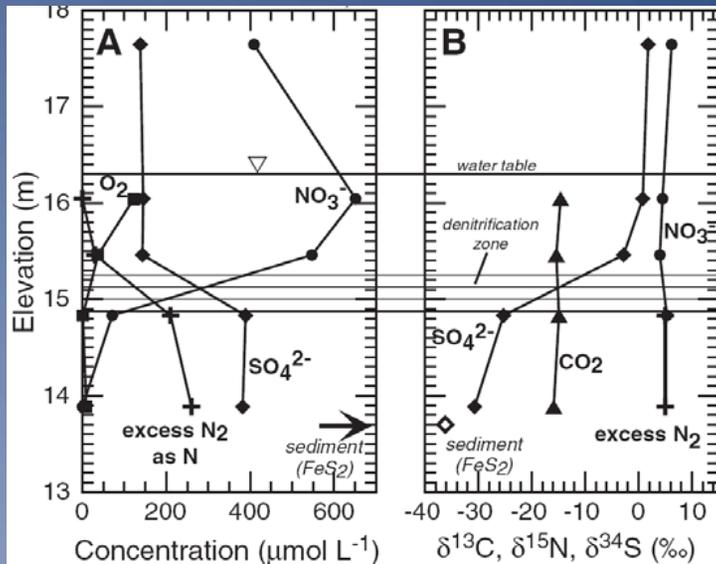


# **Redox Processes and Water Quality of Selected Principal Aquifer Systems: Implications for Ground-Water Monitoring**

**Gary Rowe  
for  
Peter McMahon and Frank Chapelle**

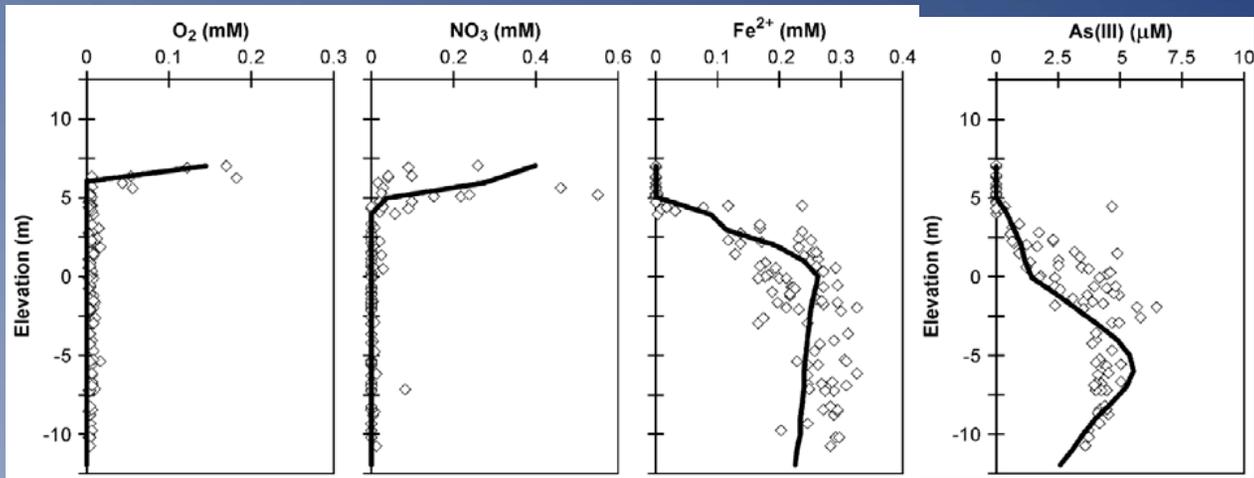
**National Water Quality Assessment Program  
U.S. Geological Survey**

