

Changes and Properties of Matter

Physical Properties of Matter

- **Physical Changes:** Changes that change only the appearance of a substance, not its chemical identity.
- **Physical Properties:** Properties that can be observed through physical change

Physical Change

Physical Property

- **Melting**
 - **Boiling**
 - **Dissolving**
 - **Evaporating**
 - **Crushing**
 - **Stretching**
- **Melting point** (the temperature at which a substance turns from a solid to a liquid)
 - **Boiling point** (the temperature at which a substance turns from a liquid to a gas)
 - **Solubility** (the amount of solute that can be dissolved in a solvent)
 - **Vapor pressure** (the pressure exerted by a vapor at vapor-liquid equilibrium)
 - **Malleability** (the ability to be hammered or rolled into thin sheets)
 - **Ductility** (the ability to be stretched into a wire)
 - **Density** (the mass of a substance per unit volume)
 - **Electrical & Heat Conductivity** (the ability to pass heat or electricity through a substance)

Chemical Properties and Changes

- **Chemical Changes:** Changes that result in changing the chemical composition of a substance. Can be reversed only by another chemical change.
- **Chemical Properties:** Properties that can only be observed through chemical change.

Chemical Change

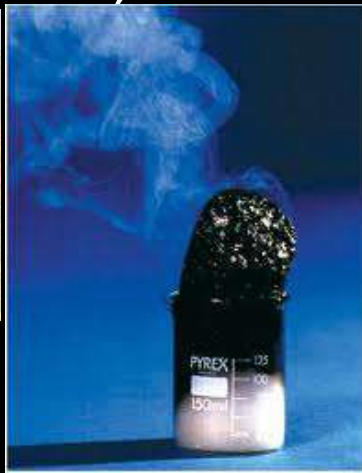
- **Corrosion of metals, flammability**
- **Chemical decomposition** e.g. hydrogen peroxide decomposes to form water and oxygen, but water does not decompose spontaneously
- **Combustion releases heat** (exothermic) Rarely do chemical changes absorb heat (endothermic)

Chemical Property

- **Reactivity** (the likelihood of one substance to undergo a chemical reaction with another substance)
- **Stability** (the likelihood that a substance will not decompose)
- **Heat of Reaction** (the energy absorbed or released by a chemical reaction)

Indications of a chemical reaction:

- Energy absorbed or released (heat and/or light)
- Color change
- Gas production (bubbling, fizzing, or odor change)
- formation of a precipitate- a solid that separates from solution (won't dissolve)
- Irreversibility- not easily reversed
- new substances produced, old substances destroyed.



Checking for understanding

Explain why are these processes physical changes:

- Boiling
- Condensation
- Shredding paper

• Explain why are these processes chemical changes:

- Burning paper
- Rusting bicycle
- Frying eggs

Change in Energy

Energy

Different types of energy exist (heat, potential, kinetic, chemical, nuclear etc.)

Heat - the energy transferred between objects that are at different temperatures. Unit of heat is joules J.

According to the Law of Conservation of Energy, energy cannot be created or destroyed. It can only be transferred or transformed. The total amount of energy in a system remains constant.

- Add a potential energy diagram
- To draw from scratch
- Do more calculations on enthalpy/energies

Potential Energy Diagram

<http://www.kentchemistry.com/links/Kinetics/PEDiagrams.htm>

Potential Energy Diagram

- Potential Energy Diagrams show changes in energy during chemical reactions.
- CHEMICAL EQUATIONS



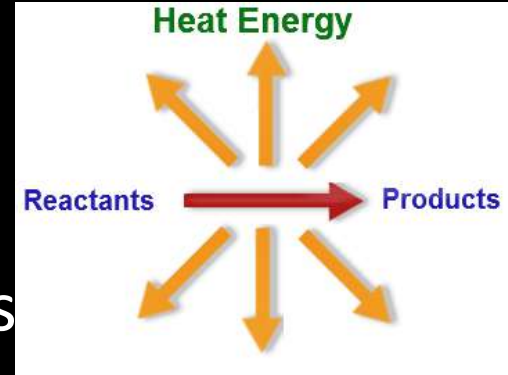
Reactant Products

- In chemical reactions, reactants react to form products with an associated **release** or **absorption** of energy.

Change in Heat Energy

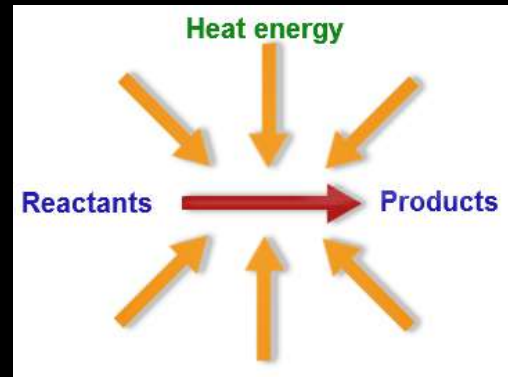
1. Exothermic Reaction

- The change in heat (ΔH) is NEGATIVE.
- Heat is released into the surroundings

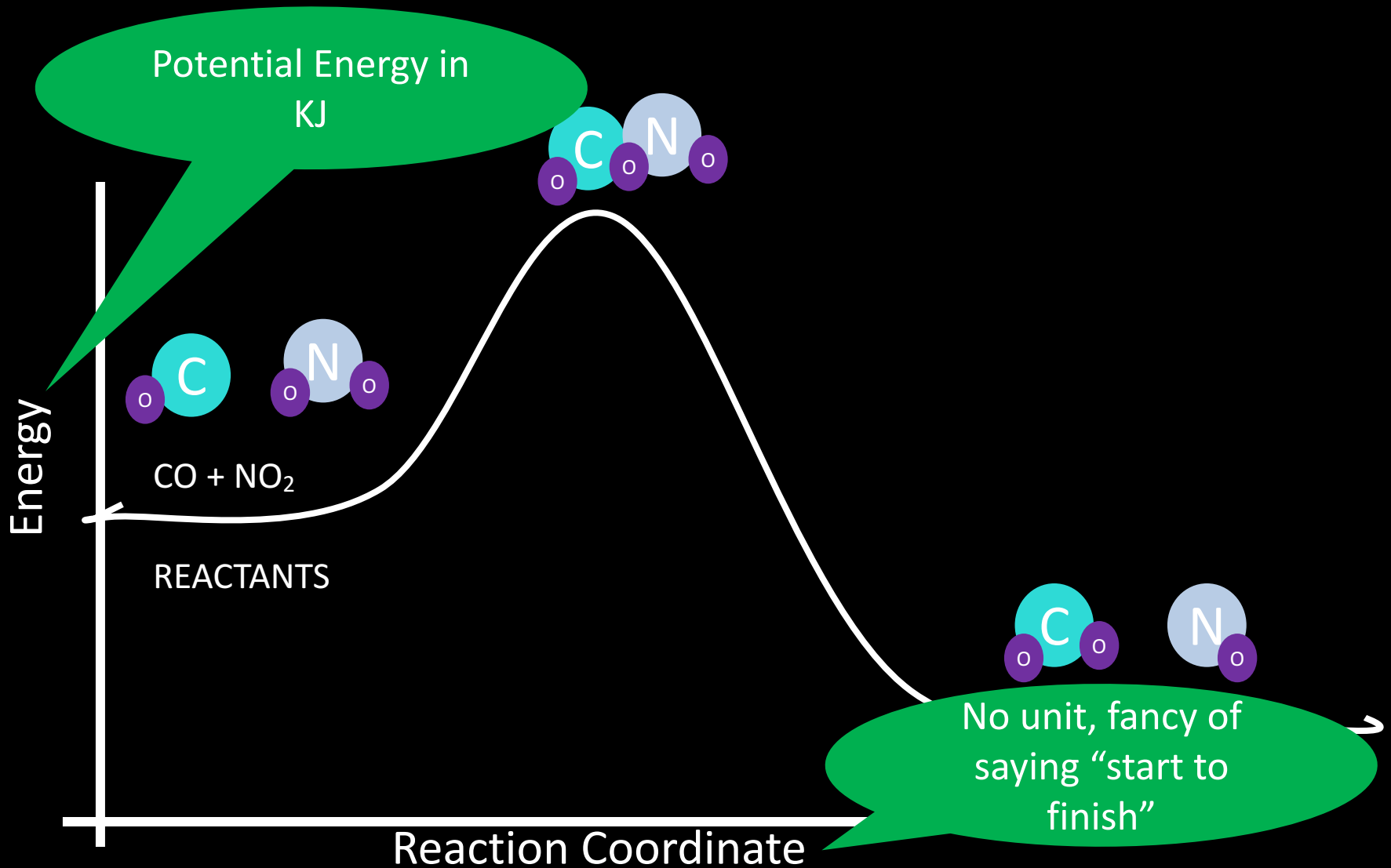


2. Endothermic Reaction

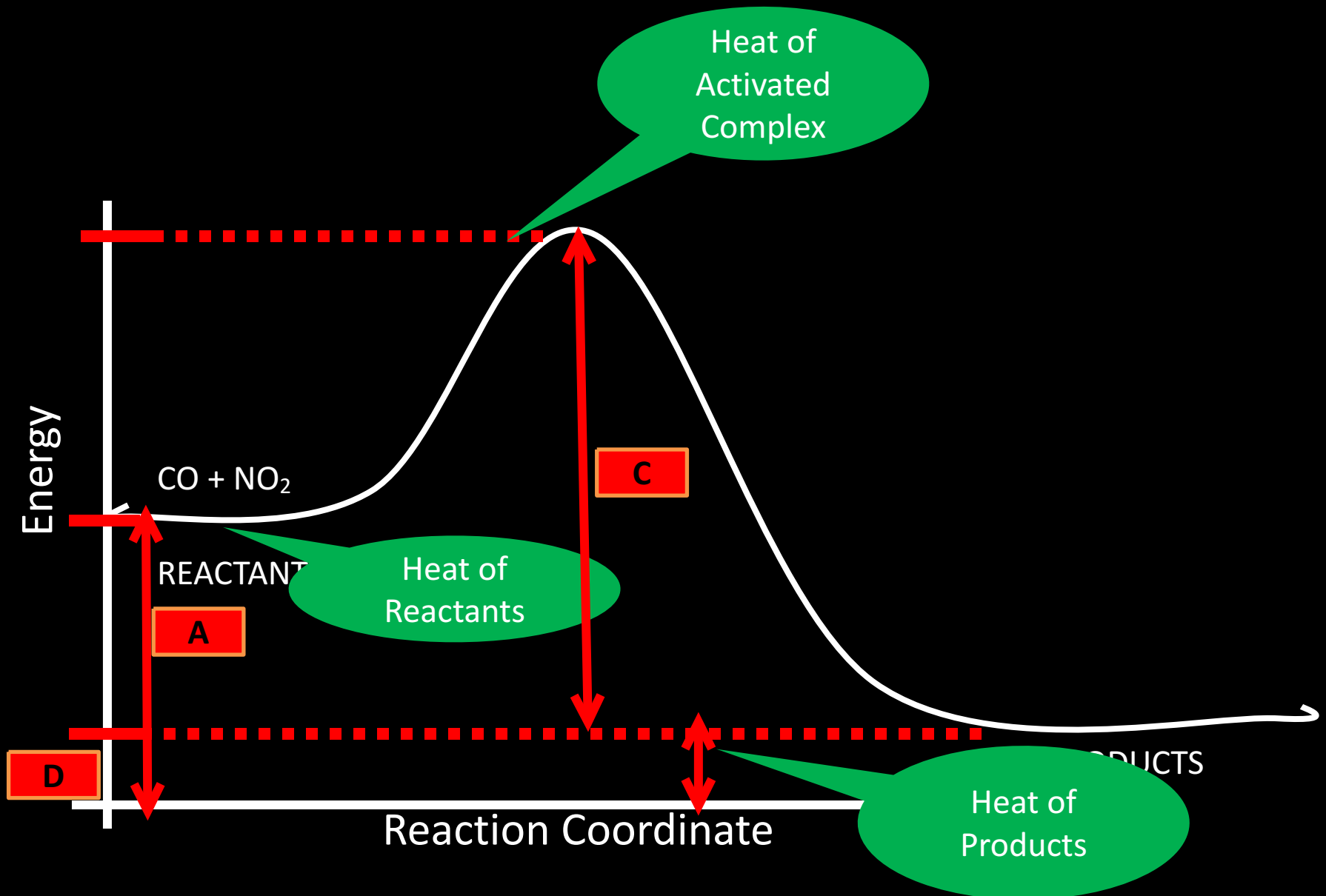
- The change in heat (ΔH) is POSITIVE.
- Heat is absorbed by the system



