

Figure 1-9
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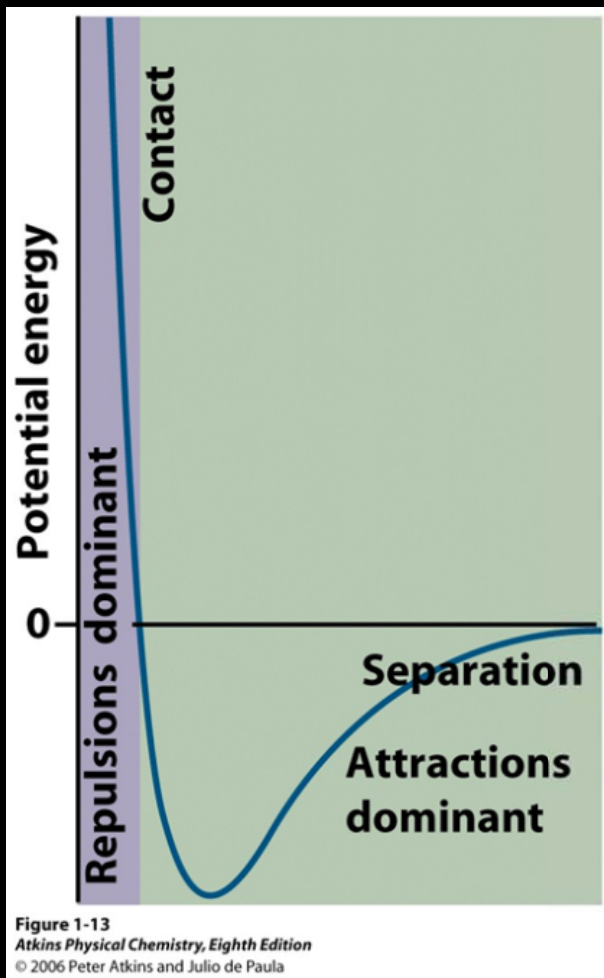


Figure 1-13
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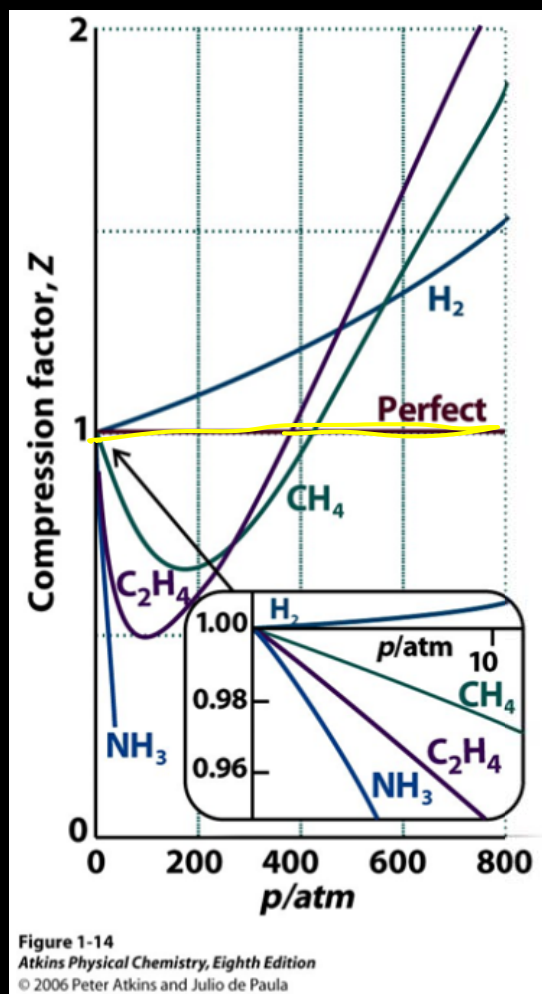


