

RDT&E for Emerging Contaminants

**Andrea Leeson, Ph.D.
Environmental Restoration Program Manager
SERDP/ESTCP**



Report Documentation Page

Form Approved
OMB No. 0704-0188

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1. REPORT DATE JUN 2010	2. REPORT TYPE	3. DATES COVERED 00-00-2010 to 00-00-2010	
4. TITLE AND SUBTITLE RDT&E for Emerging Contaminants		5a. CONTRACT NUMBER	
		5b. GRANT NUMBER	
		5c. PROGRAM ELEMENT NUMBER	
6. AUTHOR(S)		5d. PROJECT NUMBER	
		5e. TASK NUMBER	
		5f. WORK UNIT NUMBER	
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) Strategic Environmental Research and Development Program (SERDP), Environmental Security Technology Certification Program (ESTCP), 4800 Mark Center Drive, Suite 17D08, Alexandria, VA, 22350-3605		8. PERFORMING ORGANIZATION REPORT NUMBER	
		10. SPONSOR/MONITOR'S ACRONYM(S)	
9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES)		11. SPONSOR/MONITOR'S REPORT NUMBER(S)	
		12. DISTRIBUTION/AVAILABILITY STATEMENT Approved for public release; distribution unlimited	
13. SUPPLEMENTARY NOTES Presented at the NDIA Environment, Energy Security & Sustainability (E2S2) Symposium & Exhibition held 14-17 June 2010 in Denver, CO.			
14. ABSTRACT			
15. SUBJECT TERMS			
16. SECURITY CLASSIFICATION OF:			17. LIMITATION OF ABSTRACT Same as Report (SAR)
a. REPORT unclassified	b. ABSTRACT unclassified	c. THIS PAGE unclassified	
			18. NUMBER OF PAGES 45
19a. NAME OF RESPONSIBLE PERSON			



