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- Chemical Equations
- Stoichiometry

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- Lewis Model of Bonding
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- Elements and Atoms

2007-J-4:

- Stoichiometry
- Chemical Equations

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- Chemical Equilibrium

2007-J-10:

- Gas Laws
- Write balanced equations for the following nuclear reactions.

Naturally occurring thorium 232 undergoes alpha decay.

A nuclide undergoes beta decay and produces caesium 133.

$$
{ }_{90}^{232} \mathrm{Th} \rightarrow{ }_{88}^{228} \mathrm{Ra}+{ }_{2}^{4} \mathrm{He}
$$

$$
{ }_{54}^{133} \mathrm{Xe} \rightarrow{ }_{55}^{133} \mathrm{Cs}+{ }_{-1}^{0} \beta
$$

- A cook uses a microwave oven to heat up a meal. The wavelength of the radiation is 0.012 m . Calculate the frequency and energy of a photon of this radiation.

The wavelength, $\lambda$, is related to the energy and the frequency, $v$, by the equations:

$$
E=h v=\frac{h c}{\lambda} \text { and } v=\frac{c}{\lambda}
$$

Therefore with $\boldsymbol{\lambda}=\mathbf{0 . 0 1 2} \mathrm{m}$ :

$$
\begin{aligned}
& v=\frac{\left(2.998 \times 10^{8}\right)}{(0.012)}=2.5 \times 10^{10} \mathrm{~s}^{-1} \\
& \mathrm{E}=\frac{\left(6.626 \times 10^{-34}\right) \times\left(2.998 \times 10^{8}\right)}{(0.012)}=1.7 \times 10^{-23} \mathrm{~J}
\end{aligned}
$$

(As the wavelength is given to two significant figures, this limits the accuracy of the answers to also being two significant figures).
Frequency: $\mathbf{2 . 5 \times 1 0 ^ { 1 0 } \mathrm { s } ^ { - 1 }} \quad$ Energy: $1.7 \times \mathbf{1 0}^{-\mathbf{2 3}} \mathrm{J}$

ANSWER CONTINUES ON THE NEXT PAGE

- What mass of calcium chloride is required to make 250 mL of a 0.1 M solution?

The formula mass of calcium chloride, $\mathbf{C a C l}_{\mathbf{2}}$, is

$$
\text { formula mass }=40.08(\mathrm{Ca})+2 \times 35.45(\mathrm{Cl})=110.98
$$

The number of moles in the solution is given by:

$$
\text { number of moles }=\text { concentration } \times \text { volume }=0.1 \times \frac{250}{1000}=0.025 \mathrm{~mol}
$$

The mass required is therefore:

$$
\text { mass }=\text { number of moles } \times \text { formula mass }=(0.025) \times(110.98)=3 \mathrm{~g}
$$

Answer: $\mathbf{3} \mathbf{g}$
What amount of chloride ions (in mol) is present in 30.0 mL of this solution?

One moles of $\mathrm{CaCl}_{2}(\mathrm{~s})$ dissolves to give two moles of $\mathrm{Cl}^{-}(\mathrm{aq})$ ions. Therefore, the number of moles present is:

$$
\text { number of moles }=\text { concentration } \times \text { volume }=(2 \times 0.1) \times \frac{30}{1000}=0.006 \mathrm{~mol}
$$

- Complete the following table.

| Molecular formula | $\mathrm{SF}_{6}$ | $\mathrm{CIF}_{3}$ | $\mathrm{NH}_{3}$ |
| :---: | :---: | :---: | :---: |
| Name |  | chlorine trifluoride |  |
| Lewis structure |  |  |  |
| Number of bonding electron pairs on central atom | 6 | 3 | 3 |
| Number of nonbonding electron pairs on central atom | 0 | 2 | 1 |
| Molecular shape | octahedral | T-shaped | trigonal pyramidal |

- Silicon is essential to the computer industry as a major component of chips. It has three naturally occurring isotopes, the relative abundance of each being given below. Calculate the atomic mass of silicon.

