

# Classical Physics

- at end of the 19<sup>th</sup> century:

- Mechanics - Newton's Laws
- Electromagnetism - Maxwell's Eqns
- Optics - Geometric (particles) vs. Physical (waves)
- Thermodynamics - Four Laws (0-3)
- Gas Laws - Kinetic Theory

- overlaps often led to important discoveries:

• Maxwell's Eqns → EM radiation (optics)

• Newton's laws / kinetic theory



microscopic/atomic description of macroscopic gas laws

- around 1900, Theoretical problems:

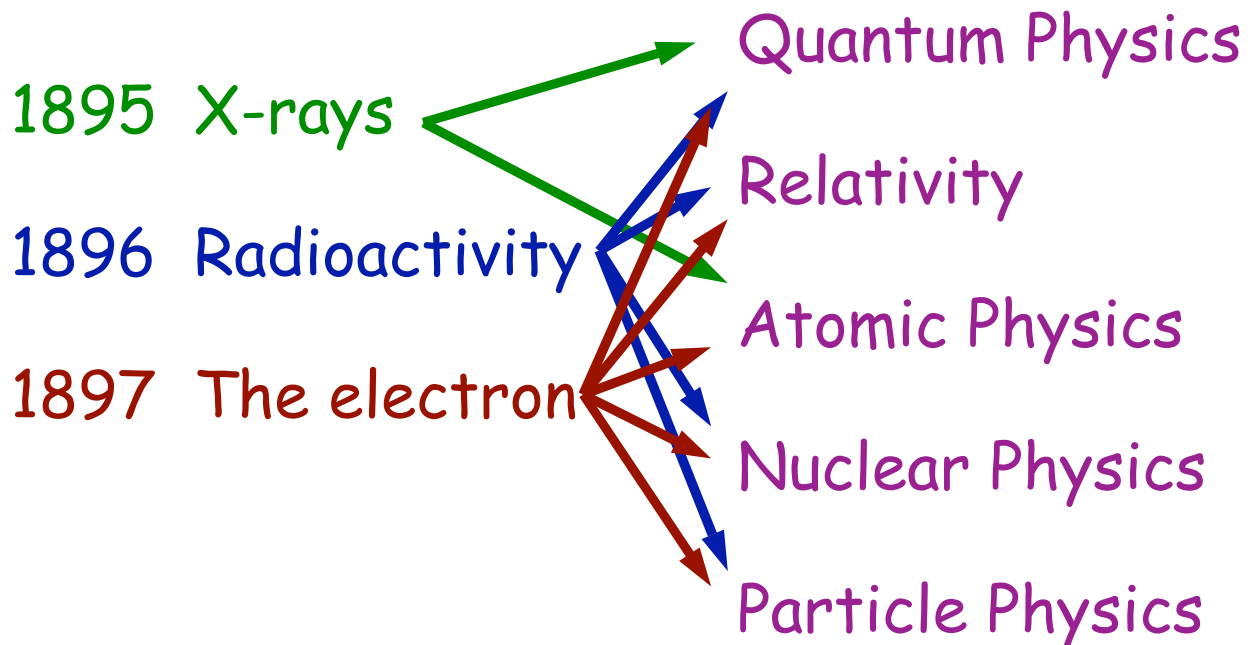
1) What is EM medium?

→ Relativity

2) Blackbody Radiation

→ Quantum Physics

- Experimental discoveries:



But first...

# Heat and Thermodynamics

- study of Thermal Energy of systems

Temperature: a measure of thermal energy, units of Kelvins

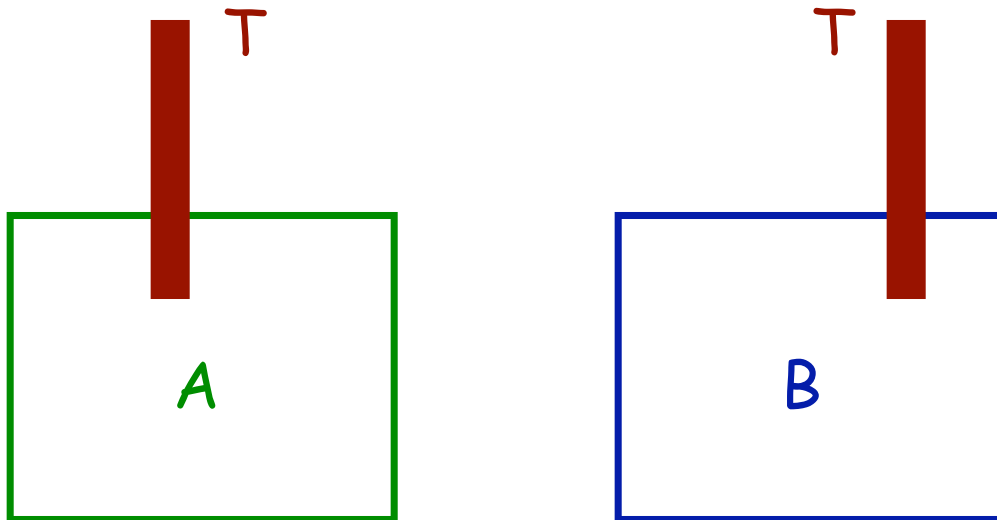
Room Temp ~ 290 K

Temperature of an object is measured by the change in some physical property.

Measuring device is called a thermometer.

# Zeroth Law of Thermodynamics

If bodies **A** and **B** are each in thermal equilibrium with a third body **T**, then they are in thermal equilibrium with each other.



**Thermal equilibrium:** all measurable properties unchanging.

Objects in thermal equilibrium are at the same temperature.

