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Physics 141. Lecture 24.

- Course Information.
- Continue our discussion of Chapter 13:
 - Equation of state
 - The energy distribution of an ideal gas and energy exchange with its environment.

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Physics 141. Course information.

- Homework 10 is due on Friday December 8 at noon.
- Homework set 11 is due on Friday December 15 at noon.
- To calculate the final homework grade, I remove the lowest homework grade and then take the average of the remaining 10 homework grades. If you are happy with homework grades 1 10, you can consider homework 11 as optional.

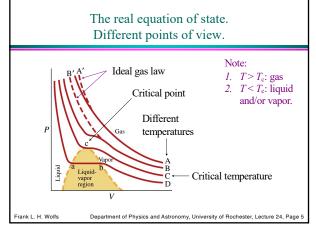
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- I will send a summary of all your grades in the course on Wednesday 12/13, including the results of a calculation that will show what grade you need to get on the final exam to get an A-, a B-, and a C- in this course.
- The final exam will take place on Wednesday 12/20 at 4 pm in Hoyt. The exam will take 3 hours and cover all the material discussed in Phy 141, except the error analysis.
- Extra office hours will be scheduled before the final exam. Details will be announced via email.

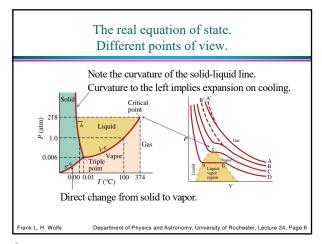
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The real equation of state. Different points of view. Note the curvature of the solid-liquid line. Curvature to the left implies expansion on cooling. Critical 218 Solid Vapor 0.00 0.01 T (°C) 56.6 T (°C) Water CO₂

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The first law of thermodynamics. Adding/removing heat from a system.

- Consider a closed system:

 - No change in
 - energy allowed (exchange with environment)
 - · Isolated system:
 - stem that does not allow an exchange of energy
- The internal energy of the system can change and will be equal to the heat added to the system minus the work done by the system: $\Delta U = Q - W$ (note: this is the work-energy theorem).

- Note: keep track of the signs:
 Heat: Q > 0 J means heat added, Q < 0 J means heat lost
 Work: W > 0 J mean work done by the system, W < 0 J means work

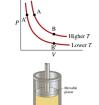
done on the system

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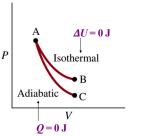
The first law of thermodynamics. Isothermal processes.

- An isothermal process is a process in which the temperature of the system is kept constant.
- This can be done by keeping the system in contact with a large heat reservoir and making all changes slowly.
- Since the temperature of the system is constant, the internal energy of the system is constant: $\Delta U = 0$ J.
- The first law of thermodynamics thus tells us that Q = W.



The first law of thermodynamics. Adiabatic processes.

- An adiabatic process is a process in which there is no flow of heat (the system is an isolated system).
- Adiabatic processes can also occur in non-isolated systems, if P the change in state is carried out rapidly. A rapid change in the state of the system does not allow sufficient time for heat flow.
- The expansion of gases differs greatly depending on the process that is followed (see Figure).



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Work done during expansion/compression.

- Consider an ideal gas at pressure p.
- ullet The gas exerts a force F on a moveable piston, and F = pA.
- If the piston moves a distance dl, the gas will do work:

$$dW = Fdl$$

Note: F and dl are parallel.

• The work done can be expressed in terms of the pressure and volume of the gas:



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Work done during expansion/compression. Isobaric and isochoric processes.

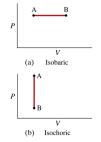
• Isobaric process:

- Processes in which the pressure is kept constant.
 W_{A>B} = pdV = p_A(V_B V_A)

· Isochoric process:

· Processes in which the volume is kept constant. • $W_{A\rightarrow B} = p_A(V_B - V_A) = 0$





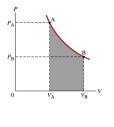
Work done during expansion/compression. Isothermal process.

• Isothermal process:

$$p = \frac{NkT}{V}$$

• The work done during the change from state A to state B is

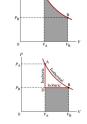
$$W = \int_{V_A}^{V_B} p dV = NkT \int_{V_A}^{V_B} \frac{1}{V} dV$$
$$= NkT \ln \left(\frac{V_B}{V_A} \right)$$



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Work done during expansion/compression.

