

# Development of an integrated conceptual model of a connected surface water-groundwater system using a hydrochemical approach at Maules Creek, NSW, Australia.

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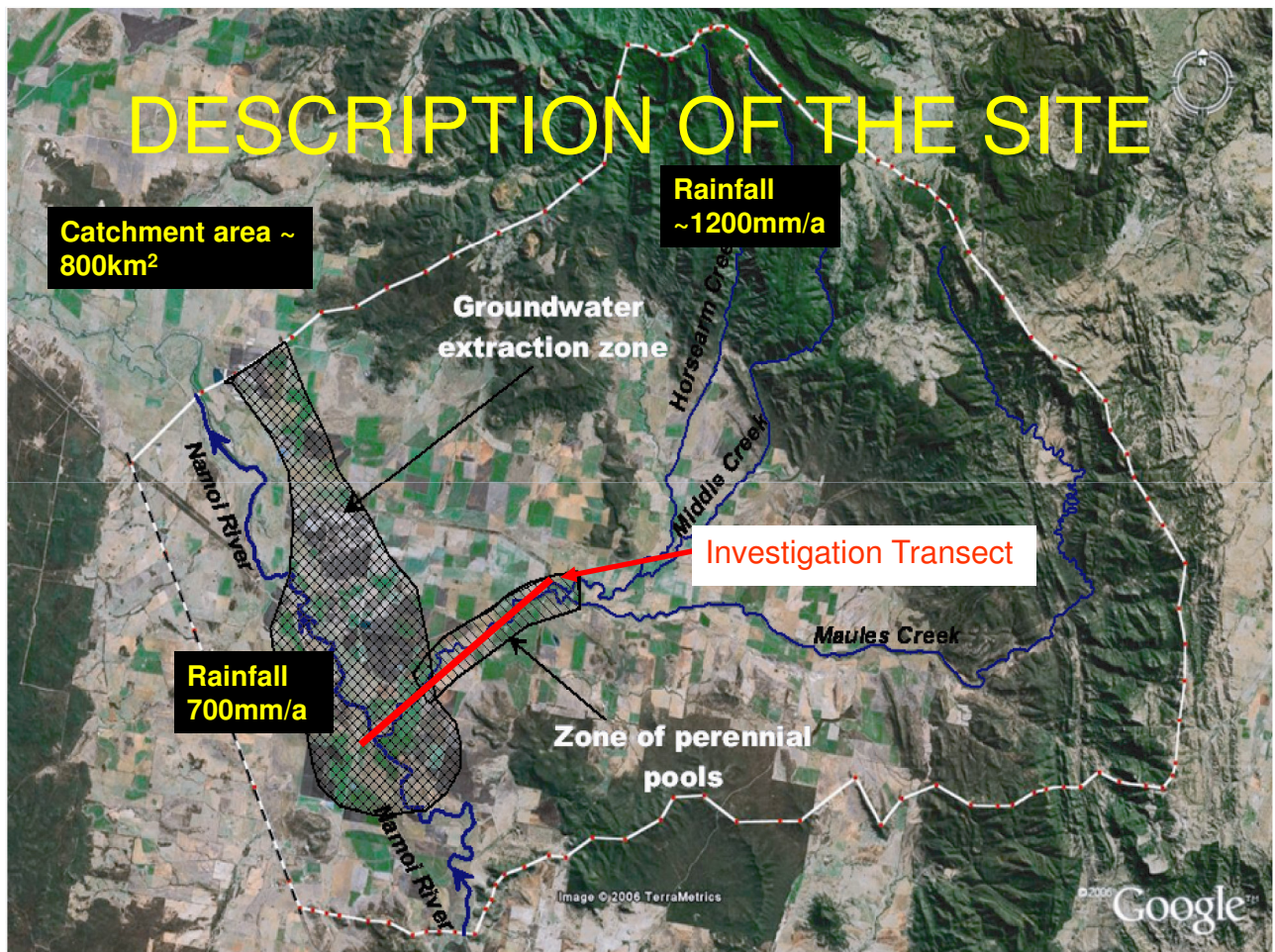