Figure 1-14
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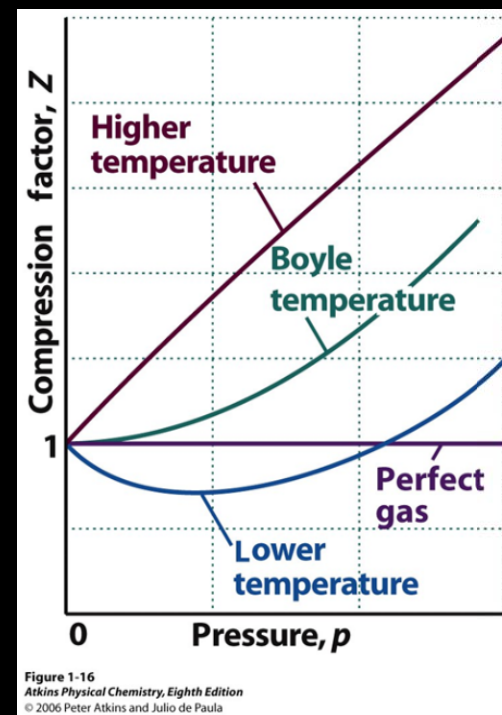


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Virial Eqn.

$$Z(T) = 1 + B'(T)p + C'(T)p^2 + \dots$$

$$= 1 + \frac{B(T)}{V_m} + \frac{C(T)}{V_m^2} + \dots$$

$$\frac{pV_m}{RT} = Z$$

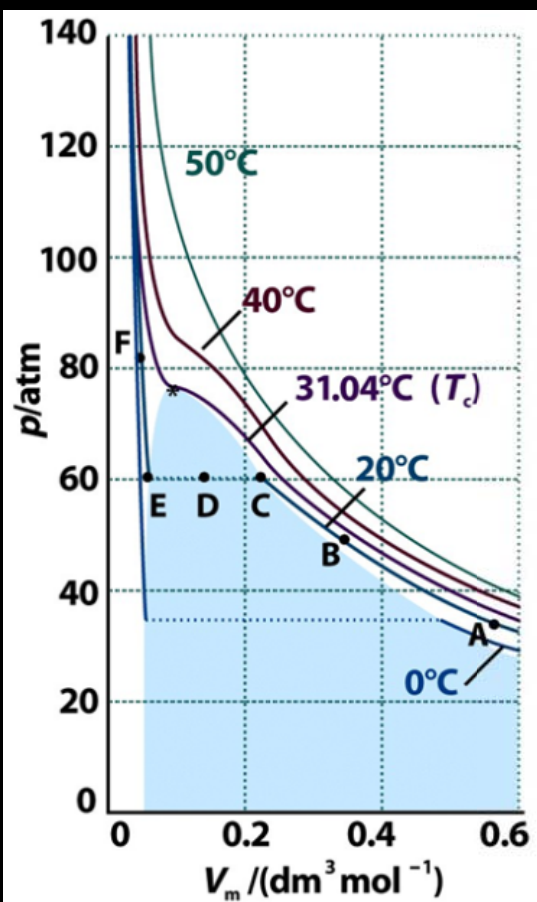


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$$\frac{dz}{dp} = B' + 2C'p + \dots$$

$p \rightarrow 0$

$$\frac{dz}{dp} = B'$$

$B'(T) \rightarrow 0$ as p is small
 $T_B \rightarrow$ Boyle's Temp.