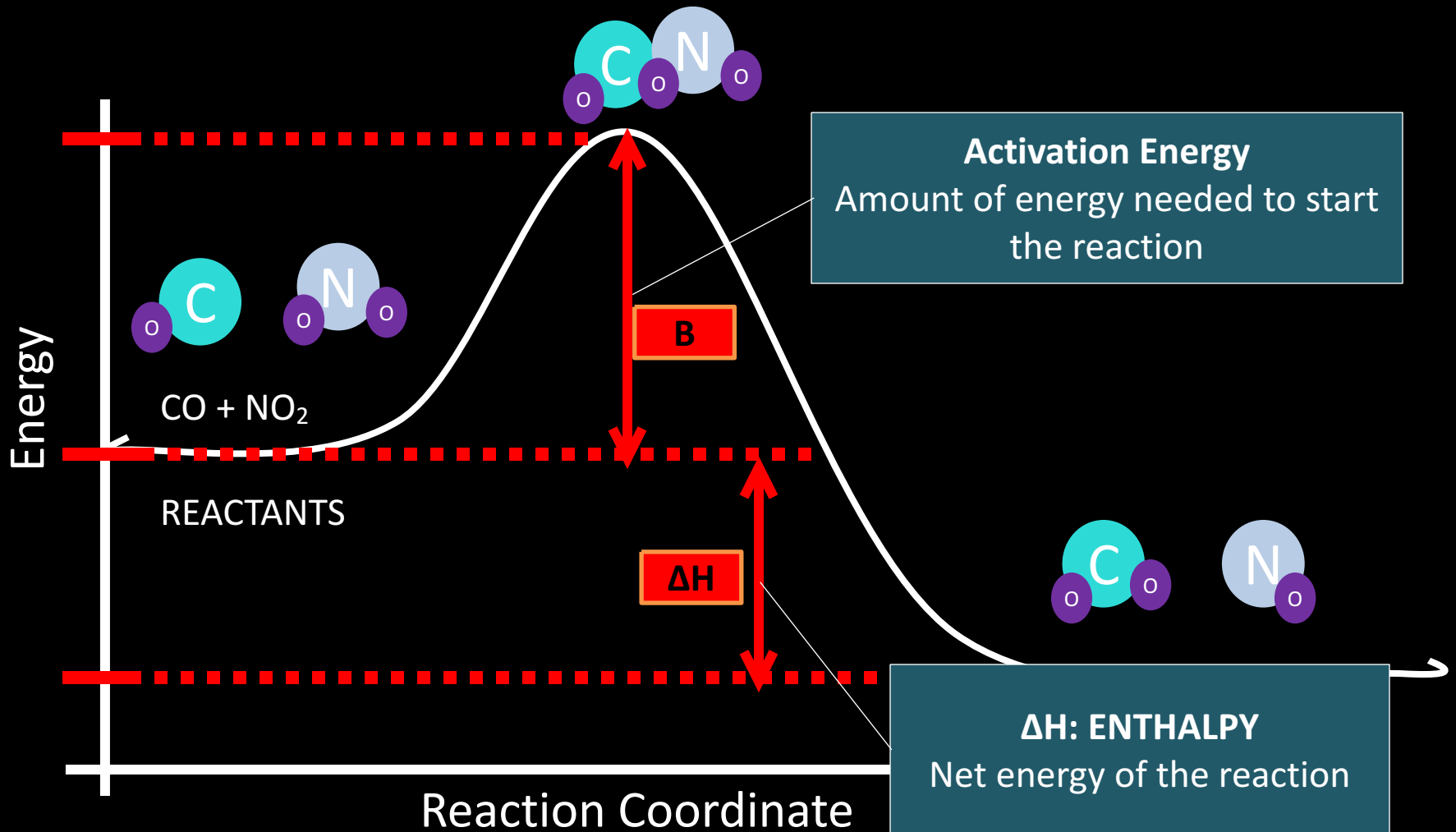
Potential Energy Diagram



Potential Energy Diagram



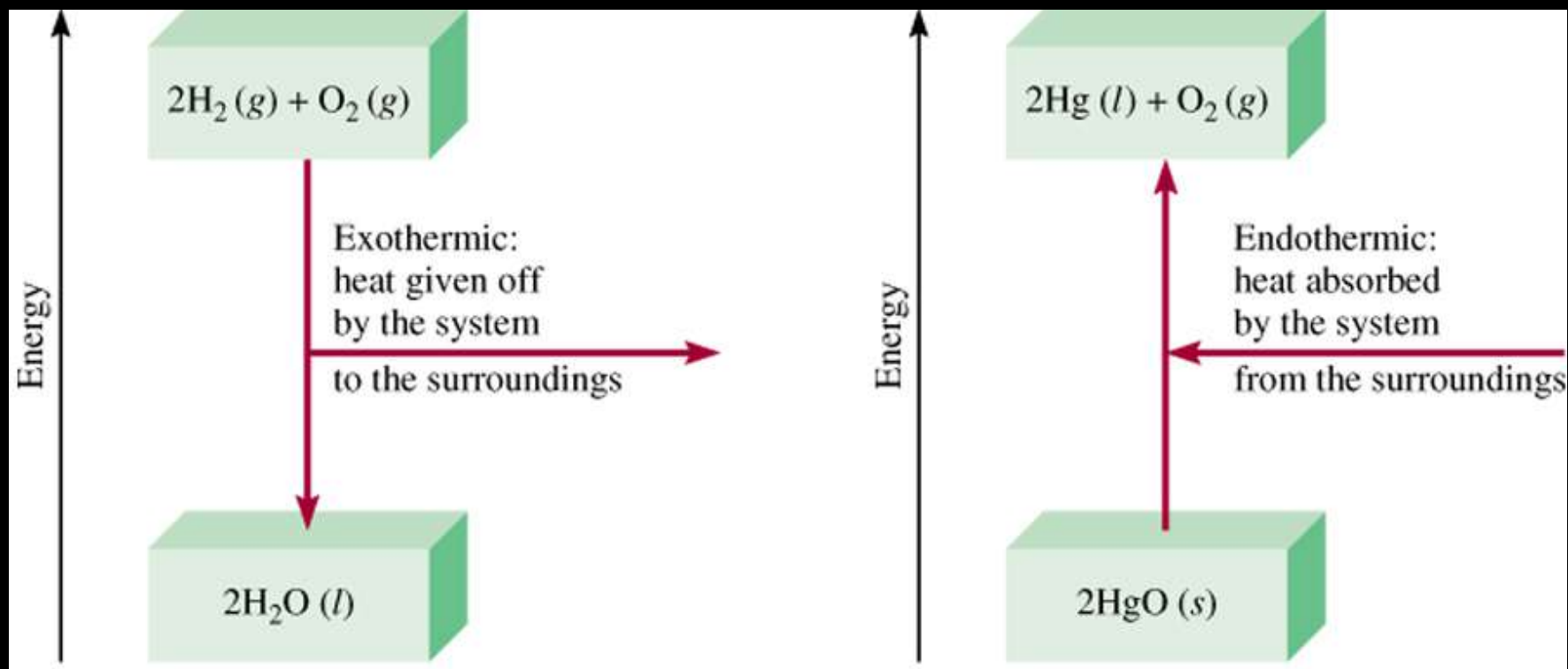
Potential Energy Diagram



Enthalpy (H) is used to quantify the heat flow into or out of a system in a process that occurs at constant pressure.

$$\Delta H = H(\text{products}) - H(\text{reactants})$$

ΔH = heat given off or absorbed during a reaction at constant pressure

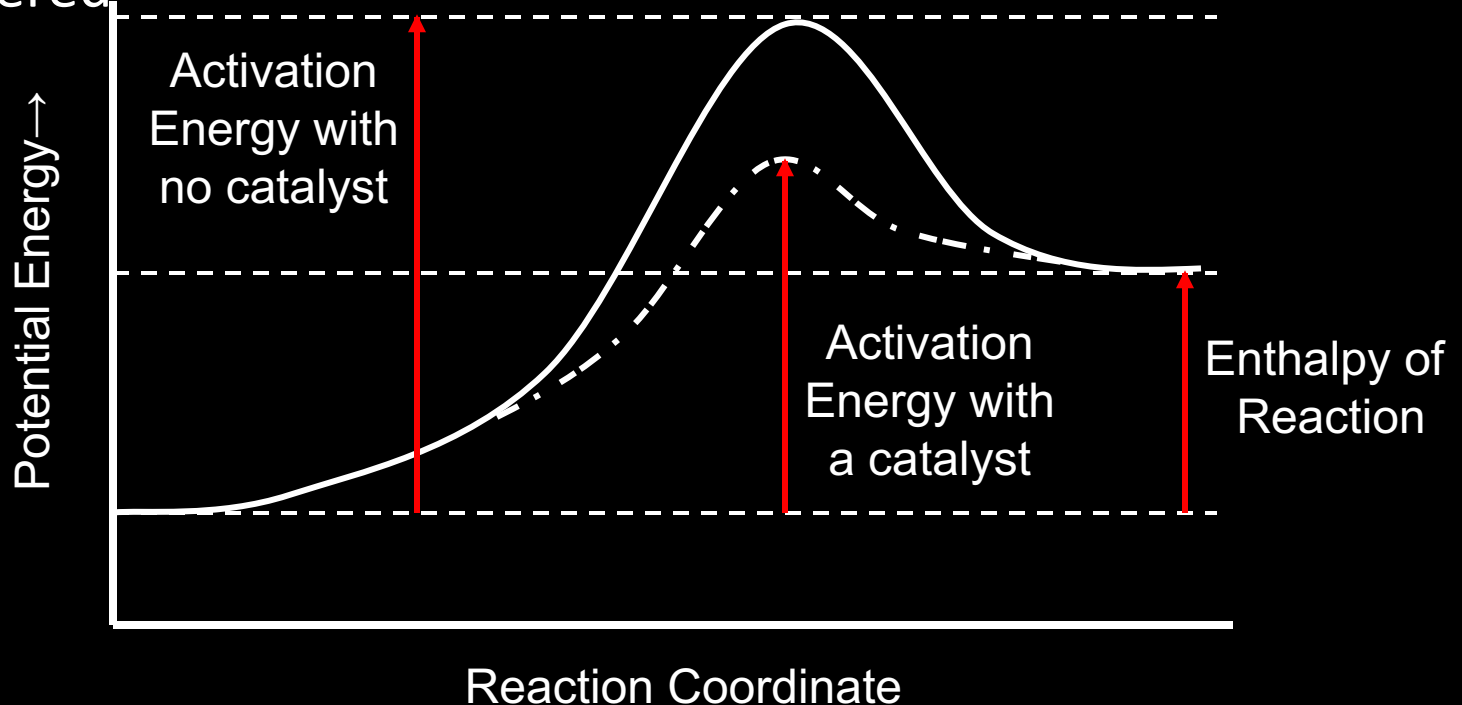


$$H_{\text{products}} < H_{\text{reactants}}$$
$$\Delta H < 0$$

$$H_{\text{products}} > H_{\text{reactants}}$$
$$\Delta H > 0$$

Potential Energy Diagrams

- A catalyst is a substance that changes the speed of reactions by changing the **activation energy**.
- Catalysts speed up chemical reaction by **lowering** the activation energy
- Catalysts are not consumed during the reaction and can be recovered