| Isotope | Mass of isotope (a.m.u.) | Relative abundance |
| :---: | :---: | :---: |
| ${ }^{28} \mathrm{Si}$ | 27.9769 | $92.23 \%$ |
| ${ }^{29} \mathrm{Si}$ | 28.9765 | $4.67 \%$ |
| ${ }^{30} \mathrm{Si}$ | 29.9738 | $3.10 \%$ |

The relative atomic mass of silicon is the weighted average of the masses of its isotopes:

$$
\begin{aligned}
\text { atomic mass } & =\left(27.9769 \times \frac{92.23}{100}\right)+\left(28.9765 \times \frac{4.67}{100}\right)+\left(29.9738 \times \frac{3.10}{100}\right) \\
& =28.09
\end{aligned}
$$

(The relative abundances are given to 4 significant figures and limit the accuracy of the answer.)

Answer: $\mathbf{2 8 . 0 9}$

- The complete combustion of butane, $\mathrm{C}_{4} \mathrm{H}_{10}$, in air gives water and carbon dioxide as the products. Write a balanced equation for this reaction.
$2 \mathrm{C}_{4} \mathrm{H}_{10}(\mathrm{~g})+\mathbf{1 3 O}_{2}(\mathrm{~g}) \rightarrow \mathbf{8} \mathrm{CO}_{2}(\mathrm{~g})+\mathbf{1 0 H}_{\mathbf{2}} \mathrm{O}(\mathrm{g})$ or
$\mathrm{C}_{4} \mathrm{H}_{10}(\mathrm{~g})+{ }^{13} / 2 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 4 \mathrm{CO}_{2}(\mathrm{~g})+5 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$

What mass of oxygen is required for the complete combustion of 454 g of butane and what masses of carbon dioxide and water are produced?

The molar mass of butane is $(4 \times 12.01(\mathrm{C}))+(10 \times 1.008(\mathrm{H}))=58.12$. Therefore, the amount of butane in $\mathbf{4 5 4} \mathrm{g}$ is:

$$
\text { number of moles of } \mathrm{C}_{4} \mathrm{H}_{10}=\frac{\text { mass }}{\text { molar mass }}=\frac{454}{58.12}=7.81 \mathrm{~mol}
$$

From the chemical equation, each mole of $\mathrm{C}_{4} \mathrm{H}_{10}$ requires ${ }^{13} / 2$ moles of $\mathrm{O}_{2}$ and produces 4 moles of $\mathrm{CO}_{2}$ and 5 moles of $\mathrm{H}_{2} \mathrm{O}$.

Therefore:
number of moles of $\mathrm{O}_{\mathbf{2}}={ }^{13} / 2 \times 7.81=50.8 \mathrm{~mol}$
number of moles of $\mathrm{CO}_{2}=4 \times 7.81=31.2 \mathrm{~mol}$
number of moles of $\mathrm{O}_{2}=\mathbf{5 \times 7 . 8 1 = 3 9 . 1} \mathbf{~ m o l}$
The molar masses of $\mathrm{O}_{2}, \mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ are:
molar mass of $\mathrm{O}_{2}=\mathbf{2 \times 1 6 . 0 0}=\mathbf{3 2 . 0 0}$
molar mass of $\mathrm{CO}_{2}=12.01(\mathrm{C})+(2 \times 16.00)=44.01$
molar mass of $\mathrm{H}_{2} \mathrm{O}=(\mathbf{2} \times \mathbf{1 . 0 0 8}(\mathrm{H}))+\mathbf{1 6 . 0 0}(\mathrm{O})=18.016$

## Therefore:

mass of $\mathrm{O}_{\mathbf{2}}=$ number of moles $\times$ molar mass $=\mathbf{5 0 . 8} \times \mathbf{3 2 . 0 0}=1620 \mathrm{~g}=1.62 \mathrm{~kg}$ mass of $\mathrm{CO}_{2}=31.2 \times 44.01=1380 \mathrm{~g}=1.38 \mathrm{~kg}$
mass of $\mathrm{H}_{2} \mathrm{O}=\mathbf{3 9 . 1} \times \mathbf{1 8 . 0 1 6}=\mathbf{7 0 4} \mathrm{g}=\mathbf{0 . 7 0 4} \mathrm{kg}$

