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Bacteria Holding Time and Degradation

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Back in 2002-03
Collaborated with water boards to provide data for discussions to overcome sample hold time road-blocks to conducting fecal indicator bacteria monitoring.

Research
Quantify the effect of sample hold time so we can standardize data and assess remote waters.

Most of the State's waters are > 6 hr from a laboratory



...and unless we can either control the weather or implement the 24 hour work day and outlaw weekends, they can be >24 hours from getting analyzed!



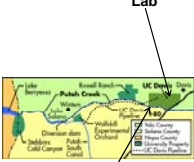

Decay Studies

- 15 streams enrolled
- Rangeland to Ag/Urban
- Joint with CVRWQCB – 5 streams, DMF & Colilert
- Samples analyzed <3 hr: then -6, 12, 24, 48, 72, 96hr, 4 C

cfu/100mL = a + b * time

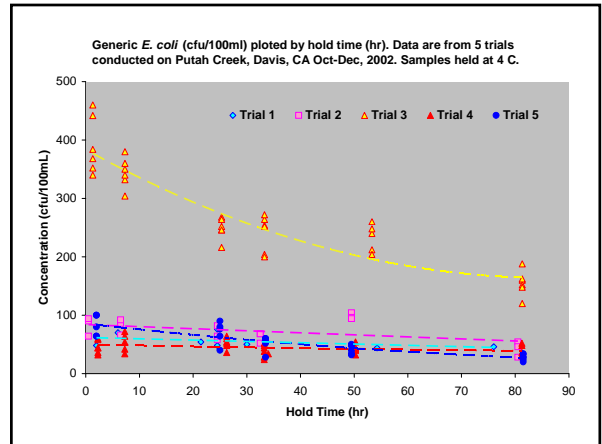



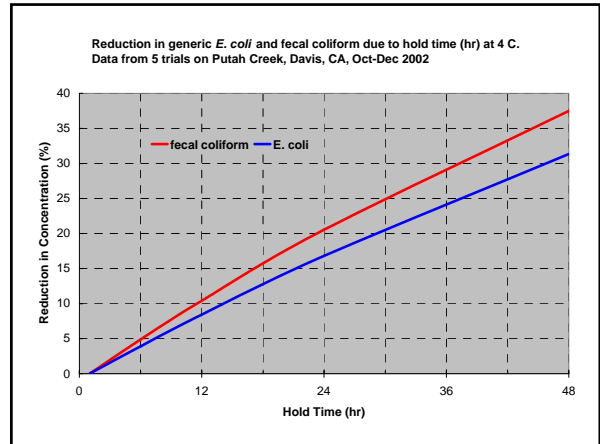
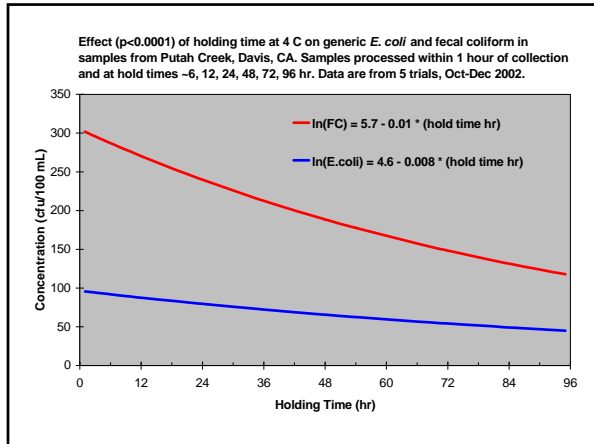
Putah Creek – Initial Study


Lab

Sample Site






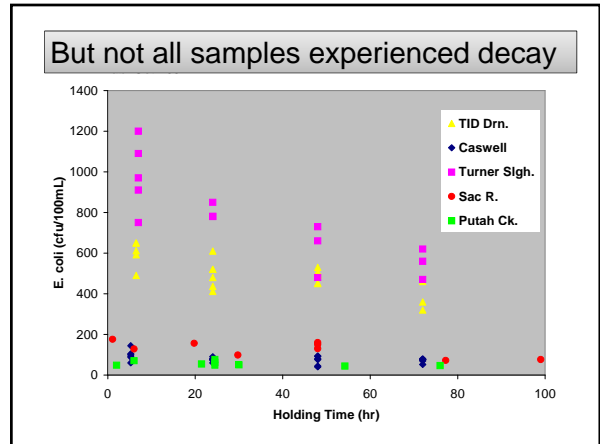
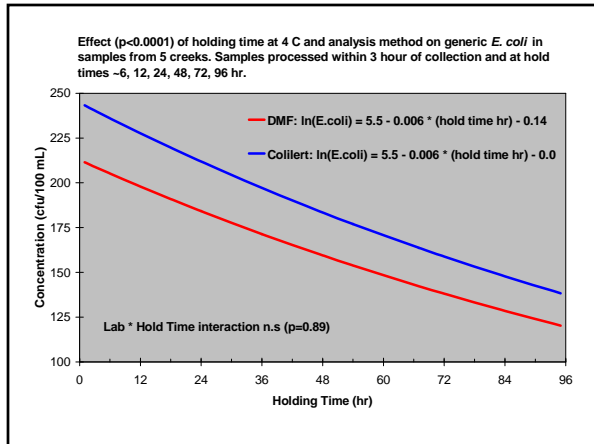
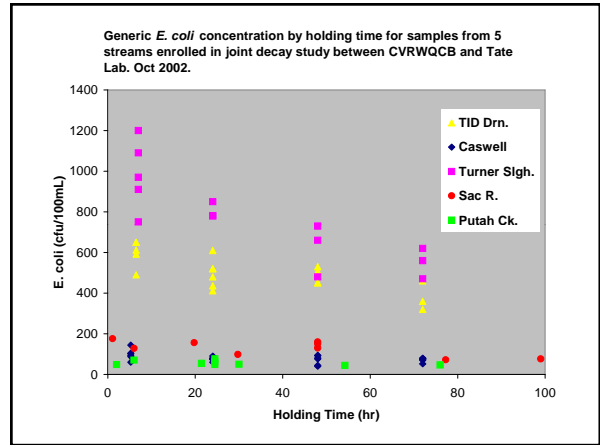
**Is Putah just weird?
Other streams?
Other methods?**

