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Outline

- 1. Heat and Internal Energy
- 2. Specific Heat and Calorimetry
- 3. Latent Heat
- 4. Work and Heat in Thermodynamics
- 5. The First Law of Thermodynamics
- 6. Some Applications of The 1st Law
- 7. Thermal Energy Transfer

1. HEAT AND INTERNAL ENERGY

Internal energy: It is the total energy that includes the translational energy, the rotational energy, and the vibrational energy of all atoms belonging to the system.

Heat: It is one kind of energy transfer process. It stands for energy transfer from high temperature substances to low temperature substances.

Heat is either a transfer of energy due to an entropy change. In a microscopic viewpoint, it originates from an increase or a decrease of the number of microstates.

Units of heat: A calorie is the heat energy to raise the temperature of 1-g water from 14.5 to 15.5°C.

Mechanical equivalent of heat – Joule's experiment: It is experimentally confirmed that the mechanical energy of 4.18 J is equivalent to 1 calorie of heat energy. It is known as the mechanical equivalence of heat.

1 cal = 4.186 J, 1 Cal = 1000 cal

1. HEAT AND INTERNAL ENERGY

Example: A student eats a dinner containing 2000 Calories of energy. He wishes to do an equivalent amount of work in the gym by lifting a 50-kg object. How many times must he raise the object to consume the earned energy? Assume that he raise it for a distance of 2 m each time.

 $E = 2000 \ Cal = 2000 \times 1000 \times 4.186 = 8.37 \times 10^6 \ J$

 $E_{lift} = 50 \times 9.8 \times 2 = 980 J$

$$N = \frac{E}{E_{lift}} = \frac{8.37 \times 10^6}{980} = 8540$$



https://giphy.com/gifs/TTPi3fB9F5Aqs

2. SPECIFIC HEAT AND CALORIMETRY

Specific heat: A quantity of energy ΔQ is transferred to a mass *m* of a substance and changing its temperature by ΔT .

The heat capacity $c: c = \Delta Q / \Delta T$

The specific heat per unit mass $C_m = c/m = \Delta Q/m\Delta T$

The specific heat: J kg⁻¹ K⁻¹

Fe	Сυ	Ag	AI	Steel	Wood	Water	Air
470	390	230	900	500	1800	4186	1000

Calorimetry: measurement of the specific heat of unknown substances

place the hot object of temperature T_h and mass m_x into a vessel containing cold water of temperature T_c , mass m_w and measure the equilibrium temperature T_{eq} .

$$m_{w}C_{m,w}(T_{eq} - T_{c}) = m_{x}C_{m,x}(T_{h} - T_{eq}) \qquad C_{m,x} = \frac{m_{w}C_{m,w}(T_{eq} - T_{c})}{m_{x}(T_{h} - T_{eq})}$$

2. SPECIFIC HEAT AND CALORIMETRY

Example: The temperature of a 0.0500-kg ingot of metal is raised to 200°C and the ingot is then dropped into a light, insulated beaker containing 0.400 kg of water initially at 20.0°C. If the final equilibrium temperature of the mixed system is 22.4°C, find the specific heat of the metal.

$$m_x = 0.05, m_w = 0.4, T_h = 200^{\circ}C, T_c = 20^{\circ}C, T_{eq} = 22.4^{\circ}C$$

$$C_{m,w} = 4186 \frac{J}{kg K}$$

$$C_{m,x} = \frac{m_w c_w (T_{eq} - T_c)}{m_x (T_h - T_{eq})} = \frac{(0.4)(4186)(22.4 - 20)}{(0.05)(200 - 22.4)} = 453 \frac{J}{kg K}$$

3. LATENT HEAT

Latent heat: It is the heat energy required for the phase transition from a solid to a liquid phase or from a liquid to a gas phase for substances.

