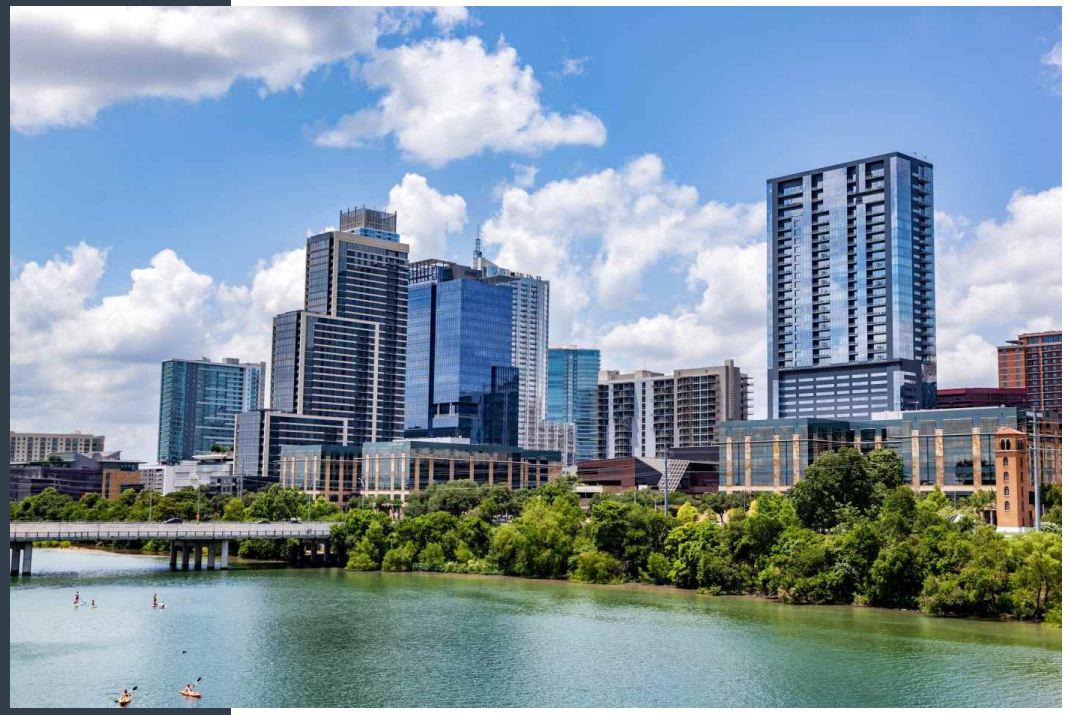
Geosyntec consultants

CREATIVE THINKING
EXCEPTIONAL SOLUTIONS



Advancements in Remedial Performance Assessment at Complex Sites Incorporating Advanced Data Analytics and Innovative Characterization Tools

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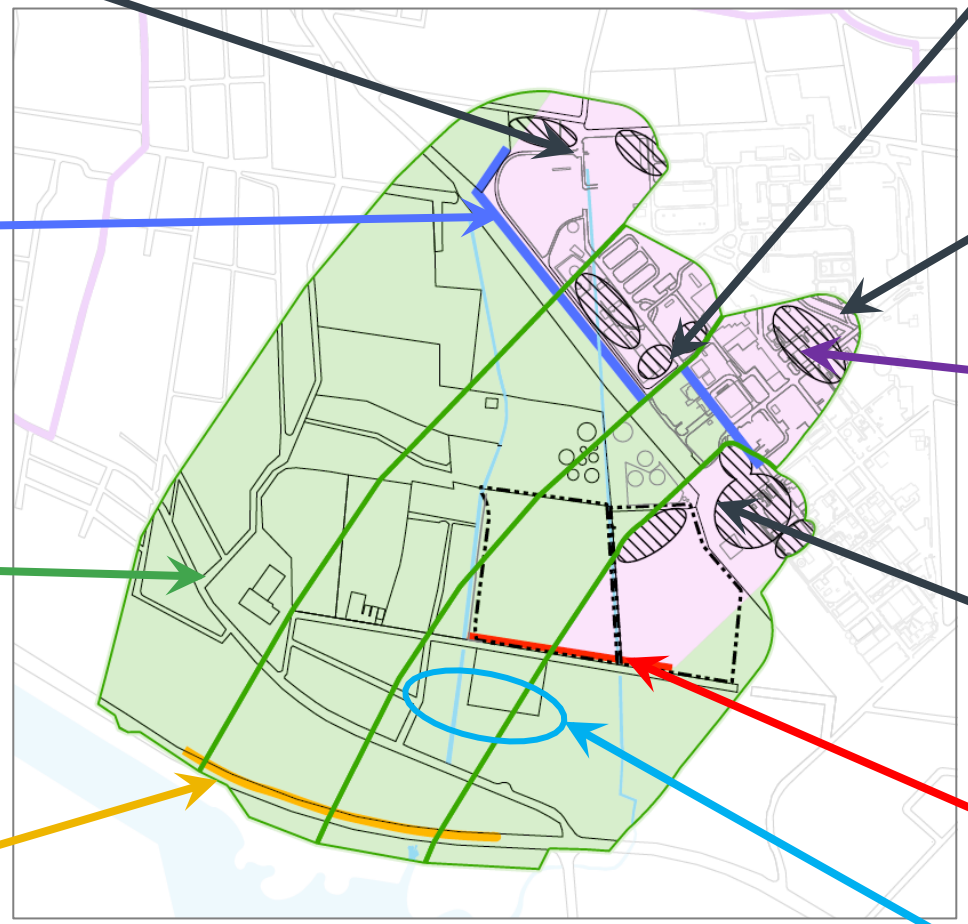
Site Background

N1/N2 SOURCES
Primarily EDC, likely residual sorbed source

ACTIVE EXTRACTION
1st Pump-and-treat containment line

PLUMES
Dissolved mass of varying chemistries, sorption/desorption, diffusion/back-diffusion, biodegradation

ACTIVE EXTRACTION
3rd containment line



N3/N4/N5 SOURCES
N3 – Primarily EDC, may be sorbed source
N4 – CTC/PCE DNAPL, some EDC
N5 – Primarily EDC

C1 SOURCE
>99% EDC DNAPL, trace others

SOURCE AREAS
DNAPL dissolution, desorption, back-diffusion, biodegradation

S1/S2/S3 SOURCES
S1 – CTC and PCE, heavy ends
S2 – CTC and PCE
S3 – TCE and CTC

ACTIVE EXTRACTION
2nd Pump-and-treat containment line

STAGNANT ZONE

Remediation Timeline

1942
Industrial operations and chemical manufacturing began on site

1960s
Larger manufacturing plants were introduced on site making a wider range of chemicals

