

# Essential Physics II

## 英語で物理学の エッセンス II

Lecture 3: 15-10-15

# Homework

If you did not do the homework... **WHY?**



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#	TITLE	CATEGORY	DUE DATE/TIME	AVAILABILITY TO STUDENTS	Edit
1	<input type="checkbox"/> <a href="#">Introduction to MasteringPhysics</a>	Homework	10/20/14 at 04:30pm	From: 09/29/14 at 06:00pm Until: 02/14/15 at 11:59pm	
2	<input type="checkbox"/> <a href="#">Week 2</a>	Homework	10/20/14 at 04:30pm	From: 10/06/14 at 06:00pm Until: 02/14/15 at 11:59pm	

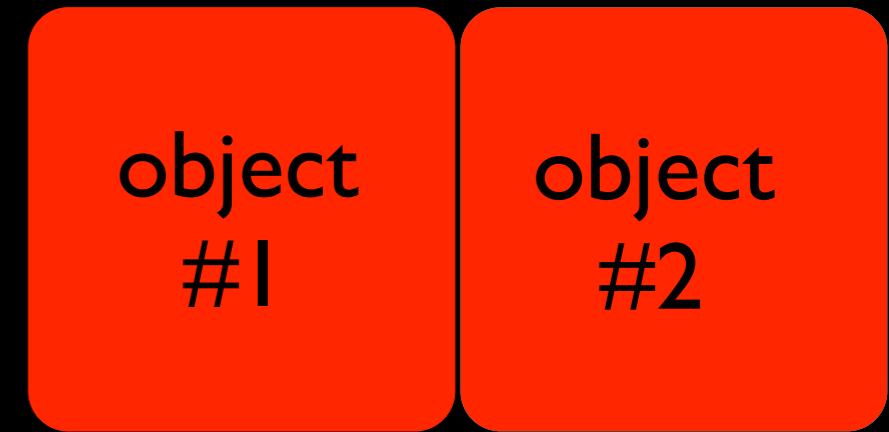
<http://masteringphysics.com>

<http://astro3.sci.hokudai.ac.jp/~tasker/teaching/ep2>

# Last Lecture

Temperature:

2 objects have the same  $T$  when in **thermodynamic equilibrium**

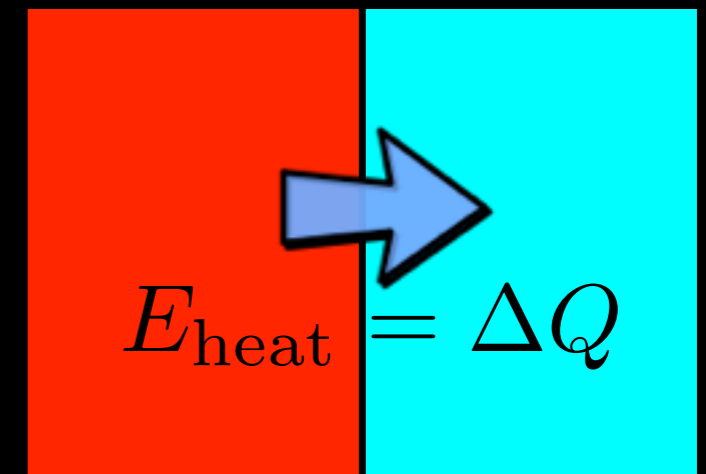


SI unit: kelvin [K]

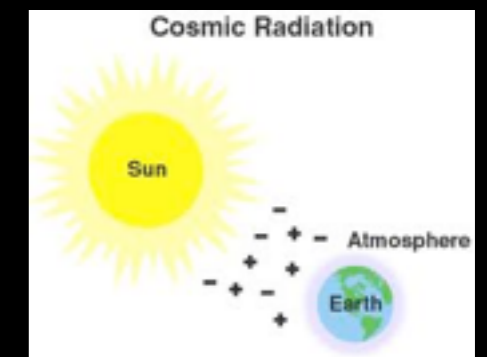
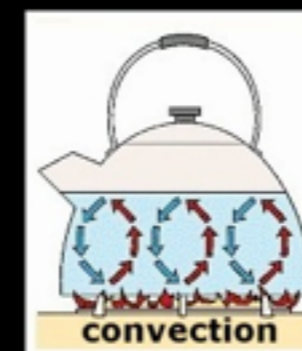
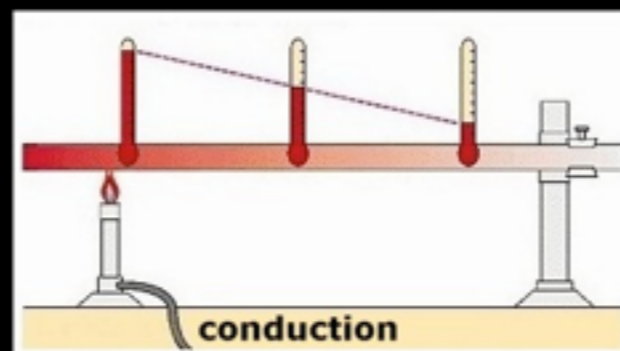
Heat is **energy** moving because of a difference in temperature ( $\Delta T$ )

$$Q = mc\Delta T$$

**specific heat**



Heat can be transferred via **conduction**, **convection** and **radiation**



# Last Lecture

The specific heat ( $c$ ) of water is greater than iron.

If you drop hot iron into a water bucket, the heat gained by the water is...

- (a) **larger** than the heat lost by iron
- (b) **smaller** than the heat lost by iron
- (c) the **same** as the heat lost by iron



$T$  is different, but energy is conserved

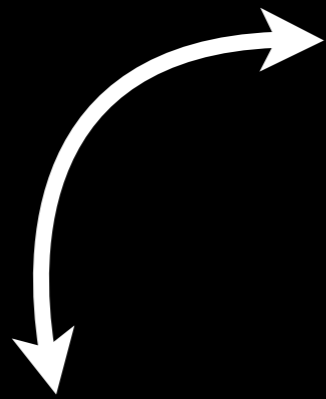
# Heat and Matter



# Ideal Gas Law

How are...

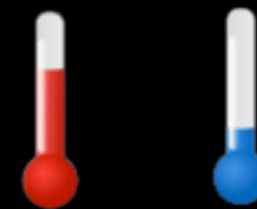
Pressure ( $p$ )



Volume ( $V$ )



Temperature ( $T$ )



... related?

# Ideal Gas Law

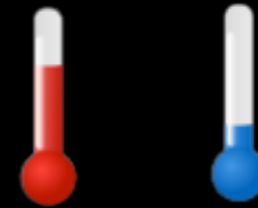
Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )

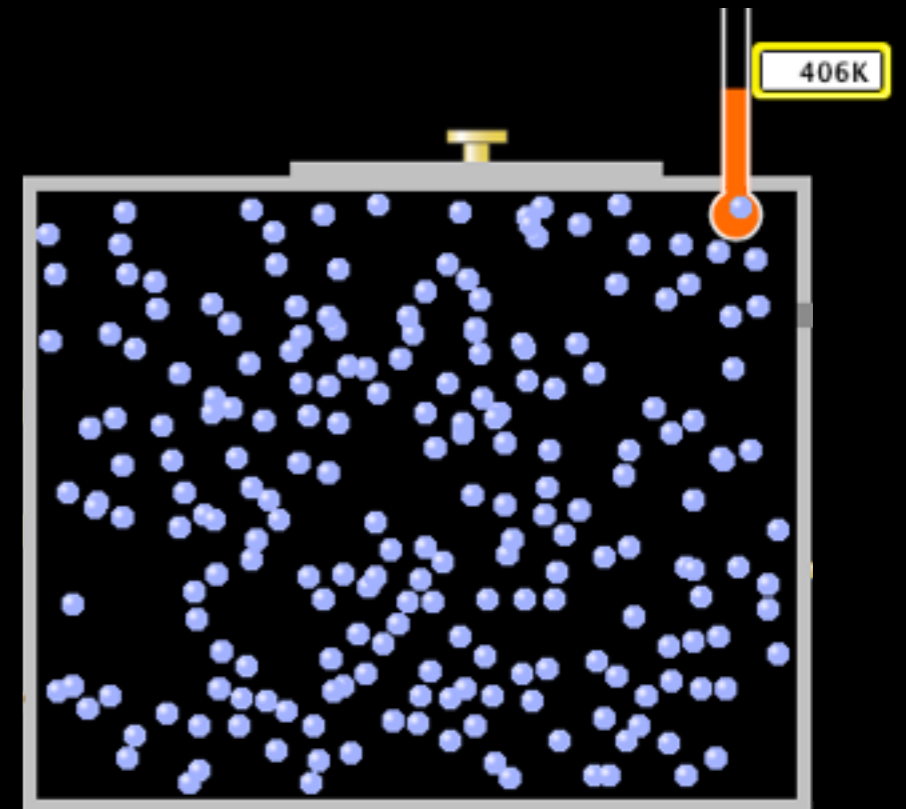


If  $V$  is constant, what happens to  $p$  if we lower  $T$ ?