## Temperature Scales

- Daniel Fahrenheit (1686-1736)
  - $0^{\circ}\text{F}$  = mixture of ice, water, salt
  - $100^{\circ}\text{F}$  = Human body temp ( $\sim 98.6^{\circ}\text{F}$ )
- Anders Celsius (1701-1744)
  - $0^{\circ}\text{C}$  = Freezing point of  $\text{H}_2\text{O}$
  - $100^{\circ}\text{C}$  = Boiling point of  $\text{H}_2\text{O}$
- Lord Kelvin (1824-1907)

$\text{H}_2\text{O}$  boil :  $100^{\circ}\text{C} = 212^{\circ}\text{F} = 373 \text{ K}$

$\text{H}_2\text{O}$  freeze :  $0^{\circ}\text{C} = 32^{\circ}\text{F} = 273 \text{ K}$

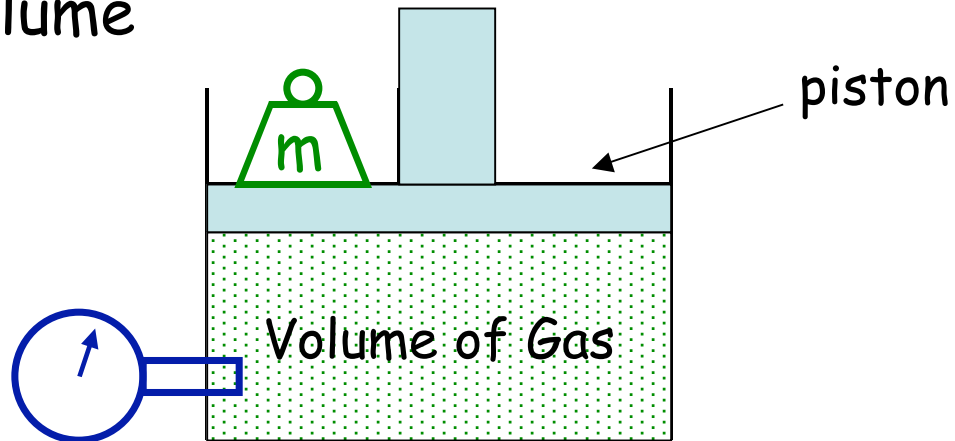
Absolute zero :  $-273^{\circ}\text{C} = -460^{\circ}\text{F} = 0 \text{ K}$

$$T_C = T_K - 273.15$$

$$T_F = (9/5)T_C + 32$$

## Constant-Volume Gas Thermometer

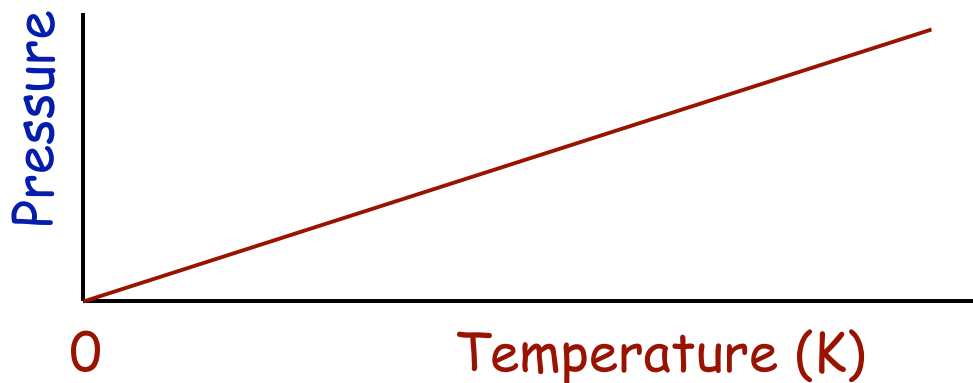
- measure pressure of gas at fixed volume



$$\underline{\text{Pressure}} = \text{Force/Area} \quad (\text{N/m}^2 = \text{Pa}) \\ (\text{Pascals})$$

$$1 \text{ atm} = 1.01 \times 10^5 \text{ Pa} = 14.7 \text{ lb/in}^2 \\ = 760 \text{ mm of Hg} = 760 \text{ torr}$$

T P at fixed V



## Ideal-Gas Temperature

$$T_k = (\text{constant}) \times P \quad \text{at fixed } V$$

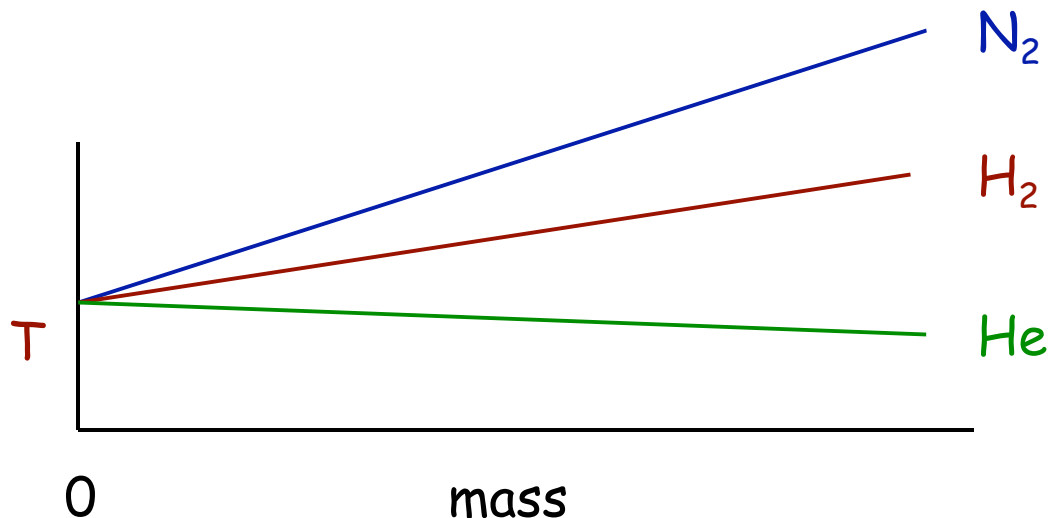
-Need one point:

Triple point of  $H_2O$   
(ice/water/steam coexist)

$$T_3 = 273.16 \text{ K}$$

-Problem: different gases give  
different  $T$

But as mass of gas reduced ( $m \rightarrow 0$ )  
and  $P_3 \rightarrow 0$ , they agree  
(approach "ideal" gas)

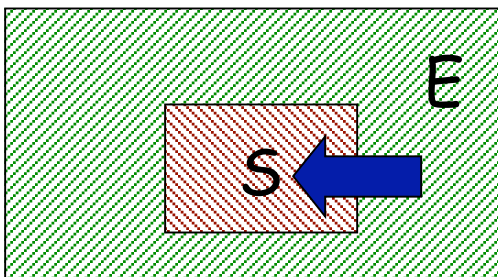


# Temperature and Heat

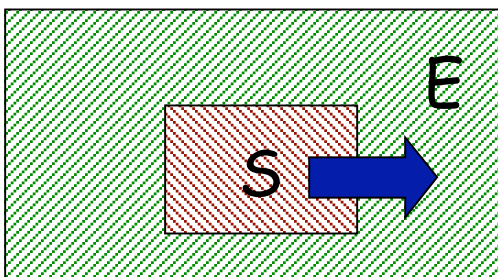
If **system S** and **environment E** are  
At different temperatures:

Energy will transfer until their  
temperatures become equal.

The transferred energy is called  
Heat (symbol  $Q$ ).



$T_E > T_S$  ,  $Q > 0$   
Heat absorbed by **S**



$T_S > T_E$  ,  $Q < 0$   
Heat lost by **S**



Defn: Require  $\Delta Q = \underline{1 \text{ calorie}}$   
to raise 1 gm of  $\text{H}_2\text{O}$  by  $\Delta T = 1^\circ\text{C}$ .

1 calorie = 4.186 joules  
(heat is a form of energy)

## Specific Heat

Amount of heat needed to raise the temperature of  $m$  grams of a substance by  $\Delta T$  is

$$\Delta Q = c m \Delta T$$

$$\Delta = \text{final} - \text{initial}$$

where  $c$  is the specific heat ( $\text{cals/g}\cdot^\circ\text{C}$ )

Specific heat of water =  $1 \text{ cal/g}\cdot^\circ\text{C}$   
=  $4186 \text{ J/kg}\cdot\text{K}$

# Molar Specific Heat

Can specify amount of substance in moles:

$$\begin{aligned} 1 \text{ mole} &= 6.02 \times 10^{23} \text{ units} \\ &= N_A \text{ units (Avogadro's number)} \end{aligned}$$

( 1 mole of Al =  $6.02 \times 10^{23}$  atoms  
1 mole of  $\text{CO}_2$  =  $6.02 \times 10^{23}$  molecules)

The mass of a substance (in grams) is

$$m = n A$$

where  $n$  = (# moles) and  
 $A$  is the atomic (molecular) weight of the substance.

1 mole of Carbon-12 has  $m = 12$  grams.

# Some Examples:

$c$  (kg)

$C$  (molar)

<u>Element</u>	<u>Spec. Heat</u> (J/kg·K)	<u>A</u> (g)	<u>Mol. Sp. Ht.</u> (J/Mole·K)
Lead	128	207	26.5
Tungsten	134	184	24.8
Silver	236	108	25.5
Copper	386	63.5	24.5
Aluminum	900	27	24.4

  
Note the relative consistency

$$\Delta Q = m c \Delta T$$

$$\Delta Q = n C \Delta T$$

$$C = c (m/n) = c A$$

$$A(\text{Pb}) = .207 \text{ kg} = 207\text{g}$$

# Heats of Transformation

Heat may also change the phase (or state) of a substance (at constant T).