# Redox Processes at Small Scales (we know a lot!)

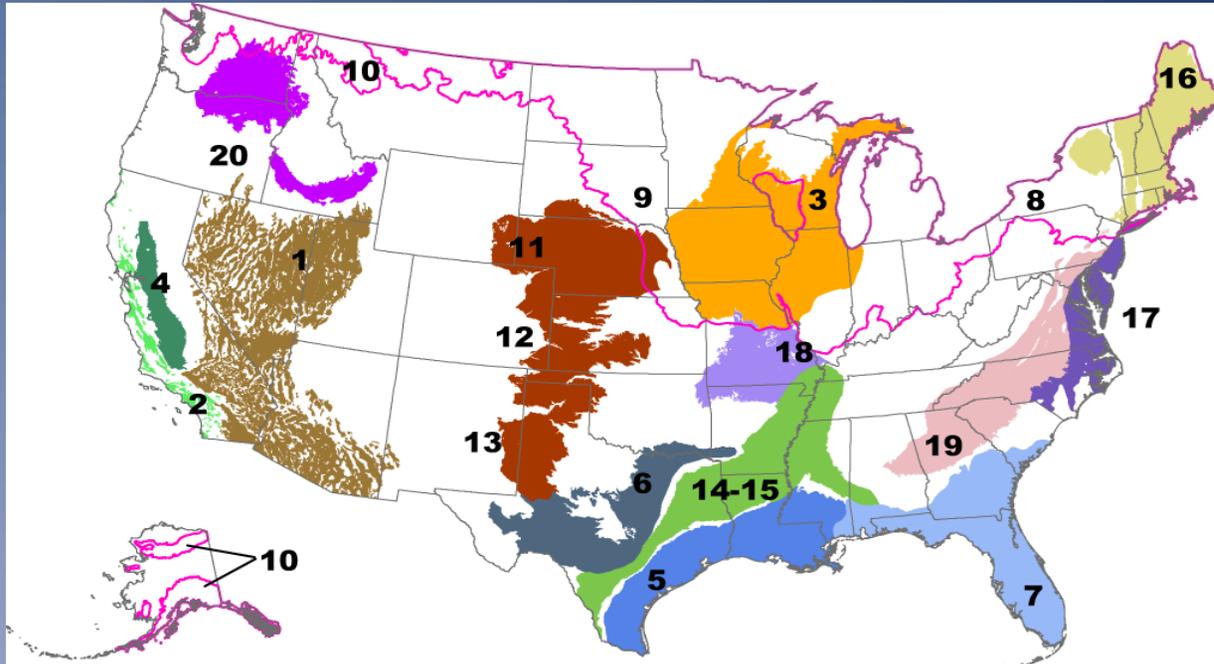


Nitrate  
(Böhlke et al., 2007, JEQ)

Arsenic  
(Postma et al., 2007, GCA)



# Redox Processes at Large Scales (there's a lot we don't know!)



## EXPLANATION

### Principal aquifers and reference number

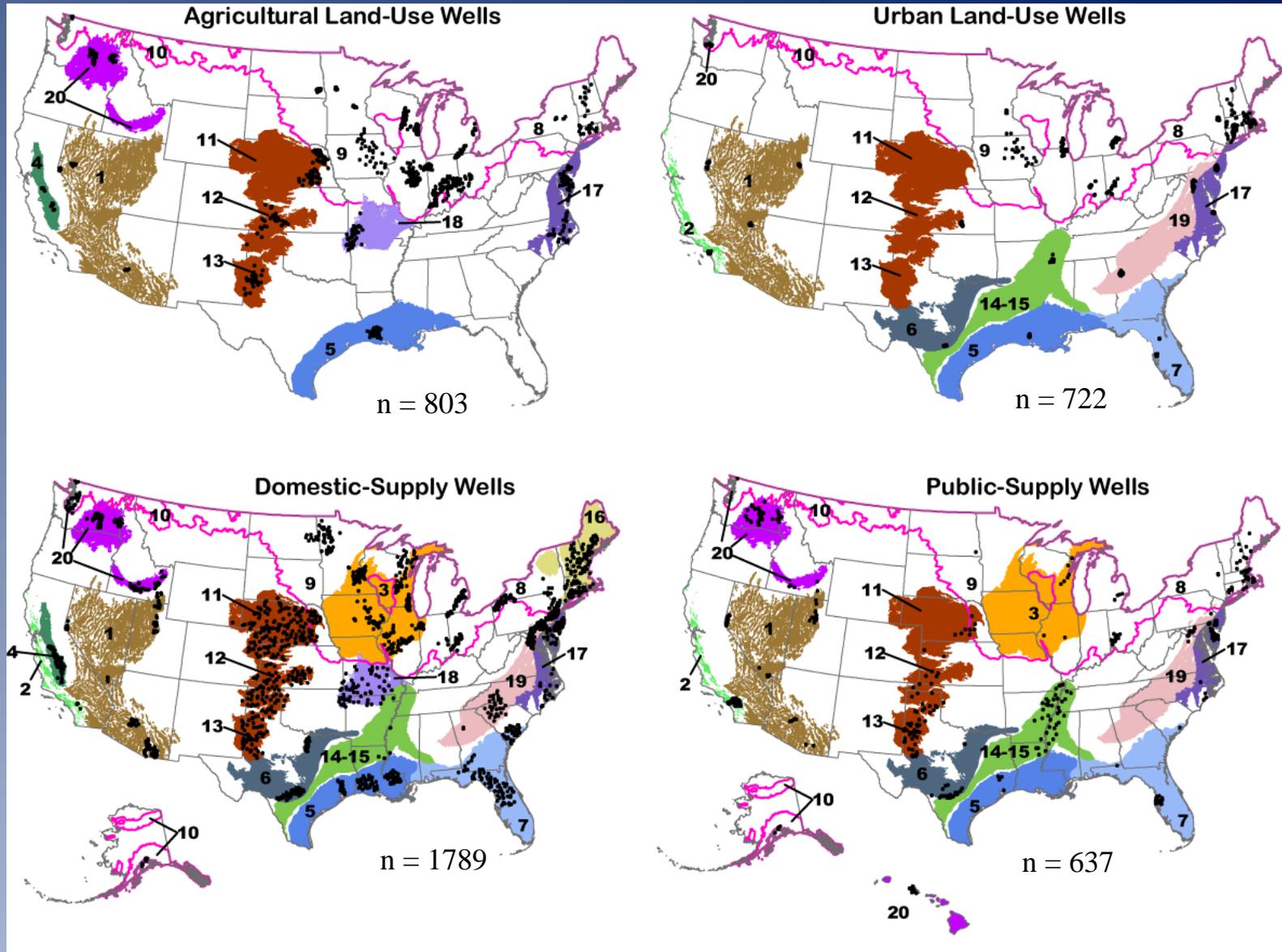
- Basin and Range (1)
- California Coastal (2)
- Cambrian-Ordovician (3)
- Central Valley (4)
- Coastal Lowlands (5)
- Edwards-Trinity (6)
- Floridan (7)
- Extent of Glacial deposits
- Glacial deposits - east (8)
- Glacial deposits - central (9)
- Glacial deposits - west (10)
- High Plains - northern (11)
- High Plains - central (12)
- High Plains - southern (13)
- Mississippi Embayment - Texas Coastal Uplands (14-15)
- New York and New England crystalline (16)
- Northern Atlantic Coastal Plain (17)
- Ozark Plateau (18)
- Piedmont and Blue Ridge crystalline and carbonates (19)
- Western Volcanics (20)