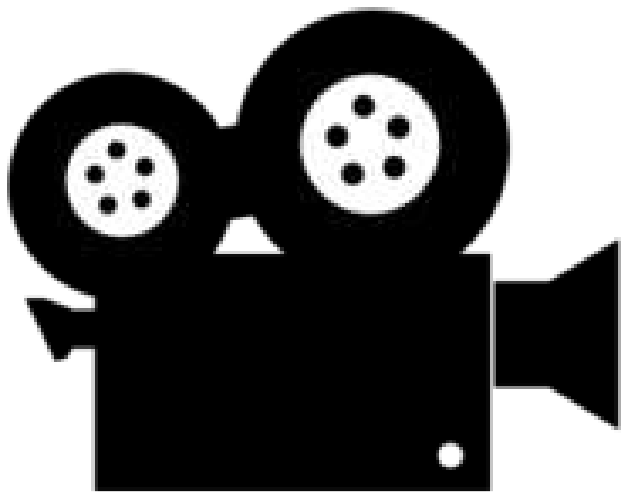


Click Below for the Video Lectures

[Activation Energy](#)

[Reaction Coordinate](#)

[Catalyst](#)



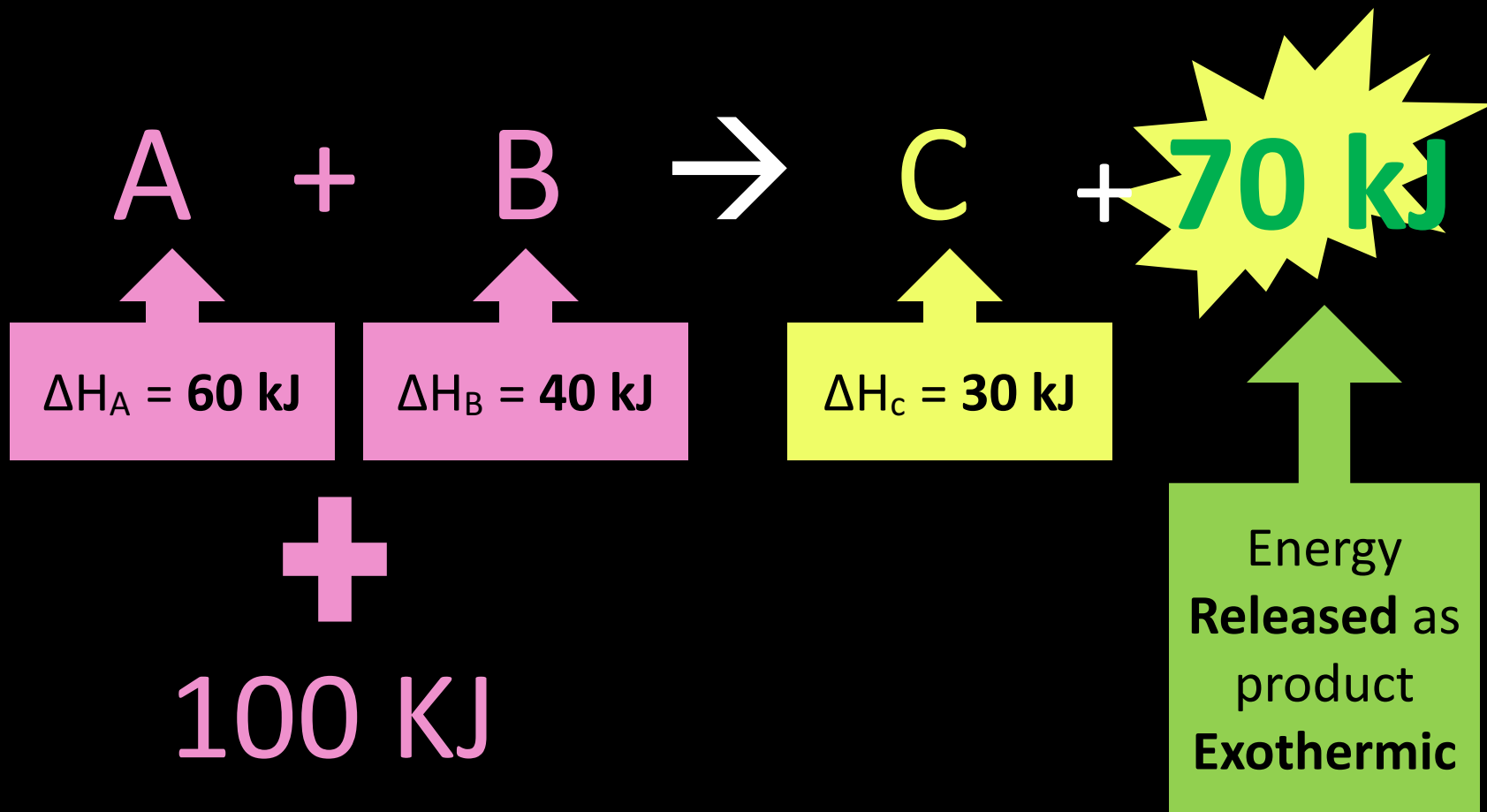
Exothermic
and
Endothermic

Exothermic Reaction



- a) **Heat is released as a product.**
- b) This heat is released into the environment surrounding the reaction, causing the temperature of the surroundings to increase.
- c) **The products have less energy than the reactants after release.**
- d) Since the products have less energy stored in their chemical bonds, they are more stable than the reactants were. Burning paper is exothermic. The ash formed by the burning is not flammable and will not burn.

General Exothermic Reaction



Law of conservation of energy states that the total energy of an isolated system cannot change.

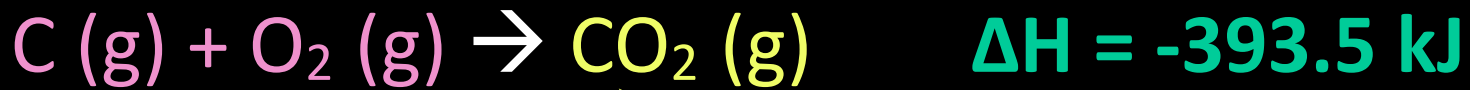
Exothermic Reaction

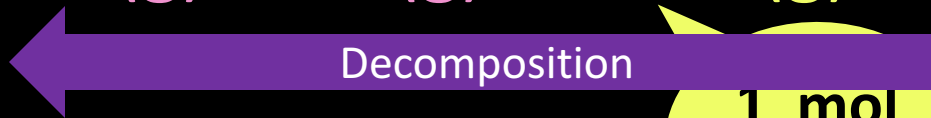


- The minus sign in front of the enthalpy 393.5 kJ indicates that this reaction is exothermic, that energy was released.
- This energy can be placed on the products side, as follows:



Synthesis 



Decomposition 

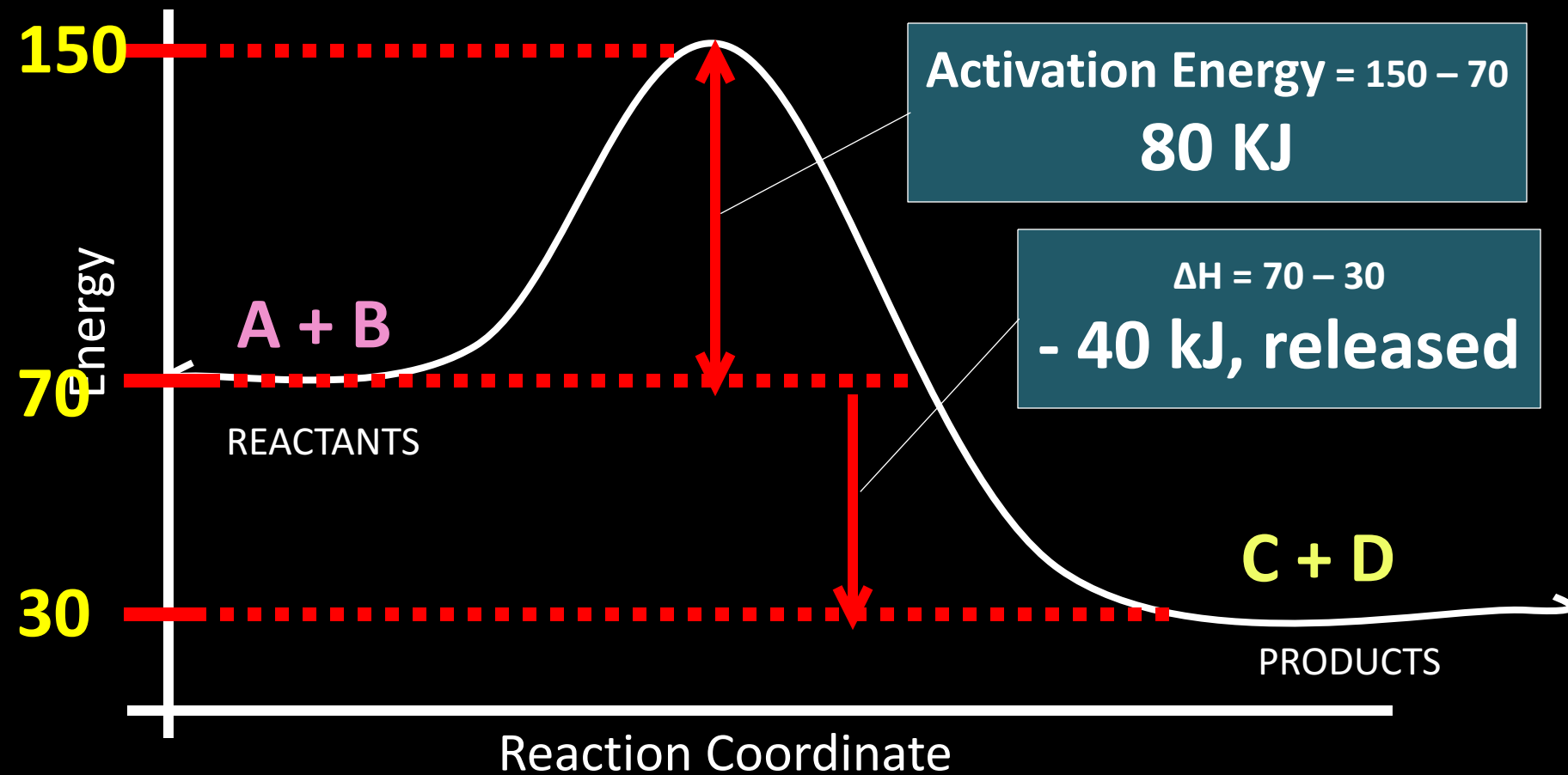
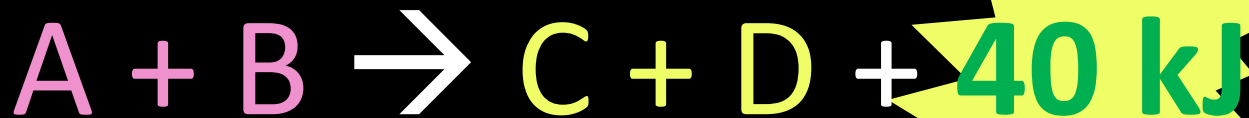
1 mol
of CO₂

