

XIV: The First Law

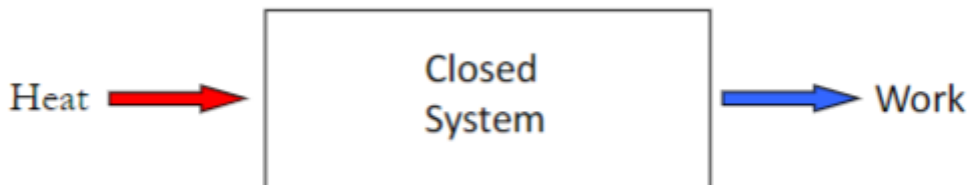
1: First Law of thermodynamics

1.1: Definition

- Usually referred as the Law of Conservation of Energy: **Energy cannot be created or destroyed, but can be transformed from one state to another.**

1.2: Balance for a closed system

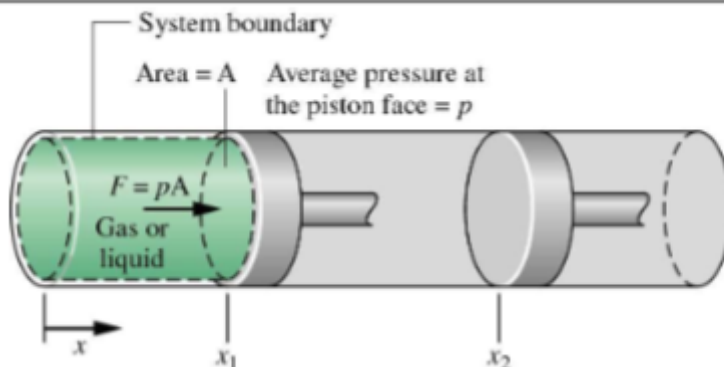
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- $\Delta E = Q - W$ or $\Delta E = Q + W$
- $\Delta E = Q - W$, which is used for sign convention of Clausius.
- $E = U + KE + PE$
- $Q - W = \Delta U$
- We use specific terms(per unit mass):
 $q - w = \Delta u$

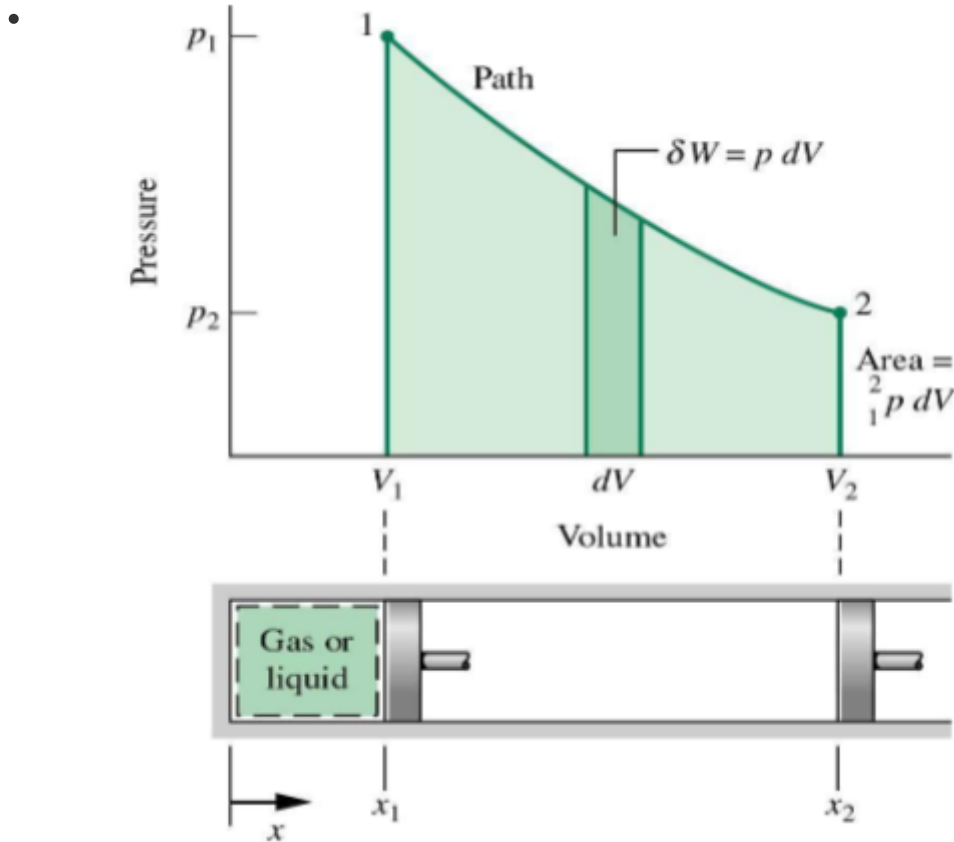
1.3: Work for a closed system

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- Total work for the sum of all the little changes.