- How does redox vary between PAs?
- Lack of consistent data across states/regions (e.g. variable reporting levels, different field & lab techniques)
- Lack of “complete” data at large scale (O<sub>2</sub> data is hard to come by!)

# Objectives

- Use NAWQA datasets to characterize ambient redox conditions in selected Principal Aquifers
- Examine water-quality issues (nitrate, arsenic, VOCs) in the redox framework

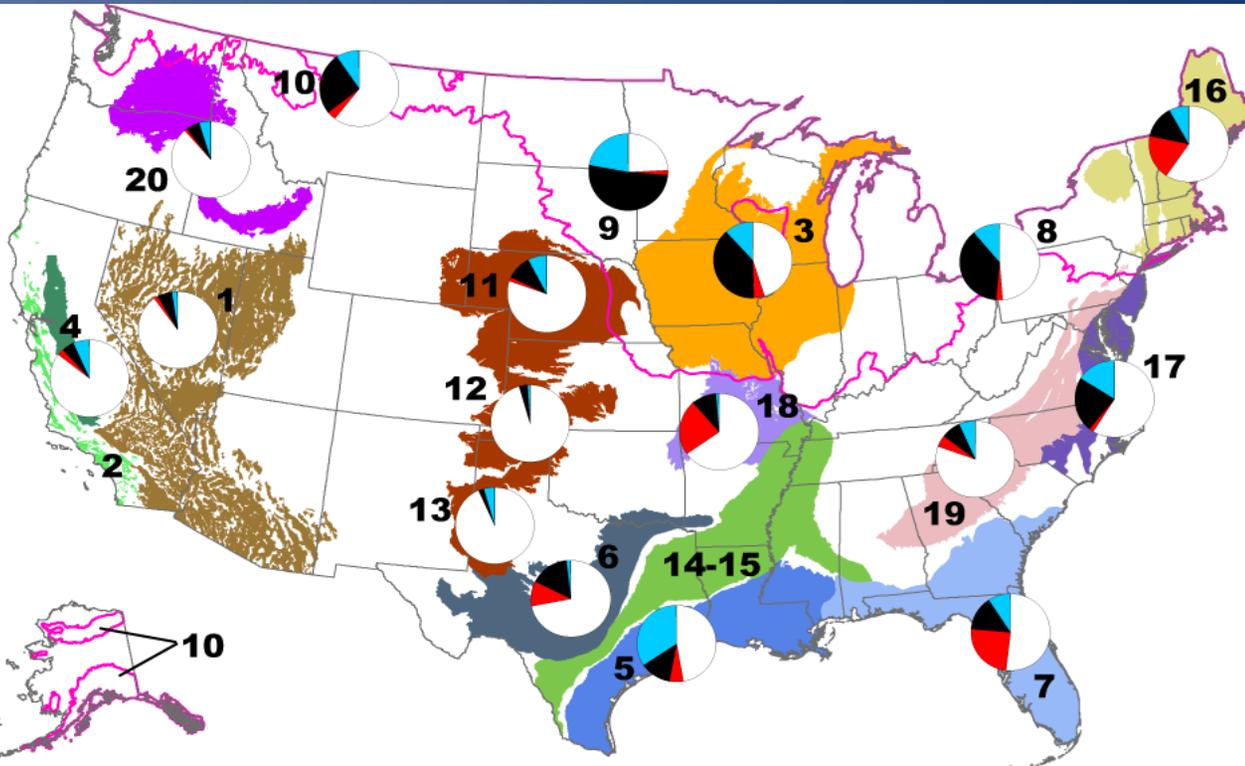
# Distribution of Sampled NAWQA Wells



# Framework for Identifying Redox Processes

Redox Process	Water-Quality Criteria (mg/L)				
	O <sub>2</sub>	NO <sub>3</sub> <sup>-</sup> -N	Mn <sup>2+</sup>	Fe <sup>2+</sup>	SO <sub>4</sub> <sup>2-</sup>
<b>Oxic</b>					
O <sub>2</sub> reduction	≥ 0.5	–	< 0.05	< 0.1	–
<b>Suboxic</b>					
–	< 0.5	< 0.5	< 0.05	< 0.1	–
<b>Anoxic</b>					
NO <sub>3</sub> <sup>-</sup> reduction	< 0.5	≥ 0.5	< 0.05	< 0.1	–
Mn(IV) reduction	< 0.5	< 0.5	≥ 0.05	< 0.1	–
Fe(III)/SO <sub>4</sub> <sup>2-</sup> reduction	< 0.5	< 0.5	–	≥ 0.1	≥ 0.5
Methanogenesis	< 0.5	< 0.5	–	≥ 0.1	< 0.5
<b>Mixed</b>	Criteria for more than one redox process are met.				