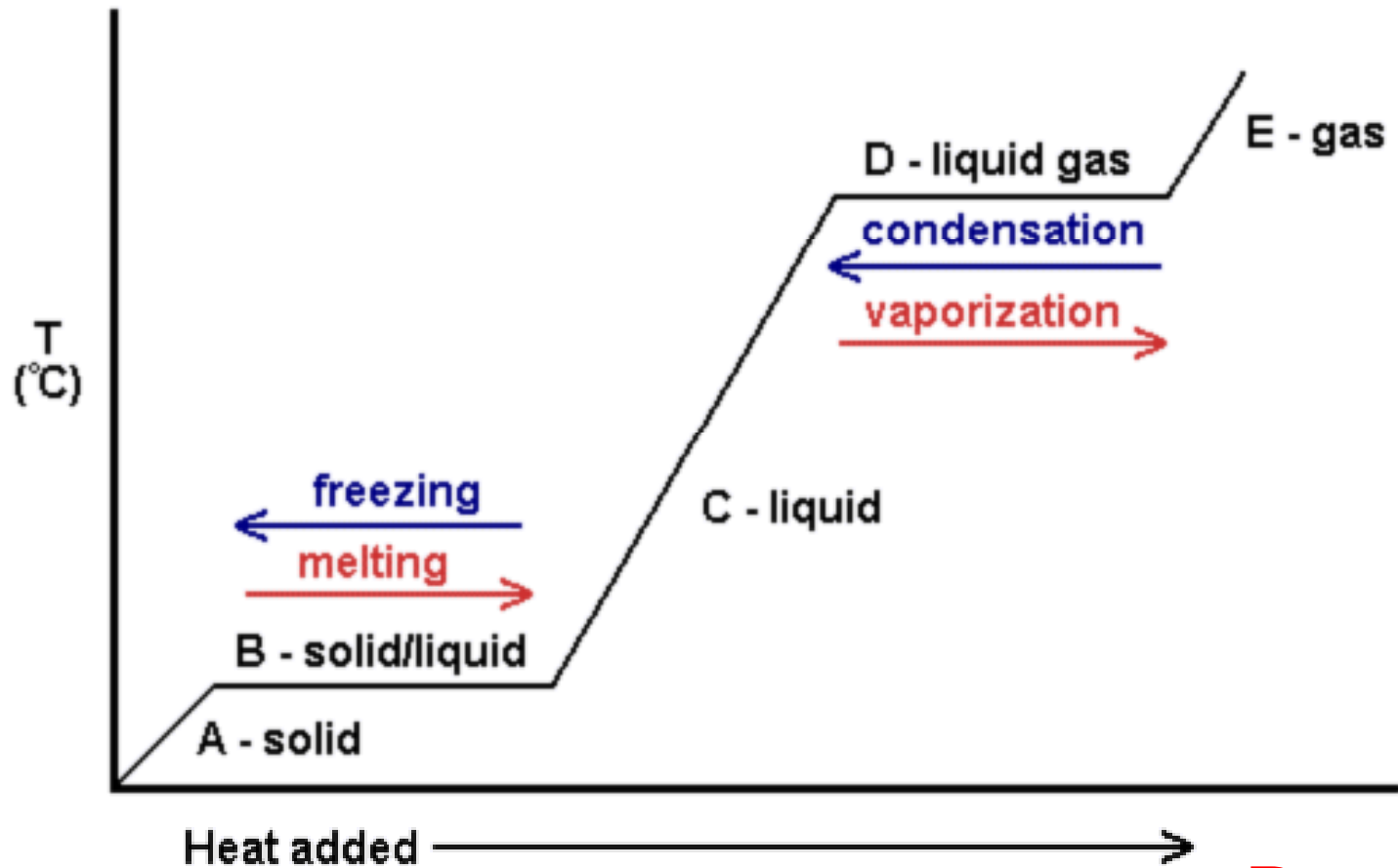
Matter exists in 3 common states:

- Solid

- Liquid

- Gas (or vapor)

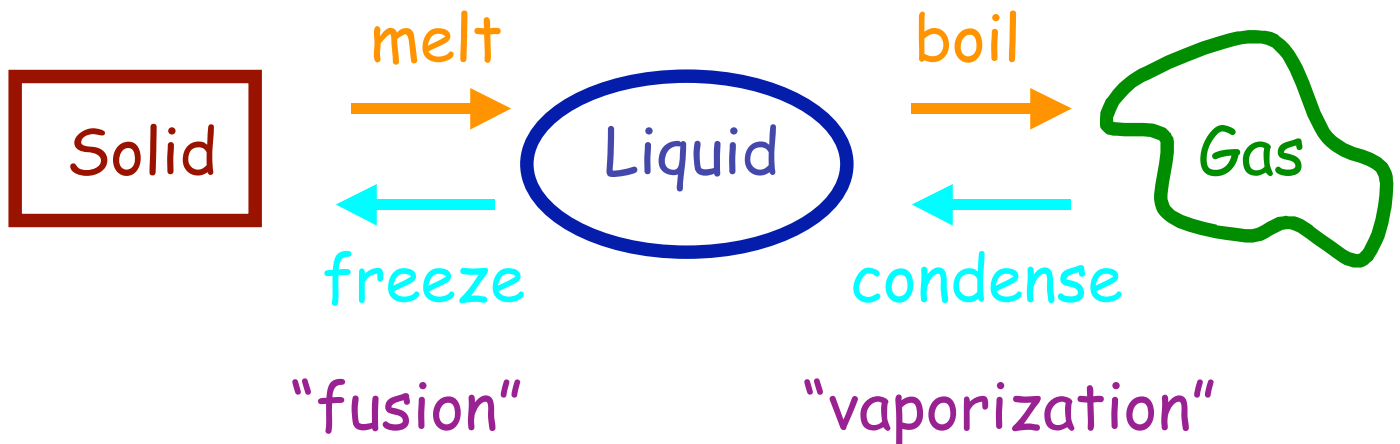
# Heating Curve





Demo

Image:

<http://library.thinkquest.org>



Requires energy   
 Releases energy 

Amount of energy/unit mass is  
Heat of transformation,  $L$ .

e.g. for water:

Heat of fusion

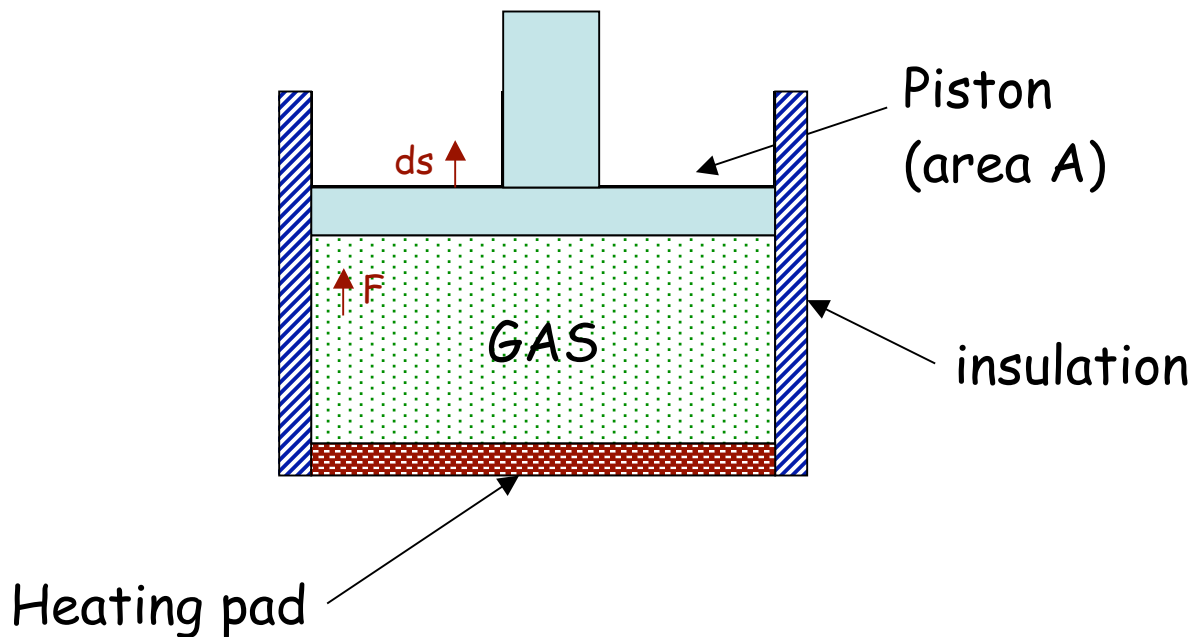
$$L_F = 79.5 \text{ cal/g} = 333 \text{ kJ/kg} = 6.01 \text{ kJ/mole}$$