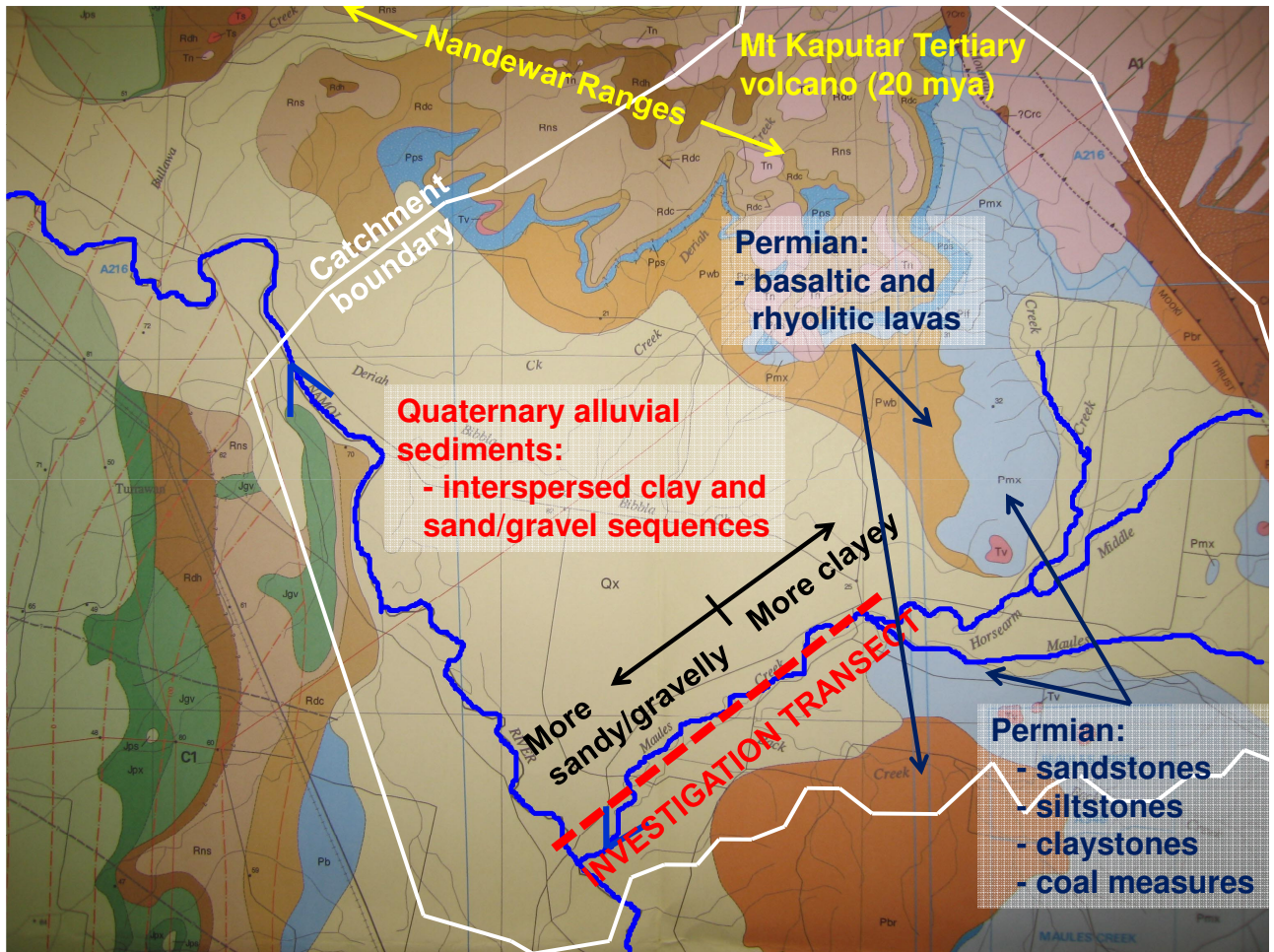
## INTRODUCTION

- Surface water-groundwater interactions important for water allocations
  - Anthropogenic needs
  - Environmental needs
- Separate allocation of surface water and groundwater resources can produce problems in connected systems
  - Reduced baseflow in streams
    - Groundwater dependant ecosystems (aquatic, terrestrial, phreatic)
  - Reduced surface water yield
- Challenge of linking SW-GW interactions into more typical hydrogeological investigations