# Spatial patterns indicate common controls on ambient redox conditions at national/PA scale

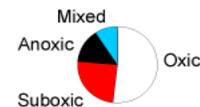


## EXPLANATION

### Principal aquifers and reference number

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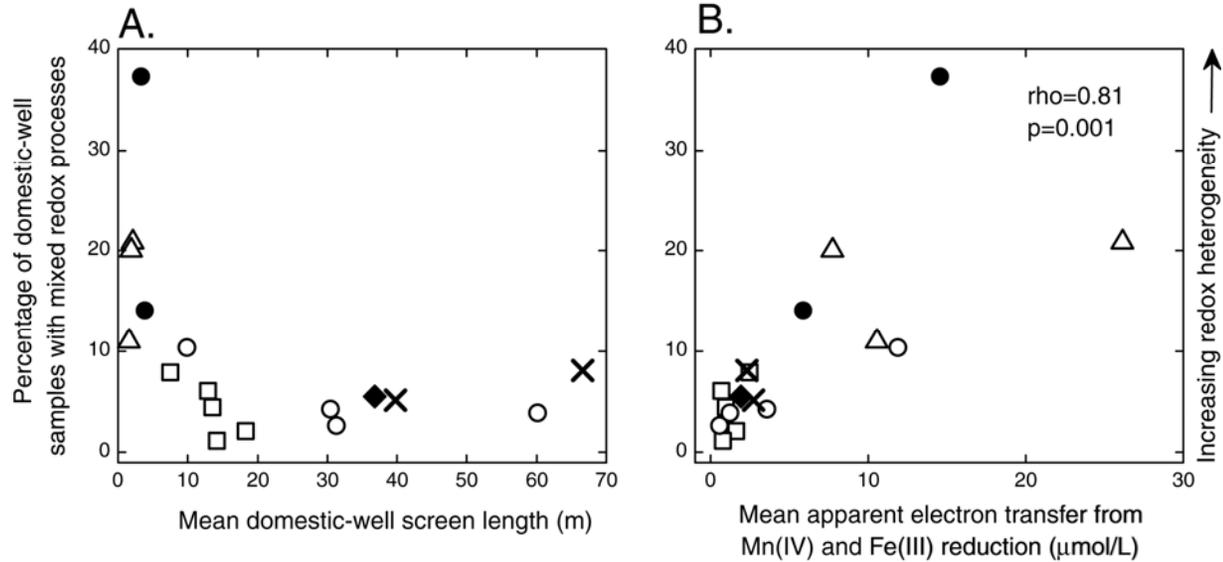
### Redox state of aquifer, as percentage of samples



Based on domestic-well data (n=1692)

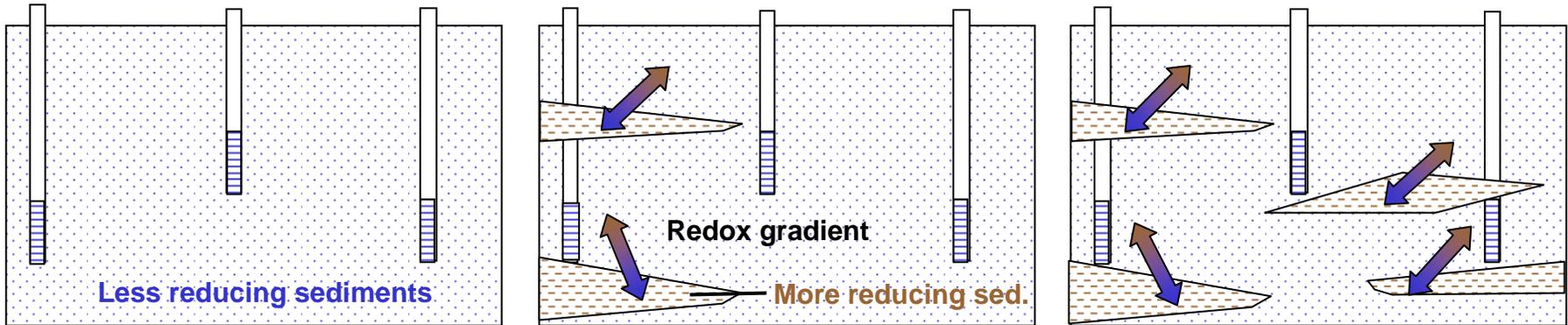
Well networks not representative of entire PAs