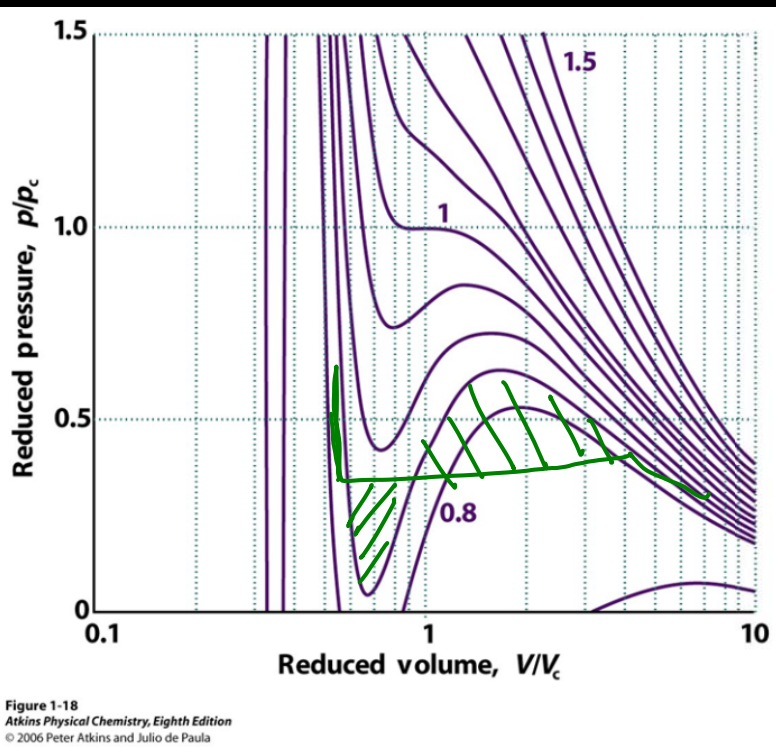
$$P_{id} V_{id} = nRT$$

$$V = V_{id} + nb$$

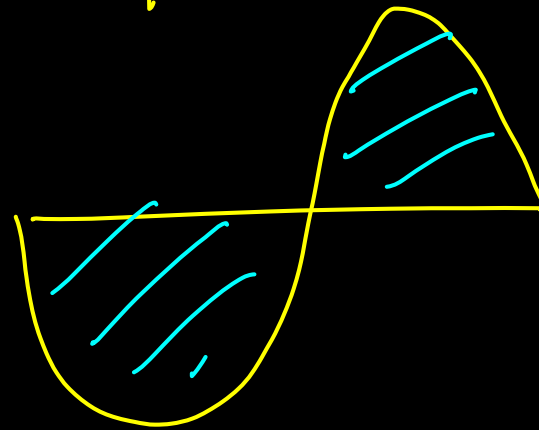
$$P = P_{id} - \frac{an^2}{V^2}$$

$$\left(P + \frac{an^2}{V^2} \right) (V - nb) = nRT$$

Van der Waals equation



Maxwell's construction
 Equal areas



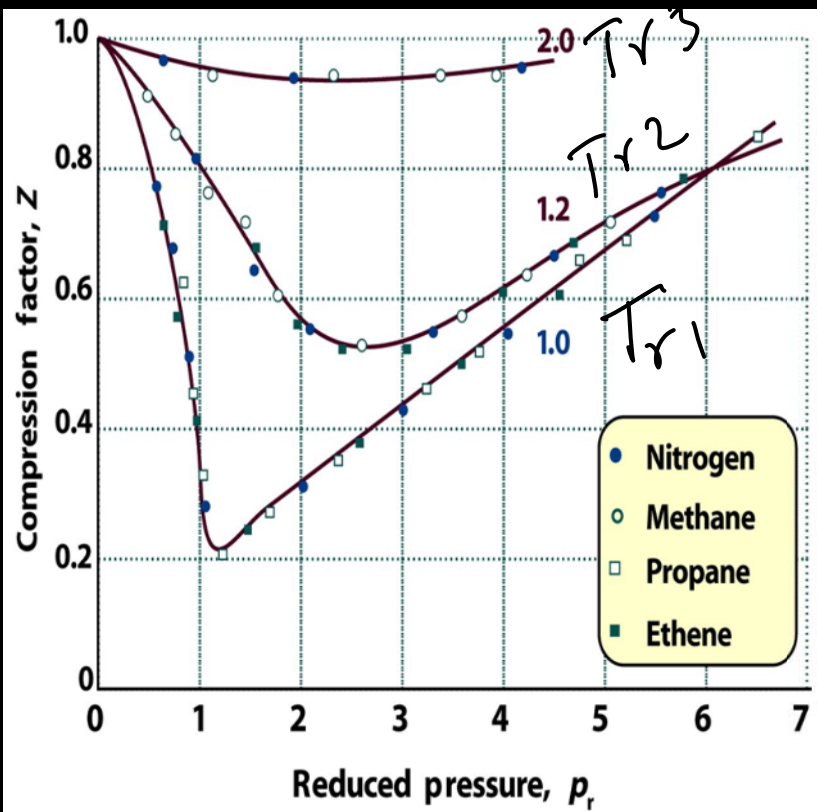


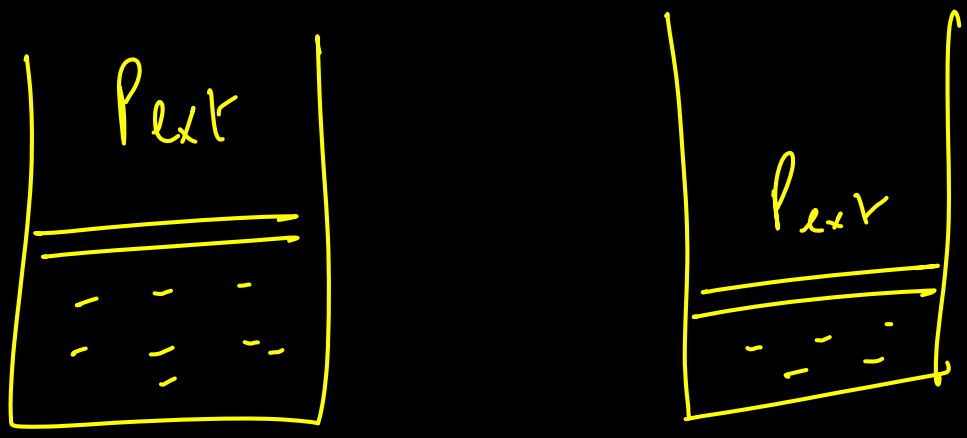
