

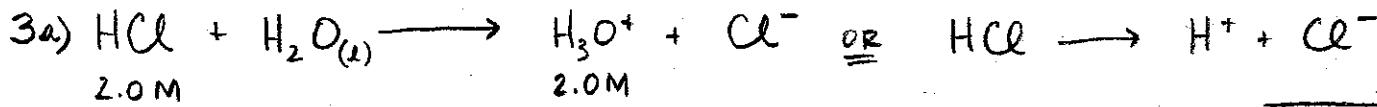
# Acid/Base I Review Key

Given data is underlined.

	$[H_3O^+]$	pH	POH	$[OH^-]$	Solution
a)	<u><math>3.0 \times 10^{-2} M</math></u>	1.52	12.48	<u><math>3.3 \times 10^{-13} M</math></u>	ACIDIC
b)	<u><math>4.6 \times 10^{-8} M</math></u>	7.34	<u><math>6.66</math></u>	<u><math>2.2 \times 10^{-7} M</math></u>	BASIC
c)	<u><math>1.9 \times 10^{-12} M</math></u>	<u>11.72</u>	<u>2.28</u>	<u><math>5.2 \times 10^{-3} M</math></u>	BASIC
d)	<u><math>5.9 \times 10^{-9} M</math></u>	8.23	5.77	<u><math>1.7 \times 10^{-6} M</math></u>	BASIC
e)	<u><math>8.0 \times 10^{-5} M</math></u>	4.10	9.90	<u><math>1.3 \times 10^{-10} M</math></u>	ACIDIC
f)	<u><math>1.0 \times 10^{-7} M</math></u>	7.00	7.00	<u><math>1.0 \times 10^{-7} M</math></u>	NEUTRAL
g)	<u><math>1.0 \times 10^{-3} M</math></u>	<u>3.00</u>	11.00	<u><math>1.0 \times 10^{-11} M</math></u>	ACIDIC
h)	10 M	-1.0	<u>15.0</u>	<u><math>1 \times 10^{-15} M</math></u>	ACIDIC
i)	<u><math>1.1 \times 10^{-12} M</math></u>	11.94	<u>2.06</u>	<u><math>8.7 \times 10^{-3} M</math></u>	BASIC
j)	<u><math>2.6 \times 10^{-6} M</math></u>	<u>5.58</u>	8.42	<u><math>3.8 \times 10^{-9} M</math></u>	ACIDIC

2. pH = 7.43  $[H_3O^+] = [H^+] = \text{inv log}(-7.43) = [3.7 \times 10^{-8} M]$

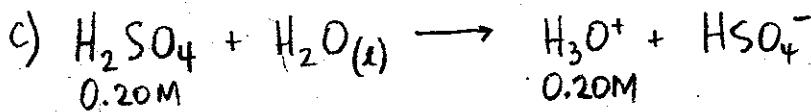
$$[OH^-] = \frac{K_w}{[H_3O^+]} = \frac{1.0 \times 10^{-14}}{3.7 \times 10^{-8} M} = [2.7 \times 10^{-7} M]$$



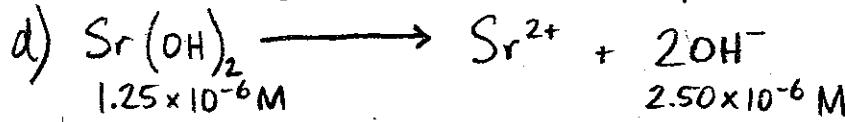
$$\text{pH} = -\log(2.0) = [-0.30] \quad \text{POH} = 14.00 - (-0.30) = [14.30]$$



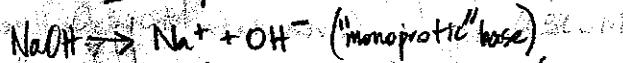
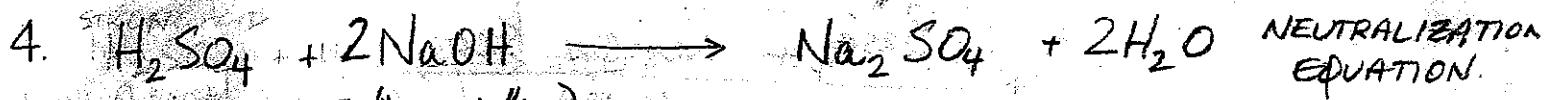
$$\text{POH} = -\log(3.5 \times 10^{-4} M) = [3.46] \quad \text{pH} = 14.00 - 3.46 = [10.54]$$



$$\text{pH} = -\log(0.20 M) = [0.70] \quad \text{POH} = 14.00 - 0.70 = [13.30]$$



$$\text{POH} = -\log(2.50 \times 10^{-6} M) = [5.602] \quad \text{pH} = 14.00 - 5.60 = [8.398]$$