1970s and 1980-1981
Groundwater contamination was first identified during site development works

1989 through 1990s
Groundwater investigations began and continued through the 1990s

1993-1996
DNAPL sources first identified, and investigations continued through 2000

1999
Zero-valent iron permeable reactive barrier trial completed

2003-2004
Bioremediation trials completed

2005-Present
Construction of the pump-and-treat system (GTP) began in 2005 and operations began in 2006

2006
Total CHCs recovered from groundwater and treated reached 200 tonnes

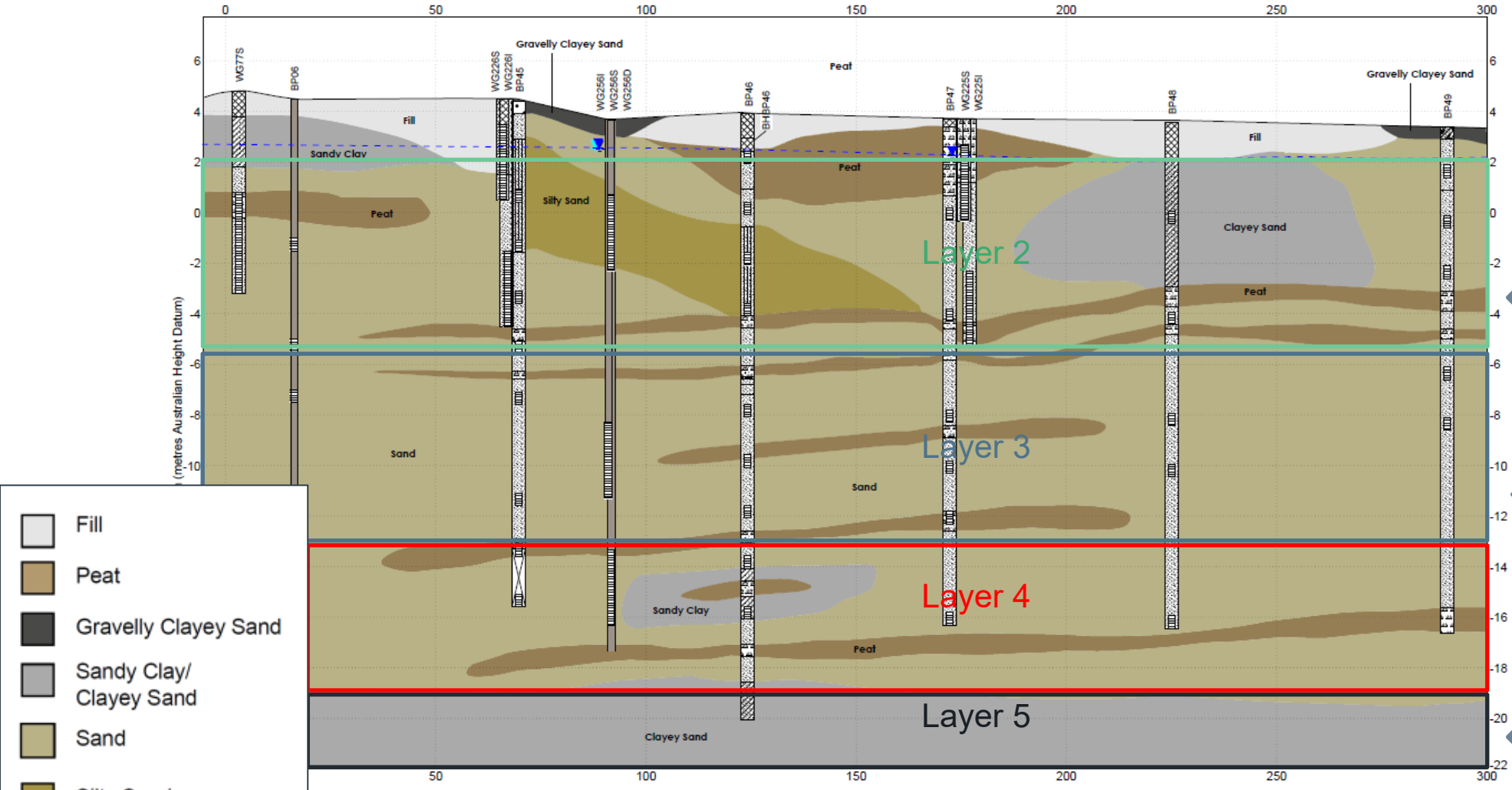
2007
Numerical modeling of GTP performance completed and results indicated GTP may need to operate for >100 years

2014
Updated modeling indicates plumes are disappearing faster than predicted in 2007

2016-Present
Attenuation studies began and continue to present day

2023
GTP has removed ~1,500 tonnes of CHCs

Geological Environment



High organic content = sorption/desorption, Low K = confining layers

Transmissive layers, becoming less transmissive with depth

Low permeability = diffusion/back-diffusion, low transmissivity

Contaminant Physical Properties

Constituent	Single-Component Solubility (mg/L)	Octanol-Water Partition Coefficient (log Kow; mL/g)	Henry's Constant (atm-m3/mole)
1,1,2,2-TeCA	2,900	2.39	3.67e-04
1,1,2-TCA	4,590	1.89	8.24e-04
EDC (1,2-DCA)	8,600	1.48	11.8e-04
TCE	1,280	2.61	98.5e-04
PCE	206	3.40	177e-04
VC	2,700	1.46	278e-04
CTC	793	2.83	276e-04
CF	7,950	1.97	36.7e-04

Legend:	
	DNAPLs dissolve readily, lower retardation (lower sorption/desorption), volatilize readily
	DNAPLs dissolve slower, moderately sorbs/desorbs, moderate volatility
	DNAPLs dissolve slowly, significant retardation/desorption, low volatility

Problem Statement

- Large, expensive pump-and-treat system currently operating
- Multiple DNAPL sources still exist on site
- Questions:
  - How long will the DNAPL sources persist for?
  - How long will the pump-and-treat system have to operate for?
  - When can it reasonably be turned off or ramped down while still managing receptor exposure risk?
  - What is driving persistence of concentrations in the plume, and is there anything we can do to accelerate attenuation?

