

# Inversion of Sucrose

## A. Reaction and Kinetics

1.  $S + H^+ + H_2O \rightleftharpoons F + G + H^+$  (*i.e.*, acid-catalyzed)
2. *Rate Law*:  $r = -d[S]/dt = k_H [H^+] [S] \equiv k_{\text{eff}} [S]$   
(*pseudo-first order* — because  $[H^+]$  doesn't change)
3. *Integrate*:  $-d[S]/[S] = k_{\text{eff}} dt \quad \square \quad \ln [S] = \text{const} - k_{\text{eff}} t$
4. *Boundary Condition*: @  $t = 0$ ,  $[S] = [S]_0 \quad \square \quad \text{const} = \ln [S]_0$   
 $\square \quad [S] = [S]_0 \exp(-k_{\text{eff}} t)$  (first-order decay law)

## B. Experiment

1. Measure  $\square$  with polarimeter. But all 3 sugars are optically active, so  $\square$  doesn't vanish when S is 100% converted.
2. Rather,  $\square = A \exp(-k_{\text{eff}} t) + B$  (exponential + background)
3.  $t = 0$ :  $\square (\equiv \square_0) = A + B$ ;  $t = \infty$ :  $\square = B$  ( $\square$  for F + G)

4. *Specific rotation*:  $[\alpha] = [\alpha]_D^T [\alpha] \ell$   
units of  $[\alpha]_D^T$ : degree L g<sup>-1</sup> dm<sup>-1</sup>
5. Need one “completion” run to get  $[\alpha]$  .

### C. Temperature Dependence

1. *Arrhenius*:  $k = A \exp(-E_a/RT)$   $\square$   $\ln k = \ln A - E_a/RT$
2. *Typical*: Have  $k$  at two  $T$ s  $\square$   $k_2/k_1 = \exp[(E_a/R)(1/T_1 - 1/T_2)]$   
 $\ln(k_2/k_1) = (E_a/R)(1/T_1 - 1/T_2)$  (don't need  $A$ )
3. **Note**: This is our second encounter with  $\ln(\ ) = a + b/T$  .  
We will see it again in analyzing the  $T$  dependence of equilibrium constants (Expt. 5) and vapor pressure (Expt. 6).

In all such thermodynamic applications,  $T$  is in K.

# Statistical Error Propagation

**A. Statistics Notes 3**

**B. Statistics Problems 11-17**

**C. Partial Differentiation**

Exercises available at

<http://www.vanderbilt.edu/AnS/Chemistry/Tellinghuisen/Chem236/PDExer.pdf>

and solutions at

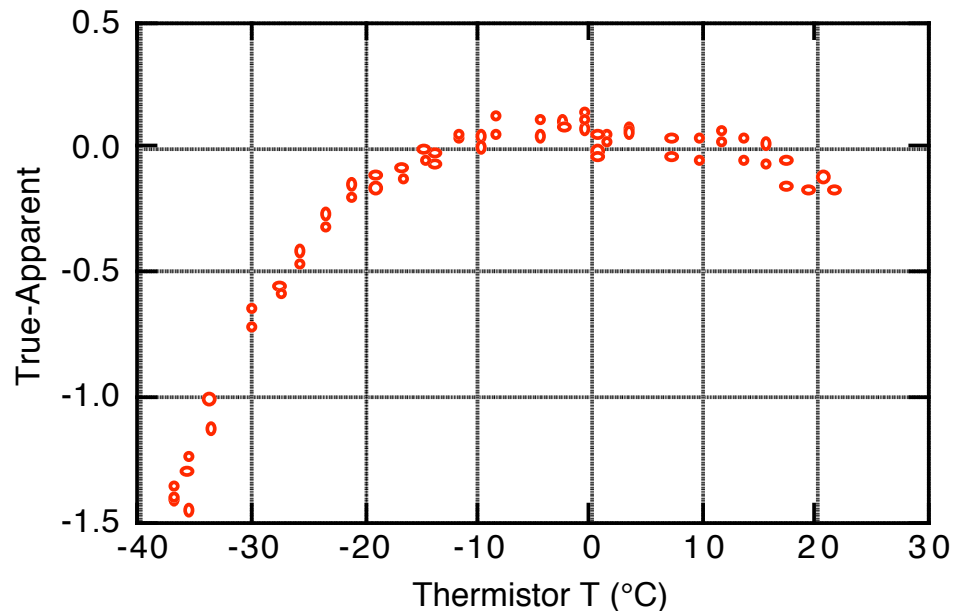
<http://www.vanderbilt.edu/AnS/Chemistry/Tellinghuisen/Chem236/PDSoln.pdf>