- If we were to synthesize 2.3 moles of CO₂, how many kJ would be released?

$$\begin{array}{r|l} 2.3 \text{ mol CO}_2 & 393.5 \text{ kJ} \\ \hline & 1 \text{ mol CO}_2 \end{array} = -910 \text{ kJ}$$

If $\Delta H_{\text{synthesis}} = -393.5 \text{ kJ/mole}$,
then $\Delta H_{\text{decomposition}} = +393.5 \text{ kJ/mole}$

Potential Energy Diagram - Exothermic

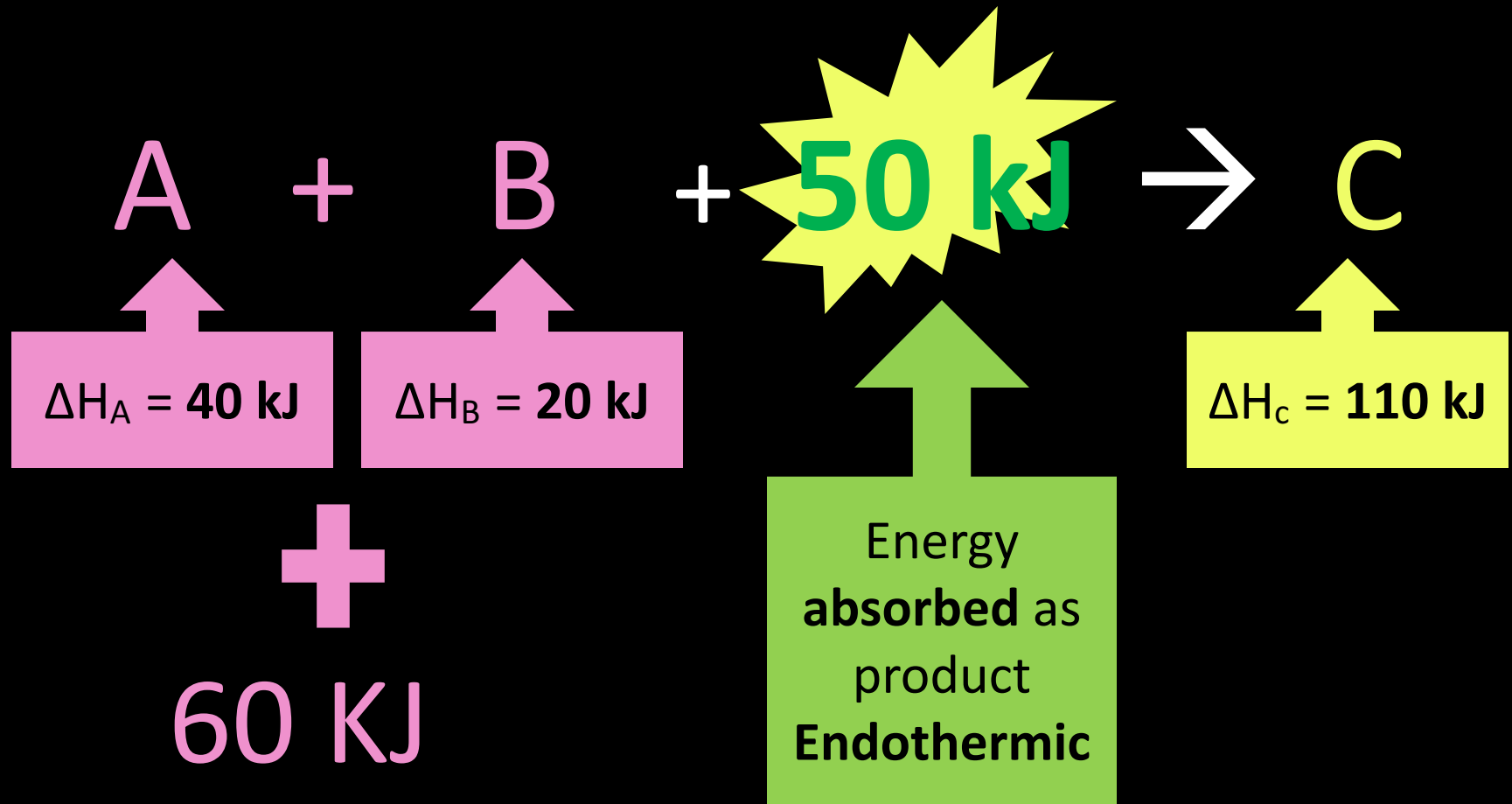


Endothermic Reaction



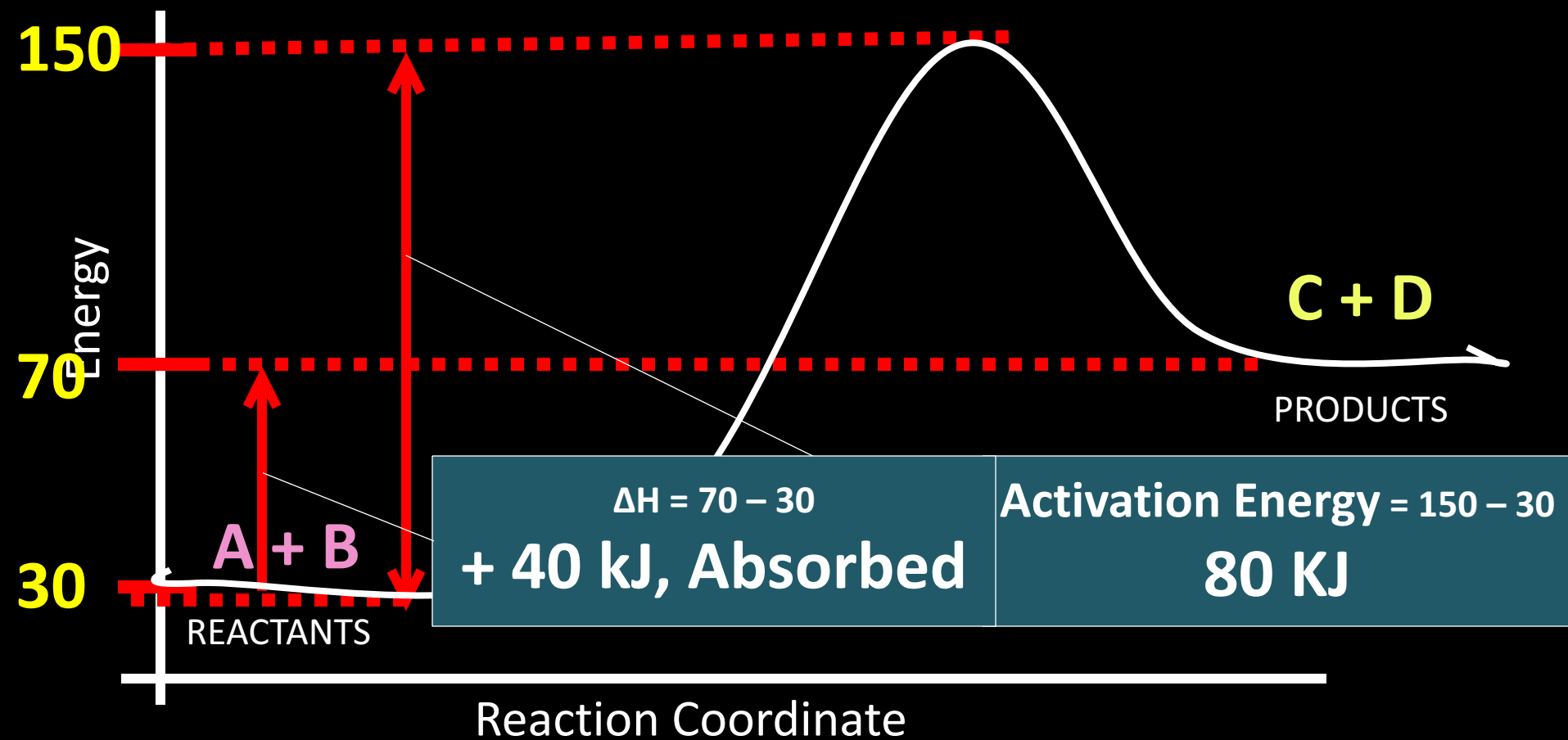
- a) **Heat is absorbed by the reactants.**
- b) This heat is absorbed from the environment surrounding the reaction, and the temperature of the surroundings decreases.
- c) **The products have more energy than the reactants after absorption.**
- d) This energy is stored in the chemical bonds of the products

General Endothermic Reaction



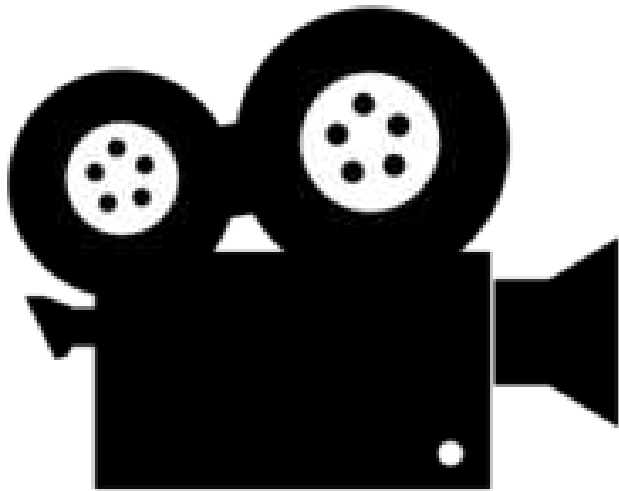
Law of conservation of energy states that the total energy of an isolated system cannot change.

Potential Energy Diagram - Endothermic



Click Below for the Video Lectures

Exothermic and Endothermic



Calorimetry

Calorimetry

$$q = m C \Delta T$$

units

Joules (J)

Grams (g)

J/g°C

°C

What Each Variable Means

q is the quantity of heat that is absorbed or released by a physical or chemical change

m is the mass of water in the calorimeter cup that absorbs heat from the change or releases heat to the change.

