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In [1]: #####
# Futures
%matplotlib inline
# from __future__ import unicode_literals
# from __future__ import print_function

# Generic/Built-in
import datetime
import argparse

# Other Libs
from IPython.display import display, Image
import matplotlib.pyplot as plt
import numpy as np

# Owned
# from nostalgia_util import log_utils
# from nostalgia_util import settings_util
__authors__ = ["Osamu Katagiri - A01212611@itesm.mx"]
__copyright__ = "None"
__credits__ = ["Marcelo Videa - mvideo@itesm.mx"]
__license__ = "None"
__status__ = "Under Work"
#####
```

## Exercise 1

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In [6]: display(Image(filename='./directions/1.jpg'))
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1. Calculate the work that 1 mol of a perfect gas ( $\bar{C}_V = 25 \text{ J/mol K}$ ) produces when it expands from an initial volume of 10 L and  $31.5^\circ\text{C}$  to:
  - (a) a final volume of 50 L, reversibly and isothermically.
  - (b) irreversibly and isothermically against an external pressure of 0.5 atm.
  - (c) a final volume of 50 L reversibly and adiabatically.
  - (d) irreversibly and adiabatically against an external pressure of 0.5 atm.
  - (e) For all cases above, determine the final state after the expansion.

### For 1.a & 1.e (Isothermal Reversible Expansion)

$$W_{\text{REV}} = -RnT \ln \left( \frac{V_f}{V_i} \right)$$

where:

$$n = 1 \text{ mol}$$

$$R = 8.314 \text{ JK}^{-1} \text{ mol}^{-1}$$

$$T = 31.5^\circ \text{C}$$

$$V_i = 10 \text{ L}$$

$$V_f = 50 \text{ L}$$

$$W_{\text{REV}} = -4076.481 \text{ J}$$

$$\begin{aligned} P_f &= P_i V_i V_f^{-1} = 50657.202 \text{ Pa} \\ V_f &= 50 \text{ L} \\ T_f &= T_i = 304.65 \text{ K} \end{aligned}$$

### For 1.b & 1.e (Isothermal Irreversible Expansion)

$$W_{\text{IRREV}} = -P_{\text{ext}} (V_f - V_i)$$

where:

$$P_{\text{ext}} = 0.5 \text{ atm}$$

$$V_f = 50 \text{ L}$$

$$V_i = 10 \text{ L}$$

$$W_{\text{IRREV}} = -20 \text{ atm L}$$

$$\text{as: } 1 \text{ atm L} = 101.3 \text{ J}$$

$$W_{\text{IRREV}} = -2027 \text{ J}$$

$$\begin{aligned} P_f &= P_{\text{ext}} = 0.5 \text{ atm} = 50700 \text{ Pa} \\ V_f &= 50 \text{ L} \\ T_f &= T_i = 304.65 \text{ K} \end{aligned}$$

### For 1.c & 1.e (Adiabatic Reversible Expansion)

$$P_i = \frac{nRT_i}{V_i} = \frac{(1 \text{ mol})(8.314 \text{ JK}^{-1} \text{ mol}^{-1})(31.5^\circ \text{C})}{(10 \text{ L})}$$