Lines of Evidence Investigated

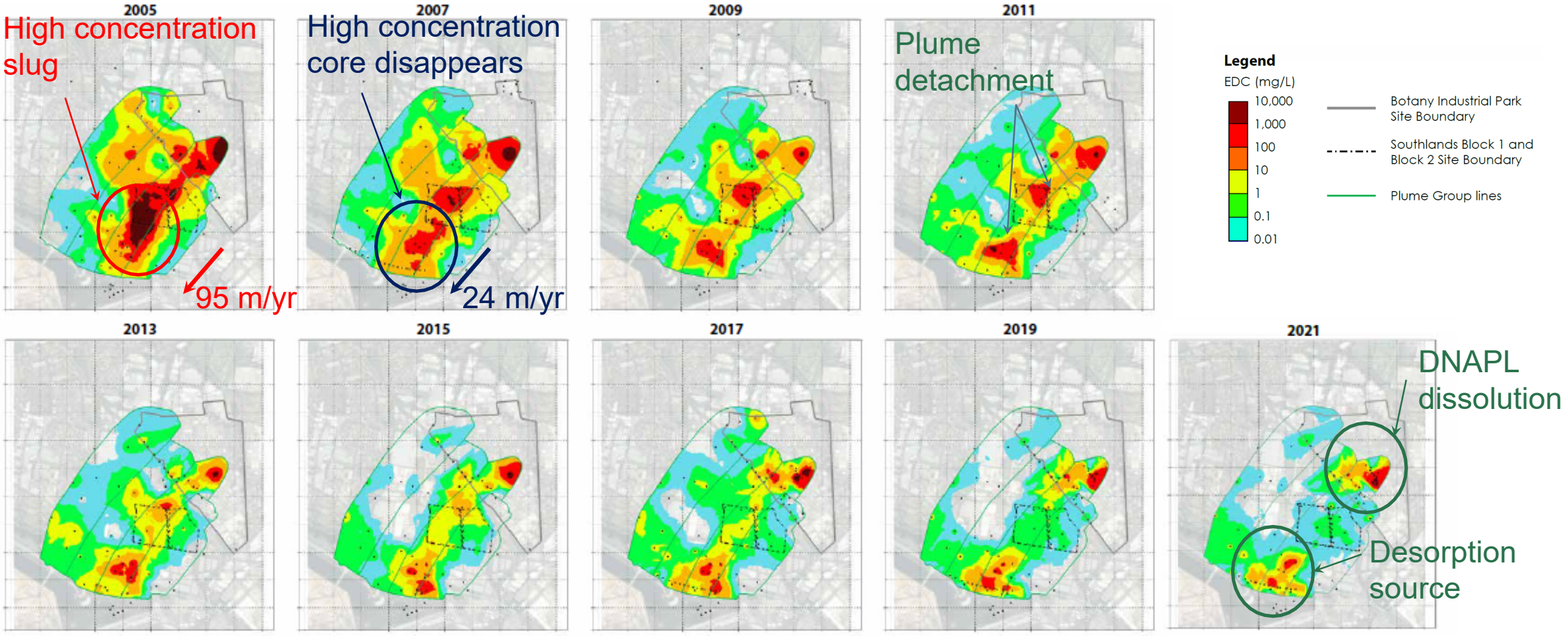
Initial evaluation included:

- Evaluating the contributions from secondary sources
- Assessing natural attenuation rates and mechanisms
- Conducting mass balances to assess impact of pump-and-treat on dissolved phase mass

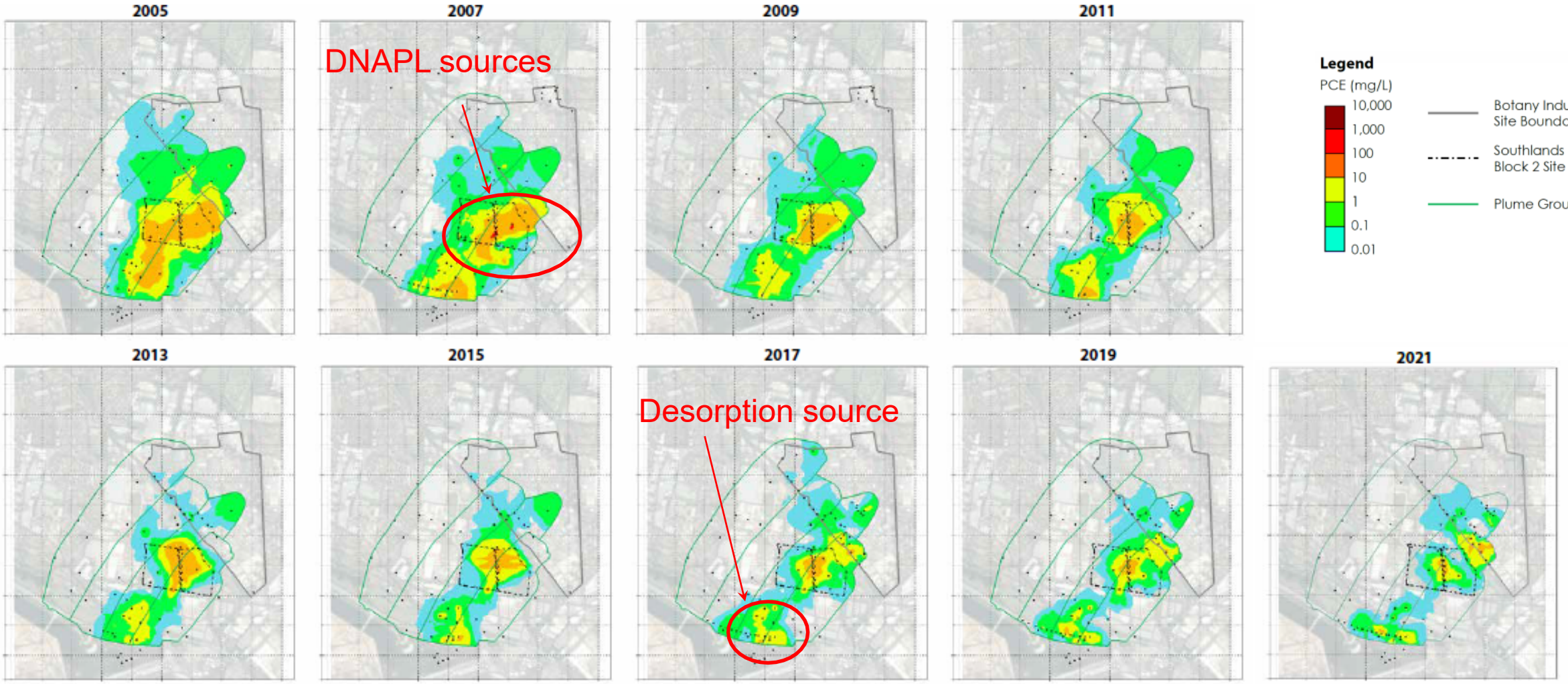
Ongoing assessments (updated every 1-2 years) include:

- Reassessing changes in natural attenuation rates and mechanisms
- Updated assessments of plume attenuation and source mass discharge rates
- Updating mass balances monitoring impact of pump-and-treat system on dissolved phase mass

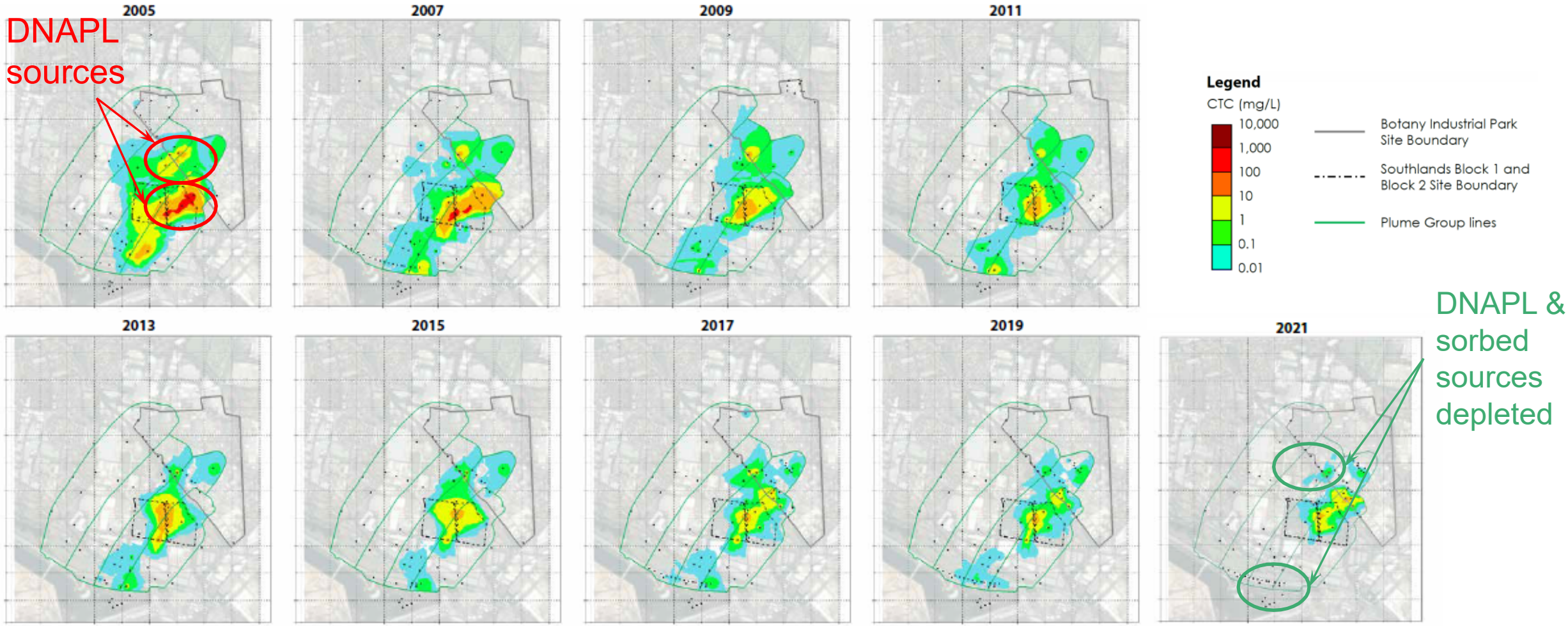
Temporal Changes in Maximum EDC (1,2-DCA) Concentrations



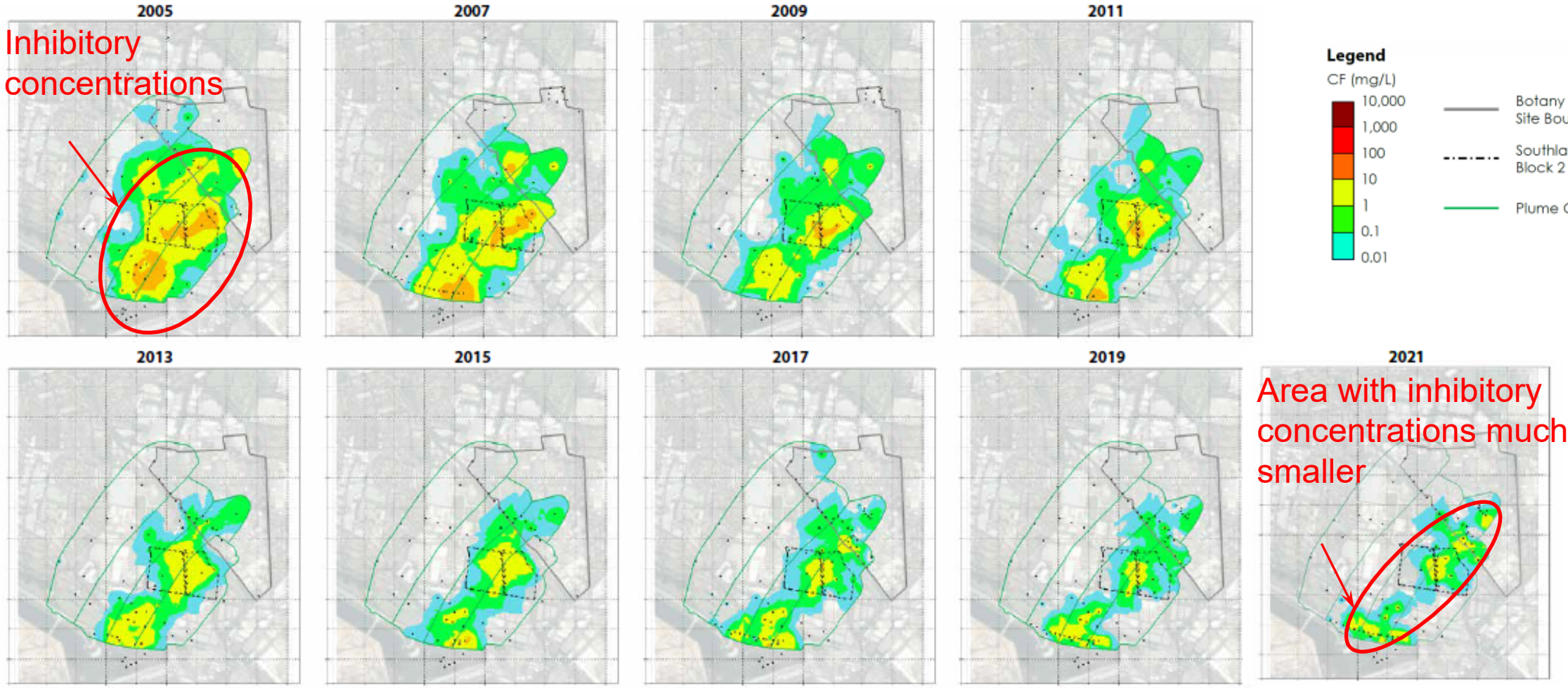
Temporal Changes in Maximum PCE Concentrations



