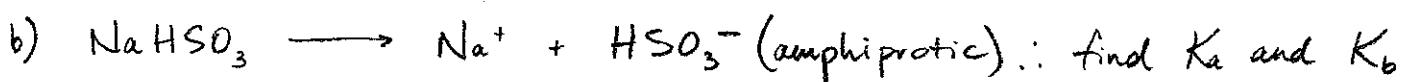
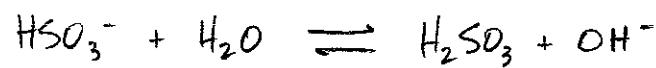
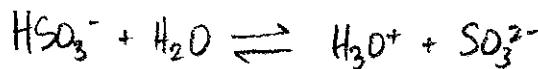


ACID/BASE II
REVIEW KEY



$$K_a(\text{HSO}_3^-) = 1.0 \times 10^{-7}$$

$$K_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{1.0 \times 10^{-7}} = 1.0 \times 10^{-7}$$



$K_a > K_b$ ∴ resulting solution is ACIDIC.



$$I \quad 0.40 \text{ M}$$

$$C \quad -x$$

$$E \quad 0.40-x$$

$$\begin{matrix} \text{H}_3\text{O}^+ & 0 \text{ M} \\ +x & +x \\ & x \end{matrix}$$

$$\begin{matrix} \text{CH}_3\text{COO}^- & 0 \text{ M} \\ +x & +x \\ & x \end{matrix}$$

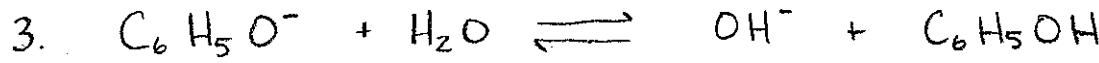
$$\text{let } x = \Delta[\text{CH}_3\text{COOH}]$$

$$K_a = 1.8 \times 10^{-5} = \frac{x^2}{0.40-x}$$

$$1.8 \times 10^{-5} = \frac{x^2}{0.40} \quad \text{Check } \checkmark$$

$$x = 2.6832816 \times 10^{-3} \text{ M}$$

$$\therefore \text{pH} = -\log(x) = 2.57$$



$$I \quad 2.00 \text{ M}$$

$$C \quad -x$$

$$E \quad 2.00-x$$

$$\begin{matrix} \text{OH}^- & 0 \text{ M} \\ +x & +x \\ & x \end{matrix}$$

$$\text{let } x = \Delta[\text{C}_6\text{H}_5\text{O}^-]$$

$$K_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{1.3 \times 10^{-10}} = 7.6923 \times 10^{-5} = \frac{x^2}{2.00}$$

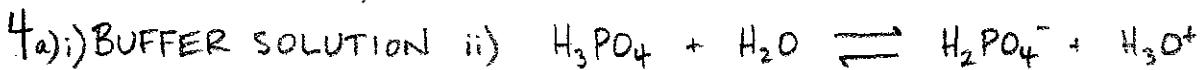
$$\text{Assume } 2.00-x \approx 2.00$$

Check ✓

$$x = 1.24035 \times 10^{-2} \text{ M}$$

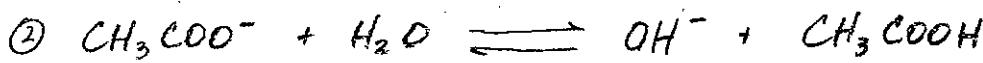
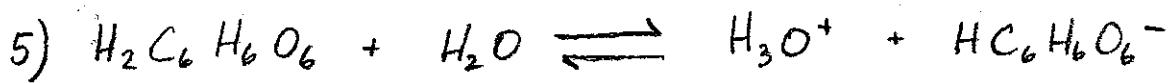
$$\text{pOH} = -\log(x) = 1.9065$$

$$\text{pH} = 14 - 1.9065 = 12.09$$

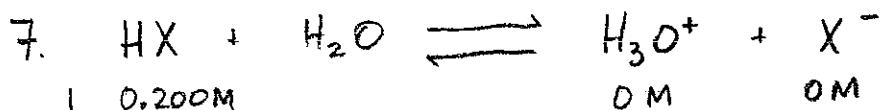


b) $\text{pH} = -\log(K_a) = -\log(7.5 \times 10^{-3}) = 2.12$

c) $\text{mol OH}^- = \text{MV} = 1.0 \times 0.0100 \text{ L} = 0.0100 \text{ mol OH}^-$ added BUT $1.00 \text{ mol H}_3\text{PO}_4$ available to neutralize. pH raises slightly as ratio of $\text{H}_3\text{PO}_4 : \text{H}_2\text{PO}_4^-$ changes but only slightly.