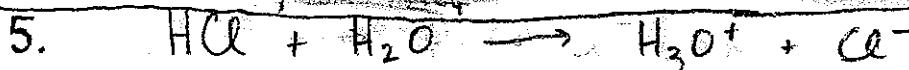
$$\text{mol H}_2\text{SO}_4 = MV = (0.500\text{M})(0.025\text{L}) = 0.0125 \text{ mol H}_2\text{SO}_4$$



$$0.0125 \text{ mol H}_2\text{SO}_4 \quad | \quad 2 \text{ mol H}_3\text{O}^+$$

$$1 \text{ mol H}_2\text{SO}_4 = 0.0250 \text{ mol H}_3\text{O}^+ = 0.0250 \text{ mol OH}^- = \text{mol NaOH}$$

$$0.0250 \text{ mol NaOH} \quad | \quad 40.0 \text{ g NaOH} \quad | \quad 1 \text{ mol NaOH} = \boxed{1.0 \text{ g NaOH}}$$



\*  $\text{H}_2\text{SO}_4$  will dissociate both protons in presence of strong Arrhenius base.

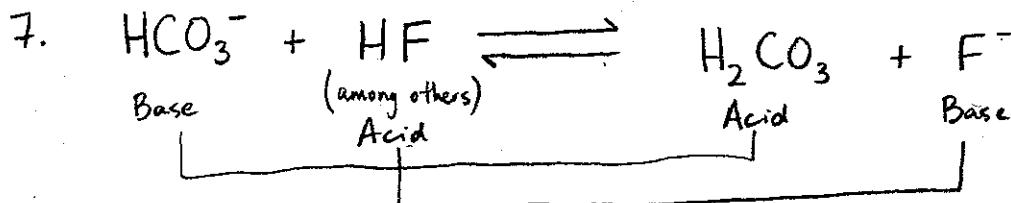
$$\text{mol HCl} = MV = (0.15\text{M})(0.02500\text{L}) = 0.00375 \text{ mol HCl} = 0.00375 \text{ mol H}_3\text{O}^+$$

$$[\text{H}_3\text{O}^+] = \frac{\text{mol}}{\text{V}} = \frac{0.00375 \text{ mol}}{0.100 \text{ L}} = 3.75 \times 10^{-2} \text{ M} \quad \text{OR} \quad M_f = \frac{M_i V_i}{V_f}$$

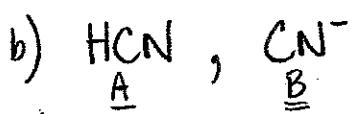
$$\text{pH} = -\log(3.75 \times 10^{-2} \text{ M}) = \boxed{1.43}$$

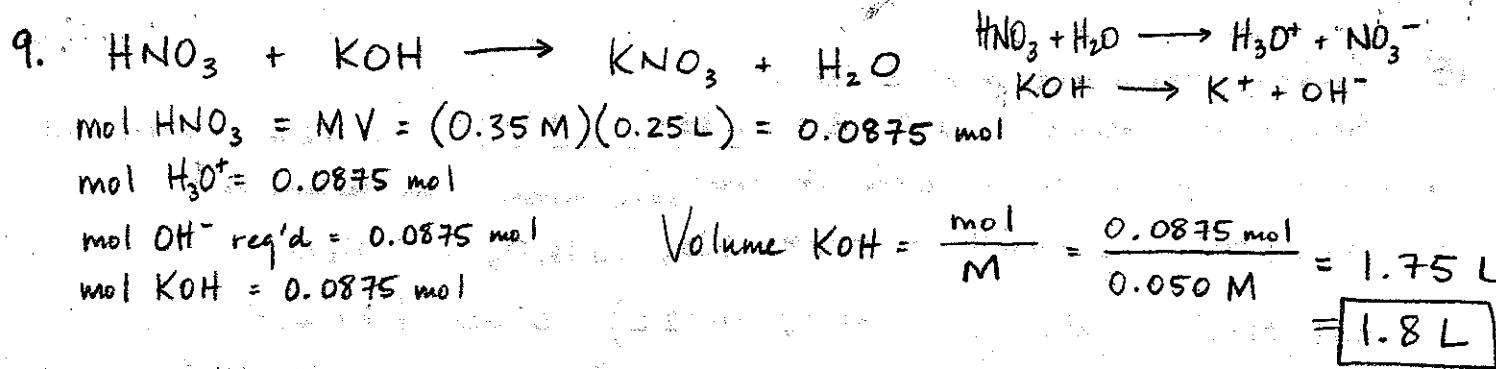
6. a)  $\text{H}_2\text{S}$  and  $\text{H}_2\text{Te}$  are the acids but since REACTANTS are favoured,  $\text{H}_2\text{Te}$  must be the stronger acid.

