# Assessment of Native Chain-Elongating Microorganisms in Aquifer Materials from a Superfund Site

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### Introduction

Chlorinated ethenes are contaminants harmful to human health due to factors including their carcinogenicity. They threaten quality of life at hundreds of sites around the US. When provided with an ample  $H_2$  source, the microbial process of reductive dechlorination can convert harmful chlorinated ethenes to a non-toxic byproduct, ethene. Microbial chain elongation (MCE) has recently shown viability as a  $H_2$ producing process for H<sub>2</sub>-consuming reductive dechlorination.. This project explores the possibility of applying MCE in-situ at a Superfund site contaminated with chlorinated ethenes through the stimulation of microorganisms native to the site.

### Objective

- Determine whether native chain-elongators are present
- Determine whether biostimulation of native microbes is ••• feasible at the site
- Inform future remedial action at the site \*\*

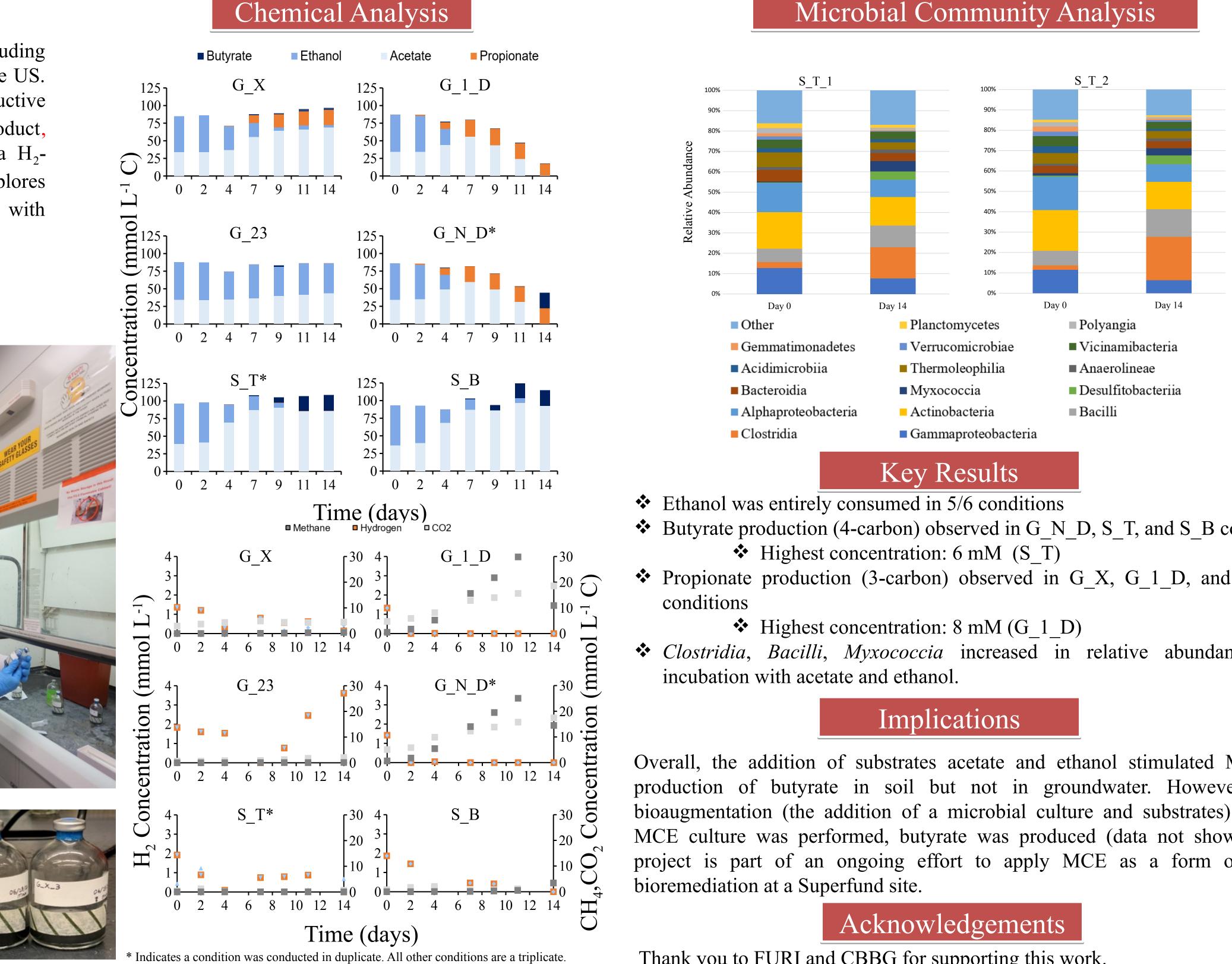
|                 | Microcosm Setup   |                          |                |  |
|-----------------|---|--------------------------|----------------|--|
| Condition label | Location  | Material                 | Nutrients      |  |
| G_X_(1-3)       | Injection/Extraction well<br>used in prior bioremediation | 50 mL <sup>a,b,c</sup>   | N              |  |
| G_I_D_(1-3)     | Monitoring well located                                   | 50 mL <sup>a,b,c,d</sup> | Y              |  |
| G_N_D_(1-3)     | within source area contamination                          | 50 mL <sup>a,b,c</sup>   | Ν              |  |
| G_23_(1-3)      | Monitoring well outside source area contamination         | 50 mL <sup>a,b,c</sup>   | Ν              |  |
| S_T_(1-3)       | Soil (0.3m depth) outside source area contamination       | $25 g^{b,c,e}$           | Y <sup>g</sup> |  |
| S_B_(1-3)       | Soil (1m depth) outside source area contamination         | 25 g <sup>b,c,e</sup>    | Y <sup>g</sup> |  |

### **Legend**

<sup>a</sup>groundwater ,<sup>b</sup>16.2 mM acetate and 27 mM ethanol. °0.2 mM sulfide and 10 mM phosphate buffer <sup>d</sup>50 uL trace A, trace B, 500 uL salt stock solution, <sup>e</sup>soil <sup>g</sup>25 mL anaerobic media containing nutrients, 10 mM phosphate, and 0.2 mM sulfide







## Microbial Community Analysis

Butyrate production (4-carbon) observed in G N D, S T, and S B conditions

✤ Propionate production (3-carbon) observed in G X, G 1 D, and G N D

Clostridia, Bacilli, Myxococcia increased in relative abundance after

Overall, the addition of substrates acetate and ethanol stimulated MCE via production of butyrate in soil but not in groundwater. However, when bioaugmentation (the addition of a microbial culture and substrates) with an MCE culture was performed, butyrate was produced (data not shown). This project is part of an ongoing effort to apply MCE as a form of in-situ

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