$$b) K_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{1.8 \times 10^{-5}} = \boxed{5.6 \times 10^{-10}}$$



I 0.200M

0M 0M

C $-3.6308 \times 10^{-3}\text{M}$

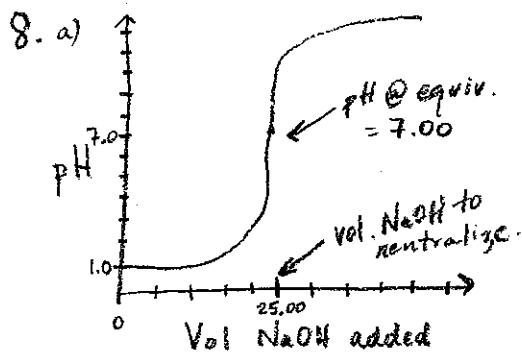
$3.6308 \times 10^{-3}\text{M} \rightarrow$

E 0.19637

$3.6308 \times 10^{-3}\text{M} \rightarrow$

$$[\text{H}_3\text{O}^+] = [\text{X}^-] = \text{invlog}(-2.44) \\ = 3.6308 \times 10^{-3}\text{M}$$

$$K_a = \frac{[\text{H}_3\text{O}^+]^2}{[\text{HX}]} = \frac{(3.6308 \times 10^{-3})^2}{0.19637} = 6.7 \times 10^{-5} \text{ The acid is } \underline{\text{BENZOIC}}$$



$$\text{pH}_{\text{initial}} = -\log(0.10\text{M})^* \text{ because HCl is strong} \\ = 1.00$$

$$\text{mol HCl} = \text{mol H}_3\text{O}^+ = MV = (0.10\text{M})(0.0250\text{L}) = 0.00250\text{ mol}$$

$$\text{mol OH}^- = \text{mol NaOH} = 0.00250 \text{ mol}$$

$$V = \frac{\text{mol}}{\text{M}} = \frac{0.00250 \text{ mol}}{0.10 \text{ M}} = 0.0250 \text{ L} \\ \approx 25.00 \text{ mL}$$

pH at equiv. point = 7.00 (Strong-Strong titration)

b) Suitable Indicator: Bromothymol blue, Phenol Red, Neutral Red

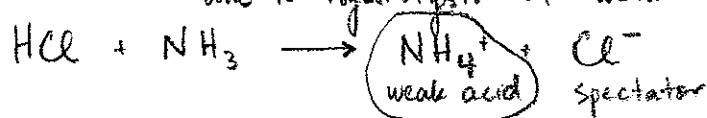
OR
* Phenolphthalein (cheap)

9. 2 characteristics of Strong Acid / Weak Base Titration Curve:

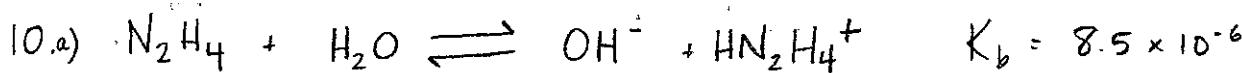
i) Initial noticeable drop in curve (pH).

ii) pH @ equiv. pt < 7.00 . (around pH 4-6)

due to hydrolysis of weak acid...

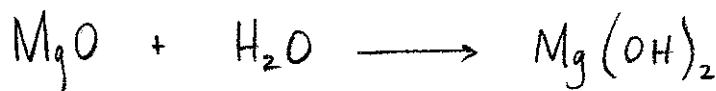


Indicator: Brom cresol Green, Methyl Red



b) CONJUGATE ACID IS HN_2H_4^+ $K_a = \frac{K_w}{K_b} = \frac{1.0 \times 10^{-14}}{8.5 \times 10^{-6}} = \boxed{1.2 \times 10^{-9}}$

11. Mg O is a metal oxide \therefore it forms a BASE w/ water.
(basic anhydride)



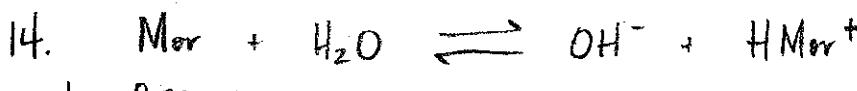
Student is wrong!