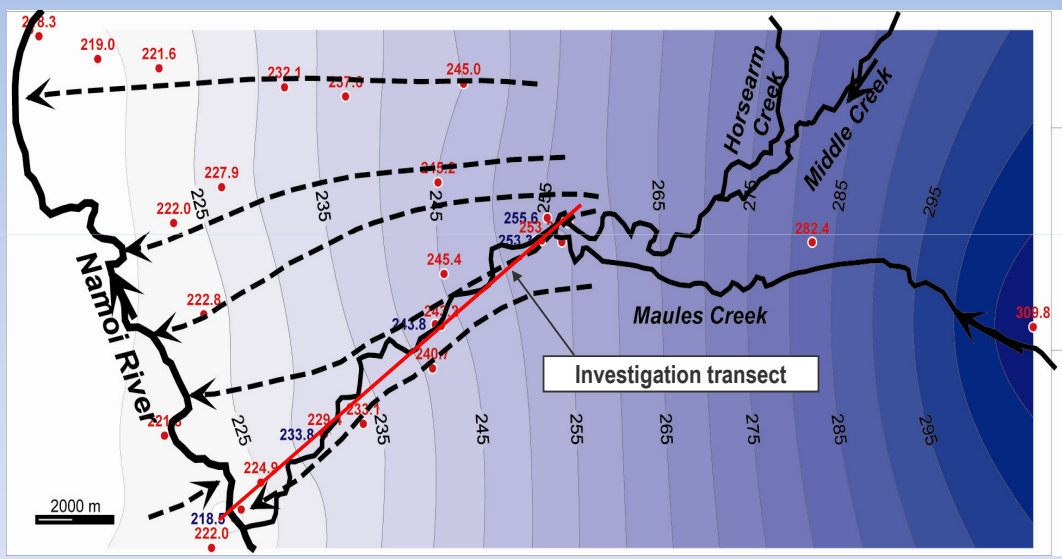
# INTRODUCTION

- Need to integrate SW-GW exchange and aquifer processes
- Hydrochemical approach used
  - Major ions ( $\text{Na}^+$ ,  $\text{Ca}^{2+}$ ,  $\text{Mg}^{2+}$ ,  $\text{K}^+$ ,  $\text{Cl}^-$ ,  $\text{HCO}_3^-$ ,  $\text{H}_4\text{SiO}_4$ ) and
  - Water characteristic parameters (pH, dissolved oxygen, dissolved  $\text{CO}_2$ , dissolved organic carbon)
  - used as **natural tracers** and to characterise **chemical processes** in the aquifer system
- Developed process-based conceptual model: hydrochemical and hydrogeological processes

