

Non Sibi High School

Andover's Chem 550/580: Advanced Chemistry

Chapter 16, Review Quiz 1 Answers

1

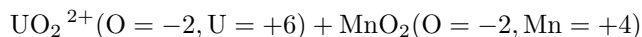
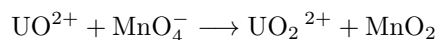
Determine all oxidation numbers in:

- a. As^{3-}
- b. F_2
- c. HO_2^-
- d. IO_4^-
- e. KH

- a. As = -3
- b. F = 0
- c. H = +1, O = -1
- d. O = -2, I = +7
- e. K = +1, H = -1

2

Determine all oxidation numbers, identify the reducing agent and oxidizing agent, and balance the following equation that occurs in aqueous acidic solution using the smallest possible whole-number coefficients:



reducing agent = UO^{2+} (U increases from +4 to +6)

oxidizing agent = MnO_4^- (Mn decreases from +7 to +4)

oxidation: $(\text{UO}^{2+} + \text{H}_2\text{O} \longrightarrow \text{UO}_2^{2+} + 2\text{H}^+ + 2\text{e}^-) \times 3$

reduction: $(\text{MnO}_4^- + 4\text{H}^+ + 3\text{e}^- \longrightarrow \text{MnO}_2 + 2\text{H}_2\text{O}) \times 2$

balanced equation: $3\text{UO}^{2+} + 2\text{MnO}_4^- + 2\text{H}^+ \longrightarrow 3\text{UO}_2^{2+} + 2\text{MnO}_2 + \text{H}_2\text{O}$

3

Rank the solid alkali metals K, Li, and Na from weakest to strongest reducing agent under standard conditions. Justify your answer using a table of standard reduction potentials.

$$\text{weakest} = \text{Na}(+2.71 \text{ V}) < \text{K}(+2.92 \text{ V}) < \text{Li}(+3.05 \text{ V}) = \text{strongest}$$

4

Rank the aqueous cations Ag^+ , Al^{3+} , and Cd^{2+} from weakest to strongest oxidizing agent under standard conditions. Justify your answer using a table of standard reduction potentials.

$$\text{weakest} = \text{Al}^{3+}(-1.66 \text{ V}) < \text{Cd}^{2+}(-0.40 \text{ V}) < \text{Ag}^+(+0.80 \text{ V}) = \text{strongest}$$

5

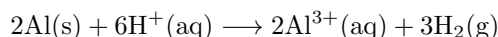
For each spontaneous reaction below, calculate E_{cell}° and then balance the equation.

- chlorine gas + aqueous potassium bromide
- solid aluminum metal + aqueous hydrochloric acid
- solid gold metal + aqueous nitric acid
- solid zinc metal + aqueous cadmium(II) nitrate

a. $E_{\text{cell}}^{\circ} = (+1.36 \text{ V}) + (-1.07 \text{ V}) = 0.29 \text{ V}$

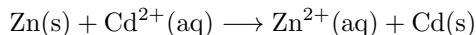


b. $E_{\text{cell}}^{\circ} = (+1.66 \text{ V}) + (0 \text{ V}) = 1.66 \text{ V}$



c. nonspontaneous

d. $E_{\text{cell}}^{\circ} = (+0.76 \text{ V}) + (-0.40 \text{ V}) = 0.36 \text{ V}$



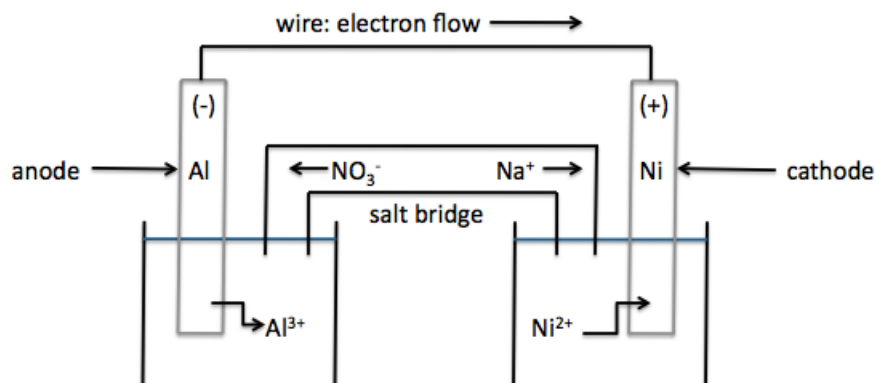
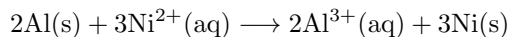
6

A galvanic cell was constructed using a strip of nickel metal and a strip of aluminum metal, a 1 M solution of NiSO_4 and a 1 M solution of $\text{Al}(\text{NO}_3)_3$, and an aqueous solution of NaNO_3 in the salt bridge. For the spontaneous reaction

that occurred, calculate E_{cell}° and ΔG° , then balance the equation. Also sketch the galvanic cell.

