Electrolytic Cells

Electrolysis

- Electrolysis → The process of supplying electrical energy to force a nonspontaneous redox reaction to occur
- Electrolytic Cell A cell that consists of a combination of two electrodes, an electrolyte, and an external power source (typically a battery)



Table 11.2 Cell Comparison

Galvanic Cell	Electrolytic Cell	
Spontaneous reaction	Non-spontaneous reaction	
Converts chemical energy to electrical energy	Converts electrical energy to chemical energy	
node (negative): Zinc Anode (positive): Copper		
Cathode (positive): Copper	Cathode (negative): Zinc	
$ \text{cidation (at anode): } \text{Zn}_{(s)} \rightarrow \text{Zn}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Oxidation (at anode): } \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2e^{-} \\ \text{Cu}_{(s)} \rightarrow \text{Cu}^{2+}_{(aq)} \rightarrow \text{Cu}^{2+}_{(aq$		
uction (at cathode): $\operatorname{Cu}^{2+}_{(aq)} + 2e^{-} \rightarrow \operatorname{Cu}_{(s)}$ Reduction (at cathode): $\operatorname{Zn}^{2+}_{(aq)} + 2e^{-} \rightarrow \operatorname{Zn}_{(s)}$		
ell reaction: $\operatorname{Zn}_{(s)} + \operatorname{Cu}^{2+}_{(aq)} \rightarrow \operatorname{Zn}^{2+}_{(aq)} + \operatorname{Cu}_{(s)}$ Cell reaction: $\operatorname{Cu}_{(s)} + \operatorname{Zn}^{2+}_{(aq)} \rightarrow \operatorname{Cu}^{2+}_{(aq)} + \operatorname{Zn}_{(s)}$		

Galvanic Cells and Electrolytic Cells are, simply, opposites of each other. Where the GC is spontaneous, the EC requires an input of energy. Where the cell potential is positive for GC, it's negative for EC. Where the Cathode is the positive electrode and the Anode is the negative electrode for GC, it's the opposite for EC.

Production of Elements

Most elements occur naturally combined with other elements in compounds. Ionic compounds of sodium (Na), potassium (K), lithium (L), etc., are not found

SOA	$Zn^{2+}_{(aq)} + 2 e^- \longrightarrow Zn_{(s)}$
	$2H_2O_{(1)} + 2e^- \longrightarrow H_{2(g)} + 2OH_{(aq)}^-$
	$Mg_{(aq)}^{2+}$ + 2 e ⁻ \longrightarrow $Mg_{(s)}$
*	$Na^+_{(aq)} + e^- \longrightarrow Na_{(s)}$

uncombined in nature, simply due to their oxidative potentials.

lonic compounds can, and are, melted, yielding molten ionic compounds that are good electrical conductors and can function as the electrolyte in a cell. In the electrolysis of molten binary ionic compounds, only one oxidizing agent and one reducing agent are present. The production of active metals from their minerals typically involves the electrolysis of molten compounds of the metal.

In molten-salt electrolysis, metal cations move to the cathode and are reduced to metals, and nonmetal anions move to the anode and are oxidized to nonmetals.

Electrolysis of Molten Salts

Cathode / Reduction / ---'ve

 $Na^+ + e^- \rightarrow Na$ $E^o = -2.71$

Anode / Oxidation / +'ve

 $2 \operatorname{Cl} + 2 \operatorname{e}^{-} \rightarrow 2 \operatorname{Cl}^{-} E^{o} = +1.36$

Final Equation:

$2[Na^+ + e^- \rightarrow Na]$	$E^{o} = -2.71$
$2 \text{ Cl}^2 \rightarrow 2 \text{ Cl} + 2 \text{ e}^2$	$E^{o} = -1.36$
$2 \operatorname{Na}^+ + 2 \operatorname{Cl}^- \rightarrow 2 \operatorname{Na} + 2 \operatorname{Cl}^-$	$E^{o} = -4.07$



Electrolysis of Water

Cathode / Reduction / -- 've

$$2 H_2 O + 2 e^- \rightarrow H_2 + 2 O H^- E^o = -0.83$$

Anode / Oxidation / +'ve

$$O_2 + 4 H^+ + 4 e^- \rightarrow 2 H_2 O E^o = +1.23$$

Final Equation:

$$2[2 H_2 0 + 2 e^- \rightarrow H_2 + 2 OH^-] \quad E^o = -0.83$$

$$2 H_2 0 \rightarrow O_2 + 4 H^+ + 4 e^- \qquad E^o = -1.23$$

$$6 H_2 0 \rightarrow 2 H_2 + O_2 + 4 H^+ + 4 OH^- E^o = -2.06$$

$$2 H_2 0 \rightarrow 2 H_2 + O_2 E^o = -2.06$$

