMS

WHEN WE TRY
TO FIGHT
MOTHER
NATURE, WE
USUALLY LOSE...

Match the remedy to the natural environment

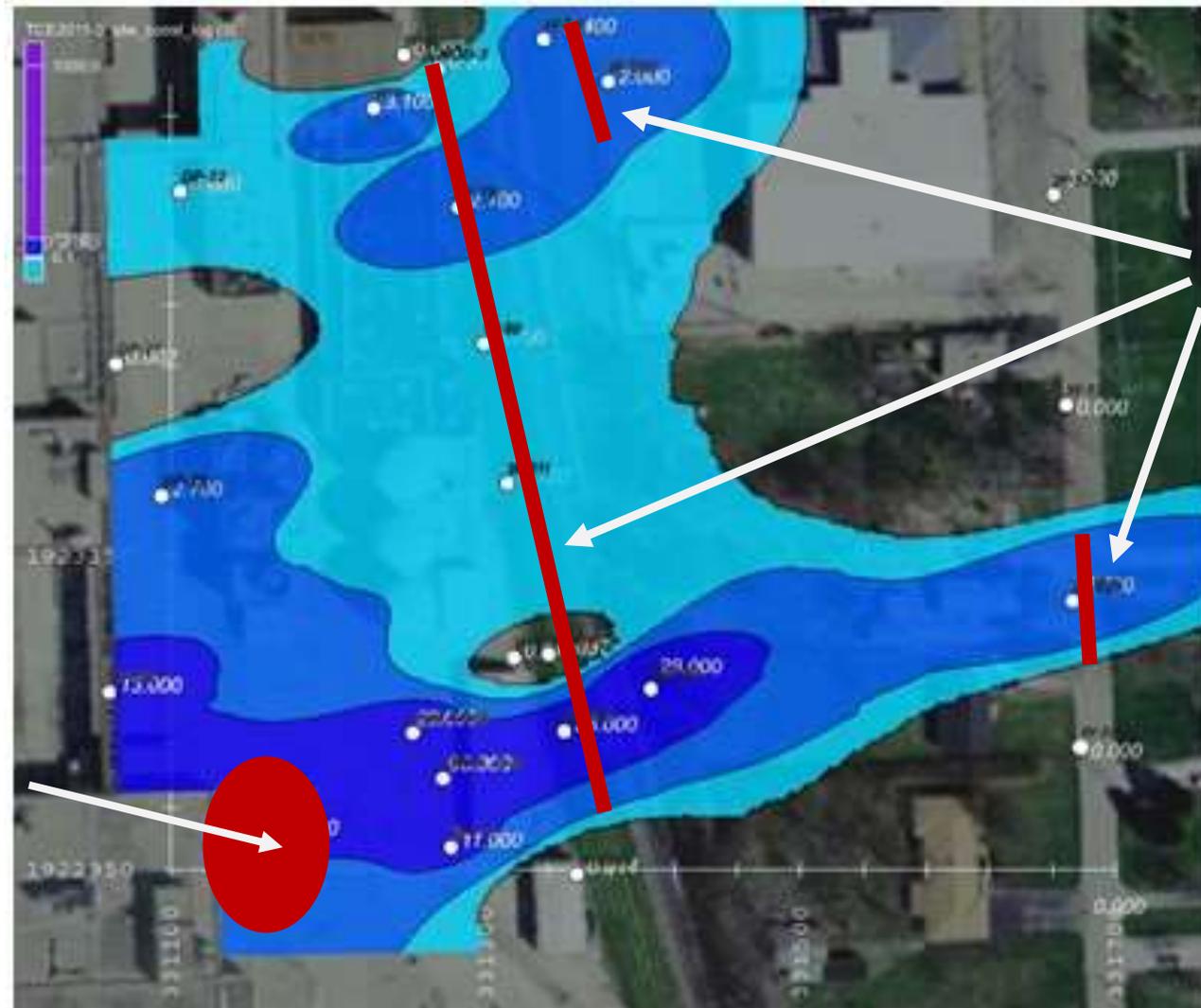
Anaerobic aquifers → EISB or ISCR
(not ISCO)

Aerobic aquifers → more options (but
avoid EISB in high-flow situations)

Low permeability soils → modify
injection method and use solid
amendments

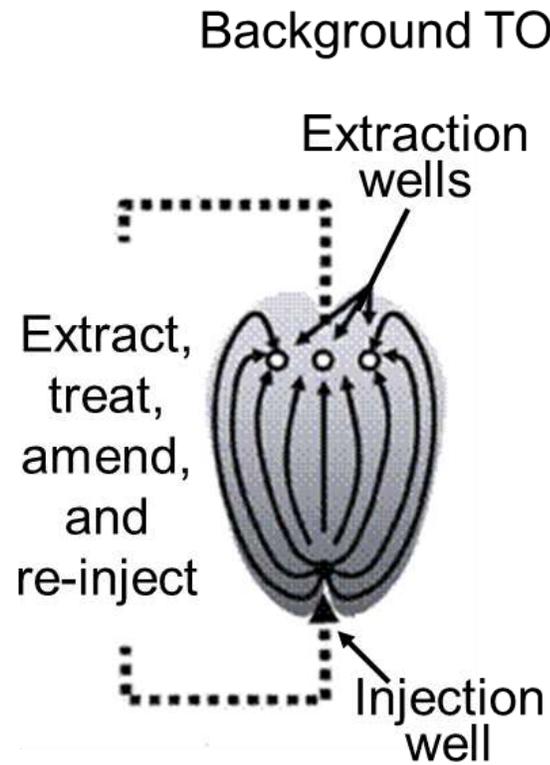
TREATMENT OPTIONS

Source
treatment

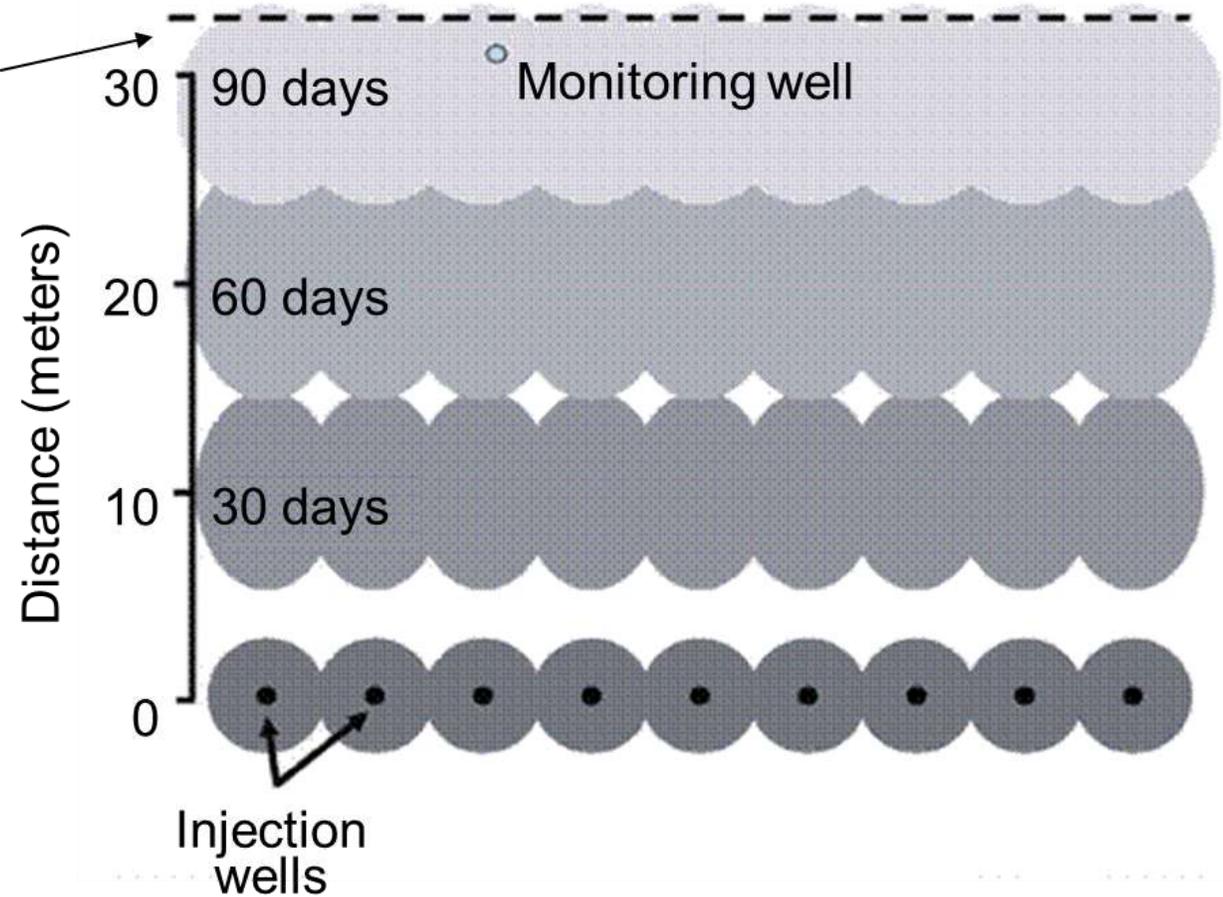


Biobarriers

TYPES OF EISB SYSTEMS



Active or
Semi-Passive



Passive

ITRC Technology Overview: In Situ Bioremediation of Chlorinated Ethene DNAPL Source Zones (BIODNAPL-1, 2005)

COMMON DELIVERY METHODS

Operation	Delivery Method	Donors	Amendment Frequency
Passive	<ul style="list-style-type: none">Discrete Injection Points (injection points, temporary or permanent injection wells)Good for reactive barriers	Solid or slow-release (e.g., EVO)	Annually to multi-year
Semi-Passive	Batch addition with intermittent recirculation	<ul style="list-style-type: none">SolubleSlow-release	Monthly to annually
Active	<ul style="list-style-type: none">Continuous recirculationCapture and recharge	Soluble	Daily to weekly

COMMON DELIVERY METHODS

	Passive	Semi-Passive	Active
# Well Required (Cap \$)	High	Med	Low
Infrastructure (Cap \$)	Low	Med	High
Operation (O&M \$)	Low	Med	High
Fouling of Wells (O&M \$)	Low	Med	High
Distribution in GW	Least	Better	Best
Control of Dose	Least	Better	Best
Maintains Water Quality	Least	Better	Best

ELECTRON DONOR CHOICES



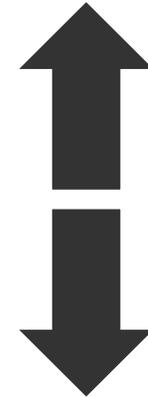
Soluble

- Lactate/other organic acids
- Methanol/ethanol
- Molasses/other carbohydrates
- Dairy whey



Slow Release

- Edible oils and oil mixtures
- Lactate polymers
- Chitin (glucosamine polymer)
- Plant matter products



Delivery methods change as a function of amendment type

Key Point: Amendment choice and injection design are closely linked.