# Today's Practice Quiz !

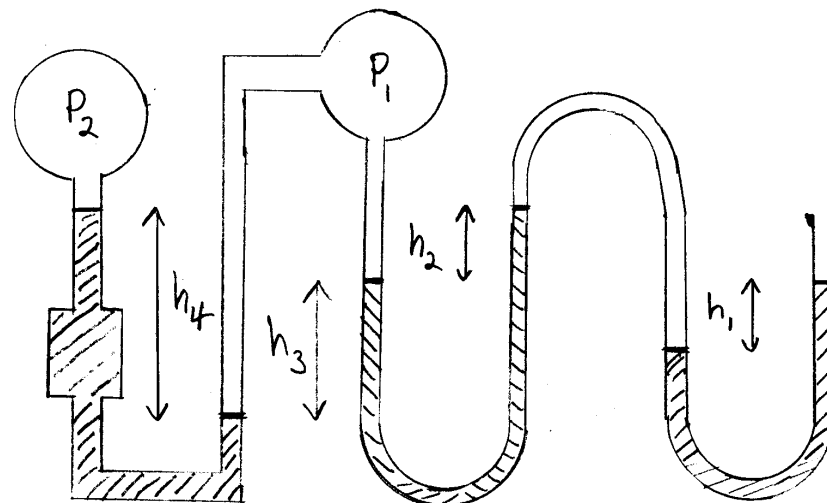
1. A quantity is known to follow a linear relationship,  $y = a + bx$ . If  $y = -1.5$  when  $x = 0.2$  and  $y = 111.7$  when  $x = 8.9$ , what are the values of  $a$  and  $b$ ?
- a.  $a = -2.5; b = 5.0$       b.  $a = -3.5; b = 10.0$       c.  $a = -4.0; b = 12.7$   
d.  $a = -4.1; b = 13.0$       e. none of these

2. Student records a data point on the sublimation curve of ice, where the thermistor reads  $-24.1^\circ\text{C}$ . If the calibration data are as shown in the accompanying graph, what is the corrected  $T$ ?

- a.  $-24.5^\circ\text{C}$       b.  $-23.7^\circ\text{C}$   
c.  $0.4^\circ\text{C}$       d.  $-0.5^\circ\text{C}$   
e. This cannot be determined w/o additional information.



3. Refer to the accompanying figure of three connected Hg manometers. If atmospheric pressure is 754 torr,  $h_1 = 111$  mm,  $h_2 = 83$  mm,  $h_3 = 192$  mm, and  $h_4 = 289$  mm, what are the pressures  $P_1$  and  $P_2$  (in Torr)?



- a. 865 & 948      **b. 948 & 659**  
 c. 948 & 851      d. 560 & 849  
 e.  $P_1 = 948$  Torr;  $P_2$  cannot be determined.

4. If barometers used water as the operating fluid, a weather barometer would need to be about how tall? [1.00 m = 39.37 in.]

- a. 5 ft      b. 15 ft      **c. 40 ft**      d. 100 ft      e. none of these

5. Besides the size problem, can you think of any other reason why water might be an unwise choice for a barometer?

- a. It is transparent.      **b. It has significant vapor pressure.**  
 c. Mosquitoes can breed in  $H_2O$  but not in Hg.      d. All of the above  
 e. None of the above