

14/15

Time: 10min

Chemistry 203

Mon., Dec. 12, 2011

Quiz 4

Name: Bilal Hammond

Section: _____

- In today's experiment you are going to use the Henderson Hasselbalch equation to determine the pKa of the indicator. Derive this equation and state how you can determine the pKa of bromthymol blue using this equation.

$$pH = pKa + \log \frac{[In^-]}{[HIn]}$$



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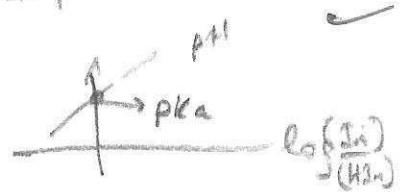
$$Ka = \frac{[In^-][H^+]}{[HIn]} = \frac{[In^-]}{[HIn]} \cdot 10^{pH}$$

$$pH = -\log [H^+] \Rightarrow [H^+] = 10^{-pH}$$

log both sides $pKa = \log \frac{[In^-]}{[HIn]} + \log 10^{pH} = \log \frac{[In^-]}{[HIn]} + pH$

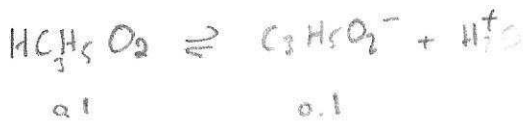
$$\Rightarrow pH = pKa + \log \frac{[In^-]}{[HIn]}$$

plot of pH versus $\log \frac{[In^-]}{[HIn]}$ and see intersect in pH



- Calculate the pH of a solution (A) containing 0.100M HC₃H₅O₂ (Ka=1.3x10⁻⁵) and 0.100M NaC₃H₅O₂.
 - Calculate the pH after adding 0.020mol NaOH to 1.00L of (A).
 - What can you say about solution (A)?

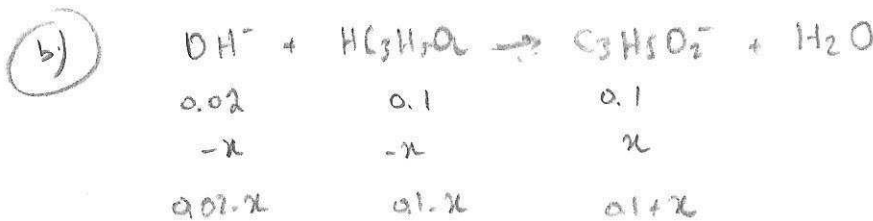
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(c) A is a buffer solution
pH is slightly change when adding OH⁻

$$a) pH = pKa + \log \frac{[C_3H_5O_2^-]}{[HC_3H_5O_2]} = pKa + \log \frac{0.1}{0.1} = pKa + \log 1 = pKa$$

$$pKa = -\log (1.3 \times 10^{-5}) = 4.89$$



$$[OH^-] \cdot \frac{1}{1} = \frac{0.02}{1} = 0.02 \text{ mol/L}$$

consider $x \approx 0.02 \text{ mol/L} \Rightarrow Ka = \frac{[C_3H_5O_2^-]}{[OH^-][HC_3H_5O_2]} \cdot Kw = \frac{(0.1+x) 10^{-14}}{(0.02-x)(0.1-x)} = 1.3 \times 10^{-5} \Rightarrow$ solve the equation

$$pH = pKa + \log \left(\frac{0.1+x}{0.1-x} \right) = 4.89 + \log \frac{0.1+0.02}{0.1-0.02} = 4.89 + 0.2 = 5.09$$

version 2

Time: 10min

Chemistry 203

Fri., Dec. 16, 2011

Quiz 4

Name: Mahmoud Tohamz

Section: 6

1. In today's experiment you are going to determine the thermodynamic functions of the dissolution of borax by a graphical method. What are going to plot and how can you determine the needed quantities from the plot?

we are going to plot $\ln K$ versus $\frac{1}{T}$

$$\Delta G_o = -RT \ln K \quad \Delta G_o = \Delta H_o - T \Delta S_o \quad \ln K = \frac{\Delta H_o - T \Delta S_o}{-RT} = \frac{\Delta S_o}{R} - \frac{\Delta H_o}{R} \cdot \frac{1}{T}$$

The slope of the curve is $-\frac{\Delta H_o}{R}$, & from it we determine ΔH_o

The y-intercept is $\frac{\Delta S_o}{R}$, & from it we determine ΔS_o

(The eqn. of the curve is $\ln K = \frac{\Delta S_o}{R} - \frac{\Delta H_o}{R} \cdot \frac{1}{T}$)

2. Consider the following reaction $\text{Br}_2(l) \rightarrow \text{Br}_2(g)$
- a. What is the sign of ΔG° for the reaction to be spontaneous?
- b. At what temperatures is the above reaction spontaneous at 1 atm?
- $\Delta H^\circ = 31.0 \text{ kJ/mol}$ $\Delta S^\circ = 93.0 \text{ J/K}\cdot\text{mol}$

a) $\Delta G^\circ < 0$ for a spontaneous reaction.

b) $\Delta G_o = \Delta H_o - T \Delta S_o$

$\Rightarrow 31 \times 10^3 - T(93.0) < 0$

$93 \times T > 31 \times 10^3$

$T > \frac{31 \times 10^3}{93} \Rightarrow T > 333.33 \text{ K}$

3. Assuming standard conditions, can the following reaction take place at room temperature, 25°C? $3\text{Cl}_2(g) + 2\text{CH}_4(g) \rightarrow \text{CH}_3\text{Cl}(g) + \text{CH}_2\text{Cl}_2(g) + 3\text{HCl}(g)$
- Given the standard free energy of formation for $\text{CH}_4(g) = -50.72 \text{ kJ/mol}$
- $\text{CH}_2\text{Cl}_2(g) = -68.85 \text{ kJ/mol}$
- $\text{CH}_3\text{Cl}(g) = -57.37 \text{ kJ/mol}$
- $\text{HCl}(g) = -95.30 \text{ kJ/mol}$

$$\Delta G^\circ_{\text{rxn}} = \sum \Delta G^\circ_{\text{products}} \times \text{stoichiometric coeff.} - \sum \Delta G^\circ_{\text{reactants}} \times \text{stoichiometric coeff.}$$

$$= \left[\Delta G^\circ(\text{CH}_3\text{Cl}) + \Delta G^\circ(\text{CH}_2\text{Cl}_2) + 3\Delta G^\circ(\text{HCl}) \right] - \left[3\Delta G^\circ(\text{Cl}_2) + 2\Delta G^\circ(\text{CH}_4) \right]$$

$$= \left[-57.37 - 68.85 + 3(-95.3) \right] - \left[0 + 2(-50.72) \right]$$

$$= -310.68 \text{ kJ/mol}$$

$\Delta G^\circ_{\text{rxn}} < 0 \Rightarrow$ Reaction is spontaneous \Rightarrow yes it can take place at 25°C