$$\int_1^2 p dV$$



1.4: Enthalpy, H

- $H = U + PV$, where U is internal energy and PV is work of the fluid.
- If we mark the q as the heat of a reaction:

$$\Delta H = q$$
- Note it is under the constant pressure conditions.

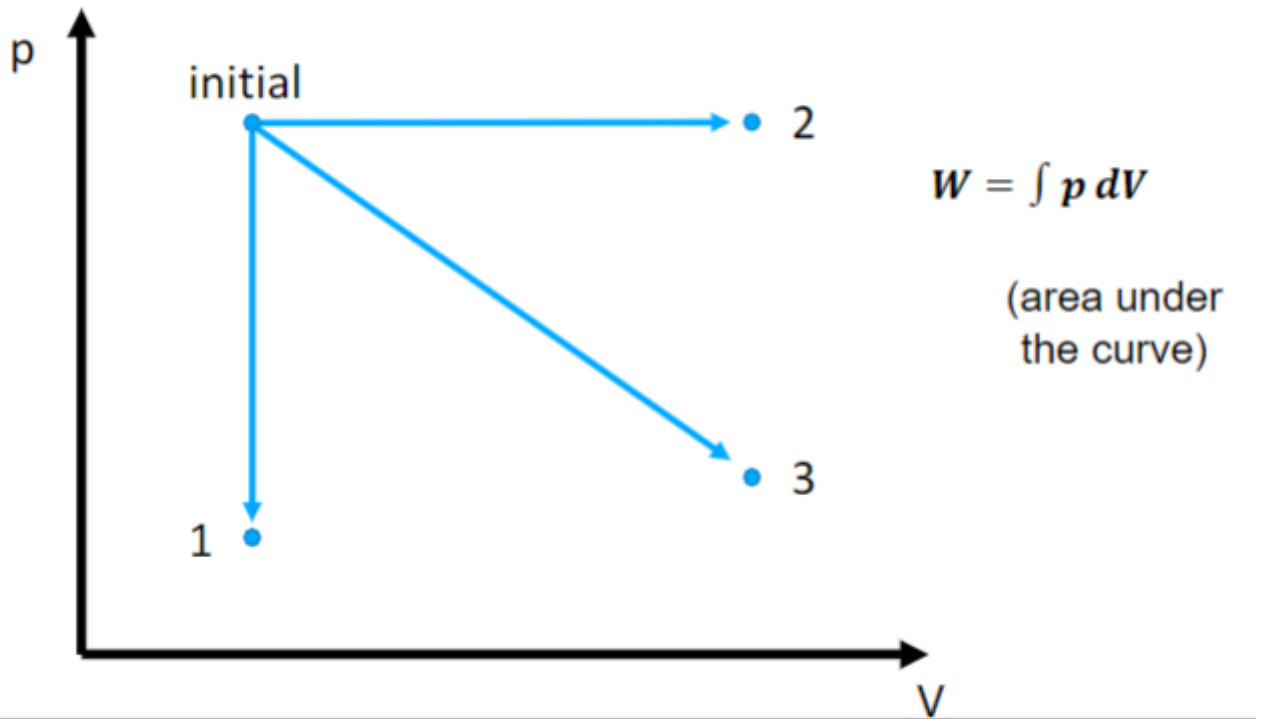
1.3: Specific heat capacity

- $Q = cm\Delta T$ or $q = c\Delta T$
- For ideal gases at constant volume: $\Delta U = c_v m\Delta T$
- Use the ideal gas law, $p\Delta V = mR\Delta T$
- Then, $Q = mc_v\Delta T + mR\Delta T$
- So we can find the specific heat capacity in constant pressure: $c_p = c_v + R = \frac{\Delta h}{\Delta T}$
- $H = U_p V$ and $h = u + pv$

