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Publication date:
2018

Document Version
Publisher's PDF, also known as Version of record

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Citation (APA):
Hyldegaard, B. H., Weeth, E. B., Jakobsen, R., Overheu, N., Gent, D., & Ottosen, L. (2018). Design and assessment of electrochemical zones for remediation of chlorinated solvents in natural groundwater aquifer settings. Poster session presented at Eleventh International Conference on Remediation of Chlorinated and Recalcitrant Compounds, Palm Springs, United States.

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Design and assessment of electrochemical zones for remediation of chlorinated solvents in natural groundwater aquifer settings



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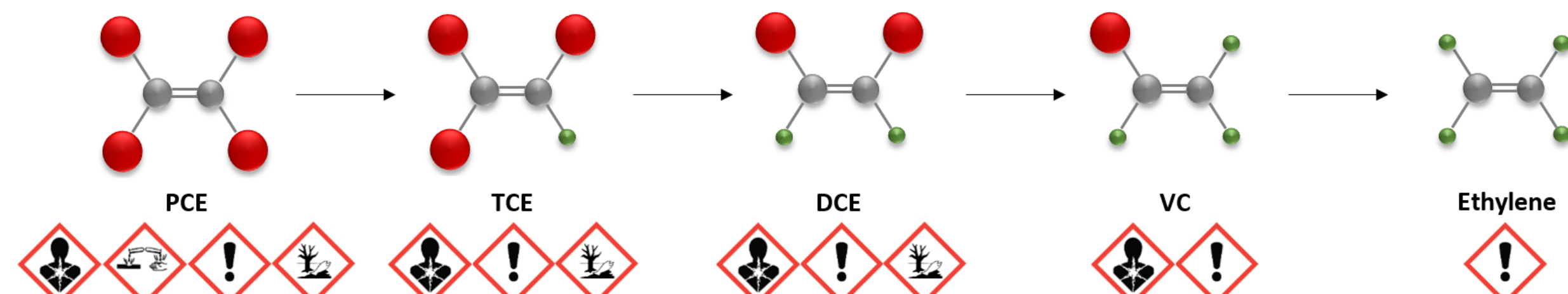
I Project objectives

- Optimization of electrochemical zone(s) for complete degradation of the harmful chlorinated solvents and their chlorinated degradation products in natural hydrogeological settings as a precautionary measure

II Motivation

• Chlorinated solvents threatens the quality of groundwater and cause health risks [1]. Consequently, extraction wells for drinking water are closed

• The compounds' properties challenge the current treatment systems



• Commonly used pump-and-treat systems for hydraulic containment are long-term solutions with substantial operation and maintenance costs

• Optimized means of protecting the groundwater from these contaminants are requested. We propose, establishment of electrochemical zones for *in situ* degradation of chlorinated solvents and degradation products.

IV State of the art

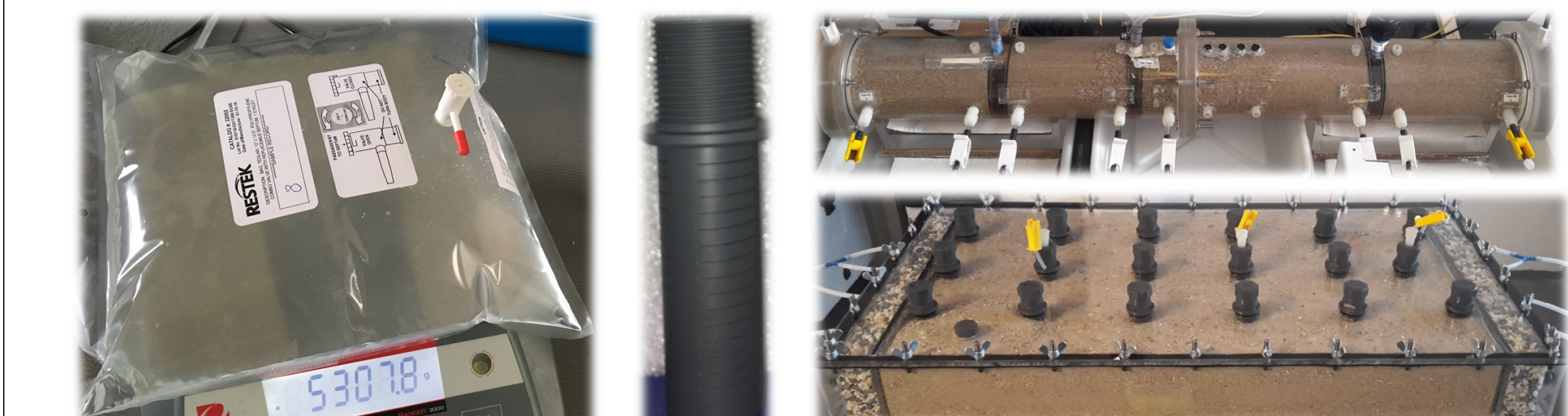
Reduction	Half-cell reaction [5]	E° [V]
PCE	$C_2Cl_4 + H^+ + 2e^- \rightarrow C_2HCl_3 + Cl^-$	0.43
TCE	$C_2HCl_3 + H^+ + 2e^- \rightarrow C_2H_2Cl_2 + Cl^-$	0.42
Cis-DCE	$C_2H_2Cl_2 + H^+ + 2e^- \rightarrow C_2H_3Cl + Cl^-$	0.31
VC	$C_2H_3Cl + H^+ + 2e^- \rightarrow C_2H_4 + Cl^-$	0.38
Oxidation	Half-cell reaction [5]	E° [V]
DCE	$C_2H_2Cl_2 + 4H_2O \rightarrow 2CO_2 + 10H^+ + 8e^- + 2Cl^-$	0.70
VC	$C_2H_3Cl + 4H_2O \rightarrow 2CO_2 + 11H^+ + 10e^- + Cl^-$	0.50