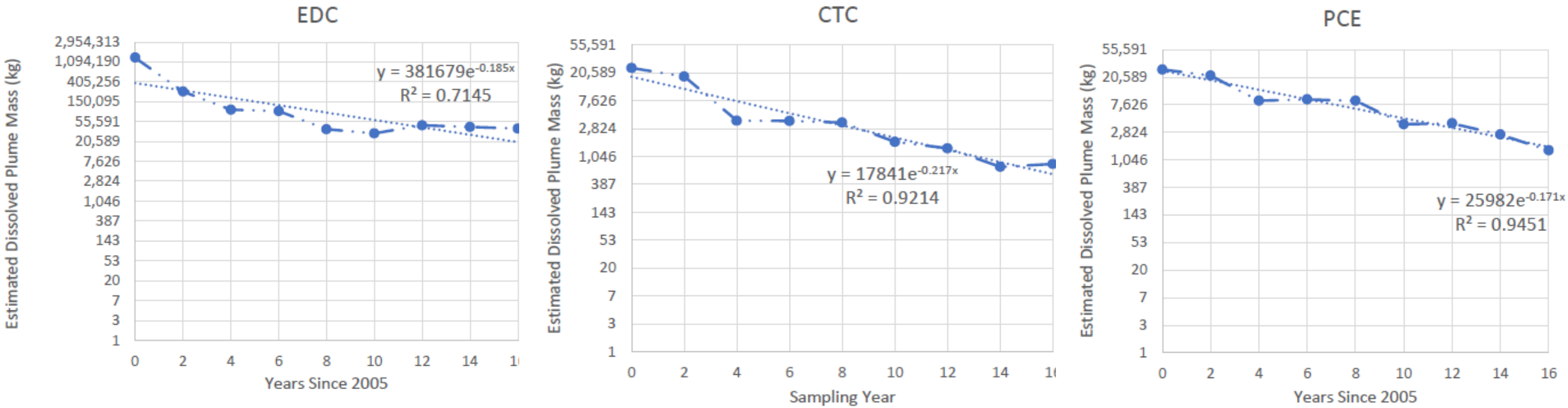
Temporal Changes in Maximum CTC Concentrations



Temporal Changes in Maximum CF Concentrations



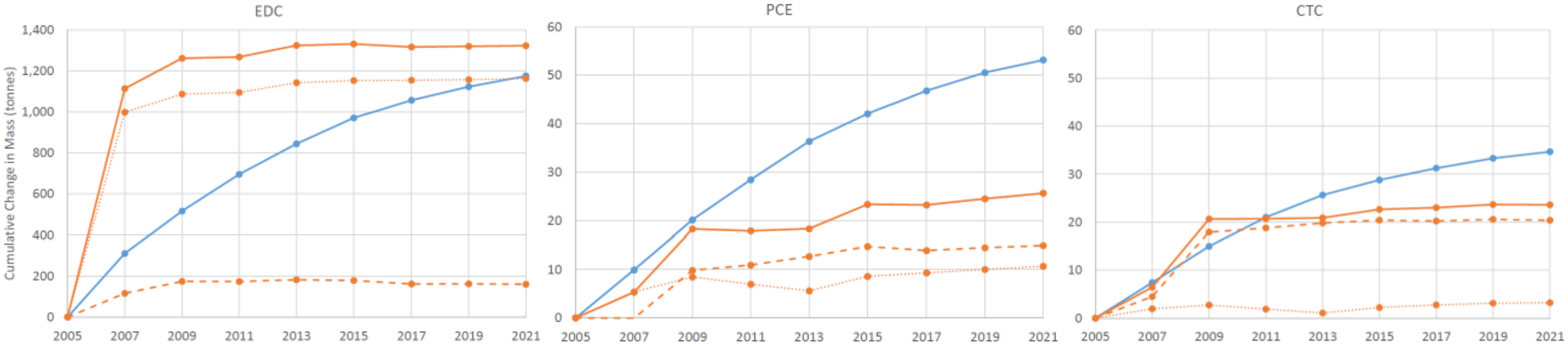
Tracking Dissolved Phase Mass Disappearance



DNAPL Constituent	Average Plume Attenuation Half-Life (years)	Range in Plume Attenuation Half-Lives (years)
EDC	3.8	2.3 to 9.7
CTC	3.2	2.5 to 4.4
PCE	4.0	3.4 to 5.0