# Mixed Redox GW Samples

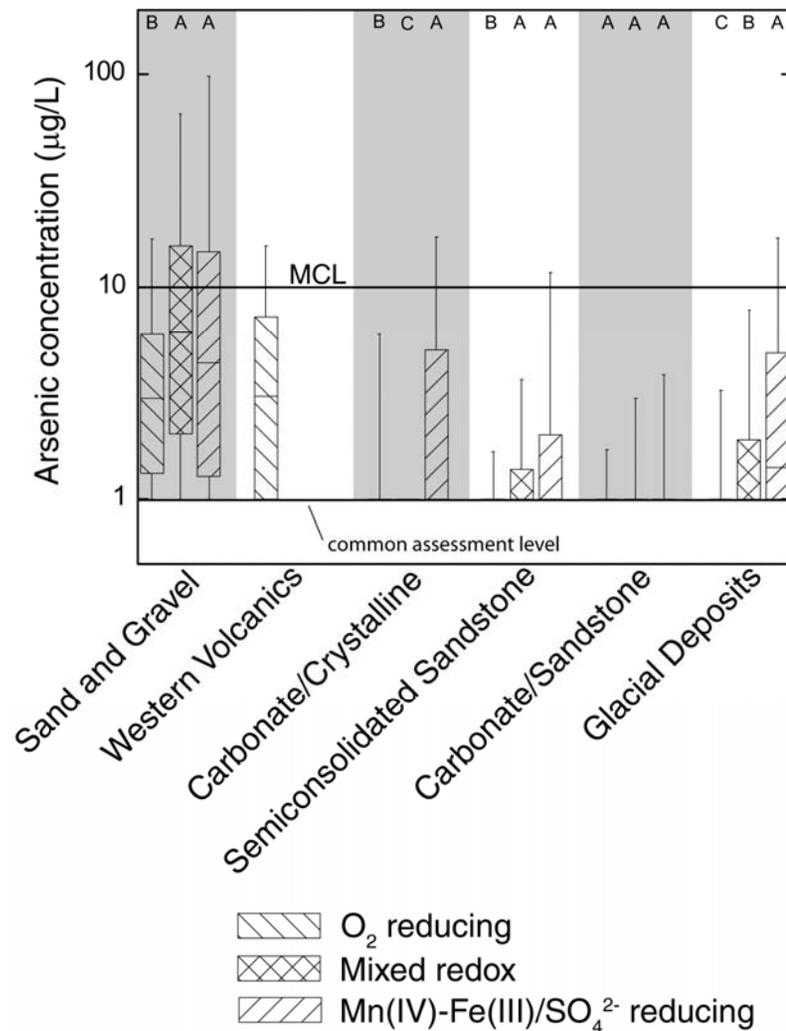
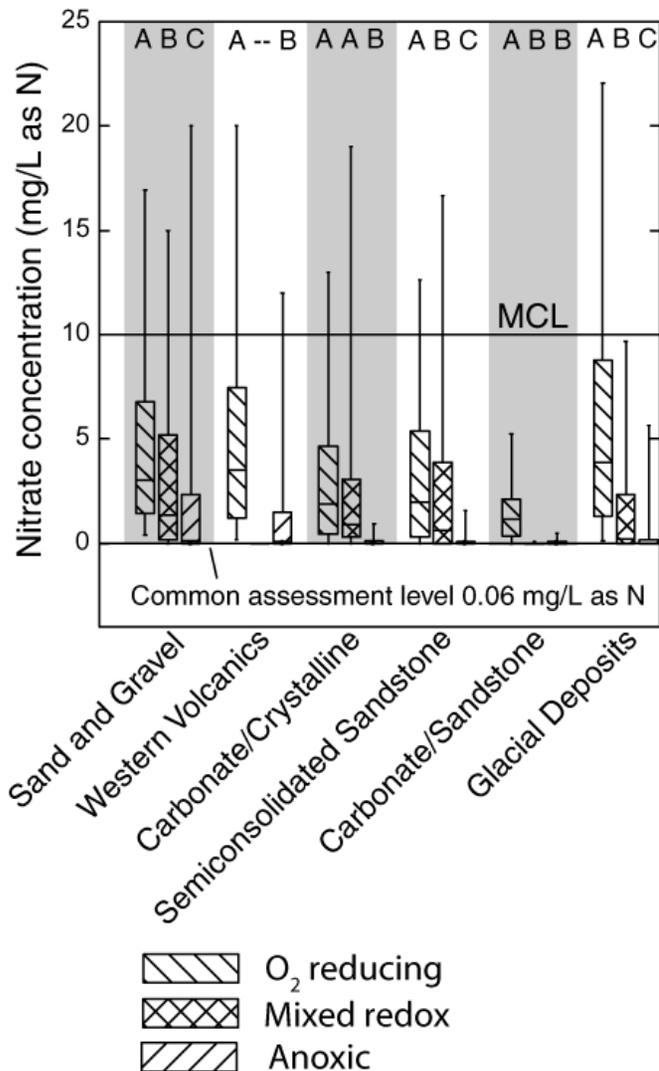


Mixed redox diagnoses may reflect inherent spatial heterogeneity of redox processes in the aquifers