$$E_{\text{cell}}^{\circ} = (+1.66 \text{ V}) + (-0.25 \text{ V}) = 1.41 \text{ V}$$

$$\Delta G^{\circ} = -6 \left(\frac{96,500 \text{ C}}{1 \text{ mol}} \right) \left(\frac{1.41 \text{ J}}{1 \text{ C}} \right) = -8.16 \times 10^5 \text{ J/mol} = -816 \text{ kJ/mol}$$

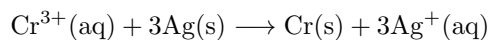


7

Calculate the minimum voltage required to bring about the reaction $\text{Cr}^{3+}(\text{aq}) + \text{Ag(s)} \longrightarrow \text{Cr(s)} + \text{Ag}^{+}(\text{aq})$ by electrolysis under standard conditions, then balance the equation.

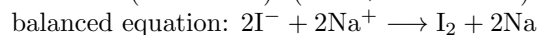
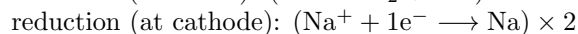
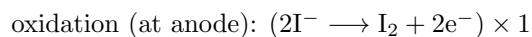
$$E_{\text{cell}}^{\circ} = (-0.74 \text{ V}) + (-0.80 \text{ V}) = -1.54 \text{ V}$$

$$\text{minimum voltage} = |E_{\text{cell}}^{\circ}| = 1.54 \text{ V}$$



8

For the electrolysis of molten NaI , write the half-reaction that occurs at the anode and the half-reaction that occurs at the cathode, then balance the equation.

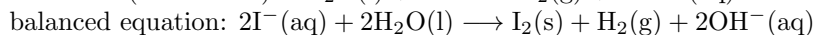
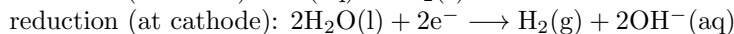
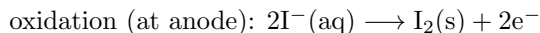


9

a. For the electrolysis of aqueous NaI, write the half-reaction that occurs at the anode and the half-reaction that occurs at the cathode, then balance the equation.

b. If the electrolysis proceeds for 2.65 days using a current of 3.75 A, how many kilograms of I_2 will be produced?

a. $Na^+(aq)$ = electrolysis spectator ion, $E_{ox}^\circ = -0.53$ V for the oxidation of $I^-(aq)$ is less negative than E_{ox}° for the oxidation of water:



b. oxidation (at anode): $2I^-(aq) \longrightarrow I_2(s) + 2e^-$

$$2.65 \text{ d} \left(\frac{24 \text{ h}}{1 \text{ d}} \right) \left(\frac{60 \text{ min}}{1 \text{ h}} \right) \left(\frac{60 \text{ s}}{1 \text{ min}} \right) \left(\frac{3.75 \text{ C}}{1 \text{ s}} \right) \left(\frac{1 \text{ mol } e^-}{96,500 \text{ C}} \right) \left(\frac{1 \text{ mol } I_2}{2 \text{ mol } e^-} \right) \left(\frac{253.8 \text{ g}}{1 \text{ mol } I_2} \right) \left(\frac{1 \text{ kg}}{1000 \text{ g}} \right) = 1.13 \text{ kg}$$

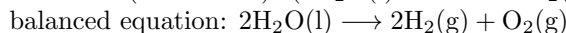
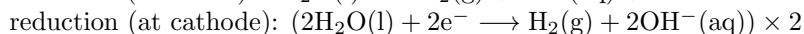
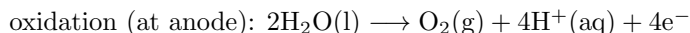
10

a. For the electrolysis of aqueous $LiNO_3$, write the half-reaction that occurs at the anode and the half-reaction that occurs at the cathode, then balance the equation.

b. Using a current of 0.285 A, how many hours must the electrolysis proceed to produce 444 mL of oxygen gas, measured at 28°C and 724 torr?

c. If 40.0 milligrams of hydrogen gas are produced when the electrolysis proceeds for 18.5 minutes, calculate the current used in the electrolysis.

