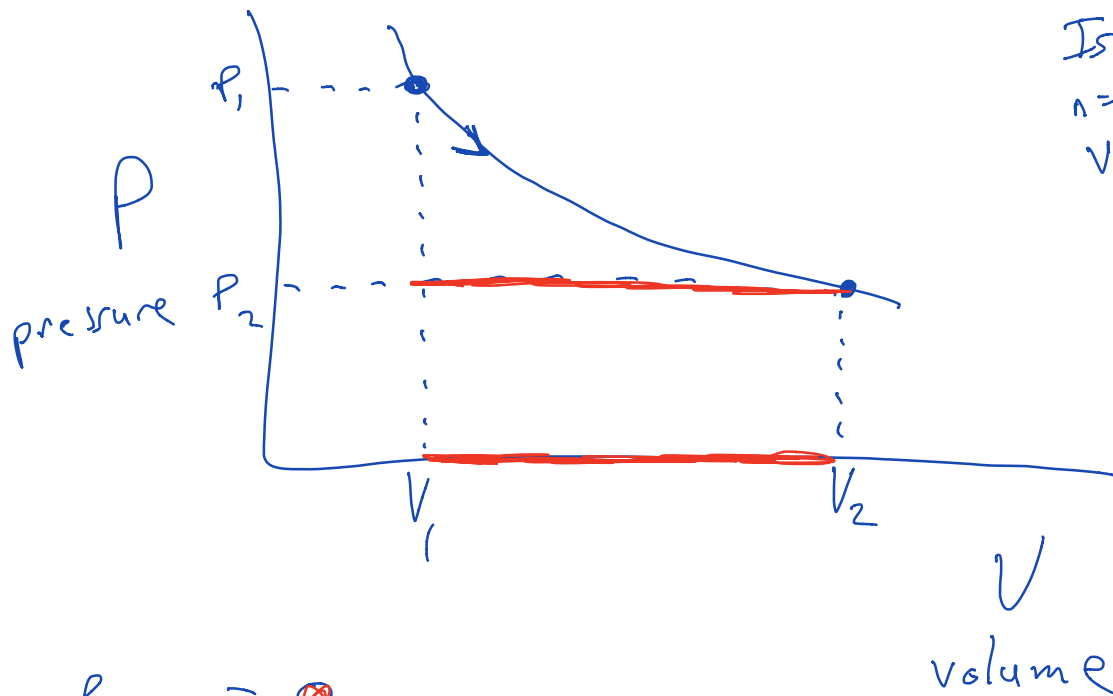


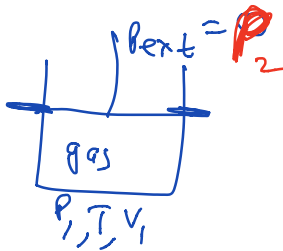
# Reversible Processes (paths)

