Worksheet for Exploration 21.4: Entropy of Expanding Ideal Gas



In this animation, N = nR (i.e., $k_B = 1$). This, then, gives the ideal gas law as PV = NT. Restart

In thermodynamic processes, the entropy depends not on the path taken but on the end points. It is a "state function" (in contrast with heat and work which depend on the process). Since $Q = \Delta U + W$

and $\Delta U = (3/2)nR\Delta T$ (for a monatomic gas),

 $\Delta S = \int dQ/T = \int (3/2)nRdT/T + \int PdV/T = nR[(3/2)In(T_f/T_i) + In(V_f/V_i)].$

Thus $\Delta S = (3/2)N \ln(Tf/Ti) + N \ln(Vf/Vi)$ for an ideal monatomic gas (note that In represents the natural log, base e).

In the animations, note that the area under the PV diagram is equal to the work.

a. What is the work done in each case?

W_{isobaric}=____

W_{isochoric}=____

W_{isothermal}=____

b. What is the heat absorbed or released in each case?

Q_{isobaric}=_____

Q_{isochoric}=____

Q_{isothermal}=____

c. What is the area under the associated TS diagram? (Note that the choice of the initial entropy is arbitrary.)

Area_{isobaric}=_____

Area_{isochoric}=_____

Area_{isothermal}=_____

d. How does the change in entropy compare for the three processes?

e. Compare your measurements from the graphs to the calculated values found using the equation above for an ideal monatomic gas.

Another way to measure the change in entropy is to use $Q = mc\Delta T$ or, for a gas, $Q = CN\Delta T$. In this case,

 $\Delta S = CN \ln(T_f/T_i).$

f. Show that for the isobaric expansion where $C = C_P = (5/2)$, you get this change in entropy.