

# Closed Systems in Thermodynamics

## Introduction

A **closed system** in thermodynamics is a system where *mass is fixed* (no matter enters or leaves), but *energy can cross the system boundary* in the form of heat or work. This is the foundation for analyzing energy conservation, processes, and cycles in engineering.

## Key Concepts

- The First Law
- Work
- Heat
- Energies
- Efficiency

## Definitions

- **Closed System:** A system where mass remains constant, but energy may cross the boundary.
- **Work ( $W$ ):** Energy transfer associated with a force acting through a distance. It is *organized* energy transfer.
- **Heat ( $Q$ ):** Energy transfer due to temperature difference. It is *disorganized* energy transfer.
- **Internal Energy ( $U$ ):** Microscopic energy associated with molecular motion and interactions.

## The First Law of Thermodynamics

For a closed system, the energy balance is:

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

where:

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

- $Q$ : heat transferred into system (positive if into system).
- $W$ : work done by system (positive if system does work).
- $\Delta U$ : change in internal energy.
- $\Delta KE$ : change in kinetic energy.
- $\Delta PE$ : change in potential energy.

Usually, we will neglect Kinetic and Potential energies. However, there are some cases in which we won't (Like a pump raising liquid, or calculating velocities through tubes).

After neglecting those energies, we get the classic equation:

$$\Delta U = Q + W$$

## Work

The work of a process will always be defined as:

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

Except there are different cases:

- **Irreversible Work (Constant  $P_{ext}$ ):**

$$W = -P_{ext}\Delta V$$

- **Isothermal Ideal Gas:**

$$W = nRT \ln \left( \frac{V_2}{V_1} \right)$$

- **Polytropic Process ( $PV^n = C$ ):**

$$W = \frac{P_2V_2 - P_1V_1}{1 - n}, \quad n \neq 1$$

- **Adiabatic Ideal gas:**

$$W = \Delta U = nC_v\Delta T = \frac{P_2V_2 - P_1V_1}{1 - \frac{C_p}{C_v}}$$

## Heat Transfer

Heat transfer ( $Q$ ) can be a lot more fluid in how it's calculated. Usually we try to limit  $Q$ , and so we can assume that the system is insulated ( $Q = 0$ ) Or we can say that our given process happens so quick that  $Q = 0$ .

- Adiabatic (Insulated):  $Q = 0$
- Isothermal:  $\Delta U = 0$  So  $Q = -W$
- Constant Pressure (Open to atmosphere):  $Q = \Delta H = nC_p\Delta T$
- Constant Volume (Rigid Container):  $W = 0$  so  $Q = \Delta U = nC_v\Delta T$

## Energies

### (i) Internal Energy

- A collection of macroscopic energies (Potential and Kinetic), at the microscopic level.
- Typically we use this energy most.
- Directly related to  $C_v$

### (ii) Kinetic Energy

- We will mostly neglect this.
- We can use this for nozzles and turbines, anything with high speeds where it can't be neglected.
- $KE = \frac{1}{2}mv^2$

### (iii) Potential Energy

- Also usually neglected.
- We can use this for pumps and different systems with water
- $PE = mgh$

### (iv) Enthalpy

- Enthalpy is the energy needed for a given system to occupy that space.
- Don't really care in closed systems, our matter isn't changing.

## Efficiency

No real world system is reversible, as it would take infinite time. Even getting close to being fully reversible is usually a waste of our time. The thing about reversible systems is that they maximize work, which is what we want. So, usually in irreversible systems, we will have a net energy loss. We like to quantify that with efficiency:

$$\eta = \frac{W_{actual}}{W_{reversible}}$$

Or, if we do it with  $Q$ :

$$\eta = \frac{Q_{actual}}{Q_{reversible}}$$

## Example Problem

Consider a piston–cylinder assembly that contains 2.5 L of an ideal gas at 30°C and 8 bar. The gas reversibly expands to 5 bar.

1. Write an energy balance for this process (you may neglect changes in potential and kinetic energy).
  - (i) The First Law of Thermodynamics states that energy balances should take the form:  $\Delta U = W + Q$   
Where  $\Delta U$  is the change in internal energy,  $W$  is the Work done by the system, and  $Q$  is the heat added to the system.
2. Suppose that the process is done isothermally. What is the change in internal energy,  $\Delta U$ , for the process? What is the work done,  $W$ , during the process? What is the heat transferred,  $Q$ ?
  - (i) We know that the process is isothermal, meaning  $\Delta T = 0$  and the process will stay at 30°C and we know that  $\Delta U = 0$  for isothermal processes.
  - (ii) For reversible processes, we know that:

$$W = - \int_{V_1}^{V_2} P_{ext} dV \quad (1)$$

With the Ideal Gas assumption we can turn this into:

$$W = -nRT \int_{V_1}^{V_2} \frac{dV}{V} \quad (2)$$

We are given  $V_1$ ,  $T$ , and  $P_1$  to which we can find  $n$ :

$$n = \frac{PV}{RT} = 0.79 \text{ mol} \quad (3)$$

Then we will solve for  $V_2$ :

$$V_2 = \frac{nRT}{P} = 4.9 \text{ L} \quad (4)$$

Now we are finally able to solve for  $W$ :

$$W = nRT \ln\left(\frac{V_1}{V_2}\right) = -13.39 \text{ bar} \cdot \text{L} = -1339.24 \text{ J} \quad (5)$$

- (iii) For isothermal processes, we know that  $W = -Q$ , so we know  $Q = 1339.24$

## Recap

Closed systems are mass-conserving but allow energy transfer as heat or work. The first law provides a universal energy balance. Understanding the different forms of work and heat is essential for applying thermodynamics to reactors, engines, and industrial processes.