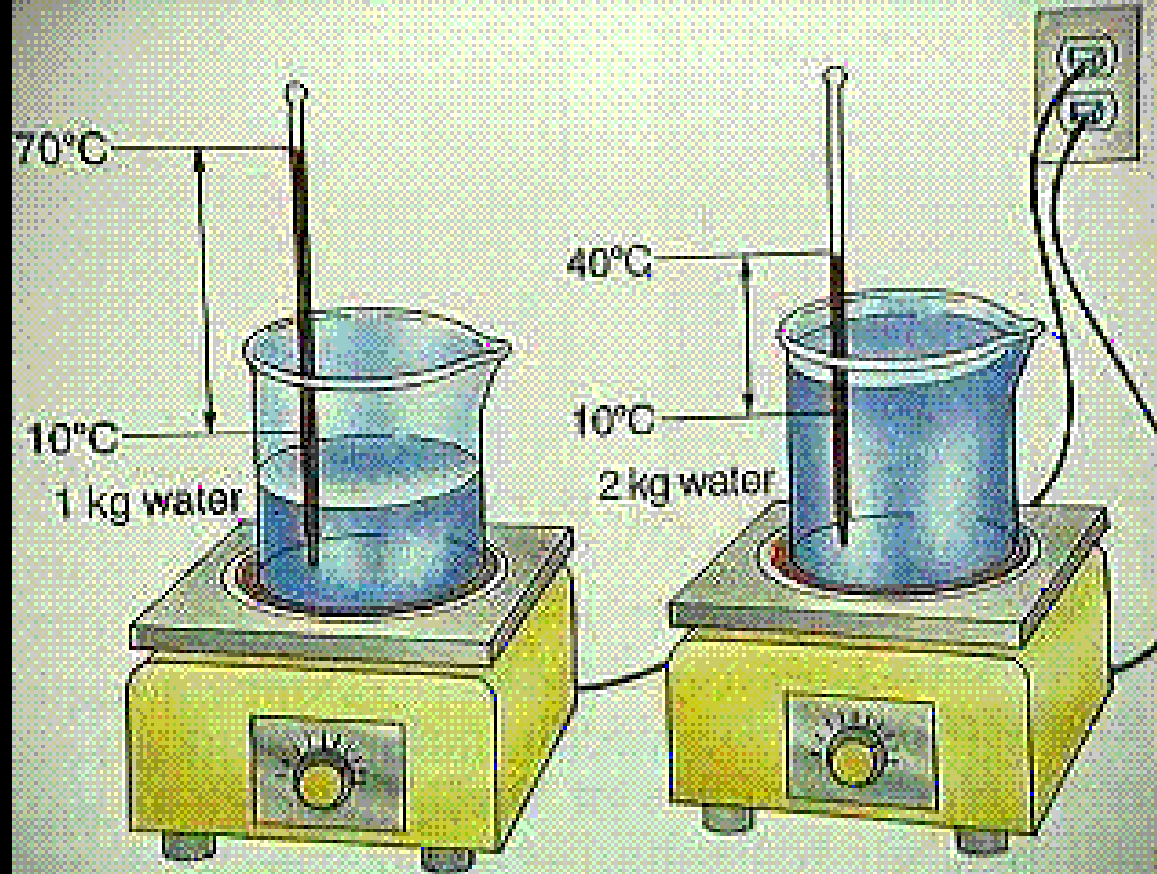
C is the specific heat of water, the rate at which water gains or loses heat if energy is absorbed or removed from it.

ΔT is the Temperature change of the water in the calorimeter cup as a result of the physical or chemical change.

C

Specific Heat

The amount of heat required to raise the temperature of **one gram** of substance by **one degree Celsius**.



	cal/g°C	J/g°C (or J/gK)
water	1.00	4.18
aluminum	0.22	0.90
copper	0.093	0.39
silver	0.057	0.24
gold	0.031	0.13

Calorimetric Formulas =

No Phase Change: $Q = m(\Delta T)C_p$

Latent Heat of Fusion: $Q = m\Delta H_{\text{fus}}$

Latent Heat of Vaporization: $Q = m\Delta H_{\text{vap}}$

Use this equation on the back of periodic table

Specific Heat of water. Use the one for correct unit! They are NOT equations!!!

Specific Heat of Water: $C_p(\text{H}_2\text{O}) = 1.00 \frac{\text{cal}}{\text{g}^\circ\text{C}} = 4.18 \frac{\text{J}}{\text{g}^\circ\text{C}}$

Latent Heat of Fusion of Water: $\Delta H_{\text{fus}}(\text{H}_2\text{O}) = 80 \frac{\text{cal}}{\text{g}} = 334 \frac{\text{J}}{\text{g}}$

Latent Heat of Vaporization of Water: $\Delta H_{\text{vap}}(\text{H}_2\text{O}) = 540 \frac{\text{cal}}{\text{g}} = 2260 \frac{\text{J}}{\text{g}}$

1. Circle the numbers, underline what you are looking for.
2. Make a list of number you circled using variables.
3. Write down the formula
4. Derive the formula to isolate the variable you are looking for.
5. Plug in the numbers

- How many joules are absorbed by 100.0 grams of water if the temperature is increased from 35.0°C to 50.0°C?

$$m = 100.0 \text{ g}$$

$$T_i = 35.0^\circ\text{C}$$

$$T_f = 50.0^\circ\text{C}$$

$$q = ?$$

$$c = 4.18 \text{ j/g}^\circ\text{C}$$

$$q = mc\Delta T$$

$$q = (100.0 \text{ g})(4.18 \text{ j/g}^\circ\text{C})(50.0^\circ\text{C} - 35.0^\circ\text{C})$$

$$q = 6270 \text{ J}$$

1. Circle the numbers, underline what you are looking for.
2. Make a list of number you circled using variables.
3. Write down the formula
4. Derive the formula to isolate the variable you are looking for.
5. Plug in the numbers

• 300. J of energy is absorbed by a 50. g sample of water in a calorimeter. How much will the temperature change by?

$$q = 300 \text{ J}$$

$$m = 50 \text{ g}$$

$$\Delta T = ?$$

$$C = 4.18 \text{ J/g}^\circ\text{C}$$

$$q = mc\Delta T$$

$$\Delta T = q / mc$$

$$\Delta T = 300 \cancel{\text{J}} / (50 \cancel{\text{g}})(4.18 \cancel{\text{J/g}^\circ\text{C}})$$

$$\Delta T = = 1.4354066 \text{ }^\circ\text{C}$$

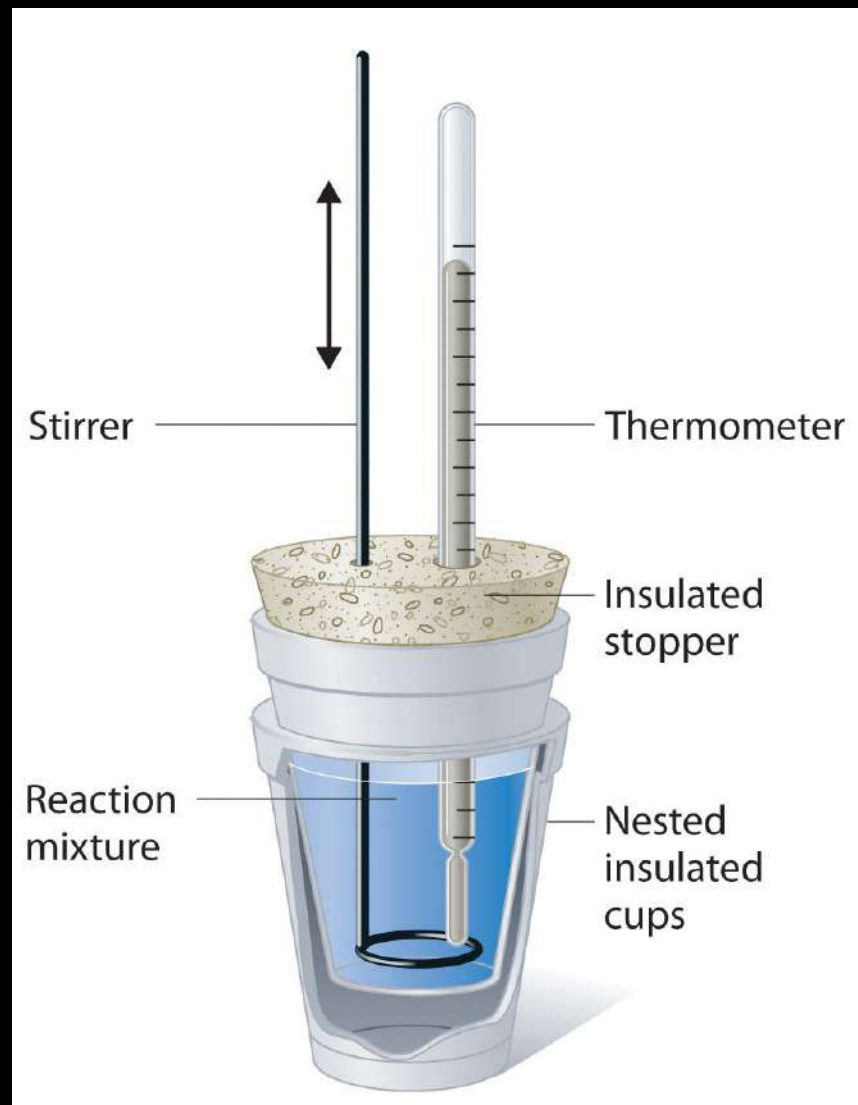
$$\Delta T = = 1.4^\circ\text{C}$$

Checking for understanding

1. How many joules are required to raise the temperature of 100. grams of water from 20.0°C to 30.0°C ?
2. How many grams of water can be heated from 20.0°C to 75.0°C using 2500. J?

Calorimetry

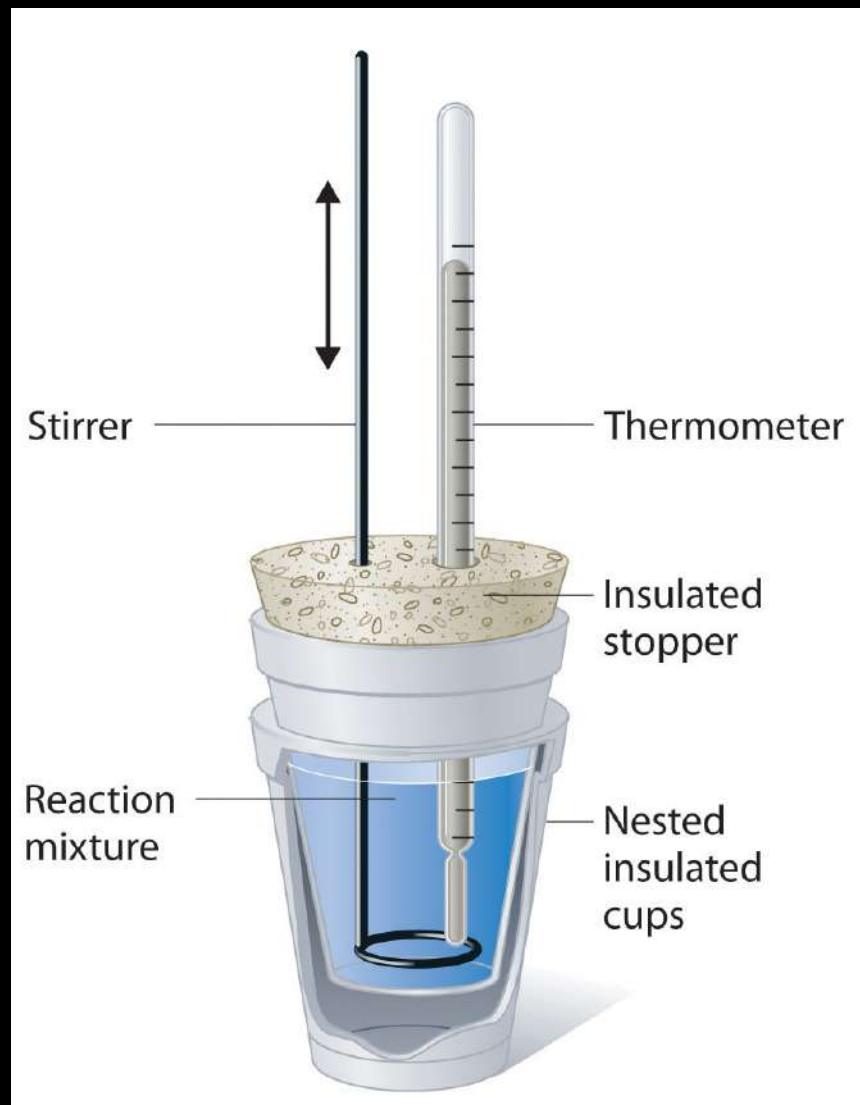
- We use an insulated device called a **calorimeter** to measure this **heat transfer**.
- A typical device is a “**coffee cup calorimeter**.”



