

Geochemical behaviour of adit drainage under saturated and unsaturated conditions at two underground mines in Northern British Columbia

John Dockrey & Alison Shaw

22nd annual BC MEND ML/ARD workshop December 2-4, 2015



Overview

- 1. Two geologically distinct mine sites with similar mining histories
- 2. Similarities in geochemical trends
 - Sulphate concentrations increase when mine workings are unsaturated
 - Metal cation (Zn and Cd) concentrations decrease when mine workings are unsaturated
- 3. Identify mechanisms that could result in decline in metal cation solubility between saturated and unsaturated time periods
- 4. Results show that metal cation behaviour is controlled by pH related solubility constraints in neutral mine drainage.



Site Naming Conventions







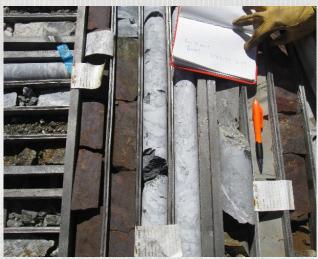
TUNNEL OF LOVE

Lorax Environmental Services Ltd.

www.lorax.ca







Site Background: Geology

Site A

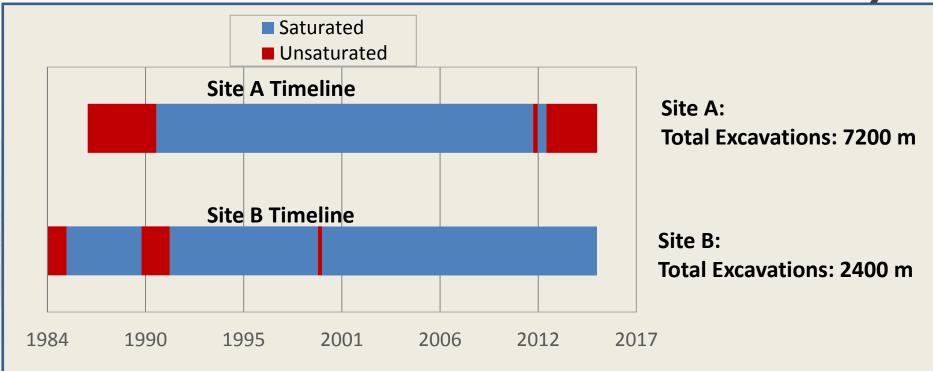
- Epithermal Au-Ag deposit hosted in stockwork veining intruding a primarily volcanic host rock.
- Existing excavations are primarily in hydrothermallyaltered PAG rock consisting of volcanic and volcaniclastic sedimentary rock units

Site B: aka the tunnel of love

- Ag-Pb-Zn carbonate replacement deposit hosted in a limestone formation
- Existing excavations are primarily in non-PAG limestone with minor amounts of mudstone and ore excavated as part of bulk sampling program.



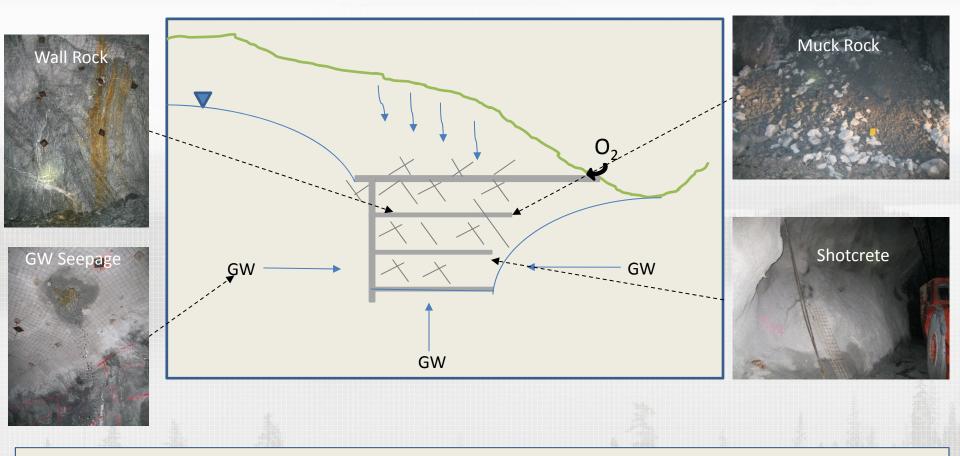
Site Background: Mining History



- Nearly all mine workings are below adit elevation in both sites
- No backfilled waste rock or tailings are present at either site