(a) Pressure goes down ↓

(b) Pressure goes up ↑

(c) Pressure stays the same



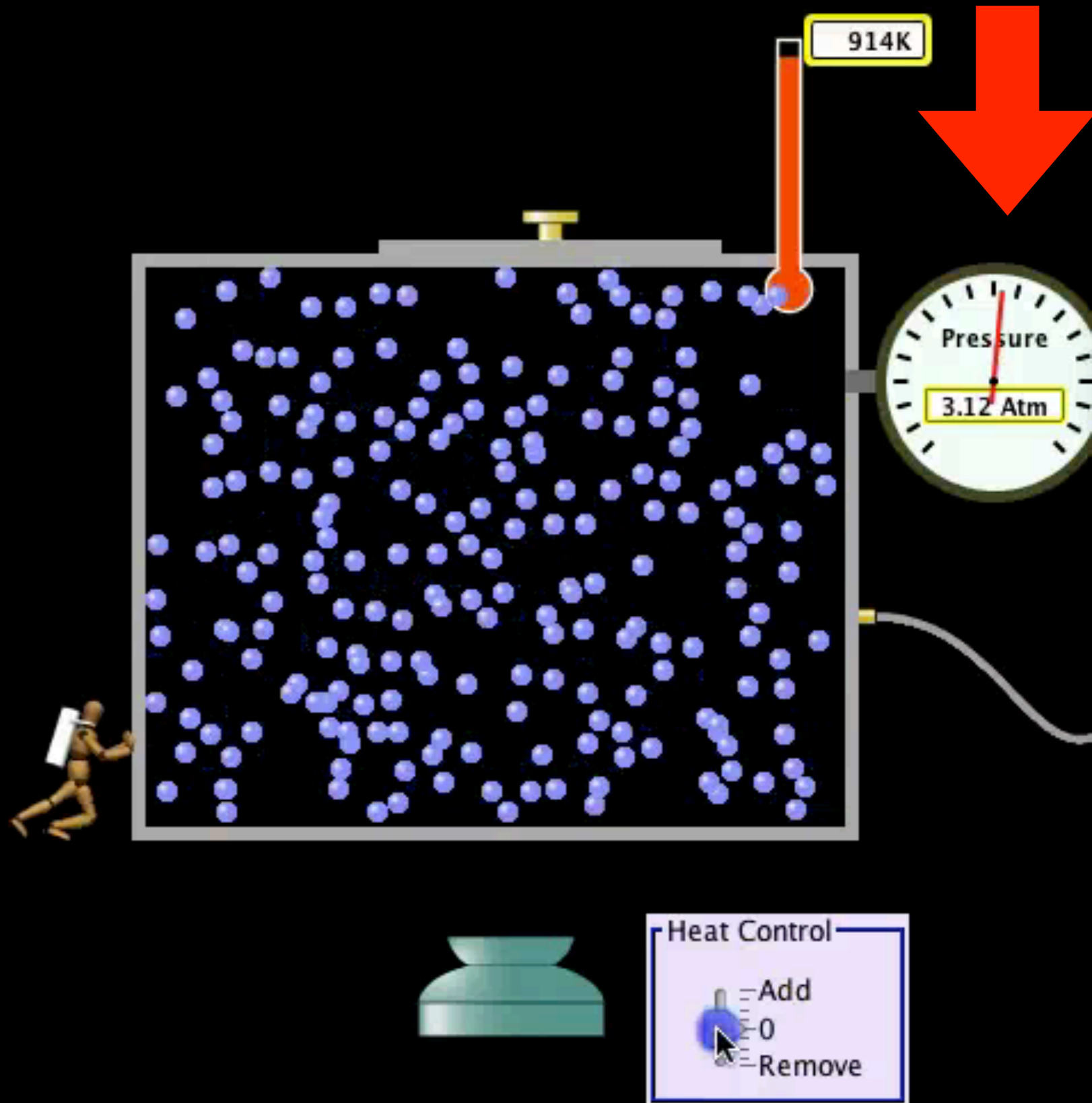
# Ideal Gas Law

V is constant, 

T goes down 

p ... .. goes down 

$$p \propto T$$





# Ideal Gas Law

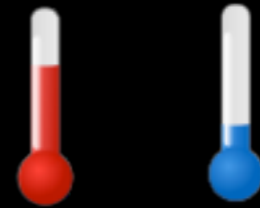
Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )

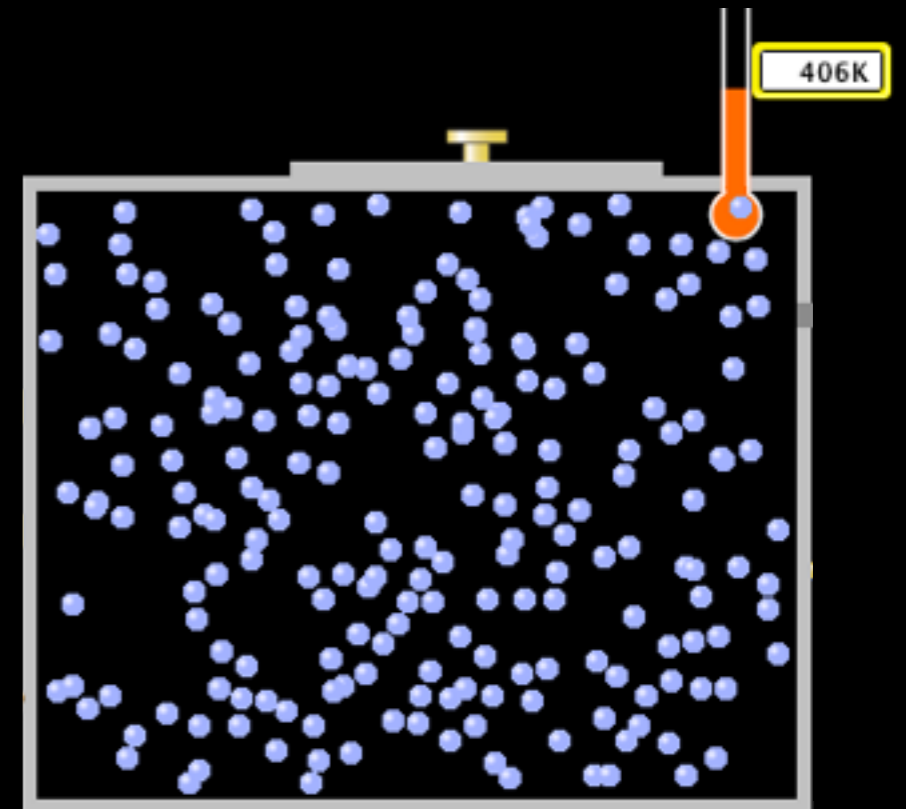


If  $V$  is constant, what happens to  $p$  if we lower  $T$ ?

(a) Pressure goes down ↓  $p \propto T$

(b) Pressure goes up ↑

(c) Pressure stays the same



# Ideal Gas Law

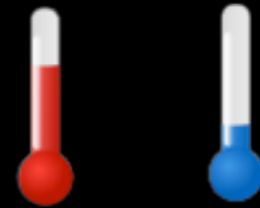
Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )

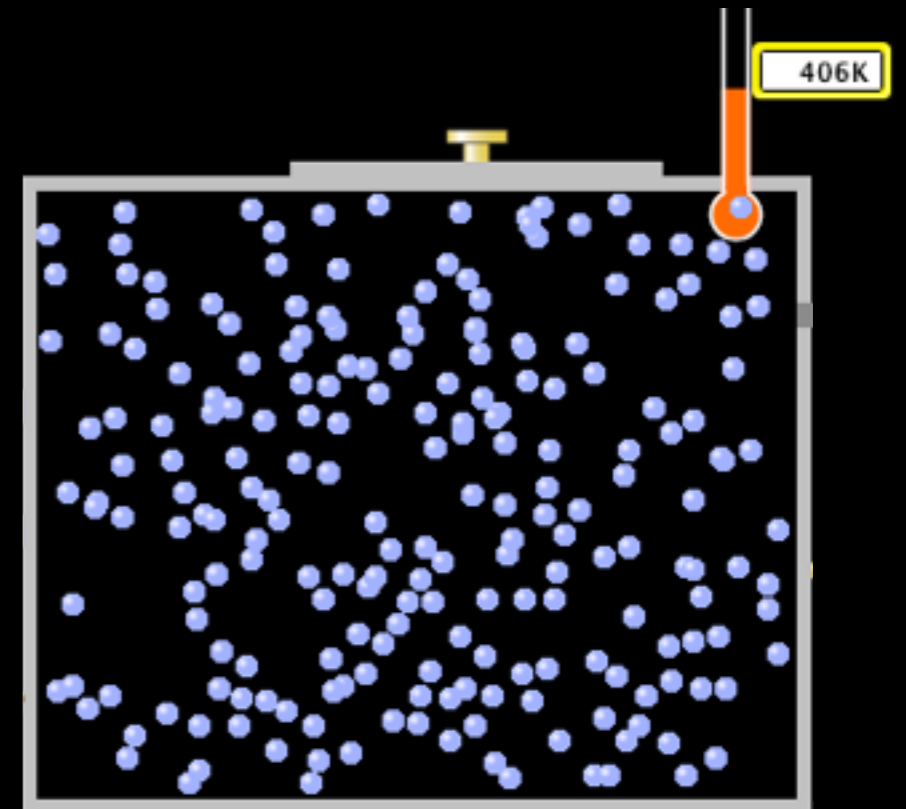


If  $T$  is constant, what happens to  $p$  if we lower  $V$ ?