$$P_i = 253286.01 \text{ Pa}$$

$$\gamma = \frac{\overline{C_p}}{\overline{C_v}} = \frac{\overline{C_v} + R}{\overline{C_v}} = \frac{(25 \text{ JK}^{-1} \text{ mol}^{-1}) + (8.314 \text{ JK}^{-1} \text{ mol}^{-1})}{25 \text{ JK}^{-1} \text{ mol}^{-1}}$$

$$\gamma = 1.33256$$

$$P_f = \frac{P_i V_i^\gamma}{V_f^\gamma} = \frac{(253286.01 \text{ Pa})(10 \text{ L})^{1.33256}}{(50 \text{ L})^{1.33256}}$$

$$P_f = 29661.406 \text{ Pa}$$

$$W_{\text{REV}} = \frac{P_f V_f - P_i V_i}{\gamma - 1} = \frac{(29661.406 \text{ Pa})(50 \times 10^{-3} \text{ m}^3) - (253286.01 \text{ Pa})(10 \times 10^{-3} \text{ m}^3)}{1.33256 - 1}$$

$$W_{\text{REV}} = -3156.693 \text{ J}$$

$$P_f = 29661.406 \text{ Pa}$$

$$V_f = 50L$$

$$T_f = T_i V_i^{\gamma-1} (V_f^{\gamma-1})^{-1} = 178.382K$$

### For 1.d & 1.e (Adiabatic Irreversible Expansion)

$$W_{\text{IRREV}} = \Delta U = -PdV = n\overline{C_v}dT$$

where:

$$n = 1mol$$

$$\overline{C_v} = 25JK^{-1}mol^{-1}$$

$$dT = T_f - T_i$$

$$T_i = 304.65K$$

$$P_i = 253286.01Pa$$

$$P_f = P_{ext} = 0.5atm = 50700Pa$$

$$T_f = \frac{T_i \left( \overline{C_v} + \frac{RP_f}{P_i} \right)}{\overline{C_p}}$$

$$T_f = 243.839K$$

$$W_{\text{IRREV}} = (1mol)(25JK^{-1}mol^{-1})(T_f - T_i)$$

$$W_{\text{IRREV}} = -1520.275J$$

$$V_f = \frac{\frac{P_f = P_{ext} = 0.5atm = 50700Pa}{(1mol)(8.314JK^{-1}mol^{-1})(243.839K)}}{\frac{50700Pa}{T_f = 243.839K}} = 40L$$

## Exercise 2

In [8]: `display(Image(filename='./directions/2.jpg'))`

2. A solar collector is used as a heat source for a Carnot engine with a heat sink at 300 K. The efficiency of the solar collector  $\varepsilon$  is defined as the fraction of the energy reaching the collector that is actually absorbed. It is related to the temperature of the collector as follows:

$$\varepsilon = 0.75 - 1.75 \left( \frac{T}{300K} - 1 \right)$$

Determine the best operating temperature of the collector.

Solar collector  $\varepsilon$  is given by:

$$\varepsilon = 0.75 - 1.75 \left( \frac{T}{300K} - 1 \right)$$

The Carnot engine efficiency  $\eta$  is given by:

$$\eta = \frac{T}{300K} - 1$$

The system's efficiency  $\epsilon$  is given by:

$$\epsilon = \varepsilon\eta$$

$$\epsilon = -\frac{1.75T^2}{(300K)^2} + \frac{4.25T}{300K} - 1.5$$

Let's find  $\frac{\partial \epsilon}{\partial T}$

$$\frac{\partial \epsilon}{\partial T} = -\frac{2(1.75)}{300^2}T + \frac{4.25}{300}$$