- Focus has been on the influence from electrode materials [6,7,9] and configurations [7,8], and of system parameters such as current density [6,7,8,9,10], flow rate [6,7] etc. in spiked, synthetic liquid phases
- Knowledge gaps between state of the art and field implementation:
 - influence of naturally occurring geochemistry and aged contamination at natural groundwater temperatures

VI The field realistic design

• The field realistic parameters

Sampled groundwater	Conc.	Sampled groundwater	Conc.	Sampled sand	
PCE [µg/l]	40	Ca ²⁺ [mg/l]	370	Porosity [%]	31
TCE [µg/l]	30	K ⁺ [mg/l]	4	Grain density [g/cm ³]	2.57
Cis-1,2-DCE [µg/l]	70	Mg ²⁺ [mg/l]	30	Carbon content [%]	0.95
Trans-1,2-DCE [µg/l]	1	Na ⁺ [mg/l]	25	Chalk content [%]	15
VC [µg/l]	0.1	Cl ⁻ [mg/l]	45	d(0.1) [mm]	0.2
pH [-]	6.9	NO ₃ ⁻ [mg/l]	4	d(0.5) [mm]	0.4
Conductivity [mS/cm]	1.7	SO ₄ ²⁻ [mg/l]	400	d(0.9) [mm]	0.9