b) Since  $\text{H}_2\text{Te}$  is the stronger acid, its conjugate base must be weaker ( $\text{HTe}^-$ ).



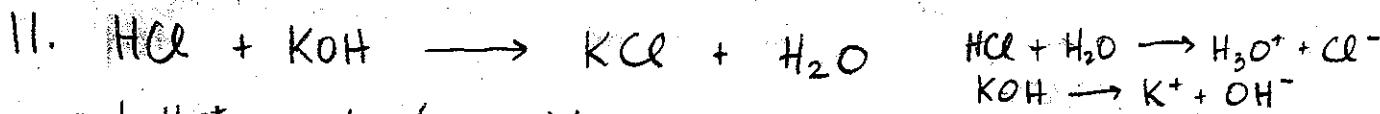
8. a) CONJUGATE BASE - The species  $\text{A}^-$  formed by loss of  $\text{H}^+$  from the acid  $\text{HA}$ .





10.  $[\text{H}_3\text{O}^+] = \text{inv log} (-4.20) = 6.3 \times 10^{-5} \text{ M}$

$$[\text{OH}^-] = \frac{K_w}{[\text{H}_3\text{O}^+]} = \frac{1.0 \times 10^{-14}}{6.3 \times 10^{-5}} = 1.6 \times 10^{-10} \text{ M}$$



mol  $\text{H}_3\text{O}^+ = MV = (0.150 \text{ M})(0.0600 \text{ L}) = 0.00900 \text{ mol H}_3\text{O}^+$

mol  $\text{OH}^- = MV = (0.100 \text{ M})(0.1400 \text{ L}) = 0.0140 \text{ mol OH}^-$

mol  $\text{OH}^- \text{ in excess} = 0.0140 \text{ mol OH}^- - 0.00900 \text{ mol H}_3\text{O}^+ = 0.0050 \text{ mol OH}^-$

$$M = \frac{\text{mol}}{V} = \frac{0.0050 \text{ mol}}{0.200 \text{ L}} = 0.025 \text{ M} \quad \text{pOH} = -\log(0.025 \text{ M}) = 1.60$$

$$\text{pH} = 12.40$$

12.a) Amphiprotic - A substance that can act as either an acid or a base.

b)  $\text{H}_2\text{PO}_4^-$ ,  $\text{HCO}_3^-$  etc...



b)  $\frac{1.00 \text{ g oxalic}}{90.0 \text{ g oxalic}} \left| \begin{array}{c} 1 \text{ mol oxalic} \\ \hline 90.0 \text{ g oxalic} \end{array} \right. = 0.01111 \text{ mol H}_2\text{C}_2\text{O}_4 \left| \begin{array}{c} 2 \text{ mol H}_3\text{O}^+ \\ \hline 1 \text{ mol H}_2\text{C}_2\text{O}_4 \end{array} \right. \xrightarrow{\text{strong base will dissociate and take both protons.}} 0.02222 \text{ mol H}_3\text{O}^+$

mol  $\text{OH}^- \text{ req'd} = 0.02222 \text{ mol OH}^- = 0.02222 \text{ mol NaOH}$

$$[\text{NaOH}] = \frac{\text{mol}}{V} = \frac{0.02222 \text{ mol}}{0.06000 \text{ L}} = 0.370 \text{ M}$$

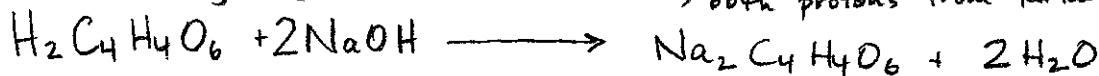
14.  $\text{inv log} (-6.51) = 3.090 \times 10^{-7} \text{ M} = [\text{H}_3\text{O}^+] = [\text{OH}^-] * \text{Water is NEUTRAL}$

$$K_w = [\text{H}_3\text{O}^+][\text{OH}^-] = (3.090 \times 10^{-7} \text{ M})^2 = 9.5 \times 10^{-14}$$