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$$p_r = \frac{p}{p_c}, \quad v_r = \frac{v}{v_c}$$

$$T_r = \frac{T}{T_c}$$

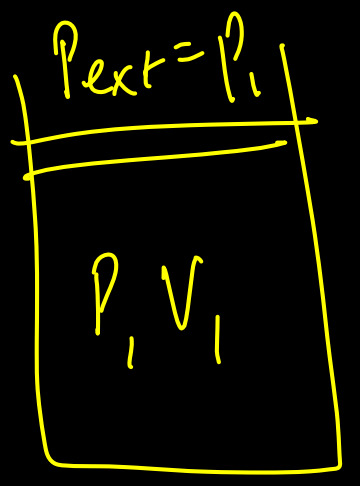
Principle of corresponding states

$$p_r = \frac{8 T_r}{3 v_r - 1} - \frac{3}{v_r^2}$$

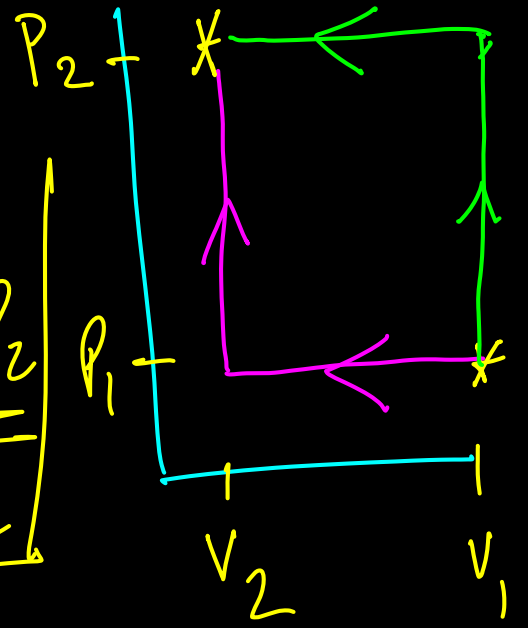
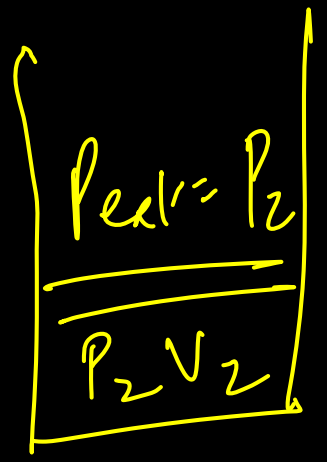