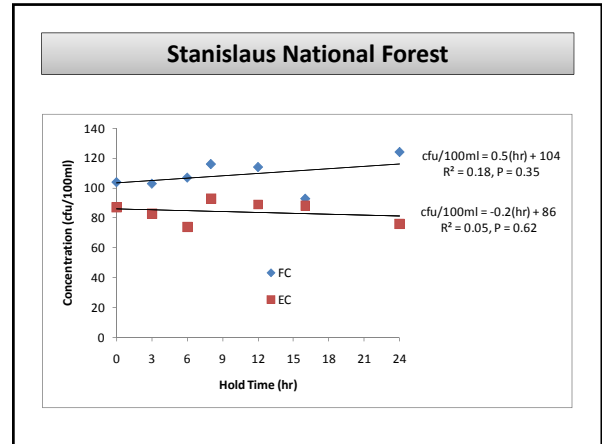
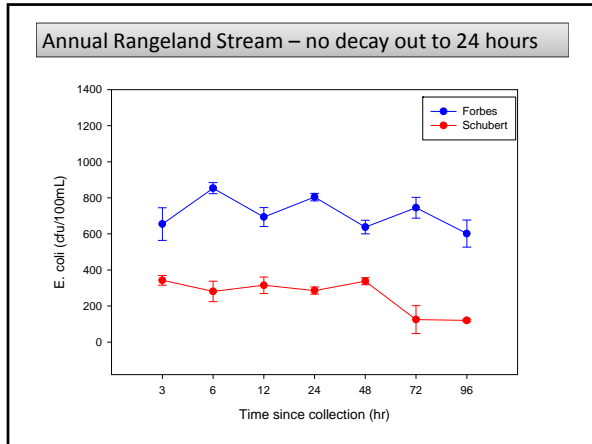


**Joint project with
CVRWQCB**

Add 4 more streams
TID, Discovery Park,
Turner Slough, Putah,
Caswell Park

2 methods
Direct Membrane Filtration,
Colilert



Tate, K.W., E.R. Atwill, J.W. Bartolome, and G.A. Nader. 2006. Significant *E. coli* Attenuation by Vegetative Buffers on Annual Grasslands. J. Environmental Quality. 35:795-805.

In order to adjust the *E. coli* concentration in each water sample tested x hours ($t=x$) after initial time of collection ($t=0$) to a single 24-hour standard ($t=24$), we first assumed the following basic model:

$$\log_{10}(EC_{24}) = \log_{10}(EC_x) + \beta(t=x) \quad [1]$$

whereby $\log_{10}(EC_x)$ was the observed \log_{10} concentration of *E. coli* determined x hours ($t=x$) after initial time of collection, $\log_{10}(EC_{24})$ was the modeled \log_{10} concentration of *E. coli* at the initial time of collection ($t=0$), and $\beta(t=x)$ was the fitted decay coefficient(s) generated by the linear mixed effects model described above. $\beta(t=x)$ was allowed to be a univariate or polynomial term depending on whether the raw data signified a first or second-order time-dependent decay process for the *E. coli* concentration in our source water. The decay process was for water samples held at approximately 4°C. Once $\beta(t=x)$ was obtained, equation (2) was used to adjust each sample to a single 24-hour standard ($t=24$).

$$EC_{24} = (EC_x) 10^{\beta(t=x)} \quad [2]$$

whereby EC_{24} was the fitted or expected concentration of *E. coli* at a 24-hour standard, EC_x was the observed concentration of *E. coli* determined x hours ($t=x$) after initial time of collection, and $10^{\beta(t=x)}$ was the expected decay coefficient adjustment factor raised to the power of 10 which allowed us to model concentrations of *E. coli* directly instead of \log_{10} values.

Summary

- We can account for the decay of indicator bacteria resulting from hold time.
- Hold times out to 24 hours may have limited decay for many streams/samples.
- Will each waterbody require a unique curve? Looks like at a minimum that unique curves will be required for “types” of streams (i.e. range v. ag v. urban).