(a) Pressure goes down ↓

(b) Pressure goes up ↑

(c) Pressure stays the same



# Ideal Gas Law

T is constant,



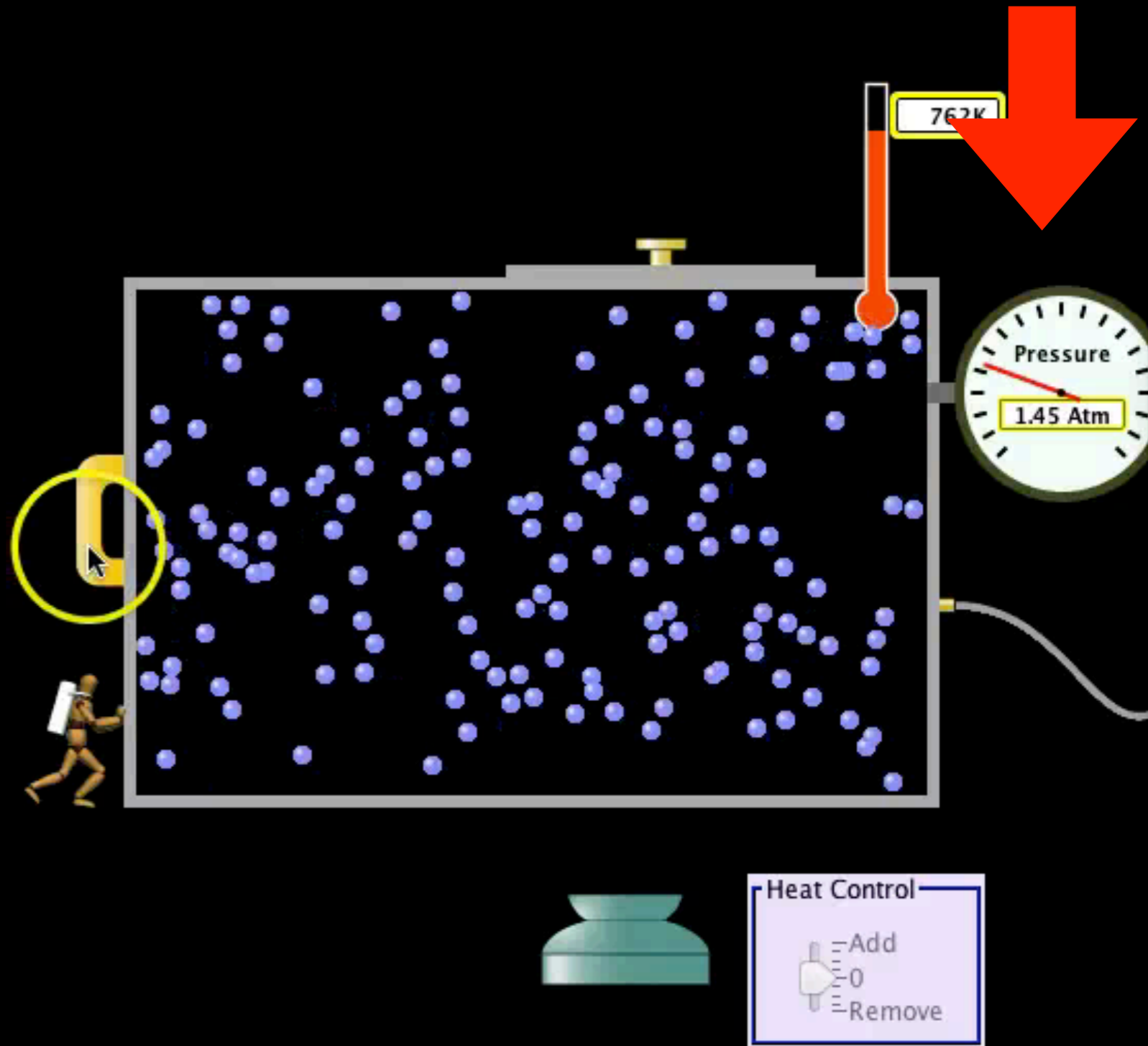
Lower V



p ... .. goes up



$$p \propto \frac{1}{V}$$



# Ideal Gas Law

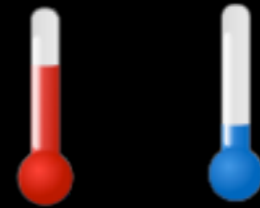
Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )



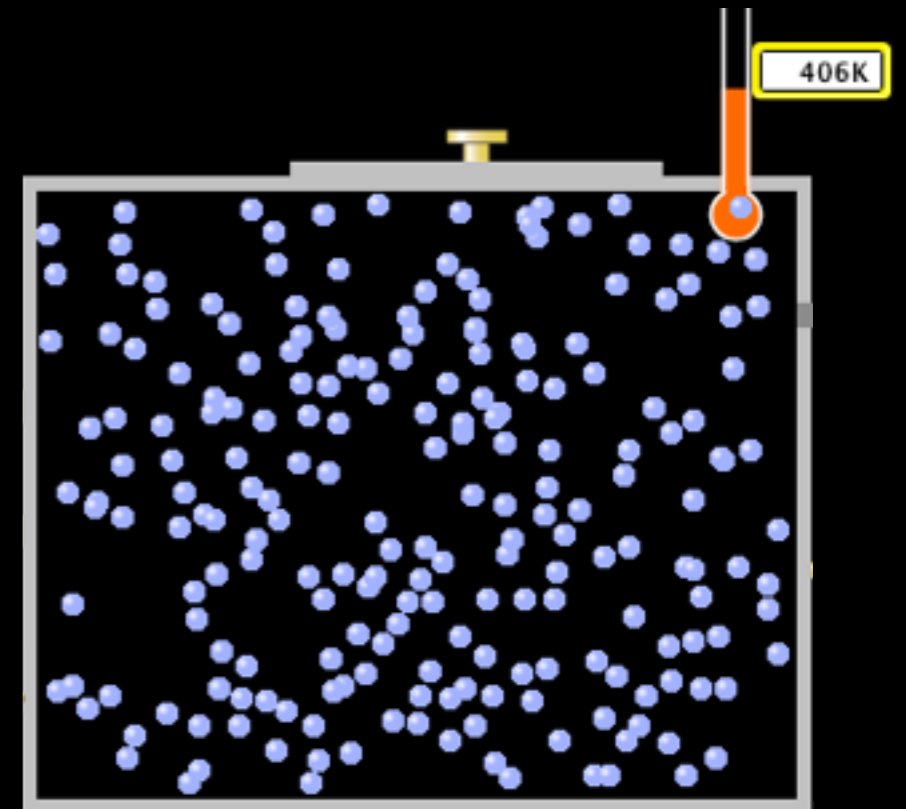
If  $T$  is constant, what happens to  $p$  if we lower  $V$ ?

(a) Pressure goes down ↓

(b) Pressure goes up ↑

(c) Pressure stays the same

$$p \propto \frac{1}{V}$$



# Ideal Gas Law

Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )

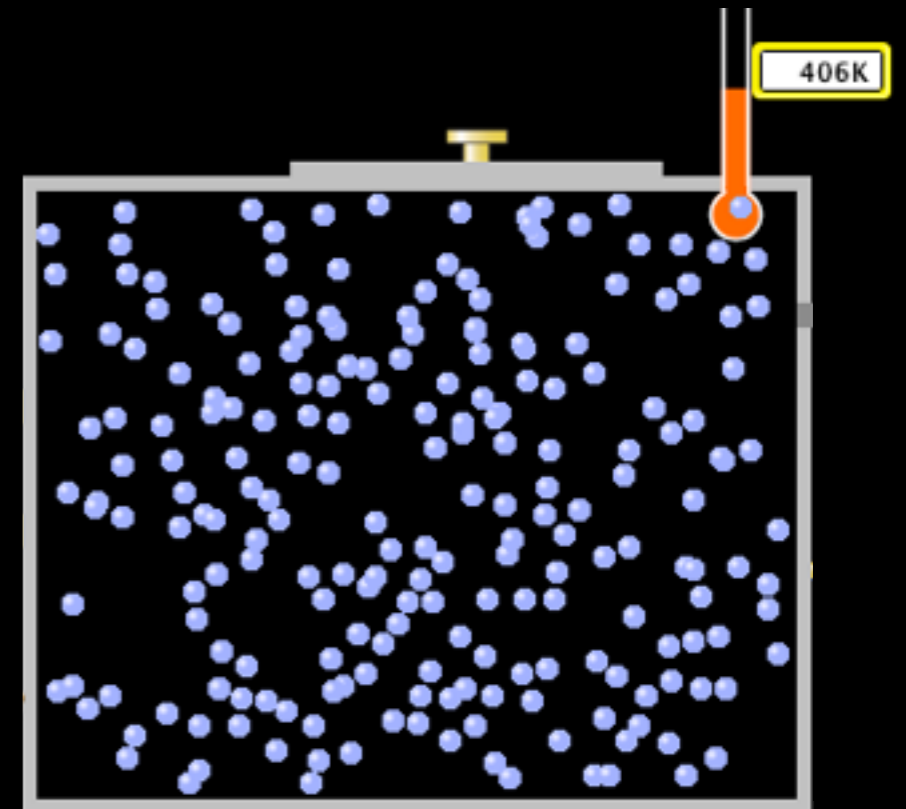


If  $p$  is constant, what happens to  $V$  if we lower  $T$ ?