- During physical activity, lactic acid forms in the muscle tissue and is responsible for muscle soreness. Elemental analysis shows that it contains by mass $40.0 \%$ C, $6.71 \% \mathrm{H}$ and $53.3 \% \mathrm{O}$. Determine the empirical formula of lactic acid.

|  | C | H | O |
| :---: | :---: | :---: | :---: |
| amount in 100 g | 40.0 | 6.71 | 53.3 |
| ratio (divide by atomic mass) | $\frac{40.0}{12.01}=3.33$ | $\frac{6.71}{1.008}=6.66$ | $\frac{53.3}{16.00}=3.33$ |
| divide by smallest | $\frac{3.33}{3.33}=1.00 \sim 1$ | $\frac{6.66}{3.33}=2.00 \sim 2$ | $\frac{3.33}{3.33}=1.00$ |

The simplest possible ratio of $\mathrm{C}: \mathrm{H}: \mathrm{O}$ is thus $1: 2: 1$ and the empirical formula is $\mathrm{CH}_{2} \mathrm{O}$.

$$
\text { Answer: } \mathbf{C H}_{\mathbf{2}} \mathbf{O}
$$

Given that lactic acid has a molar mass of $90.08 \mathrm{~g} \mathrm{~mol}^{-1}$, determine its molecular formula.

The molecular formula is $\left(\mathrm{CH}_{2} \mathrm{O}\right)_{\mathrm{n}}$ so the molar mass is:

$$
\begin{aligned}
\text { molar mass } & =n \times(12.01(\mathrm{C})+2 \times 1.008(\mathrm{H})+16.00(\mathrm{O})) \\
& =30.026 \mathrm{n}=90.08 \text { so } \mathrm{n}=3
\end{aligned}
$$

The molecular formula is thus $\left(\mathrm{CH}_{2} \mathrm{O}\right)_{3}$ or $\mathrm{C}_{3} \mathrm{H}_{6} \mathrm{O}_{3}$

- If 50 mL of a 0.10 M solution of $\mathrm{AgNO}_{3}$ is mixed with 50 mL of a 0.40 M solution of $\mathrm{Na}_{2} \mathrm{CO}_{3}$, what mass of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$ will precipitate from the reaction?

The ionic equation for the precipitation reaction is:

$$
2 \mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq}) \rightarrow \mathrm{Ag}_{2} \mathrm{CO}_{3}(\mathrm{~s})
$$

Thus, two moles of $\mathrm{Ag}^{+}(\mathrm{aq})$ are required for every one mole of $\mathrm{CO}_{3}{ }^{\mathbf{2 -}}(\mathrm{aq})$.
The number of moles of $\mathrm{Ag}^{+}(\mathrm{aq})$ and $\mathrm{CO}_{3}{ }^{\mathbf{2 -}}(\mathrm{aq})$ are given by:

$$
\begin{aligned}
& n\left(\mathrm{Ag}^{+}(\mathrm{aq})\right)=\text { concentration } \times \text { volume }=0.10 \times \frac{50}{1000}=0.0050 \mathrm{~mol} \\
& n\left(\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})\right)=0.40 \times \frac{50}{1000}=0.020 \mathrm{~mol}
\end{aligned}
$$

There is insufficient $\mathrm{Ag}^{+}(\mathrm{aq})$ to react with all of the $\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ and so it is $\mathrm{Ag}^{+}(\mathrm{aq})$ is the limiting reagent. From the chemical equation, 1 mole of $\mathrm{Ag}_{2} \mathbf{C O}_{3}(\mathrm{~s})$ is produced from every two moles of $\mathrm{Ag}^{+}(\mathrm{aq})$ ions. The amount of $\mathrm{Ag}_{2} \mathbf{C O}_{3}(\mathrm{~s})$ produced is therefore:

$$
\mathbf{n}\left(\mathrm{Ag}_{2} \mathrm{CO}_{3}(\mathrm{~s})\right)=1 / 2 \times \mathbf{n}\left(\mathrm{Ag}^{+}(\mathbf{a q})\right)=1 / 2 \times 0.0050=0.0025 \mathrm{~mol}
$$