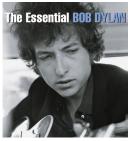
- The work done during the expansion of a gas is equal to the area under the pV curve.
- ullet Since the shape of the pV curve depends on the nature of the expansion, so does the work
 - Isothermal: $W = NkT \ln(V_B/V_A)$
 - Isochoric: W = 0
 - Isobaric: $W = p_B (V_B V_A)$
- The work done to move state A to state B can take on any value!



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2 Minute 48 Second Intermission.



- · Since paying attention for 1 hour and 15 minutes is hard when the topic is physics, let's take a 2 minute 48 second intermission.
- You can:
 Stretch out.
- Talk to your neighbors.
 Ask me a quick question.
 Enjoy the fantastic music.
- Solve a WeBWorK problem.



First law of thermodynamics. Molecular specific heat.

- When we add heat to a system, its temperature will increase.
- · For solids and liquids, the increase in temperature is proportional to the heat added, and the constant of proportionality is called the specific heat of the solid or liquid.
- When we add heat to a gas, the increase in temperature will depend on the other parameters of the system. For example, keeping the volume constant will results in a temperature rise that is different from the rise we see when we keep the pressure constant (the heat capacities will differ):

(Constant Volume) (Constant Pressure)

Here, C_V and C_P are the molecular specific heats for constant volume and constant pressure.

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First law of thermodynamics. Molecular specific heat (p = constant).

- · Consider what happens when we add Q_p to the system while keeping its pressure constant (p = NkT/V =constant).
- The work done by the gas will be $p\Delta V$.
- \bullet Using the ideal gas law, we can rewrite Pthe work done by the gas as $W = p\Delta V = Nk\Delta T.$
- The change in the internal energy of the gas is thus equal to

 $\Delta U = Q_p - W = Q_p - Nk\Delta T$

• Using the definition of C_P we can rewrite this relation as

 $\Delta U = NC_P \Delta T \text{-} Nk \Delta T = N(C_P \text{-} k) \ \Delta T$

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V

Isobaric

(a)

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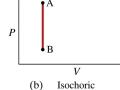
First law of thermodynamics. Molecular specific heat (V = constant).

- Consider what happens when we add Q_V to the system while keeping its volume constant (V =NkT/p = constant).
- The work done by the gas will be $p\Delta V = 0$ J.
- The change in the internal energy of the gas is thus equal to

 $\Delta U = Q_V = NC_V \Delta T$

· Note: we also know from the Boltzmann distribution that

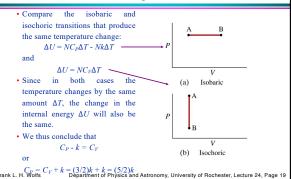
 $\Delta U = (3/2)Nk\Delta T$ · We thus conclude that



Note: if the molecules have more than 3 degrees of freedom, C_V will increase!

 $C_V=(3/2)k.$

First law of thermodynamics. Molecular specific heat.



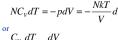
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Adiabatic processes (Q = 0 J). What is the shape of the pV curve?

- The change in the internal energy of
- the gas is $N(3/2k) \Delta T = NC_V \Delta T$.
- The first law of thermodynamics thus tells us that

$$NC_V \Delta T = NC_V \int_{V_A}^{V_B} dT = -\int_{V_A}^{V_B} p dV$$

 Comparing the integrands we must require that



 $\frac{C_V}{k}\frac{dT}{T} + \frac{dV}{V} = 0$

A Isothermal B Adiabatic C

What is the shape of the pV curve?

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Adiabatic processes (Q = 0 J).

• Integrating each term in the previous expression shows that

$$\frac{C_V}{k}\ln T + \ln V = \ln T^{\frac{C_V}{k}} + \ln V = \ln V T^{\frac{C_V}{k}} = \text{constant}$$

or $VT^{\frac{C_V}{k}} = \left(TV^{\frac{k}{C_V}}\right)^{\frac{C_V}{k}} = \text{constant}$

 This expression can also be written in terms of the pressure and volume (which is of course what we need to defined the curve in the pressure versus volume graph):

curve in the pressure versus volume graph):
$$TV^{\frac{k}{C_{V}}} = \left(\frac{pV}{Nk}\right)V^{\frac{k}{C_{V}}} = \frac{pV^{\frac{k}{C_{V}}}}{Nk} = \frac{pV^{\frac{k}{C_{V}}}}{Nk} = \frac{pV^{\frac{k}{C_{V}}}}{Nk} = \text{constant}$$
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