

(45)

$$c. [\text{OH}^-] = \frac{1.0 \times 10^{-14}}{12} = 8.3 \times 10^{-16} \text{ M}; \text{ the solution is acidic.}$$

$$d. [\text{OH}^-] = \frac{1.0 \times 10^{-14}}{5.4 \times 10^{-5}} = 1.9 \times 10^{-10} \text{ M}; \text{ the solution is acidic.}$$

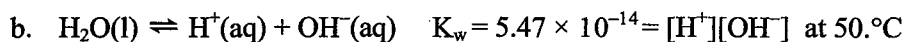
$$45 \rightarrow [\text{OH}^-] = \frac{1.0 \times 10^{-14}}{10 \times 10^{-7}} = 10^{-7} \text{ M}$$

$$b) [\text{OH}^-] = \frac{10^{-14}}{5.3 \times 10^{-10}} = 1.9 \times 10^{-5} \text{ M}$$

Basic

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a. Because the value of the equilibrium constant increases as the temperature increases, the reaction is endothermic. In endothermic reactions, heat is a reactant, so an increase in temperature (heat) shifts the reaction to produce more products and increases K in the process.



In pure water $[\text{H}^+] = [\text{OH}^-]$, so $5.47 \times 10^{-14} = [\text{H}^+]^2$, $[\text{H}^+] = 2.34 \times 10^{-7} \text{ M} = [\text{OH}^-]$

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$\text{pH} = -\log[\text{H}^+]$; $\text{pOH} = -\log[\text{OH}^-]$; at 25°C , $\text{pH} + \text{pOH} = 14.00$; for Exercise 45:

$$a. \text{pH} = -\log[\text{H}^+] = -\log(1.0 \times 10^{-7}) = 7.00; \text{pOH} = 14.00 - \text{pH} = 14.00 - 7.00 = 7.00$$

$$b. \text{pH} = -\log(8.3 \times 10^{-16}) = 15.08; \text{pOH} = 14.00 - 15.08 = -1.08$$

$$c. \text{pH} = -\log(12) = -1.08; \text{pOH} = 14.00 - (-1.08) = 15.08$$

$$d. \text{pH} = -\log(5.4 \times 10^{-5}) = 4.27; \text{pOH} = 14.00 - 4.27 = 9.73$$

Note that pH is less than zero when $[\text{H}^+]$ is greater than 1.0 M (an extremely acidic solution). For Exercise 46:

$$a. \text{pOH} = -\log[\text{OH}^-] = -\log(1.5) = -0.18; \text{pH} = 14.00 - \text{pOH} = 14.00 - (-0.18) = 14.18$$

$$b. \text{pOH} = -\log(3.6 \times 10^{-15}) = 14.44; \text{pH} = 14.00 - 14.44 = -0.44$$

$$c. \text{pOH} = -\log(1.0 \times 10^{-7}) = 7.00; \text{pH} = 14.00 - 7.00 = 7.00$$

$$d. \text{pOH} = -\log(7.3 \times 10^{-4}) = 3.14; \text{pH} = 14.00 - 3.14 = 10.86$$

Note that pH is greater than 14.00 when $[\text{OH}^-]$ is greater than 1.0 M (an extremely basic solution).

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$$a. \text{pOH} = 14.00 - 6.88 = 7.12; [\text{H}^+] = 10^{-6.88} = 1.3 \times 10^{-7} \text{ M}$$

$$[\text{OH}^-] = 10^{-7.12} = 7.6 \times 10^{-8} \text{ M}; \text{ acidic}$$

$$b. [\text{H}^+] = \frac{1.0 \times 10^{-14}}{8.4 \times 10^{-14}} = 0.12 \text{ M}; \text{pH} = -\log(0.12) = 0.92$$

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$$\text{pOH} = 14.00 - 0.92 = 13.08; \text{ acidic}$$

$$\text{c. } \text{pH} = 14.00 - 3.11 = 10.89; [\text{H}^+] = 10^{-10.89} = 1.3 \times 10^{-11} \text{ M}$$

$$[\text{OH}^-] = 10^{-3.11} = 7.8 \times 10^{-4} \text{ M}; \text{ basic}$$

$$\text{d. } \text{pH} = -\log(1.0 \times 10^{-7}) = 7.00; \text{ pOH} = 14.00 - 7.00 = 7.00$$

$$[\text{OH}^-] = 10^{-7.00} = 1.0 \times 10^{-7} \text{ M}; \text{ neutral}$$

53.

