

Thermodynamics I Chapter 3 Energy, Heat and Work

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Energy, Heat and Work (Motivation)

- A system changes due to interaction with its surroundings.
- Energy interaction is a major factor.
- This chapter studies the nature of energy and its various forms and transfers so that we are able to follow its interaction with a system.



ENERGY & THE 1st LAW OF THERMO.

1st Law : concerning **quantity of energy**

Energy is **conserved**

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(<u>Amount</u> of energy is constant, but can <u>change forms</u>)

(e.g. potential, kinetic, electrical, chemical, etc.)

$$E = Total system energy[J,$$
$$e = \frac{E}{m} \left[\frac{kJ}{kg} \right] (specific energ)$$





<u>Macroscopic</u> – system energy which value depends on a *reference point* (<u>Kinetic</u> <u>Energy (KE), Potential Energy</u> (PE) <u>Microscopic</u> – energy due to *molecular interactions* & *activity* (independent of any reference point) (<u>Internal Energy</u>, U)

Total System Energy

$$E = KE + PE + U [kJ]$$
$$e = ke + pe + u [kJ/kg]$$



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Kinetic Energy (KE)



Potential Energy (PE)

PE = mgz
$$\Delta$$
PE = mg($z_2 - z_1$)
pe = $\frac{PE}{m}$ = gz Δ pe = $\frac{\Delta PE}{m}$ = g($z_2 - z_1$)



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Internal Energy, U

- Molecular movement (vibration, collision, etc) <u>sensible energy</u> (molecular activity temperature)
 - Bond Energy between molecules (phase change)
 - *latent energy* (constant temperature)
 - Bond Energy between atoms in a molecule chemical energy
 - Bond Energy between protons & neutrons in the nucleus
 - <u>nuclear energy</u>





Modes of Energy Transfer

Energy Interaction between System and Surrounding

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Energy can cross the boundary (transferred) of a closed system by **2 methods**;

HEAT & WORK



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Heat - Energy that is *being transferred* due to a <u>temperature difference</u>

- •Heat is a mode of energy transfer
- •Heat is not a property
- •Energy is related to <u>states</u> (property)
- •Heat is related to <u>processes</u> (not a property, depends on the path)

$$q = \frac{Q}{m}$$
$$\dot{Q} = \frac{Q}{t} \left[\frac{kJ}{s} = kW \right]$$
$$\int_{1}^{2} \delta Q = Q_{12}$$

- = rate of heat transfer
- Amount of heat transferred during a process (depends on the path)





Adiabatic ≠ Isothermal! (T can change by other methods; energy can enter system by <u>work</u>)





WORK, W [J, kJ]

<u>Work</u> -- Energy that is *crossing the boundary* <u>other than heat</u> (electrical, stirrer, shaft, moving piston, etc.)

-Not a property (related to *process*) -Mode of energy *transfer*

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Types of Work







Boundary Work for Polytropic Processes

Boundary Work

$$W_B = \int_{V_1}^{V_2} P dV$$

Area under P-v graph

General Polytropic Work

$$W = \frac{P_2 V_2 - P_1 V_1}{1 - n} \qquad (n \neq 1)$$

Const. Pressure Work (n=0)

$$W = p(V_2 - V_1)$$

Isothermal Work (n=1) *ideal gas*

$$W = mRTln\left(\frac{V_2}{V_1}\right) = mRTln\left(\frac{P_1}{P_2}\right)$$

Const. Volume Work (dv=0)

$$W_{b-constvol} = 0$$







1st Law Closed System





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- Recall that System Energy consists of: $E_{system} = U + KE + PE$

or

$$\Delta E_{\text{system}} = \Delta U + \Delta KE + \Delta PE$$

thus;



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1st Law for Closed System in different forms;

$$Q - W = \Delta U + \Delta KE + \Delta PE \qquad [kJ]$$

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per unit mass;

 $q - w = \Delta u + \Delta ke + \Delta pe$ [kJ/kg]

Rate form;

$$\dot{Q} - \dot{W} = \frac{dU}{dt} + \frac{d(KE)}{dt} + \frac{d(PE)}{dt}$$
[kW]

<u>Cyclic Process</u> $\Delta E = 0$

$$Q - W = 0$$
$$Q_{cycle} = W_{cycle}$$
$$\sum Q = \sum W$$





summary;



etc...



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