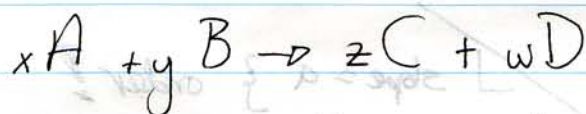


# Methylene Glycol Clock Reaction

\* Gather Data in Pairs.

\* Work calculations & process data separately

## Rate Law



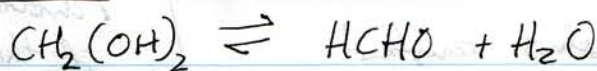
$$\text{rate} = k [A]^a [B]^b$$

• Powers are orders of reaction  
not to be confused with  
stoichiometric coefficients.

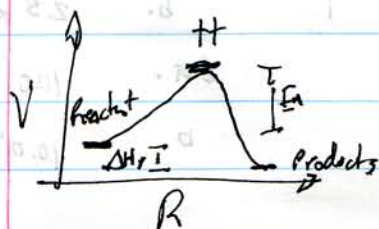
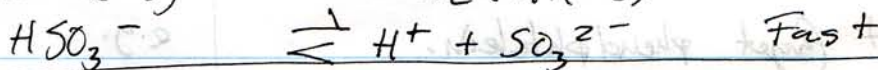
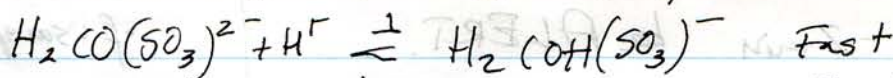
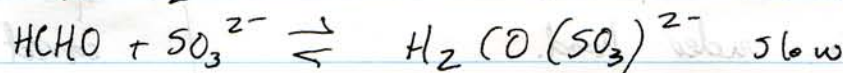
• Overall order is sum.

$$\text{rate} = \frac{d[\text{product}]}{dt} = - \frac{d[\text{reactant}]}{dt} \quad [ ] = \frac{\text{moles}}{L}$$

## Mechanism



Slowest  $\Rightarrow$  rate determining



$$k = A e^{-E_a/RT}$$

Arrhenius Eq.

A = collision frequency factor

k = specific rate constant

$$\ln k = \ln A - \frac{E_a}{R} \cdot \frac{1}{T}$$

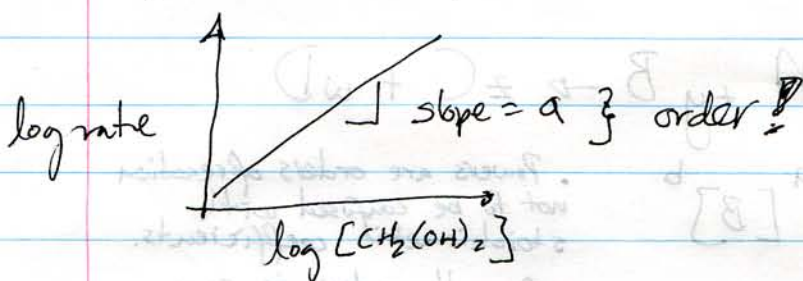
$$pH = pK_a + \log \left( \frac{[base]}{[acid]} \right) \left\{ \frac{[A^-]}{[AH]} \frac{[salt]}{[acid]} \frac{[Na_2SO_3]}{[NaHSO_3]} \right\}$$

$$rate = k [CH_2(OH)_2]^a [NaHSO_3]^b$$

$$\log rate = \underbrace{\log k}_{\text{constant}} + a \log [CH_2(OH)_2] + b \log [NaHSO_3]$$

constant  
initial rates method.

$$\log rate = a \log [CH_2(OH)_2] + \text{const.}$$



$$rate = k [A]^a [B]^b \quad K = \frac{rate}{[A]^a [B]^b}$$

- Do twice
- Try to keep T of solns. constant.
- Reaction under hood.
- 10s - 2min be ALERT.
- Don't forget phenolphthalein.
- Use white paper under beaker.

• check to see if their books are correct on preparation of samples.

e.g. A  $\xrightarrow{[HSO_3^-]}$  B  $\xrightarrow{gluc}$

Sol.	2.5	10.0
b.	2.5 <sup>not</sup>	10.0
6a.	10.0	10.0
b.	10.0 <sup>not</sup>	10.0