DoD's Environmental Technology Programs



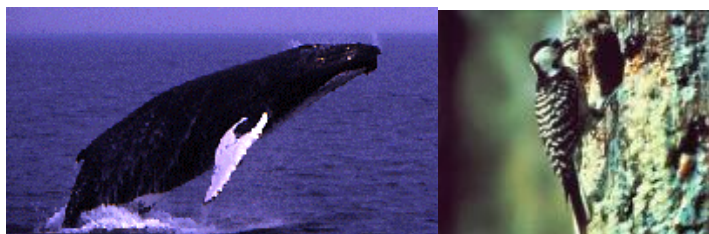
Science and Technology

Demonstration/ Validation



Environmental Drivers

Sustainability of Ranges, Facilities, and Operations



Maritime Sustainability
Threatened and Endangered Species



Toxic Air Emissions and Dust

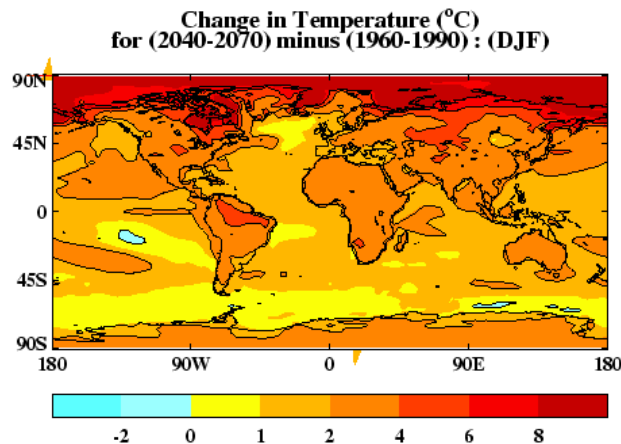


UXO & Munitions
Constituents

Noise



Urban Growth &
Encroachment



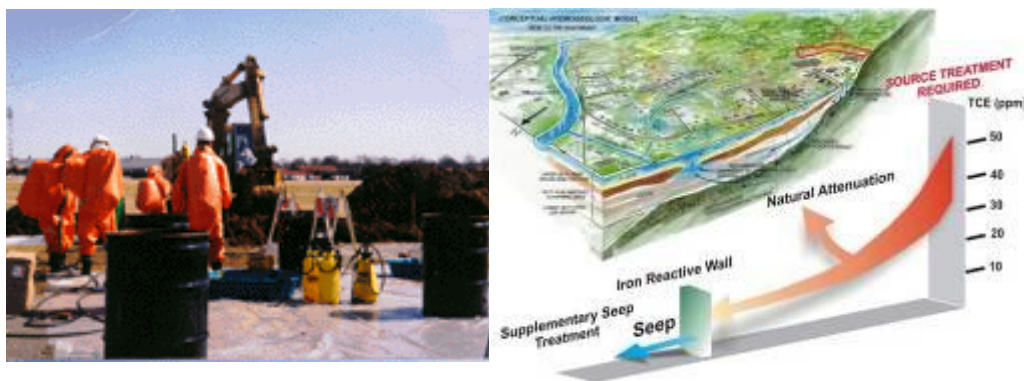
Climate Change
& GHG



Environmental Drivers

Reduction of Current and Future Liability

Contamination from Past Practices



- Groundwater, Soils and Sediments
- Large UXO Liability
- Emerging Contaminants

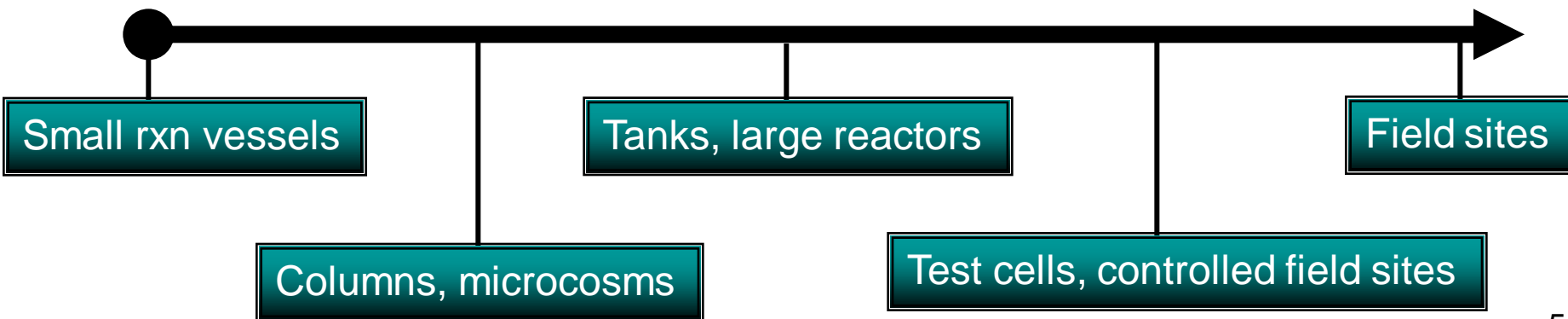
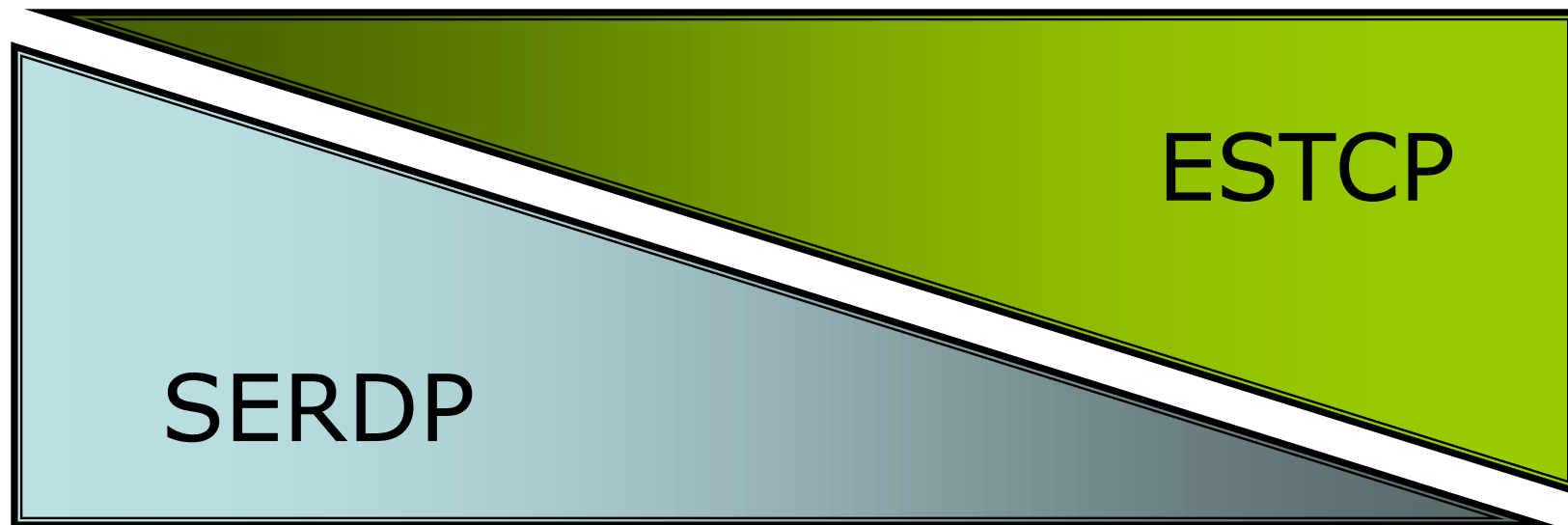
Pollution Prevention to Control Life Cycle Costs



- Elimination of Pollutants and Hazardous Materials in Manufacturing Maintenance & Operations
- Achieve Compliance Through Pollution Prevention



Scales of Research





Emerging Contaminant Defined

- Synthetic or naturally-occurring chemical or microbe
- Not commonly monitored
- Potential to enter the environment and cause known or suspected adverse environmental or health effects
- Sometimes heretofore undetectable



Current Research on Emerging Contaminants

- Perchlorate
- NDMA
- 1,4-Dioxane
- PFCs



Perchlorate Issue




- Broad Use & Occurrence
 - ◆ DoD
 - Rocket propellant
 - Insensitive munitions
 - ◆ Pyrotechnics and flares
 - ◆ Agricultural
 - ◆ Natural deposition





Perchlorate RDT&E

	FY00	FY01	FY02	FY03	FY04	FY05	FY07	FY09
In-Situ Remediation	SERDP	SERDP	SERDP	ESTCP	ESTCP	ESTCP	ESTCP	ESTCP
Eco-toxicology	SERDP	SERDP	SERDP	SERDP	SERDP			
Alternatives			SERDP	SERDP	SERDP	SERDP	SERDP	SERDP
Ex-Situ Treatment	AWWARF	AWWARF	AWWARF	AWWARF	AWWARF	ESTCP	ESTCP	ESTCP
Sources						ESTCP	ESTCP	ESTCP

 SERDP  ESTCP  AWWARF



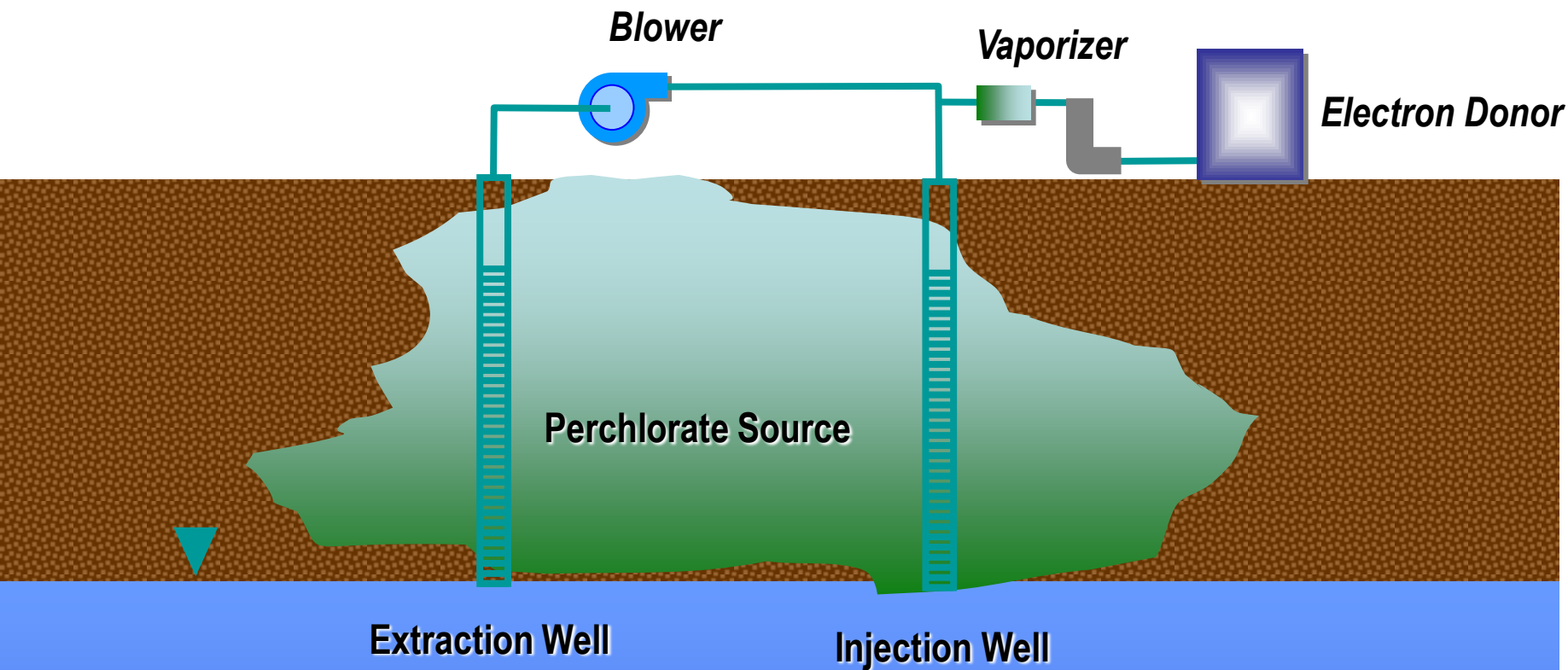
Eco-Toxicology

- SERDP initiated studies in 1998
- A comprehensive program
 - ◆ amphibians
 - ◆ fish
 - ◆ invertebrates
 - ◆ birds
 - ◆ small mammals
- Laboratory and field studies
- Work is the basis for EPA eco-risk assessment
- Investment Completed
- Comprehensive book being written





