

**Table 18.1 Selected Standard Electrode Potentials at 25 °C**

Reduction Half-Reaction	$E^\circ$ Volts
<b>Acidic Solution</b>	
$F_2(g) + 2 e^- \longrightarrow 2 F^-(aq)$	+2.866
$O_3(g) + 2 H^+(aq) + 2 e^- \longrightarrow O_2(g) + H_2O(l)$	+2.075
$S_2O_8^{2-}(aq) + 2 e^- \longrightarrow 2 SO_4^{2-}(aq)$	+2.01
$H_2O_2(aq) + 2 H^+(aq) + 2 e^- \longrightarrow 2 H_2O(l)$	+1.763
$MnO_4^-(aq) + 8 H^+(aq) + 5 e^- \longrightarrow Mn^{2+}(aq) + 4 H_2O(l)$	+1.51
$PbO_2(s) + 4 H^+(aq) + 2 e^- \longrightarrow Pb^{2+}(aq) + 2 H_2O(l)$	+1.455
$Cl_2(g) + 2 e^- \longrightarrow 2 Cl^-(aq)$	+1.358
$Cr_2O_7^{2-}(aq) + 14 H^+(aq) + 6 e^- \longrightarrow 2 Cr^{3+}(aq) + 7 H_2O(l)$	+1.33
$MnO_2(s) + 4 H^+(aq) + 2 e^- \longrightarrow Mn^{2+}(aq) + 2 H_2O(l)$	+1.23
$O_2(g) + 4 H^+(aq) + 4 e^- \longrightarrow 2 H_2O(l)$	+1.229
$2 IO_3^-(aq) + 12 H^+(aq) + 10 e^- \longrightarrow I_2(s) + 6 H_2O(l)$	+1.20
$Br_2(l) + 2 e^- \longrightarrow 2 Br^-(aq)$	+1.065
$NO_3^-(aq) + 4 H^+(aq) + 3 e^- \longrightarrow NO(g) + 2 H_2O(l)$	+0.956
$Ag^+(aq) + e^- \longrightarrow Ag(s)$	+0.800
$Fe^{3+}(aq) + e^- \longrightarrow Fe^{2+}(aq)$	+0.771
$O_2(g) + 2 H^+(aq) + 2 e^- \longrightarrow H_2O_2(aq)$	+0.695
$I_2(s) + 2 e^- \longrightarrow 2 I^-(aq)$	+0.535
$Cu^{2+}(aq) + 2 e^- \longrightarrow Cu(s)$	+0.340
$SO_4^{2-}(aq) + 4 H^+(aq) + 2 e^- \longrightarrow 2 H_2O(l) + SO_2(g)$	+0.17
$Sn^{4+}(aq) + 2 e^- \longrightarrow Sn^{2+}(aq)$	+0.154
$S(s) + 2 H^+(aq) + 2 e^- \longrightarrow H_2S(g)$	+0.14
$2 H^+(aq) + 2 e^- \longrightarrow H_2(g)$	0
$Pb^{2+}(aq) + 2 e^- \longrightarrow Pb(s)$	-0.125
$Sn^{2+}(aq) + 2 e^- \longrightarrow Sn(s)$	-0.137
$Co^{2+}(aq) + 2 e^- \longrightarrow Co(s)$	-0.277
$Fe^{2+}(aq) + 2 e^- \longrightarrow Fe(s)$	-0.440
$Zn^{2+}(aq) + 2 e^- \longrightarrow Zn(s)$	-0.763
$Al^{3+}(aq) + 3 e^- \longrightarrow Al(s)$	-1.676
$Mg^{2+}(aq) + 2 e^- \longrightarrow Mg(s)$	-2.356
$Na^+(aq) + e^- \longrightarrow Na(s)$	-2.713
$Ca^{2+}(aq) + 2 e^- \longrightarrow Ca(s)$	-2.84
$K^+(aq) + e^- \longrightarrow K(s)$	-2.924
$Li^+(aq) + e^- \longrightarrow Li(s)$	-3.040
<b>Basic Solution</b>	
$O_3(g) + H_2O(l) + 2 e^- \longrightarrow O_2(g) + 2 OH^-(aq)$	+1.246
$OCl^-(aq) + H_2O(l) + 2 e^- \longrightarrow Cl^-(aq) + 2 OH^-(aq)$	+0.890
$O_2(g) + 2 H_2O(l) + 4 e^- \longrightarrow 4 OH^-(aq)$	+0.401
$2 H_2O(l) + 2 e^- \longrightarrow H_2(g) + 2 OH^-(aq)$	-0.828

## IMPORTANT FORMULAS

$$\Delta G^\circ = \sum n \Delta G^\circ_{\text{prod}} - \sum m \Delta G^\circ_{\text{react}} \text{ (similar formulas for } \Delta H^\circ \text{ and } \Delta S^\circ)$$

$$\Delta G^\circ = -nFE^\circ$$

F is the Faraday constant = 96.5 kC (gives an answer in kJ)

$$E^\circ_{\text{cell}} = \frac{0.0592}{n} \log K$$

$$E_{\text{cell}} = E^\circ_{\text{cell}} - \frac{0.0592}{n} \log Q$$

$$E^\circ_{\text{cell}} = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$\Delta G^\circ = -5.71 \log K \text{ (at } 25^\circ \text{ C)}$$

$$T_{1/2} = \frac{0.693}{k} \text{ first order}$$

$$\text{pH} = \text{pK}_a + \log \frac{[\text{base}]}{[\text{acid}]}$$

$$\log k_2/k_1 = \frac{E_a}{19.14} \times \left( \frac{1}{T_1} - \frac{1}{T_2} \right) \text{ (gives answer in J)}$$

$$1 \text{ COULOMB} = 1 \text{ AMP} \times 1 \text{ SEC}$$

- rate =  $k[A]^x[B]^y$  (14.1) Rate law expressions. The sum ( $x + y$ ) gives the overall order of the reaction.
- $\ln \frac{[A]_t}{[A]_0} = -kt$  (14.3) Relationship between concentration and time for a first-order reaction.
- $\ln [A]_t = -kt + \ln [A]_0$  (14.4) Equation for the graphical determination of  $k$  for a first-order reaction.
- $t_{1/2} = \frac{0.693}{k}$  (14.5) Half-life for a first-order reaction.
- $\frac{1}{[A]_t} = kt + \frac{1}{[A]_0}$  (14.6) Relationship between concentration and time for a second-order reaction.
- $[A]_t = -kt + [A]_0$  (14.8) Relationship between concentration and time for a zero-order reaction.
- $k = Ae^{-E_a/RT}$  (14.10) The Arrhenius equation expressing the dependence of the rate constant on activation energy and temperature.
- $\ln k = \left(-\frac{E_a}{R}\right)\left(\frac{1}{T}\right) + \ln A$  (14.12) Equation for the graphical determination of activation energy.
- $\ln \frac{k_1}{k_2} = \frac{E_a}{R} \left(\frac{T_1 - T_2}{T_1 T_2}\right)$  (14.13) Relationship of rate constants at two different temperatures.  $R = 8,31 \text{ J/K}$