INJECTION MATRIX

Low pressure fluid injection is not appropriate for low permeability soils

Table 4-4. Site-Specific Impacts on Reagent Distribution Technique

Parameter	Vertical Injection Wells	Vertical Recirculation Wells	Horizontal Wells	Direct-push Technology Injection	Hydraulic Fracture	Pneumatic Fracture
<u>Amenability to Media Type</u>						
Unconsolidated media	Excellent	Excellent	Excellent	Excellent	Excellent	Excellent
Consolidated media	Excellent	Good	Excellent	Not recommended	Excellent	Excellent
<u>Fracture Continuity</u>						
Good fracture continuity	Good	Good	Fair	Not recommended	Good	Good
Poor fracture continuity	Fair	Poor	Poor	Not recommended	Good	Good
<u>Hydraulic Conductivity</u>						
$>10^{-3}$ cm/sec	Excellent	Excellent	Excellent	Excellent	Poor	Poor
$<10^{-3}$ but $>10^{-4}$ cm/sec	Good	Fair	Fair	Excellent	Fair	Fair
$<10^{-4}$ but $>10^{-5}$ cm/sec	Fair	Poor	Poor	Good	Good	Good
$<10^{-5}$ but $>10^{-6}$ cm/sec	Poor	Not recommended	Not recommended	Fair	Excellent	Excellent
$<10^{-6}$ cm/sec	Not recommended	Not recommended	Not recommended	Not recommended	Excellent	Excellent
<u>Lithology</u>						
Homogeneous ($K_{max}/K_{min} < 1,000$)	Excellent	Excellent	Excellent	Excellent	Excellent	Fair
Heterogeneous ($K_{max}/K_{min} > 1,000$)	Fair	Fair	poor	Good	Fair	Fair

Source: Best Practices for Injection and Distribution of Amendments, Technical Report TR-NAVFAC-EXWC-EX-1303, March 2013

Donor Emplacement by Direct Injection





ESIB DESIGN PROCESS



What is a treatability test?

- Laboratory based “bench-scale” testing
- Uses site soil, sediment or rock and groundwater, typically in batch bottles
- Used to assess biodegradation potential under site-specific conditions
- Usually 4-12 months long
- Column studies can also be performed, but are much less common

Why do Treatability Studies?



Relatively low cost



Test multiple variables at the same time – narrows potential options prior to going to the field



Identify potential complications and address them before they cause problems in the field



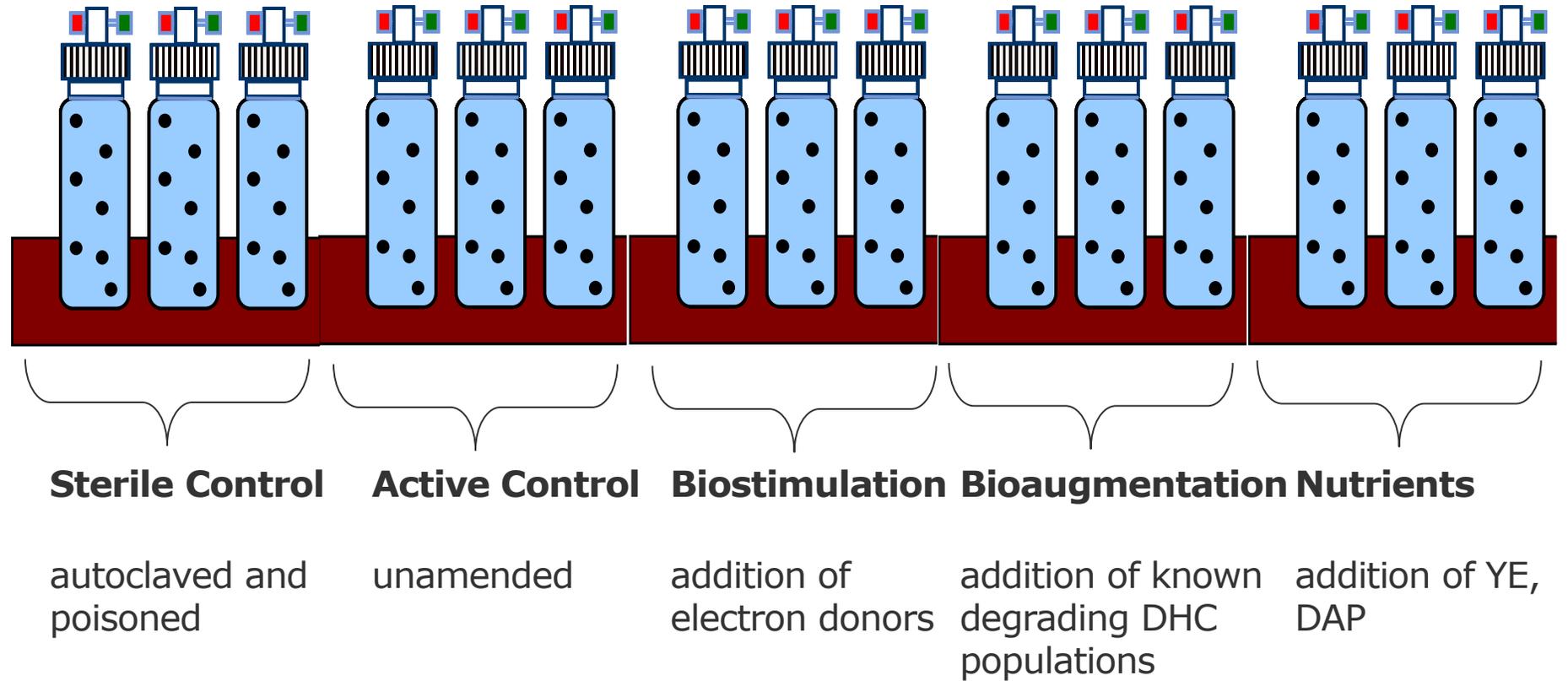
Obtain regulator or client buy-in prior to investing in field-scale tests

Typical Cost

Lab treatability study - \$20-30 K

Field pilot test - \$100-300 K

BIOTREAT- ABILITY STUDY DESIGN





Importance of Sample Collection

- The sample is “alive”
- Collect using core tube
- Minimize field disturbance
- Cap and seal ends, store on ice
- Ship to lab quickly
- Lab should transfer soil to glass container and store under anaerobic conditions
- Set up study quickly
- Understand that soil has a “shelf life”



PILOT TESTING

Why Perform a Pilot Test?



Reduce scale up uncertainty and better estimate full-scale project cost



Verify amendment distribution

- Injection well/injection point spacing
- Fluid injection rates
- Amendment transport
- Potential for surfacing



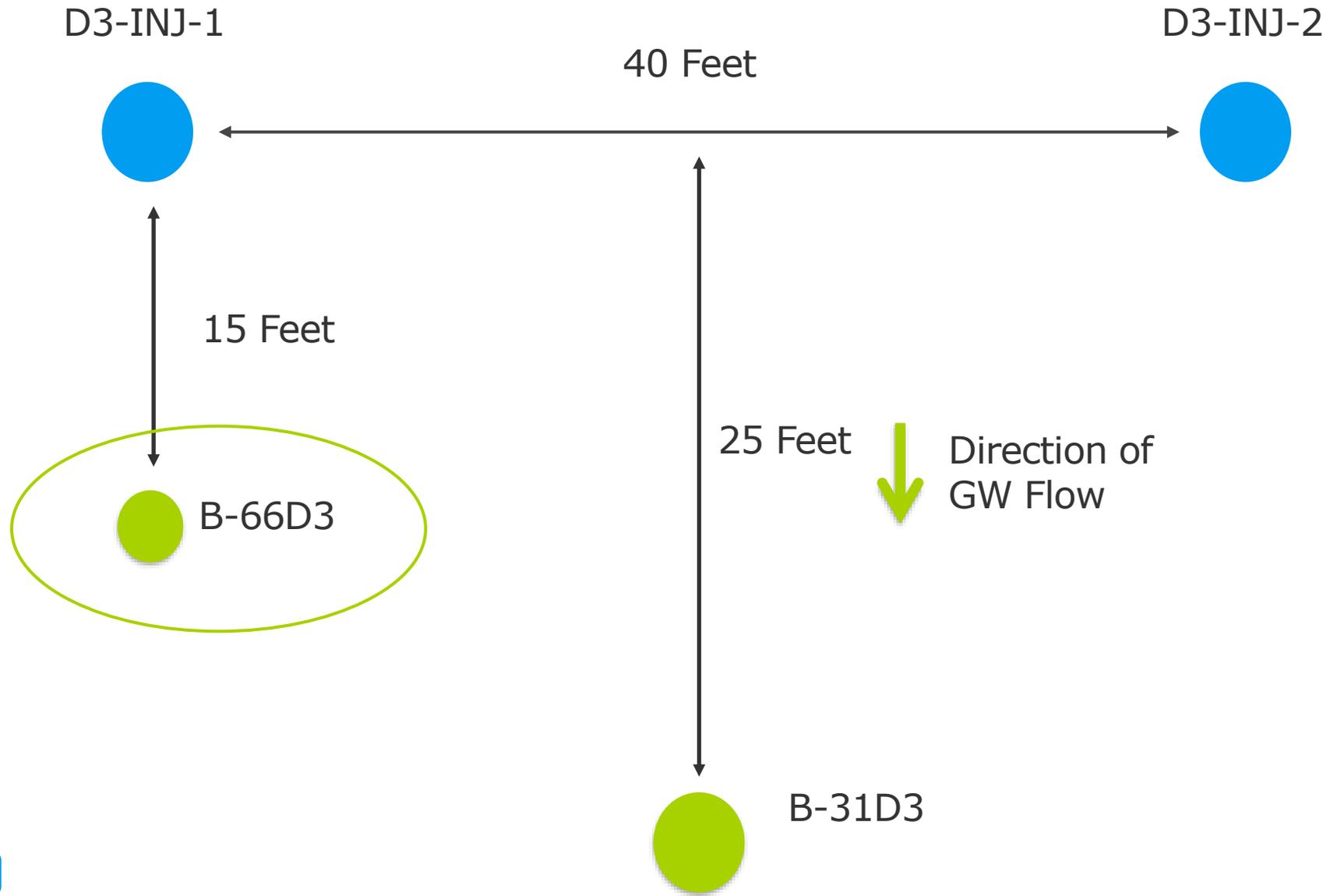
Identify implementation problems (biofouling, aquifer plugging)