Vadose Zone Treatment





Ex Situ Treatment

- 1998 drinking water treatment R&D was initiated by an industry consortium (AWWARF)
 - ◆ Completed in 2004
- Successful ESTCP waste water bio-treatment transitioned in 2000
- Only ion-exchange currently used for drinking water
- FY2005 initiatives
 - ◆ ESTCP Congressional program to dem/val new approaches (ion exchange, biotreatment, tailored GAC)
 - ◆ SERDP develop program for next generation treatment

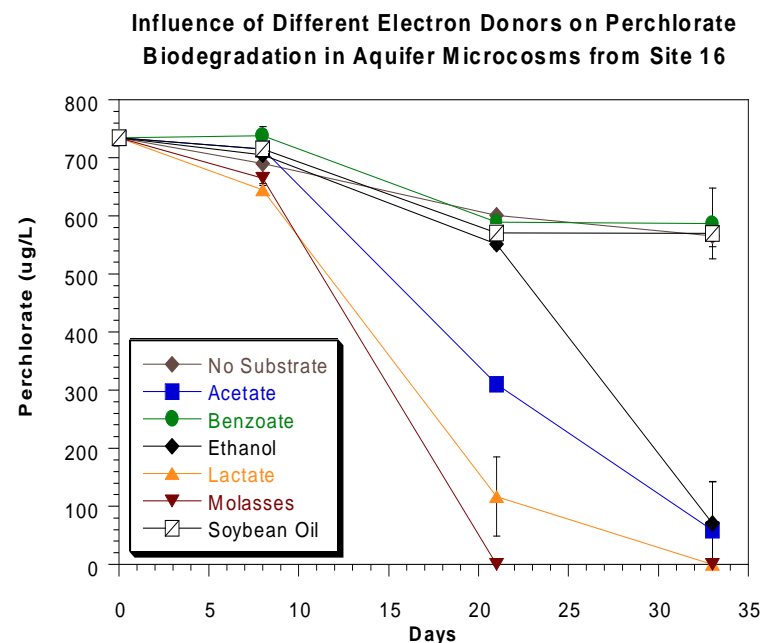


Ex-Situ Bio-Reactor



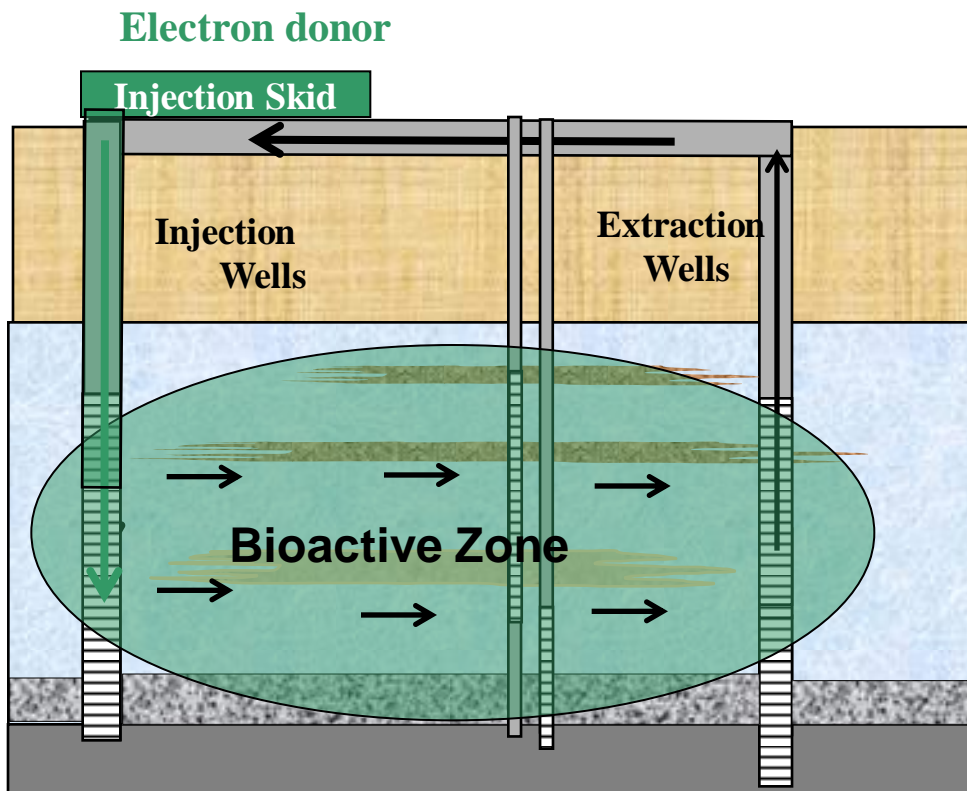
In Situ Treatment

- SERDP initiated bioremediation R&D in 1998
 - ◆ Fundamental and applied studies
 - ◆ Showed potential and method for cost effective treatment
 - ◆ Investment completed
- Dozens of field demonstrations ongoing across DoD
- Fully commercialized
 - ◆ Two full-scale applications



Microbial Biodegradation of Perchlorate

Treatment Approaches



- **Active Treatment**
 - Soluble Electron Donor
 - Continuous pumping
- **Semi-Passive Treatment**
 - Soluble Electron Donor
 - Intermittent Pumping
- **Passive Treatment**
 - Slow Release Electron Donor
 - No Pumping

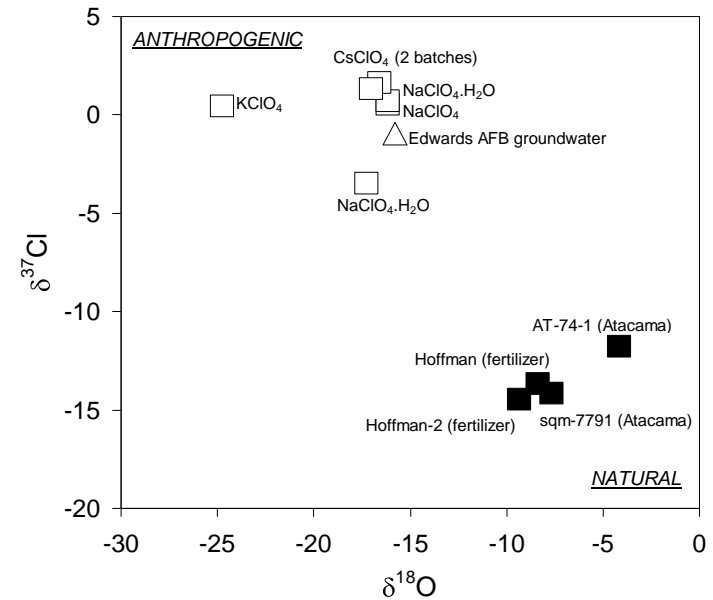
Considerations:

Mixing, O&M Costs, Biofouling,
Secondary Groundwater impacts



Perchlorate Sources

- DoD Sources
 - ◆ Manufacturing
 - ◆ Demilitarization
 - ◆ Test and Training Ranges
- Natural Sources (FY05 Start)
 - ◆ Cause
 - ◆ Distribution
 - ◆ Fate
 - ◆ Identification
- Non Military Sources (FY05 Start)
 - ◆ Magnitude
 - ◆ Extent
 - ◆ Identification



Isotopic Identification of Perchlorate Sources

Road Flares

- **Background**
 - ◆ 20-40 million flares sold annually
- **Laboratory**
 - ◆ Lab studies showed 5-6% potassium perchlorate in unburned flares (10g for a 15 min flare)
 - ◆ Complete burning reduced perchlorate by 99% - still have up to 66 mg perchlorate in flare residue
- **Field**
 - ◆ Monitoring of background levels of perchlorate in highway runoff
 - ◆ Monitored highway run-off near a road flare deployed by State Police at an accident scene (I-95 MA)
 - ◆ Max ClO_4^- concentration leaving highway: ~ 314,000 PPB
 - ◆ Peak load of ClO_4^- leaving highway : 32.4 mg/min.
 - ◆ Total ClO_4^- load to receiving waters :1.3 g
 - ◆ Flares can be a significant point source of perchlorate





Fireworks

- Background

- ◆ 221 million pounds consumed in U.S. in 2003
- ◆ May contain up to 70 wt% potassium perchlorate
- ◆ Case studies discussing contamination at display sites are limited

- Field Study

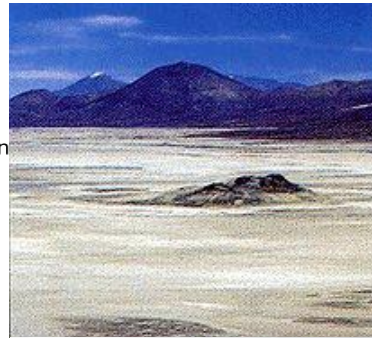
- ◆ Concentration of perchlorate increased from ND to 5 mg/kg after firework display

Perchlorate and Metals Concentrations in Firework Charges

Parameter (mg/kg)	Charge 1	Charge 2
Perchlorate	389,000,000	355,000,000
Aluminum	77,000	120,000
Antimony	ND	ND
Barium	440	190
Calcium	1,700	720
Magnesium	80,000	120,000
Potassium	160,000	160,000
Sodium	ND	150
Strontium	18	22

- Perchlorate concentration in fireworks charge was 389 g/kg. Aluminum, magnesium and potassium were also present at high concentrations

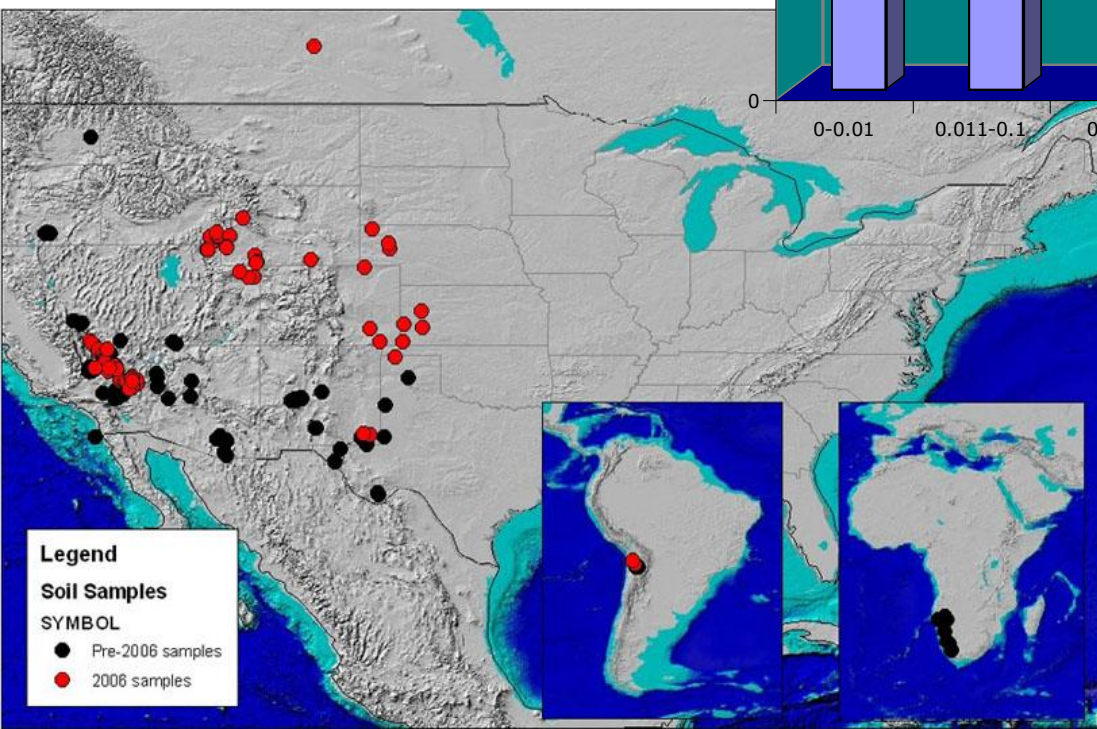
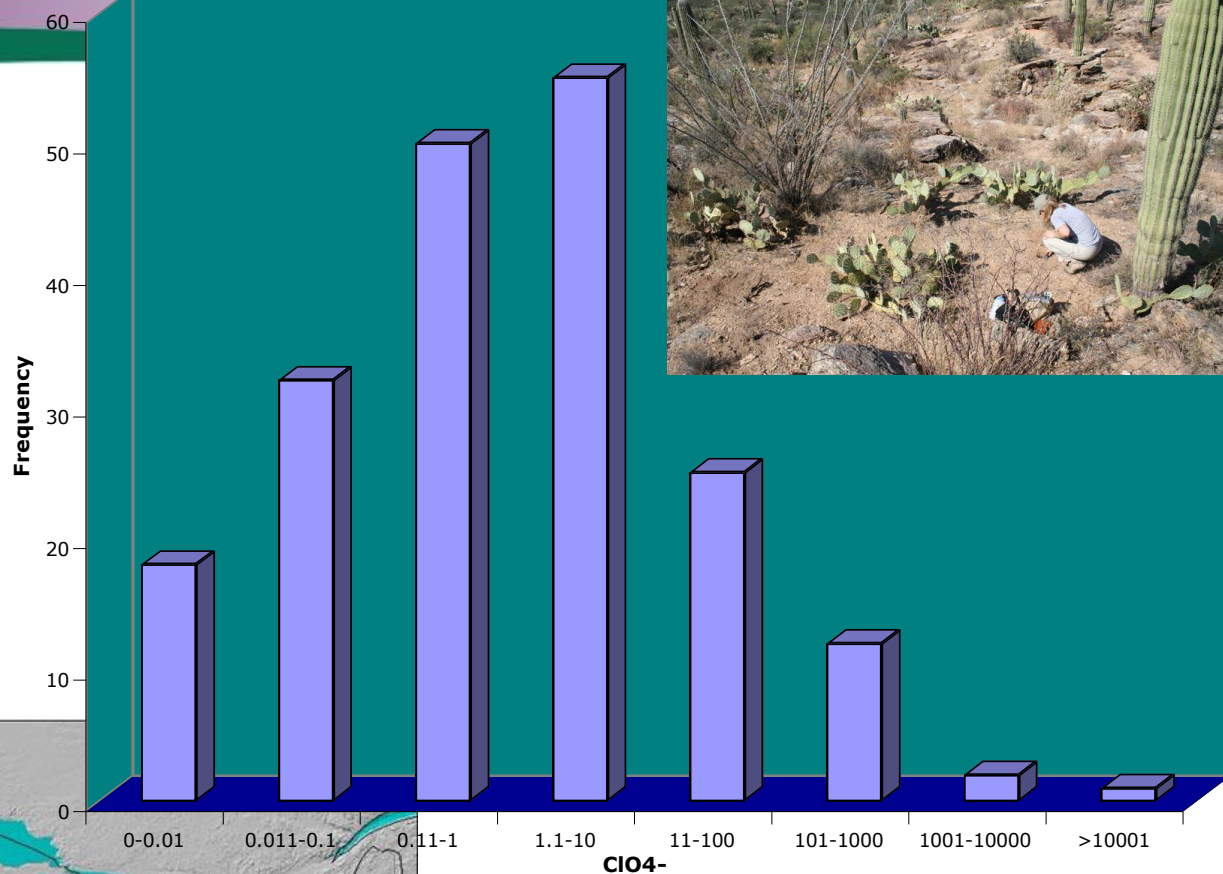
Natural Sources: Where it all started



