Thermodynamic Processes

• States of a thermodynamic system can be changed by interacting with its surrounding through work and heat. When this change occurs in a system, it is said that the system is undergoing a **process**.

• A thermodynamic **cycle** is a sequence of different processes that begins and ends at the same thermodynamic state.

- Some sample processes:
 - \rightarrow Isothermal process: temperature is constant T=C
 - \rightarrow Isobaric process: pressure is constant, P=C
 - Isentropic process: entropy is constant, s=C
 - \rightarrow Constant-volume process, v=C
 - \rightarrow Adiabatic process: no heat transfer, Q=0

• Use ideal gas assumption (closed system):

 \rightarrow Isothermal process: T=constant Energy balance $\Delta U=Q-W$, for ideal gas $\Delta U=\Delta H=0$ since both are functions of temperature only Q=W, W= $\int PdV = \int \frac{mRT}{V} dV = mRT \int_{1}^{2} \frac{dV}{V}$ $= mRT \ln\left(\frac{V_2}{V_1}\right) = mRT \ln\left(\frac{P_1}{P_2}\right)$ \longrightarrow Isobaric process: P=constant $\Delta U = Q - W, W = \int P dV = P \int dV = P(V_2 - V_1)$ $Q = \Delta U + P(V_2 - V_1) = (U_2 - U_1) + P(V_2 - V_1)$ $= (U_2 + PV_2) - (U_1 + PV_1) = H_2 - H_1 = \Delta H$

 \rightarrow Constant volume process: V=constant Q-W= ΔU , W= $\int P dV = 0$, no work done $Q = \Delta U = m \Delta u = m \int c_v dT$ \rightarrow Adiabatic process: Q=0 $Q-W=\Delta U, -W=\Delta U$ $-\delta W = dU$ (infinitesimal increment of work and energy) dU+PdV=0, $\operatorname{mc}_{v} dT + \left(\frac{mRT}{V}\right) dV = 0$ $c_v dT + \left(\frac{RT}{V}\right) dV = 0, \frac{c_v}{R} \frac{dT}{T} = -\frac{dV}{V}$, integrate and assume

c_v=constant

$$\frac{c_{v}}{R} \ln\left(\frac{T_{2}}{T_{1}}\right) = -\ln\left(\frac{V_{2}}{V_{1}}\right), \quad \frac{T_{2}}{T_{1}} = \left(\frac{V_{1}}{V_{2}}\right)^{R/C_{v}} = \left(\frac{V_{1}}{V_{2}}\right)^{R/C_{v}}$$

$$\frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{k-1}, \text{ from ideal gas relation}$$

$$PV = RT, \left(\frac{V_1}{V_2}\right) = \left(\frac{T_1}{T_2}\right) \frac{P_2}{P_1}, \text{ substitute}$$

$$\frac{T_2}{T_1} = \left(\left(\frac{T_1}{T_2}\right) \left(\frac{P_2}{P_1}\right)\right)^{k-1}, \text{ multiply } \left(\frac{T_2}{T_1}\right)^{k-1} \text{ from both sides}$$

$$\frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{(k-1)/k}, \text{ and } \left(\frac{P_2}{P_1}\right) = \left(\frac{V_1}{V_2}\right)^k$$
Also $P_1V_1^k = P_2V_2^k \text{ and } pV^k = cons \tan t$
For an ideal gas undergoing adiabatic process

Polytropic Process: its P-V relation can be expressed as $PV^n = constant$, where n is a constant for a specific process

➤ Isothermal, T=constant, if the gas is an ideal gas then PV=RT=constant, n=1

Isobaric, P=constant, n=0 (for all substances)

➤ Constant-volume, V=constant, V=constant(P)^(1/n), n=∞ (for all substances)

> Adiabatic process, n=k for an ideal gas

$$P_{1}V_{1}^{n} = P_{2}V_{2}^{n} = PV^{n}$$

$$W = \int_{1}^{2} PdV = \int_{1}^{2} (P_{1}V_{1}^{n})V^{-n}dV$$

$$= (P_{1}V_{1}^{n})\int_{1}^{2} V^{-n}dV = \frac{(P_{1}V_{1}^{n})}{1-n}(V_{2}^{1-n} - V_{1}^{1-n}) = \frac{P_{2}V_{2} - P_{1}V_{1}}{1-n}$$