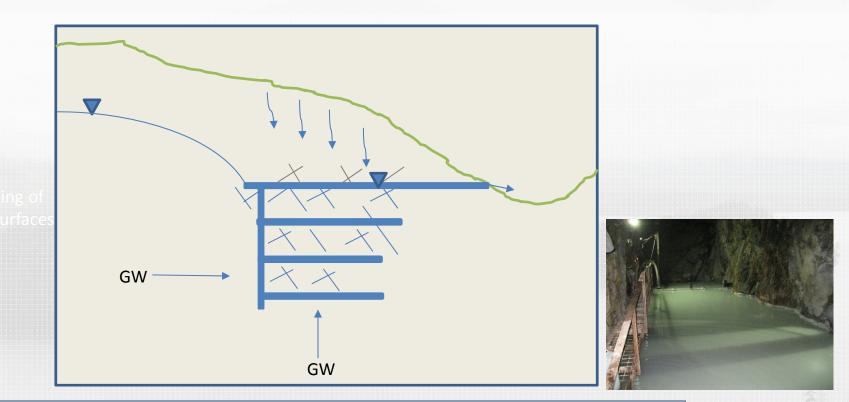
Conceptual Model of Geochemical Loading: Unsaturated



Unsaturated Geochemical Load = Wall Rock Oxidation + Rinsing of Freshly Exposed Surfaces + Additives + Groundwater