(a) Volume goes down ↓

(b) Volume goes up ↑

(c) Volume stays the same



# Ideal Gas Law

$p$  is constant,



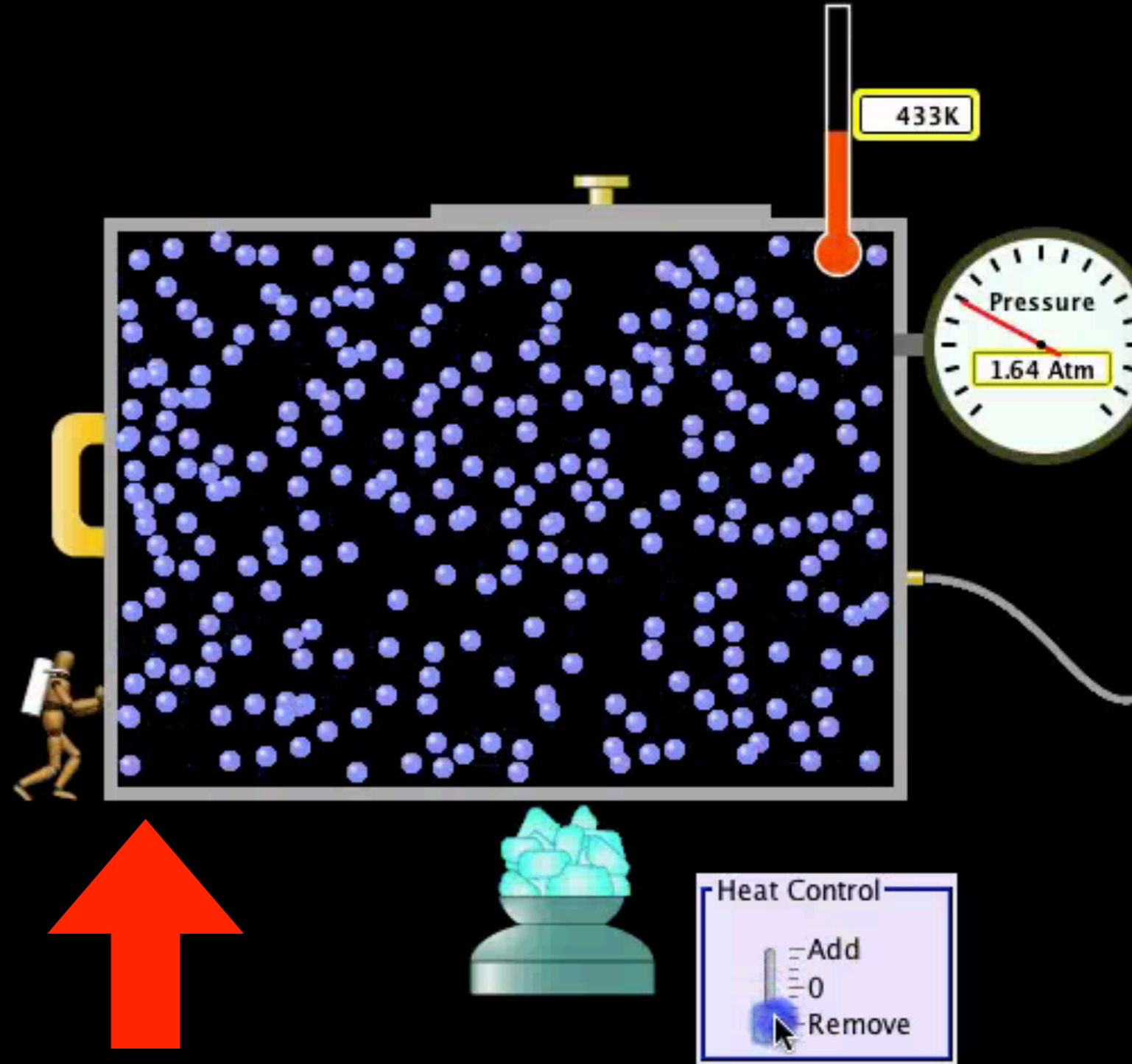
$T$  goes down ↓



$V$  ... .. goes down ↓



$$T \propto V$$



# Ideal Gas Law

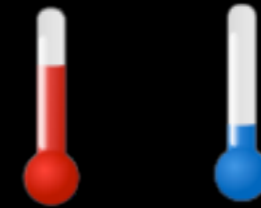
Volume ( $V$ )



Pressure ( $p$ )



Temperature ( $T$ )

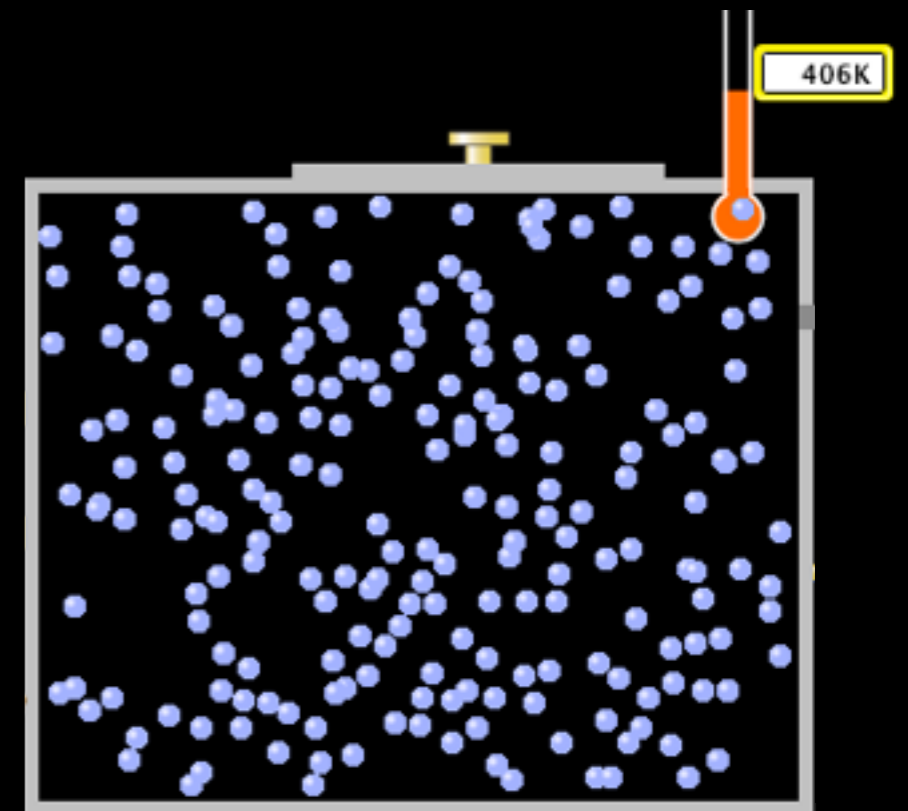


If  $p$  is constant, what happens to  $V$  if we lower  $T$ ?

(a) Volume goes down ↓  $T \propto V$

(b) Volume goes up ↑

(c) Volume stays the same



# Ideal Gas Law

$$p \propto T \quad p \propto \frac{1}{V} \quad T \propto V$$

Therefore:  $pV \propto T$       What else affects  $p, V$  and  $T$ ?

*Bonus question:* ★

If  $V$  and  $T$  are constant, what happens to  $p$  if we lower  $N$ ?

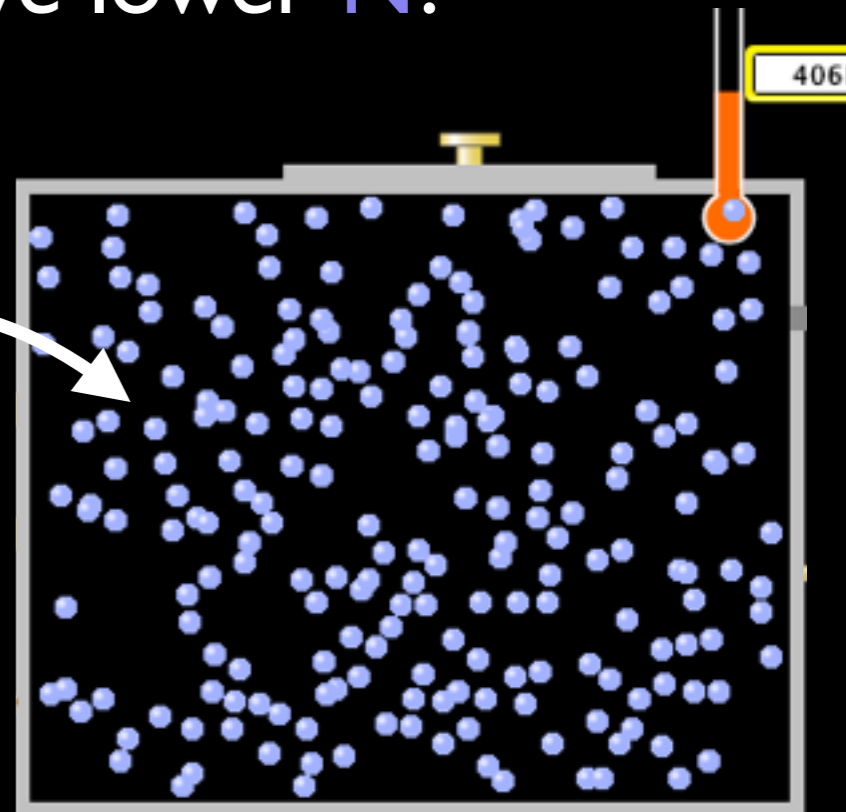
(a) Pressure goes down

$$p \propto N$$

(b) Pressure goes up

(c) Pressure stays the same

$N$  gas molecules





# Ideal Gas Law

$$p \propto T$$

$$p \propto \frac{1}{V}$$

$$T \propto V$$

$$p \propto N$$

Therefore:  $pV = NkT$  ideal-gas law

$N = \#$  of gas molecules



$$k = 1.38 \times 10^{-23} \text{ J/K}$$

Boltzmann's constant

very large! ↓

count # of moles: 1 mole,  $N_A = 6.023 \times 10^{23}$  atoms  
Avogadro's number

→  $pV = nN_AkT = nRT$

# of moles

$$R = N_Ak = 8.314 \text{ J/K} \cdot \text{mol}$$

universal gas constant

# Ideal Gas Law

Ex.

What is the volume,  $V$ , of 1.00 mol of an ideal gas at  $T = 0^\circ\text{C}$  and  $p = 101.3\text{ kPa}$ ?

$$R = 8.314\text{ J/K} \cdot \text{mol}$$

Ideal gas law:  $pV = nRT$

$$\begin{aligned} V &= \frac{nRT}{p} = \frac{(1.00\text{ mol})(8.314\text{ J/K} \cdot \text{mol})(273\text{ K})}{1.01 \times 10^5\text{ Pa}} \\ &= 22.4 \times 10^{-3}\text{ m}^3 \\ &= 22.4\text{ L} \end{aligned}$$

# Ideal Gas Law

# Quiz

An ideal gas has  $T = 109^\circ\text{C}$  and  $p = 1.2 \times 10^4 \text{ Pa}$ .

