

1) 31 200 J

7) 0.0456 J/g °C

2) -31 700 J

8) 424 g

3) 120°C

9) 2.60 J/g °C

4) 28°C

10) 6.21 J

5) 1100 J

11) 42.6 L

6) 14 900 J

Homework - Worksheet

$$\begin{aligned} m &= 50.0\text{g} & q &= mC\Delta T \\ T_i &= 140^\circ\text{C} & q &= mC(T_f - T_i) \\ q &= -2500\text{J} & -2500\text{J} &= (50.0\text{g})(2.01\frac{\text{J}}{\text{g}\cdot^\circ\text{C}})(T_f - 140^\circ\text{C}) \\ C &= 2.01\frac{\text{J}}{\text{g}\cdot^\circ\text{C}} & \frac{-2500\text{J}}{(50.0\text{g})(2.01\frac{\text{J}}{\text{g}\cdot^\circ\text{C}})} &= T_f - 140^\circ\text{C} \\ T_f &= ? & -24.876^\circ\text{C} &= T_f - 140^\circ\text{C} \\ & & T_f &= -24.876^\circ\text{C} + 140^\circ\text{C} \\ & & \boxed{T_f = 120^\circ\text{C}} & \end{aligned}$$

$$\begin{aligned} \textcircled{\text{II}} \quad T_i &= 75.0^\circ\text{C} & q &= vC\Delta T \\ T_f &= 10.0^\circ\text{C} & -3.32\text{kJ} &= v(0.0012\frac{\text{kJ}}{\text{L}\cdot^\circ\text{C}})(-65.0^\circ\text{C}) \\ q &= -3.32\text{kJ} & v &= \frac{-3.32\text{kJ}}{(0.0012\frac{\text{kJ}}{\text{L}\cdot^\circ\text{C}})(-65.0^\circ\text{C})} \\ C &= 0.0012\frac{\text{kJ}}{\text{L}\cdot^\circ\text{C}} & & \\ v &= ? & \boxed{v = 42.6\text{L}} & \end{aligned}$$

PHASE CHANGE AND ENTHALPY

Classifying types of systems:

1. Open system - a system where both matter and energy can flow into or out of the system.
2. Closed system - a system where energy is allowed to be transferred into and out but matter cannot be transferred.
3. Isolated system - a system where neither matter nor energy is allowed to enter or leave the system.

ENTHALPY (H) - The total internal (potential) energy and kinetic energy of a system under constant pressure.

⇒ Enthalpy is usually expressed in kJ.

ENTHALPY CHANGE (ΔH) - A change under constant pressure where the surroundings of a system absorb energy or release it to the system.

PHASE CHANGE - is a change in the state of matter without a change in the chemical composition of the system.

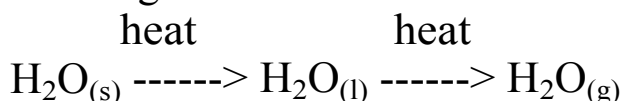
Ex. $\text{H}_2\text{O}_{(l)} \longrightarrow \text{H}_2\text{O}_{(g)}$

⇒ **always involve a change in energy but never involve a change in temperature.**

Question:

- (i) What is the temperature where water just starts boiling?
- (ii) What is the temperature when water is boiling violently?
- (iii) If energy is still going into the water and the temperature is not increasing, where is the energy going?

Consider melting ice to water and then boiling water to steam:



MOLAR ENTHALPY

For any system:

- an exothermic change involves a decrease in enthalpy

⇒ gives off energy to the surroundings

⇒ ΔH is negative.

- *solidification, condensation*

- an endothermic change involves an increase in enthalpy.

⇒ takes in energy from the surroundings

⇒ ΔH is positive.

- *fusion, vaporization*

The enthalpies for substances undergoing phase changes have been measured experimentally. (TABLE 17.3 p. 522)

- enthalpies are reported as molar enthalpies and are expressed as kJ/mol.

Solid

liquid

gas

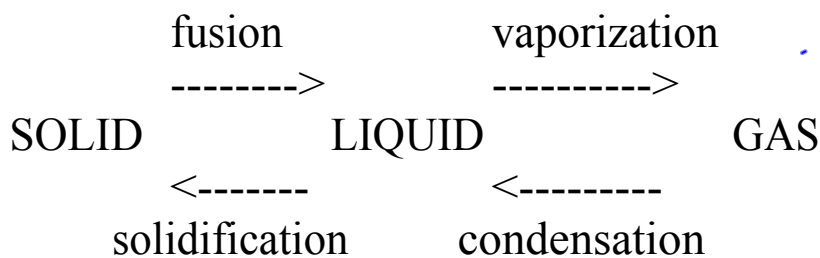
Endothermic Phase Changes

- the molar enthalpy of fusion (H_{fus}) represents the quantity of heat that the substance absorbs per mole as it changes state from **solid to liquid**.

- the molar enthalpy of vaporization (H_{vap}) represents the quantity of heat that the substance absorbs per mole as it changes state from **liquid to gas**.

Exothermic Phase Changes

- the molar enthalpy of condensation (H_{cond}) represents the quantity of heat that the substance releases per mole as it changes state from **gas to liquid**
- the molar enthalpy of solidification (H_{solid}) represents the quantity of heat that the substance releases per mole as it changes state from **liquid to solid**.



$$\Delta H_{\text{fus}} = - \Delta H_{\text{solid}}$$

$$\Delta H_{\text{vap}} = - \Delta H_{\text{cond}}$$

	H_{fus} (kJ/mol)	H_{vap} (kJ/mol)
H_2O	6.01	40.7
	H_{solid}	$H_{cond.}$
	-6.01	-40.7

Example

If 500. g of $\text{CCl}_2\text{F}_2(l)$ is vaporized at SATP, find the enthalpy change of the system ($H_{\text{vap}} = 34.99 \text{ kJ/mol}$).

$$m = 500. \text{ g}$$



$$H_{\text{vap}} = 34.99 \frac{\text{kJ}}{\text{mol}}$$

$$\Delta H_{\text{vap}} = ?$$

$$\Delta H_{\text{vap}} = nH_{\text{vap}}$$

$$\Delta H_{\text{vap}} = \left(\frac{500 \text{ g}}{120.91 \text{ g/mol}} \right) \left(34.99 \frac{\text{kJ}}{\text{mol}} \right)$$

$$\Delta H_{\text{vap}} = 145 \text{ kJ}$$

Worksheet