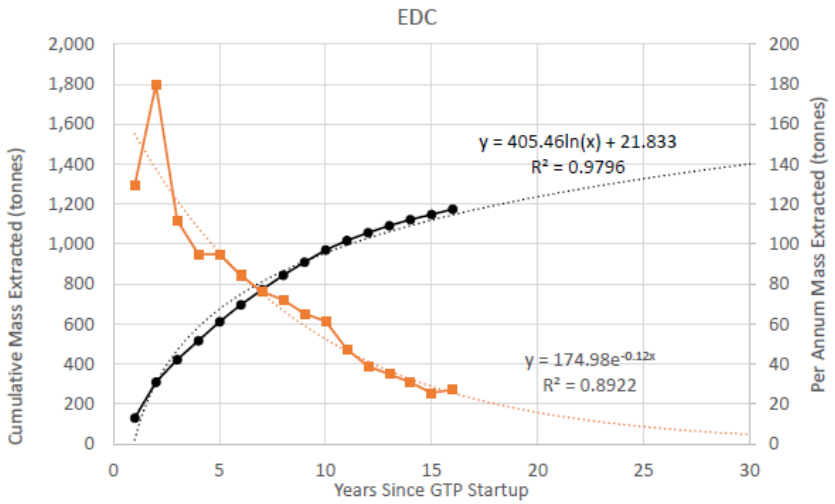
← Beginning to increase

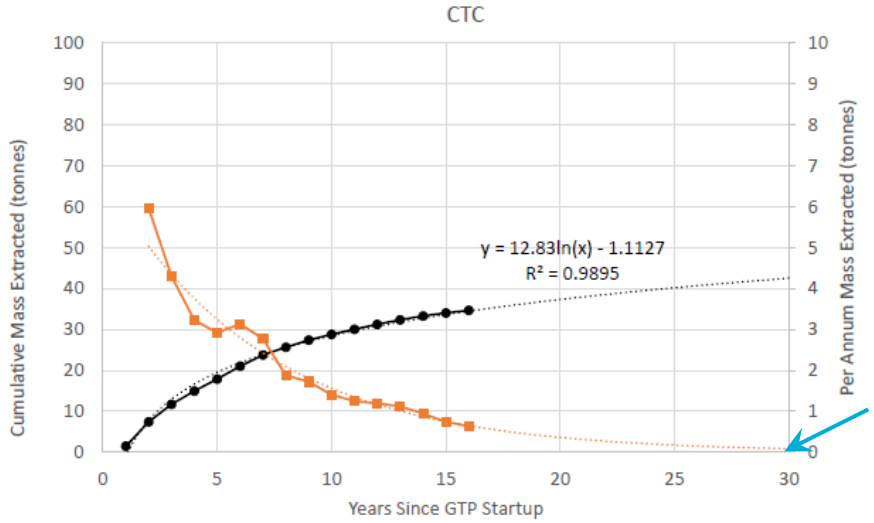
GTP Mass Extraction vs. Changes in Plume Mass

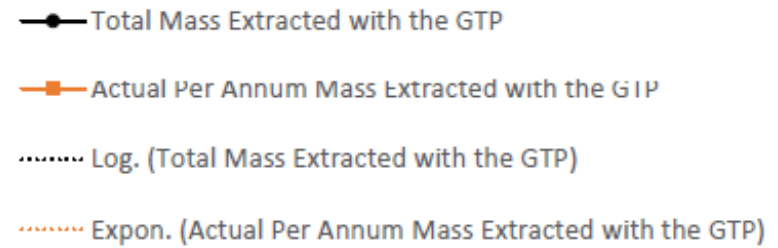


- More EDC (1,2-DCA) mass has disappeared from the plume than can be explained by GTP operation, especially in first two years
- Expanding EDC plume resulted in early-time plume losses due to sorption
- Continuing operation of the GTP is removing PCE and CTC, but having minimal effect on plume mass
- Rates of mass removal are slowing down

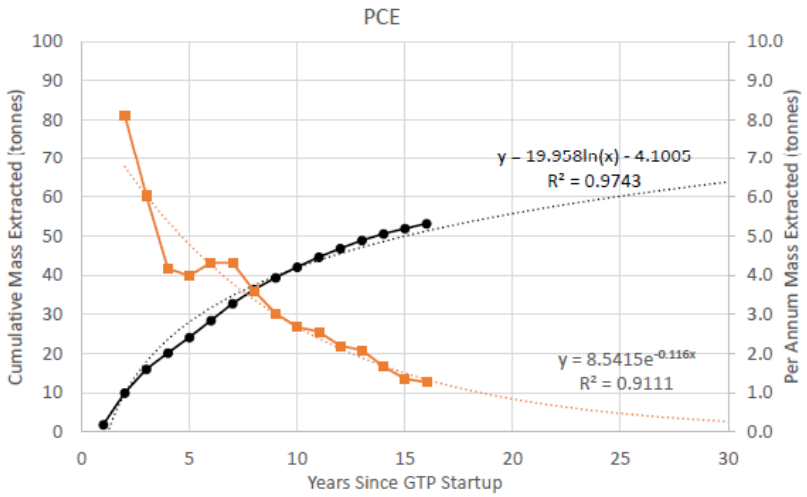
Forward Projections of Total Mass Extracted with the GTP