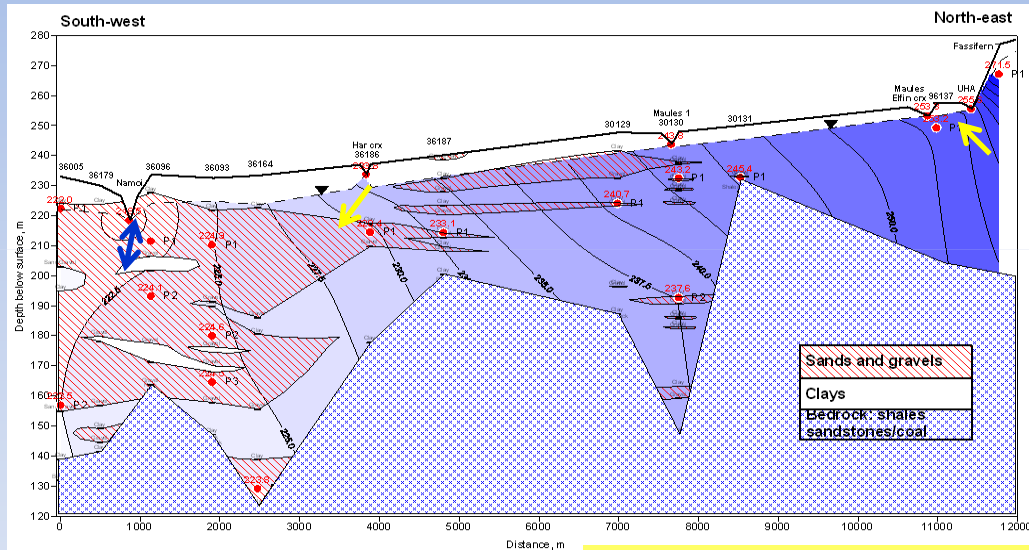




# HEAD DISTRIBUTION AND GROUNDWATER FLOW



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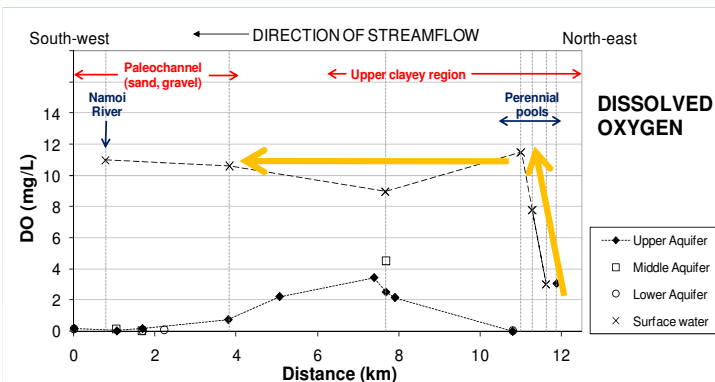
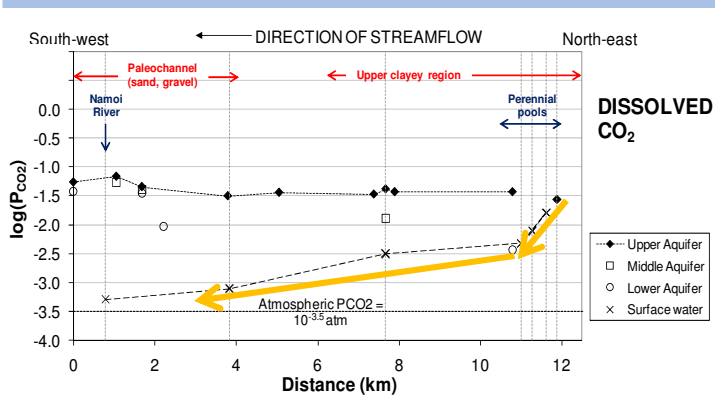
## Defined:

Upper aquifer: <30m

Middle aquifer: 30-60m

Lower aquifer: >60m

# RESULTS: PCO<sub>2</sub> and DO

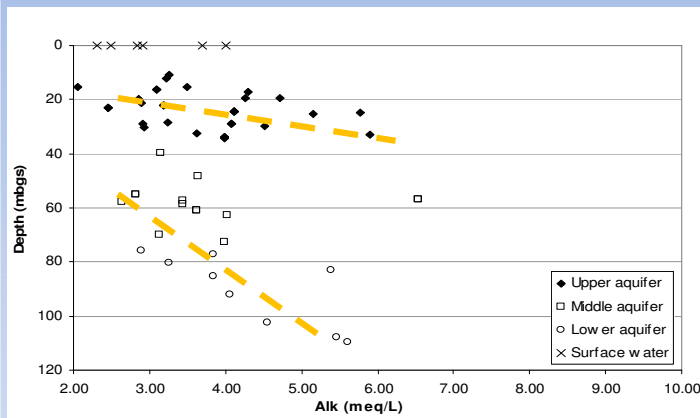


- Dissolved CO<sub>2</sub>:
  - Relatively constant in Upper aquifer
  - Decreases in Middle and Lower aquifer
  - Surface water trend