Conceptual Model of Geochemical Loading: Saturated

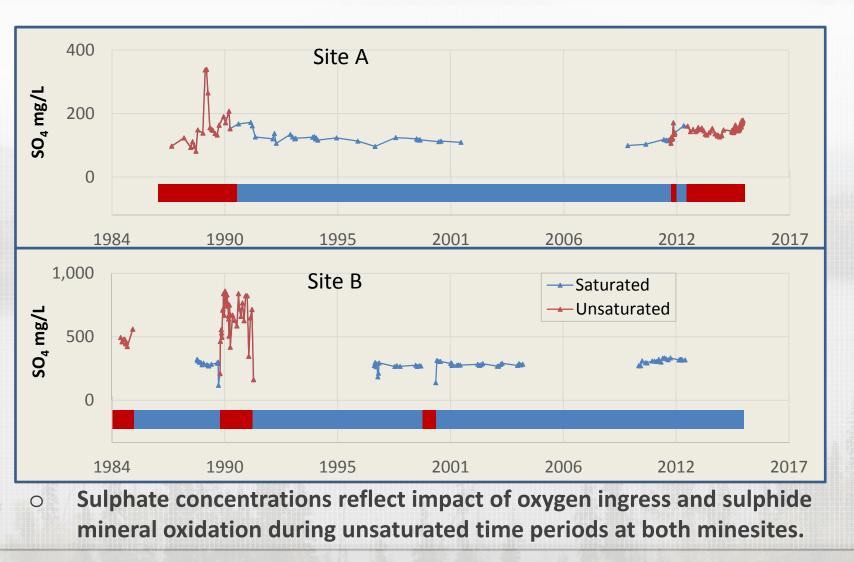


Short-Term Geochemical Load = Flushing Load + Groundwater

Long-Term Geochemical Load = Groundwater

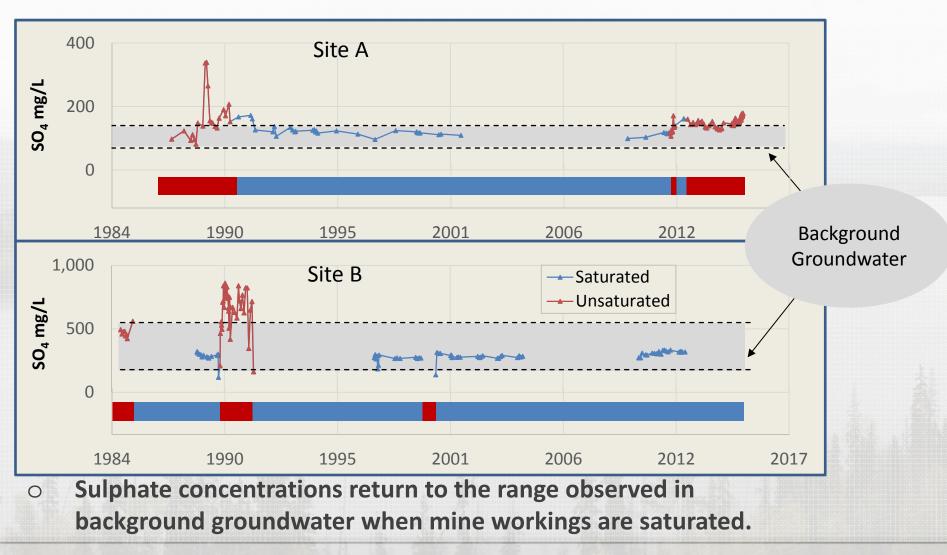


Sulphate Results





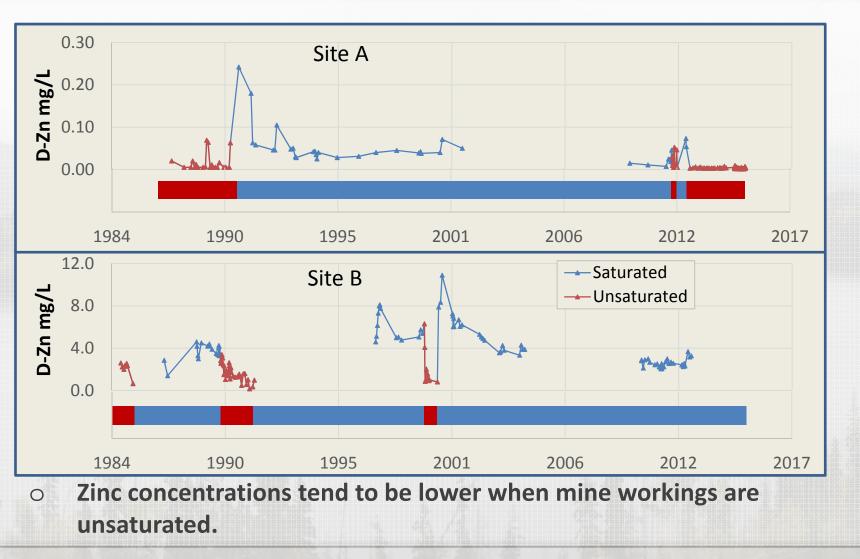
Sulphate Results



Lorax Environmental Services Ltd.