Obtain regulator and/or client buy-in

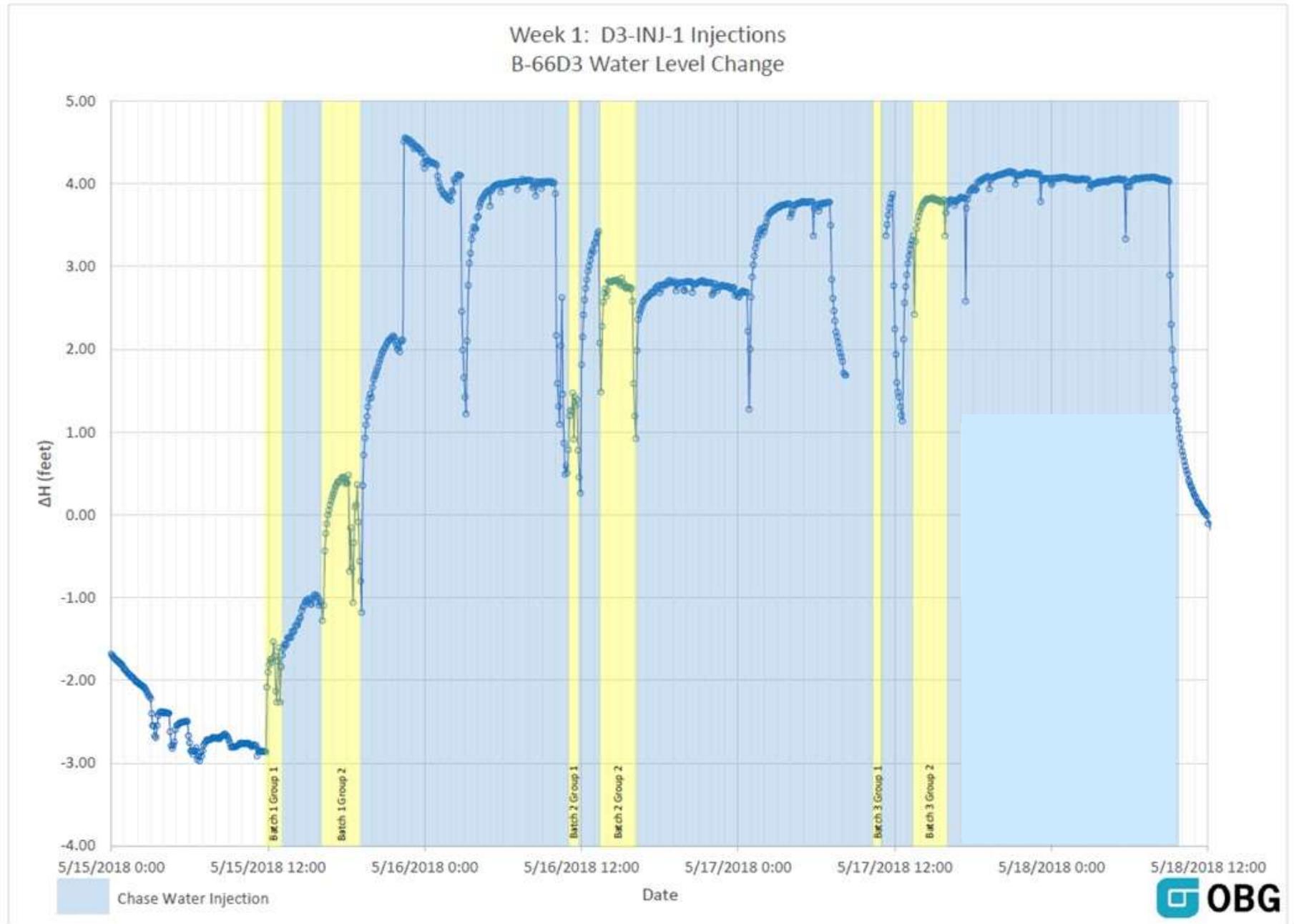
Important to measure what we put into the ground