15. Firstly; throw out trial 2 data.

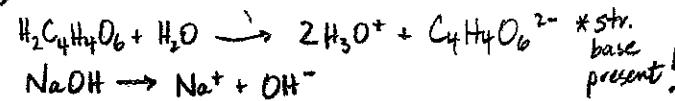
Secondly, average trial 1 + 3:  $\frac{11.33 \text{ mL} + 11.31 \text{ mL}}{2} = 11.32 \text{ mL NaOH added}$

\* NaOH is a strong base that will dissociate (take) both <sup>2</sup> protons from tartaric acid.



$$\text{mol NaOH} = MV = (0.104 \text{ M})(0.01132 \text{ L}) = 0.0011773 \text{ mol}$$

$$\text{mol OH}^- = 0.0011773 \text{ mol}$$



$$\text{mol H}_3\text{O}^{\text{req'd}} = 0.0011773 \text{ mol}$$

$$\text{mol H}_2\text{C}_4\text{H}_4\text{O}_6 = \frac{0.0011773 \text{ mol H}^+}{2 \text{ mol H}_3\text{O}^+} = 5.8864 \times 10^{-4} \text{ mol tartaric}$$

$$\frac{5.8864 \times 10^{-4} \text{ mol tartaric acid}}{1 \text{ mol tartaric acid}} \left| \begin{array}{l} 150.0 \text{ g tartaric acid} \\ 1 \text{ mol tartaric acid} \end{array} \right. = 8.8296 \times 10^{-2} \text{ g tartaric acid (PURE)}$$

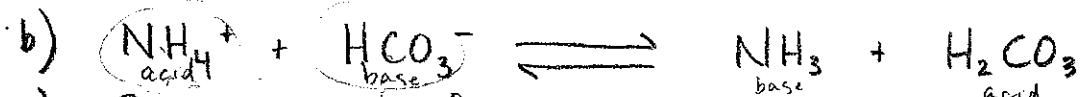
$$\frac{10.00 \text{ mL}}{1 \text{ mL}} \left| \begin{array}{l} 1.0 \text{ g} \\ 1 \text{ mL} \end{array} \right. = 10.0 \text{ g tartaric (IMPURE.)}$$

$$\% \text{ PURITY} = \frac{\text{PURE MASS}}{\text{IMPURE MASS}} = \frac{8.8296 \times 10^{-2} \text{ g}}{10.0 \text{ g}} = \boxed{0.88 \%} \quad (2 \text{ sig figs due to DENSITY value})$$

16. ① HBr (STRONG ACID = 100% dissociation).

② CH<sub>3</sub>COOH (higher than HCN in acid column)

③ HCN. → greater dissociation.



c) Reactants are favoured b/c H<sub>2</sub>CO<sub>3</sub> is a stronger acid than NH<sub>4</sub><sup>+</sup>.

18. \* H<sub>3</sub>O<sup>+</sup> is the DOMINANT species; focus calcs. on it!

$$\text{pH } 4.50: [\text{H}_3\text{O}^+] = \text{avlog}(-4.50) = 3.1623 \times 10^{-5} \text{ M H}_3\text{O}^+$$

$$\text{mol H}_3\text{O}^+ = MV = (3.1623 \times 10^{-5} \text{ M})(140 \text{ L}) = 4.427 \times 10^{-3} \text{ mol H}_3\text{O}^+$$

$$\text{pH } 6.80: [\text{H}_3\text{O}^+] = \text{avlog}(-6.80) = 1.5849 \times 10^{-7} \text{ M H}_3\text{O}^+$$

$$\text{mol H}_3\text{O}^+ = MV = (1.5849 \times 10^{-7} \text{ M})(140 \text{ L}) = 2.2189 \times 10^{-5} \text{ mol H}_3\text{O}^+$$

$$\text{mol OH}^{\text{req'd.}} = 4.427 \times 10^{-3} \text{ mol H}_3\text{O}^+ - 2.2189 \times 10^{-5} \text{ mol H}_3\text{O}^+$$

$$(\text{= mol H}_3\text{O}^+ \text{ added}) = 4.4048 \times 10^{-3} \text{ mol OH}^- \quad \text{DIPROTIC BASE}$$

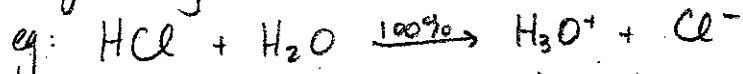


$$\frac{4.4028 \times 10^{-3} \text{ mol OH}^-}{2 \text{ mol OH}^-} \left| \begin{array}{l} 1 \text{ mol Ca(OH)}_2 \\ 74.1 \text{ g Ca(OH)}_2 \end{array} \right. = \boxed{1.6 \times 10^{-1} \text{ g Ca(OH)}_2}$$

$$= 0.16 \text{ g Ca(OH)}_2$$

19.  $\text{H}_3\text{O}^+$  is the strongest weak acid.

