

```
In [1]: import numpy as np
import matplotlib.pyplot as plt
```

Problem 3.1

Part A

```
In [2]: n = 0.05 #num moles
Va = 0.0001 #Volume (m^3)
Th = 400 #Temp (K)
Vb = 0.0003 #Volume (m^3)
Vc = 0.0005 #Volume (m^3)
Tc = 284.5 #Temp (K)
Vd = 0.000167 #Volume (m^3)
R = 8.3145 #IDG const. J/m^3*K

#Heatflow into engine:
q_out = n*R*Th*np.log(Va/Vb) #Heat in Joules
q_in = -q_out
print(q_in)
```

182.68823748261997

Part B

```
In [3]: #work done by engine
w_carnot_cycle = -n*R*Th*np.log(Va/Vb) - n*R*Tc*np.log(Vc/Vd) #Work in Joules
print(w_carnot_cycle)
```

52.98753986550585

Part D

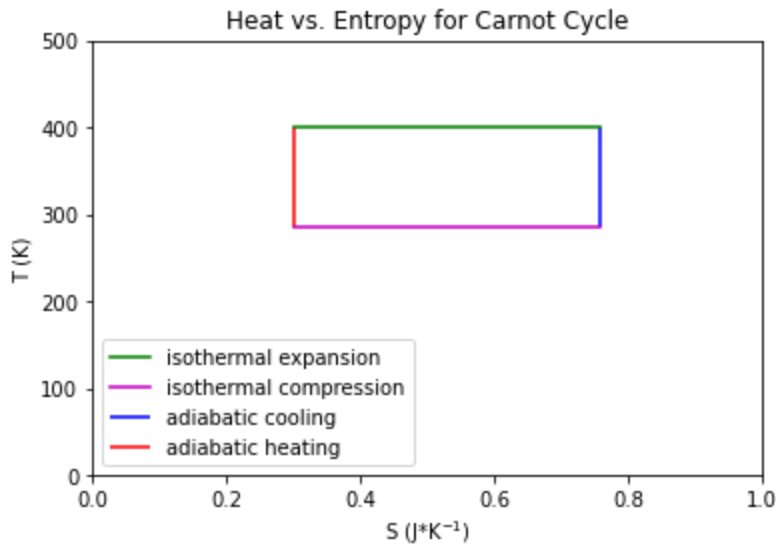
```
In [4]: S_a = 0.3 #J/K
DeltS_Th = n*R*np.log(Vb/Va) #J/K
S_b = DeltS_Th + S_a #J/K
DeltS_Tc = n*R*np.log(Vd/Vc) #J/K
S_c = S_b
S_d = DeltS_Tc + S_c

Sh = np.linspace(S_a, S_b, 100)
Ths = [Th for x in range(100)]

Sc = np.linspace(S_c, S_d, 100)
Tcs = [Tc for x in range(100)]

plt.plot(Sh, Ths, label="isothermal expansion", color = 'g')
plt.vlines(x = S_b, ymax = Th, ymin = Tc, color = 'b', label="adiabatic cooling")
plt.plot(Sc, Tcs, label="isothermal compression", color = 'm')
plt.vlines(x = S_d, ymax = Th, ymin = Tc, color = 'r', label="adiabatic heating")
plt.axis([0,1,0,500])
plt.ylabel("T (K)")
plt.xlabel("S (J*K$^{-1}$)")
plt.legend()
plt.title("Heat vs. Entropy for Carnot Cycle")
```

Out[4]: Text(0.5, 1.0, 'Heat vs. Entropy for Carnot Cycle')



Part E

```
In [5]: A = (S_b - S_a) * (Th - Tc) #Simple area formula: Delta_Y * Delta_X
print(A) #J (about equal to work!)
```

52.75122857310652

Problem 3.2

```
In [6]: T_initial = 25 + 273.15 #C -> K
T_final = 100 + 273.15 #C -> K
V_initial = 0.5 * 0.001 #L -> m^3
V_final = 1 * 0.001 #L -> m^3
P_initial = 1 * 101325 #Atm -> Pa
n = P_initial * V_initial / (R * T_initial)

Cv = (3/2) * n * R #Assume Ar behaves as a monoatomic idg
Cp = Cv + n*R

Del_S_isochoric = Cv*np.log(T_final / T_initial)
Del_S_isobaric = Cp*np.log(V_final / V_initial)
Del_S_tot = Del_S_isochoric + Del_S_isobaric
print(Del_S_tot) #Total Entropy Change in J/K

0.3516457707793893
```

