

Bioremediation Study



Santa Susana Field Laboratory Area IV
Energy Technology Engineering Center

Research Team:

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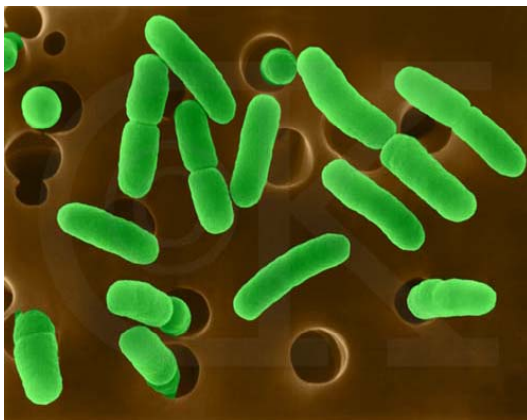
California Polytechnic State University at San Luis Obispo

STIG Presentation 30 May 2013

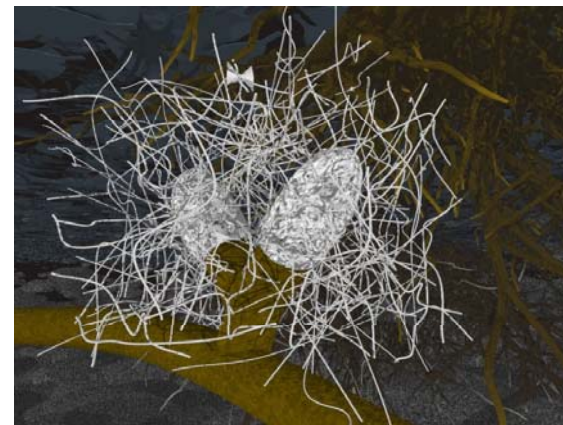


Soil Bacteria and Fungi Biodegrade Contaminants

- Site-specific subsurface conditions determine biodegradability and rate of biodegradation
 - Nutrient availability, oxygen availability, contaminant concentrations, microbial population, pH, etc.



http://www.woodrow.org/teachers/help/te mp_presentations/kim/bioremediation.htm



<http://www.scivit.de/blog/?p=97>

Project Goals:

- Establish a baseline of contaminant biodegradation rates in the field
- Determine which methods could effectively increase biodegradation rates
- Determine which soils in Area IV could be left in place to biodegrade

1. Establish Natural Attenuation

Baseline:

- Identify bacteria and fungi at the site capable of degrading contaminants of concern: PAH, TPH, PCB, dioxins, mercury
- Determine contaminant biodegradation rates in laboratory treatability studies without amendments.

2. Investigate *potential* methods to increase biodegradation rates:

- Biostimulation
 - Soil aeration
 - Nutrient fertilization
- Surfactants to increase bioavailability
- Bioaugmentation (add foreign microorganisms)



Research Plan – Part 1: Field Studies and literature review: Identify contaminant-degrading microbial communities in Area IV soils

1. Collect soil samples from Area IV and culture microbes on plates with contaminants
 - DNA sequencing of isolated organisms
 - Bacteria: 16S rDNA
 - Fungi: ITS rDNA
2. Identify hard-to-cultivate microorganisms with molecular biological tools
 - Extract DNA from soils and amplify with polymerase chain reaction
 - ID with terminal fragment length polymorphism
3. Literature review of organisms identified

Research Plan- Part 2:

Laboratory Treatability Tests:

Soil Microcosms

- Collect soil cores from Area IV
- Incubate in microcosms
- Baseline microcosms: Match field conditions
- Test potential of methods to accelerate biodegradation:
 - Add nutrients (biostimulation)
 - Add surfactant (increase bioavailability)
 - Add microorganisms (bioaugmentation)

Soil Microcosms: Methods

- 500 g contaminated soil (homogenized contaminant concentrations)
- 4-L glass jars
- Teflon-lined lids
- 15% soil moisture content
- 5 replicate microcosms for each condition tested
- Sterile controls: to differentiate between biotic and abiotic transformations



<http://www.loftfield.de/gas/kosmen2.jpg>

Soil Microcosms: Data Collection and Analysis

- Collect 25-g composite samples
 - 0, 6, 12 months after incubation
- Data analysis in California state-certified laboratory
 - Quality Control procedures:
 - Blind duplicates
 - Blanks
 - Contaminant-spiked soils

Expected Outcomes

- Estimate for natural biodegradation rates in the field
- Knowledge of the benefits of acceleration methods
 - Nutrients
 - Surfactants
 - Bioaugmentation
- Estimate potential for accelerated biodegradation rates in the field
- Identification of soils at Area IV which may be amenable to bioremediation without hauling from the site.

Project Timeline

TASK	2013			2014			
	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Review sampling plan with DOE/DTSC/STIG		■					
Field sampling for microbial assays		■					
Microbe isolation and DNA analysis		■					
Literature review of isolated microbes			■				
Laboratory Treatability Testing		■					
Data analysis and interpretation				■			
Presentations to STIG		■		■			■

Questions?