D3 PILOT TEST – PHASE 1 LAYOUT



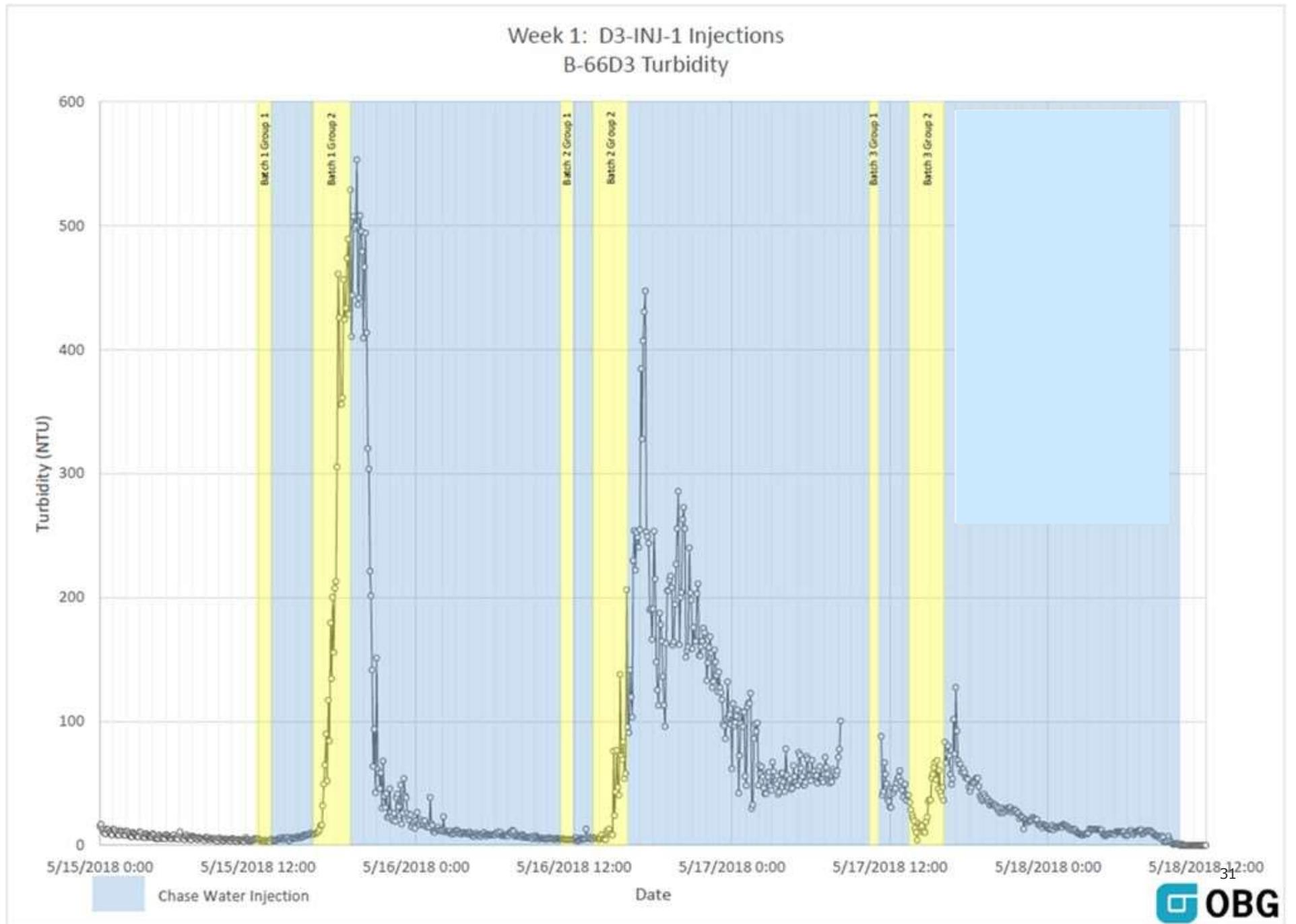
PRESSURE TRANSDUCER DATA

Phase 1 Pilot -
Week 1



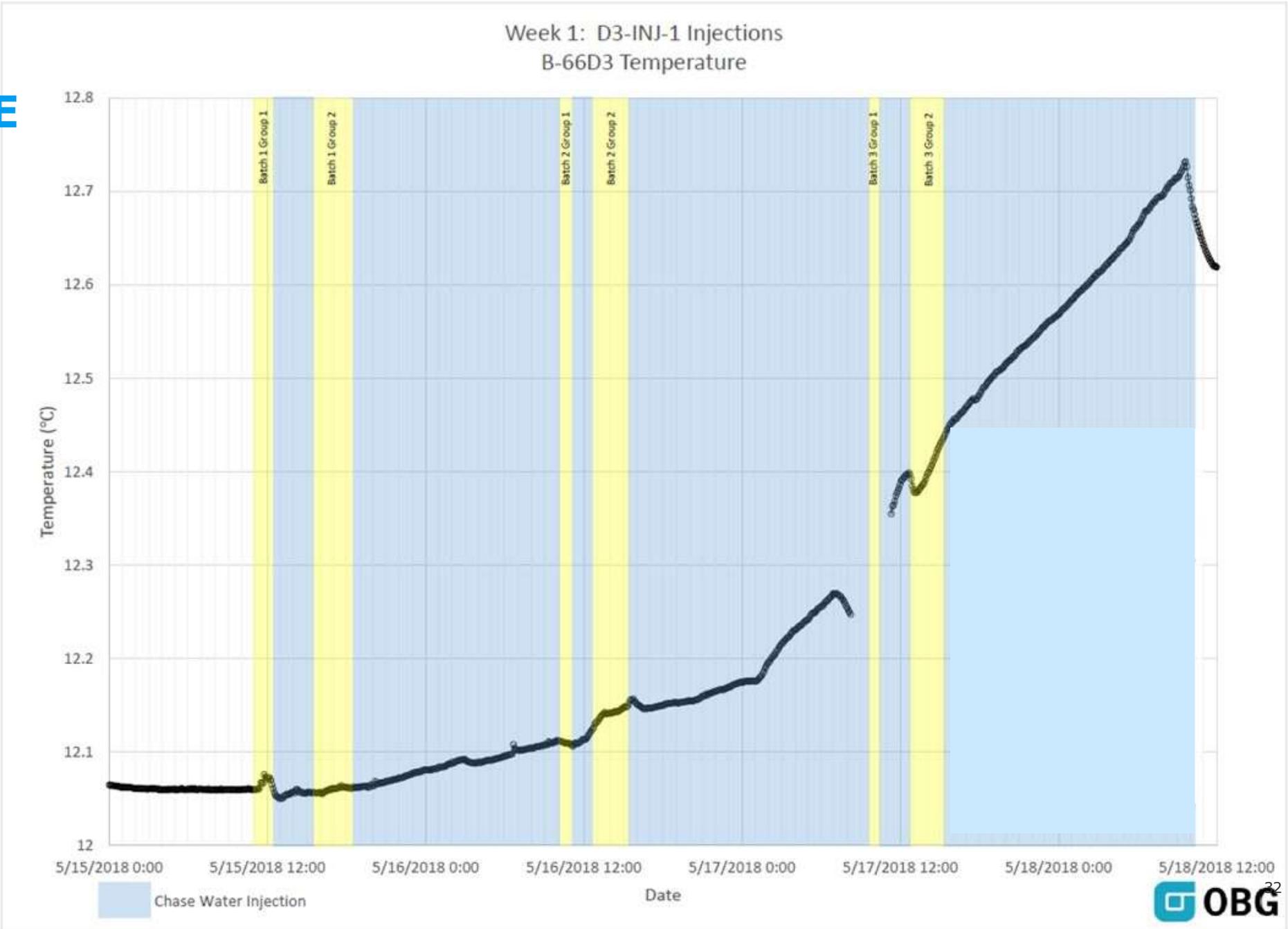
PROBE TURBIDITY DATA

Phase 1 Pilot -
Week 1

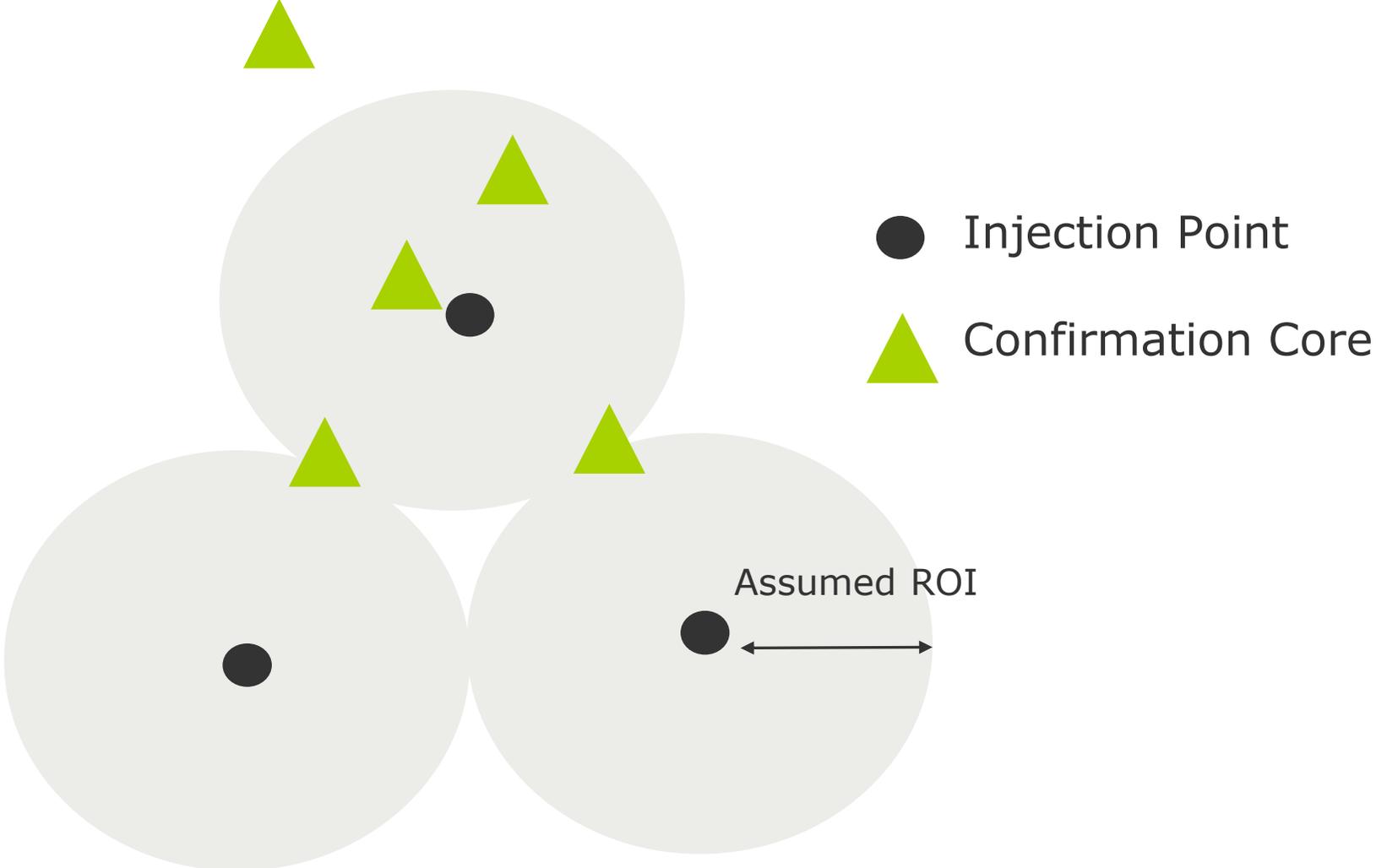


PROBE TEMPERATURE DATA

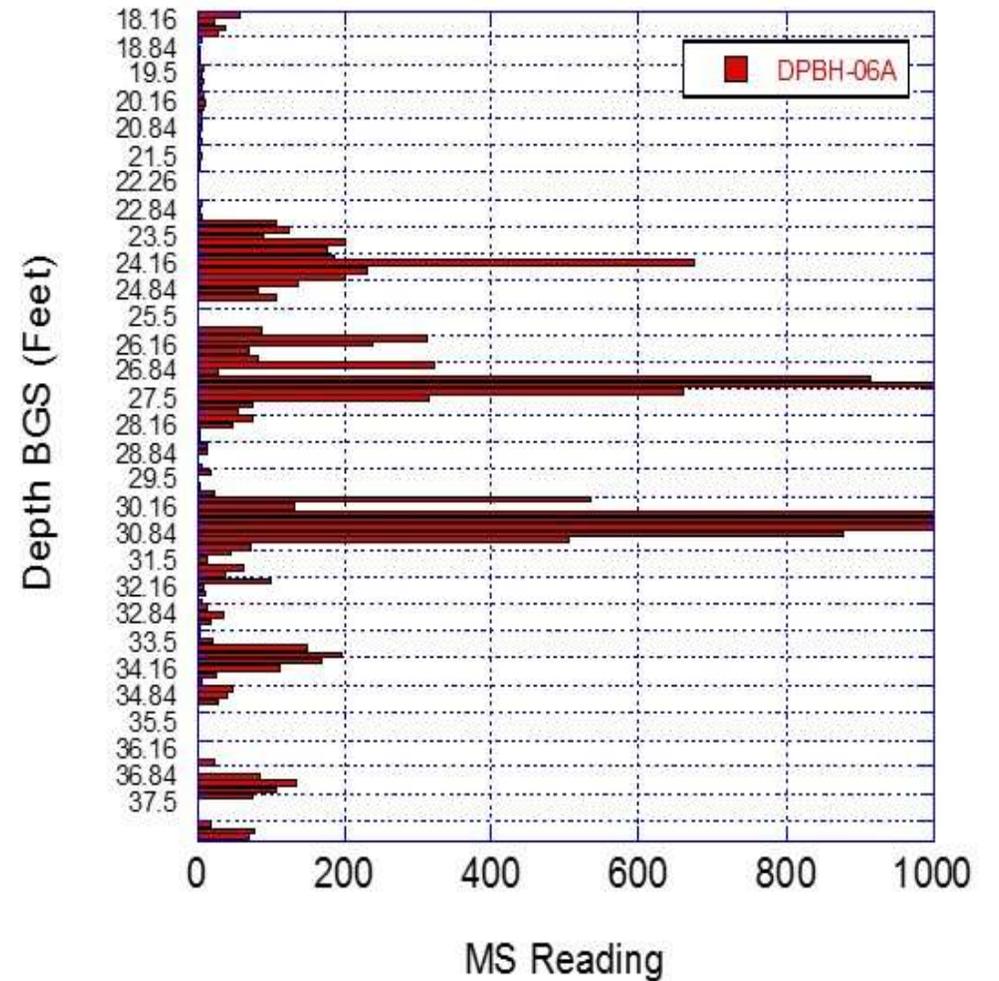
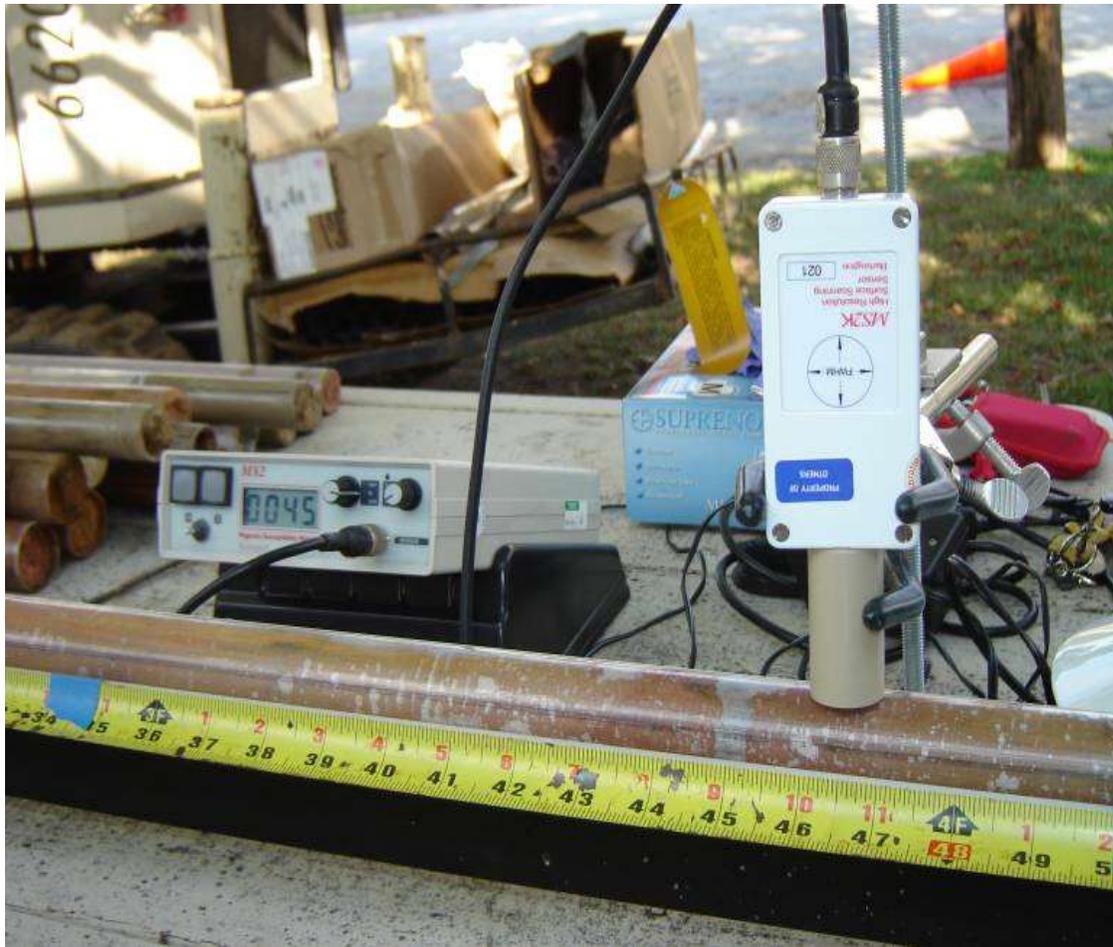
Phase 1 Pilot - Week 1



ROI DETERMINATION FOR DONOR INJECTION PROGRAMS



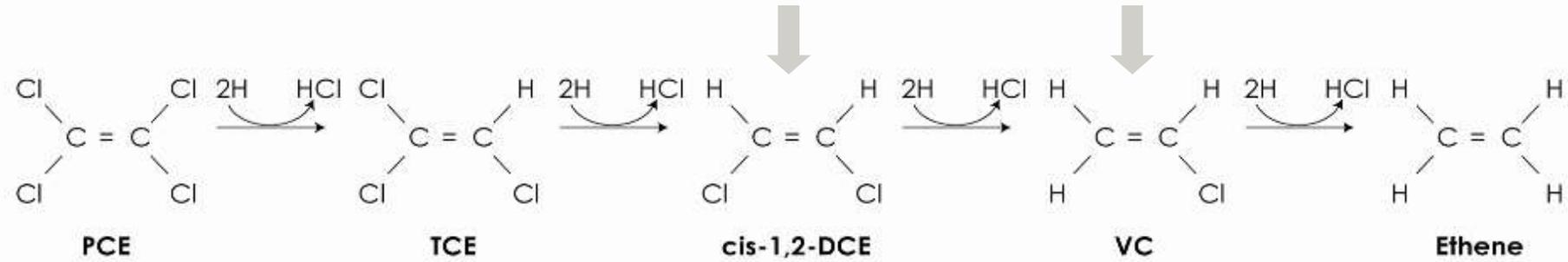
MAGNETIC SUSCEPTIBILITY MEASUREMENT





OTHER COMMON OPTIONS

Can accumulate if requisite bacteria are absent



REDUCTIVE DECHLORINATION PATHWAY

Dehalobacter
Dehalospirillum
Desulfitobacterium
Desulfuromonas
Dehalococcoides
Geobacter