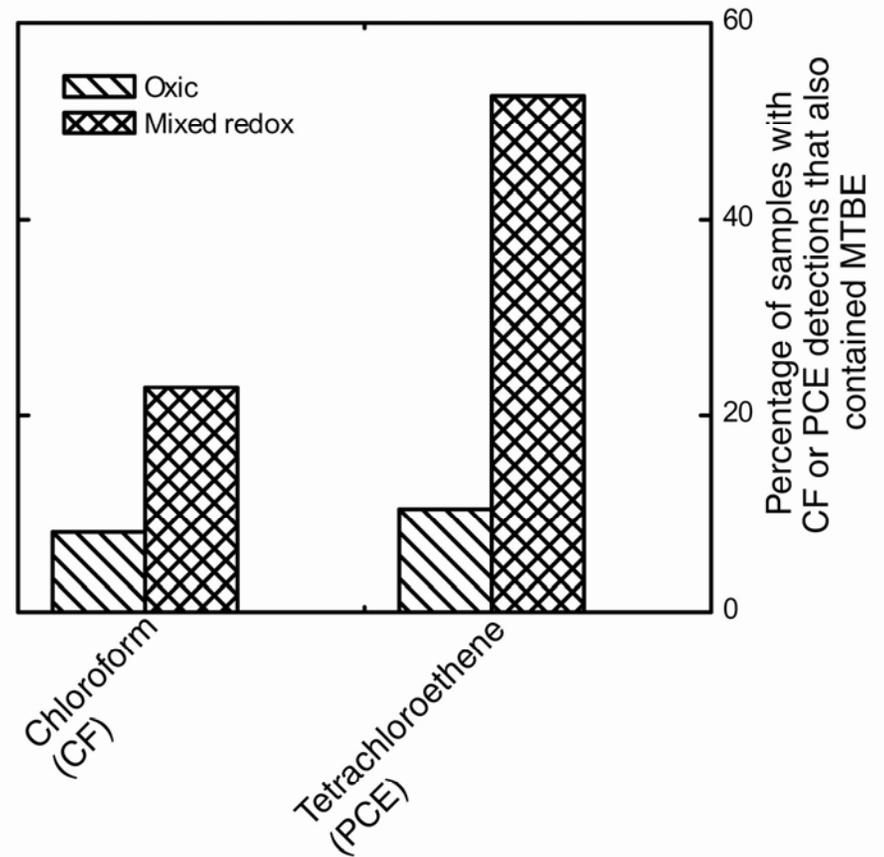
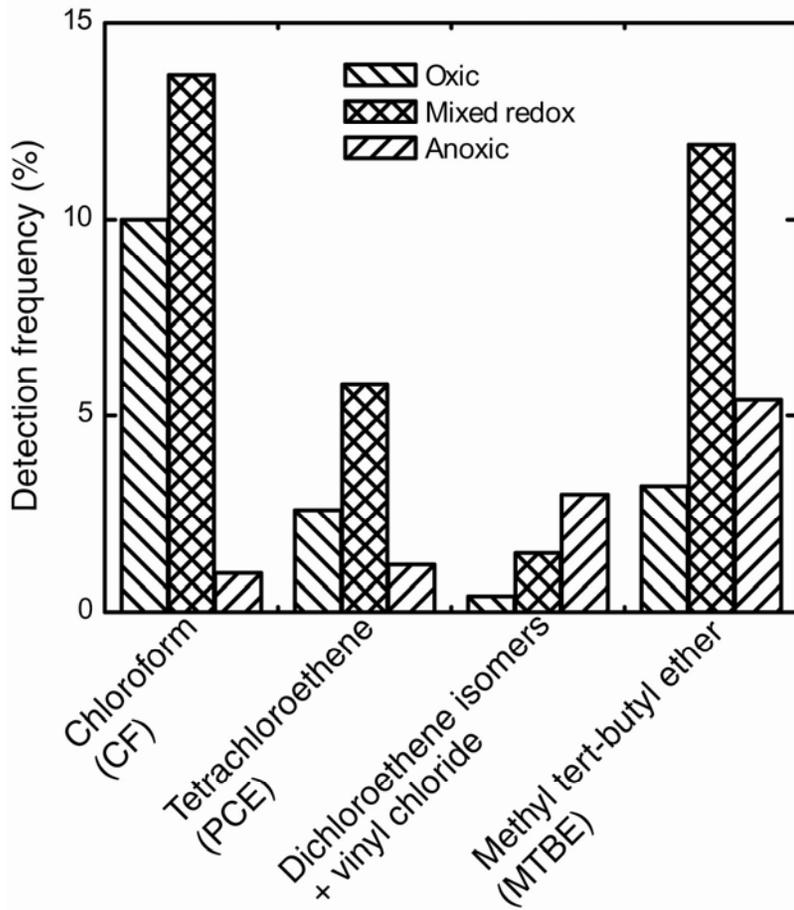
- Sand and Gravel
- Carbonate/Sandstone
- △ Glacial Deposit
- × Carbonate/Crystalline
- Semiconsolidated Sandstone
- ◆ Western Volcanics



# Redox Controls on Nitrate and Arsenic



# Redox and VOCs



data from all well types  
except agri. monitoring wells

# Next Steps

**Produce companion analysis of public supply well data set**

**Extend analysis to other PA's using other USGS data sets**

# Lack of DO data a problem!

- Have queried USGS NWIS data base to get additional data to extend redox comparison to other PA's.
- **Colorado example:**
- # samples in initial retrieval = **6,457**
- # samples after dropping samples with no data for
  - $O_2$  = **552**
  - $NO_3$ , Mn, Fe, or  $SO_4$  = **386**
  - Aquifer code = **353**
- # samples after retaining only most recent sample for wells with multiple samples = **200**