12. a) pH of distilled water = 7.00 \therefore Bromoresol green is BLUE.

b) Acetic acid produces some H_3O^+ which shifts the indicator's equilibrium left. $\text{[HInd]} > \text{[Ind}^-]$ \therefore more yellow than blue...
pH must be ≤ 3.8 . (see ind. table)

13. Transition Point = End Point $\therefore \text{[HInd]} = \text{[Ind}^-]$ COMBINATION OF THE 2 COLOURS

$$K_a = \frac{[\text{H}_3\text{O}^+][\text{Ind}^-]}{\cancel{[\text{HInd}]}}$$



I 0.010 M

0

C $-1.2589 \times 10^{-4}\text{ M}$

$1.2589 \times 10^{-4}\text{ M} \longrightarrow$

E 0.009874 M

$1.2589 \times 10^{-4}\text{ M} \longrightarrow$

$$\text{pH} = 10.10 \quad \text{pOH} = 3.90$$

$$[\text{OH}^-]_{\text{eq}} = \text{inv log}(-3.90) \\ = 1.2589 \times 10^{-4}\text{ M}$$

$$K_b = \frac{[\text{OH}^-]^2}{[\text{Mor}]} = \frac{(1.2589 \times 10^{-4})^2}{0.009874} = \boxed{2 \times 10^{-6}}$$

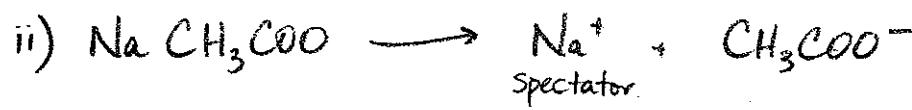
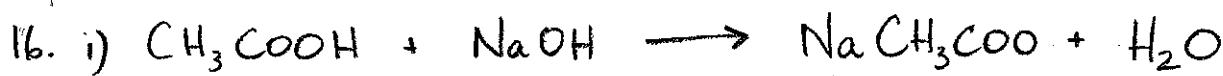
1 sig fig.



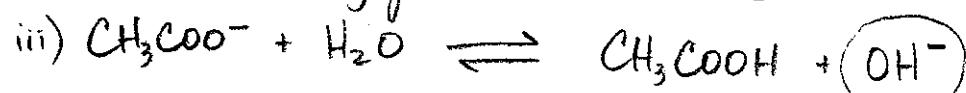
$$K_a \text{ of } \text{NH}_4^+ = 5.6 \times 10^{-10}$$

$$K_b \text{ of } \text{C}_2\text{O}_4^{2-} = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{5.6 \times 10^{-10}} = 1.6 \times 10^{-5}$$

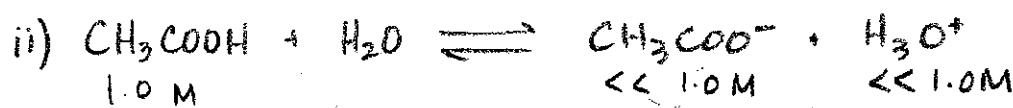
$K_a > K_b$ so ACIDIC



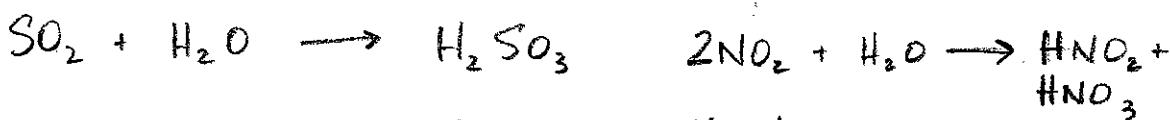
Na^+ is conjugate acid of strong base; cannot act as an acid!



basic $\therefore \text{pH} > 7.00$.



NaNO_2 would have greater conductivity (2.0 M total ions)!