When any strong acid reacts w/ water, it dissociates 100% to form  $\text{H}_3\text{O}^+$ .



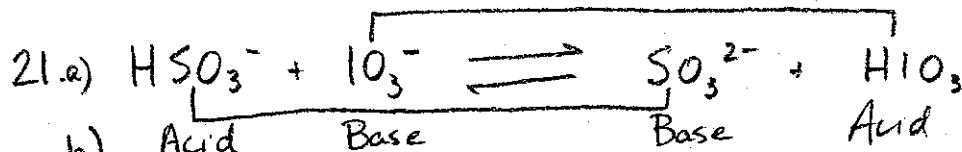
∴ no strong acids exist in water.

20.a)  $K_w = 2.95 \times 10^{-15} = [\text{H}_3\text{O}^+][\text{OH}^-] * [\text{H}_3\text{O}^+] = [\text{OH}^-] \Rightarrow \text{WATER!}$

$$2.95 \times 10^{-15} = [\text{H}_3\text{O}^+]^2$$

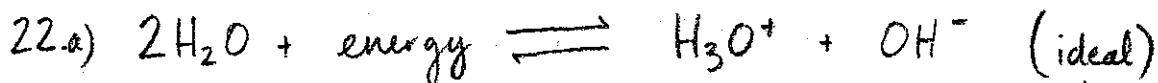
$$[\text{H}_3\text{O}^+] = 5.4314 \times 10^{-8} \text{ M} \quad \text{pH} = -\log(5.4314 \times 10^{-8} \text{ M}) \\ = 7.265$$

b) Water is NEUTRAL!  $[\text{H}_3\text{O}^+] = [\text{OH}^-]$

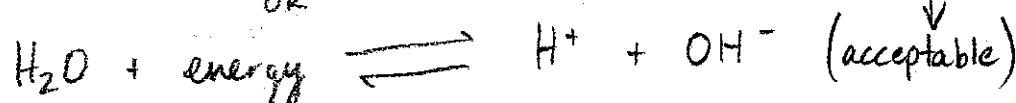


b) Acid Base Base Acid

c) Reactants are favoured b/c  $\text{HIO}_3$  is higher than  $\text{HSO}_3^-$  in acid column on table



OR



b)  $K_w = [\text{H}_3\text{O}^+][\text{OH}^-]$

c)  $K_w = 1.0 \times 10^{-14} \quad \text{p}K_w = 14$

d) If temp is increased,  $K_w$  increases and  $\text{p}K_w$  decreases.

23.a) Blood = 7-8 b) Rainwater = 5-7 c) Household cleaning products = 10-14

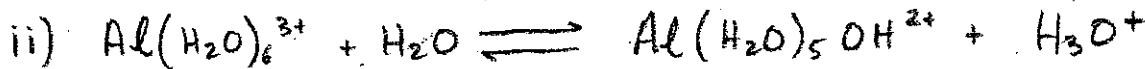
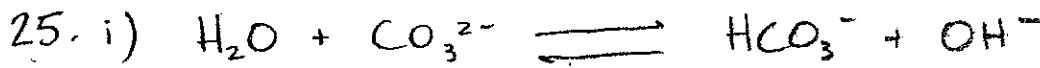
d) Battery Acid = 1-3 e) Coffee = 4-6

24.a) Arrhenius Acid - Any substance that releases  $\text{H}^+$  in water.

b) Arrhenius Base - Any substance that releases  $\text{OH}^-$  in water.

c) B-L Acid - A substance that donates a proton ( $\text{H}^+$ ) to another substance.

d) B-L Base - A substance that accepts a proton ( $\text{H}^+$ ) from another substance.



26. i) Measure electrical conductivity → Strong acid conducts better due to 100% dissociation.

ii) Measure the pH ⇒ Lower pH = Strong acid due to higher  $[\text{H}_3\text{O}^+]$ .