Chilean NO_3^- Deposits (Atacama Desert)

- ◆ Desert for at least last 1 MY
- ◆ ClO_4^- (>.1%) identified over 100 years ago
- ◆ Deposits also contain IO_3^- , CrO_7 (mg/kg in some strata)

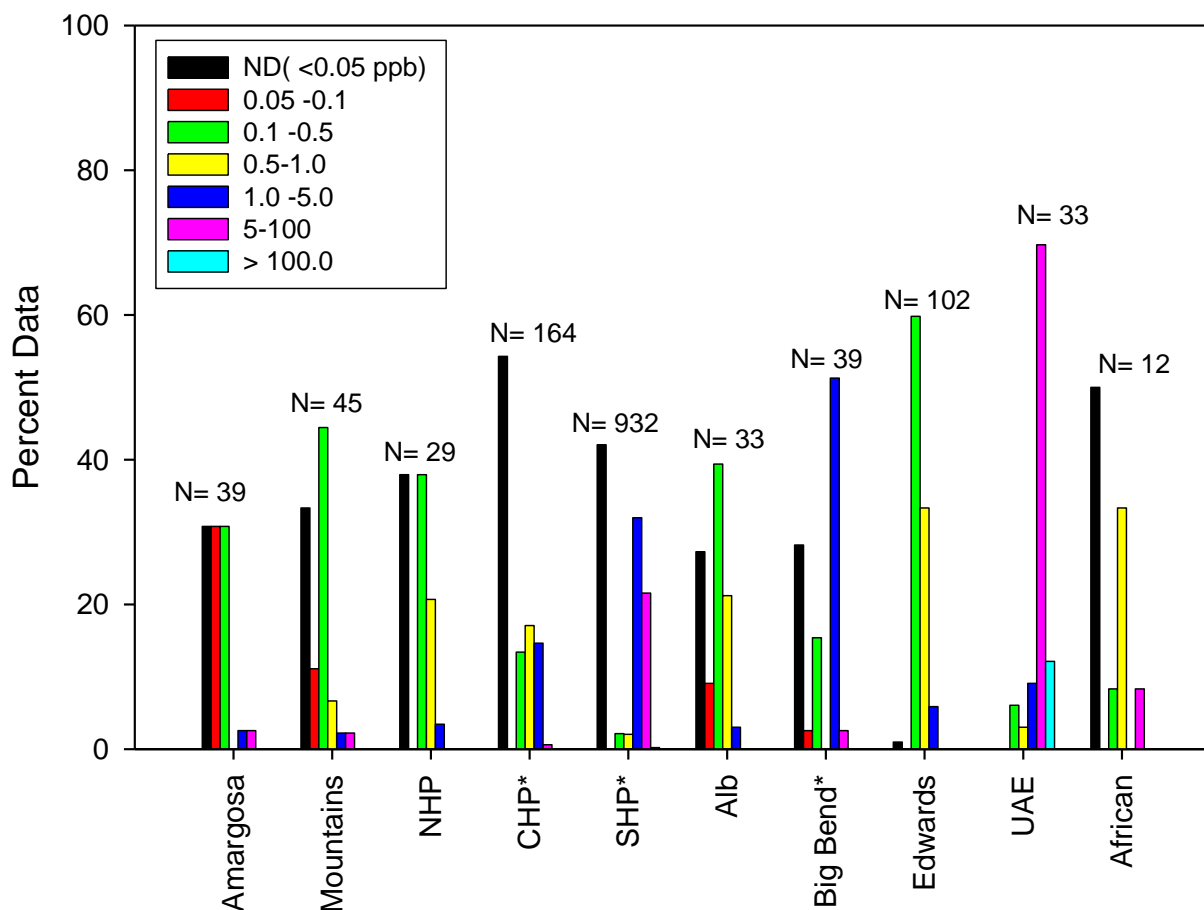
Does Natural Perchlorate Impact other Areas?