19. Any weak acid in equal conc. with its conjugate base
(and vice versa)

e.g. 1.0 M CH_3COOH and 1.0 M CH_3COO^- (as NaCH_3COO)

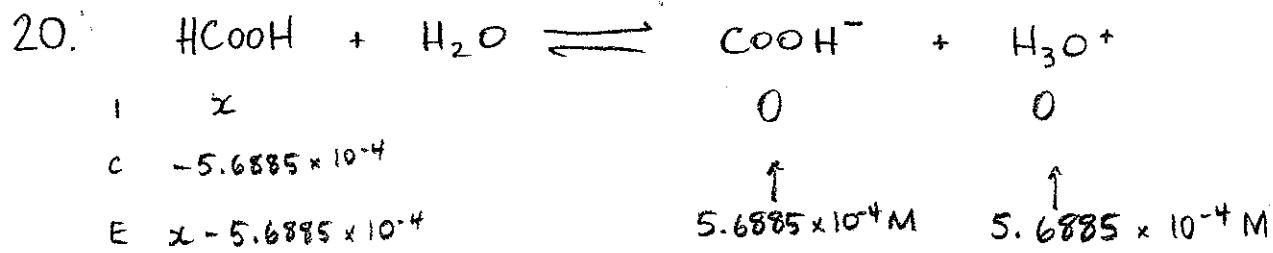


add HCl $\rightarrow \uparrow [\text{H}_3\text{O}^+] \therefore [\text{CH}_3\text{COO}^-] \downarrow$ (to neutralize) $\therefore [\text{CH}_3\text{COOH}] \uparrow$
EQUIL. shifts LEFT

Therefore, there exists more CH_3COOH after neutralization than CH_3COO^- .

But pH only decreases slightly ... a fact?

Buffer works though b/c without it, 'raw' HCl would affect pH much much more.

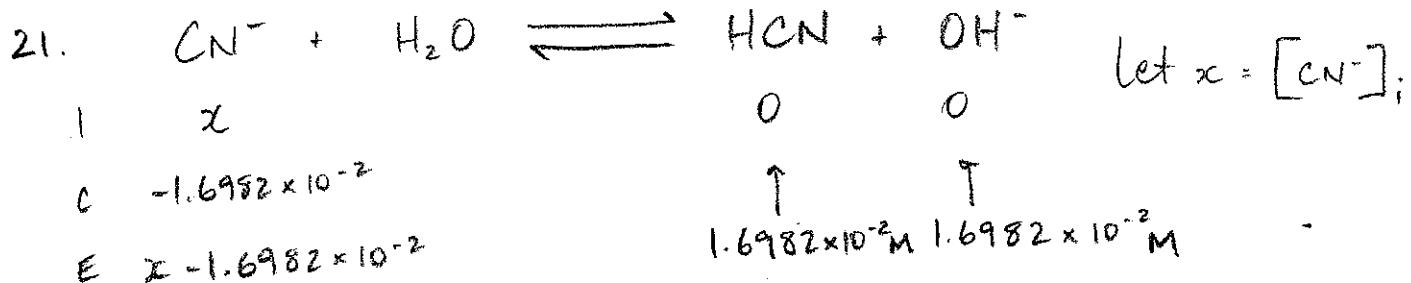


Let $x = [\text{HCOOH}]_i$; $[\text{H}_3\text{O}^+]_{eq} = \text{invlog}(-\text{pH})$
 $= \text{invlog}(-3.245)$
 $= 5.6885 \times 10^{-4} \text{ M}$

$$K_a = 1.8 \times 10^{-4} = \frac{[\text{H}_3\text{O}^+]^2}{[\text{HCOOH}]} = \frac{(5.6885 \times 10^{-4})^2}{x - 5.6885 \times 10^{-4}}$$

$$x = 2.3666 \times 10^{-3} \text{ M}$$

$$[\text{HCOOH}]_i = \boxed{2.4 \times 10^{-3} \text{ M}}$$



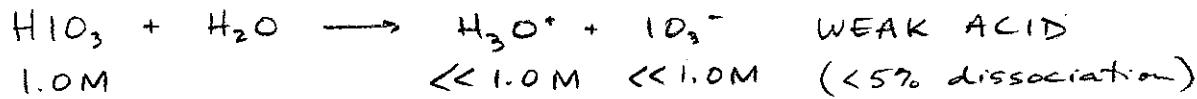
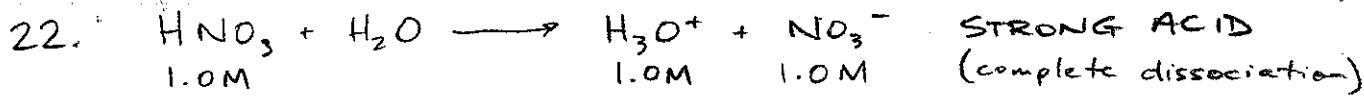
$$[\text{OH}^-]_{eq} = \text{invlog}(-\text{pOH}) \\ = \text{invlog}(-1.77) \\ = 1.6982 \times 10^{-2} \text{ M}$$

