

Integrated Biogeochemical / Electrochemical Method for Remediation of Contaminated Groundwater

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Presentation Outline

◆ Problem Statement

- Overview of Available ISCO Technologies
- Why Do We Need Another?

◆ What is Provect-"EBR^{®"?}

- What is Provect-EBR?
- How does it Work / Mode of Action?
- Remote System Control and Real-Time Monitoring
- Applications to Date

◆ Case Studies

- CHCs: Confidential Site (Tel Aviv, Israel)
- MTBE: Operating Gas Station, Sonol Kiryon Site
- MTBE/BTEX: Operating Gas Station, Neve Tdizek Site
- Summary and Conclusions / R&D Needs

ISCO = Breaking Chemical Bonds

- ◆ Oxidant must be able to accept electrons
	- Capacity = Equivalent weight (MW / No. electrons)
- **◆ Ultimate end point is mineralization**
	- Partial oxidation is common

Summary of ISCO Technologies

the hydroxyl **influence**

https://sites.google.com/site/ecpreparation/ferrate-vi

- •**Treats limited range of contaminants**
- **Partial oxidation of TPHs, etc**
- **Long subsurface lifetime**

Fenton's

• **Potential effects on hydrogeology**

Reactive Oxidant Species (ROS) Higher oxidation potential = stronger the oxidizer

Why We Need A New ISCO Technology

- **Longevity**: Conventional ISCO amendments and means of generating ROS are limited by distribution, kinetics, and short environmental half-lives (10E-9 to $10E^{-6}$ seconds) = need to be continuously generated / applied.
- **ISCO PRBs:** PRB applications using existing ISCO (candles, KPS, etc) are limited
- **Sustained,** *In Situ* **Production of ROS could yield effective PRBs** especially for COIs not conducive to ISCR/ZVI such as 1,4-dioxane, MTBE/TBA, perchlorate, (PFAS?) plumes.

APPENDIX A. Comparative Analysis of Various Options for an Example PRB @ 50 m long x 5 m deep (4 to 9 m bgs) x 3 m wide.

Provect- "EBR®" ISCO PRB

In Situ ISCO Generator to continuously produce Fenton's type ROS yields an effective PRB technology for:

- Challenging lithologies (deep aquifers, clayey soils, fractured rock)
- Situations where sorption/sequestration is not considered an effective response
- Alternatives to hydraulic containment (long term O&M&M)

> 100 ppm 50 ppm 10 ppm 1 ppm 0.5 ppm 0.1 ppm < 0.05 ppm source plume down gradient EZVI ISCR ISCO OR ISCR EBR® ISCO

Contaminant Concentration

What is Provect-"EBR®"

Electro Bioremediation (EBR) well(s) contain an air sparge plus 3 electrodes:

- \bullet H₂O₂ production
- \triangle Fe²⁺ release
- \bullet O₂ production

Computerized controller Electrolytic cells for O, production Reactor Electrolytic cells for H₂O₂ production Electrolytic cells for Fe²⁺ production

Computerized control panel for remote system / adjustment and real-time performance monitoring

US Patent No. 9,975,156 B2

How Does EBR Work?

The EBR Well Generates Reactive Oxidant Species (ROS) in a manner similar to other Electro-Fenton's (EF) type systems (Nazari *et al*., 2019; Rosales, *et.al,* 2012; Sires *et al*., 2014; Yuan et al., 2013):

Production of O₂: electrolytic reduction of water on a catalytic electrode yields molecular oxygen, $O₂$

Production of H₂O₂. two-electron reduction of oxygen on a cathode surface generates H_2O_2

Release of Iron: H_2O_2 interacts with ferrous iron (Fe²⁺) released from a third cell to yield hydroperoxyl (HO₂∙)/superoxide (O₂⋅) and hydroxyl radicals (OH*∙*), and likely ferrates

$$
O_2 + 2H^+ + 2e^- \rightarrow H_2O_2
$$

Fe²⁺ + H₂O₂ \rightarrow Fe³⁺ + HO' + OH⁻
Fe³⁺ + H₂O₂ \rightarrow Fe²⁺ + HO₂ + H⁺
Fe³⁺ + HO₂⁺ \rightarrow Fe²⁺ + O₂ + H⁺

How Does EBR Differ From EF?

Fe2+/3+ Nanoclusters: At neutral pH EBR uniquely generates "low" Fermi Level (highly oxidized) FeII/III oxyhydroxide nanoclusters (2 nM) as the sacrificial Fe source corrodes within the well (Ai *et al.,* 2013; Elgressy 2019).

Subsurface distribution of Fe nanoclusters throughout the aquifer is driven by:

- Induced redox fronts
- Electro-redox current densities
- Electroosmosis
- **Electrophoresis**
- Dynamic coupling between EBR wells
- **Equilibration of differences in Fermi level energies self-generated self-propagated**

How Does EBR Differ From EF?

Fe2+/3+ Nanoclusters: A critical and unique feature of the EBR is use of geophysical mechanisms to enhance subsurface distribution of low Fermi level Fe nanoclusters and propagate catalysis *in situ* to continuously generate reactive oxidants throughout its effective ROI.