Energy into sys
+ve
Energy out of sys
-ve

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



compress →



$$w = - \int_{V_1}^{V_2} P_1 dV = - \int_{V_1}^{V_2} P dV = -P_1 (V_2 - V_1)$$

$$w = - \int_{V_1}^{V_2} P dV = - \int_{V_1}^{V_2} P_2 dV = -P_2 (V_2 - V_1)$$

dW, dq inexact

$(dW + dq)$ exact

dU

'U'
Energy
internal
energy

$$dU_{\text{sys}} = dW + dq, \quad dU_{\text{surr}} = - (dW + dq)$$

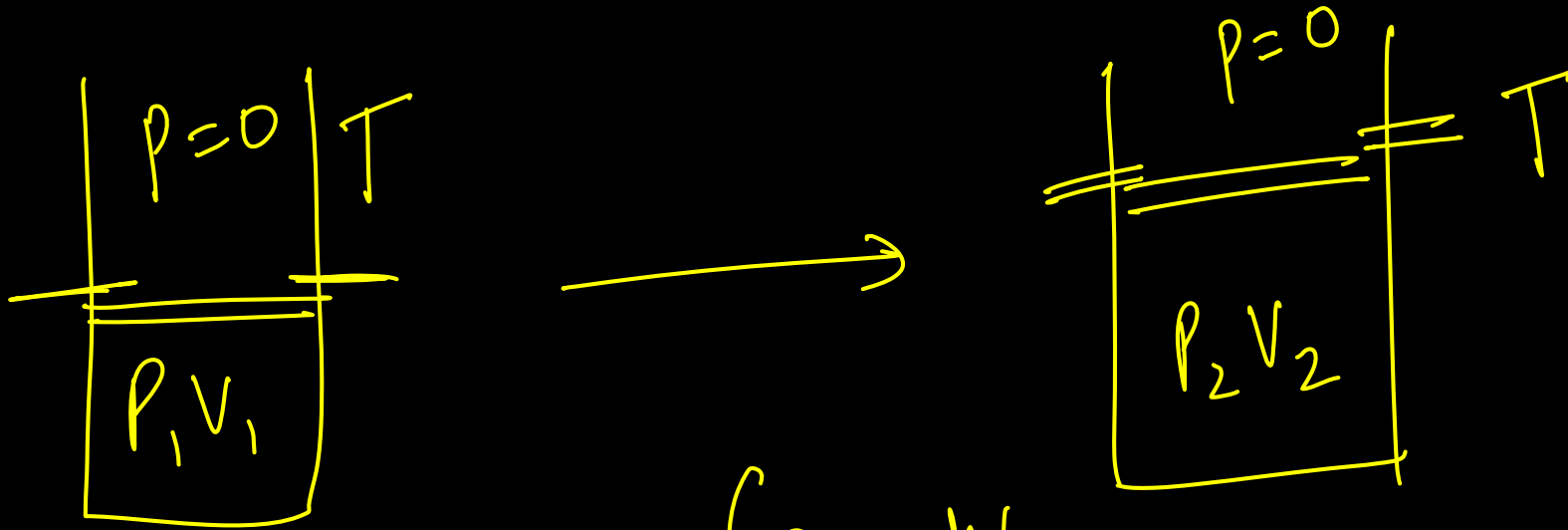
$$dU_{\text{sys}} + dU_{\text{surr}} = 0 = \text{Energy Conservation}$$

