# Thermodynamic Processes Solutions

GSI: Caleb Eades

9/13

## 1 Cycle Through the Cycles

This problem will develop a useful reference: a list of all quantities associated with thermodynamic processes of ideal gases. Suppose that there are N molecules of an ideal gas with d degrees of freedom (use  $\gamma = \frac{d+2}{d}$  where it is more convenient). Suppose the gas starts at  $(P_0, V_0)$ . Then  $T_0 = P_0 V_0/(Nk)$ . Complete the following table and  $draw\ each\ process\ on\ a\ P-V\ diagram$ .

Table 1: This table is also available in the workbook on pg. 153.

| Quantity   | Isochoric   | Isovolumetric                                      | Isothermal                              | Adiabatic  |
|------------|---|--|---|--|
| $P_f$      | $P_0$   | $P_f$ (given)                                      | $P_0 rac{V_0}{V_f}$                    | $P_0\left(rac{T_f}{T_0} ight)^{rac{\gamma}{\gamma-1}}$ |
| $V_f$      | $V_f$ (given)   | $V_0$  | $V_f$ (given)                           | $V_0 \left(\frac{T_0}{T_f}\right)^{\frac{1}{\gamma-1}}$  |
| $T_f$      | $T_0rac{V_f}{V_0}$   | $T_0 rac{P_f}{P_0}$                               | $T_0$                                   | $T_f$ (given)  |
| $\Delta E$ | $rac{d}{2}Nk_BT_0\left(rac{V_f}{V_0}-1 ight)$                   | $\frac{d}{2}Nk_BT_0\left(\frac{P_f}{P_0}-1\right)$ | 0 J                                     | $\frac{d}{2}Nk_B(T_f - T_0)$                             |
| Q          | $Nk_BT_0\left(\frac{V_f}{V_0}-1\right)\left(\frac{d}{2}-1\right)$ | $\frac{d}{2}Nk_BT_0\left(\frac{P_f}{P_0}-1\right)$ | $Nk_BT_0ln\left(\frac{V_f}{V_0}\right)$ | 0  |
| W          | $P_0(V_f-V_0)$  | 0 J  | $Nk_BT_0ln\left(rac{V_f}{V_0} ight)$   | $-\frac{d}{2}Nk_B(T_f-T_0)$                              |
| $\Delta S$ |   |  | $Nk_Bln\left(rac{V_f}{V_0} ight)$      | 0  |

## 2 Problems

#### 2.1 Heat from the Ocean

(a) Maximum efficiency is through Carnot's theorem combined with the efficiency of a Carnot cycle to get

$$e = 1 - \frac{T_L}{T_H} \tag{1}$$

So in this case evaluating yields

$$e = 1 - (273.15 + 4K)/(273.15 + 22K)$$

$$\approx 0.061$$

Thus, the maximum theoretical efficiency would be about 6.1%.

(b) The efficiency can also be written as

$$e = \frac{W}{Q_H} \tag{2}$$

Producing GW of power means 1 GJ of work is being done every second since  $[W] = \frac{[J]}{[s]}$ . Inverting the above equation,  $Q_H = \frac{W}{e}$ , so the heat per second is  $\frac{dQ_H}{dt} = \frac{P}{e}$ . The heat input is related to the volume by

$$\begin{split} Q &= mc\Delta T \\ &= \rho cV\Delta T \\ V &= \frac{Q}{\rho c\Delta T} \\ \frac{dV}{dt} &= \frac{P}{e\rho c\Delta T} \end{split}$$

Plugging in numbers, the volume processed in one second is

$$\frac{VolumeProcessed}{1s} = \frac{1 \times 10^{9} J/s}{(0.061)(1000 kg/m^{3})(480 J/kg * K)(22 - 4K)}$$

$$\approx 218 m^{3}$$

$$= 2.18 \times 10^{5} L$$

### 2.2 Challenge: Adiabatic Atmosphere

From the Ideal Gas Law, PV = nRT = NkT. Notice that  $\rho = \frac{Nm}{V}$ , so

$$P = \rho \frac{kT}{m} \tag{3}$$

Differentiating the given equation of  $P\rho^{-\gamma} = constant$ , we have

$$\begin{split} \frac{dP}{dh}\rho^{-\gamma} - \gamma\rho^{-\gamma-1}\frac{d\rho}{dh}P &= 0\\ \frac{dP}{dh} - \gamma\rho^{-1}\frac{d\rho}{dh}P &= 0\\ \frac{d\rho}{dh} &= \frac{\rho}{P}\frac{1}{\gamma}\frac{dP}{dh} \end{split}$$

From Eq.  $\ref{eq:posterior}, \, \frac{\rho}{P} = \frac{m}{kT}, \, \text{so}$ 

$$\frac{d\rho}{dh} = \frac{m}{kT} \frac{1}{\gamma} \frac{dP}{dh} \tag{4}$$

Now, differentiating Eq. ??,

$$\frac{dP}{dh} = \frac{kT}{m}\frac{d\rho}{dh} + \frac{\rho k}{m}\frac{dT}{dh} \tag{5}$$

Plutting into Eq. ??, we have

$$\begin{split} \frac{dP}{dh} &= \frac{kT}{m} \left( \frac{m}{kT} \frac{1}{\gamma} \frac{dP}{dh} \right) + \frac{\rho k}{m} \frac{dT}{dh} \\ \frac{\rho k}{m} \frac{dT}{dh} &= \frac{dP}{dh} \left( 1 - \frac{1}{\gamma} \right) \end{split}$$

Now, for any substance, the hydrodynamic condition posits that  $\frac{dP}{dh}=-\rho g$  (e.g., think of pressure in a lake). Plugging this into the above equation,

$$\frac{dT}{dh} = \frac{m}{\rho k} (-\rho g) \left( \frac{\gamma - 1}{\gamma} \right)$$
$$= -\frac{mg}{k} \frac{\gamma - 1}{\gamma}$$

If we assume the atmosphere is monoatomic,  $\gamma = \frac{5}{3}$ , so  $\frac{\gamma - 1}{\gamma} = \frac{2}{5}$  and we have

$$\frac{dT}{dh} = -2mg/5k\tag{6}$$