Non Sibi High School

Andover's Chem 300: Accelerated/Honors 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:

$$\begin{split} & UO^{2+} + MnO_4^- \longrightarrow UO_2^{\ 2+} + MnO_2 \\ & UO^{2+}(O=-2,U=+4) + MnO_4^-(O=-2,Mn=+7) \longrightarrow \\ & UO_2^{\ 2+}(O=-2,U=+6) + MnO_2(O=-2,Mn=+4) \end{split}$$

reducing agent = $\rm UO^{2+}$ (U increases from +4 to +6) oxidizing agent = $\rm MnO_4$ $^-$ (Mn decreases from +7 to +4)

oxidation: (UO²⁺ + H₂O
$$\longrightarrow$$
 UO₂ ²⁺ + 2H⁺ + 2e⁻) × 3 reduction: (MnO₄ ⁻ + 4H⁺ + 3e⁻ \longrightarrow MnO₂ + 2H₂O) × 2 balanced equation: 3UO²⁺ + 2MnO₄ ⁻ + 2H⁺ \longrightarrow 3UO₂ ²⁺ + 2MnO₂ + H₂O

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.

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

4

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

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

5

For each spontaneous reaction below, calculate $E_{\rm cell}^{\circ}$ and then balance the equation.

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

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

 $Cl_2(g) + 2Br^-(aq) \longrightarrow 2Cl^-(aq) + Br_2(l)$

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

$$2Al(s) + 6H^{+}(aq) \longrightarrow 2Al^{3+}(aq) + 3H_{2}(g)$$

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

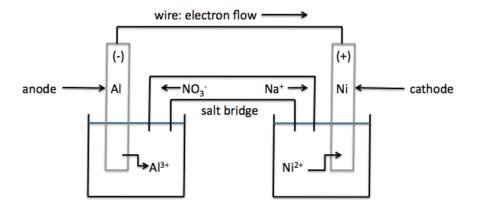
$$\operatorname{Zn}(s) + \operatorname{Cd}^{2+}(\operatorname{aq}) \longrightarrow \operatorname{Zn}^{2+}(\operatorname{aq}) + \operatorname{Cd}(s)$$

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 $Al(NO_3)_3$, 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.

$$\begin{split} E_{cell}^{\circ} &= (+1.66\,V) + (-0.25\,V) = 1.41\,V \\ \Delta G^{\circ} &= -6\left(\frac{96,500\,C}{1\,mol}\right)\left(\frac{1.41\,J}{1\,C}\right) = -8.16\times10^5\,J/mol = -816\,kJ/mol \\ &\quad 2Al(s) + 3Ni^{2+}(aq) \longrightarrow 2Al^{3+}(aq) + 3Ni(s) \end{split}$$



7

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

$$\begin{split} E_{cell}^\circ &= (-0.74\,V) + (-0.80\,V) = -1.54\,V \\ \\ minimum voltage &= |E_{cell}^\circ| = 1.54\,V \\ \\ Cr^{3+}(aq) + 3Ag(s) &\longrightarrow Cr(s) + 3Ag^+(aq) \end{split}$$

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.

oxidation (at anode): $(2I^- \longrightarrow I_2 + 2e^-) \times 1$ reduction (at cathode): $(Na^+ + 1e^- \longrightarrow Na) \times 2$ balanced equation: $2I^- + 2Na^+ \longrightarrow I_2 + 2Na$ 9

For the electrolysis of molten NaI, if the electrolysis proceeds for 2.65 days using a current of 3.75 A, how many kilograms of I_2 will be produced?

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

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

10

For the electrolysis of molten KCl, using a current of 0.285 A, how many hours must the electrolysis proceed to produce 888 mL of $\rm Cl_2$ gas, measured at 28°C and 724 torr?

oxidation (at anode): $2Cl^- \longrightarrow Cl_2 + 2e^-$

$$n = \frac{\frac{724}{760} \operatorname{atm} \times \frac{888}{1000} L}{0.0821 \frac{L \cdot \operatorname{atm}}{\operatorname{mol \cdot K}} \times (28 + 273) \, K} = 0.03423 \, \mathrm{mol \, Cl_2}$$

$$0.03423\,\mathrm{mol}\,\mathrm{Cl_2}\left(\frac{2\,\mathrm{mol}\,\mathrm{e^-}}{1\,\mathrm{mol}\,\mathrm{Cl_2}}\right)\left(\frac{96,500\,\mathrm{C}}{1\,\mathrm{mol}\,\mathrm{e^-}}\right)\left(\frac{1\,\mathrm{s}}{0.285\,\mathrm{C}}\right)\left(\frac{1\,\mathrm{min}}{60\,\mathrm{s}}\right)\left(\frac{1\,\mathrm{h}}{60\,\mathrm{min}}\right) = 6.44\,\mathrm{h}$$



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