$$I.G. P = \frac{nRT}{V}$$

Iso T.  
n = constant  
 $V_2 > V_1$



$$P_{\text{ext}} = P_2$$



$$W = 0$$

$$dw = -\underbrace{P_{\text{ext}}}_{0} dV$$

$$\therefore dW = 0$$

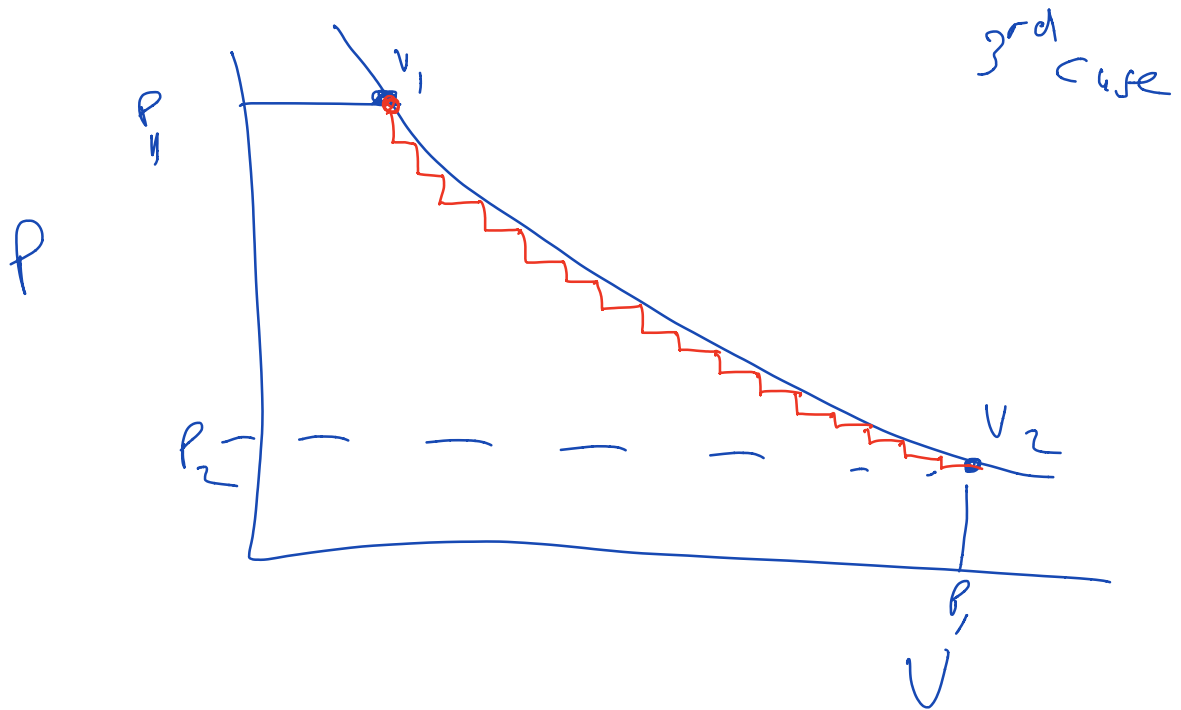
$$dw = -P_{\text{ext}} dV$$

$$\therefore \int dw = 0$$

$$W = -\int P_{\text{ext}} dV = -P_2(V_2 - V_1)$$

2<sup>nd</sup> case

$$\frac{= W}{1^{\text{st}} \text{ case}}$$



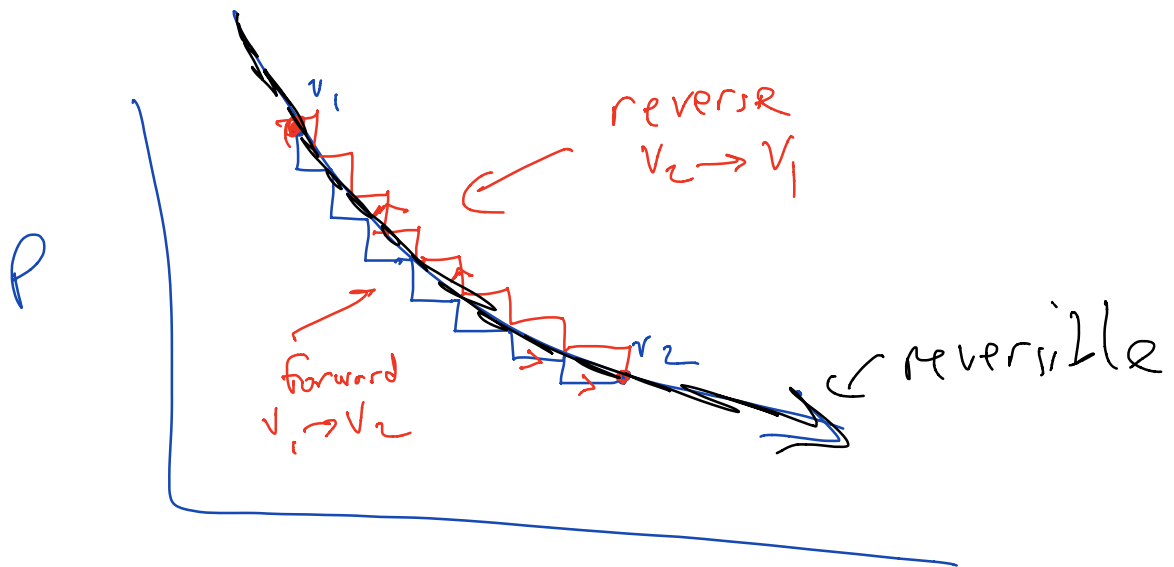
Reversible process

→ isothermal expansion of I.G.

→  $W = W_{max}$  done by system

→ system is always infinitesimally away from equilibrium

~~XXXX~~ → effectively: system always at equilibrium |||||



$V_1 \rightarrow V_2 \rightarrow V_1$

environment did net work

### Implication :

consider : process  $\underline{1}$   $\rightarrow$   $\underline{2}$   
 equil equil

$$\Delta U = U_2 - U_1 \leftarrow \underline{\text{state function}}$$

Reversible Process  $1 \rightarrow 2$   
Irreversible Process  $1 \rightarrow 2$

$$dU_{rev} = dq_{rev} + dW_{rev}$$

$$dU_{irrev} = dq_{irrev} + dW_{irrev}$$

$$0 = dq_{rev} - dq_{irrev} + dW_{rev} - dW_{irrev}$$

$$\underline{dW_{irrev} - dW_{rev} = dq_{rev} - dq_{irrev}}$$

$$0 < dq_{rev} - dq_{irrev}$$

$dq_{rev} > dq_{irrev}$