$$K_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{4.9 \times 10^{-10}} = 2.0408 \times 10^{-5}$$

$$2.0408 \times 10^{-5} = \frac{[\text{OH}^-]^2}{[\text{CN}^-]} = \frac{(1.6982 \times 10^{-2})^2}{x - 1.6982 \times 10^{-2}}$$

$$x = 14.15 \text{ M}$$

$$[\text{CN}^-] = \boxed{14 \text{ M}}$$

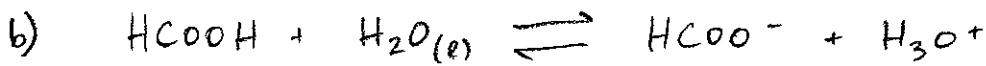


LESS IONS = LESS CONDUCTIVITY.

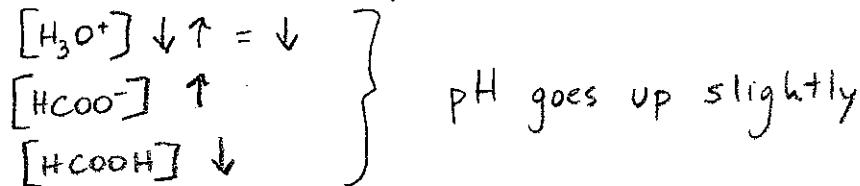
23.a) since $[\text{HCOOH}] = [\text{HCOO}^-]$, we have a 'perfect' buffer so

$$K_a = [\text{H}_3\text{O}^+] \text{ and } \text{pH} = \text{p}K_a$$

$$\begin{aligned} \text{pH} &= -\log(1.8 \times 10^{-4}) \\ &= \boxed{3.74} \end{aligned}$$



Adding OH^- will do the following:

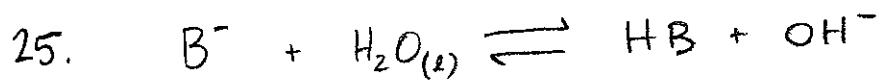


24. i) use a pH meter

ii) use an indicator or a combination of indicators

iii) perform a titration with a known [base].

i)



$$\text{pH} = 11.78 \quad \text{pOH} = 2.22$$

$$[\text{OH}^-]_{eq} = \text{inv log}(-2.22) = 6.0256 \times 10^{-3} \text{ M}$$

$$\begin{array}{l} \text{from titr. data E} \\ \text{C} \rightarrow 0.2956 \text{ M} \\ \text{C} - 0.0060256 \\ \text{E} 0.28957 \end{array}$$

$$\begin{array}{ccc} \text{O} & & \text{O} \\ \leftarrow +6.0256 \times 10^{-3} & & \text{from pH data} \\ \leftarrow 6.0256 \times 10^{-3} \text{ M} & & \end{array}$$

Titration:

$$K_b = \frac{[\text{OH}^-]^2}{[\text{B}^-]}$$

$$= \frac{(6.0256 \times 10^{-3})^2}{0.28957}$$

$$K_b = 1.3 \times 10^{-4}$$

$$\begin{aligned} \text{mol HCl added} &= \text{MV} \\ &= (0.25 \text{ M})(0.02956 \text{ L}) \\ &= 0.00739 \text{ mol HCl} \\ &= 0.00739 \text{ mol H}_3\text{O}^+ \\ &= 0.00739 \text{ mol OH}^- \\ &= 0.00739 \text{ mol B}^- \end{aligned}$$

$$[\text{B}^-]_i = \frac{\text{mol}}{\text{V}} = \frac{0.00739 \text{ mol}}{0.02500 \text{ L}} = 0.2956 \text{ M}$$