$$\text{pOH} = 14.0 - \text{pH} = 14.0 - 2.1 = 11.9; [\text{H}^+] = 10^{-\text{pH}} = 10^{-2.1} = 8 \times 10^{-3} \text{ M (1 sig. fig.)}$$

$$[\text{OH}^-] = \frac{K_w}{[\text{H}^+]} = \frac{1.0 \times 10^{-14}}{8 \times 10^{-3}} = 1 \times 10^{-12} \text{ M or } [\text{OH}^-] = 10^{-\text{pOH}} = 10^{-11.9} = 1 \times 10^{-12} \text{ M}$$

The sample of gastric juice is acidic because the pH is less than 7.00 at 25°C.

Solutions of Acids

55. All the acids in this problem are strong acids that are always assumed to completely dissociate in water. The general dissociation reaction for a strong acid is $\text{HA}(\text{aq}) \rightarrow \text{H}^+(\text{aq}) + \text{A}^-(\text{aq})$, where A^- is the conjugate base of the strong acid HA . For 0.250 M solutions of these strong acids, 0.250 M H^+ and 0.250 M A^- are present when the acids completely dissociate. The amount of H^+ donated from water will be insignificant in this problem since H_2O is a very weak acid.

- a. Major species present after dissociation = H^+ , ClO_4^- , and H_2O ;

$$\text{pH} = -\log[\text{H}^+] = -\log(0.250) = 0.602$$

- b. Major species = H^+ , NO_3^- , and H_2O ; $\text{pH} = 0.602$

57. Strong acids are assumed to completely dissociate in water; for example; $\text{HCl}(\text{aq}) + \text{H}_2\text{O}(\text{l}) \rightarrow \text{H}_3\text{O}^+(\text{aq}) + \text{Cl}^-(\text{aq})$ or $\text{HCl}(\text{aq}) \rightarrow \text{H}^+(\text{aq}) + \text{Cl}^-(\text{aq})$.

- a. A 0.10 M HCl solution gives 0.10 M H^+ and 0.10 M Cl^- because HCl completely dissociates. The amount of H^+ from H_2O will be insignificant.

$$\text{pH} = -\log[\text{H}^+] = -\log(0.10) = 1.00$$

- b. 5.0 M H^+ is produced when 5.0 M HClO_4 completely dissociates. The amount of H^+ from H_2O will be insignificant. $\text{pH} = -\log(5.0) = -0.70$ (Negative pH values just indicate very concentrated acid solutions.)

- c. 1.0×10^{-11} M H^+ is produced when 1.0×10^{-11} M HI completely dissociates. If you take the negative log of 1.0×10^{-11} , this gives $\text{pH} = 11.00$. This is impossible! We dissolved an acid in water and got a basic pH. What we must consider in this problem is that water

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14.3c

$$46) a) [H^+] = \frac{10^{-14}}{1.5 \text{ M}} = 6.67 \times 10^{-15} \text{ M} \quad \text{Basic}$$

$$b) [H^+] = \frac{10^{-14}}{3.6 \times 10^{-15} \text{ M}} = 2.78 \text{ M} \quad \text{Acidic}$$

$$c) [H^+] = \frac{10^{-14}}{1.0 \times 10^{-7} \text{ M}} = 10^{-7} \text{ M} \quad \text{Neutral}$$

$$d) [H^+] = \frac{10^{-14} \text{ M}}{7.3 \times 10^{-4} \text{ M}} = 1.37 \times 10^{-11} \text{ M} \quad \text{Basic}$$

$$48) K_w = 2.92 \times 10^{-14} = [H^+][OH^-]$$

$$a) [H^+] = [OH^-] = \sqrt{2.92 \times 10^{-14}} = 1.71 \times 10^{-7} \text{ M}$$

$$b) \underline{6.77} = -\log 1.71 \times 10^{-7} \text{ M} = \text{pH}$$

$$c) [OH^-] = 0.1 \text{ M} \quad \frac{2.92 \times 10^{-14}}{(0.1 \text{ M})} = [H^+]$$

$$[H^+] = 2.92 \times 10^{-13} \text{ M}$$

$$\text{pH} = -\log 2.92 \times 10^{-13}$$

$$\underline{\text{pH} = 12.5}$$