$$\Delta U_{\text{sys}} + \Delta U_{\text{surr}} = 0$$

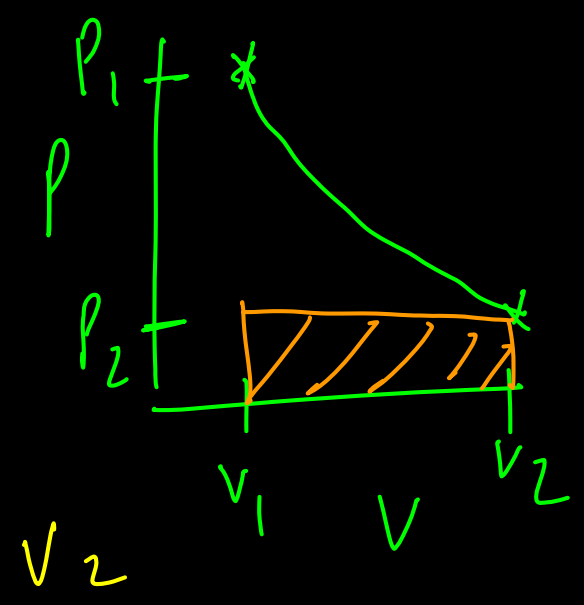
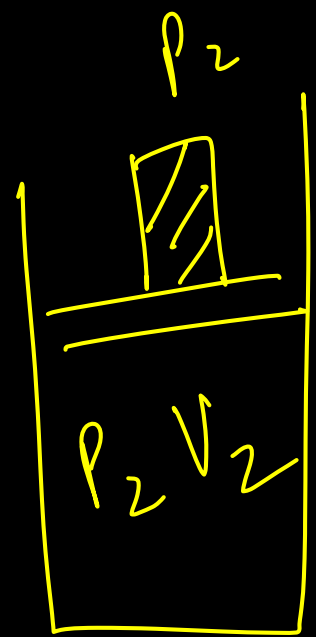
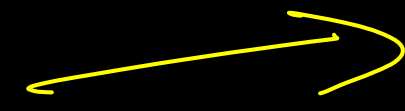
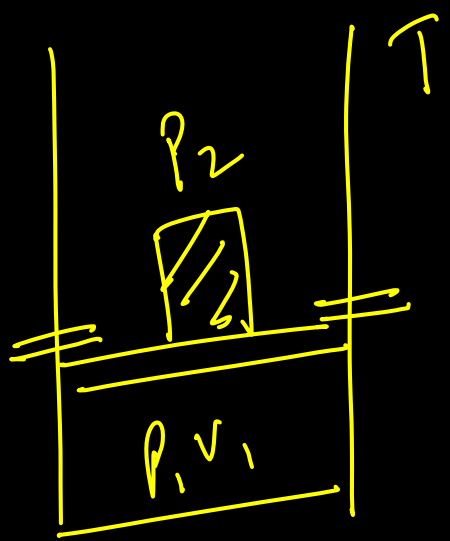
FIRST LAW

Isothermal Gas Expansion $\Delta T = 0$

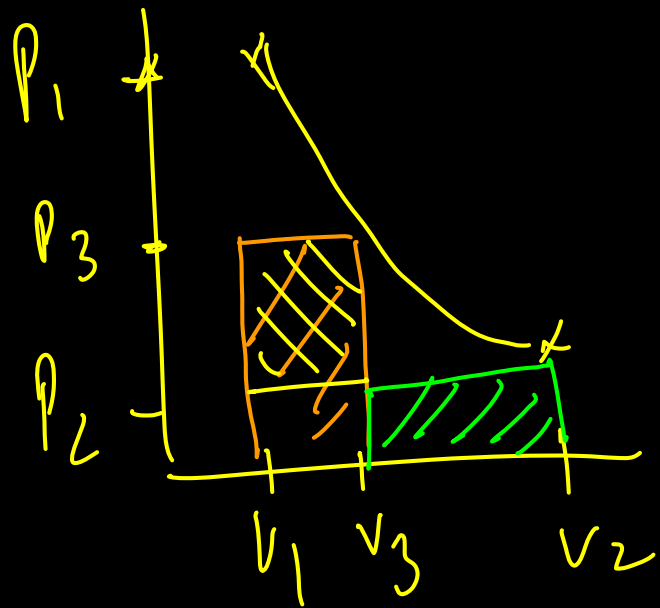
gas A (P_1, V_1, T) \longrightarrow A (P_2, V_2, T)



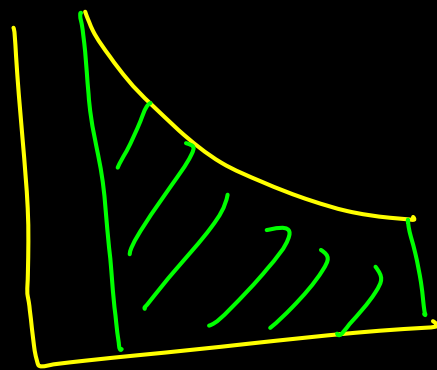
$$W = - \int P_{\text{ext}} dV = 0$$



$$\begin{aligned}
 W &= - \int P_{\text{ext}} dV = - P_2 \int_{V_1}^{V_2} dV \\
 &= - P_2 (V_2 - V_1)
 \end{aligned}$$



$$W = -P_3(V_3 - V_1) - P_2(V_2 - V_3)$$



Max work obtained under reversible conditions.

$$W = - \int P dV = - \int \frac{RT}{V} dV = RT \ln \frac{V_1}{V_2}$$

