

16.14 (a) According to the Arrhenius definition, a *base* when dissolved in water increases $[\text{OH}^-]$. According to the Brønsted-Lowry theory, a *base* is an H^+ acceptor regardless of physical state. A Brønsted-Lowry base is not limited to aqueous solution and need not contain OH^- or produce it in aqueous solution.

(b) $\text{NH}_3(\text{g}) + \text{H}_2\text{O}(\text{l}) \rightleftharpoons \text{NH}_4^+(\text{aq}) + \text{OH}^-(\text{aq})$ When NH_3 dissolves in water, it accepts H^+ from H_2O (B-L definition). In doing so, OH^- is produced (Arrhenius definition).

Note that the OH^- produced was originally part of the H_2O molecule, not part of the NH_3 molecule.

16.16 A conjugate acid has one more H^+ than its conjugate base.

- (a) HCN (b) OH^- (c) H_2PO_4^- (d) $\text{C}_2\text{H}_5\text{NH}_3^+$

	<u>B-L acid</u>	+	<u>B-L base</u>	\rightleftharpoons	<u>Conjugate acid</u>	+	<u>Conjugate base</u>
(a)	$\text{HBrO}(\text{aq})$		$\text{H}_2\text{O}(\text{l})$		$\text{H}_3\text{O}^+(\text{aq})$		$\text{BrO}^-(\text{aq})$
(b)	$\text{HSO}_4^-(\text{aq})$		$\text{HCO}_3^-(\text{aq})$		$\text{H}_2\text{CO}_3(\text{aq})$		$\text{SO}_4^{2-}(\text{aq})$
(c)	$\text{H}_3\text{O}^+(\text{aq})$		$\text{HSO}_3^-(\text{aq})$		$\text{H}_2\text{SO}_3(\text{aq})$		$\text{H}_2\text{O}(\text{l})$

- 16.22 (a) $C_2H_3O_2^-$, weak base; $HC_2H_3O_2$, weak acid
 (b) HCO_3^- , weak base; H_2CO_3 , weak acid
 (c) O_2^{2-} , strong base; OH^- , negligible acid
 (d) Cl^- , negligible base; HCl , strong acid
 (e) NH_3 , weak base; NH_4^+ , weak acid

16.44 For a strong acid, which is completely ionized, $[H^+] =$ the initial acid concentration.

(a) $0.00835 M HNO_3 = 0.00835 M H^+$; $pH = -\log(0.00835) = 2.08$

(b) $\frac{0.525 \text{ g } HClO_4}{2.00 \text{ L soln}} \times \frac{1 \text{ mol } HClO_4}{100.5 \text{ g } HClO_4} = 2.612 \times 10^{-3} = 2.61 \times 10^{-3} M HClO_4$

$[H^+] = 2.61 \times 10^{-3} M$; $pH = -\log(2.61 \times 10^{-3}) = 2.583$

(c) $M_c \times V_c = M_d \times V_d$; $0.500 \text{ L} = 500 \text{ mL}$

$1.00 M HCl \times 5.00 \text{ mL HCl} = M_d HCl \times 500 \text{ mL HCl}$

$M_d HCl = \frac{1.00 M \times 5.00 \text{ mL}}{500 \text{ mL}} = 1.00 \times 10^{-2} M HCl = 1.00 \times 10^{-2} M H^+$

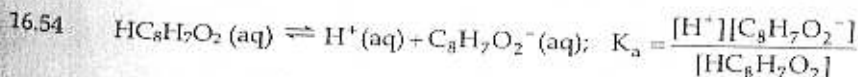
$pH = -\log(1.00 \times 10^{-2}) = 2.000$

(d) $[H^+]_{\text{total}} = \frac{\text{mol } H^+ \text{ from HCl} + \text{mol } H^+ \text{ from HI}}{\text{total L solution}}$; $\text{mol} = M \times L$

$[H^+]_{\text{total}} = \frac{(0.020 M HCl \times 0.0500 L) + (0.010 M HI \times 0.150 L)}{0.200 L}$