- Dissolved oxygen:
  - Oxic to anoxic along transect
  - Low DO in upstream surface water

➤ GW discharge to SW upstream of perennial pools

# RESULTS: $\text{HCO}_3^-$ with depth

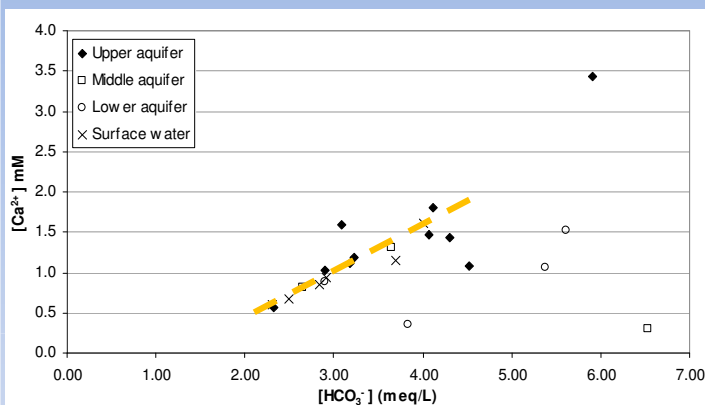


- Relatively linear increase in  $\text{HCO}_3^-$  with depth in Middle and Lower aquifers

- $\text{HCO}_3^-$  variable in Upper aquifer

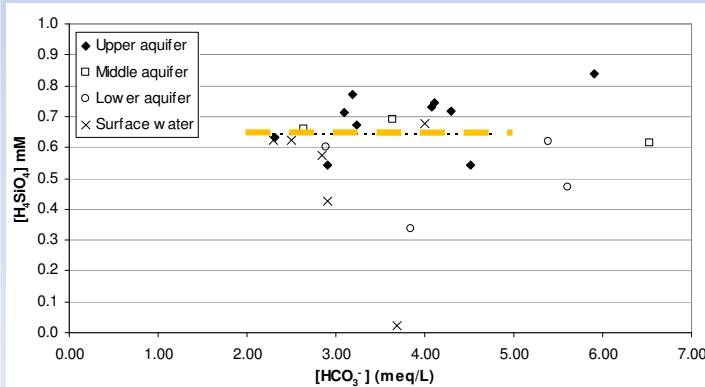
➤ Return flow of irrigation water sourced from aquifer

# RESULTS: $\text{Ca}^{2+}$ and dissolved silica



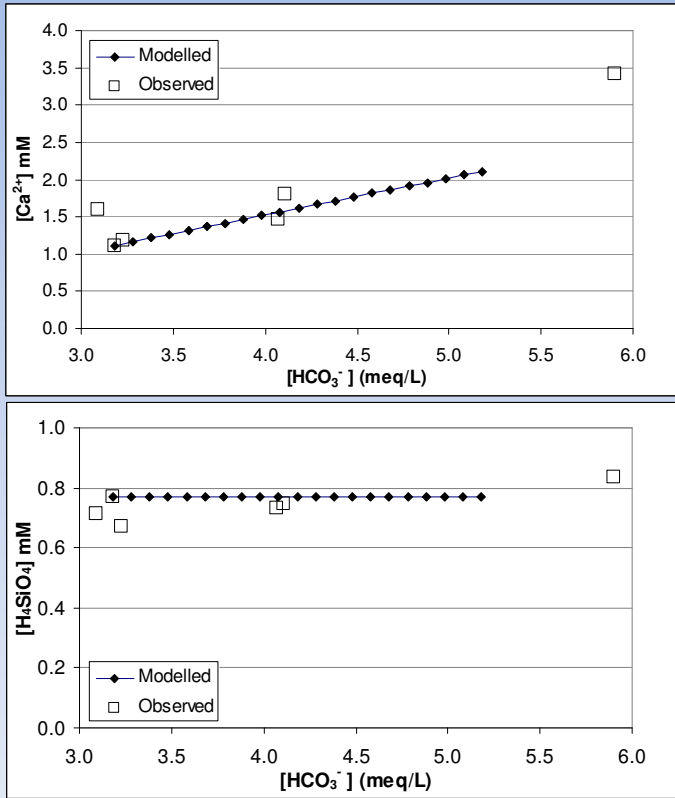
- Ca acquired relatively linearly against  $\text{HCO}_3^-$