Multiple strains within
a single group
(*Dehalococcoides*)

Bioaugmentation Culture Application

Dhc requires anaerobic conditions and neutral pH

Bioaugmentation culture should be added after the amendment and when ORP measurements indicate the subsurface is anaerobic

Some advocate short-cutting the process by injecting bioaugmentation culture with the amendments (*not recommended*)

Chase water should be used to “push” bacteria out into the formation

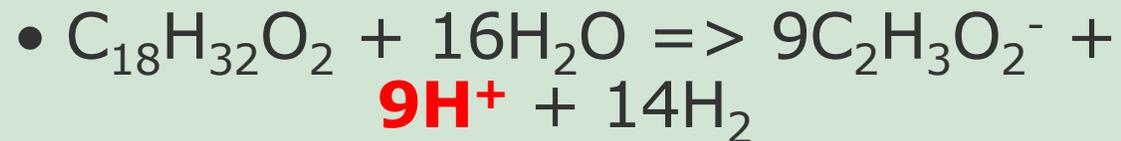
Chase water should be anaerobic and neutral pH

Both biological and chemical amendments are available to create anaerobic chase water

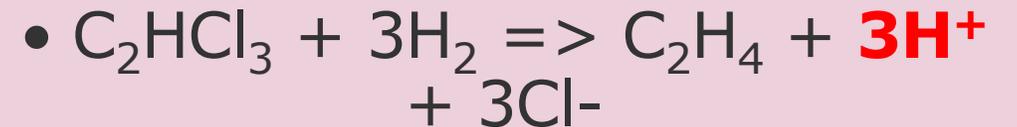
PH EFFECTS IN EISB

- EISB processes produce acid as the result of donor fermentation and reductive dichlorination.

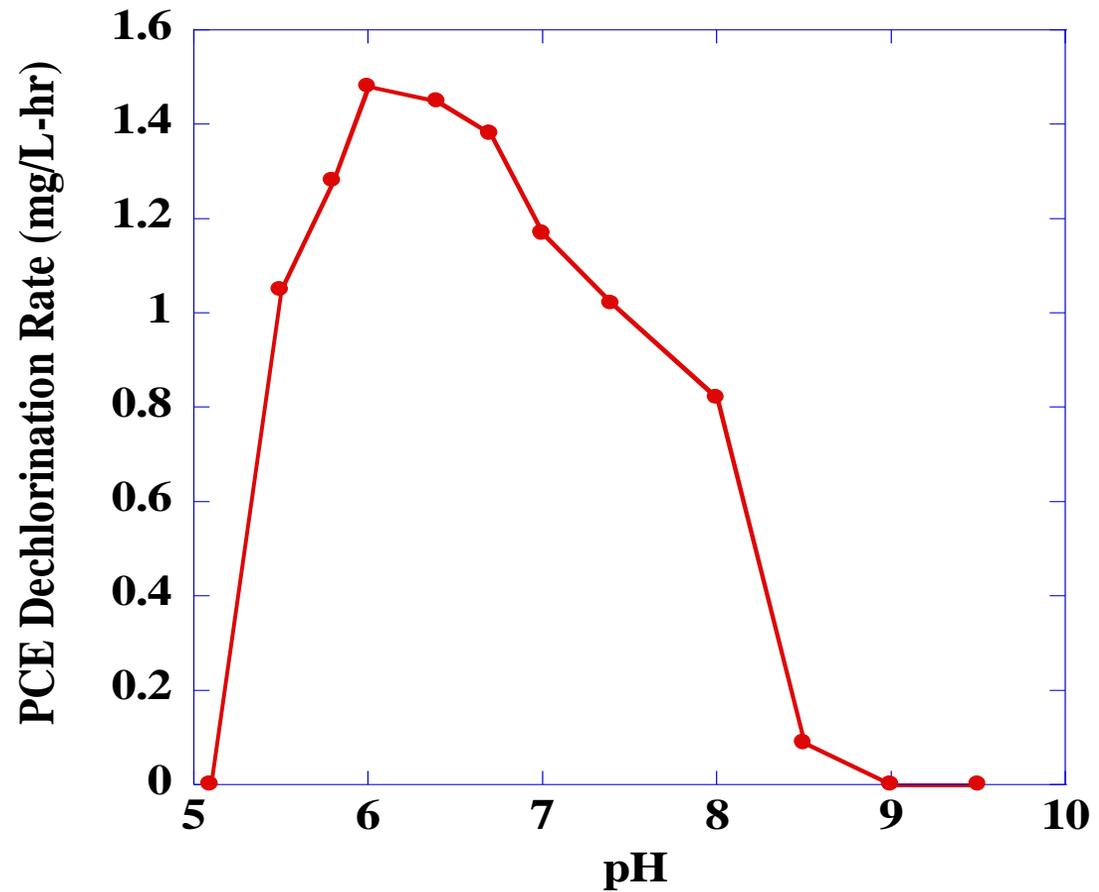
Fermentation of EVO



Dechlorination of TCE to Ethene



IMPACT OF PH ON DECHLORINATION BY DHC

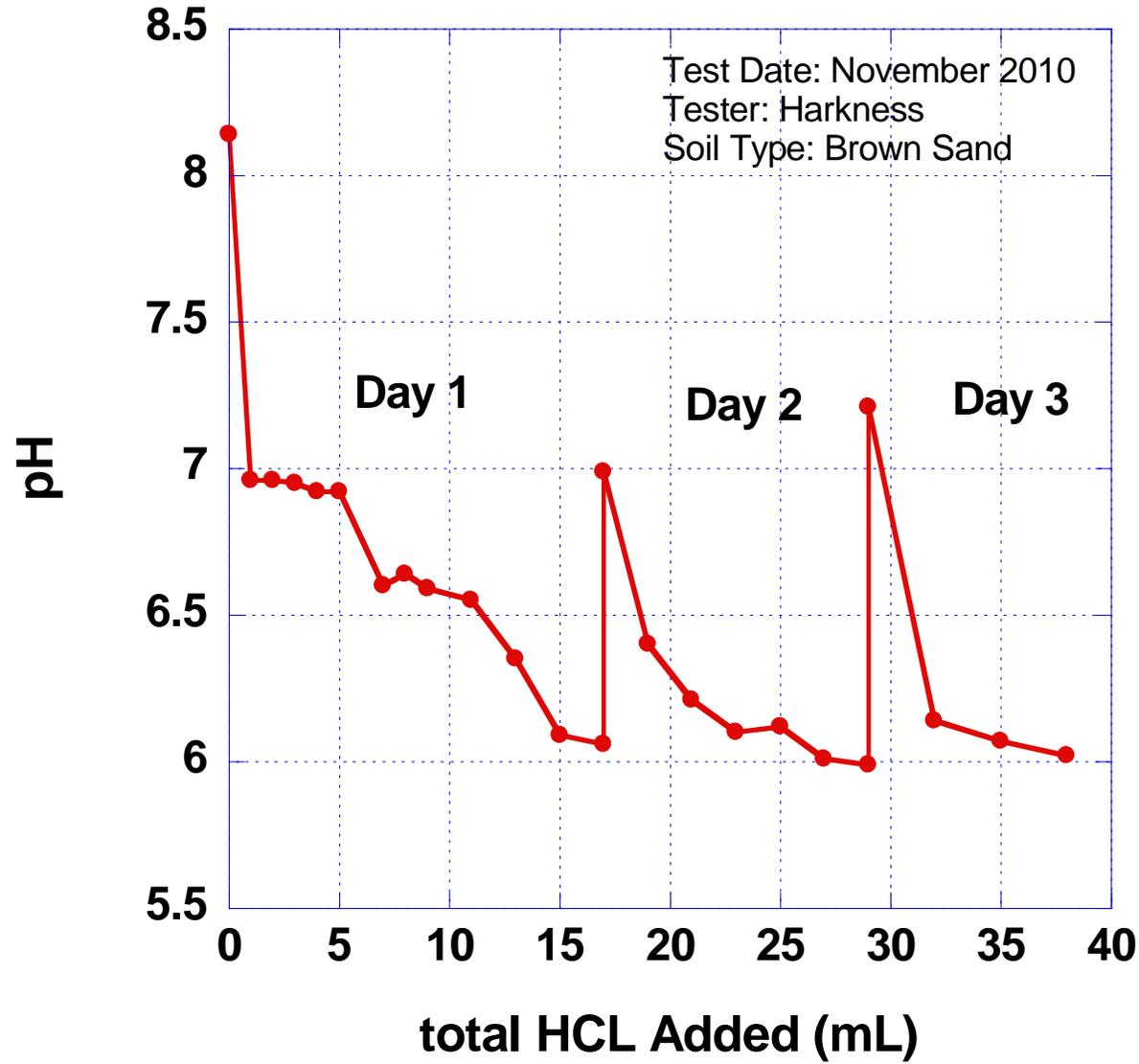


(Shaw Environmental)

Dhc have an optimum pH range of 6 - 8. Groundwater pH outside of this range will slow or inhibit reductive dechlorination activity.