Problem 3.3

Part A

```
In [7]: Ntot = 10
Vtot = 60
nL = 10
VL = 10
nR = 20
VR = Vtot - VL

def n_choose_k(n, k):
    return np.math.factorial(n) / (np.math.factorial(k) * np.math.factorial(n-k))

def Boltzmann_Entropy(W):
    Avog = 6.022e23
    return (R/Avog)*np.log(W)

def Find_Entropy_Stats(nL, ntot, VL, Vtot):
    W_Left = n_choose_k(VL, nL) #Multiplicity of Left
    W_Right = n_choose_k(Vtot-VL, ntot-nL) #Multiplicity of Right
    W_Tot = W_Left * W_Right

    S_Left = Boltzmann_Entropy(W_Left)
    S_Right = Boltzmann_Entropy(W_Right)
    S_Total = S_Left + S_Right
    return (W_Left, W_Right, W_Tot, S_Left, S_Right, S_Total)

print("Left Mult.: \t{}\nRight Mult.: \t{}\nTotal Mult.: \t{}\n\nLeft S: \t\t{}\nRight S: \t\t{}\nTotal S: \t\t{}".format(*Find_Entropy_Stats(nL, nL+nR, VL, Vtot)))

Left Mult.:      1.0
Right Mult.:     47129212243960.0
Total Mult.:     47129212243960.0

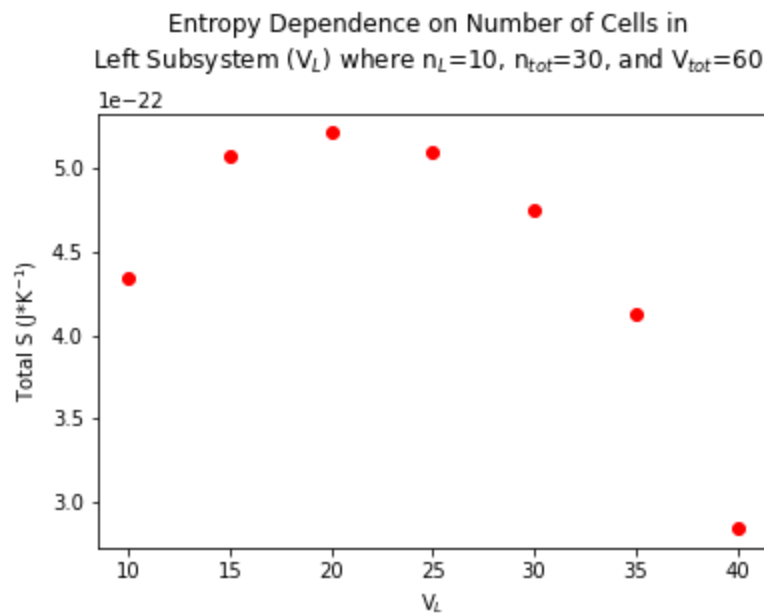
Left S:          0.0
Right S:         4.346944605367547e-22
Total S:         4.346944605367547e-22
```

Part B

The equilibrium position will lie where entropy (S) is maximized. From the plot shown below, this is at $V_L=20$ and $V_R=40$

```
In [8]: Vls = [x*5 + 10 for x in range(0,7)]
S_total_vals = [Find_Entropy_Stats(nL, nL+nR, y, Vtot)[5] for y in Vls]
plt.plot(Vls, S_total_vals, 'or')
plt.xlabel("V$_L$")
plt.ylabel("Total S (J*K$^{-1}$)")
plt.title("Entropy Dependence on Number of Cells in \nLeft Subsystem (V$_L$) where n$_L$=10, n$_{tot}$=30, and V$_{tot}$=60 \n")
```

```
Out[8]: Text(0.5, 1.0, 'Entropy Dependence on Number of Cells in \nLeft Subsystem (V$_L$) where n$_L$=10, n$_{tot}$=30, and V$_{tot}$=60 \n')
```



Problem 3.5

Part C

Interpretation:

The entropy of the system increases up until around $E_{tot}=6$ and then begins to decrease. The corresponding Temperatures for a change in E (always 1) and the change in entropy calculated below is maximized when Entropy is maximized which is reflected in the dS being smallest around $E_{tot}=6$. Since internal energy changes are constantly 1 on the Y axis, for this system, large changes in S correspond to low temperatures and smaller changes in S correspond to high temperatures. This is consistent with what we should expect from the given relationship $T = dU/dS$. Small entropy changes reflect high temperature when internal energy is low, but when it is high, larger entropy changes are required to achieve the same temperature.

```

In [9]: E_tots = [3, 4, 5, 6, 7, 8, 9]
W_tots = [1, 3, 6, 7, 6, 3, 1]
T_tots = []

Boltzmann_S = [R/6.022e23 * np.log(x) for x in W_tots]
prev = 0
dS = []
for i in Boltzmann_S:
    sub = i - prev
    dS.append(sub)
    prev = i

T = [abs(1/j) for j in dS if j != 0 ]

plt.plot(Boltzmann_S, E_tots, 'r-o')
plt.ylabel("E$_{tot}$")
plt.xlabel("S (J*K$^{-1}$)")
plt.title("Plot of E$_{tot}$ [U] vs. Entropy for System of 3 Molecules and Energy Levels")

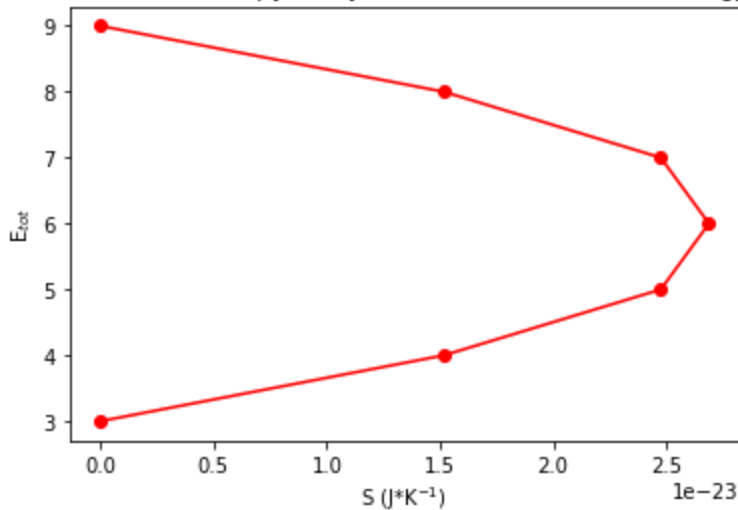
print("Temperature From Changes:")
print(T)

```

Temperature From Changes:

```
[6.592652141135143e+22, 1.0449106423998241e+23, 4.698499328891361e+23, 4.698499328891361e+23, 1.0449106423998241e+23, 6.592652141135143e+22]
```

Plot of E_{tot} [U] vs. Entropy for System of 3 Molecules and Energy Levels



Secret Code 20:

3.1 A)	Isothermal Expansion Adiabatic expansion Isothermal Compression Adiabatic Compression	$\frac{q_{\text{carnot}}}{nRT_H \ln(V_A/V_B) = q_{\text{in}}}$ 0 $nRT_C \ln(V_C/V_D)$ 0	$\frac{w}{nRT_H \ln(V_A/V_B)}$ $\frac{C_V(T_H - T_C)}{nRT_C \ln(V_C/V_D)}$ $C_V(T_C - T_H)$
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$$q_{\text{out}} = C_{T_H} = nRT_H \ln(V_A/V_B) = -q_{\text{in}}$$

$$|q_{\text{in}} = 182.7 \text{ J}|$$

$$3.1 B) \quad W_{\text{tot}} = nRT_H \ln(V_A/V_B) + nRT_C \ln(V_C/V_D)$$

$$= -52.99 \text{ J}$$

$$= 52.99 \text{ J done by system}$$

$$3.1 C) \quad \text{Total Area: (assuming axes the same as above)}$$

$$(0.0008 \text{ m}^3)(2.0 \times 10^6 \text{ Pa}) = 1000 \text{ J}$$

$$\therefore \text{Paper energy density: } \frac{1000 \text{ J}}{1467.8 \text{ mg}} = 0.6813 \text{ J/mg}$$

$$\text{Carnot weight} = 78.7 \text{ mg}$$

$$\text{Carnot energy} = 0.6813 \frac{\text{J}}{\text{mg}} \times 78.7 \text{ mg} = |53.62 \text{ J}|$$