$[H^+]_{\text{total}} = \frac{1.0 \times 10^{-3} \text{ mol } H^+ + 1.50 \times 10^{-3} \text{ mol } H^+}{0.200 L} = 0.0125 = 0.013 M$

$pH = -\log(0.0125) = 1.90$



$[H^+] = [C_8H_7O_2^-] = 10^{-2.68} = 2.09 \times 10^{-3} = 2.1 \times 10^{-3} M$

$[HC_8H_7O_2] = 0.085 - 2.09 \times 10^{-3} = 0.0829 = 0.083 M$

$K_a = \frac{(2.09 \times 10^{-3})^2}{0.0829} = 5.3 \times 10^{-5}$

16.60



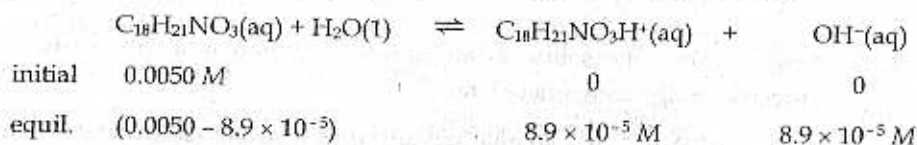
$$K_a = \frac{[\text{H}^+][\text{ClO}^-]}{[\text{HClO}]} = \frac{x^2}{(0.0075 - x)} \approx \frac{x^2}{0.0075} = 3.0 \times 10^{-5}$$

$$x^2 = 0.0075 (3.0 \times 10^{-5}); x = 1.5 \times 10^{-5} \text{ M} = [\text{H}^+] = [\text{H}_3\text{O}^+] = [\text{ClO}^-]$$

$$[\text{HClO}] = 7.5 \times 10^{-3} - 1.5 \times 10^{-5} = 7.485 \times 10^{-3} = 7.5 \times 10^{-3} \text{ M}$$

Check. $\frac{4.7 \times 10^{-5} \text{ M H}^+}{0.0075 \text{ M HClO}} \times 100 = 0.20\%$ ionization; the assumption is valid

16.78 (a) $\text{pOH} = 14.00 - 9.95 = 4.05$; $[\text{OH}^-] = 10^{-4.05} = 8.91 \times 10^{-5} = 8.9 \times 10^{-5} \text{ M}$



$$K_b = \frac{[\text{C}_{18}\text{H}_{21}\text{NO}_3\text{H}^+][\text{OH}^-]}{[\text{C}_{18}\text{H}_{21}\text{NO}_3]} = \frac{(8.91 \times 10^{-5})^2}{(0.0050 - 8.91 \times 10^{-5})} = 1.62 \times 10^{-5} = 1.6 \times 10^{-6}$$

(b) $\text{p}K_b = -\log(K_b) = -\log(1.62 \times 10^{-5}) = 5.79$

- 16.86 (a) acidic; Cr^{3+} is a highly charged metal cation and a Lewis acid; Br^- is negligible.
 (b) neutral; both Li^+ and I^- are negligible.
 (c) basic; PO_4^{3-} is the conjugate base of HPO_4^{2-} ; K^+ is negligible.
 (d) acidic; CH_3NH_3^+ is the conjugate acid of CH_3NH_2 ; Cl^- is negligible.
 (e) acidic; HSO_4^- is a negligible base, but a fairly strong acid ($K_a = 1.2 \times 10^{-2}$).
 K^+ is negligible.

16.96

- (a) NO_2^- (HNO_3 is the stronger acid because it has more nonprotonated O atoms, so NO_2^- is the stronger base.)
 (b) PO_4^{3-} (K_a for HAsO_4^{2-} is greater than K_a for HPO_4^{2-} , so K_b for PO_4^{3-} is greater and PO_4^{3-} is the stronger base. Note that P is more electronegative than As and H_3PO_4 is a stronger acid than H_3AsO_4 , which could lead to the conclusion that AsO_4^{3-} is the stronger base. As in all cases, the measurement of base strength, K_b , supercedes the prediction. Chemistry is an experimental science.)
 (c) CO_3^{2-} (The more negative the anion, the stronger the attraction for H^+ .)