TITRATION TEST RESULTS

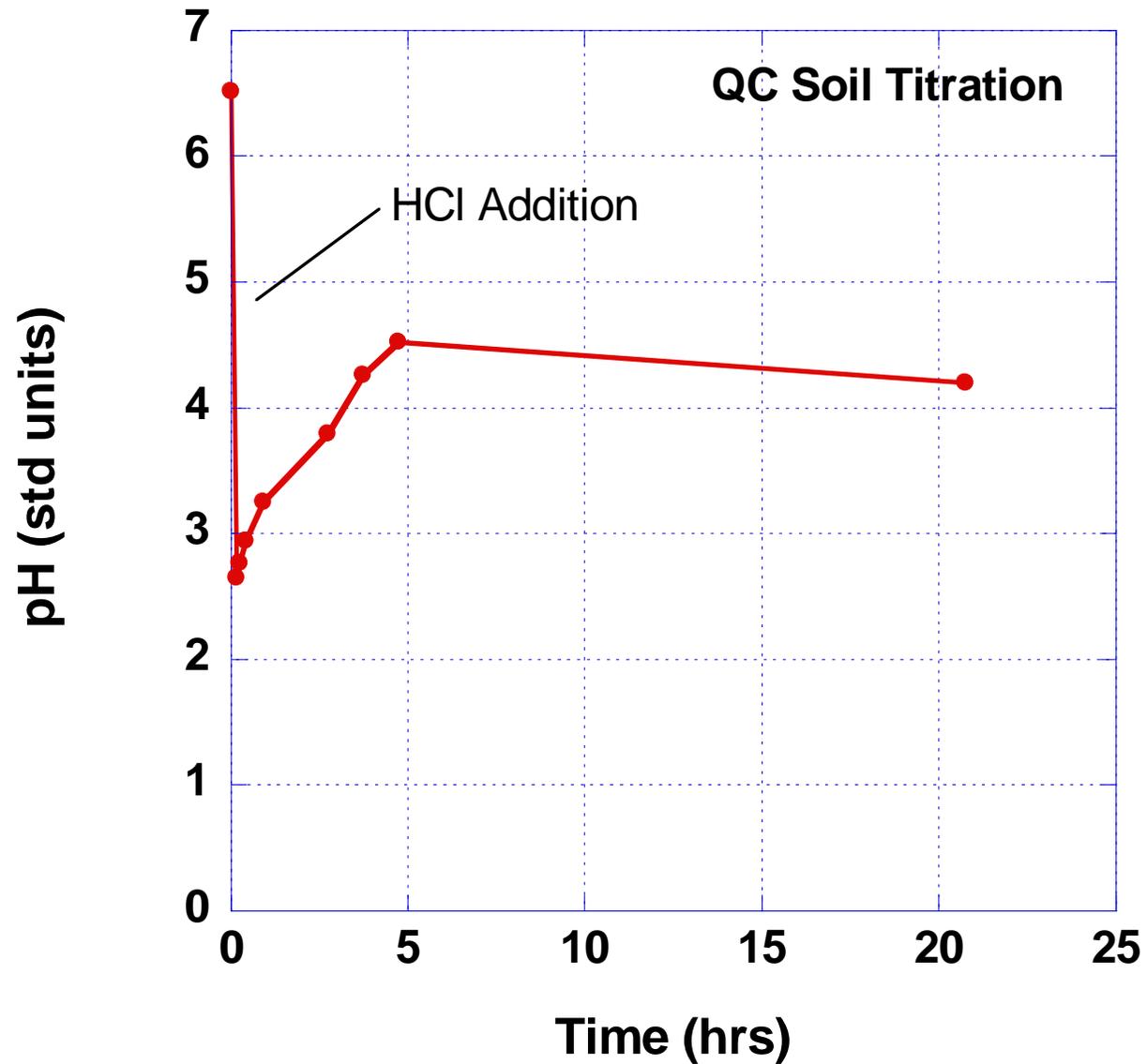
Indianapolis Soil Titration Results



Response of well buffered soil to 0.725 N HCl addition

Note – periodic rebound reflects rate limited dissolution of calcite from soil

TITRATION TEST RESULTS



Response of
poorly buffered
soil to addition
of 1 mL of
0.725 N HCl

BUFFERING OPTIONS IN EISB SYSTEMS



Need to add base to counteract the impact of acid generation due to donor fermentation and reductive dichlorination.



Addition of strong bases like sodium hydroxide can raise the pH of the system too much (e.g., produce a pH of 10 or higher), which can kill the bacteria.



Addition of weak bases are preferred since the pH rise is limited to 8.5.

BUFFERING OPTIONS IN EISB SYSTEMS



Sodium or potassium bicarbonate are effective buffers. These are soluble compounds that may require periodic addition.



CAUTION: Extensive use of bicarbonate buffers will alter groundwater geo-chemistry and may result in precipitation of insoluble residuals (e.g., calcium sulfate), with potential for aquifer plugging.



Commercial forms of slower release buffers are also available (e.g., calcium carbonate, magnesium hydroxide). These are typically combined with dispersants to allow them to travel in the aquifer.



APPLICATION OF EISB TO SOURCE AREAS

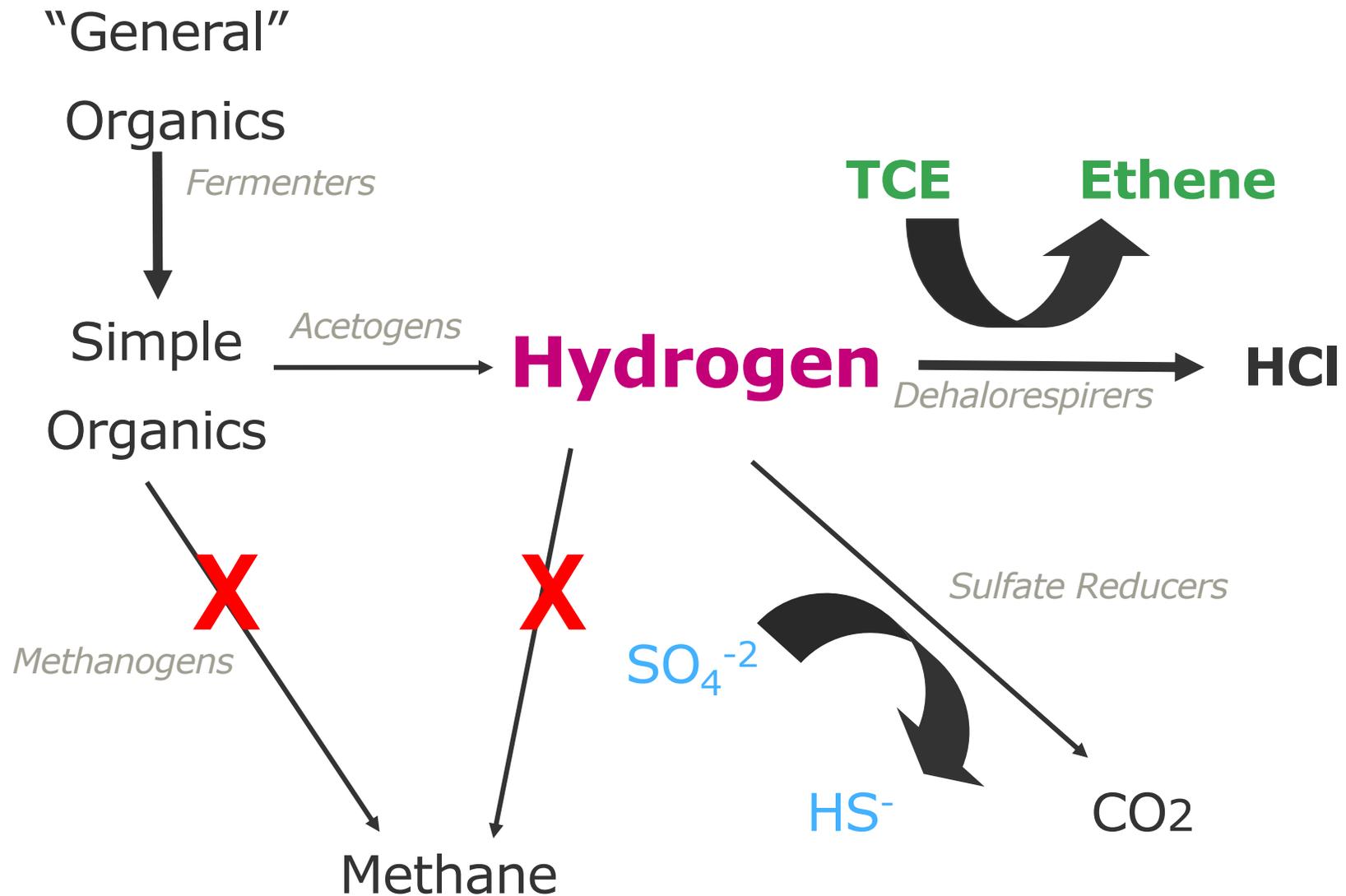
APPLICATION OF EISB TO SOURCE AREAS

EISB initially was applied to almost exclusively to plumes due to concern over microbial inhibition at high source area contaminant concentrations.

We now know that dechlorinating bacteria are active at near saturation concentrations for PCE and TCE.

In fact, there are several benefits to treating source areas using EISB

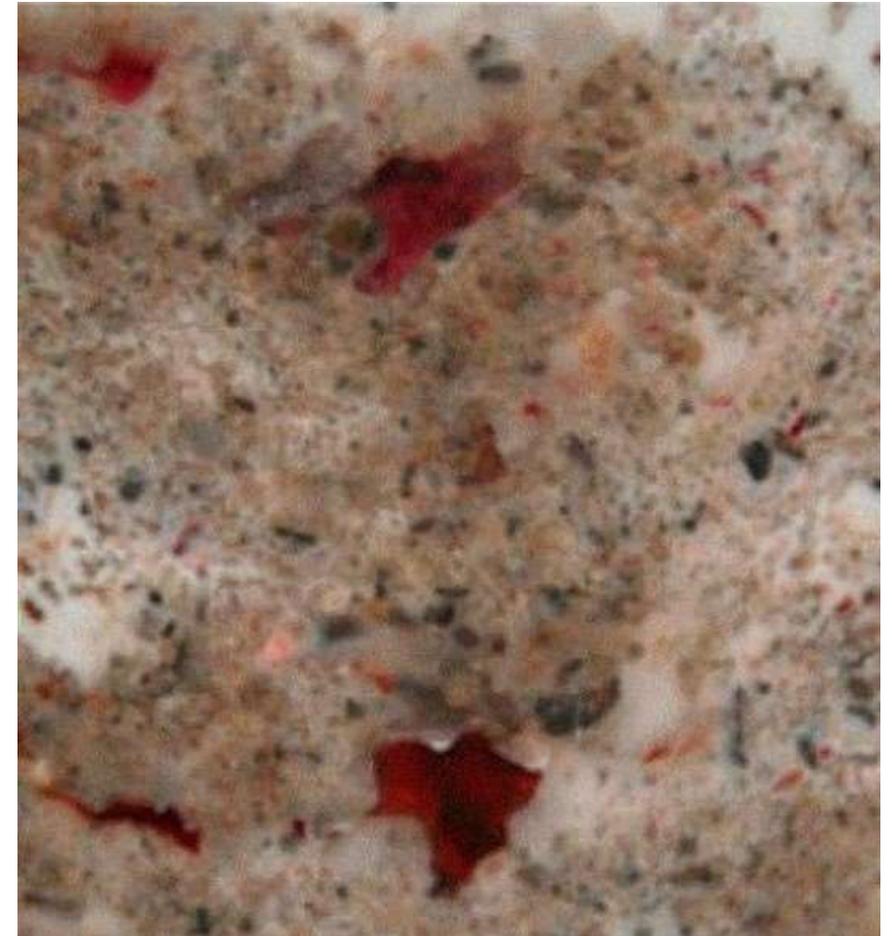
BENEFITS OF
SOURCE AREA
BIOLOGICAL
TREATMENT:
METHANOGEN
INHIBITION



BENEFITS OF
SOURCE AREA
BIOLOGICAL
TREATMENT:
PARTITIONING
DONOR
BEHAVIOR
(EVO)