Calorimetry

To measure ΔH for a reaction

1. dissolve the reacting chemicals in known volumes of water
2. measure the initial temperatures of the solutions
3. mix the solutions
4. measure the final temperature of the mixed solution



Calorimetry

- The heat generated by the **reactants** is absorbed by the **water**.
- We know the mass of the water, m_{water} .
- We know the change in temperature, ΔT_{water} .
- We also know that water has a specific heat of $C_{\text{water}} = 4.18 \text{ J/}^\circ\text{C-g}$.
- We can calculate the heat of reaction by:
- $q_{\text{sys}} = \Delta H = -q_{\text{surr}} = -m_{\text{water}} \times C_{\text{water}} \times \Delta T_{\text{water}}$

When 25.0 mL of water containing 0.025 mol of HCl at 25.0°C is added to 25.0 mL of water containing 0.025 mol of NaOH at 25.0°C in a coffee cup calorimeter, a reaction occurs. Calculate ΔH (in kJ) during this reaction if the highest temperature observed is 32.0°C. Assume the densities of the solutions are 1.00 g/mL.

Knowns: $V_{\text{final}} = V_{\text{HCl}} + V_{\text{NaOH}} = (25.0 + 25.0) \text{ mL} = 50.0 \text{ mL}$

$$D_{\text{water}} = 1.00 \text{ g/mL}$$

$$\Delta T_{\text{water}} = T_{\text{final}} - T_{\text{initial}} = 32.0^\circ\text{C} - 25.0^\circ\text{C} = +7.0^\circ\text{C}$$

$$C_{\text{water}} = 4.18 \text{ J/}^\circ\text{C-g}$$

Calculation: $m_{\text{water}} = 50.0 \text{ g}$

$$\Delta H = -m \times C \times \Delta T$$

$$= -(50.0 \text{ g})(4.18 \text{ J/}^\circ\text{C-g})(7.0^\circ\text{C})$$

$$= -1463 \text{ J}$$

$$= -1.5 \times 10^3 \text{ J}$$

Phase Change Heating and Cooling Curve

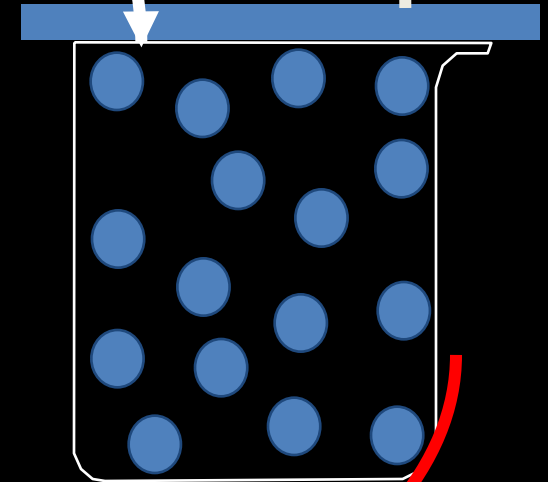
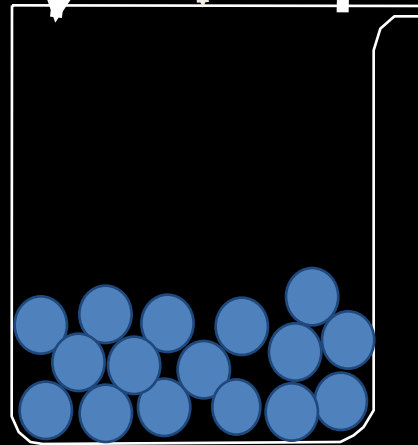
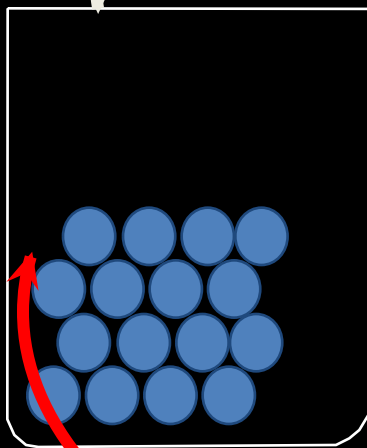
Sublime

Freeze

Condense

Melt

Evaporate



Solid

Liquid

Gas

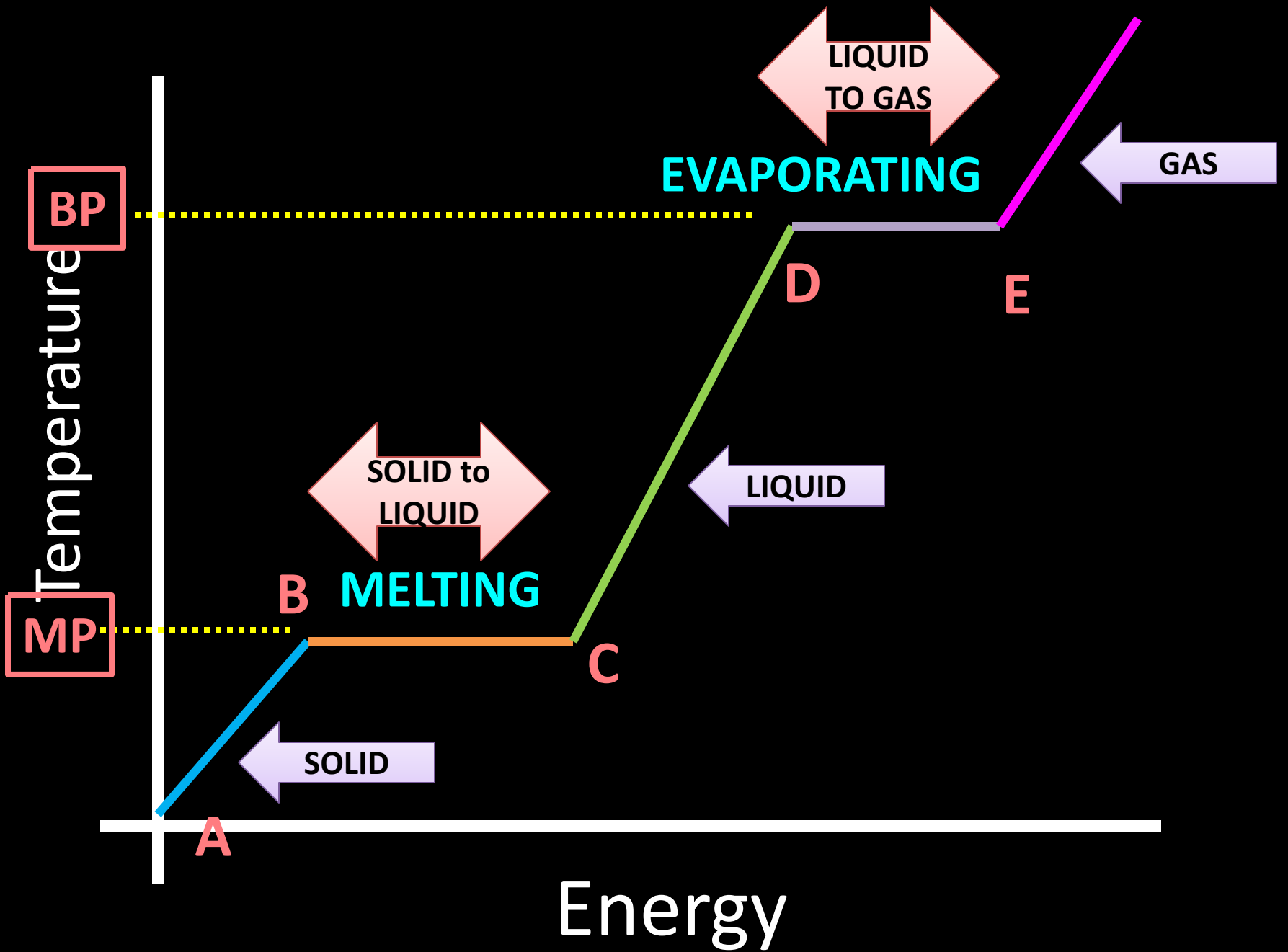
Phase Change

- During a phase change, there is a change in heat energy but no change in temperature.

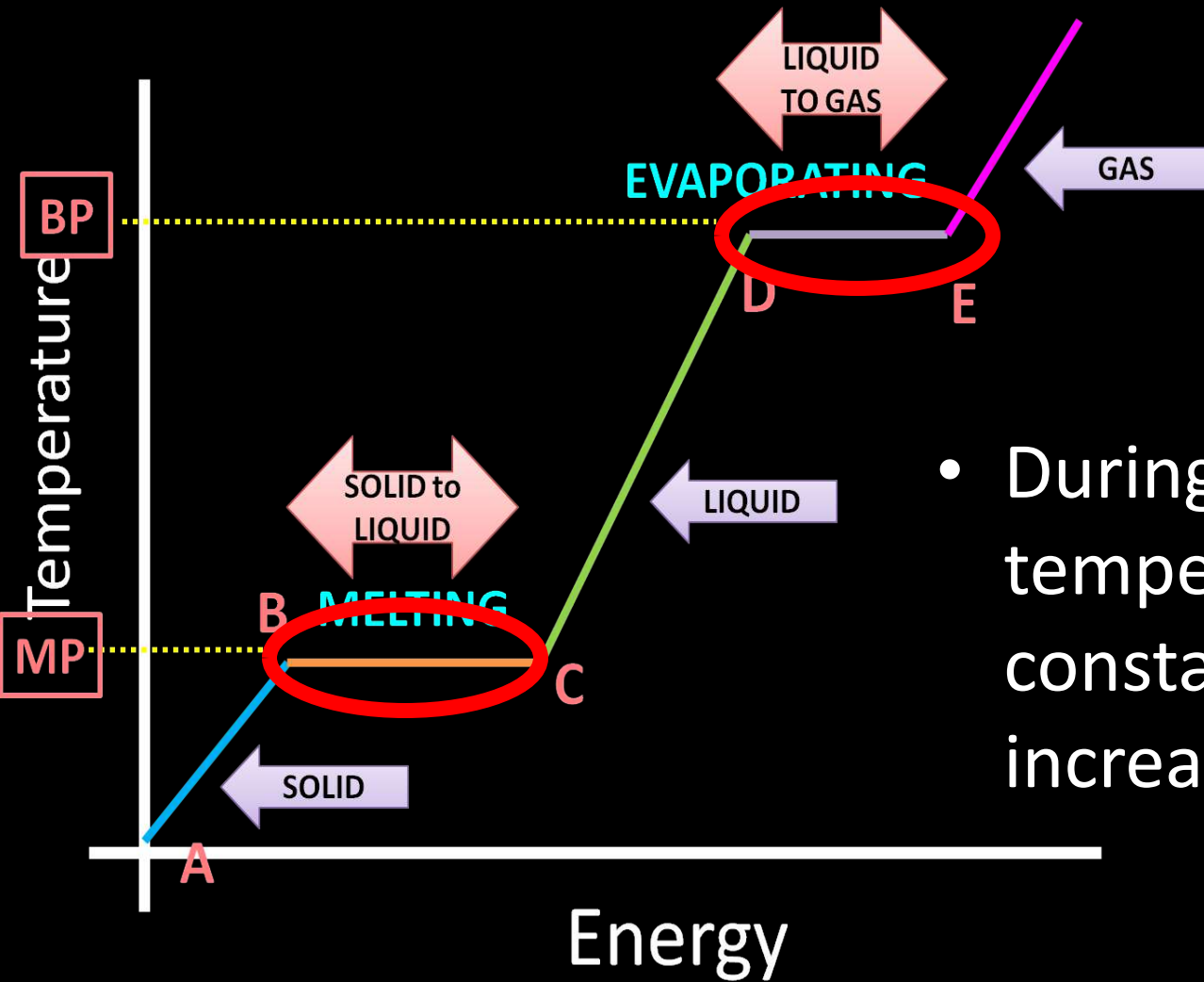
At the melting point, heat energy is being used to break down the crystalline lattice.