Electrochemical Potential of an e- is the difference in potential between the oxidized and reduced species (Peljo *et al.,* 2017; Scanlon *et al.,* 2015)

Fermi Level is a thermodynamic "value" to define the electrochemical potential of an electron in a redox couple in solution

At +850mV ("low" Fermi Level electrochemical potential) electrons are essentially freely transferred from $Fe³⁺$ to $Fe²⁺$

Scheme 3 Redox equilibria for metallic NPs in solution showing the capabilities of metallic NPs to be (A) charged and (B) discharged upon Fermi level equilibration with an excess of a single dominant redox couple in solution.

In Situ **Generation of ROS**

As Fe (hydro)oxides within the aquifer ROI equilibrate their Fermi level electrochemical potentials they continuously catalyze *in situ* generation of new ROS from dissolved molecular $O₂$ via two kinds of molecular oxygen activation pathways (Ai *et al*., 2013):

- On the Fe core via rapid two-electron-reduction molecular oxygen activation (may eventually be blocked by the formation of iron oxide coatings), then
- Surface bound ferrous ions catalyze the singleelectron-reduction molecular oxygen activation pathway

Summary of EBR Reactions

- Generation of H_2O_2
- Release of $Fe²⁺$
- H2O2 interacts Fe2+ to yield ROS HO2∙/O2∙ and OH*∙* (ferrate?)
- Release of $O₂$ and low Fermi Level Fe²⁺/Fe³⁺ nanoclusters
- Self-propagation throughout ROI (less confined by lithology)
- Continuous *in situ* production of ROS catalyzed by O₂ activation from equilibration of Fermi levels of Fe BEFLORESSY
	- **Transition from ISCO to** bioremediation (using oxygen and iron as electron acceptors) and RNA using abiotic transformations.
	- **Process controlled remotely with** real-time monitoring

Where has it been Used?

- ♦ In 2017, Israel had 27 gas stations undergoing active remediation
- \triangle EBR technology was employed at 9 (33%) + 2 chlorinated solvent sites
- Today, 7 sites are in clean-closure monitoring after 1 year of operation
- EBR is ISO-certified and approved by the Israeli Water Authority
- ◆ No PRB Applications. No USA applications.

Case Study - Solvent Site

◆ DTW 19.5 m bgs

- **◆ Sandy aquifer impacts**
	- ◆ PCE max. 257 ug/L
	- ◆ TCE max. 25,146 ug/L
	- ◆ DCE max. 47 ug/L

 $W-C$

CVOC Removal (60 days; ppb)

Single EBR Well + Control Panel and remote monitoring < \$45K installed

- ROI observed 20 m downgradient within 30 to 60 days.
- ◆ >99% CVOC removal within 30 days

Case Study – Neve Tzedik Site

◆ Operating Gasoline Station

- ◆ Groundwater at 7 to 8 m bgs
- ♦ sandy aquifer with si cl lenses
- MTBE >50 mg/L; TPH >100 mg/L
- \triangle 242 m² impacted area

Monitoring wells

MTBE Concentration (mg/L) in Water

 \bullet MW-6 ca. 130 ppm to \lt 5 ppb within 12 months

BTEX Concentrations (mg/L) in Water

 \bullet MW-5 ca. 25 ppm to \lt 5 ppb within 12 months

Dissolved Oxygen (DO)

GW field parameters (ORP)

Soil / Groundwater BTEX (18 mo)

Soil / Groundwater MTBE (18 mo)

- From >50 ppm to < 0.05 ppm
- 5 EBR Wells, Control Panel, O&M < \$150K

Case Study – Sonol Kiryon Site

◆ Operating Gasoline Station

- ◆ Groundwater at 3 m bgs
- **◆ sandy aquifer**
- \triangle MTBE >17 mg/L
- \bullet ca. 300 m² impacted area

Case Study – Sonol Kiryon Site

Groundwater MTBE (5 months)

GSI MANN-KENDALL TOOLKIT

for Constituent Trend Analysis

5 months EBR operation (as of December, 2018)

Groundwater COIs ug/L (7 months)

- Toluene, Ethylbenzene <5 ppb
- 6 EBR Wells, Control Panel, O&M < \$180K

Learn More About EBR MOA

Provectus Environmental Products

- **Complimentary Site Evaluation**
- **Complimentary review of quarterly field performance data with every project**
- **Laboratory Treatability Studies**
- **Turn-Key, Pay-for-Performance Contracting Options**
- **Project Specific Guarantees and Warranties**

- **USA (Florida, Illinois, New Jersey, Ohio, Pennsylvania, Wisconsin)**
- **Australia, Brazil, Canada, China, Colombia, Germany, Israel, Italy, Spain and Taiwan**

Future R&D / Continued Studies

Validate ROI and Effective Propagation Time, Vertically and Horizontally (ESTCP submittal Mueller, Shi, Ginn, and Tratnyek 2019)

- **ORP / Measurements (indirect)**
- **↓ COI Reductions (indirect)**
- Fe2+/Fe3+ measurements: Particle size (BEM) and mineralogy (XRD patterns, TEM micrographs, XPS spectra and high-resolution scan); possible using variations of Bradley and Tratnyek (2019).
- ♦ Self-Potential Method (direct): passive geophysical analysis based on the natural occurrence of electrical fields resulting from the existence of source currents in the conductive subsurface (Fachin *et al*., 2012)
- ♦ Electrical Resistivity Tomography (direct): measures variations in electrical conductivity associated with changes in pore water ionic strength or water phase saturation.
- Lab-fabricated oxygen microprobes/sensors (direct): validate the distribution of ROS.
- Simple and Predictive Models: facilitate PRB design and implementation