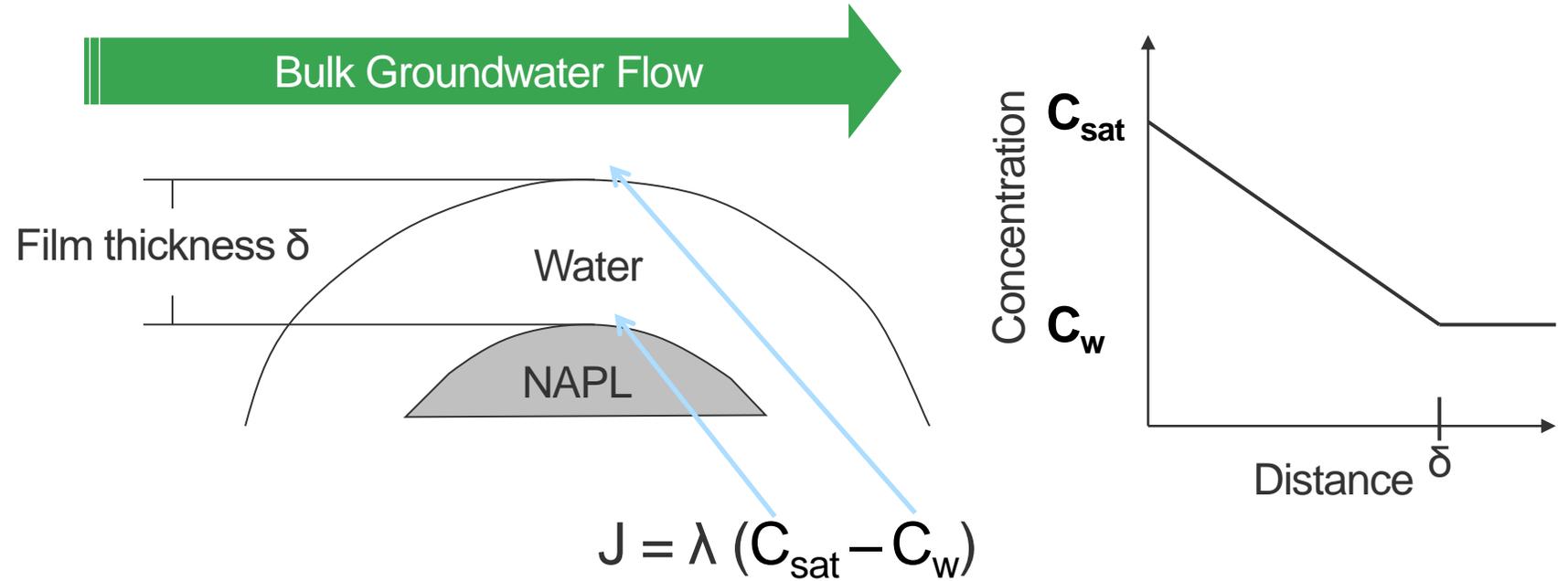


Three days



Six days

DNAPL DISSOLUTION AND MASS REMOVAL



$$\lambda = f \text{ (surface area, velocity)}$$

J = flux

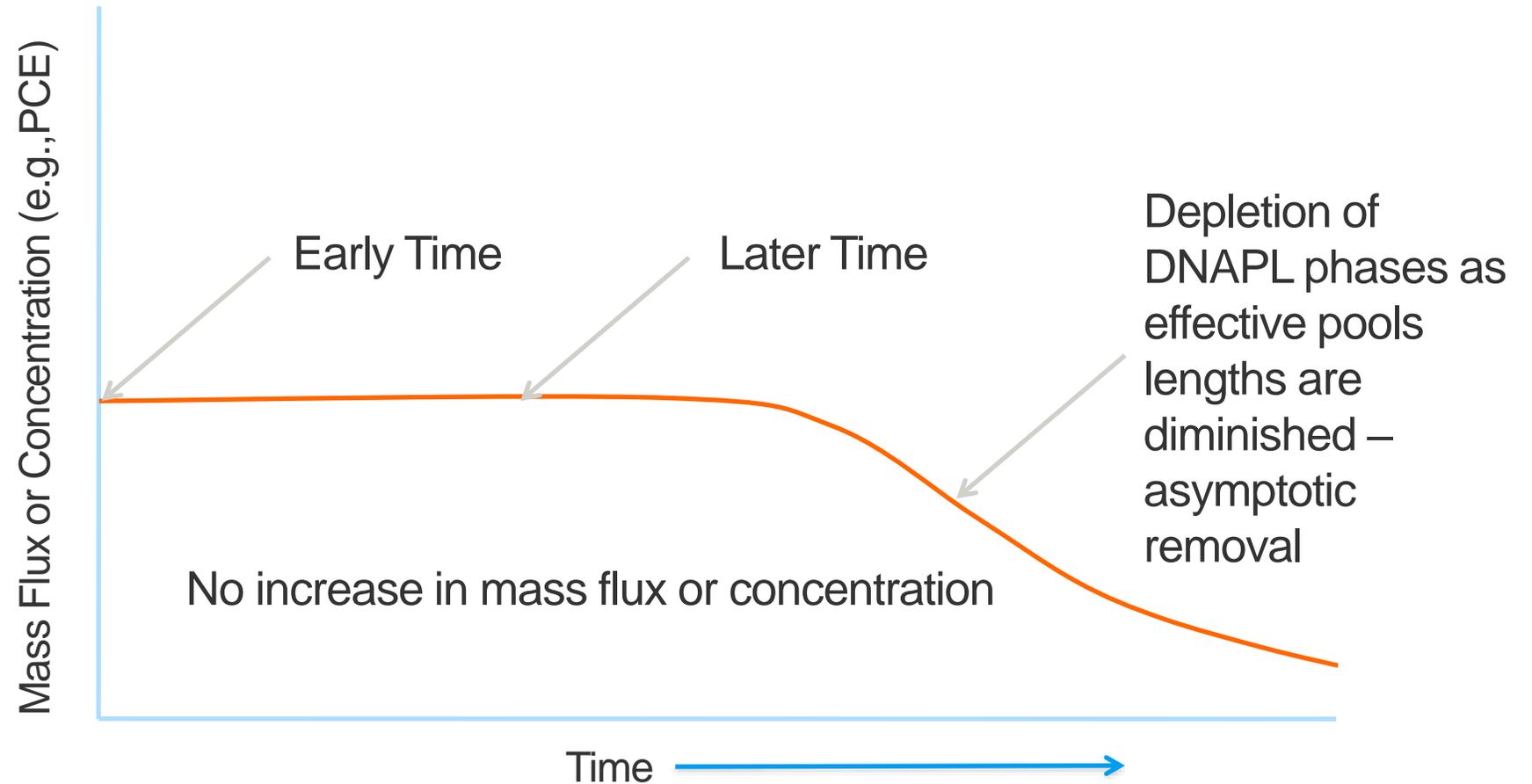
λ = mass transfer rate coefficient

C_{sat} = saturated concentration at the DNAPL/water Interface

C_w = bulk water concentration

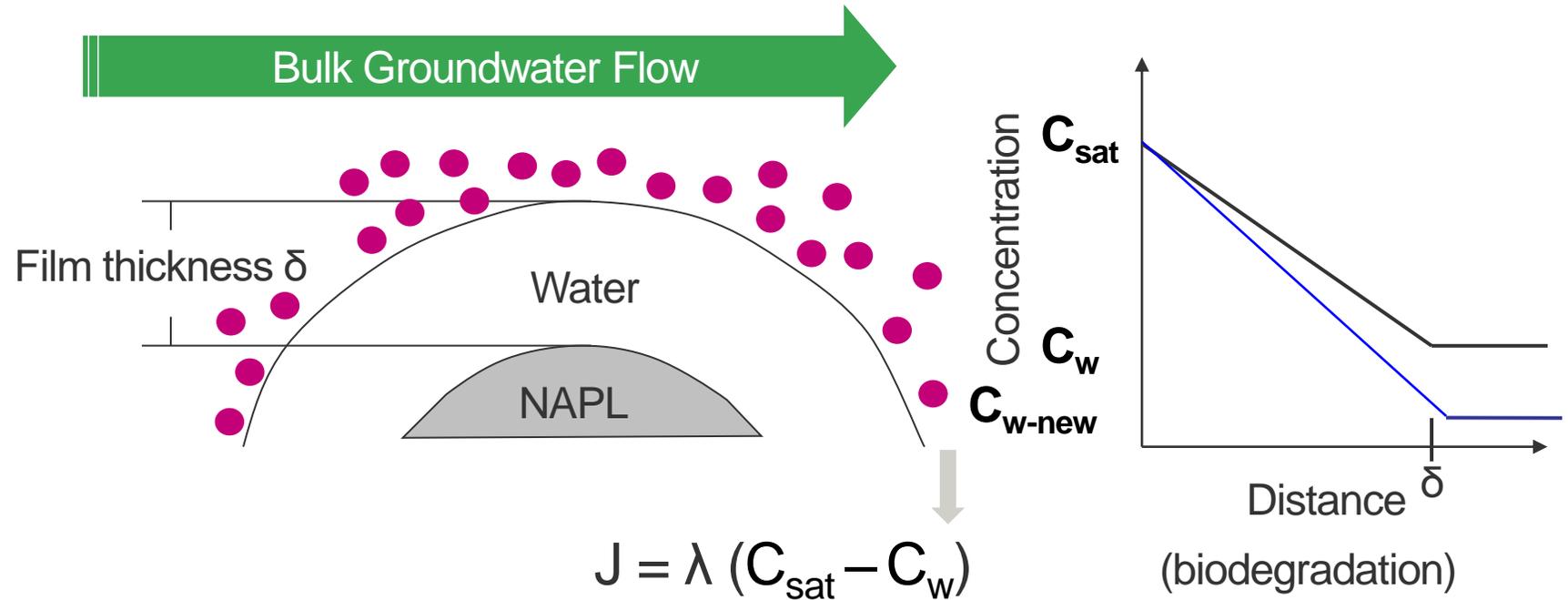
(ITRC training)

MASS REMOVAL OVER TIME WITHOUT EISB



(ITRC Modeling Study)

