Phosphorus sequestration by oxidizing iron in groundwater fed lowland catchments

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# Oxidation of Fe-rich groundwater → Fe oxyhydroxide particles JUST AFTER SAMPLING 2 DAYS LATER

Reduced Fe(II) (soluble)





### $Fe^{2+} + 3H_2O \rightarrow Fe(OH)_3 + 3H^+ + e^-$

- Oxidation rate at circumneutral pH: hours...days
- In the presence of PO<sub>4</sub>
  - First, Fe hydroxyphosphate formation until near depletion of PO<sub>4</sub>
  - Then, continued formation of Fe oxyhydroxides

#### The P bound to Fe oxyhydroxides is poorly available to algae



Algal growth and P uptake is reduced by adding Fe oxyhydroxide colloids. After time, part of the P slowly becomes available.

## What if Fe-rich groundwater surfaces across a large area?

- Vac 8453,222

### STUDY AREA

- Groundwater fed lowland catchments
- Flat topography, shallow phreatic aquifers
- Underground: permeable marine deposits, some rich in glauconite
- Average shallow groundwater composition: 20 mg Fe/L; 0.4 mg P/L
  - Fe source: glauconite weathering
  - P sources: geogenic, leaching from agricultural land



300 km<sup>2</sup>





The Fe(II) is oxidized and removed from solution (<0.45  $\mu$ m) as the groundwater surfaces and flows through the catchment into increasingly larger streams



Before the water reaches the headwater streams, 90% of the P is removed from the <0.45  $\mu m$  fraction

### Drainage ditches with Fe-rich sediments seasonally evacuate groundwater





### What happens as Fe and P bearing groundwater flows through Ferich sediments and into drainage ditches?



- As groundwater surfaces and flows through the catchment, the Fe is oxidized and Fe oxyhydroxide particles and colloids are produced.
- These particles strongly bind P, and thereby remove it from solution
- This study area is extreme in terms of its low P:Fe ratio
  - to what extent does this process occur in other lowlands?