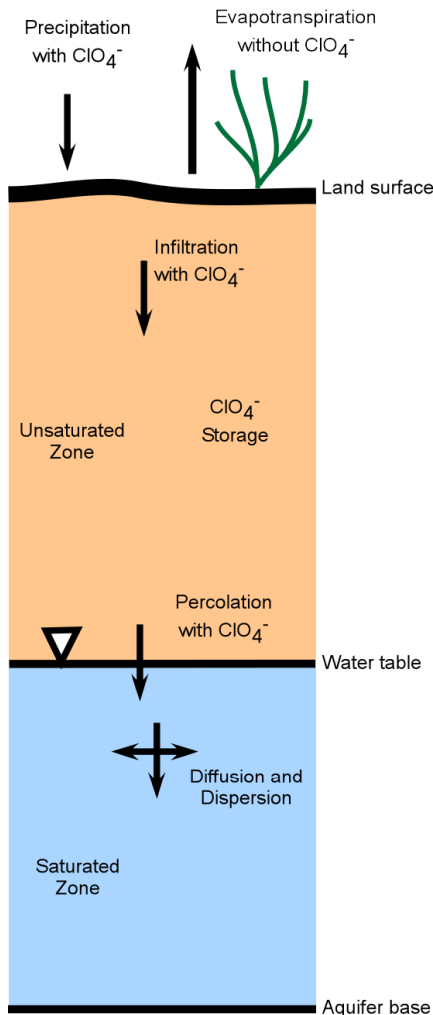
Concentration (ppb)
Distribution of Perchlorate
in Surface Soils



ClO₄⁻ Concentration Distribution in Groundwater from Selected Areas

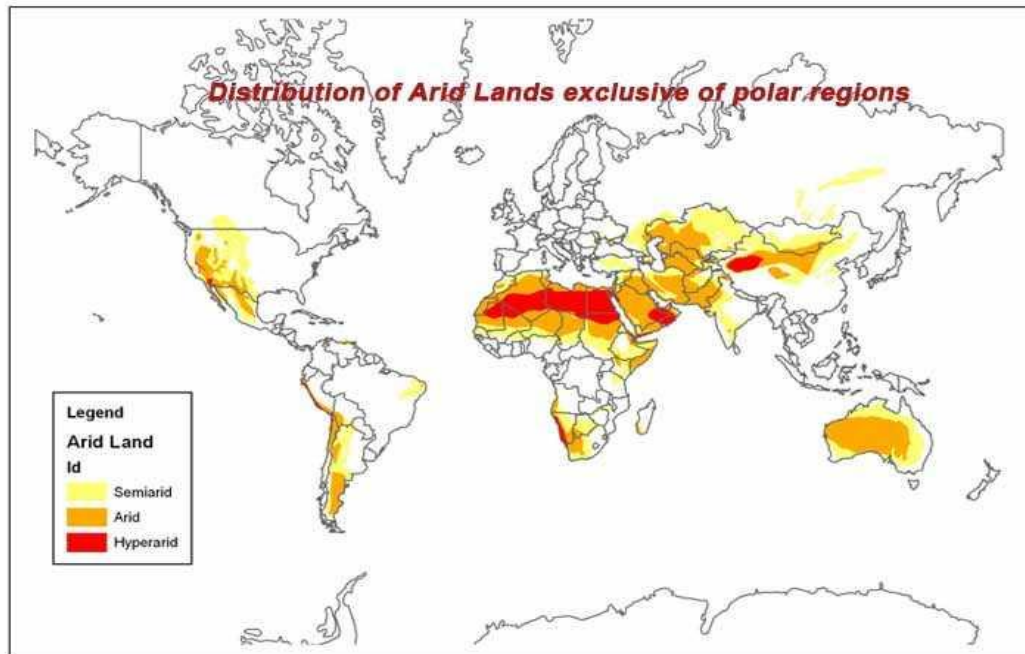


Proposed Perchlorate Accumulation Mechanisms



- ◆ Atmospheric Production and Deposition
- ◆ Partial Transport in Undisturbed Arid Areas
- ◆ Accumulation over long Periods
- ◆ Flushing Possible from Irrigation or Climate Shifts
- ◆ Not Stable in Anaerobic Environments and Some Plant Uptake

What's the Overall Significance?



- Exposure
 - ◆ Plants?
 - ◆ Milk?
 - ◆ GW?
- Future GW impacts
 - ◆ Desert Urbanization
 - ◆ Climate Change
 - ◆ Irrigation
- Site Assessment
 - ◆ Establish Background
 - ◆ Isotopic Differentiation



Natural vs. Anthropogenic Perchlorate

Key Question: Can You Distinguish Natural from Man-Made Perchlorate?



??





Isotope Ratio Analysis to Differentiate Perchlorate Sources

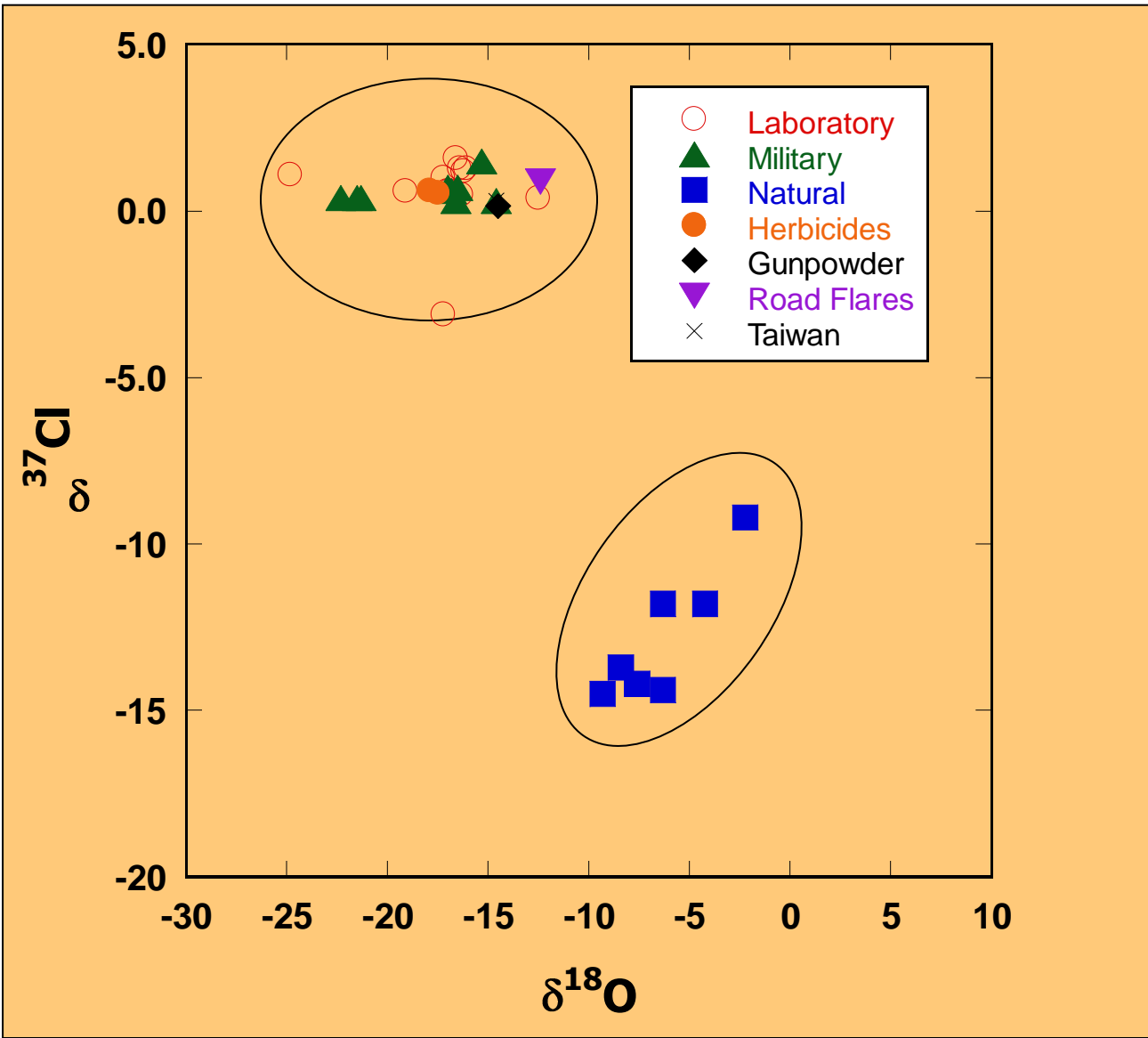
- Objectives
 - ◆ Analyze Isotope Ratios in Commercial, Military, and Natural Perchlorate Sources.
 - Develop broad database quantifying difference between natural and anthropogenic perchlorate.
 - ◆ Analyze Isotope Ratios of Perchlorate in Groundwater Plumes with Anthropogenic Origin and Suspected Natural Sources.
 - Demonstrate/validate isotopic procedure for forensic analysis.
- Elements in a compound can have widely different isotopic ratios based on mode of formation (e.g., ^{18}O in NO_3 from nitrification vs. atmospheric).
- Stable isotope ratios provide a unique “fingerprint” of a chemical compound, another dimension of information invisible from dissolved concentrations.