Heat of vaporization

$$L_V = 539 \text{ cal/g} = 2256 \text{ kJ/kg} = 40.7 \text{ kJ/mole}$$

# Heat and Work

Consider this system:



Pressure = Force/Area  $(P=F/A)$

If piston moves  $ds$ , then work done by the gas:

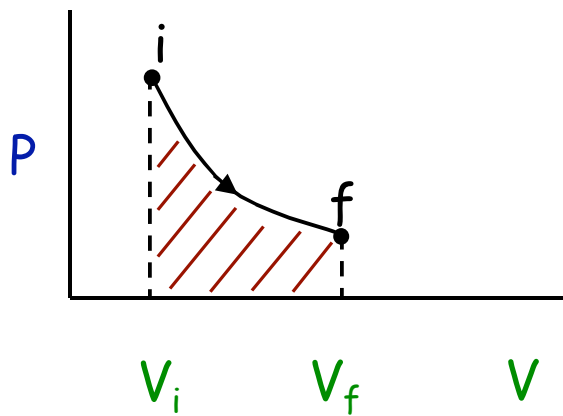
$$dW = F ds = P A ds = P dV$$

Total work done by the gas in moving from  $V_i$  to  $V_f$ :

$$\square W = \int_{V_i}^{V_f} P dV$$

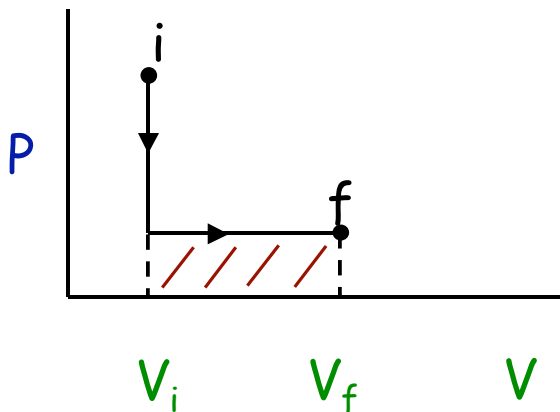
# P-V Diagrams

Study effects of heat added/work done by plotting  $P$  vs  $V$  of gas:

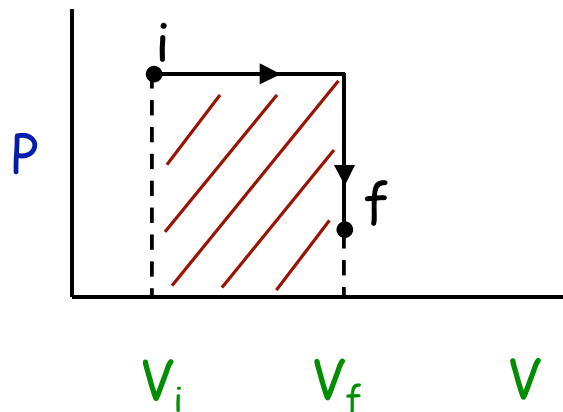


The area under the curve is the work done.

The work done depends on the specific path from  $i$  to  $f$ .