Latent heat: $L = \Delta Q/m$

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Latent heat of fusion: L_f
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Latent heat of vaporization: L_v



Energy Input

Substance	Melting point	L_f	Boiling point	<i>L_v</i> (J / kg)
Не	0.95 K		4.22 K	2.09X10 ⁴
H ₂	14.15 K		20.15 K	
N ₂	63.15 K		77.35 K	2.01X10 ⁵
water	0°C	3.33X10 ⁵	100°C	2.26X10 ⁶
Pb	327.3°C		1750°C	
Al	660°C		2450°C	

4. WORK AND HEAT IN THERMODYNAMICS

If the gas is compressed quasi-statically (slowly enough) to be remained in thermal equilibrium at all times, the work done on the gas is

 $dW = \vec{F} \cdot d\vec{r} = -Fdy = -PAdy = -PdV$

dy is in the direction of volume expansion F is in the volume compression direction

$$\Delta W = W = -\int_{V_i}^{V_f} P dV$$



Consider a compression of the volume of gas, its work is positive. A positive work means an increase of energy in the gas system.

4. WORK AND HEAT IN THERMODYNAMICS

Example: An ideal gas is taken through two processes in which $P_f = 1.0 \times 10^5 \text{ Pa}$, $V_f = 2.0 \text{ m}^3$, $P_i = 0.20 \times 10^5 \text{ Pa}$, and $V_i = 10 \text{ m}^3$. For process 1, the temperature remains constant. For process 2, the pressure remains constant and then the volume remains constant. What is the ratio of the work W_1 done on the gas in the first process to the work W_2 done in the second process?

$$P_{i}V_{i} = P_{f}V_{f} = 2.0 \times 10^{5} J$$
Process 1: $\Delta W = -\int_{V_{i}}^{V_{f}} P dV = -\int_{10}^{2} \frac{2.0 \times 10^{5}}{V} dV$

$$P_{i}$$

$$P_{i}$$

$$P_{i}$$

$$V_{f}$$

4. WORK AND HEAT IN THERMODYNAMICS

Isothermal expansion: A gas at temperature T expands slowly while absorbing energy from a reservoir to maintain the constant temperature.

Free expansion: A gas expands rapidly into an evacuated region after a membrane is broken.

$$PV = nRT = const \rightarrow P = \frac{nRT}{V}$$



The two kinds of expansions give the same results of a decrease of work energy and an increase of heat energy.

Note that here the absorption of heat energy originates from an increase of microstates due to the volume expansion.

$$\Delta W = W = -\int_{V_i}^{V_f} \frac{nRT}{V} dV = nRT \ln\left(\frac{V_i}{V_f}\right), \Delta E = 0 \to \Delta Q = -\Delta W$$

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5. THE FIRST LAW OF THERMODYNAMICS

The first law of thermodynamics: It is a special case of the law of conservation of energy.

 $\Delta E_{int} = \Delta Q + \Delta W$

The change of internal energy is independent of the path.

A cyclic process which starts and ends at the same state gives no changes of its internal energy.

 $\Delta E_{int} = 0 \rightarrow \Delta Q + \Delta W = 0 \rightarrow \Delta Q = -\Delta W$

In the cyclic process, the net work done on the system per cycle equals the area enclosed by the path representing the process on the P-V diagram

6. SOME APPLICATIONS OF THE 1ST LAW

Equations for an ideal gas:

 $PV = Nk_BT = nRT$ $\Delta W = -P\Delta V = -\int PdV$ $\Delta E_{int} = \Delta Q + \Delta W$

In Chapter 20, we will learn that the total energy is $E = 3Nk_BT/2$.

Adiabatic process: $\Delta Q = 0 \rightarrow \Delta E_{int} = \Delta W$

Isothermal process: $\Delta E_{int} = 0$, $PV = const \rightarrow \Delta Q = -\Delta W$

Isovolumetric process: $\Delta V = 0 \rightarrow \Delta W = 0 \rightarrow \Delta E_{int} = \Delta Q$

Isobaric process: $\Delta P = 0 \rightarrow \Delta W = -P\Delta V, \Delta E_{int} = \Delta Q + \Delta W$

The three typical types of energy transfer are conduction, convection, and radiation. In all mechanisms of heat transfer, the rate of cooling of a body is approximately proportional to the temperature difference between the body and its surrounding.

$$R_{heat} = I_{heat} = \frac{\Delta Q}{\Delta t} = P \propto \Delta T = (T_{body} - T_{surrounding})$$

Thermal conduction:

$$I_{heat} = P = kA \frac{\Delta T}{L}$$

k: thermal conductivity, 1/k: thermal resistivity

$$\Delta T = I_{heat} \left(\frac{1}{k}\right) \frac{L}{A}$$
 compare with $V = IR, R = \rho \frac{L}{A}$



