The worksheets are available in the tutorials and form an integral part of the learning outcomes and experience for this unit.

**Model 1: The Effect of Concentration on the Cell Potential**

1. If $[H^+(aq)] = 1.0 \text{ M}$ then $\text{pH} = 0$.
2. At a lower $[H^+(aq)]$, the reaction is less favourable and so $E$ will be less positive.

**Model 2: The Nernst Equation**

1. For reaction (1), $n = 2$. For reaction (2), $n = 2$.

2. 
   $$Q = \frac{[H_2O][NAD^+]}{[O_2]^{1/2}[H^+][NADH]} = \frac{(1)(1)}{(1)(10^{-7})(1)} = 10^7$$

   With $T = 37 \degree C$ and $n = 2$:
   $$E_{\text{cell}} = E_{\text{cell}}^0 - \frac{RT}{nF} \ln Q = (+1.335 \text{ V}) - \frac{(8.314 \text{ J K}^{-1} \text{ mol}^{-1})(310 \text{ K})}{(2)(96485 \text{ C mol}^{-1})} \ln(10^7) = 1.116 \text{ V}$$

3. $E_{\text{cell}}$ is less positive $E_{\text{cell}}^0$ as predicted.

**Model 3: Concentration Cells**

1. (a) Cu$^{2+}$ ions will move from the more concentrated to the less concentrated solution until the concentrations are equal. The colour of the two solutions will become indistinguishable.
   (b) The entropy will increase.
   (c) The enthalpy will stay the same.

2. (a) No.
   (b) Cell A: [Cu$^{2+}$(aq)] must decrease.
   Cell B: [Cu$^{2+}$(aq)] must increase.
   (c) Cell A: Cu$^{2+}$(aq) + 2e$^- \rightarrow$ Cu(s)
   Cell B: Cu(s) $\rightarrow$ Cu$^{2+}$(aq) + 2e$^-$
   (d) Electrons will flow through the wire from B to A.
   (e) A = cathode. B = anode.
   (f) $E_{\text{cell}}^0 = 0 \text{ V}$. If the concentrations are both 1 M, the system is at equilibrium.
   (g) 
   $$Q = \frac{[\text{Cu}^{2+}(aq)]_B}{[\text{Cu}^{2+}(aq)]_A} = \frac{0.1}{1.0} = 0.1$$
   $$E_{\text{cell}} = 0 - \frac{(8.314 \text{ J K}^{-1} \text{ mol}^{-1})(298 \text{ K})}{(2)(96485 \text{ C mol}^{-1})} \ln(0.1) = 0.030 \text{ V}$$

**Model 4: Voltaic Cells**

1. The zinc electrode will lose mass and the tin electrode will gain mass.
2. Sn$^{2+}$(aq) + Zn(s) $\rightarrow$ Sn(s) + Zn$^{2+}$(aq)
3. Zn is being oxidised and Sn$^{2+}$ is being reduced. The Zn electrode is the anode. The Sn electrode is the cathode.

4. Electrons flow through the wire, from the zinc electrode towards the tin electrode.

5. The anode is negative and the cathode is positive.

6. SO$_4^{2-}$(aq) moves into the zinc half cell (as cations are being made in the oxidation reaction in this cell). Na$^+$ (aq) moves into the tin half cell (as cations are being lost in this cell).

**Model 5: Electrolytic Cells**

1. The zinc electrode will *gain* mass and the tin electrode will *lose* mass.

2. Sn(s) + Zn$^{2+}$(aq) $\rightarrow$ Sn$^{2+}$(aq) + Zn(s)

3. Zn is being reduced and Sn$^{2+}$ is being oxidised. The Zn electrode is the cathode. The Sn electrode is the anode.

4. Electrons flow through the wire, from the tin electrode towards the zinc electrode.

5. The anode is positive and the cathode is negative. The power source pumps electrons to the cathode from the anode.

6. SO$_4^{2-}$(aq) moves into the tin half cell (as cations are being made in the oxidation reaction in this cell). Na$^+$ (aq) moves into the zinc half cell (as cations are being lost in this cell).

**Model 6: Electrolysis of Water**

1. $2\text{H}_2\text{O}(l) \rightarrow 2\text{H}_2(g) + \text{O}_2(g)$.

2. $F = (1.602 \times 10^{-19} \text{ C}) \times (6.022 \times 10^{23}) = 96485 \text{ C mol}^{-1}$.

3. Number of moles of electrons = $I \times t / F = (10.0 \text{ A}) \times (2.00 \times 60 \times 60 \text{ s}) / (96485 \text{ C mol}^{-1}) = 0.746 \text{ mol}$

4. From the half cell equation for the reduction of H$_2$O, 2e$^-$ are required for each H$_2$. Therefore, 0.746 mol will produce $\frac{1}{2} \times 0.746 \text{ mol} = 0.373 \text{ mol of H}_2$.

5. From Q1, half as much O$_2$ will be produced: 0.187 mol.

Alternatively, from the half cell equation for the oxidation of H$_2$O, 4e$^-$ are produced for each O$_2$. Therefore, 0.746 mol will have been produced by $\frac{1}{4} \times 0.746 \text{ mol} = 0.187 \text{ mol of O}_2$.

6. The reduction potential of water is -0.82 V so a cation with a more negative reduction potential should be used: Cr$^{3+}$, Al$^{3+}$, Mg$^{2+}$, Na$^+$, Ca$^{2+}$ or Li$^+$.

The oxidation potential of water is -1.23 V so an anion with a more negative oxidation potential should be used: Cl$^-$ or SO$_4^{2-}$.

Na$_2$SO$_4$ or K$_2$SO$_4$ are commonly used.