- Dissolved silica relatively high (17mg/L), constant, and supersaturated with respect to some silica oxide phases



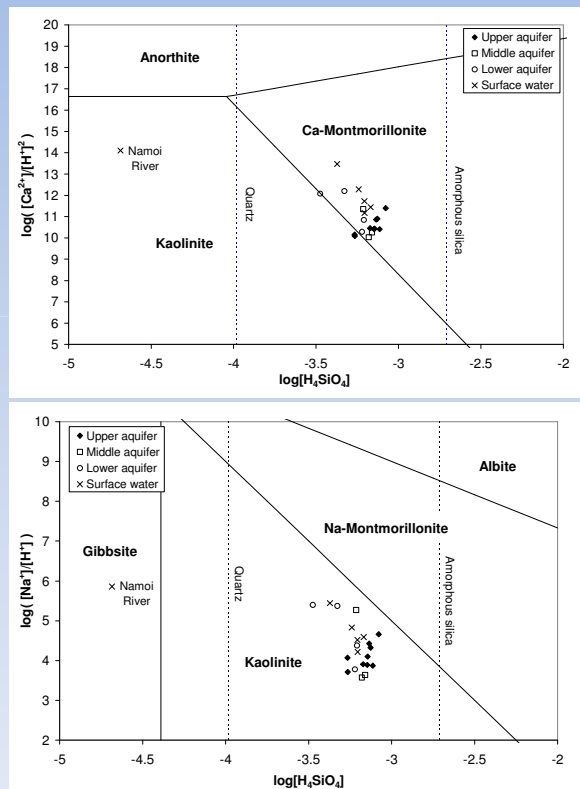
➤ Appears that substantial primary silicate weathering is occurring and releasing cations to solution

# RESULTS: PHREEQC



- PHREEQC Batch reaction model to test hypothesis of primary silicate weathering
  - Upper aquifer water
  - Dissolve 1 mM Anorthite (an idealised pure-phase Ca-plagioclase feldspar)
  - **Look at general trends – are they reasonable?**
- Appears plausible process
- Silica conserved in solid phase with Kaolinite precipitation indicated

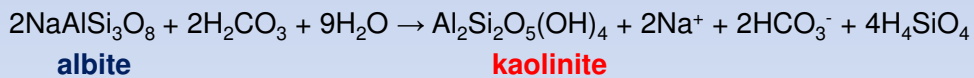
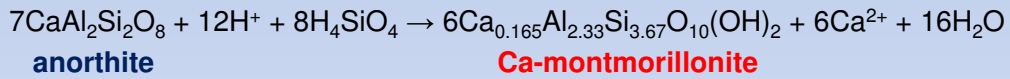
# RESULTS: STABILITY DIAGRAMS