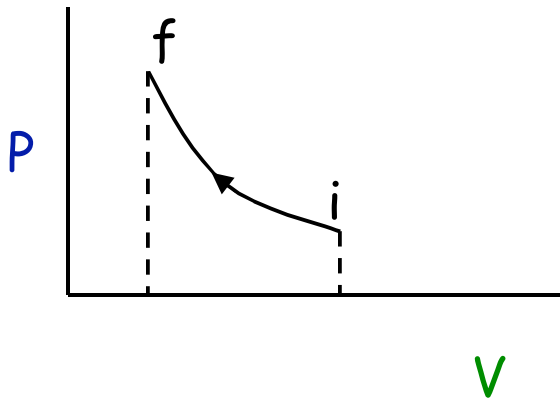
**A**



**B**

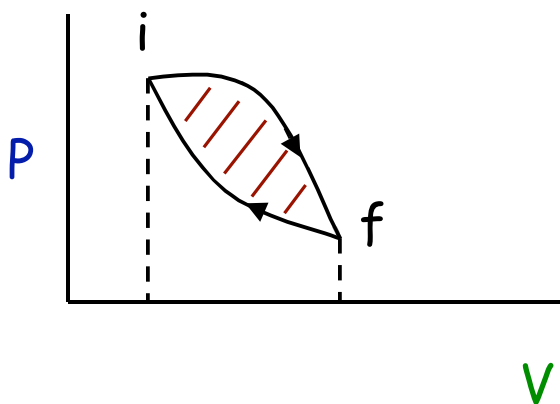


# Thermodynamic Cycles

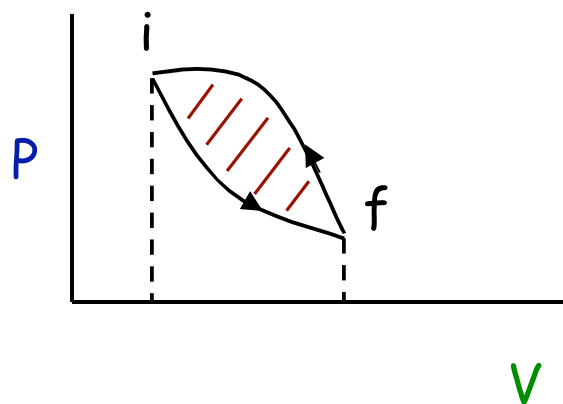


If volume decreases, the work done (by the gas) is negative.

If we go from  $i \rightarrow f$  and back to  $i$ , the net work done by the gas is the area inside the curve.



+ work



- work

# First Law of Thermodynamics

Heat  $\Delta Q$  added to the system can have two effects:

- Increase the internal energy of the system
- Cause the gas to do work

Conservation of Energy says:

$$\Delta Q = \Delta U + \Delta W$$

where

$U$  is the internal energy of the system.

□ 1<sup>st</sup> Law of Thermodynamics.

# Energy Conservation: A Parable from Feynman

**Count** sugar cubes directly

Deduce number hidden in a box

Even dissolved in a pond

Count those that passed through a window

(named  $W$  or  $Q$ )

That's defining the system boundary!

Locality is assumed: cubes don't just re-materialize  
outside the windows: must pass through the windows

Learned to calculate various forms of internal energy.

Can't *derive* conservation of  $E$ : must deduce that it's plausible,  
learn to account for different forms of  $E$ , and then measure  
that it works, with your chosen definitions and calculation  
methods.

# State Functions

A property of the state of the system is often called a "State Function".

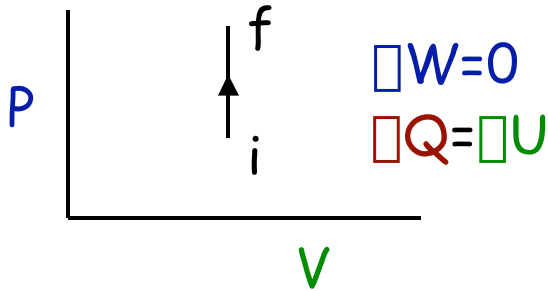
$P$ ,  $V$ , and  $T$  are state functions.  
So is  $U$  (the internal energy).

Heat and Work are not.  
They are path-dependent, i.e. they depend on how we go from  $i$  to  $f$ .

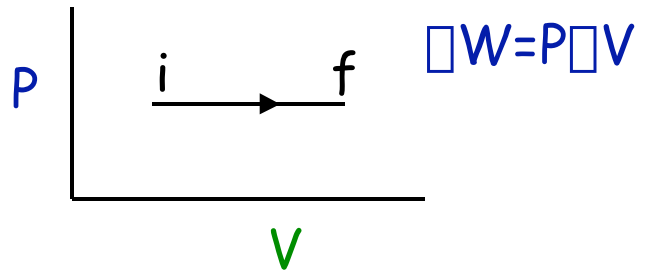
However the combination  
 $\Delta Q - \Delta W = \Delta U$  does not depend on the path.

# Various System Changes

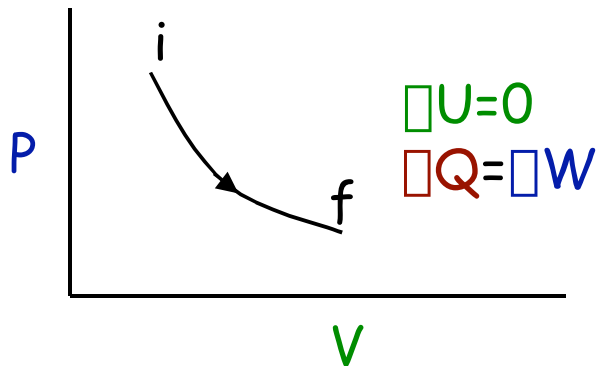
Constant Volume  
(isochoric)



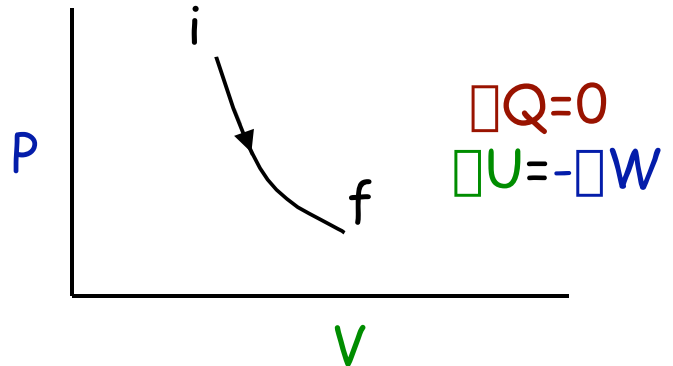
Constant Pressure  
(isobaric)



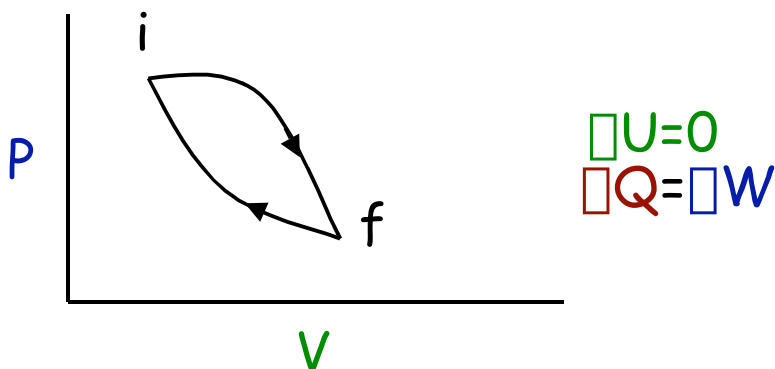
Constant Temp  
(isothermal)