How many gas molecules are in a cubic centimeter ( $1 \text{ cm}^3$ )?

$$R = 8.314 \text{ J/K} \cdot \text{mol} \quad N_A = 6.023 \times 10^{23}$$

$$pV = nRT \rightarrow n = \frac{pV}{RT}$$

(a)  $7.98 \times 10^{18}$  molecules

$$= \frac{(1.2 \times 10^4 \text{ Pa})(0.01^3 \text{ m}^3)}{(8.314 \text{ J/K} \cdot \text{mol})(109 + 273 \text{ K})}$$

(b)  $3.8 \times 10^{-6}$  molecules

$$= 3.8 \times 10^{-6} \text{ moles}$$

(c)  $2.3 \times 10^{18}$  molecules

$$N = 3.8 \times 10^{-6} \times 6.023 \times 10^{23}$$

(d)  $1.3 \times 10^5$  molecules

$$= 2.27 \times 10^{18} \text{ molecules}$$

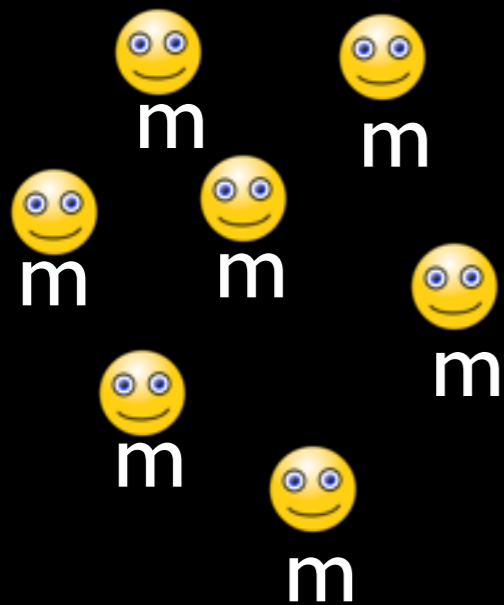
# Kinetic Theory of Gases

$pV = nRT$  is a very simple relation! ... why does it work?



Assume:

(I) Gas is made of identical molecules



→ Mass,  $m$

→ Size  $\sim 0$



nothing inside  
(no internal structure)

~ True, if distance between gas molecules  $\gg$  molecule size



# Kinetic Theory of Gases

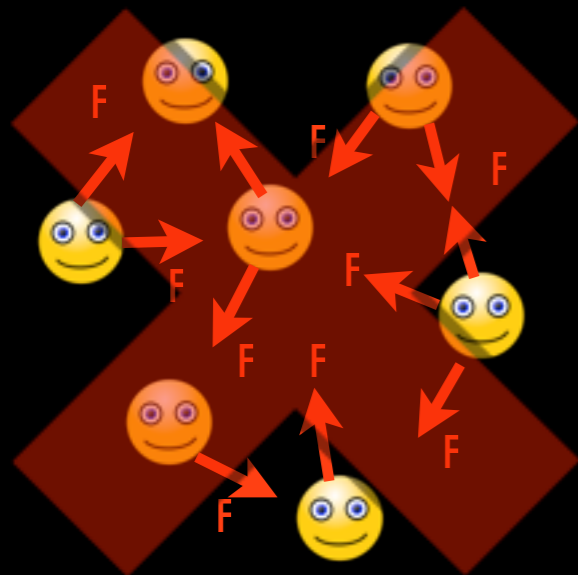
$pV = nRT$  is a very simple relation!

... why does it work?



Assume:

(II) No forces between molecules



only kinetic energy (K)

A yellow emoji-like molecule with a green arrow pointing upwards and to the right, labeled with the symbol  $\bar{v}$ , representing its velocity.
$$K = \frac{1}{2}mv^2$$

no potential energy (U)



# Kinetic Theory of Gases

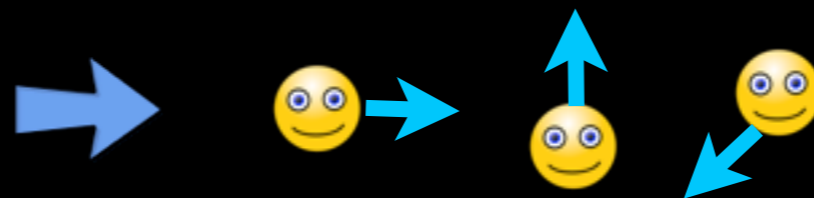
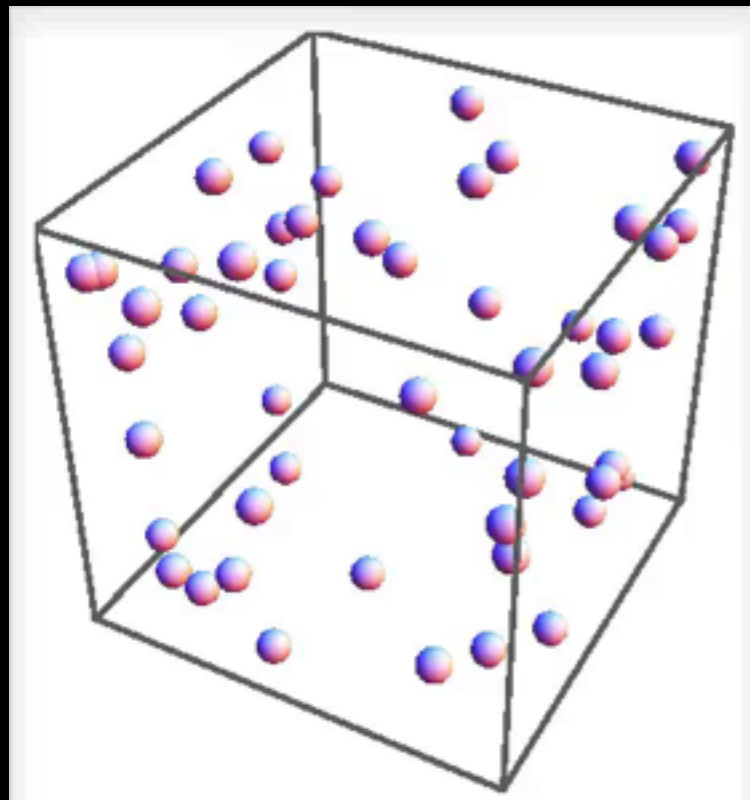
$pV = nRT$  is a very simple relation!

... why does it work?



Assume:

(III) molecules moves in random directions



speed  $\propto$  direction



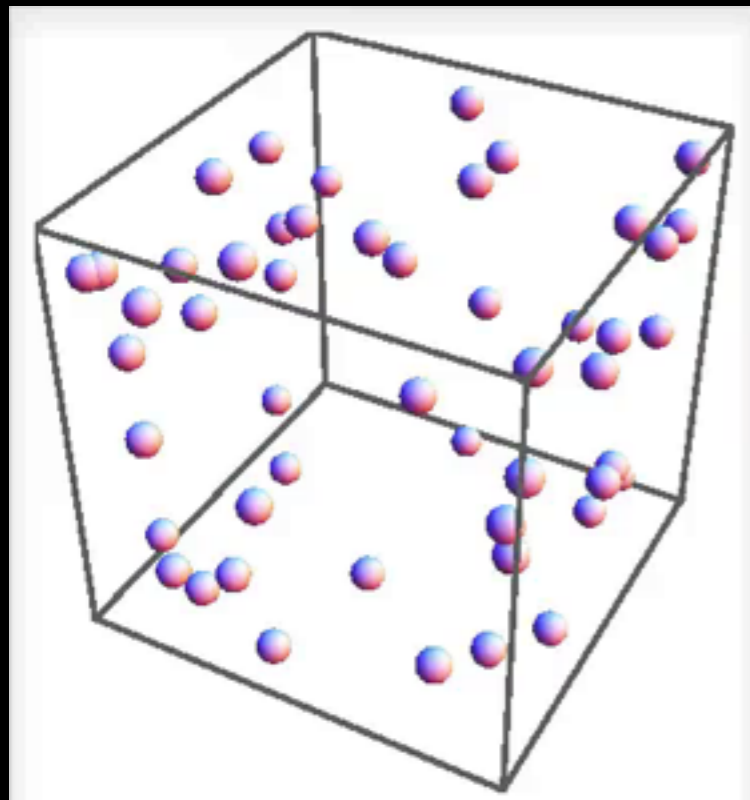
# Kinetic Theory of Gases

$pV = nRT$  is a very simple relation! ... why does it work?

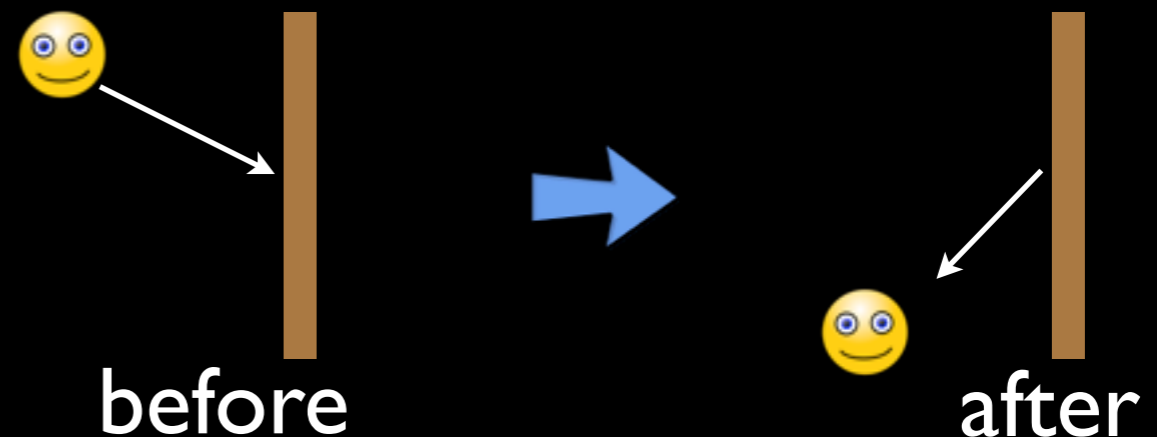


Assume:

(IV) collisions with container wall are elastic



conserve molecule's momentum  
energy



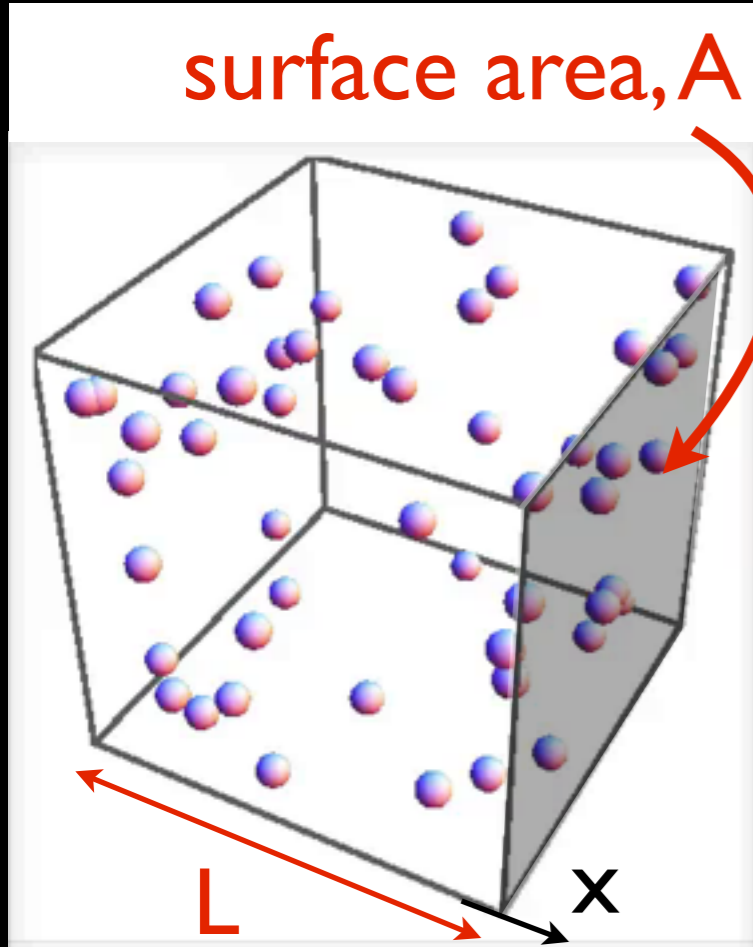
$$E_{\text{before}} = E_{\text{after}}$$

$$p_{m,\text{before}} = p_{m,\text{after}}$$

# Kinetic Theory of Gases

**N molecules in box**

**surface area, A**



Molecules collide with wall with force,  $\bar{F}_i$



many molecules  $\rightarrow$  continuous F

$$\text{gas pressure, } p = \frac{\bar{F}}{A}$$

F? 🤔

1 particle's collision with right wall:

Force,  $\bar{F}$ , only in x-direction: only  $\bar{v}_x$  changes

$$\bar{v}_{a,y} = \bar{v}_{b,y}$$

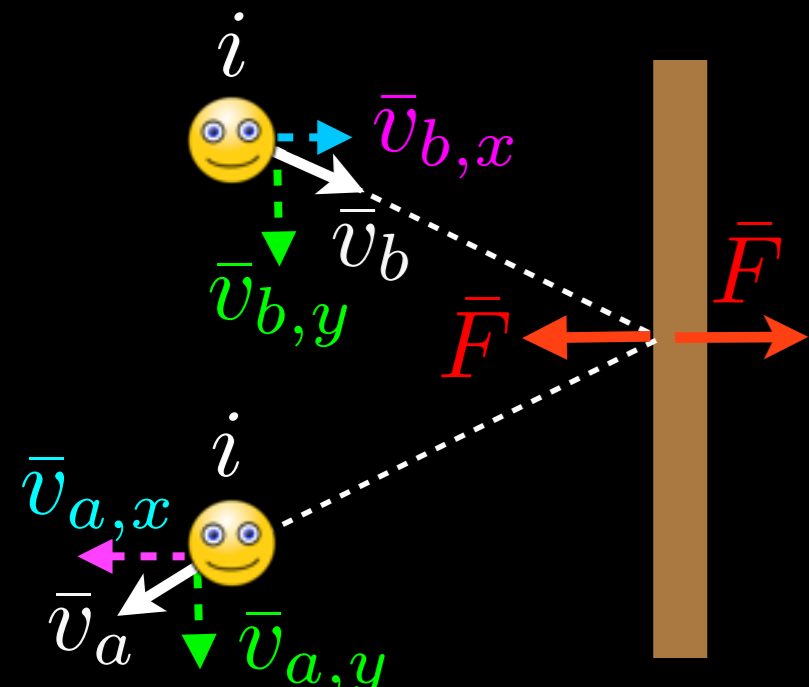
$$\bar{v}_{a,x} = -\bar{v}_{b,x} \rightarrow \Delta v = 2v_x$$

velocity x-component

Newton's 2nd law for particle  $i$

$$\bar{F}_i = \frac{\Delta \bar{p}_{mi}}{\Delta t_i}$$

change in momentum,  $\Delta p_{mi} = m\Delta v_i$   
 time between collisions,  $\Delta t_i = 2L/v_{x,i}$

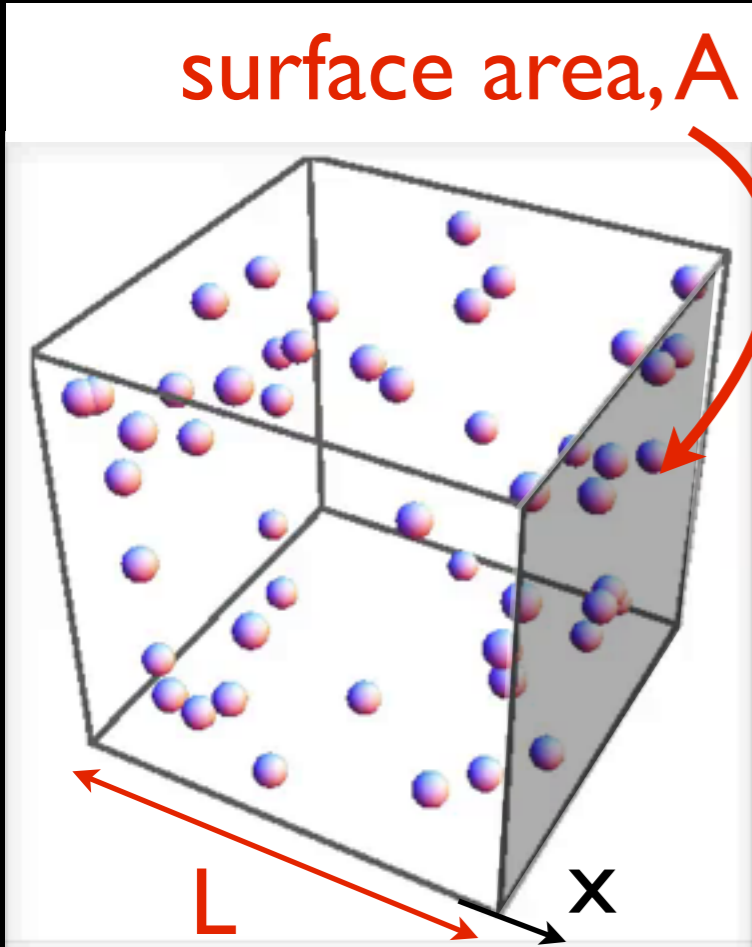




# Kinetic Theory of Gases

**N molecules in box**

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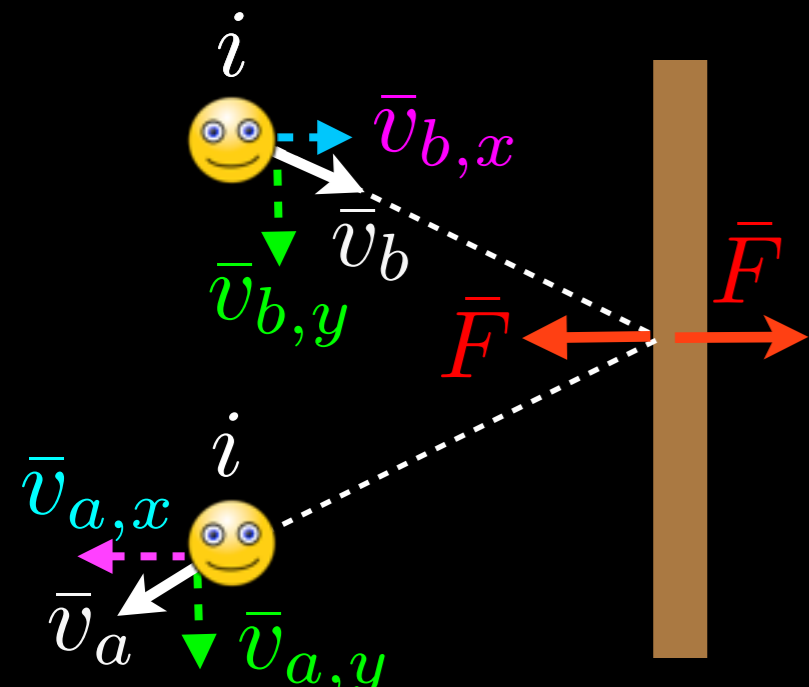
$$\bar{v}_{a,y} = \bar{v}_{b,y}$$

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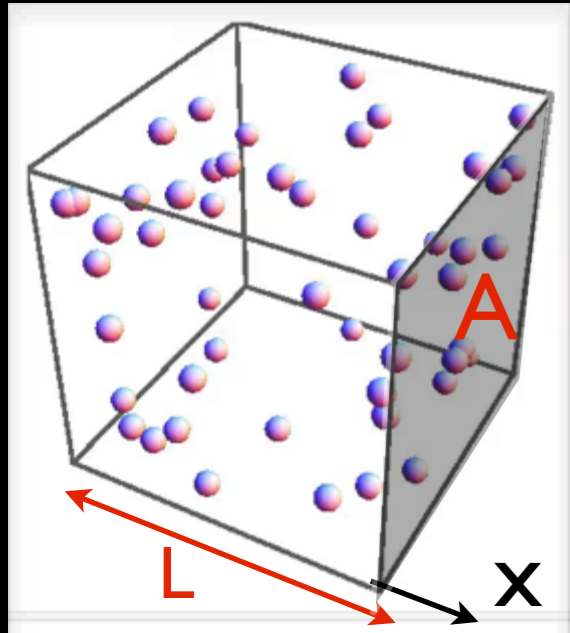
Newton's 2nd law for particle  $i$

$$\bar{F}_i = \frac{\Delta \bar{p}_{mi}}{\Delta t_i} = \frac{2mv_{x,i}}{2L/v_{x,i}} = \frac{mv_{x,i}^2}{L}$$