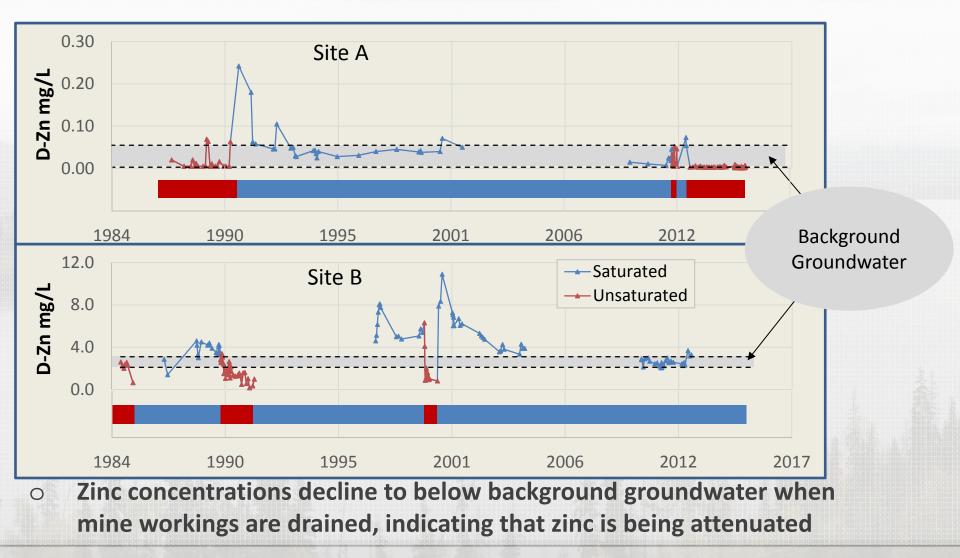


Zinc Results





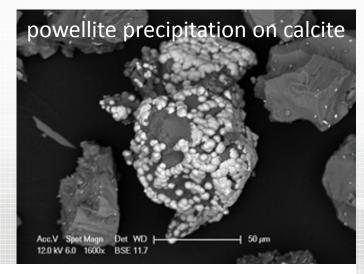
Zinc Results

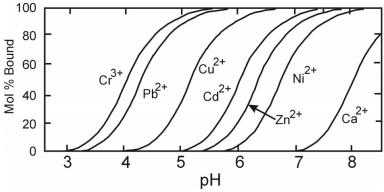




Potential Attenuation Mechanisms

- 1. Secondary Mineral Precipitation
 - o solution chemistry
 - equilibrium concentrations can be calculated
 - thermodynamic data only available for end-members
 - potential rate limitations
- 2. Sorption to Particle Surfaces
 - o solution chemistry
 - availability of surfaces
 - mineralogy of surfaces
 - must be empirically determined





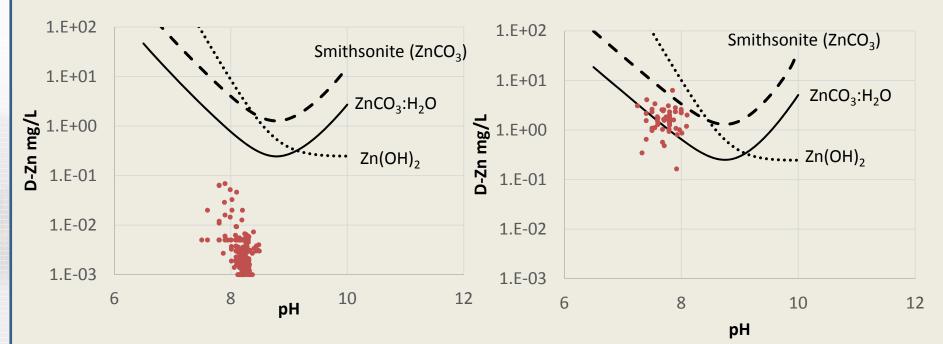
The pH Dependence of Metal Sorption on Fe-oxide (After Stumm, 1992)



Potential Attenuation Mechanisms

Site **B**

Site A



- Site A: sorption controlled
- Site B: secondary mineral control



Factors influencing sorption and solubility during active mining

- 1. Elevated TSS: increased particle surfaces available for sorption
- 2. Cement in additives: dissolution of CaO
- 3. Cation exchange: increase in carbonate mineral equilibrium pH
- 4. Aeration of mine water: leading to degassing of carbon dioxide



Elevated TSS

- During active mining TSS is observed to increase at both minesites
- Blasting and ground disturbance releases and suspends silt and clay particles





Cement in Additives

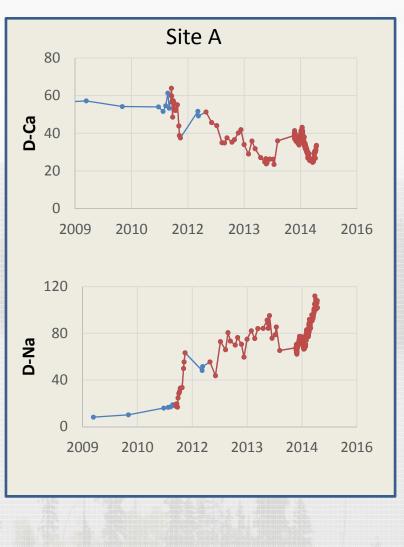
Cement Dissolution: CaO + H₂O \rightarrow 2OH⁻ + Ca²⁺