Constant Heat  
(adiabatic)



Cyclical Process  
(returns to original state)



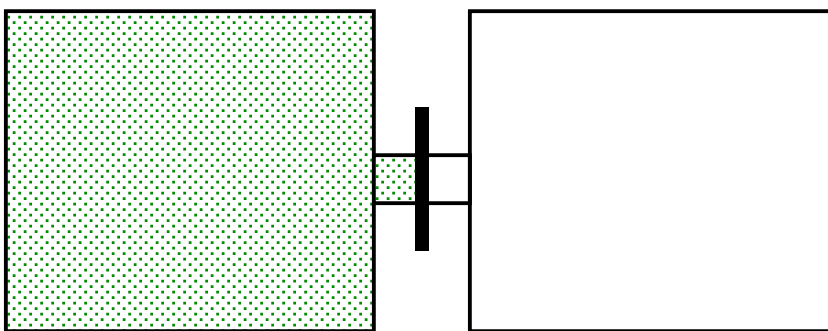
All previous cases are "quasi-static":

Change occurs slow enough that thermal equilibrium can be considered true at all times.

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A non-quasi-static process:

Adiabatic, free expansion:



$$\square Q = 0 \quad (\text{adiabatic})$$

$$\square W = 0 \quad (\text{nothing to work against})$$

$$\square \square U = 0$$

# Heat Transfer Mechanisms

How does heat exchange occur?

- Conduction

- Convection

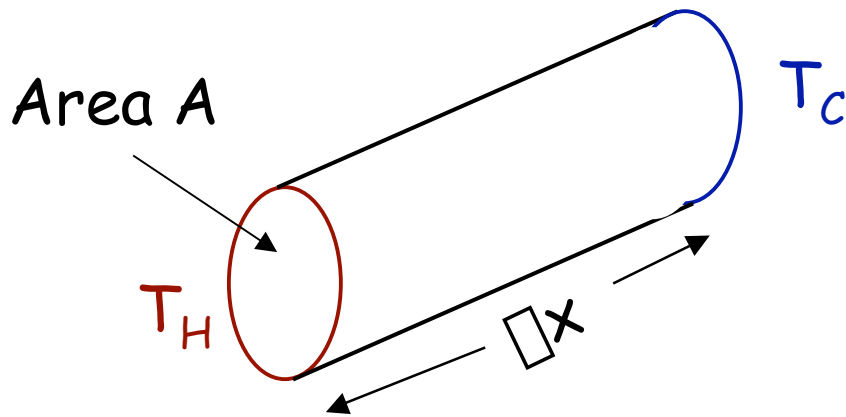
- Radiation

# Conduction

- Occurs in systems where atoms stay in a fixed region.
- Heat energy causes them to move, rotate, and/or vibrate.
- Energy is transferred to adjacent atoms by interactions/collisions.

Energy moves, not the atoms





Heat conduction rate is

$$P_{\text{cond}} = \kappa Q / \Delta t = \kappa A \Delta T / \Delta x$$

- $\Delta T / \Delta x = (T_H - T_C) / \Delta x$   
is the Temperature gradient.
- $P_{\text{cond}}$  is the Energy transferred per time (SI units: Watts), sometimes called thermal current,  $I$ .
- $\kappa$  is the Thermal Conductivity (SI units: Watts/m·K).

## Some Thermal Conductivities

Silver	428	W/m·K
Copper	402	
Aluminum	235	
Lead	35	
Stainless Steel	14	
Hydrogen	0.18	
Helium	0.15	
Dry Air	0.026	
Window Glass	1.0	
White Pine	0.11	
Fiberglass	0.048	
Polyurethane Foam	0.024	

Using the notation  $I$  for  $P_{\text{cond}}$ , we can write

$$\Delta T = I \Delta x / (\kappa A)$$

or

$$\Delta T = I R$$

where  $R = \Delta x / (\kappa A)$  is the thermal resistance.

Note the analogy with Ohm's Law for electricity.

$$\text{Note: } R_{\text{fishbane}} = \Delta x / \kappa = R A$$

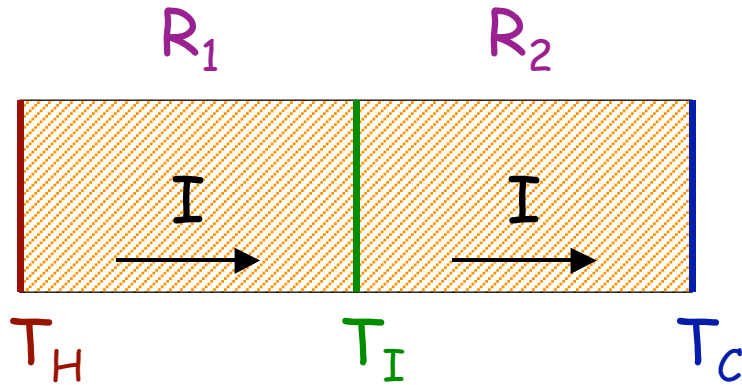
$R_{\text{fishbane}}$  = insulation "R-value"

= material and thickness only

can't change size of attic

can add more, or better, insulation

Two or more conductors (or insulators) in series:



Steady state  $\square$  thermal current is same through both slabs

$$T_H - T_I = I R_1$$

$$T_I - T_C = I R_2$$

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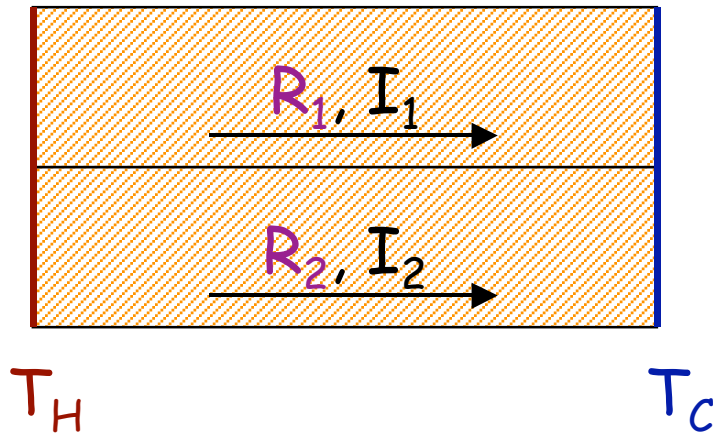
$$T_H - T_C = I (R_1 + R_2) = I R_{\text{equiv}}$$

where

$$R_{\text{equiv}} = R_1 + R_2 + \dots$$

(like resistances in series)

Conductors in parallel:  
(multiple paths for heat flow)



$T_H - T_C = \Delta T$ : same for all paths  
but current flows ( $I$ ) are different.

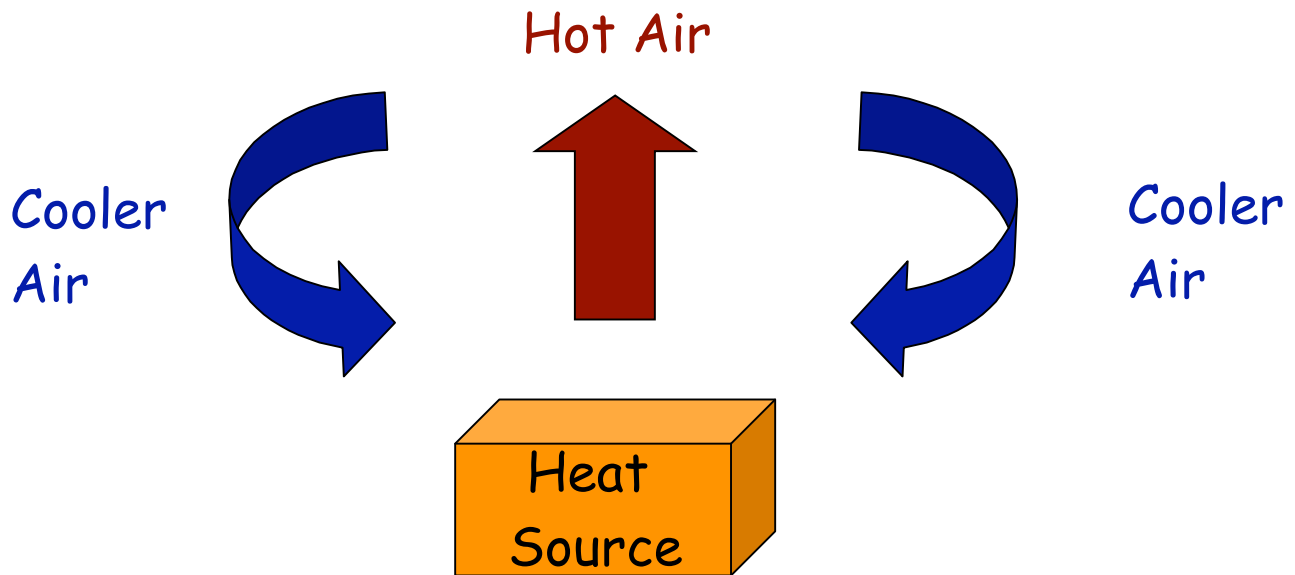
$$\begin{aligned} I_{\text{total}} &= I_1 + I_2 + \dots \\ &= \Delta T / R_1 + \Delta T / R_2 + \dots \\ &= \Delta T (1/R_1 + 1/R_2 + \dots) \\ &= \Delta T / R_{\text{equiv}} \end{aligned}$$

with

$$1/R_{\text{equiv}} = 1/R_1 + 1/R_2 + \dots$$

(like resistances in parallel)

# Convection



- Occurs in fluid systems.  
The energy flows along with the medium.
- Fluid near heat source becomes hot, expands, and rises. Surrounding cooler fluid takes its place. Etc.

# Radiation

Here the energy is carried by electromagnetic waves. Called Thermal Radiation.

The Rate at which an object radiates is given by the **Stefan-Boltzman Law**:

$$P_{\text{rad}} = \varepsilon \sigma A T^4$$

where

- $P_{\text{rad}}$  : Power radiated in Watts
- $A$  : Area of emitter (or absorber!)
- $T$  : Temperature of emitter in K
- $\sigma$  : Universal constant  
(S-B's constant)  
 $\sigma = 5.6703 \times 10^{-8} \text{ W}/(\text{m}^2\text{K}^4)$
- $\varepsilon$  : the emissivity of the emitter  
( $0 < \varepsilon < 1$ , depending on the composition of the surface)

The rate an object absorbs thermal radiation is given by the same formula:

$$P_{\text{abs}} = \epsilon A (T_{\text{env}})^4$$

except that now  $T_{\text{env}}$  is the temperature of the environment.

The **emissivity**  $\epsilon$  of an object is the same for radiation and absorption.

- Lighter objects reflect more.  
(smaller  $\epsilon$ )
- Darker objects absorb more.  
(larger  $\epsilon$ )  
They also emit more.

A surface with  $\epsilon = 1$  is called a Blackbody radiator.