Diminishing returns over time

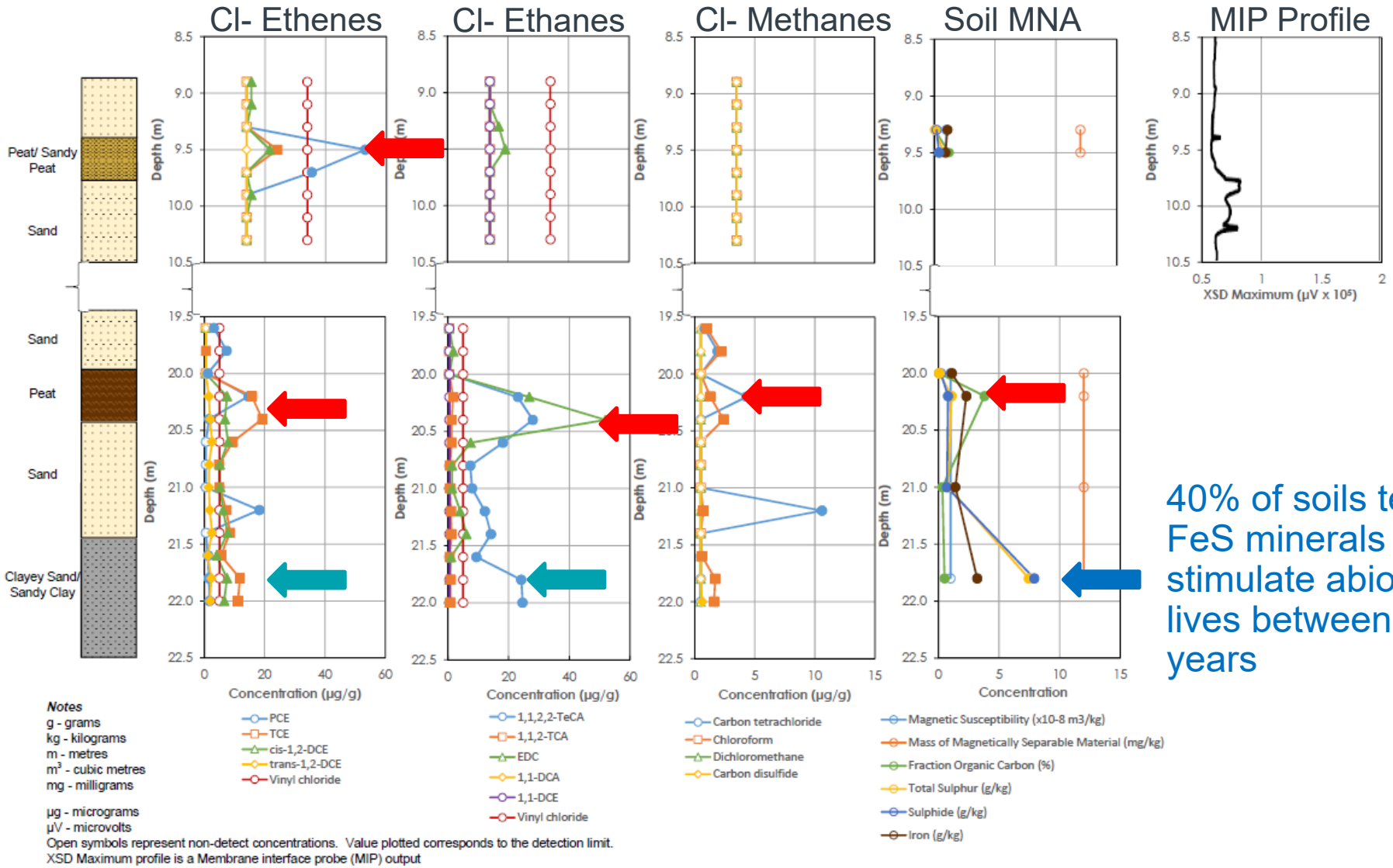


Constituent	30-Year Flushing "Efficiency" (% of total initial mass)
EDC	29% (~1,400 tonnes)
CTC	1.6% (~43 tonnes)
PCE	1.1% (~63 tonnes)

Bulk Attenuation Rates at Individual Locations

DNAPL Constituent	Average Location-Specific Attenuation Half-Life (years)	Range in Location-Specific Attenuation Half-Lives (years)	Number of Locations with Increasing Concentrations
EDC	2.2	0.3 to 10	14 (2%)
CTC	1.8	0.2 to 4.7	1 (0.1%)
PCE	2.3	0.2 to 10	7 (1%)

Sorption and Diffusion as Secondary Sources

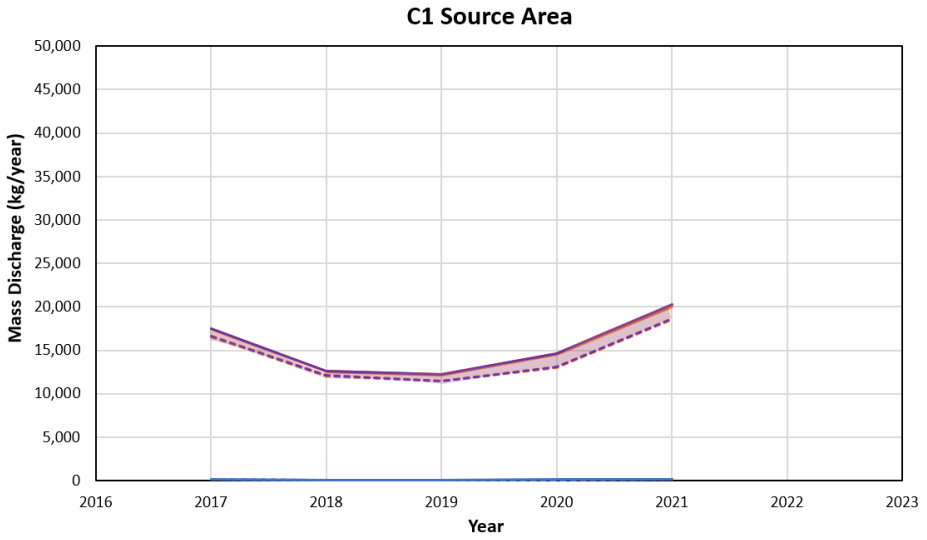


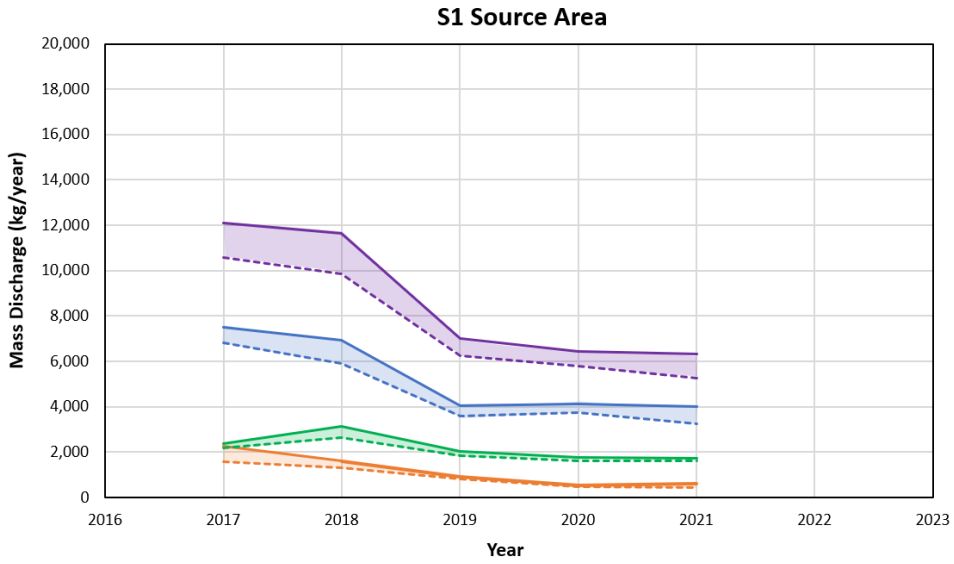
Notes
g - grams
kg - kilograms
m - metres
m^3 - cubic metres
mg - milligrams

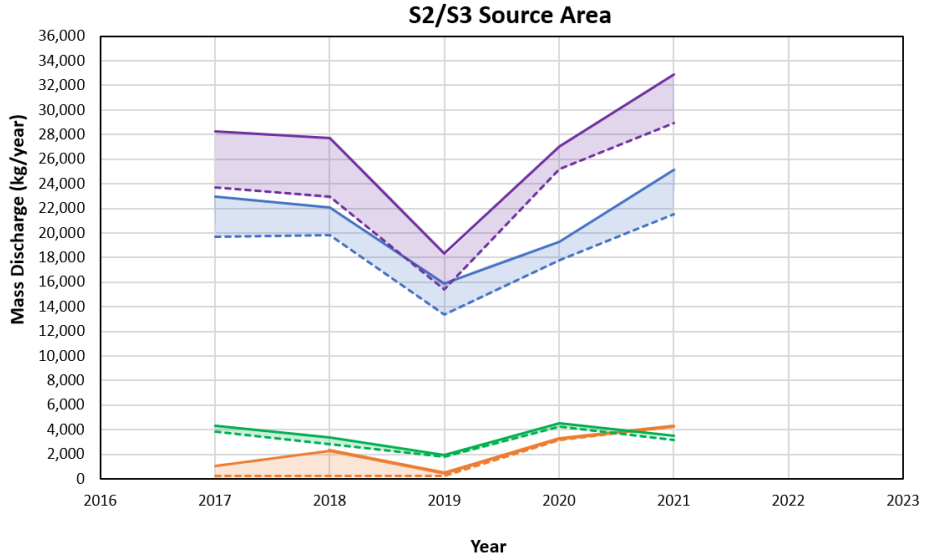
µg - micrograms
µV - microvolts
Open symbols represent non-detect concentrations. Value plotted corresponds to the detection limit.
XSD Maximum profile is a Membrane interface probe (MIP) output