The formula mass of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$ is $(2 \times 107.87(\mathrm{Ag}))+12.01(\mathrm{C})+(3 \times 16.00(\mathrm{O}))=$ 275.75. This number of moles thus corresponds to a mass of:
mass of $\mathrm{Ag}_{2} \mathrm{CO}_{3}=$ number of moles $\times$ formula mass $=0.0025 \times 275.75=\mathbf{0 . 6 9} \mathrm{g}$

Answer: $0.69 \mathbf{g}$
What is the final concentration of $\mathrm{CO}_{3}{ }^{2-}$ ions in the solution after the above reaction?

From the chemical equation, one mole of $\mathrm{Ag}_{2} \mathrm{CO}_{3}(\mathrm{~s})$ is produced from every mole of $\mathrm{CO}_{3}{ }^{2-}$ which reacts. Therefore 0.0025 mol of $\mathrm{CO}_{3}{ }^{2-}$ reacts. This leaves:

$$
\text { number of moles of unreacted } \mathrm{CO}_{3}{ }^{2-}=0.020-0.0025=0.018 \mathrm{~mol}
$$

The total volume of the solution after mixing is $(50+50)=100 \mathrm{~mL}$. The final concentration is therefore:

$$
\text { concentration }=\frac{\text { number of moles }}{\text { volume }}=\frac{0.018}{100 / 1000}=0.18 \mathrm{M}
$$

- Give balanced ionic equations for the reactions that occur in each of the following cases.

Sodium metal is added to excess water.

$$
2 \mathrm{Na}(\mathrm{~s})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightarrow 2 \mathrm{Na}^{+}(\mathrm{aq})+2 \mathrm{OH}^{-}(\mathrm{aq})+\mathrm{H}_{2}(\mathrm{~g})
$$

Solutions of cobalt(II) nitrate and sodium phosphate are mixed.

$$
3 \mathrm{Co}^{2+}(\mathrm{aq})+2 \mathrm{PO}_{4}{ }^{3-}(\mathrm{aq}) \rightarrow \mathrm{Co}_{3}\left(\mathrm{PO}_{4}\right)_{2}(\mathrm{~s})
$$

Solid calcium carbonate is dissolved in dilute nitric acid.

$$
\mathrm{CaCO}_{3}(\mathrm{~s})+2 \mathrm{H}^{+}(\mathrm{aq}) \rightarrow \mathrm{Ca}^{2+}(\mathrm{aq})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})
$$

- A 60.0 g piece of Ag metal is heated to $90.0^{\circ} \mathrm{C}$ and dropped into 120.0 g of water at $25.0^{\circ} \mathrm{C}$ in a well insulated container. The final temperature of the $\mathrm{Ag}-\mathrm{H}_{2} \mathrm{O}$ mixture is $26.7^{\circ} \mathrm{C}$. Calculate the specific heat of silver.
Data: The specific heat of water is $4.18 \mathrm{~J} \mathrm{~g}^{-1} \mathrm{~K}^{-1}$.

The temperature of the Ag metal decreases from $90.0^{\circ} \mathrm{C}$ to $26.7^{\circ} \mathrm{C}$. The heat change is given by:

$$
\text { heat change of } A g=q_{A g}=c_{A g} \times m_{A g} \times \Delta T=c_{A g} \times(60.0) \times(26.7-90.0)
$$

The temperature of the water changes from $25.0^{\circ} \mathrm{C}$ to $26.7^{\circ} \mathrm{C}$. The heat change is given by:

$$
\begin{aligned}
\text { heat change of water }=\mathbf{q}_{\text {water }} & =\mathbf{c}_{\text {water }} \times \mathbf{m}_{\text {water }} \times \Delta \mathrm{T} \\
& =(4.18) \times(\mathbf{1 2 0 . 0}) \times(\mathbf{2 6 . 7} \mathbf{- 2 5 . 0})
\end{aligned}
$$