$$\frac{\partial \epsilon}{\partial T} = \frac{17}{1200} - \frac{7}{180000}T$$

Let's find where the function is the maximum by calculating T for  $\frac{\partial \epsilon}{\partial T} = 0$

$$\frac{17}{1200} - \frac{7}{180000}T = 0$$

$$T = 354.286K$$

$$\varepsilon = 0.75 - 1.75 \left( \frac{T}{300K} - 1 \right); \text{ with } T = 354.286K$$

$$\varepsilon = 0.433$$

The operating temperature shall be  $T = 354.286K$  to yield the maximum power from the Carnot engine, with an efficiency of  $\varepsilon = 0.433$

## Exercise 3

In [9]: `display(Image(filename='./directions/3.jpg'))`

3. Produce a single graph of  $P_f/P_i$  vs.  $V_f/V_i$  for the following expansion processes:

- (a) Reversible isothermic process.
- (b) Reversible adiabatic processes for monoatomic, diatomic and polyatomic perfect gases.
- (c) Irreversible adiabatic processes for monoatomic, diatomic and polyatomic perfect gases.

Compare the plots in the graph and discuss the different behaviors observed. Choose the representation of your axes wisely.

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**Function to compute the "next" pressure value for REVERSIBLE processes**

$$P = \frac{nRT^\gamma}{V^\gamma}$$

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**Function to compute the "next" pressure value for IRREVERSIBLE processes**

$$\begin{aligned}dU &= -PdV = n\overline{C}_v dT \\ n\overline{C}_v \int_{T_i}^{T_f} dT &= -P_f \int_{V_i}^{V_f} dV \\ n\overline{C}_v (T_f - T_i) &= -P_f (V_f - V_i) \\ n\overline{C}_v \left( \frac{P_f V_f}{nR} - \frac{P_i V_i}{nR} \right) &= -P_f (V_f - V_i) \\ \frac{n\overline{C}_v P_f V_f}{nR} - \frac{n\overline{C}_v P_i V_i}{nR} &= -P_f (V_f - V_i) \\ P_f \left( \frac{n\overline{C}_v V_f}{nR} + V_f - V_i \right) &= \frac{n\overline{C}_v P_i V_i}{nR}\end{aligned}$$

$$P_f = \frac{\overline{C}_v P_i V_i}{\overline{C}_v V_f + V_f R - V_i R}$$

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In [5]: # Function to compute the "next" pressure value for REVERSIBLE processes
def P_rev(n, R, T, V, gamma):
    return (n*R*(T**gamma))/(V**gamma);

# Function to compute the "next" pressure value for IRREVERSIBLE processes
def P_irrev(Cv, Pi, Vi, Vf, R):
    return (Cv*Pi*Vi)/(Cv*Vf+Vf*R-Vi*R);

# Draw the plot's workspace
n = 6;
plt.subplots(figsize=(3*n, 2*n))

# Initial State, Constants & Final Volume
Ti = 304.65; #K
