

# CHEM2011 Chapter 1 problems

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## Question 1a,b

One mole of an ideal **monoatomic** gas initially at  $300\text{K}$  and pressure of  $15.0\text{atm}$  expands to a final pressure of  $1.00\text{atm}$ . The expansion can occur via any of five different paths. For each case, calculate the value of  $q$ ,  $w$ ,  $\Delta U$  and  $\Delta H$ .

a) Isothermal and reversible

### Solution::

Assume  $T_2 = T_1 \implies \Delta T = 0$  by isothermal. Because  $\Delta T = 0$ , we have that  $\Delta H = 0$  and  $\Delta U = 0$  as a consequence. So,  $\Delta U = q + w = 0 \implies w = -q$ .

We can now find the value of  $w$  by using the formula::

$$w = -nRT \ln\left(\frac{P_1}{P_2}\right) = -(1)(8.314)(300) \ln\left(\frac{15}{1}\right) = -6.75\text{E} - 3\text{J} \implies q = 6.75\text{E} - 3\text{J}$$

b) Isothermal and irreversible

### Solution::

$\Delta U$  and  $\Delta H$  are still 0 since  $\Delta T = 0$ . Because this process is irreversible,  $P_2 = P_{\text{external}}$ . So we have that

$$w = -P_2 \Delta V = -P_2(V_2 - V_1)$$

Apply ideal gas law to find  $V_1$  and  $V_2$ .

$$V_1 = \frac{nRT}{P_1} = \frac{(1)(0.08206)(300)}{15} = 1.641L, V_2 = \frac{nRT}{P_2} = 24.62L$$

We can now find our work for the irreversible process.

$$w = -1(24.62 - 1.641)\left(\frac{101.325J}{1Latm}\right) = -2.33E3J \implies q = 2.33E3J$$

- c) Isothermal and irreversible in a 2 step process. **Step 1::**  $P = 7atm$ , **Step 2::**  $P=1atm$ .

### Solution::

The process is still isothermal, so assume  $\Delta T, \Delta U, \Delta H = 0$  and  $q = -w$ .

Let  $P' = 7atm$  denote the intermediate pressure between the two steps. So we want to find the work between  $V_1$  and  $V'$  along with  $V'$  and  $V_2$ .

$$V_1 = 1.641L, V' = \frac{(1)(0.08206)(300)}{7} = 3.52L, V_2 = 24.62L$$

Now let us calculate the work done for both steps.

$$w = w_1 + w_2 = -P_2\Delta V = -((7)(3.52 - 1.641) + (1)(24.62 - 3.52)) \approx -34.251Latm\left(\frac{101.325}{Latm}\right) =$$

### Question 4

The constant pressure heat capacity of an ideal gas,  $A \in \mathbb{C}$  was found to vary with temperature according to the expression

$$\bar{C}_P = 22.17 + 0.32T \text{ in units of } JK^{-1}mol^{-1}$$

- a) Calculate  $q, w, \Delta U, \Delta H$  when the temperature of  $2mol$  of  $A$  is raised from  $0^\circ C \rightarrow 50^\circ C (273K \rightarrow 323K)$ . Assume  $P \equiv const$ .

### Solution::

First, we can find  $\Delta H$ .

$$\begin{aligned}\Delta H &= \int_{273}^{323} C_P dT = n \int_{273}^{323} \bar{C}_P dT = (2) \int_{273}^{323} 21.17 + 0.32T = 2 \left[ 21.17T + \frac{0.32}{2}T^2 \right]_{273}^{323} \\ &= 2 \left[ \left( 22.17(323) + \frac{0.32}{2}(323)^2 \right) - \left( 22.17(273) + \frac{0.32}{2}(273)^2 \right) \right] = 11753J = 11.75kJ\end{aligned}$$

Because we assumed  $P \equiv \text{const}$ ,  $\Delta H = q = 11.75kJ$ . Now we find  $\Delta U$ .

$$\Delta U = \Delta H - \Delta(PV) = \Delta H - nR\Delta T = 11753 - (2)(8.314)(323 - 273) = 10.92kJ$$

And so,  $w = \Delta U - q$ .

$$w = 10.92kJ - 11.75kJ = -0.83kJ$$

Because work is negative, work was done by the system, whereby we have an expansion.

b) Now assume  $V \equiv \text{const}$ .

### Solution

Because  $V \equiv \text{const}$ , we can carry over our  $\Delta H$  and  $\Delta U$  values since they're only dependent on temperature.

When volume is constant,  $w = 0$  since there is no expansion or compression. This implies that  $q = \Delta U = 10.92kJ$ .