When the Ag metal is dropped into the water, the heat lost by the silver is gained by the water until both reach the same temperature: $-\mathbf{q}_{\mathrm{Ag}}=\mathbf{q}_{\text {water }}$ :

$$
\begin{aligned}
& -\mathrm{c}_{\mathrm{Ag}} \times(60.0) \times(26.7-90.0)=(4.18) \times(120.0) \times(26.7-25.0) \\
& \mathbf{c}_{\mathrm{Ag}}=\frac{(4.18) \times(120.0) \times(26.7-25.0)}{(60.0) \times(90.0-26.7)}=0.22 \mathrm{~J} \mathrm{~g}^{-1} \mathrm{~K}^{-1}
\end{aligned}
$$

Answer: $\mathbf{0 . 2 2} \mathbf{J ~ g}^{\mathbf{- 1}} \mathbf{K}^{\mathbf{- 1}}$

- Determine $K_{\mathrm{c}}$ for the reaction $1 / 2 \mathrm{O}_{2}(\mathrm{~g})+\mathrm{Na}_{2} \mathrm{O}(\mathrm{s}) \rightleftharpoons \mathrm{Na}_{2} \mathrm{O}_{2}(\mathrm{~s})$ at $25^{\circ} \mathrm{C}$.

Data: $\quad \mathrm{Na}_{2} \mathrm{O}(\mathrm{s}) \rightleftharpoons 2 \mathrm{Na}(\mathrm{s})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \quad K_{\mathrm{c}}=2 \times 10^{-25}$ at $25^{\circ} \mathrm{C}$.

$$
\mathrm{Na}_{2} \mathrm{O}_{2}(\mathrm{~s}) \rightleftharpoons 2 \mathrm{Na}(\mathrm{~s})+\mathrm{O}_{2}(\mathrm{~g}) \quad K_{\mathrm{c}}=5 \times 10^{-29} \text { at } 25^{\circ} \mathrm{C}
$$

The reaction involves the formation of $\mathrm{Na}_{2} \mathrm{O}_{2}(\mathrm{~s})$ from $\mathrm{Na}_{2} \mathrm{O}(\mathrm{s})$ and thus involves the first reaction and the reverse of the second reaction. The reactions can be combined:

$$
\begin{array}{ll}
\mathrm{Na}_{2} \mathrm{O}(\mathrm{~s}) \rightleftharpoons 2 \mathrm{Na}(\mathrm{~s})+1 / 2 \mathrm{O}_{2}(\mathrm{~s}) & K_{\mathrm{c}}=2 \times 10^{-25} \\
2 \mathrm{Na}(\mathrm{~s})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{Na}_{2} \mathrm{O}_{2}(\mathrm{~s}) & K_{\mathrm{c}}=\frac{1}{\left(5 \times 10^{-29}\right)}=2 \times 10^{-30}
\end{array}
$$

$$
\mathrm{Na}_{2} \mathrm{O}(\mathrm{~s})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{Na}_{2} \mathrm{O}_{2}(\mathrm{~g}) \quad K_{\mathrm{c}}=\left(2 \times 10^{-25}\right) \times \frac{1}{\left(5 \times 10^{-29}\right)}=4 \times 10^{3}
$$

- Consider a cell composed of the following half-reactions.

$$
\begin{aligned}
& \mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{e}^{-} \rightarrow \mathrm{Ag}(\mathrm{~s}) \\
& \mathrm{Cr}(\mathrm{~s}) \rightarrow \mathrm{Cr}^{3+}(\mathrm{aq})+3 \mathrm{e}^{-}
\end{aligned}
$$