Vi = 0.01; #10 L
Vf = 0.05; #50 L
V = np.linspace(Vi,Vf,1000);
n = 1; #mol
R = 8.314; #J/K mol)

# Reversible Isothemic Process
gamma = 1;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_rev(n, R, Ti, V, gamma);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Reversible Isothemic Process');

# Reversible Adiabatic Process (monoatomic)
Cv = (3/2)*R;
Cp = Cv + R;
gamma = Cp/Cv;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_rev(n, R, Ti, V, gamma);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Reversible Adiabatic Process (monoatomic)');

# Reversible Adiabatic Process (diatomic)
Cv = (5/2)*R;
Cp = Cv + R;
gamma = Cp/Cv;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_rev(n, R, Ti, V, gamma);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Reversible Adiabatic Process (diatomic)');

# Reversible Adiabatic Process (polyatomic)
Cv = 3*R;
Cp = Cv + R;
gamma = Cp/Cv;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_rev(n, R, Ti, V, gamma);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Reversible Adiabatic Process (polyatomic)');

# Irreversible Adiabatic Process (monoatomic)
Cv = (3/2)*R;
Cp = Cv + R;
gamma = Cp/Cv;
#Tf = 243.839; #K (from exercise 1.d)
#Vf = (n*R*Tf)/(50700); #L
#V = np.linspace(Vi,Vf,1000);
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_irrev(Cv, Pi, Vi, Vf, R);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Irreversible Adiabatic Process (monoatomic)');

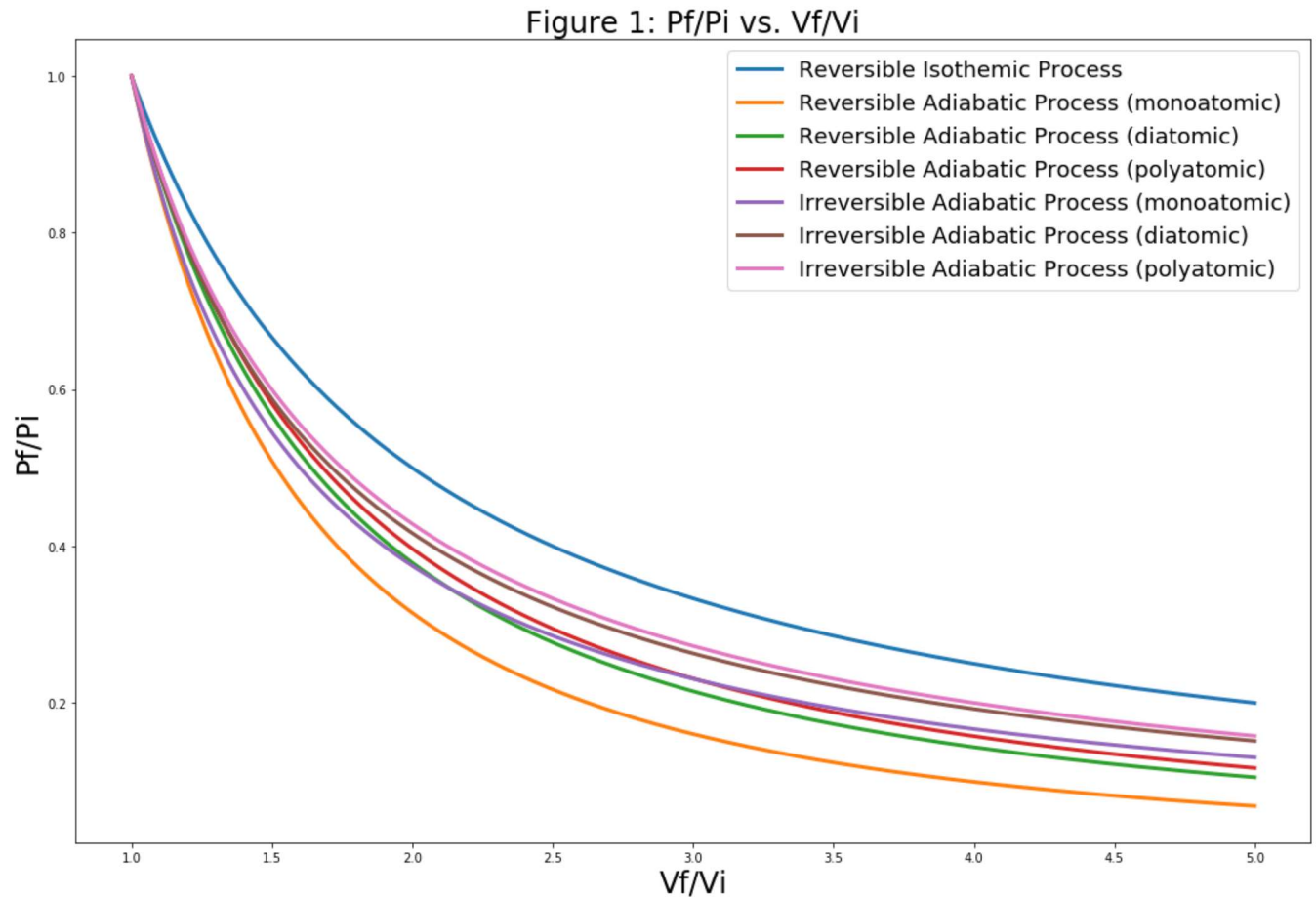
# Irreversible Adiabatic Process (diatomic)
Cv = (5/2)*R;
Cp = Cv + R;
gamma = Cp/Cv;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_irrev(Cv, Pi, Vi, Vf, R);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Irreversible Adiabatic Process (diatomic)');

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```
# Irreversible Adiabatic Process (polyatomic)
Cv = 3*R;
Cp = Cv + R;
gamma = Cp/Cv;
Vf = V;
Pi = P_rev(n, R, Ti, Vi, gamma);
Pf = P_irrev(Cv, Pi, Vi, Vf, R);
plt.plot(Vf/Vi, Pf/Pi, '-', linewidth=3, label='Irreversible Adiabatic Process (polyatomic)');

plt.xlabel('Vf/Vi', fontsize=24);
plt.ylabel('Pf/Pi', fontsize=24);
plt.title("Figure 1: Pf/Pi vs. Vf/Vi", size=24)
plt.legend(prop={'size': 18})
display(plt);

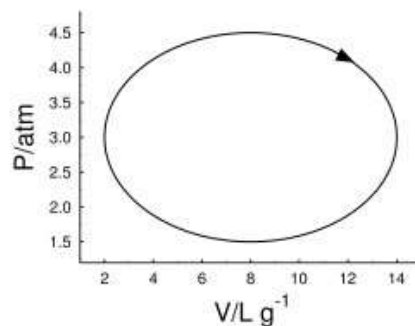
<module 'matplotlib.pyplot' from 'C:\\Users\\oskat\\Anaconda3\\lib\\site-packages\\matplotlib\\pyplot.py'>
```