DNAPL DISSOLUTION AND MASS REMOVAL



$$\lambda = f \text{ (surface area, velocity)}$$

J = flux

λ = mass transfer rate coefficient

C_{sat} = saturated concentration at the DNAPL/water Interface

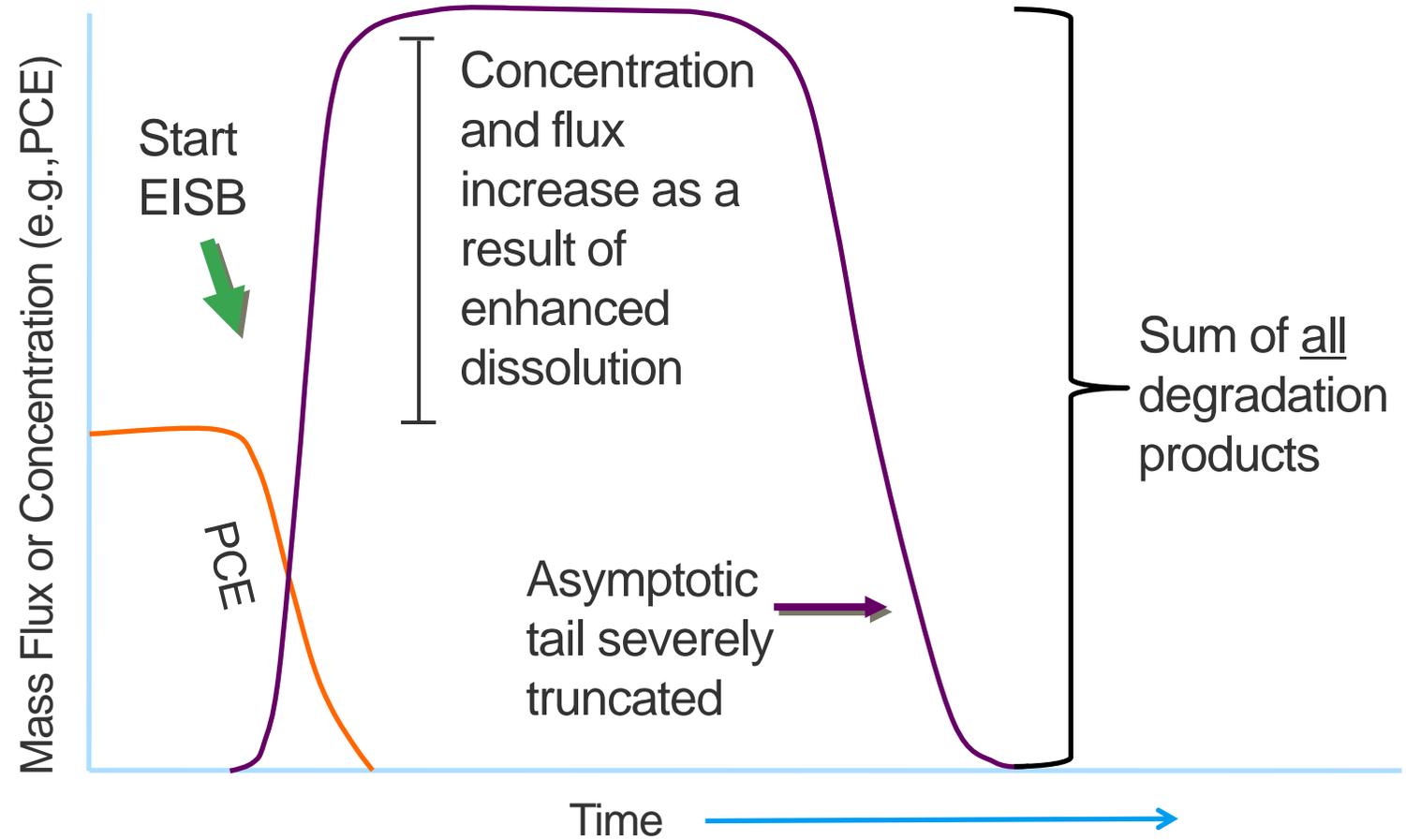
C_w = bulk water concentration

(ITRC training)

WATER SOLUBILITY

Compound	mg/L	mM
PCE	150	0.9
TCE	1,100	8.4
cDCE	3,500	36
VC	2,700	43

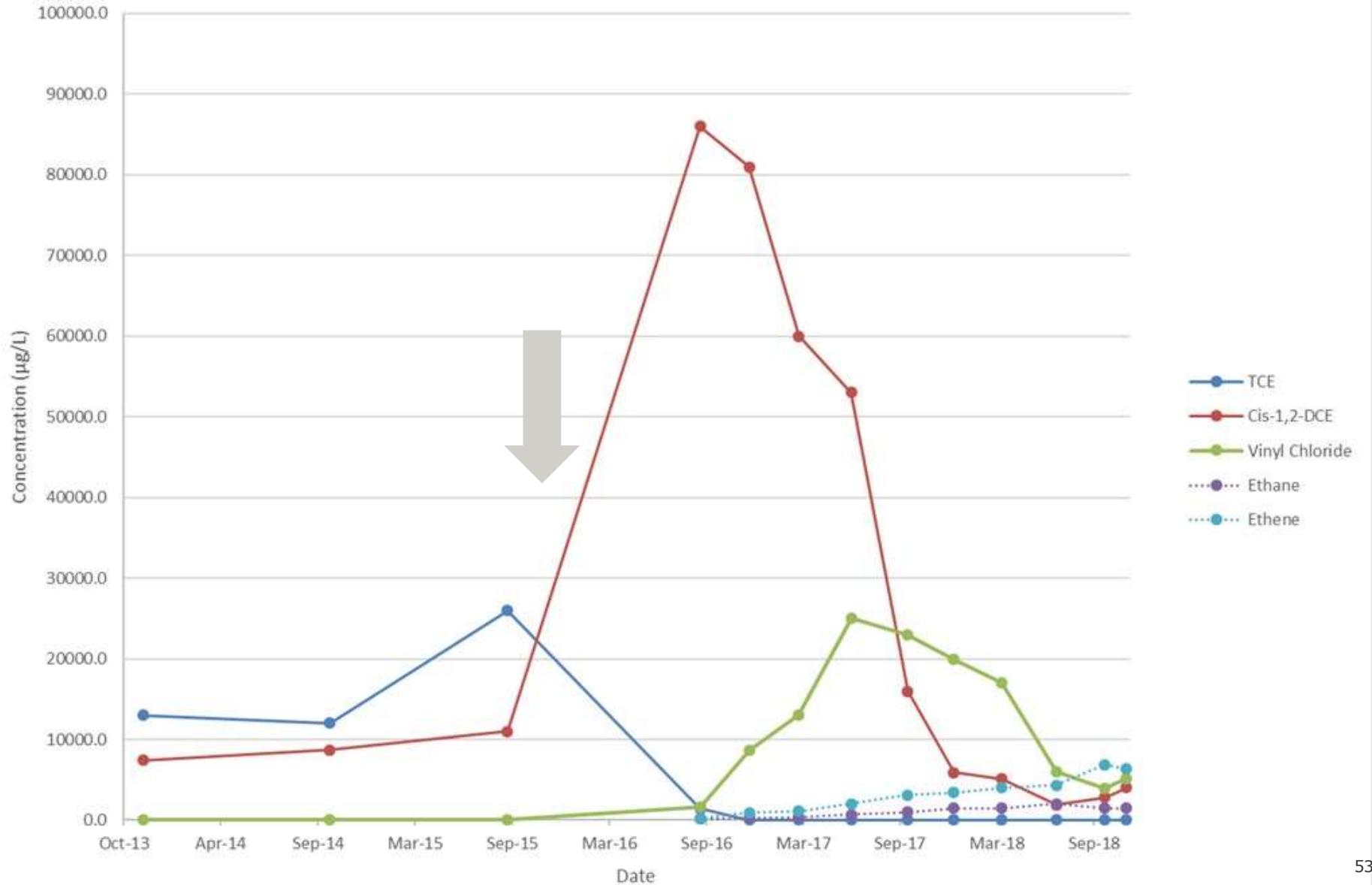
MASS REMOVAL OVER TIME WITH EISB



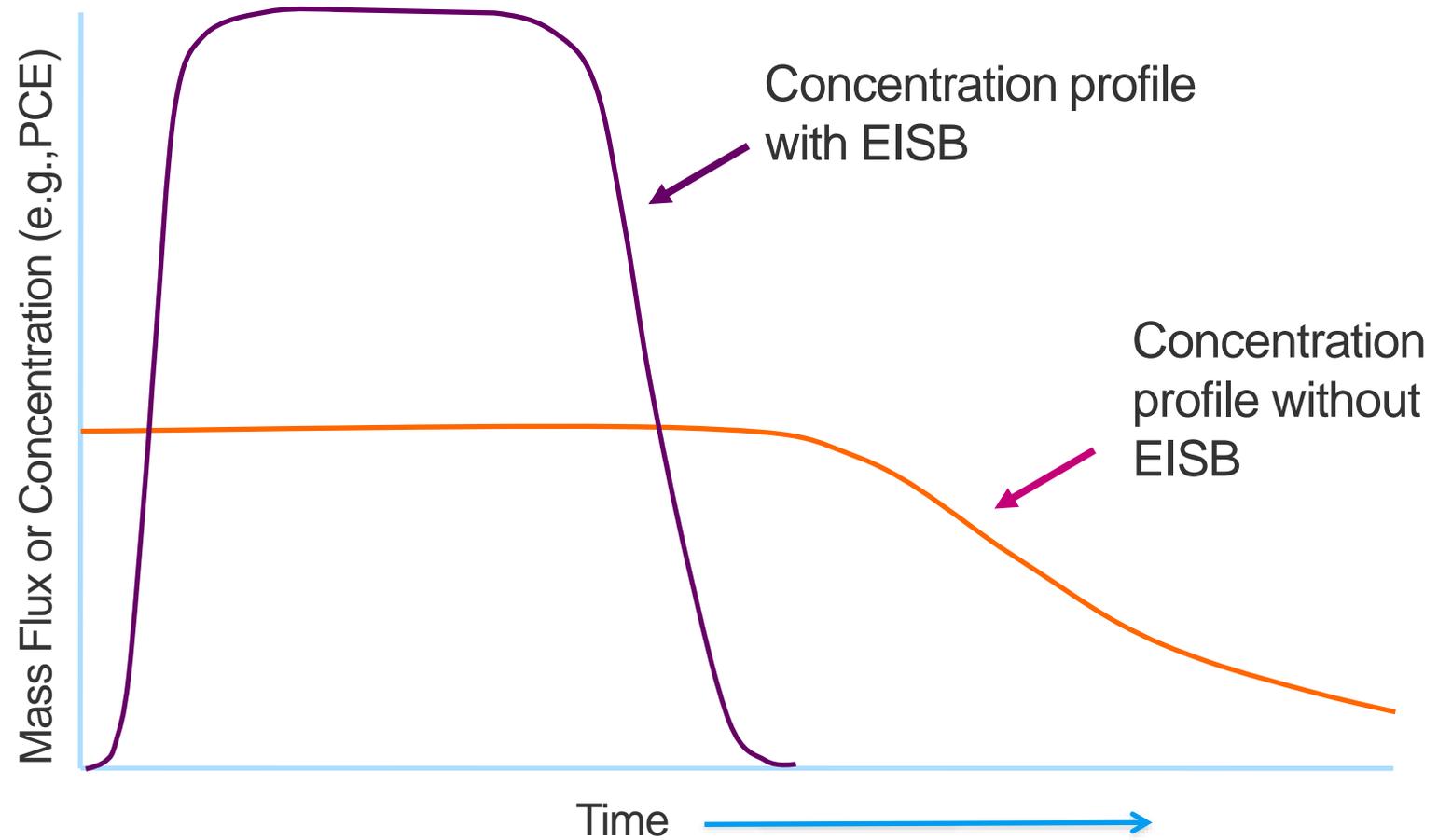
(ITRC Modeling Study)

PERFORMANCE MONITORING

Former Wire Mill Area
DM-421-G



COMPARISON OF MASS REMOVAL OVER TIME



DISSOLUTION ENHANCEMENT

How much should we expect?

	Compound	
	PCE	TCE
Solubility (mg/L)	150	1,100
Enhancement in Lab	5-15	~2
Enhancement in Field	3-5	~1.5

Course Code - MHSB



QUESTIONS