2: Polytropic processes

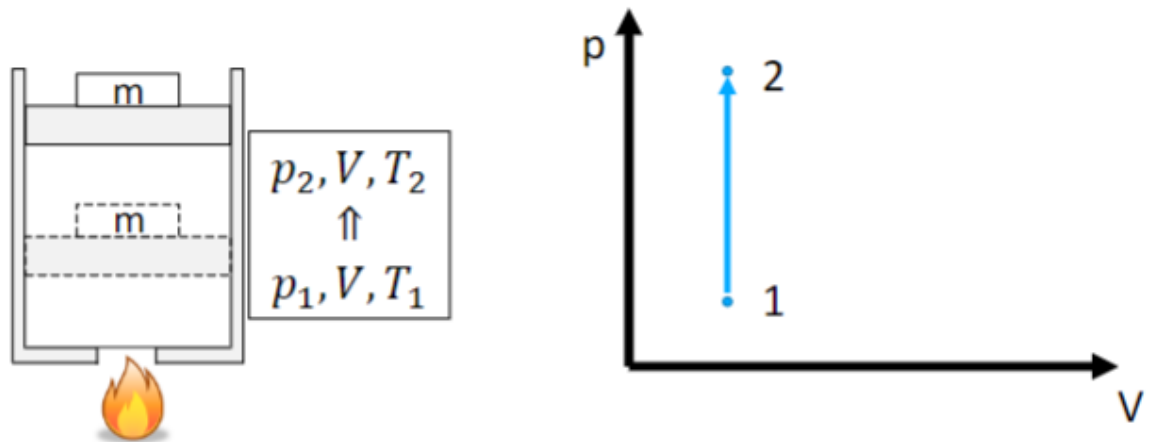
2.1: Boundary work

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2.2: Isochoric process (V=constant)

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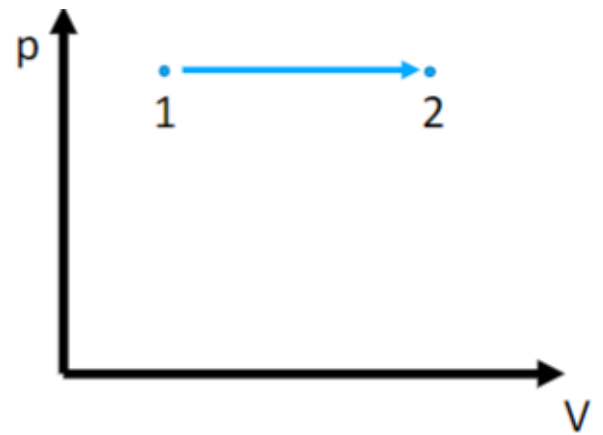
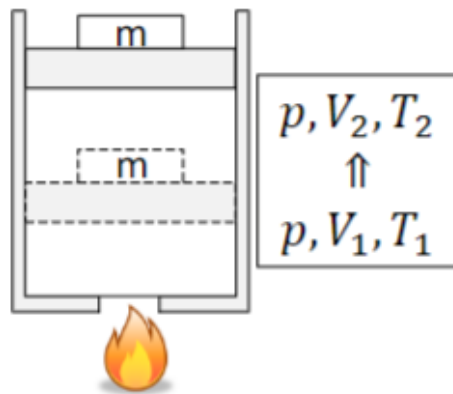
- Work done
- Heat transfer
- For gases with constant c_v

$$W_{12} = \int_{V_1}^{V_2} p \, dV = 0$$

$$Q = \Delta U = mc_v \Delta T$$

$$Q = mc_v(T_2 - T_1)$$

2.3: Isobaric process (P=constant)

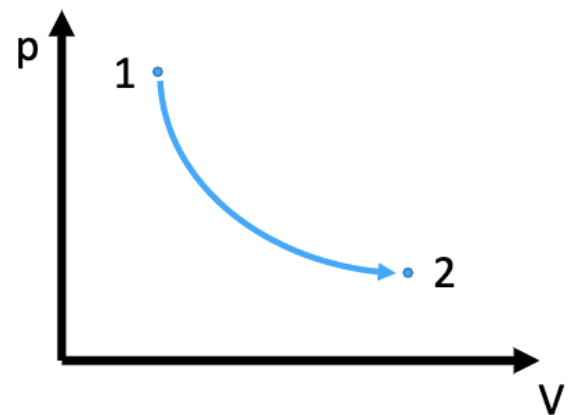
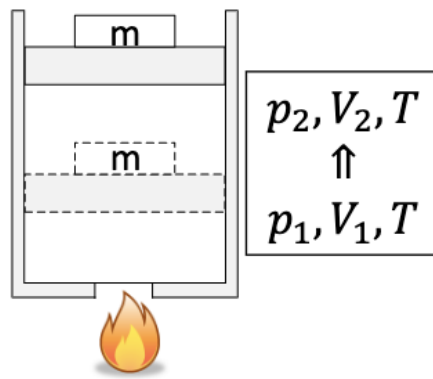