What is the balanced equation for the spontaneous reaction?
The $\mathbf{E}^{0}$ values for the $\mathrm{Ag}^{+} / \mathbf{A g}$ and $\mathbf{C r} / \mathrm{Cr}^{3+}$ cells are +0.80 and $-\mathbf{0 . 7 4} \mathrm{V}$ respectively. The least positive $\left(\mathbf{C r} / \mathbf{C r}^{3+}\right)$ is reversed and is the oxidation half cell:

$$
3 \mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{Cr}(\mathrm{~s}) \rightarrow \mathbf{3 A g}(\mathrm{s})+\mathrm{Cr}^{3+}(\mathrm{aq})
$$

What is the value of $E^{\circ}$ for the cell? Relevant standard reduction potentials are on the data sheet.
$\mathbf{E}_{\text {cell }}^{0}=\mathbf{E}_{\text {red }}^{0}+\mathbf{E}_{\text {ox }}^{0}=(+\mathbf{0 . 8 0})+(\mathbf{0 . 7 4})=+\mathbf{1 . 5 4} \mathbf{V}$
The potential for the $\mathbf{C r} / \mathbf{C r}^{3+}$ cell is reversed as it is the oxidation half cell.
Answer: +1.54 V

- Calculate the standard heat of reaction for the following reaction.

$$
\mathrm{Zn}(\mathrm{~s})+2 \mathrm{Cu}^{+}(\mathrm{aq}) \rightarrow 2 \mathrm{Cu}(\mathrm{~s})+\mathrm{Zn}^{2+}(\mathrm{aq})
$$

Data: $\quad \Delta H_{\mathrm{f}}^{\mathrm{o}}=+51.9 \mathrm{~kJ} \mathrm{~mol}^{-1}$ for $\mathrm{Cu}^{+}(\mathrm{aq})$

$$
\Delta H_{\mathrm{f}}^{\mathrm{o}}=-152.4 \mathrm{~kJ} \mathrm{~mol}^{-1} \text { for } \mathrm{Zn}^{2+}(\mathrm{aq})
$$

Using $\Delta_{\mathrm{rxn}} \mathrm{H}^{\mathbf{0}}=\sum \mathrm{m} \Delta_{\mathrm{f}} \mathrm{H}^{\mathbf{0}}$ (products) $-\sum \mathrm{n} \Delta_{\mathrm{f}} \mathrm{H}^{\mathbf{0}}$ (reactants), the heat of the reaction as written is:

$$
\begin{aligned}
\Delta_{\mathrm{rxn}} \mathbf{H}^{0} & =\left[2 \Delta_{\mathrm{f}} \mathbf{H}^{0}(\mathrm{Cu}(\mathrm{~s}))+\Delta_{\mathrm{f}} \mathbf{H}^{0}\left(\mathrm{Zn}^{2+}(\mathrm{aq})\right)\right]-\left[\Delta_{\mathrm{f}} \mathrm{H}^{0}(\mathrm{Zn}(\mathrm{~s}))+2 \Delta_{\mathrm{f}} \mathbf{H}^{0}\left(\mathrm{Cu}^{+}(\mathrm{a}\right.\right. \\
& =[(2 \times 0)+(-152.4)]-[(\mathbf{0})+(2 \times+51.9)]=-256.2 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{aligned}
$$

$\Delta_{f} H^{\mathbf{0}}(\mathbf{C u}(\mathrm{s}))=\Delta_{\mathrm{f}} \mathrm{H}^{\mathbf{0}}(\mathrm{Zn}(\mathrm{s}))=0$ for elements already in their standard states.

Answer: - $\mathbf{2 5 6 . 2} \mathbf{~ k J ~ m o l}{ }^{\mathbf{- 1}}$

- An electrolytic cell contains a solution of $\mathrm{MCl}_{3}$. A total charge of 3600 C is passed through the cell, depositing 0.65 g of the metal, M , at the cathode. What is the identity of the metal, M ?