First Objective: Analyze Isotope Ratios in Commercial, Military, & Natural Perchlorate Sources

- Military sources
 - ◆ Propellant-grade perchlorate
 - ◆ Demilitarization activities
- Commercial sources
 - ◆ Reagent grade perchlorate
 - ◆ Fireworks
 - ◆ Emergency flares
 - ◆ Cotton defoliants
 - ◆ Bleach
- Natural sources
 - ◆ Chilean caliche
 - ◆ Natural fertilizers with Chilean nitrate
 - ◆ Southwest US: Evaporites
 - ◆ Potash salt

Results: Forensic Isotopic Analysis of Perchlorate $\delta^{37}\text{Cl}$ and $\delta^{18}\text{O}$



Chlorine markedly
“heavier” in anthropogenic
Perchlorate (n = 25).

$$\delta^{37}\text{Cl}: 0.6 \pm 0.9$$
$$\text{Range: } -3.1 \text{ to } 1.6$$

$$\delta^{18}\text{O}: -17.2 \pm 2.8$$
$$\text{Range: } -24.8 \text{ to } -12.5$$

Oxygen consistently
“heavier” in natural
Perchlorate (n = 7).

$$\delta^{37}\text{Cl}: -12.8 \pm 2.0$$
$$\text{Range: } -14.5 \text{ to } -9.2$$

$$\delta^{18}\text{O}: -6.3 \pm 2.5$$
$$\text{Range: } -9.3 \text{ to } -2.2$$

NDMA

- Toxicology

- ◆ NDMA is a potent mutagen, teratogen, & carcinogen.
- ◆ EPA 10^{-6} Lifetime Cancer Risk = 0.7 ng/L.
- ◆ California DHS; 10 ng/L Action Level;
California OEEHA 3 ng/L PHG (12/2006)

- Sources

- ◆ 1,1-Dimethylhydrazine Rocket Fuel[(CH_3)₂NNH₂]
- ◆ Aerozine 50 (Mixture of Hydrazine and 1,1DMH)
- ◆ Disinfection Byproduct (Chloramine)
- ◆ Industrial, Agricultural and Food Sources.

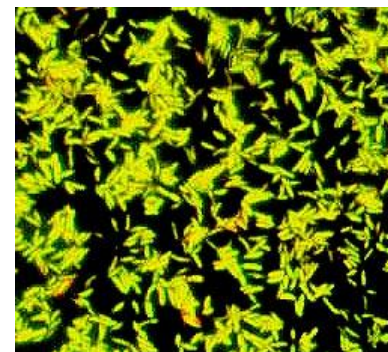
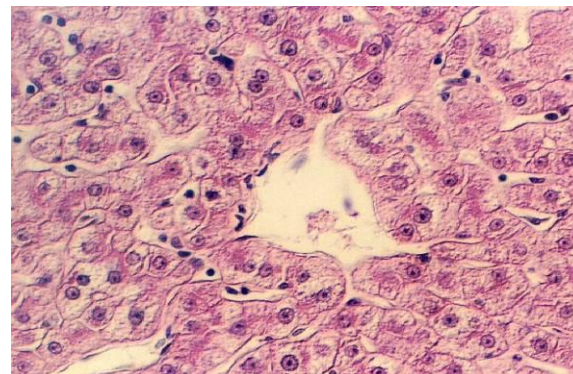
- Treatment

- ◆ Pump-and-Treat with UV Irradiation
- ◆ 1000 mj/cm² for 10-fold reduction
- ◆ (10X for *Cryptosporidium*)



Biological Degradation of NDMA

- Summary of Previous Research:
 - ◆ **Mammalian Metabolism**
 - Cytochrome P-450 System
 - ◆ **Biological Degradation**
 - Several Papers 1970's – 1980's
 - Biodegradation Observed in Soils and Lake Water, Intestinal Bacteria
 - Persistent in Groundwater
 - ◆ **No Environmental Isolates Capable of Growth on NDMA**
 - ◆ **One Isolate Capable of Cometabolism**
 - *Methylosinus trichosporium* OB3b



Potential Remedial Applications



Ex Situ



In Situ





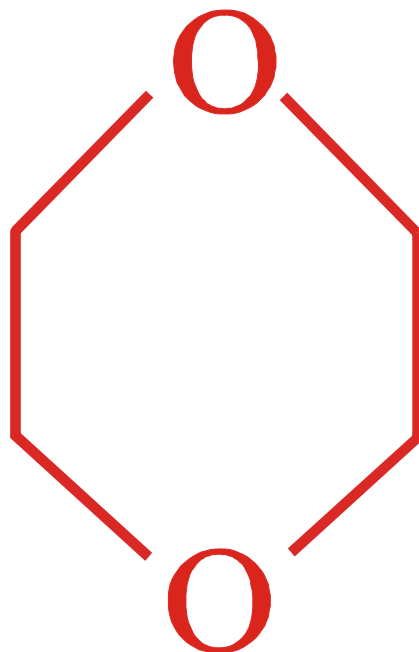
NDMA Summary

- Treatable by UV Oxidation
- *In Situ* and *Ex Situ* Biotreatment Possible
 - ◆ May require propane biostimulation to reach low levels
- *Ex Situ* Metal Catalyst Treatment Showing Promise (*Data not shown*)

1,4-Dioxane



1,4-Dioxane



1,1,1-Trichloroethane



The 1,4-Dioxane Problem

- Used extensively as a stabilizer in chlorinated solvents
 - ◆ Primarily used with 1,1,1-TCA
 - ◆ 1,1,1-TCA found at 809 NPL sites (www.atsdr.gov; 2004)
- 1,4-Dioxane has recently emerged as a contaminant of concern
 - ◆ Low action levels in several states: California (3 ppb); Florida (5ppb); Maine (70 ppb); Massachusetts (50 ppb); Michigan (1 ppb); North Carolina (7ppb)
 - ◆ Risk of closed sites being re-opened
- Little detailed information on the fate of 1,4-dioxane in groundwater
 - ◆ Few biodegradation studies