$$3.1 D) \quad \text{Isothermal } \Delta S: nR \ln(V_F/V_I) = S_F - S_I$$

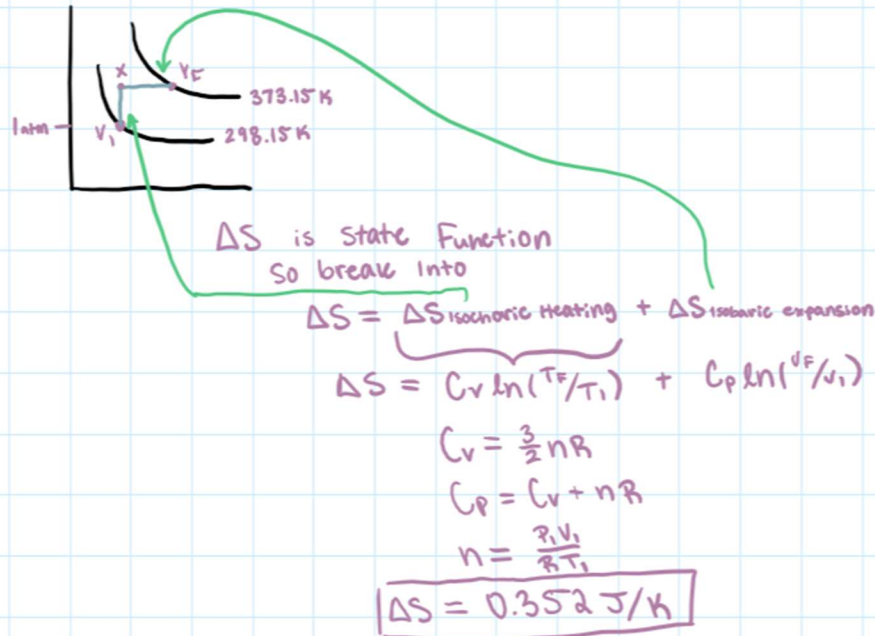
$$\text{given } S_I = 0.3 \text{ J/K}$$

$$S_F = nR \ln(V_F/V_I) + S_I$$

$$\text{Adiabatic } \Delta S: 0 \quad \Delta T = T_H - T_C$$

$$\begin{aligned}
 3.1E) \quad A &= \Delta S \times \Delta T \\
 &= (S_F - S_i) \cdot (T_H - T_C) = [J/K \cdot K] \\
 &= \underline{152.8 \text{ J} = W \text{ done by cycle}}
 \end{aligned}$$

3.2)



$$3.4A) \quad dB = e dK - J dL$$

$$dB = \left(\frac{\partial B}{\partial K} \right) dK - \left(\frac{\partial B}{\partial L} \right) dL$$

$$3.4B) \quad e = \left(\frac{\partial B}{\partial K} \right)$$

$$J = \left(\frac{\partial B}{\partial L} \right)$$

3.4C)

$$\left(\frac{\partial e}{\partial L} \right)_K = \left(- \frac{\partial J}{\partial K} \right)_L$$

$$\begin{aligned}
 dB &= e dK + -J dL \\
 \parallel & \quad \frac{\partial e}{\partial K} \quad \quad \frac{-\partial J}{\partial K} \\
 e dK & \quad \quad \parallel \\
 + & \quad \quad \frac{\partial e}{\partial K} \quad \quad \frac{-\partial J}{\partial L} \\
 -J dL & \quad \quad \frac{\partial e}{\partial K} \quad \quad \frac{-\partial J}{\partial L}
 \end{aligned}$$

3.5A)

$\epsilon_3 = 3$							
$\epsilon_2 = 2$							
$\epsilon_1 = 1$							
	$W_{3,1} = 1$	$W_{4,1} = \frac{3!}{2!}$	$W_{5,1} = \frac{3!}{2!}$	$W_{6,1} = 3!$	$W_{7,1} = \frac{3!}{2!}$	$W_{8,1} = \frac{3!}{2!}$	$W_{9,1} = 1$
	$W_{3,2} = 0$	$W_{4,2} = 0$	$W_{5,2} = \frac{3!}{2!}$	$W_{6,2} = 1$	$W_{7,2} = \frac{3!}{2!}$	$W_{8,2} = 0$	$W_{9,2} = 0$
	$W_{3,3} = 0$	$W_{4,3} = 0$	$W_{5,3} = 0$	$W_{6,3} = 0$	$W_{7,3} = 0$	$W_{8,3} = 0$	$W_{9,3} = 0$

$E_{\text{tot}} = 3 \quad E_{\text{tot}} = 4 \quad E_{\text{tot}} = 5 \quad E_{\text{tot}} = 6 \quad E_{\text{tot}} = 7 \quad E_{\text{tot}} = 8 \quad E_{\text{tot}} = 9$

3.5B)

$W_{3,\text{tot}} = 1 \quad W_{4,\text{tot}} = 3 \quad W_{5,\text{tot}} = 6 \quad W_{6,\text{tot}} = 7 \quad W_{7,\text{tot}} = 6 \quad W_{8,\text{tot}} = 3 \quad W_{9,\text{tot}} = 1$