# Kinetic Theory of Gases

N molecules in box



Total force on wall:  $\bar{F} = \sum \bar{F}_i$


$$p = \frac{\bar{F}}{A} = \frac{\sum \bar{F}_i}{A} = \frac{\sum m v_{x,i}^2 / L}{A} = \frac{m \sum v_{x,i}^2}{\underbrace{AL}_{\text{volume, } V}}$$

$$= \frac{\underbrace{N}_{=1} m \sum v_{x,i}^2}{V} = \frac{mN}{V} \frac{\sum v_{x,i}^2}{N} = \frac{mN}{V} \bar{v}_x^2$$

average  $\bar{v}_x^2$  of all molecules  
 $= \bar{v}_x^2$

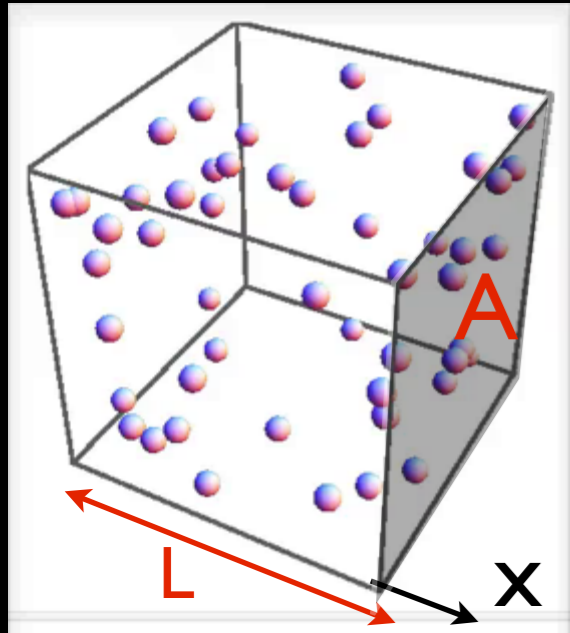
 But molecules move in random directions (III) :  $\bar{v}_x^2 = \bar{v}_y^2 = \bar{v}_z^2$

Average speed:  $\bar{v}^2 = \bar{v}_x^2 + \bar{v}_y^2 + \bar{v}_z^2 = 3\bar{v}_x^2 \rightarrow \bar{v}_x^2 = \frac{1}{3}\bar{v}^2$

  $p = \frac{mN}{3V} \bar{v}^2$

# Kinetic Theory of Gases

N molecules in box



$$p = \frac{mN}{3V} \bar{v}^2 \quad \rightarrow \quad pV = \overset{=1}{\frac{2}{2}} \frac{mN}{3} \bar{v}^2$$

$$= \frac{2}{3} N \left( \frac{1}{2} m \bar{v}^2 \right)$$

Average kinetic energy of molecule

But... ideal gas law:  $pV = NkT$

$$kT = \frac{2}{3} \left( \frac{1}{2} m \bar{v}^2 \right)$$



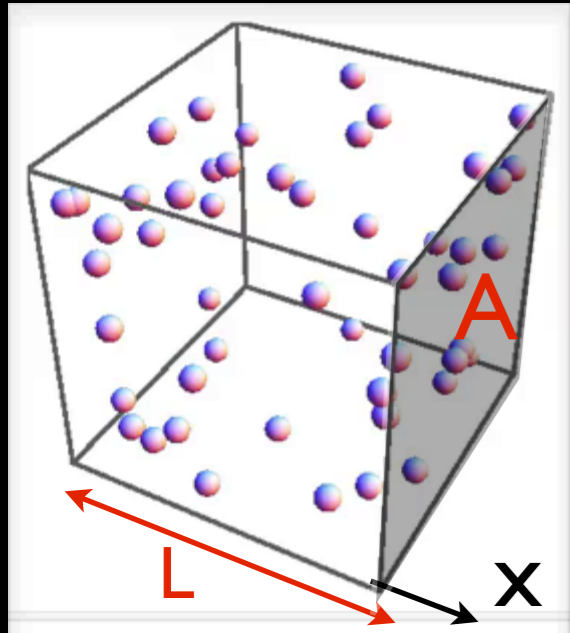
macroscopic property  
(Temperature, T)

microscopic property  
(molecule's energy) 😊

Laws of mechanics give us the ideal gas law!

# Kinetic Theory of Gases

N molecules in box



$$p = \frac{mN}{3V} \bar{v}^2 \quad \rightarrow \quad pV = \frac{2}{2} \frac{mN}{3} \bar{v}^2$$

$$= \frac{2}{3} N \left( \frac{1}{2} m \bar{v}^2 \right)$$

Average kinetic energy of molecule

But... ideal gas law:  $pV = NkT$

$$kT = \frac{2}{3} \left( \frac{1}{2} m \bar{v}^2 \right)$$



macroscopic property (Temperature, T)

microscopic property (molecule's energy) 🤖

$$\frac{3}{2} kT = \frac{1}{2} m \bar{v}^2$$

T measures average K of molecules

# Kinetic Theory of Gases

Ex.

Find the average kinetic energy of a molecule at 20° C.

$$\begin{aligned}\text{Average kinetic energy: } \bar{K} &= \frac{1}{2} m \bar{v}^2 = \frac{3}{2} kT \\ &= \frac{3}{2} (1.38 \times 10^{-23} \text{ J/K})(293 \text{ K}) \\ &= 6.07 \times 10^{-21} \text{ J}\end{aligned}$$

# Kinetic Theory of Gases

Ex.

... what is the speed of a nitrogen molecule ( $\text{N}_2$ ) with this energy?

$$1u = 1.66 \times 10^{-27} \text{ kg}$$

14 u

14 u

$$\begin{aligned} m &= 2(14u)(1.66 \times 10^{-27} \text{ kg/u}) \\ &= 4.65 \times 10^{-26} \text{ kg} \end{aligned}$$

$$\begin{aligned} \bar{K} &= \frac{1}{2} m \bar{v}^2 \quad \rightarrow \quad v = \sqrt{\frac{2\bar{K}}{m}} = \sqrt{\frac{2(6.07 \times 10^{-21} \text{ J})}{4.65 \times 10^{-26} \text{ kg}}} \\ &= 511 \text{ m/s} \end{aligned}$$

# Kinetic Theory of Gases

# Quiz

A 5.0 litre gas tank holds 1.4 moles of helium (He) and 0.70 moles of oxygen ( $O_2$ ) at  $T = 260$  K.

Find the TOTAL (translational) kinetic energy of the gas.

(a) 6.8 kJ

(b) 6.1 kJ

(c) 7.6 kJ

(d) 8.3 kJ

$$N_A = 6.023 \times 10^{23}$$

$$k = 1.38 \times 10^{-23} \text{ J/K}$$

$$1u = 1.66 \times 10^{-27} \text{ kg}$$

$$m_{\text{He}} = 4u$$

$$m_{\text{O}} = 16u$$

# Kinetic Theory of Gases

# Quiz

A 5.0 litre gas tank holds 1.4 moles of helium (He) and 0.70 moles of oxygen ( $O_2$ ) at  $T = 260$  K.

Find the TOTAL (translational) kinetic energy of the gas.

K for one molecule:  $K = \frac{3}{2}kT$

(a) 6.8 kJ

$$\frac{3}{2}(1.38 \times 10^{-23} \text{ J/K})(260 \text{ K})$$

$$= 5.38 \times 10^{-21} \text{ J}$$

All molecules:

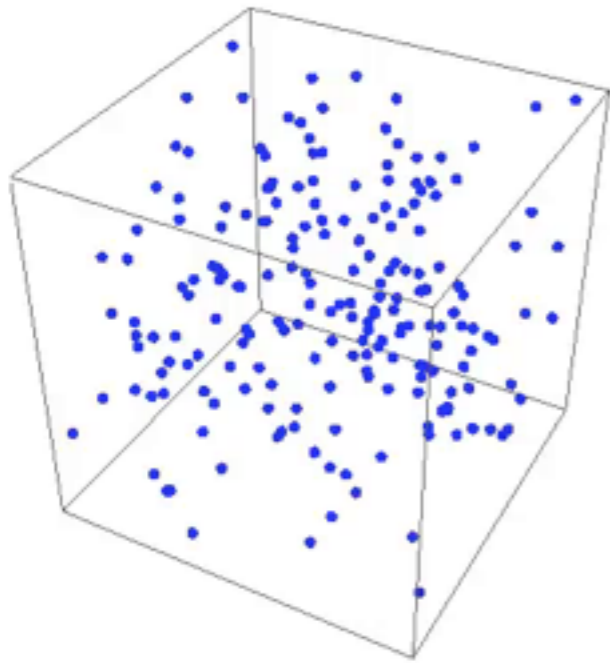
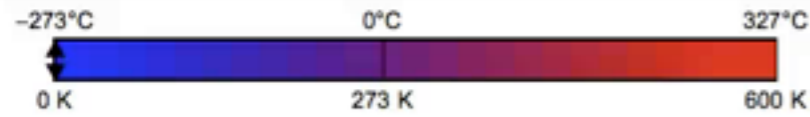
$$(1.4 \times 6.023 \times 10^{23} + 0.7 \times 6.023 \times 10^{23}) \times 5.38 \times 10^{-21} \text{ J}$$

He  $O_2$

$$= 6.8 \text{ kJ}$$



# Molecule speeds



$$\frac{3}{2}kT = \frac{1}{2}mv^2$$



$$v_{\text{th}} = \sqrt{\frac{3kT}{m}}$$

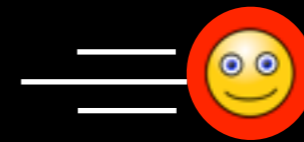
thermal speed

Typical molecular speed

but ... what are the range of speeds?