Current Treatment Options for 1,4-Dioxane

- *In situ* oxidation
 - ◆ Reported to work in some cases
- Advanced Oxidation (HiPOx)
 - ◆ Some full-scale systems in place
- Biological Treatment
 - ◆ Co-metabolic process (propane/THF)
 - ◆ Biological treatment has proven to be challenging
- No universal solution yet available



Perfluoroalkyl Contaminated Groundwater

- FY11 SON: In Situ Remediation of Perfluoroalkyl Contaminated Groundwater
- Objectives:
 - ◆ Improve understanding of mechanisms involved in F&T processes in groundwater under varying natural & engineered conditions.
 - ◆ Determine impact of co-contaminants on F&T processes.
 - ◆ Improve understanding of behavior of perfluoroalkyl contaminants under typical remedial technologies for co-contaminants.
 - ◆ Develop remedial strategies for perfluoroalkyl contaminants, including consideration of the necessity for treatment train approaches to facilitate treatment of co-contaminants.



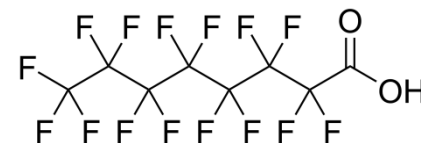
What Are Perfluorochemicals (PFCs)?

- General formula: $F(CF_2)_n-R$
 - ◆ Hydrophobic alkyl chain of varying length (typically C_4 to C_{16})
 - ◆ Hydrophilic end group
- Man-made compounds with unique chemical properties
 - ◆ Very stable and persistent in the environment
 - ◆ Ionic form of PFCs – highly soluble, non-volatile, and poorly sorb to soil
- Primary PFCs of interest

- Perfluorooctane sulfonate (PFOS)



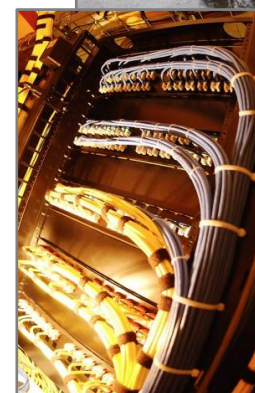
- Perfluorooctanoic acid (PFOA)





What Are PFCs Used For?

- Used to make:
 - ◆ Fluoropolymer coatings and products that resist heat, oil, stains, and grease.
 - Clothing
 - Furniture
 - Food packaging
 - Heat resistant non-stick cooking surfaces
 - Electrical wire insulation
 - ◆ Fluorosurfactants
 - Aqueous film forming foam (AFFF)
 - Chromium plating mist suppressants
 - Stain repellants
 - Photolithographic chemicals





Aqueous Film Forming Foam

- AFFF
 - ◆ Developed in 1960s by 3M and U.S. Navy for use on Class B fires (flammable liquids)
 - ◆ Contains fluorosurfactants other compounds as required) per MILSPEC MIL-F-24385F(SH)
 - ◆ Low surface tension and positive spreading coefficient enable film formation on top of lighter
- PFCs in AFFF
 - ◆ Historically, AFFF contained PFOS and small percentage of PFO (disassociated form of PFOA)
 - ◆ 3M, sole producer of PFOS in the U.S., discontinued production of PFOS in 2001
 - ◆ Continued use of stockpiled PFOS-based AFFF not currently restricted under U.S. regulations
 - ◆ AFFF now produced using smaller chain PFCs (<math><C_6</math>) fuels





Growing Regulatory Interest in PFCs

- Interest driven by findings of PFCs in :
 - ◆ Occurrence in biological organisms and environmental media
 - ◆ Groundwater near PFC manufacturing and disposal facilities
 - DuPont Washington Works Facility, West Virginia
 - 3M Cottage Grove Facility, Minnesota
 - Numerous landfills and disposal sites in Minnesota
 - ◆ Soil and groundwater near fire training facilities in Minnesota
 - ◆ Soil and compost at north Georgia wastewater treatment facility
 - ◆ Sewage sludge and agricultural soils in Alabama
 - ◆ Public water supply systems in New Jersey



Federal Regulation Related to Cleanup

- CERCLA - not a hazardous substance, pollutant, or contaminant
- Not RCRA regulated waste (listed or characteristic)
- PFOA/PFOS not currently regulated under the USEPA Safe Drinking Water Act
 - ◆ Recently included on the USEPA Drinking Water Contaminant Candidate List (CCL3)
- USEPA Provisional Health Advisory Values
 - ◆ PFOA – 0.4 µg/L
 - ◆ PFOS – 0.2 µg/L
 - ◆ Developed in response to contaminated agricultural sites in Alabama but values can be used to assess exposure at other sites
 - ◆ Based on
 - 10-kg child consuming 1 L drinking water per day.
 - Default relative source contribution (RSC) – 20%



State Environmental Guidelines/Action Levels

Guideline / Action Level	Media	PFOA	PFOS
Minnesota Health Risk Limit	Groundwater	0.3 µg/L	0.3 µg/L
North Carolina Interim Maximum Allowable Concentration	Groundwater	2 µg/L	-----
New Jersey Preliminary Guidance Value	Drinking Water	0.04 µg/L	-----
California – under review for possible Prop. 65 listing	NA	√	-----
Washington Persistent Bioaccumulative Toxins Rule	NA	-----	√

Environmental release of PFCs

- Historical testing or emergency activation of fire suppression systems in hangars
- Leaks from storage tanks and pipelines
- Historical fire fighter training exercises





Scope

- Scope of potential impact difficult to define
- Site investigations have not typically included analysis for PFCs, given their emerging status
- Scope of potential problem can be estimated using the number of “Fire/Crash/Training” sites as a surrogate for actual site data
 - ◆ May underestimate problem by not including AFFF spills, pipeline leaks, or testing/emergency activation of aircraft hangar fire suppression systems



Potential Impacts to DoD Restoration Program

DoD Fire/Crash/Training Sites

Service	Total Sites	Remedy in Place (RIP)		Response Complete (RC)	
		RIP ≤ 2008	RIP > 2009	RC ≤ 2008	RC > 2009
<i>Air Force</i>	353	296	47	249	104
Army	94	7	6	79	15
Navy	132	115	17	51	56
DLA	3	1		3	
FUDS	12		1	7	5
<i>Total</i>	594	419	71	389	180



Cleanup Challenges

- Many conventional treatment approaches are not effective for PFCs in water (e.g., direct oxidation, air stripping, vapor extraction)
- Technologies currently available to treat PFCs in water include
 - ◆ Granular activated carbon (GAC) is most effective method
 - Drinking water treatment (municipal and private wells)
 - Landfill water treatment
 - ◆ Reverse osmosis is effective for higher concentration industrial waste streams
- Bench-scale research to develop alternative treatment approaches continues



Home Pages



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