## Exercise 4

In [10]: `display(Image(filename='./directions/4.jpg'))`

4. A  $PV$  diagram of an imaginary heat engine is represented in the following figure.



- Find the work performed per cycle by 1 g of working fluid.
- Find the engine efficiency if it rejects 5.7 kJ/g during each cycle

The work performed per cycle is given by the area within the PV curve.

$$W_{performed} = \pi(14 - 8)(4.5 - 3.0)atmLg^{-1}$$
$$W_{performed} = 9\pi atmLg^{-1}$$

as:  $1atmL = 101.3J$

$$W_{performed} = \frac{9117}{10}\pi Jg^{-1} = 2864.190Jg^{-1}$$

$$\eta_{engine} = \frac{W_{performed}}{W_{performed} + W_{rejected}}$$

$$\eta_{engine} = \frac{2864.190Jg^{-1}}{2864.190Jg^{-1} + 5700Jg^{-1}} = 0.334$$

## Exercise 5

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In [3]: display(Image(filename='./directions/5.jpg'))
```

5. A heat engine, of which all steps in a cycle are reversible, absorbs thermal energy from a high-temperature reservoir, performs an amount of net work  $w_{\text{net}}$ , and rejects thermal energy into a low-temperature reservoir. Initially, the reservoirs are at temperatures  $T_1$  and  $T_2$ . Their heat capacities are constant with values  $C_1$  and  $C_2$ , respectively. Calculate what will be the final temperatures for the heat reservoirs and the maximum amount of work produced by the engine.



Let's denote some variables:

$dQ_H$  = heat transfer out of the hot reservoir

$dQ_C$  = heat transfer into the cold reservoir

$dW$  = work output by the engine

$dT_H$  = change in temperature of the hot reservoir ( $dT_H < 0$  as it's cooling down)

$T_{Ho} = T_1$  = initial temperature of the hot reservoir

$dT_C$  = change in temperature of the cold reservoir ( $dT_C > 0$  as it warms up)

$T_{Co} = T_2$  = initial temperature of the cold reservoir

$T$  = final temperature of both reservoirs

$\eta$  = 2nd law efficiency given

As both reservoirs have temperature-independent heat capacities  $C_1$  and  $C_2$ , then it follows that:

$$\begin{aligned} dQ_H &= -C_1 dT_H \\ dQ_C &= C_2 dT_C \end{aligned}$$

The engine efficiency can be defined as:

$$\varepsilon = \frac{dW}{dQ_H} = 1 - \frac{dQ_C}{dQ_H} = 1 + \frac{dT_C}{dT_H}$$

With the following Carnot limiting efficiency:

$$\varepsilon_c = 1 - \frac{T_C}{T_H}$$

The 2nd law efficiency  $\eta$  is assumed to be constant.

$$\eta = \frac{W_{net,out}}{Q_H} = \frac{Q_H - Q_C}{Q_H} = 1 - \frac{Q_C}{Q_H}$$

Rearranging terms,  $T_C$  can be defined as a function of  $T_H$ :

$$T_H \frac{dT_C}{dT_H} + \eta T_C = (\eta - 1)T_H$$

The complementary solution of the previous is  $AT_H^{-\eta}$  where  $A$  is a constant to tailor the initial conditions. On the other hand, by trying a particular solution of the equation that is proportional to  $T_H$ , the following is obtained  $(\eta - 1)T_H/(\eta + 1)$ . Adding together these complementary and particular solutions and fitting to the initial temperatures of the two reservoirs gives:

$$\frac{T_C}{T_{Ho}} = \left( \frac{T_H}{T_{Ho}} \right)^\eta \left\{ \frac{T_{Co}}{T_{Ho}} + \frac{1 - \eta}{1 + \eta} \left( 1 - \left( \frac{T_H}{T_{Ho}} \right)^{\eta+1} \right) \right\}$$

Let's set  $T_C = T_H = T$  to find the common final temperature of the two reservoirs:

$$\frac{T}{T_{Ho}} = \left( \frac{(T_{Co} + T_{Ho}) + \eta(T_{Co} - T_{Ho})}{2T_{Ho}} \right)^{\frac{1}{1+\eta}}$$

$$T = T_{Ho} \left( \frac{(T_{Co} + T_{Ho}) + \eta(T_{Co} - T_{Ho})}{2T_{Ho}} \right)^{\frac{1}{1+\eta}}$$

where:

$T_{\min} = \sqrt{T_{Co} T_{Ho}}$  is the geometric average of the initial temperatures

$T_{\max} = \frac{T_{Co} + T_{Ho}}{2}$  is their arithmetic average

The total work output by the engine is the difference in heat transfers for the two reservoirs, as follows:

$$W = Q_H - Q_C = C_1(T_{Ho} - T) - C_2(T - T_{Co})$$

And the maximum amount of work is given by:

$$W_{max} = C_1 C_2 (T_{max} - T_{min})$$

The maximum work is achieved for reversible operation of the engine when  $\eta = 1$ . On the other hand, zero work is output for maximally irreversible operation of the engine when  $\eta = 0$ .

## Comment Summary