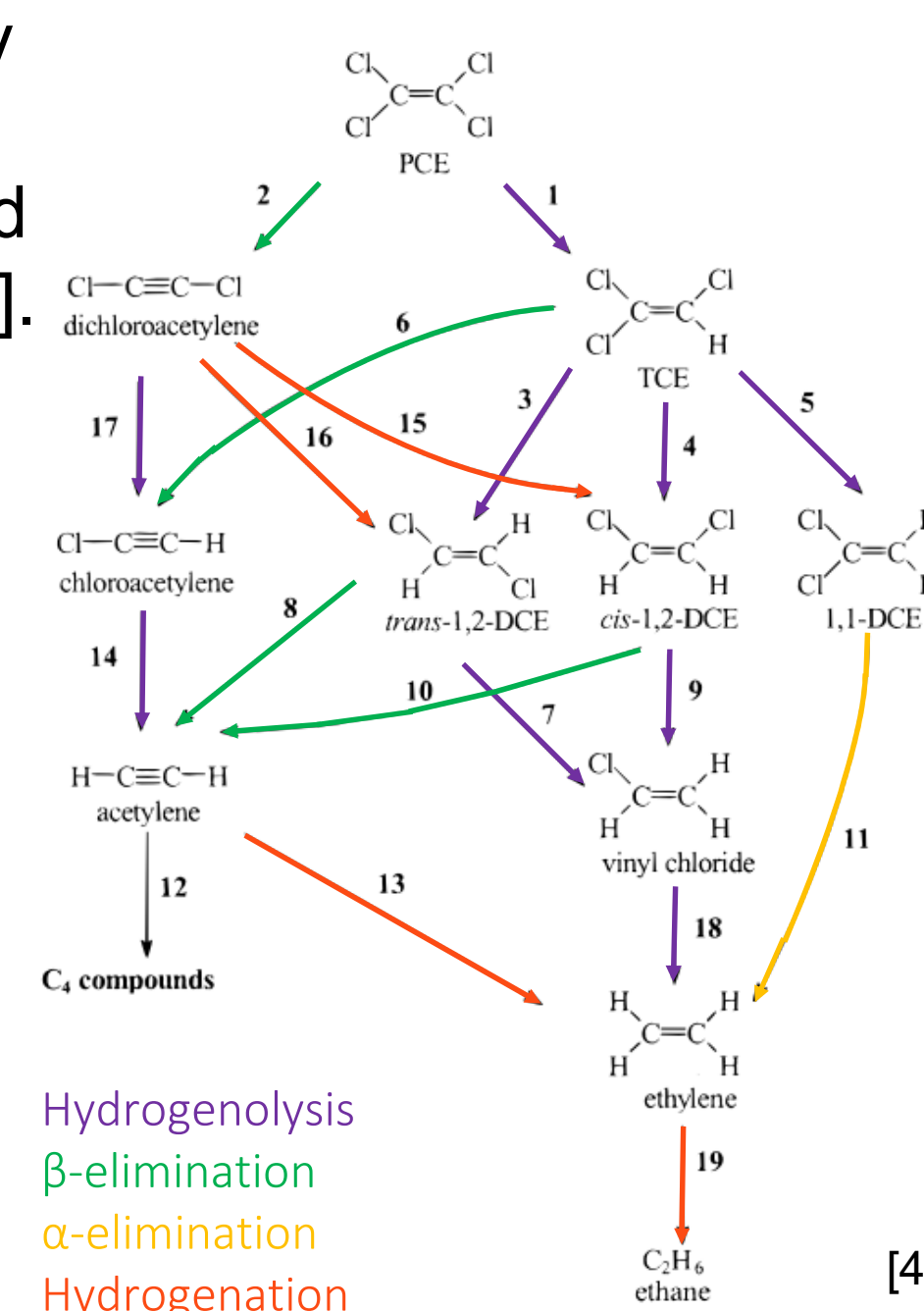
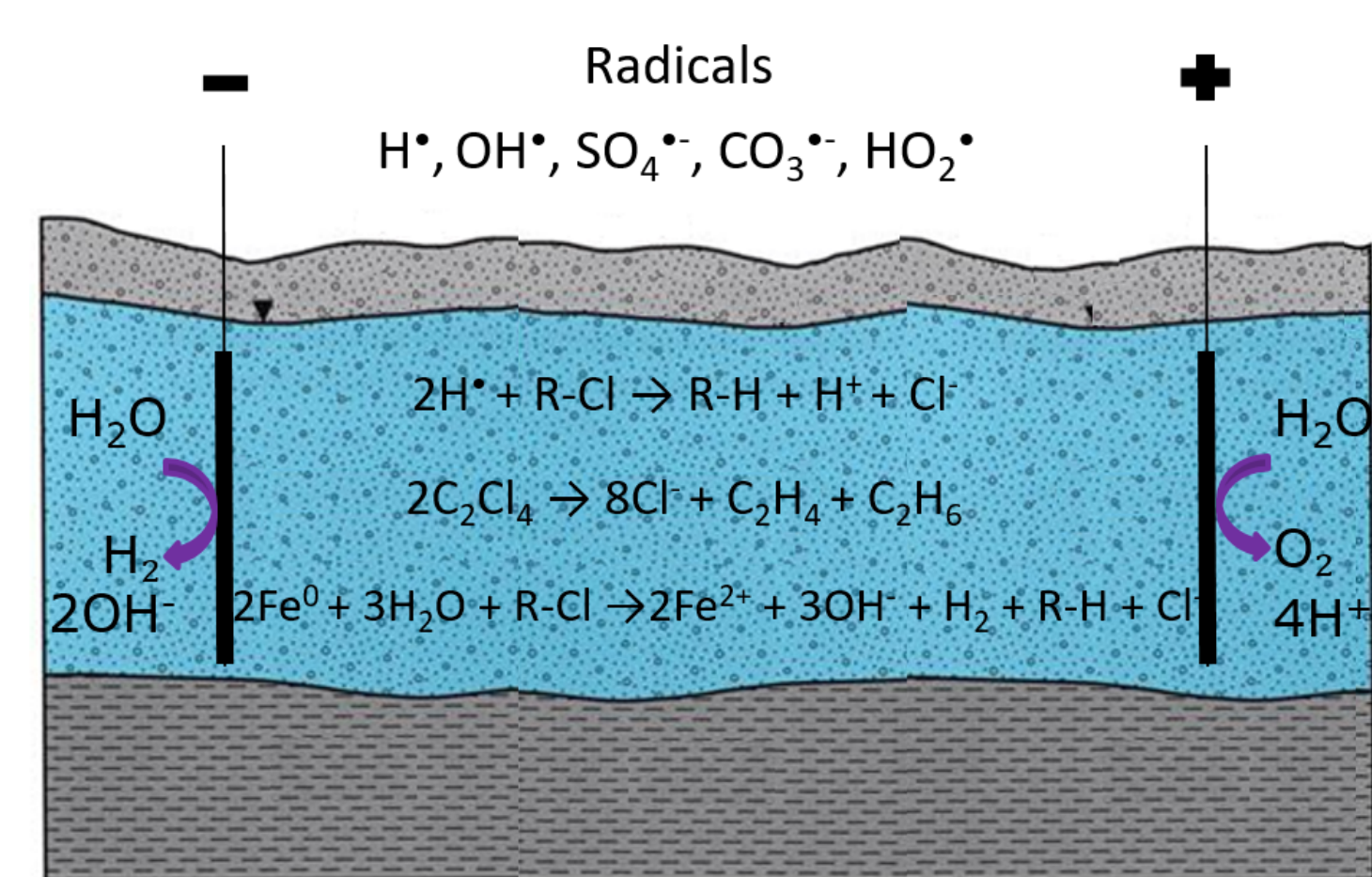
IX Acknowledgement