Connected in series: $R = R_1 + R_2 + \cdots$ Connected in parallel: total conductance = sum of each conductance, $G = G_1 + G_2 + \cdots, \left(\frac{1}{R}\right) = \left(\frac{1}{R_1}\right) + \left(\frac{1}{R_2}\right) + \cdots$

Thermal conductivity of materials in the unit of Wm⁻¹K⁻¹:

Fe	Cu	Ag	AI	Steel	Wood	Water	Air
80	401	429	210	46	0.13	0.58	0.026

Note that the mass specific heat for Fe and Water are 470 and 4186 J kg⁻¹K⁻¹.

Thermal radiation: Stephan-Boltzmann law: $P = \sigma A e T^4$, where $\sigma = 5.67 \times 10^{-8} \frac{W}{m^2 K^4}$ is the universal constant (Stephan's constant) and *e* is the emissivity of the object. The value of the emissivity is between 0 and 1.

Newton's coffee cooling:

 $T = T_0 + \Delta T$

 $\Delta P = \sigma A e \left(T^4 - T_0^4 \right) \cong 4 \sigma A e T_0^3 \Delta T$



Example: Two slabs of the same area A, thickness L_1 and L_2 and thermal conductivities k_1 and k_2 are in thermal contact with each other. The temperature of their outer surfaces are T_c and T_h , respectively, and $T_h > T_c$. Determine the temperature at the interface and the rate of energy transfer by conduction through the slabs in the steady-state condition.

-T)

 T_h

 T_c

assume the temperature *T* at the interface between the two slabs

$$\begin{split} P_{1} &= \frac{k_{1}A(T-T_{c})}{L_{1}} = P_{2} = \frac{k_{2}A(T_{h}-T)}{L_{2}} \rightarrow k_{1}L_{2}(T-T_{c}) = k_{2}L_{1}(T_{h}-T_{c}) \\ T &= \frac{k_{1}L_{2}T_{c} + k_{2}L_{1}T_{h}}{k_{1}L_{2} + k_{2}L_{1}}, P_{1} = P_{2} = \frac{A(T_{h}-T_{c})}{\frac{L_{1}}{k_{1}} + \frac{L_{2}}{k_{2}}} \\ I_{heat} &= \frac{\Delta T}{R_{heat}}, R_{heat} = \frac{L}{kA} \qquad R_{t} = R_{1} + R_{2} = \frac{L_{1}}{k_{1}A} + \frac{L_{2}}{k_{2}A} \\ I_{heat} &= P_{1} = P_{2} = \frac{T_{h} - T_{c}}{\frac{L_{1}}{k_{1}A} + \frac{L_{2}}{k_{2}A}} \end{split}$$

Example: The inside (radius a) of a hollow cylinder is maintained at a temperature T_a while the outside (radius b) is a lower temperature, T_b . The cylinder has a length of L and thermal conductivity k. Ignoring end effects, please calculate the rate of energy transfer from the inner to the outer surface.

confirm that the thermal resistances are connected in series

$$\Delta T = I_{heat} \left(\frac{1}{k}\right) \frac{L}{A}$$

$$R_{heat} = \frac{L}{kA}$$

$$dR = \frac{dr}{k(2\pi rL)} \rightarrow R = \int_{a}^{b} \frac{dr}{k(2\pi rL)} = \frac{\ln(b/a)}{2\pi kL}$$

$$P = I_{heat} = \frac{\Delta T}{R} = \frac{2\pi k L (T_a - T_b)}{\ln(b/a)}$$



Example: Estimate the order of magnitude of the temperature of the filament of a 100 W lightbulb when it is operating. To model the filament as a cylinder 10.0 cm long with a radius of 0.0500 mm. The emissivity of the filament is 1.

 $P = \sigma A e T^4$

$$\sigma = 5.67 \times 10^{-8} \frac{W}{m^2 K^4}, e = 1$$

$$A = 2\pi rL = 2\pi (5.0 \times 10^{-5})(1.0 \times 10^{-1}) = 3.14 \times 10^{-5} m^2$$

$$T = \left(\frac{P}{\sigma A e}\right)^{1/4} = 2740 \ K$$

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