a. $Li^+(aq)$ and $NO_3^-(aq)$ = electrolysis spectator ions:



b. oxidation (at anode): $2H_2O(l) \longrightarrow O_2(g) + 4H^+(aq) + 4e^-$

$$n = \frac{\frac{724}{760} \text{ atm} \times \frac{444}{1000} \text{ L}}{0.0821 \frac{\text{L}\cdot\text{atm}}{\text{mol}\cdot\text{K}} \times (28 + 273) \text{ K}} = 0.01712 \text{ mol } O_2$$

$$0.01712 \text{ mol } O_2 \left(\frac{4 \text{ mol } e^-}{1 \text{ mol } O_2} \right) \left(\frac{96,500 \text{ C}}{1 \text{ mol } e^-} \right) \left(\frac{1 \text{ s}}{0.285 \text{ C}} \right) \left(\frac{1 \text{ min}}{60 \text{ s}} \right) \left(\frac{1 \text{ h}}{60 \text{ min}} \right) = 6.44 \text{ h}$$

c. reduction (at cathode): $2\text{H}_2\text{O}(l) + 2e^- \longrightarrow \text{H}_2(g) + 2\text{OH}^-(aq)$

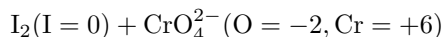
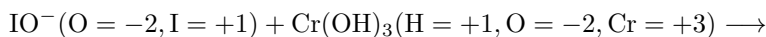
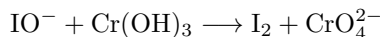
$$18.5 \text{ min} \left(\frac{60 \text{ s}}{1 \text{ min}} \right) = 1110 \text{ s}$$

$$40.0 \text{ mg} \left(\frac{1 \text{ g}}{1000 \text{ mg}} \right) \left(\frac{1 \text{ mol H}_2}{2.016 \text{ g}} \right) \left(\frac{2 \text{ mol e}^-}{1 \text{ mol H}_2} \right) \left(\frac{96,500 \text{ C}}{1 \text{ mol e}^-} \right) = 3829 \text{ C}$$

$$\text{current} = \frac{3829 \text{ C}}{1110 \text{ s}} = 3.45 \text{ C/s} = 3.45 \text{ A}$$

11

Determine all oxidation numbers, identify the reducing agent and oxidizing agent, and balance the following equation that occurs in aqueous basic solution using the smallest possible whole-number coefficients:



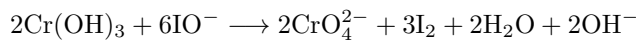
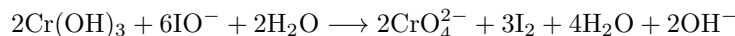
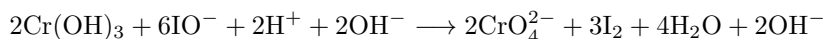
reducing agent = $\text{Cr}(\text{OH})_3$ (U increases from +3 to +6)

oxidizing agent = IO^- (I decreases from +1 to 0)

oxidation: $(\text{Cr}(\text{OH})_3 + \text{H}_2\text{O} \longrightarrow \text{CrO}_4^{2-} + 5\text{H}^+ + 3e^-) \times 2$

reduction: $(\text{IO}^- + 4\text{H}^+ + 2e^- \longrightarrow \text{I}_2 + 2\text{H}_2\text{O}) \times 3$

balanced equation (acidic): $2\text{Cr}(\text{OH})_3 + 6\text{IO}^- + 2\text{H}^+ \longrightarrow 2\text{CrO}_4^{2-} + 3\text{I}_2 + 4\text{H}_2\text{O}$



12

Calculate K_c for the reaction $\text{Fe}^{2+}(aq) + \text{Cd}(s) \longrightarrow \text{Cd}^{2+}(aq) + \text{Fe}(s)$ at 25°C .

$$E_{\text{cell}}^\circ = (-0.44 \text{ V}) + 0.40 \text{ V} = -0.04 \text{ V}$$

$$K_{\text{eq}} = K_c :$$

$$-0.04 \text{ V} = \frac{0.0257 \text{ V}}{2} \ln K_c$$

$$K_c = 0.04$$

13

A galvanic cell is constructed at 25°C that utilizes the reaction $2\text{Fe}^{3+}(\text{aq}) + \text{Cu}(\text{s}) \longrightarrow 2\text{Fe}^{2+}(\text{aq}) + \text{Cu}^{2+}(\text{aq})$.

a. If all aqueous ions in the cell have an initial concentration of 1 M, calculate E_{cell}° .

b. Predict whether E_{cell} will increase, decrease, or remain unchanged if the initial concentration of Fe^{3+} is 1 M, but the initial concentrations of Fe^{2+} and Cu^{2+} are both changed to 0.075 M.

c. Calculate E_{cell} if the initial concentrations of all aqueous ions are 0.025 M.

a. $E_{\text{cell}}^{\circ} = 0.77 \text{ V} + (-0.34 \text{ V}) = 0.43 \text{ V}$

b. decrease product concentrations only = Q_c decreases = E_{cell} increases

c.

$$E_{\text{cell}} = 0.43 \text{ V} - \frac{0.0257 \text{ V}}{2} \ln \frac{(0.025)^2(0.025)}{(0.025)^2} = 0.48 \text{ V}$$



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