

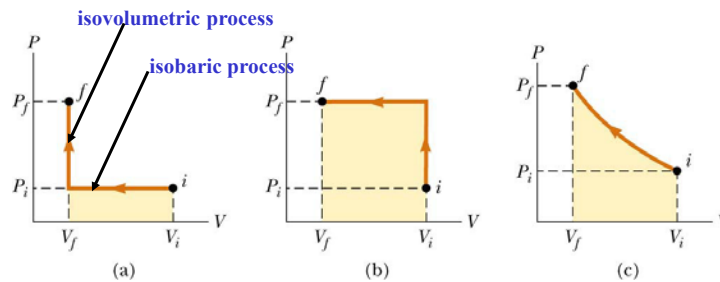
Chap. 15, Thermodynamics

Work in Thermodynamic Processes

Mechanical work done on a system is

$$W = -P \Delta V$$

For a gas, the work done can be determined from a PV diagram.



First Law of Thermodynamics

First Law of Thermodynamics

The internal energy of a system changes from an initial value U_i to a final value of U_f due to heat Q and work W :

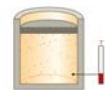
$$\Delta U = U_f - U_i = Q + W$$

The internal energy depends only on the state of a system, not on the method (path) by which the system arrives at a given state.

For an isolated system, for a cyclic process, or for an isothermal (constant temperature) process,

$$\begin{aligned} \Delta U &= 0 \\ W &= -Q \end{aligned}$$

Work
Isothermal



Ideal Gas

monatomic molecules $U = \frac{3}{2} nRT$

$$\Delta U = \frac{3}{2} nR \Delta T$$

molar specific heat at constant volume $C_v \equiv \frac{3}{2} R$

$$\Delta U = nC_v \Delta T$$

Ideal Gas At Constant Pressure

isobaric

$$Q = \Delta U - W = \Delta U + P\Delta V$$

$$\Delta U = \frac{3}{2} nR\Delta T$$

$$P\Delta V = nR\Delta T$$

$$Q = \frac{5}{2} nR\Delta T$$

molar specific heat at constant volume

$$Q = nC_p \Delta T$$

$$C_v \equiv \frac{3}{2} R$$

$$C_p = C_v + R$$

$$\gamma = \frac{C_p}{C_v}$$

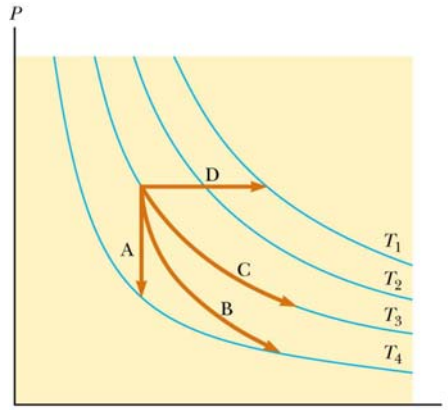
Adiabatic Process

An adiabatic process is one which involves no heat flow with the outside environment.

$$Q = 0$$

$$\Delta U = W$$

$$PV^\gamma = \text{const.}$$



Adiabatic Process

Step #1: Isobaric $P, V, T (= PV / nR) \Rightarrow P, V + \Delta V, T + \Delta T_1$

$$\Delta U_1 = nC_p \Delta T_1 - P\Delta V \quad nC_v \Delta T_1 = nC_p \Delta T_1 - P\Delta V$$

$$P\Delta V = n(\gamma - 1)C_v \Delta T_1$$

Step 2: Lose the same Heat at Constant Volume $P, V + \Delta V, T + \Delta T_1 \Rightarrow P + \Delta P, V + \Delta V, T + \Delta T_1 + \Delta T_2$

$$\Delta U_2 = nC_v \Delta T_2 = -nC_p \Delta T_1 \quad \Delta T = \Delta T_1 + \Delta T_2 = (1 - \gamma)\Delta T_1$$

$$\Delta U = \Delta U_1 + \Delta U_2 = -P\Delta V$$

$$(P + \Delta P)(V + \Delta V) = nR(T + \Delta T_1 + \Delta T_2) \quad \leftarrow \text{Gas Law}$$

$$V\Delta P + P\Delta V = nR(1 - \gamma)\Delta T_1 = -\frac{R}{C_v}P\Delta V \quad \gamma P\Delta V = -V\Delta P$$

$$\frac{\gamma \Delta V}{V} = -\frac{\Delta P}{P} \quad \rightarrow \quad \boxed{PV^\gamma = \text{const.}}$$

Reversible and Irreversible Processes

A reversible process is one in which every state along some path is an equilibrium state. The system can be returned to its initial conditions along the same path.