At the boiling point, heat energy is being used to convert the liquid to a gas.

During a phase change, heat is being used to change phase, not increase the kinetic energy of the particles. It is for this reason that **Temperature remains constant in a phase change while heat energy changes.**

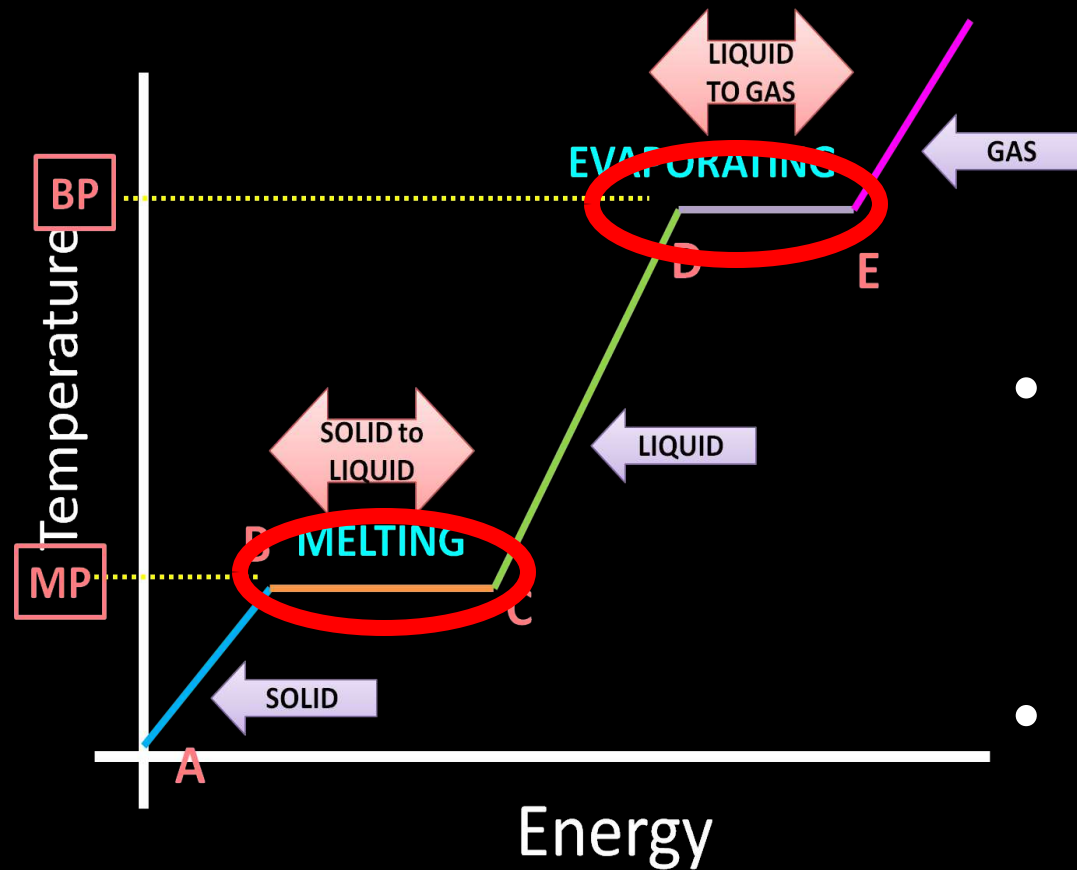


Phase Change



- During phase change, temperature stays constant but energy increases.

Phase Change



- Energy needed to melt is called **Latent Heat of Fusion** (ΔH_{fus})
- Energy need to vaporize is called **Latent Heat of Vaporization** ($\Delta H_{vaporization}$)

Constants

Volume of Ideal Gas at STP: $22.4 \frac{\text{L}}{\text{mol}}$

Speed of Light in a Vacuum: $c = 3.00 \times 10^8 \frac{\text{m}}{\text{s}}$

Specific Heat of Water: $C_p(\text{H}_2\text{O}) = 1.00 \frac{\text{cal}}{\text{g} \cdot ^\circ\text{C}} = 4.18 \frac{\text{J}}{\text{g} \cdot ^\circ\text{C}}$

Latent Heat of Fusion of Water: $\Delta H_{\text{fus}}(\text{H}_2\text{O}) = 80 \frac{\text{cal}}{\text{g}} = 334 \frac{\text{J}}{\text{g}}$

Latent Heat of Vaporization of Water: $\Delta H_{\text{vap}}(\text{H}_2\text{O}) = 540 \frac{\text{cal}}{\text{g}} = 2260 \frac{\text{J}}{\text{g}}$