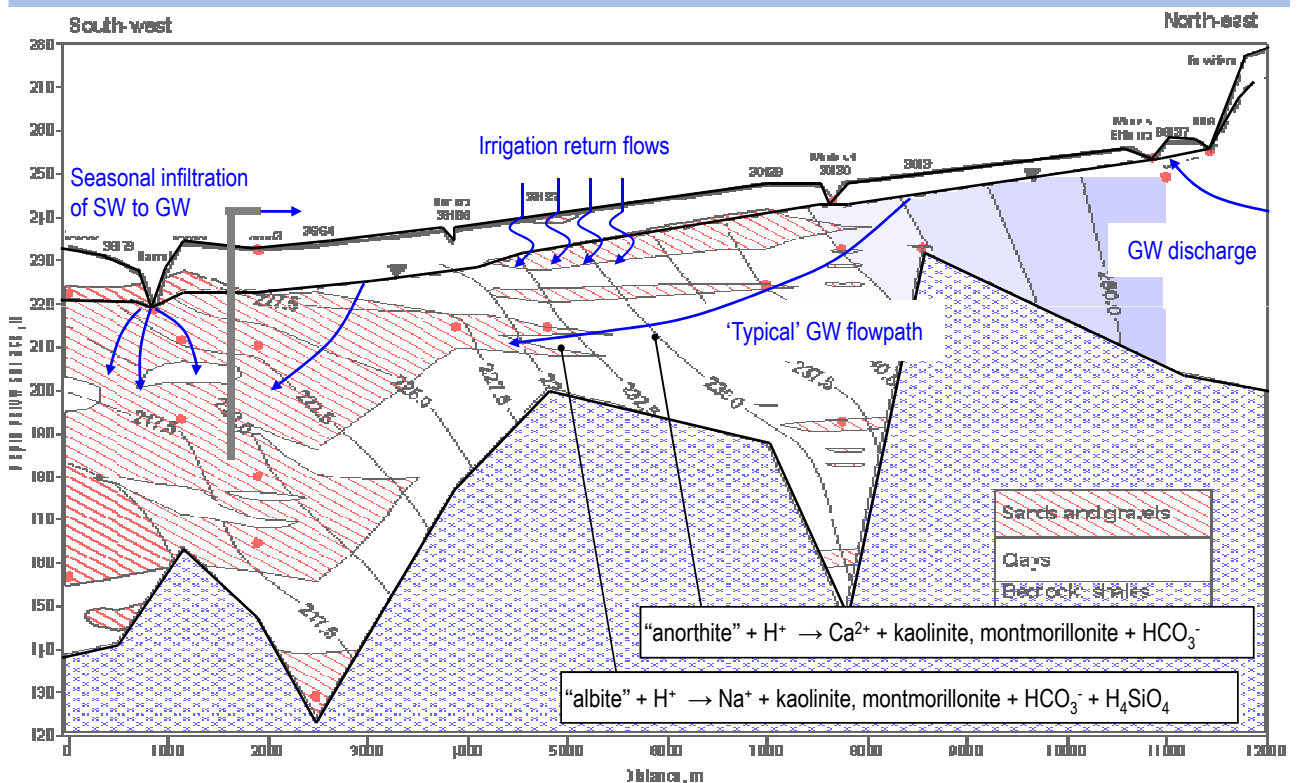
- Secondary weathering products important from process-perspective
- Stability diagrams very idealised, give indication only
- Montmorillonite and kaolinite weathering products (vs PHREEQC model indicated kaolinite)
- Likely to be heterogeneous mixture of co-existing secondary products

# GEOCHEMICAL PROCESSES

- Weathering of primary silicate reactions important eg the pure-phase Ca- and Na-plagioclases were idealised cases considered:



# CONCEPTUAL MODEL



## CONCLUSIONS

- Some key geochemical processes deduced
- Linked with previous investigations into SW-GW interactions and strengthened knowledge of hydrogeological processes
- Presented results as a conceptual model that integrates key hydrochemical and hydrogeological processes
- Useful for future aquifer system modelling

## ACKNOWLEDGEMENTS

- CCC-CRC for funding
- Department of Natural Resources access to sample monitoring wells and for supplying basic data from their archive



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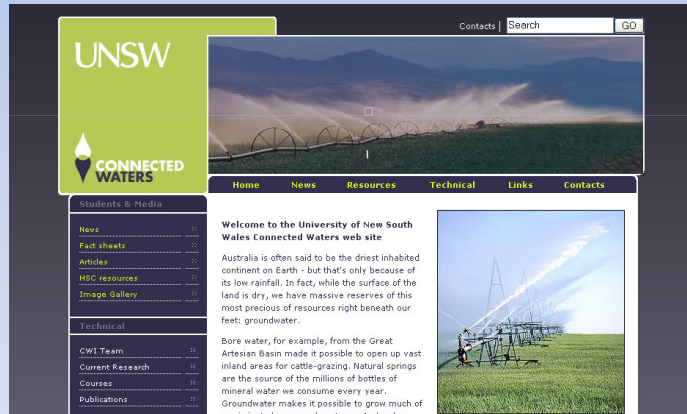
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