A process that does not satisfy these requirements is irreversible.

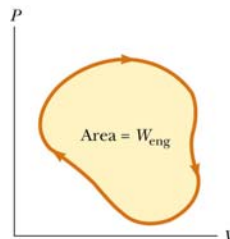
Heat Engines

A heat engine is a device that converts internal energy to other useful forms of energy.

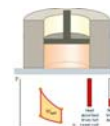
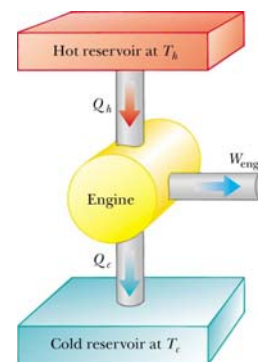
In a complete cycle

$$W = |Q_h| - |Q_c|$$

Efficiency
$$e = \frac{W}{|Q_h|} = 1 - \frac{|Q_c|}{|Q_h|}$$



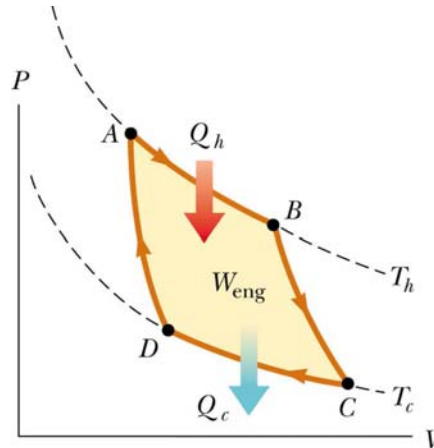
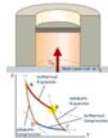
second law of thermodynamics:
it is impossible to construct a heat engine that is 100% efficient.



Carnot Engine

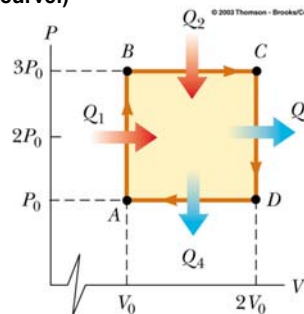
No real engine operating between two energy reservoirs can be more efficient than a Carnot engine operating between the same two reservoirs.

$$e = 1 - \frac{T_C}{T_H}$$



Example Problem

- 58.** One mole of an ideal gas is taken through the cycle shown in Figure P12.58. At point A, the pressure, volume, and temperature are P_0 , V_0 , and T_0 . In terms of R and T_0 , find
- the total energy entering the system by heat per cycle,
 - the total energy leaving the system by heat per cycle,
 - the efficiency of an engine operating in this cycle, and
 - the efficiency of an engine operating in a Carnot cycle between the temperature extremes for this process. (*Hint:* Recall that work done on the gas is the negative of the area under a PV curve.)



Entropy And Disorder

The change in entropy for reversible processes is $\Delta S = Q/T$ with T expressed in Kelvin scale. Entropy is a measure of the “disorder” of a system. It is related to the number of possible ways the total energy of a system can be subdivided into its individual components.

$$S = k_B \ln W$$

Another version of the second law of thermodynamics

The total entropy of the universe does not change when a reversible process occurs ($\Delta S_{\text{universe}} = 0$) and increases when an irreversible process occurs ($\Delta S_{\text{universe}} > 0$).

Third Law of Thermodynamics

Third Law of Thermodynamics

It is not possible to lower the temperature of any system to absolute zero in a finite number of steps.

Wise Guy's Interpretation of Thermodynamics Laws

Zeroth Law: Everyone is treated the same way. No exceptions.

First Law: You can't win.

Second Law: You must lose.

Third Law: You can't get out of the game.