

# AP CHEMISTRY CHEAT SHEET

## Unit 6 : THERMODYNAMICS

### Quick Overview

- **Focus:** energy transfer, enthalpy, entropy, Gibbs free energy, and spontaneity.
- **Exam Lens:** connect energy changes → entropy → temperature → reaction feasibility.

### System vs Surroundings

- **System:** part of the universe being studied.
- **Surroundings:** everything else.
- **Universe:** system + surroundings.

### Energy flow:

- **Exothermic:** energy released to surroundings ( $\Delta H < 0$ ).
- **Endothermic:** energy absorbed from surroundings ( $\Delta H > 0$ ).

### Heat, Work & Energy

- **Heat (q):** energy transferred due to temperature difference.
- **Work (w):** energy from force acting over distance.

#### Mini formula box

$$q = mC\Delta T$$
$$w = -P\Delta V$$
$$\Delta E = q + w$$

### Enthalpy ( $\Delta H$ )

- $\Delta H = H_{\text{products}} - H_{\text{reactants}}$
- Negative  $\Delta H$  → exothermic
- Positive  $\Delta H$  → endothermic

### Standard enthalpy of formation ( $\Delta H^{\circ}_f$ ):

- Enthalpy change when 1 mol of compound forms from elements in standard states.
- $\Delta H^{\circ}_f$  for elements in standard state = 0.

#### Mini formula box

$$\Delta G^{\circ} = -RT \ln K$$

- $R = 8.314 \text{ J}\cdot\text{mol}^{-1}\cdot\text{K}^{-1}$
- Large  $K$  → negative  $\Delta G^{\circ}$

### Calorimetry

Used to measure heat changes experimentally.

- $q_{\text{reaction}} = -q_{\text{solution}}$
- Constant pressure calorimeter → measures  $\Delta H$ .

#### Mini formula box

$$q = mC\Delta T$$

**C = specific heat capacity**

### Hess's Law

- Overall  $\Delta H$  is independent of reaction pathway.
- Reverse reaction → flip sign of  $\Delta H$ .
- Multiply reaction → multiply  $\Delta H$ .

**Mnemonic:** "Flip flips, scale scales."

### Entropy ( $\Delta S$ )

Entropy = measure of disorder or randomness.

- $\Delta S > 0$  → disorder increases.
- $\Delta S < 0$  → disorder decreases.

### Entropy increases when:

- Solids → liquids → gases
- Number of gas particles increases
- Temperature increases
- Mixing occurs

### Common AP shortcut:

More gas particles = higher entropy.

### Gibbs Free Energy ( $\Delta G$ )

Predicts spontaneity.

- $\Delta G < 0$  → spontaneous
- $\Delta G = 0$  → equilibrium
- $\Delta G > 0$  → nonspontaneous

**Mnemonic:** "Good Horses Take Sugar" ( $G = H - T\cdot S$ ).

#### Mini formula box

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

### Temperature & Spontaneity

- At equilibrium:  $\Delta G = 0$ .
- Standard free energy change relates to equilibrium constant.

#### Spontaneity

$\Delta H$	$\Delta S$	Spontaneity
-	+	Always spontaneous
+	-	Never spontaneous
-	-	Low temperature
+	+	High temperature

### Free Energy & Equilibrium

- At equilibrium:  $\Delta G = 0$ .
- Standard free energy change relates to equilibrium constant.

### Reaction Energy Diagrams

- Peak height = activation energy ( $E_a$ ).
- Difference between products and reactants =  $\Delta H$ .
- Catalyst lowers  $E_a$ , not  $\Delta H$  or  $\Delta G$ .

### Key distinction:

- Thermodynamics → if reaction occurs.
- Kinetics → how fast it occurs.

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