**This is less than 3% of starting number!**

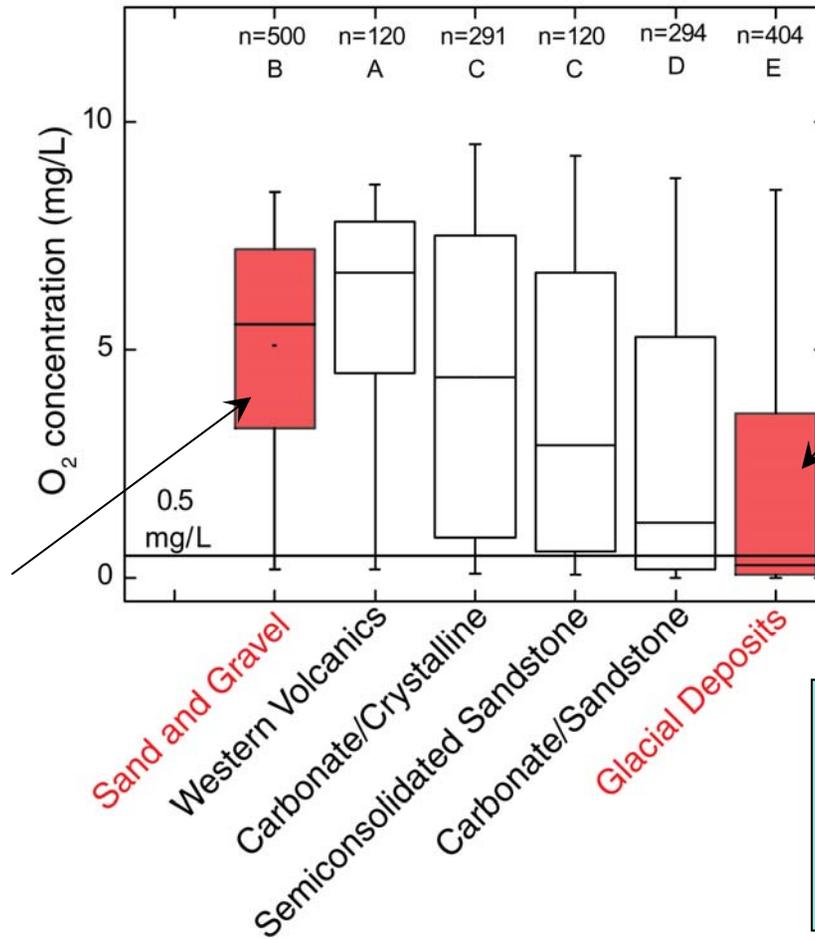
# Take Home Messages

- Spatial patterns in redox at the Principal Aquifer scale reflect the influence of climate, geology, and hydrology
- Mixed redox ground-water samples provide information on redox heterogeneity that can be related to the occurrence of selected contaminants
- Relatively inexpensive and easy to measure redox parameters such as DO, NO<sub>3</sub>, Fe, Mn, and SO<sub>4</sub> should be included in routine water-quality monitoring programs whenever possible
- **Require sampling of these parameters in the National Ground Water Monitoring Network!**

## Reference for this Talk

- P.B. McMahon, and F.H. Chapelle, 2008, Redox Processes and Water Quality of Selected Principal Aquifer Systems, Ground Water 46 (2) , 259–271

# Geologic Controls on Redox

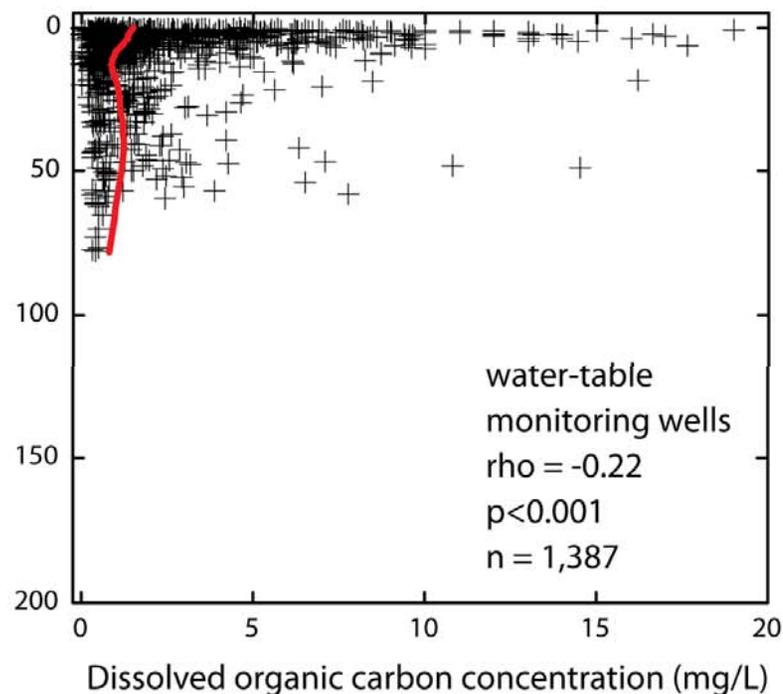
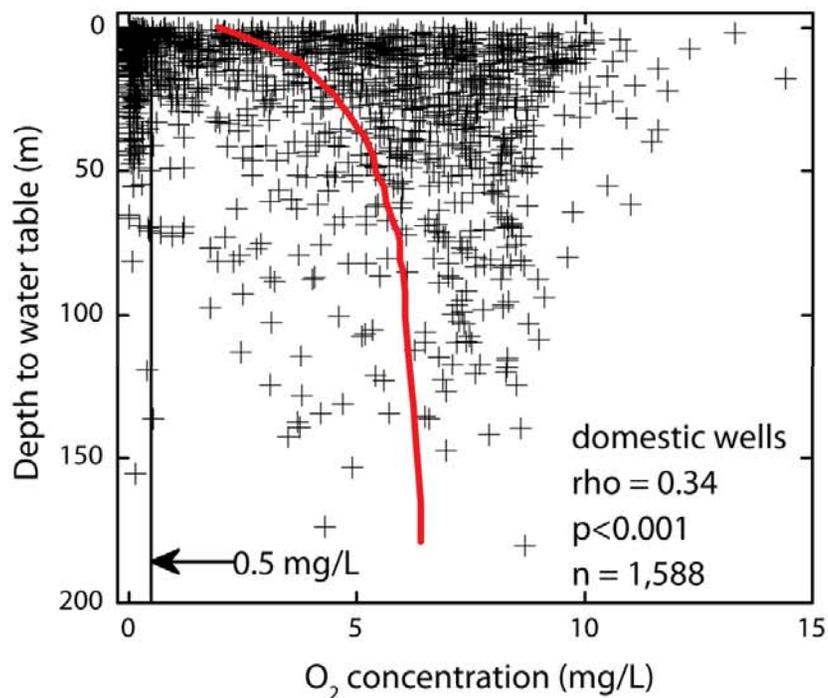