- Work done
- Change in internal energy
- Heat transfer

$$W_{12} = \int_{V_1}^{V_2} p \, dV = p(V_2 - V_1)$$

$$\Delta U = mc_v \Delta T$$

$$Q = mc_p \Delta T$$



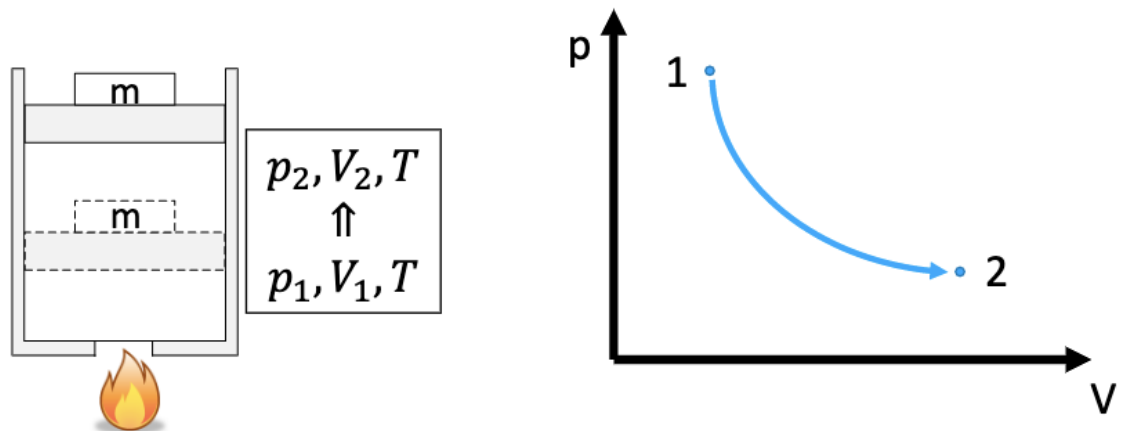
- Work done
- For an ideal gas

$$W_{12} = \int_{V_1}^{V_2} p \, dV$$

$$pV = nR_0T$$

$$W_{12} = \int_{V_1}^{V_2} \frac{nR_0T}{V} dV = nR_0T \int_{V_1}^{V_2} \frac{1}{V} dV = nR_0T \ln \frac{V_2}{V_1}$$

2.4: Isothermal process (T=constant)



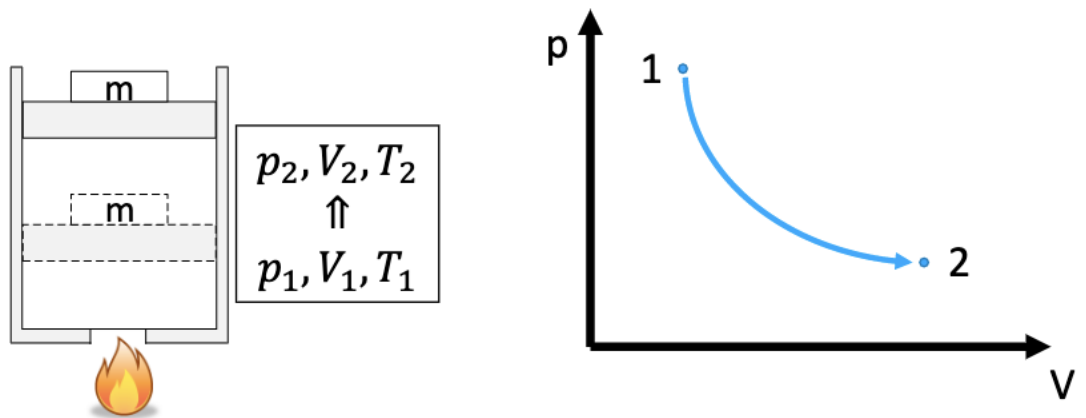
- Work done
- For an ideal gas

$$W_{12} = \int_{V_1}^{V_2} p dV$$

$$pV = nR_0T$$

$$W_{12} = \int_{V_1}^{V_2} \frac{nR_0T}{V} dV = nR_0T \int_{V_1}^{V_2} \frac{1}{V} dV = nR_0T \ln \frac{V_2}{V_1}$$