• Dissolution of cement in grout or shotcrete can drive up pH

Lorax Environmental Services Ltd.

www.lorax.ca





Cation Exchange

Cation Exchange:

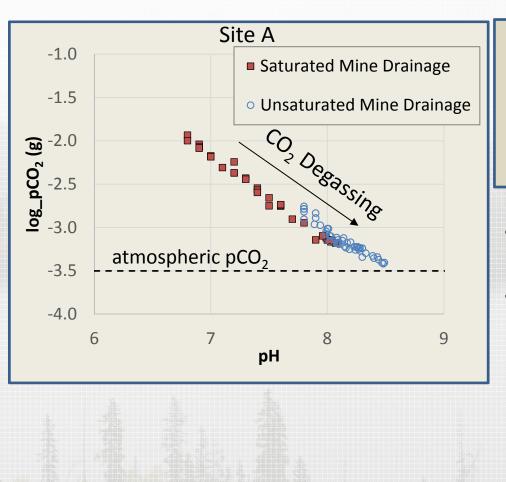
 $Ca^{2+} + Na_2:Ex \rightarrow 2Na^+ + Ca:Ex$

 $2CaCO_3 + 2H_2O \rightarrow 2HCO_3^- + 2OH^- + Ca^{2+}$

- Water quality trends provide evidence that Na:Ca exchange reactions are occurring and potentially driving up pH at site A.
- Similar trend observed in early humidity cell data from freshly crushed drill core samples



Aeration of Mine Drainage

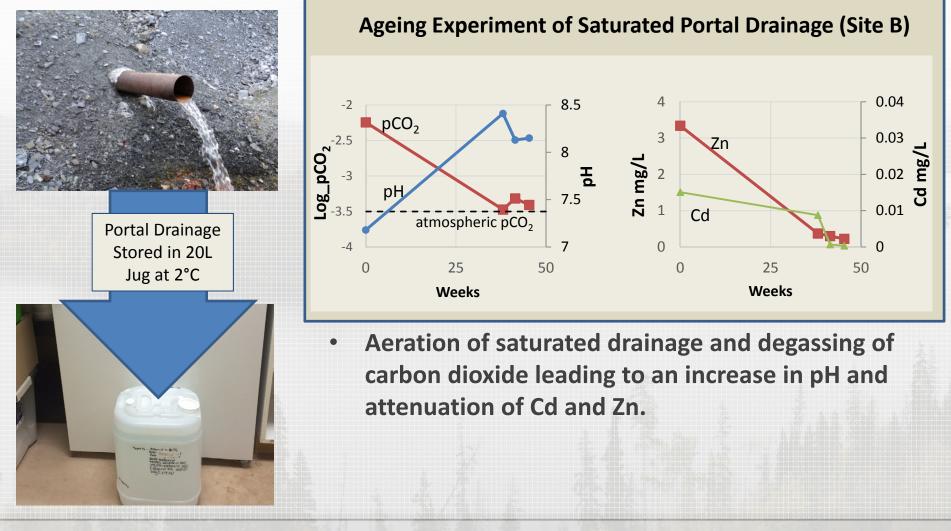


Carbon Dioxide Degassing: (1) $CO_3^{2-} + H_2O = HCO_3^{-} + OH^{-}$ (2) $HCO_3^{-} + H_2O = H_2CO_3 + OH^{-}$ (3) $H_2CO_3 = H_2O + CO_{2(g)}$

- A number of reactions are caused by mine water aeration
- Mine drainage at unsaturated time points is closer to atmospheric CO₂ equilibrium, indicating that degassing has occurred.



Aeration of Mine Drainage





Conclusions

- Geology is not the dominant control of metal cation behaviour.
- Solubility controls dictate metal cation behaviour in neutral pH settings. Solubility decreases when mine workings are drained due to:
 - Elevated TSS
 - Dissolution of cement
 - Cation exchange reactions
 - Aeration
- Interpreting neutral mine drainage datasets and predicting future water quality must take into account geochemical reactions which are highly sensitive to pH in addition to the intrinsic metal leaching potential of mine rock.



Questions?

