# **Chapter Three: isothermal expansion**

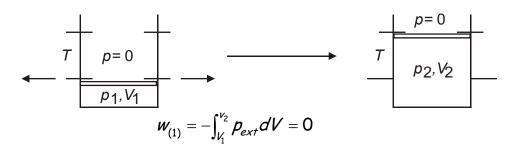
Isothermal Gas Expansion

$$(\Delta T=0)$$

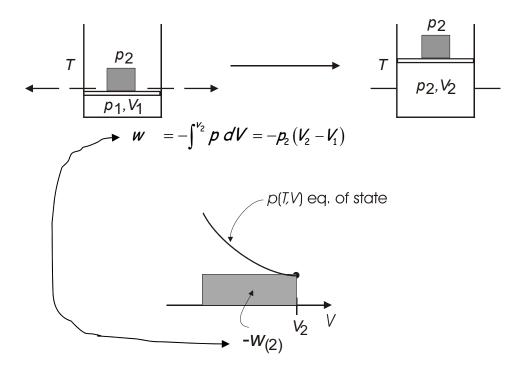
gas 
$$(p1, V1, T) = gas (p2, V2, T)$$

Irreversibly (many ways possible) (1)

# 1- Set pext = 0



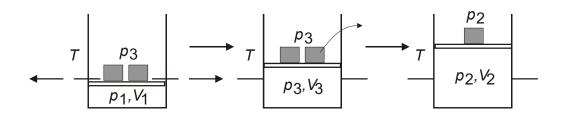
### 2- set Pext=P2



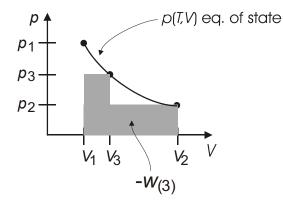
Note, work is negative: system expands against surroundings

#### (3) Carry out change in two steps

gas 
$$(p_1, V_1, T) = gas (p_3, V_3, T) = gas (p_2, V_2, T)$$
  
 $p_1 > p_3 > p_2$ 



$$w_{(3)} = -\int_{V_1}^{V_3} p_3 dV - \int_{V_3}^{V_2} p_2 dV = -p_3 (V_3 - V_1) - p_2 (V_2 - V_3)$$



More work delivered to surroundings in this case.

## For ideal gas

$$w_{rev} = -\int_{V_1}^{V_2} \frac{nRT}{V} dV = -nRT \ln \frac{V_2}{V_1} = nRT \ln \frac{p_2}{p_1}$$

# The Internal Energy U

$$dU = d^{T}q + d^{T}w$$
 (First Law)  
 $dU = C_{path}dT - p_{ext}dV$ 

And U (T,V) 
$$\Rightarrow dU = (\frac{\partial U}{\partial T})_V + (\frac{\partial U}{\partial V})_T dV$$

Some frequent constraints:

Reversible 
$$\rightarrow dU = \eth q_{rev} + \eth w_{rev} = \eth q_{rev} - PdV$$
,  $p = p_{ext}$ 

Isolated  $\rightarrow \eth q = \eth w = 0$ 

Adiabatic  $\rightarrow \eth q = 0 \quad \rightarrow dU = \eth w = -pdV$  (reversible)

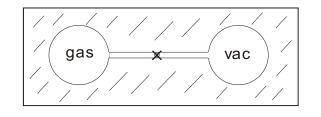
constant  $V \rightarrow w = 0 \quad \rightarrow dU = \eth q_V$ 

$$dU = (\frac{\partial U}{\partial T})_V dT + (\frac{\partial U}{\partial V})_T dV \qquad constant V$$
But also  $\delta q_V = (\frac{\partial U}{\partial T})_V dT$ 

$$\delta q_V = C_V dT \qquad \rightarrow (\frac{\partial U}{\partial T})_V = C_V \qquad \text{very important result}$$

so 
$$dU = c_V dT + (\frac{\partial U}{\partial V})_T dV$$

Joule free expansion of a gas  $(to get (\frac{\partial U}{\partial V})_T)$ 



Adiabatic 
$$q = 0$$

Expansion into Vac. 
$$w = 0$$
  
( $p_{ext}=0$ )

gas 
$$(p_1, T_1, V_1) = gas (p_2, T_2, V_2)$$
  
Since  $q = w = 0 \Rightarrow dU$  or  $\Delta U = 0$ 

$$recall dU = C_V dT + (\frac{\partial U}{\partial V})_T dV = 0$$

$$\left(\frac{\partial U}{\partial V}\right)_T dV_U = C_V dT_U$$

$$(\frac{\partial U}{\partial V})_T = -C_V(\frac{\partial T}{\partial V})_U$$
 measure in joule  $exp't$   $(\frac{\Delta T}{\Delta V})_U$ 

Joule did this 
$$\lim_{\Delta V \to 0} (\frac{\Delta T}{\Delta V})_U = (\frac{\Delta T}{\Delta V})_U = n_J$$

$$\therefore \quad dU = C_V dT - C_V n_J dV \qquad \qquad n_J \quad is \ a \ Joule \ coefficient$$

For Ideal gas 
$$\Rightarrow \eta_J = 0$$
 exactly

$$dU = C_V dT$$
 Always for ideal gas

U(T) only depends on T

The internal energy of an ideal gas depends only on temperature

#### Consequences

 $\Rightarrow \Delta U = 0$  For <u>all isothermal</u> expansions or compressions of <u>ideal gases</u>

 $\Rightarrow \Delta U = \int C_V dT$  For <u>any ideal gas</u> change in state