O<sub>2</sub> can persist for 1000s of years along kilometer-scale flow paths

Can transition from O<sub>2</sub> reducing to methanogenic conditions along meter-scale flowpaths

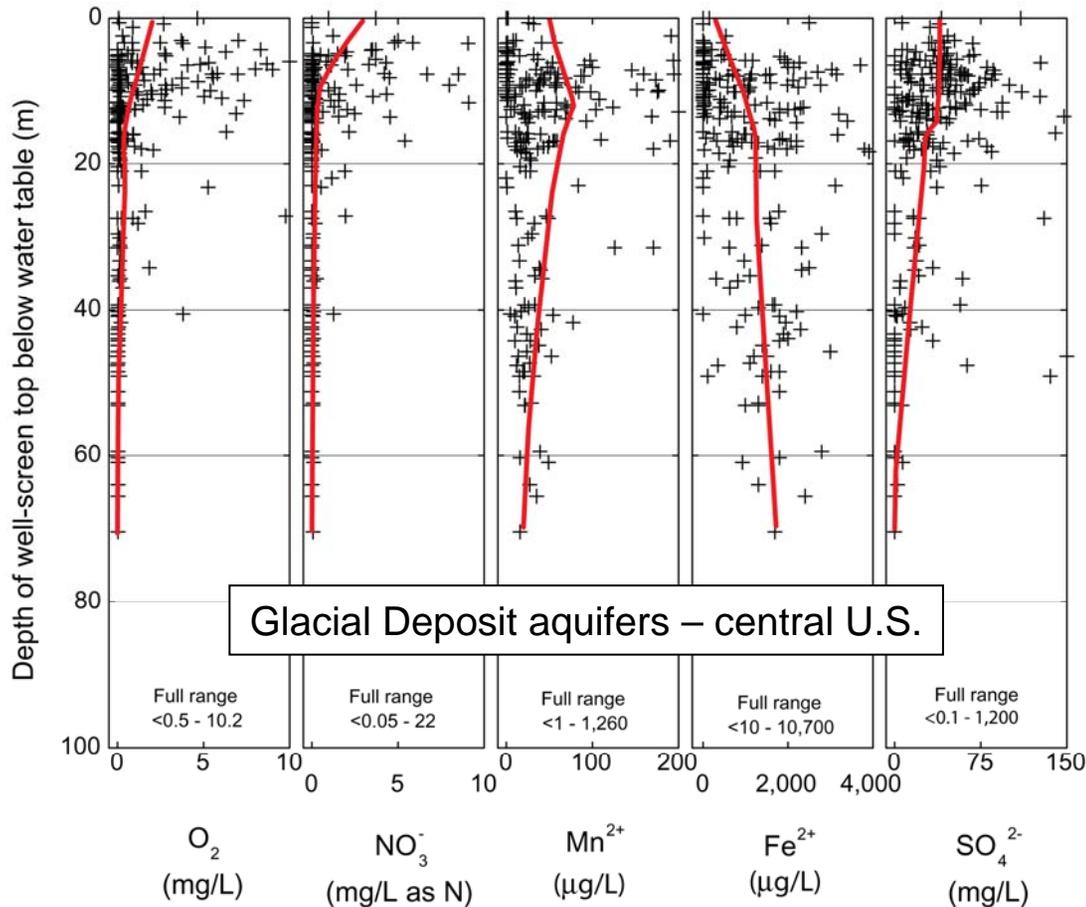
Sediment source and depositional environment influence e-acceptor/donor availability

# Climatic Controls on Redox



Soil development, vegetation, and recharge influence e- acceptor/donor availability

# Hydrologic Controls on Redox



Water residence time influences reaction progress

Anthropogenic factors also influence e-acceptor/donor availability and water residence time