• This project and conference attendance is funded by



III Relevant chemical processes

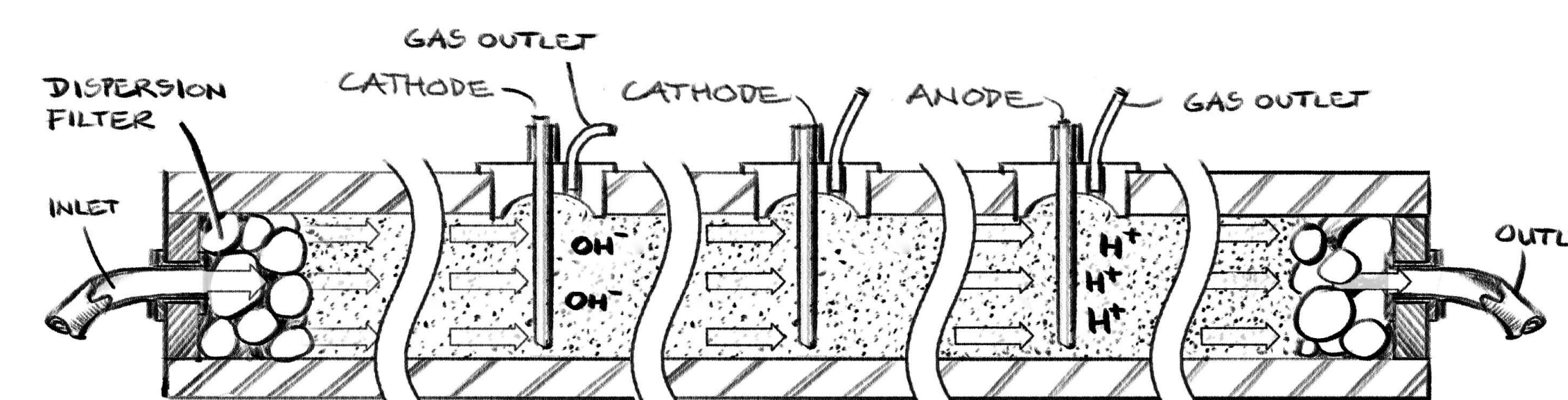
• Reactants can be generated and subsequently reduce or oxidize the chlorinated solvents [3] and fast electrochemical reduction of chlorinated solvents near the electrodes can be obtained [2].



V Method

• We have designed 1D and 2D experimental set-ups targeting electrochemical plume control in field realistic designs

- allows for assessment of single parameters; current density, flow and electrode material, and power consumption, lateral dispersion of reactants, electrode configuration and spacing
- replicates site conditions: Flow-through of natural groundwater with an aged contamination of PCE in a sandy aquifer material at common groundwater flow rates and temperatures



VII Challenges and opportunities

- Contaminant fate when no current is applied is unexpected; upon test completion, dissolved and gaseous fractions are low. When current is applied, these fractions are high in proportion.
- Alterations in redox conditions induce e.g. deposition and eventually clogging of the pore spaces in the geological matrix. One design solution may be short-term polarity reversal.
- Present geochemistry competes with dechlorination for electrons, e.g. reduction of carbon dioxide to methane followed by polymerization to ethane and oxidative dehydrogenation to ethylene [11]. These hydrocarbons may interfere with the resulting mass balances of dechlorination.
- Some precipitates formed may improve the abiotic dechlorination of chlorinated ethylenes, e.g. magnetite and green rust [12].
- The reduced conditions in the proximity of cathodes visually appear to enhance microbial growth. Dependent on the microbial culture, biodegradation of the chlorinated ethylenes may establish [13].

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