2.5: Polytropic process (multiple process)



- Many real processes follow
- Equivalent to
- Or using the ideal gas law

$$pV^n = \text{const}$$

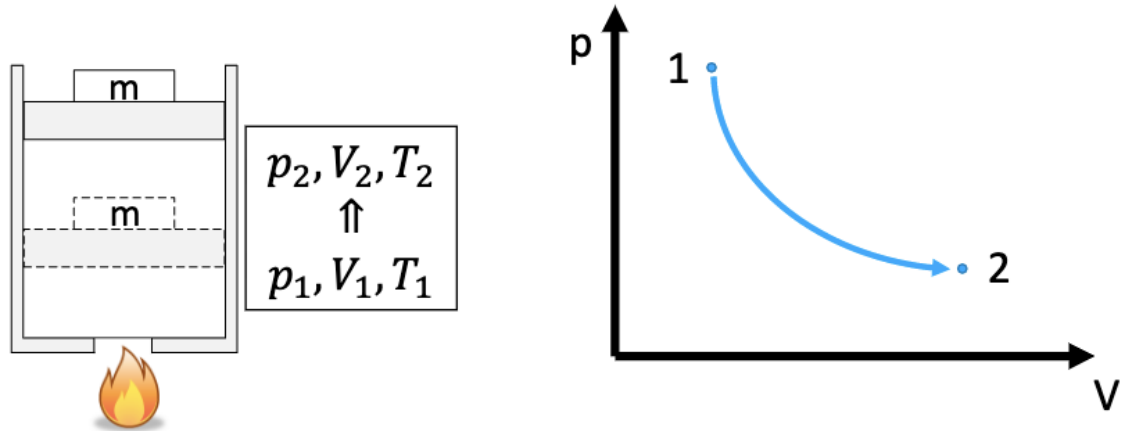
$$pv^n = \text{const}$$

$$TV^{n-1} = \text{const} \quad pT^{-n/(n-1)} = \text{const}$$

- Work done (using $pV^n = \alpha$)

$$W_{12} = \int_{V_1}^{V_2} p dV = \alpha \int_{V_1}^{V_2} V^{-n} dV = \alpha \left[\frac{V^{-n+1}}{1-n} \right]_{V_1}^{V_2} = \frac{p_2 V_2 - p_1 V_1}{1-n}$$

2.6: Isentropic process (entropy process)



- Adiabatic means no heat crosses the system boundary
- An adiabatic process that is reversible is called isentropic
- Special case of a polytropic process where

$$n = \gamma = \frac{c_p}{c_v} \quad (\text{note } \gamma = 1.4 \text{ for air})$$

- Adiabatic means

$$Q_{12} = 0$$

- From polytropic work (with $n = \gamma$)

$$W_{12} = \frac{p_2 V_2 - p_1 V_1}{1 - \gamma}$$

Value of n	Process	Description	Result of IGL
∞	<i>isochoric</i>	constant volume ($V_1 = V_2$)	$\frac{p_1}{T_1} = \frac{p_2}{T_2}$
0	<i>isobaric</i>	constant pressure ($p_1 = p_2$)	$\frac{V_1}{T_1} = \frac{V_2}{T_2}$
1	<i>isothermal</i>	constant temperature ($T_1 = T_2$)	$p_1 V_1 = p_2 V_2$
$1 < n < \gamma$	polytropic	-none-	$\frac{p_1}{p_2} = \left(\frac{V_2}{V_1}\right)^n = \left(\frac{T_1}{T_2}\right)^{\frac{n}{n-1}}$
γ	<i>isentropic</i>	constant entropy ($S_1 = S_2$) adiabatic & reversible	

2.7: Polytropic work

Process	Boundary Work
<i>isochoric</i>	$W_{12} = p(V_2 - V_1) = 0$
<i>isobaric</i>	$W_{12} = p(V_2 - V_1)$
<i>isothermal</i>	$W_{12} = RT \ln \frac{V_2}{V_1}$
polytropic	$W_{12} = \frac{p_2 V_2 - p_1 V_1}{1 - n}$
<i>isentropic</i>	