The total number of electrons passed is given by:

$$
\text { number of moles of electrons }=\frac{I t}{F}=\frac{Q}{F}=\frac{3600}{96485}=0.037 \mathrm{~mol} .
$$

As the solution contains $\mathbf{M C l}_{3}$, these electrons are reducing $\mathbf{M}^{3+}(\mathrm{aq})$ ions, each of which requires $3 e^{-}$. Hence, the number of moles of metal, $M$, formed at the cathode is:

$$
\text { number of moles of } M=1 / 3 \times 0.037=0.012
$$

This amount of $M$ has a mass of 0.65 g . The atomic mass of $M$ is therefore:

$$
\text { atomic mass }=\frac{\text { mass }}{\text { number of moles }}=\frac{0.65 \mathrm{~g}}{0.012 \mathrm{~mol}}=52 \mathrm{~g} \mathrm{~mol}^{-1}
$$

This atomic mass corresponds to that of chromium (Cr).

Answer: Chromium (Cr)

- A metal-metal hydride battery has the following shorthand notation:

$$
\mathrm{MH}(\mathrm{~s}), \mathrm{M}(\mathrm{~s})\left|\mathrm{OH}^{-}(\mathrm{aq}) \| \mathrm{OH}^{-}(\mathrm{aq})\right| \mathrm{NiO}(\mathrm{OH})(\mathrm{s}), \mathrm{Ni}(\mathrm{OH})_{2}(\mathrm{~s})
$$

Which component of the battery is the cathode?
$\mathrm{NiO}(\mathrm{OH})(\mathrm{s}) / \mathrm{Ni}(\mathrm{OH})_{2}(\mathrm{~s})$
(the right hand side of the cell notation)

Give the balanced half equation of the reaction that takes place at the cathode.
$\mathrm{NiO}(\mathrm{OH})(\mathrm{s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})+\mathrm{e}^{-} \rightarrow \mathrm{Ni}(\mathrm{OH})_{2}(\mathrm{~s})+\mathrm{OH}^{-}(\mathrm{aq})$

Why is it important that all redox active species are solids in this reaction?

As all redox active species are solids, their concentrations do not change during the reaction. They therefore do not appear in the equilibrium expression or in the Nernst equation.

The battery therefore maintains a constant voltage.

- $K_{\mathrm{p}}=7.0$ for the reaction $\mathrm{Br}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{BrCl}(\mathrm{g}) \quad$ at 400 K .

Suppose a 1.0 L flask is filled with $0.30 \mathrm{~atm} \mathrm{Br}_{2}(\mathrm{~g})$ and $0.30 \mathrm{~atm} \mathrm{Cl}_{2}(\mathrm{~g})$ at 400 K . Find the pressures of all three gases at equilibrium.

The equilibrium constant in terms of partial pressures, $K_{p}$, is given by:

$$
K_{p}=\frac{\left(\mathbf{P}_{\mathrm{BrCl}}\right)^{2}}{\left(\mathbf{P}_{\mathrm{Br}_{2}}\right)\left(\mathbf{P}_{\mathrm{Cl}_{2}}\right)}
$$

The reaction table is:

| pressure | $\mathrm{Br}_{\mathbf{2}}(\mathrm{g})$ | $\mathrm{Cl}_{\mathbf{2}}(\mathrm{g})$ | $\rightleftharpoons$ | $2 \mathrm{FrCl}(\mathrm{g})$ |
| :---: | :---: | :---: | :---: | :---: |
| start | 0.30 | 0.30 |  | 0 |
| change | $-\mathbf{x}$ | $-\mathbf{x}$ |  | $+2 \mathbf{x}$ |
| equilibrium | $0.30-\mathbf{x}$ | $0.30-\mathbf{x}$ |  | $\mathbf{2 x}$ |

The equilibrium constant in terms of partial pressures, $K_{p}$, is given by:

$$
K_{p}=\frac{\left(P_{\mathrm{BrCl}}\right)^{2}}{\left(\mathrm{P}_{\mathrm{Br}_{2}}\right)\left(\mathrm{P}_{\mathrm{Cl}_{2}}\right)}=\frac{(2 x)^{2}}{(0.30-x)(0.30-x)}=\frac{(2 x)^{2}}{(0.30-x)^{2}}=7.0
$$