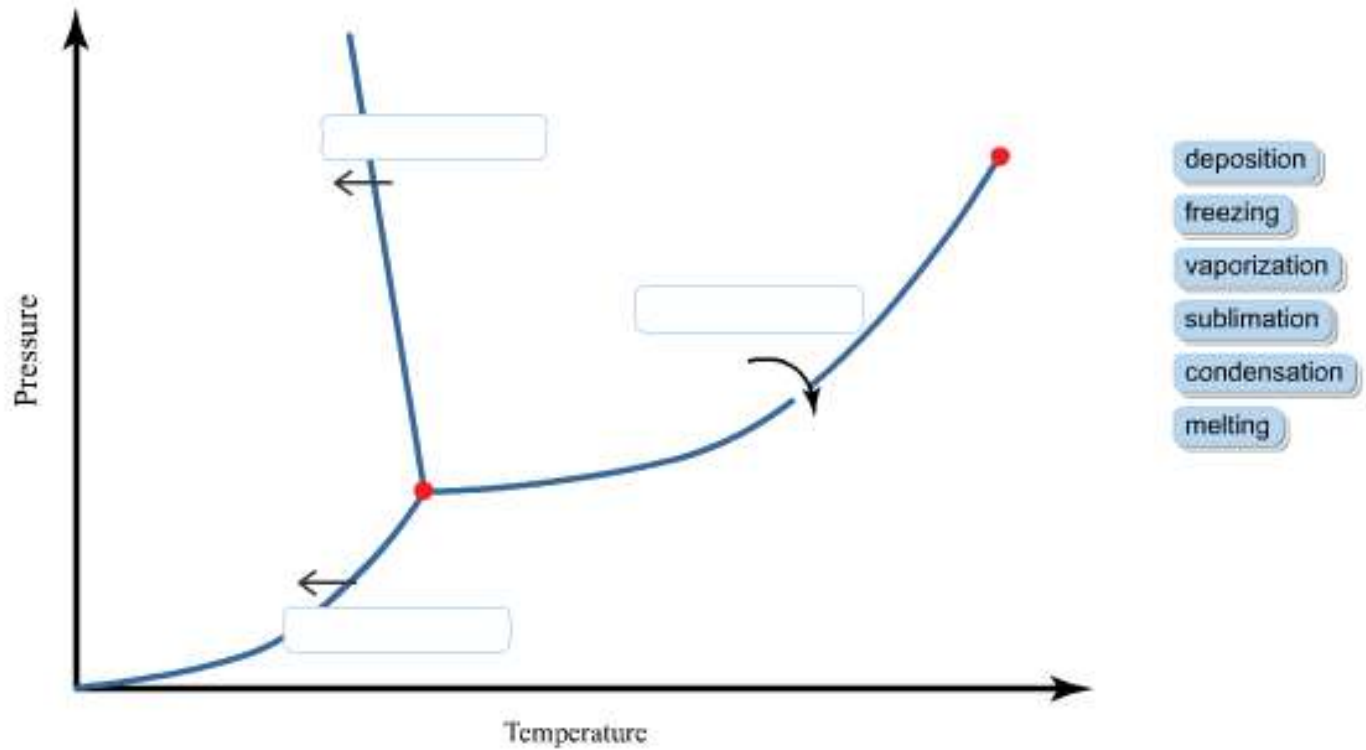
Calorimetric Formulas –

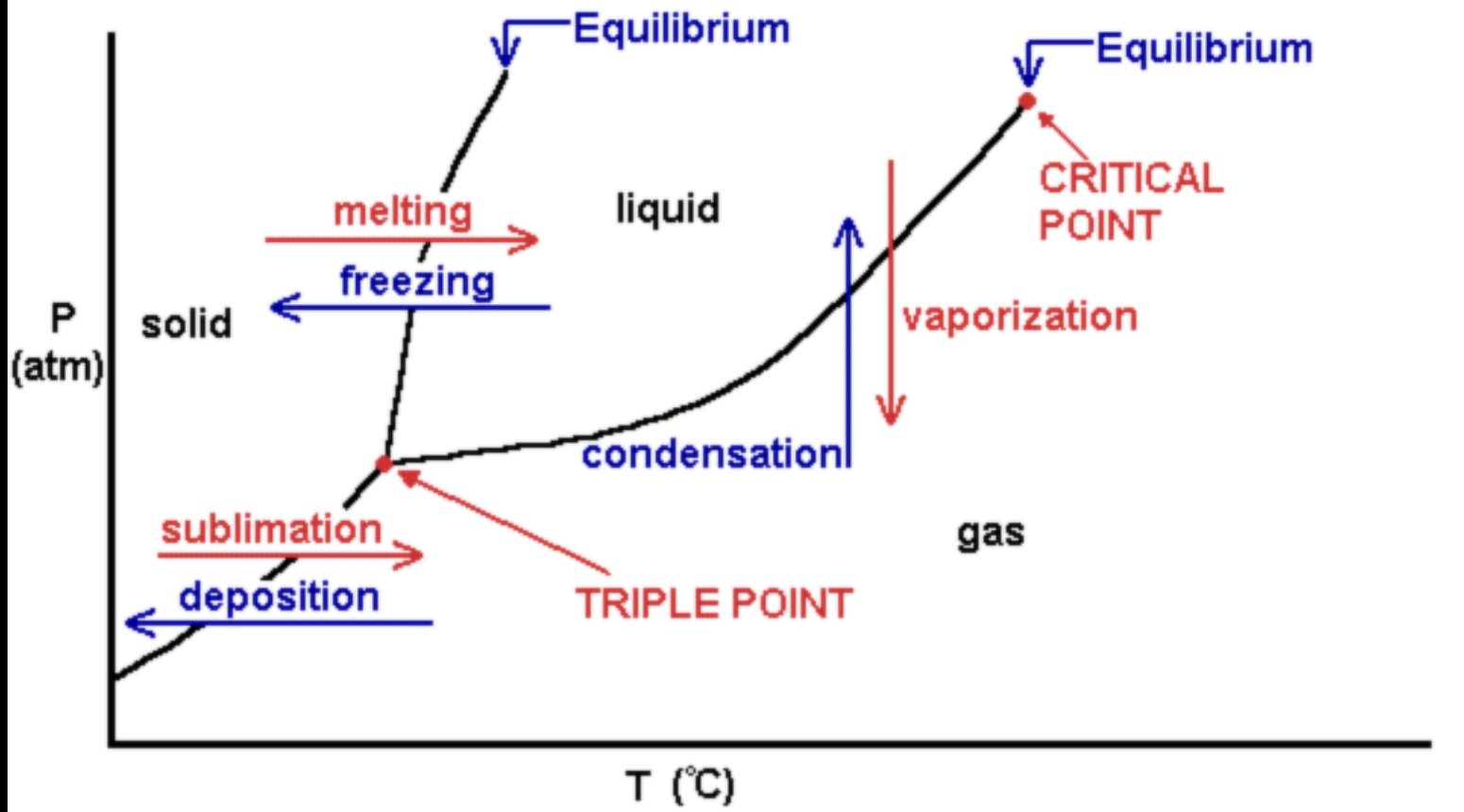
No Phase Change: $Q = m(\Delta T)C_p$

Latent Heat of Fusion: $Q = m\Delta H_{\text{fus}}$

Latent Heat of Vaporization: $Q = m\Delta H_{\text{vap}}$

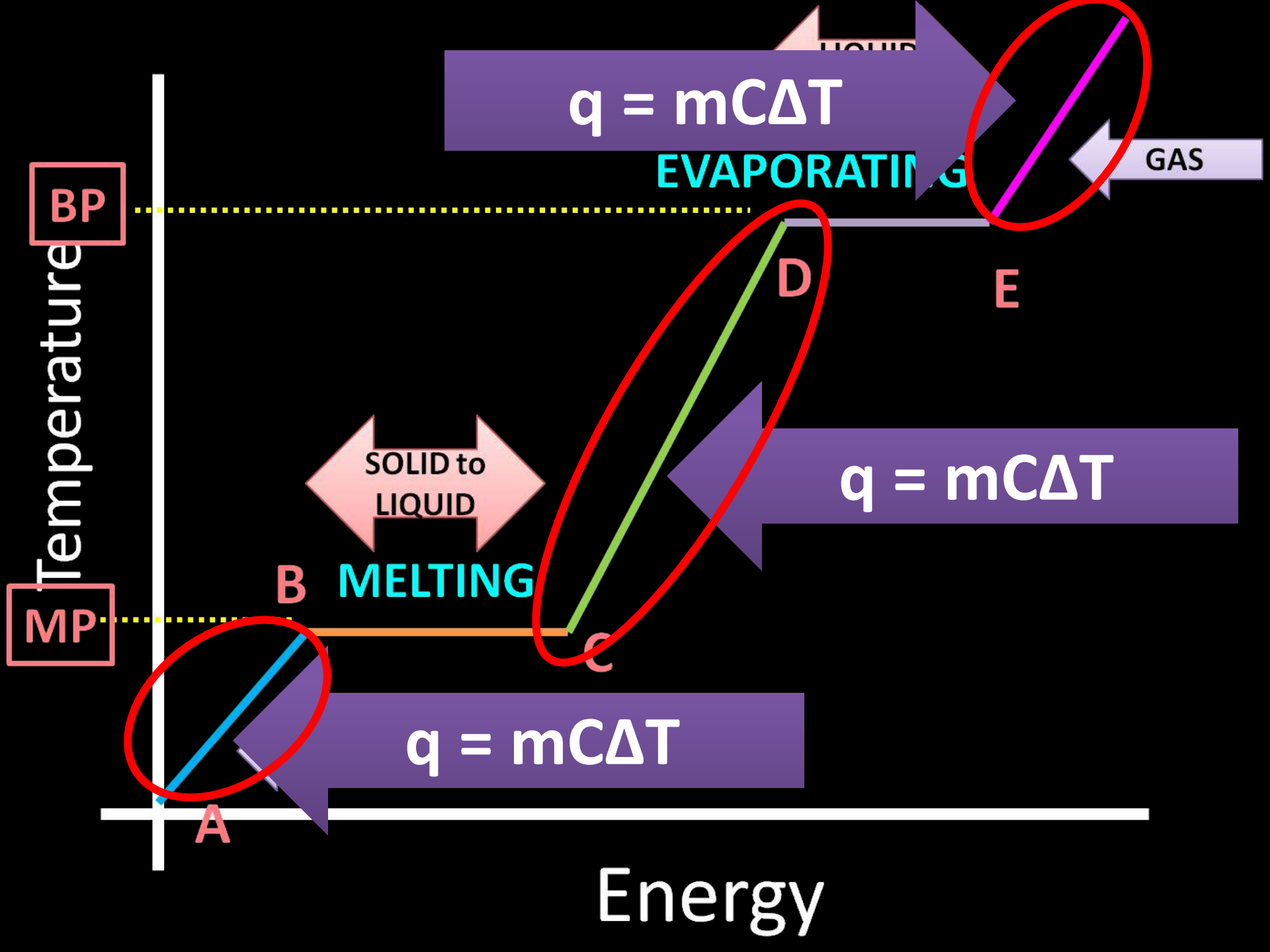
Identify the three phase changes indicated by each arrow below in the phase diagram of water.

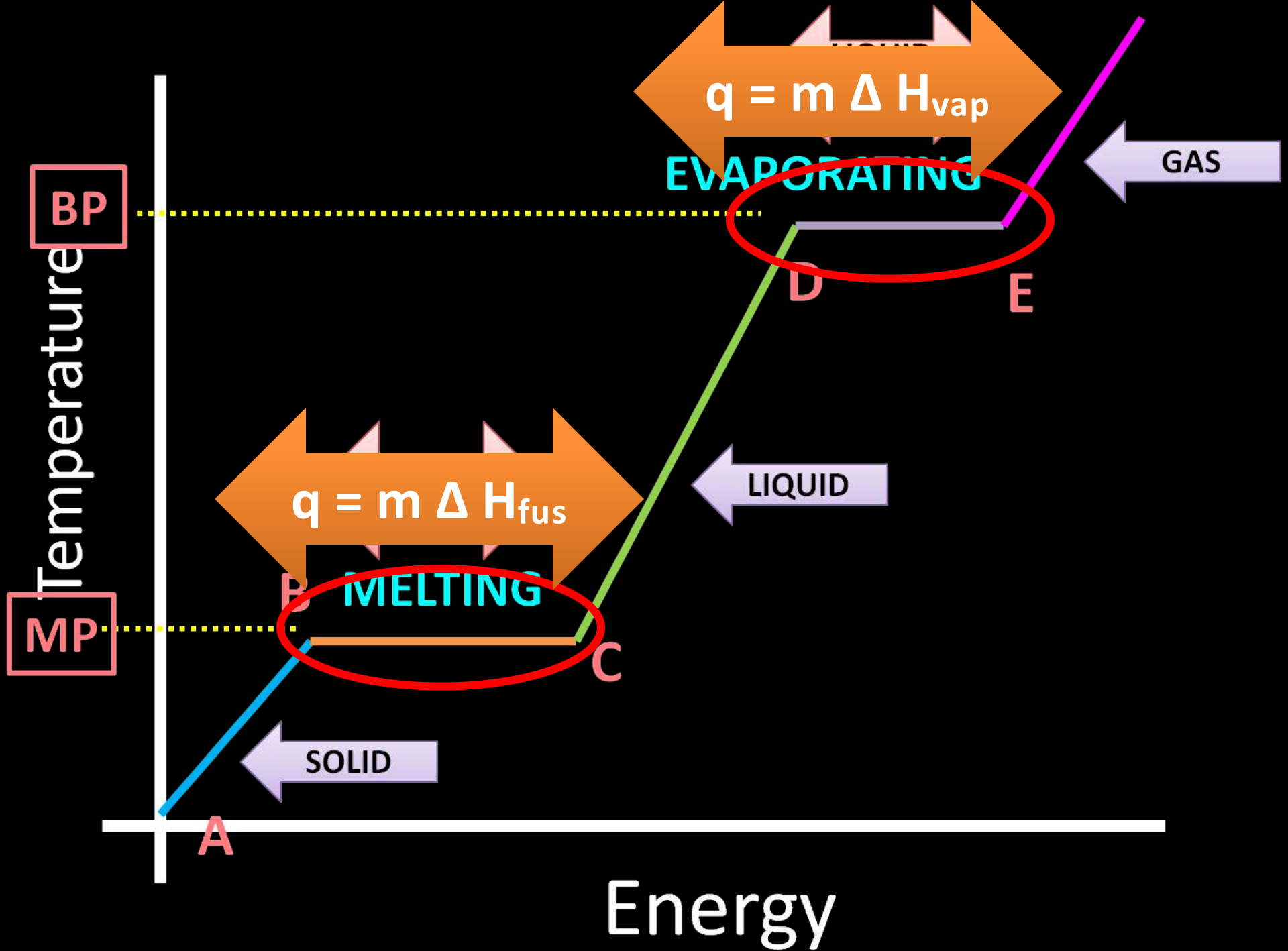




Triple point – the point on a phase diagram at which the three states of matter: gas, liquid, and solid coexist

Critical point – the point on a phase diagram at which the substance is indistinguishable between liquid and gaseous states





How many joules does it take to melt 100. grams of water at its melting point?

$$q = m \Delta H_{\text{fus}}$$

$$= 100. \text{ grams} \times 334 \text{ J/gram}$$

$$= 33400 \text{ Joules}$$

Constants

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How many joules does it take to condense 50.0 grams of water at its boiling point?

$$\begin{aligned}q &= m \Delta H_{\text{vap}} \\ &= 50.0 \text{ grams} \times 2260 \text{ J/gram} \\ &= 113\,000 \text{ Joules}\end{aligned}$$

Constants

Volume of Ideal Gas at STP: $22.4 \frac{\text{L}}{\text{mol}}$

Speed of Light in a Vacuum: $c = 3.00 \times 10^8 \frac{\text{m}}{\text{s}}$

Specific Heat of Water: $C_p(\text{H}_2\text{O}) = 1.00 \frac{\text{cal}}{(\text{g } ^\circ\text{C})} = 4.18 \frac{\text{J}}{(\text{g } ^\circ\text{C})}$

Latent Heat of Fusion of Water: $\Delta H_{\text{fus}}(\text{H}_2\text{O}) = 80 \frac{\text{cal}}{\text{g}} = 334 \frac{\text{J}}{\text{g}}$

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