

Influence of Wetting and Mass Transfer Properties of Organic Chemical Mixtures in Vadose Zone Materials on Groundwater Contamination by Nonaqueous Phase Liquids



1. Motivation

 Approximately 750,000 kg of CCl₄ was discharged to the vadose zone with large volumes of wastewater (13,400,000L) at the Hanford site.
Assuming equilibrium partitioning, an estimated 64% remains as NAPL in the vadose zone.

•The presence of NAPL has not been observed in soil cores. •Efforts to remove CCl₄ using soil vapor extraction have resulted in removal of only 11% of the estimated original mass. •Either the conceptual model used to calculate CCl₄ mass is incorrect, or NAPL volatilization during SVE is far below what one would expect. •It is difficult to determine if CCl₄ in the vadose zone contributes to groundwater contamination at the Hanford site, or if continued SVE efforts will mitigate this contribution.

2. Objectives

- Overall Objective: Develop a new conceptual model for CCl₄ at the Hanford site based on new information about process coupling in chemical mixtures
- Critical information on NAPL mixtures must be determined before an accurate conceptual model at the Hanford site is possible.
 Organic acids, organic bases, and detergent-like chemicals change surface wettability: such chemicals were present in the wastewater and
- NAPL mixtures discharged at the Hanford site

Specific Objectives

- •Determine the effect of organic chemical mixtures on surface wettability
- •Determine the effect of organic chemical mixtures on CCl₄

volatilization rates from NAPL

•Determine the migration, entrapment, and volatilization of organic chemical mixtures.

3. Background (results from our prior support)

 Vapor density gradients induce gas flow from the atmosphere to the vadose zone and downward liquid flow enhances gas flow
Induced gas flow enhances NAPL evaporation and mass losses to the atmosphere.

•After 38 years, only 54.3 % of total CCl₄ mass discharged was retained in the vadose zone; 42.7% flowed out of the model domain in the vapor phase through the ground surface, and only 2.9% moved across the water table.

•Changes in NAPL wettability and composition of mixtures will alter NAPL migration rates and change these results



Figure 1. (left) Numerical domain, boundary conditions, and distribution of heterogeneous permeability in stratigraphic units. (right) Snapshots of total NAPL saturation (color bar), gas concentration (g/L, contour lines), and gas velocity (vectors).

 SVE removes mass from high permeability zones relatively auickly.

•Residual and trapped NAPL in low permeability zones is removed more slowly.

 The amount of residual and trapped NAPL will increase if conditions change from water to NAPL wetting.



Figure 2. (a) Snapshots of NAPL saturation (color bar) and gas concentrations (g/L, contour lines) at different times. (b) The spatial distributions of total NAPL saturation (color bar) and trapped NAPL saturation (contour lines) at different times.

4. Approach

•Task 1: Prepare and age NAPL mixtures

MIX1 (NAPL) - carbon tetrachloride, lard oil, tributyl phosphate, dibutyl phosphate, dibutyl butyl phosphonate

·Wastewater1 - nitric acid, fluoride, nitrate, phosphate

•MIX2: MIX1

 $\bullet Wastewater2: Wastewatewr1 + tributyl phosphoric acid + bis-(2-ethylhexl) phosphate$

•MIX3: MIX2 + dodecylamine •Wastewater3: Wastewater2 •Task 2: Measure interfacial tension, contact angle, and capillary pressure-saturation curves •Interfacial tension with fresh and reaction-aged NAPL.

Contact angles on quartz and calcite crystals.
Measure two-phase capillary pressure-saturation curves



Figure 4. Interfacial tension data for pure CCl4, pure water, and mixtures

•Organic-water interfacial tension is affected by the additional of DBBP to MIX1 •The water-air interfacial tension is affected by the addition of nitric acid and nitrate salts to the solution





•Chemical aging over more than a month does not appreciably change the interfacial tension results for MIX1 or Wwater1.

 Pre-equilibrating quartz slides in MIX1 markedly reduces the organic-water-quartz contact angle



Figure 6. (left) Contact angle drops and (right) measurements for pure CCl4 and mixtures.

Task 3: Measure interface mass transfer rates

 Image entrapped NAPL and measure volatilization rates of residual and trapped NAPL from micromodels in the presence of residual wastewater



Figure 7. a) Schematic of micromodel. b) Stereomicroscope image of etched micromodel pore network. c&d) Composite order image of NAPL trapped in a homogeneous and heterogeneous pore network.

•Task 4: Intermediate-Scale Column and Flow-Cell Experiment

•Test P-S and K-S relationships with column experiments •Evaluate effects of heterogeneity in flow cell experiments



Figure 8. (left) Photograph of the meter-long column and (right) photograph of intermediate-scale flow-cell with layered sand packing

Task 5: Modeling

•Modify a three-dimensional multiphase flow simulator (STOMP) to model phase partitioning and mass transfer of NAPLs mixtures.

 Adapt and further refine the k-S-P model proposed by Bradford and coworkers and Lenhard and Oostrom

•Use intermediate-scale experimental system to test the STOMP simulator.

•Apply the modified STOMP model to different spill scenarios at the Hanford Site.

 Use the modified model to evaluate how different remediation scenarios can mitigate the impact of CCl₄ in the vadose zone upon groundwater contamination.