40% of soils tested had FeS minerals sufficient to stimulate abiotic half-lives between 1 to 5 years

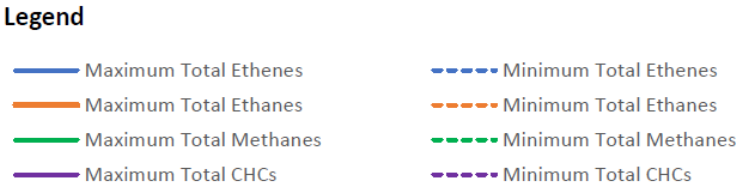
Temporal Trends in Source Mass Discharge



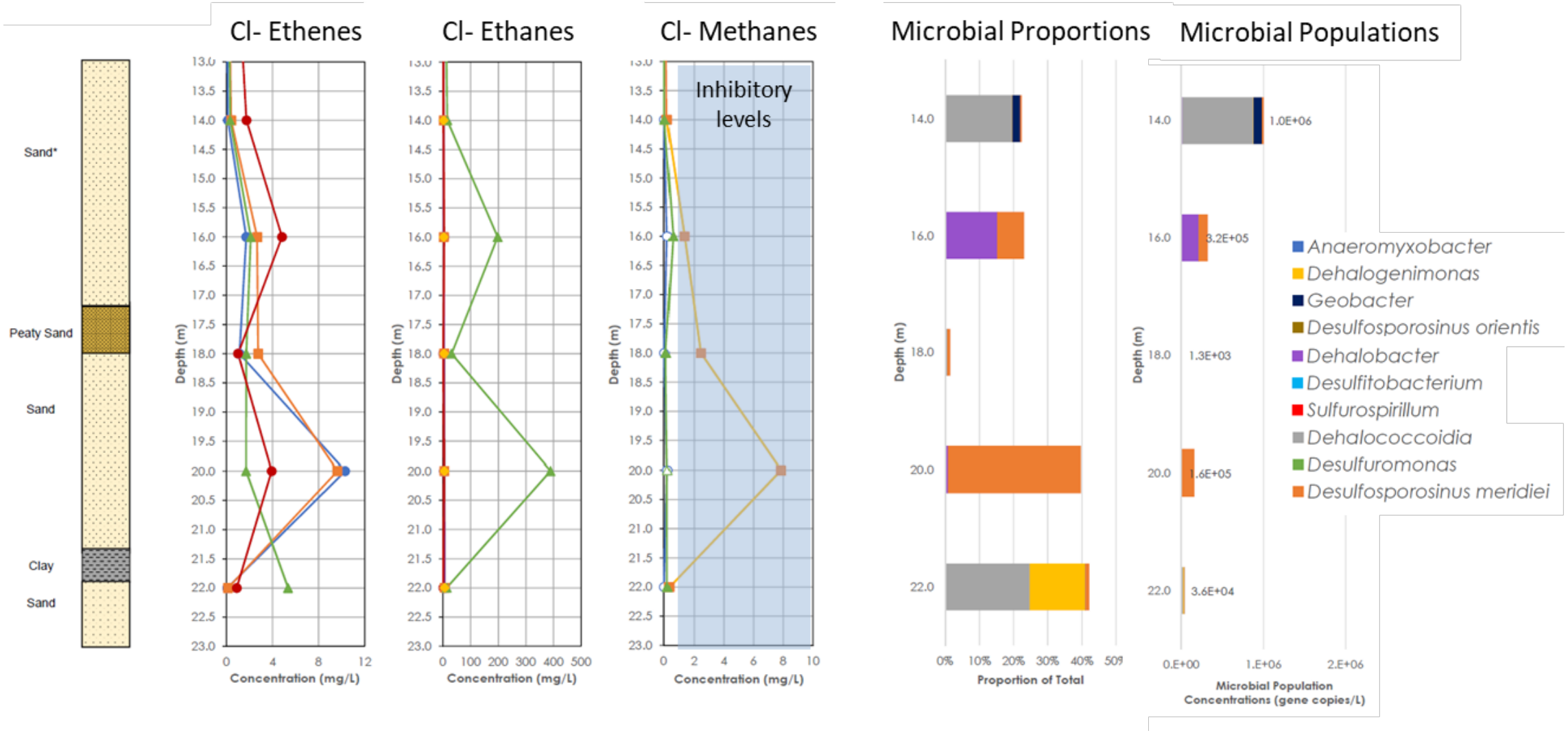




Source Area	Trend in Mass Discharge	Change in Total Mass Discharge (%)
C1	Stable to increasing	+14%
S1	Decreasing	-49%
S2/S3	Fluctuating	+19%



Spatial Variability in Microbial Activity Can Be Significant



Overall Picture

- Bulk mass attenuation behaviour is a complex balance of competing attenuation mechanisms
- Flushing and GTP operation alone will not likely achieve end goals
  - GTP has greater impact on EDC removal than other DNAPL constituents
- In situ degradation is an important mechanism contributing to faster decay of the plumes and sources
  - Abiotic degradation is slow but is occurring
  - Biological degradation is rapid with CTC, contributing to its faster disappearance, slower with other constituents but still substantial
  - Elevated CF is inhibiting bioactivity in some areas of the site, but that volume of aquifer is getting smaller every year

What does the future look like?

- Slowing mass removal with the GTP, but recovery continues to be substantial (10s tonnes/year)
- Desorption/back-diffusion may drive long-term persistence of plumes
- CTC will disappear first
- As CF concentrations reduce below inhibitory levels, microbial populations will likely increase, enhancing rate of mass removal *in situ*
- PCE may be long-term driver for site cleanup

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Questions?

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