As $K_{p}$ is not small, the assumption that $0.30-x \sim x$ cannot be made. Hence,

$$
\begin{aligned}
& \frac{(2 x)}{(0.30-x)}=\sqrt{7.0} \\
& 2 x=(0.30-x) \times \sqrt{7.0}=(0.30 \times \sqrt{7})-x \sqrt{7} \\
& x=\frac{(0.30 \times \sqrt{7})}{(2+\sqrt{7})}=0.17
\end{aligned}
$$

The partial pressures at equilibrium are therefore:

| $\mathbf{P}_{\mathbf{B r}_{2}}=\mathbf{P}_{\mathrm{Cl}_{2}}=\mathbf{0 . 3 0 - \mathbf { x } = \mathbf { 0 . 1 3 } \mathbf { ~ a t m } , \mathbf { P } _ { \mathrm { BrCl } } = \mathbf { 2 x } = \mathbf { 0 . 3 4 } \mathbf { ~ a t m }}$ |  |  |
| :--- | :--- | :--- |
| $p\left(\mathrm{Br}_{2}\right): \mathbf{0 . 1 3} \mathbf{~ a t m}$ | $p\left(\mathrm{Cl}_{2}\right): \mathbf{0 . 1 3} \mathbf{~ a t m}$ | $p(\mathrm{BrCl}): \mathbf{0 . 3 4} \mathbf{~ a t m}$ |

- The Voyager I spacecraft determined that the atmospheric pressure at the surface of Saturn's moon, Titan, is 1.6 times that of earth and that the atmosphere contains $6.0 \mathrm{~mol} \%$ methane, $\mathrm{CH}_{4}$. What is the partial pressure of methane on Titan in mmHg ?

If the atmospheric pressure on Titan is $\mathbf{1 . 6}$ times that on earth, $\mathbf{P}_{\text {total }}=\mathbf{1 . 6} \mathbf{~ a t m}$.
If the atmosphere contains $6.0 \mathrm{~mol} \%$ methane, then:

$$
P_{\mathrm{CH}_{4}}=0.06 \times 1.6=0.096 \mathrm{~atm}=0.096 \times 760=73 \mathrm{mmHg}
$$

## Answer: $\mathbf{7 3} \mathbf{~ m m H g}$

- Many gases are available for use in compressed gas cylinders, in which they are stored at high pressures. Calculate the mass of $\mathrm{O}_{2}$ that can be stored at $20 .{ }^{\circ} \mathrm{C}$ and 170. atm pressure in a cylinder with a volume of 60.0 L .

Using the ideal gas law, $\mathrm{PV}=\mathrm{nRT}$, the number of moles that can be stored is:

$$
\mathrm{n}=\frac{\mathrm{PV}}{\mathrm{RT}}=\frac{(170 .) \times(60.0)}{(0.08206) \times(20 .+273)}=424 \mathrm{~mol}
$$

As the molar mass of $\mathrm{O}_{\mathbf{2}}$ is $(\mathbf{2} \times \mathbf{1 6 . 0 0})=\mathbf{3 2 . 0 0}$, this corresponds to a mass of:

$$
\text { mass }=\text { number of moles } \times \text { molar mass }=424 \times 32.00=13600 \mathrm{~g}=13.6 \mathrm{~kg}
$$

Answer: 13,600 g or $\mathbf{1 3 . 6} \mathbf{~ k g}$
What volume would this mass of oxygen occupy at 1.00 atm pressure and $20^{\circ} \mathrm{C}$ ?

Using the ideal gas law, the volume of 424 mol at 1.00 atm and $20^{\circ} \mathrm{C}$ is:

$$
V=\frac{n R T}{P}=\frac{(424) \times(0.08206) \times(20+273)}{(1.00)}=10.2 \times 10^{4} \mathrm{~L}
$$

Equivalently, $\mathbf{P}_{1} \mathbf{V}_{\mathbf{1}}=\mathbf{P}_{2} \mathbf{V}_{2}$ gives:

$$
V_{2}=\frac{P_{1} V_{1}}{P_{2}}=\frac{(170) \times(60.0)}{(1.00)}=1.02 \times 10^{4} \mathrm{~L}
$$

