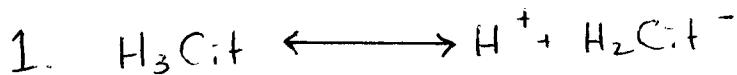


Acid #6

Answer key.



$$K_a = \frac{[\text{H}^+][\text{H}_2\text{Cit}^-]}{[\text{H}_3\text{Cit}]} = 8.4 \times 10^{-4}$$

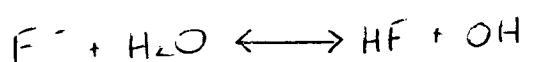
$$[\text{H}^+] = \text{antilog}(-2.10) = 7.94 \times 10^{-3} \text{ M}$$

$$[\text{H}_2\text{Cit}^-] = [\text{H}^+] = 7.94 \times 10^{-3} \text{ M}$$

$$[\text{H}_3\text{Cit}] = [\text{H}_3\text{Cit}]_{\text{initial}} - [\text{H}^+]$$

$$K_a = \frac{(7.94 \times 10^{-3})^2}{[\text{H}_3\text{Cit}]_{\text{initial}} - 7.94 \times 10^{-3}} = 8.4 \times 10^{-4}$$

$$[\text{H}_3\text{Cit}]_{\text{initial}} = \underline{\underline{8.3 \times 10^{-2} \text{ M}}}$$

2. NaF:  $\text{Na}^+$  does not hydrolyze

$$K_b = \frac{[\text{HF}][\text{OH}^-]}{[\text{F}^-]} = \frac{k_w}{k_c(\text{HF})} = \frac{1.0 \times 10^{-14}}{3.5 \times 10^{-4}} = 2.86 \times 10^{-11}$$

$$\text{let } x = [\text{OH}^-]$$

$$\text{then } [\text{HF}] = x$$

$$\text{and } [\text{F}^-] = 1.0 \text{ M} - x$$

$\approx 1.0 \text{ M}$  if  $x$  is small

$$2.86 \times 10^{-11} = \frac{x^2}{1.0}$$

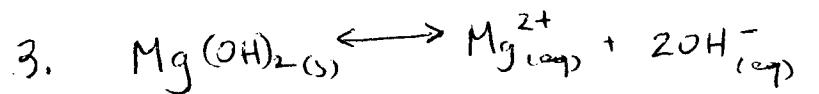
$$[\text{OH}^-] = x = 5.35 \times 10^{-6} \text{ M}$$

$$\text{pOH} = 5.272$$

$$\underline{\underline{\text{pH} = 8.73}}$$

Acid b cont'd.

Answer key:



$$K_{sp} = [Mg^{2+}][OH^-]^2 = 1.8 \times 10^{-11}$$

$$\text{let } [Mg^{2+}] = x$$

$$\text{then } [OH^-] = 2x$$

$$K_{sp} = 4x^3 = 1.8 \times 10^{-11}$$

$$x = 1.65 \times 10^{-4} M$$

$$[OH^-] = 2x = 3.30 \times 10^{-4} M$$

$$pOH = 3.481$$

$$\underline{pH = 10.52}$$



$$K_a = \frac{[H_3O^+][F^-]}{[HF]} = 3.5 \times 10^{-4}$$

$$[H_3O^+] = \text{anti-log}(-2.21) = 6.17 \times 10^{-3} M$$

$$[F^-] = [H_3O^+] = 6.17 \times 10^{-3} M$$

$$[HF] = \left(2.09/L \div 20 \text{ g/mol}\right) = 6.17 \times 10^{-3} M = 9.38 \times 10^{-2} M$$

$$K_a = \frac{(6.17 \times 10^{-3})^2}{9.38 \times 10^{-2}} = \underline{\underline{4.1 \times 10^{-4}}}$$

b) % dissociation =  $\frac{6.17 \times 10^{-3}}{(2.09/L \div 20 \text{ g/mol})} \times 100 = \underline{\underline{6.2\%}}$

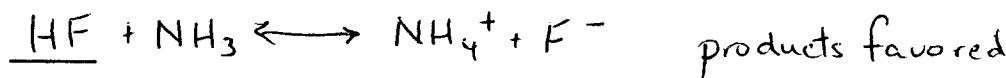
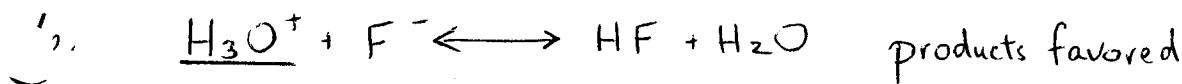
5. pH range 8.2 - 10.0

pOH range 5.8 - 4.0

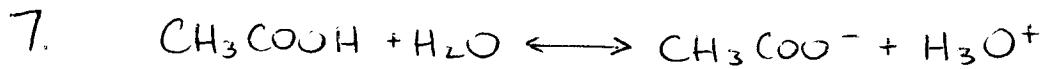
[OH<sup>-</sup>] range  $1.5 \times 10^{-6} M$  to  $1.0 \times 10^{-4} M$  a difference of  $9.85 \times 10^{-5} M$   
or  $1 \times 10^{-4} M$

Acid #6 cont'd

Answer key



Stronger acid underlined



$$K_a = \frac{[\text{CH}_3\text{COO}^-][\text{H}_3\text{O}^+]}{[\text{CH}_3\text{COOH}]} = 1.8 \times 10^{-5}$$

$$[\text{H}_3\text{O}^+] = \text{anti log } (-3.2) = 6.3 \times 10^{-4} \text{ M}$$

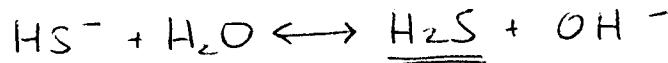
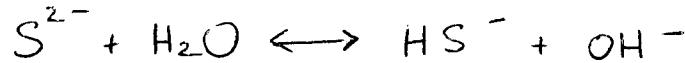
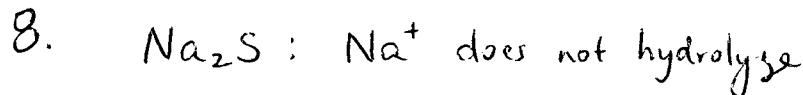
$$[\text{CH}_3\text{COO}^-] = [\text{H}_3\text{O}^+] = 6.3 \times 10^{-4} \text{ M}$$

$$[\text{CH}_3\text{COOH}] = [\text{CH}_3\text{COOH}]_{\text{initial}} - 6.3 \times 10^{-4} \text{ M}$$

$$1.8 \times 10^{-5} = \frac{(6.3 \times 10^{-4})^2}{[\text{CH}_3\text{COOH}]_{\text{ini}} - 6.3 \times 10^{-4}}$$

$$[\text{CH}_3\text{COOH}]_{\text{initial}} = 2.3 \times 10^{-2} \text{ M}$$

$$\text{mass} = 2.3 \times 10^{-2} \text{ mol/L} \times 60.0 \text{ g/mol} = \underline{1.4 \text{ g per litre}} = \underline{1 \text{ g/L}}$$



as  $\text{H}_2\text{S}$  gas escapes the reactions proceed in the forward direction to produce more  $\text{OH}^-$  (strongly basic) and  $\text{H}_2\text{S}$  gas (odour.)