$v_{\text{min}}$



$v_{\text{max}}$

As T ↑

Typical speed ↑

But not all molecules have the same speed

# Molecule speeds

Maxwell-Boltzmann distribution:  
range of molecule speeds  
peak at typical speed,  $v_{th}$   
peak position depends on T

Higher T

Higher  $v_{th}$

Larger range: more molecules at  
**high** and **low** speeds

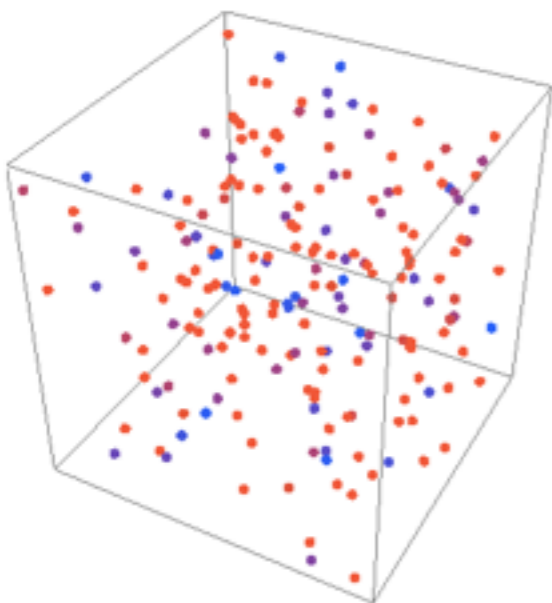
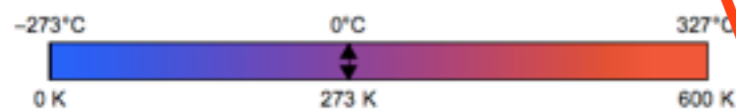
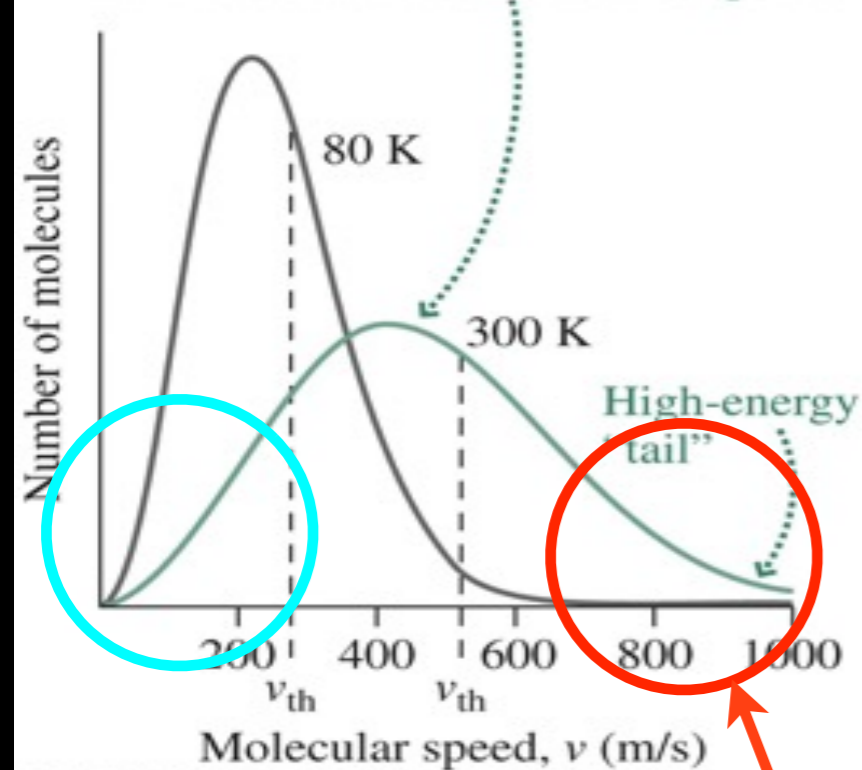


reaction rates  
faster at high T

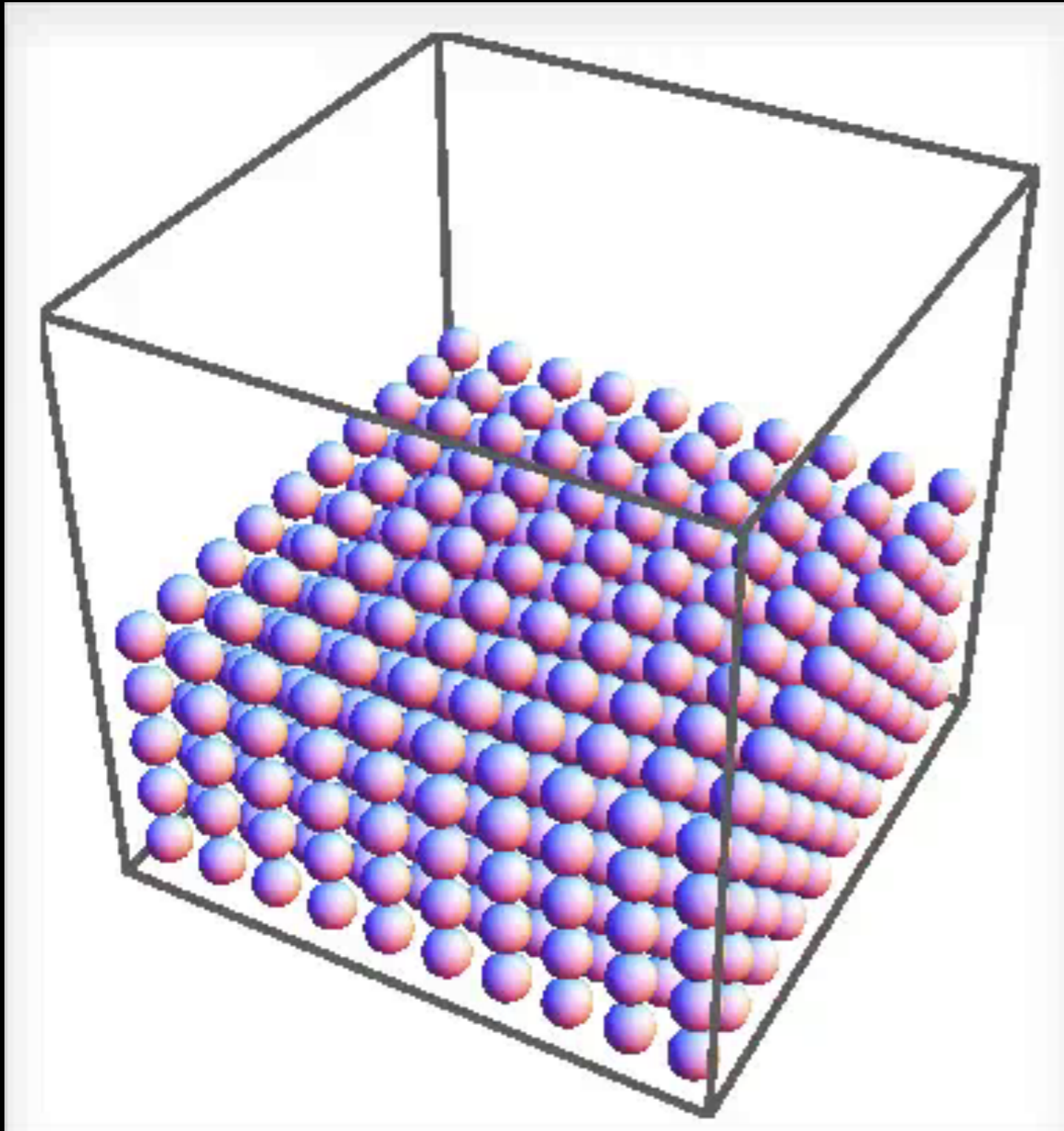
foods keep  
longer at low T



Molecules at a higher temperature  
have a broader distribution of speeds.



# Phase changes



++ heat energy



T increases...



... molecules move faster  
(++ kinetic energy)

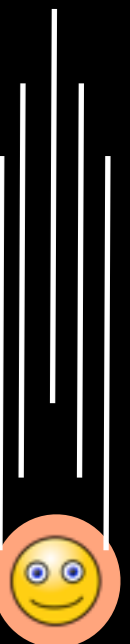


*But, if we add more heat*

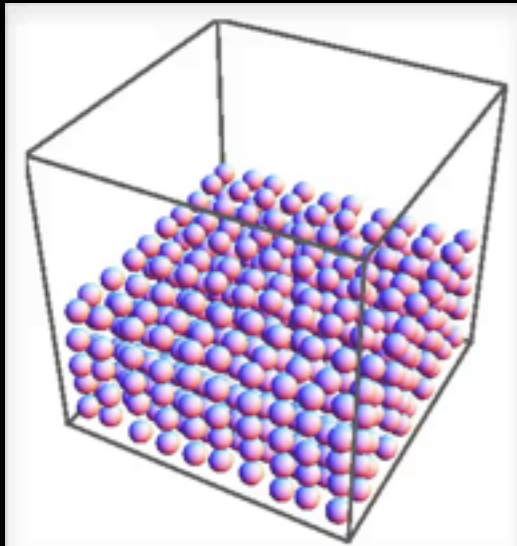
suddenly molecules  
move **freely** ...

... at the same T (and v)

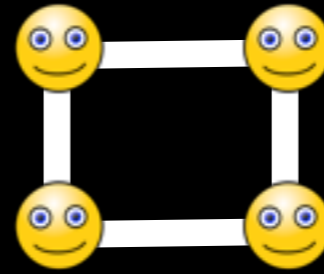
This is a **phase change**



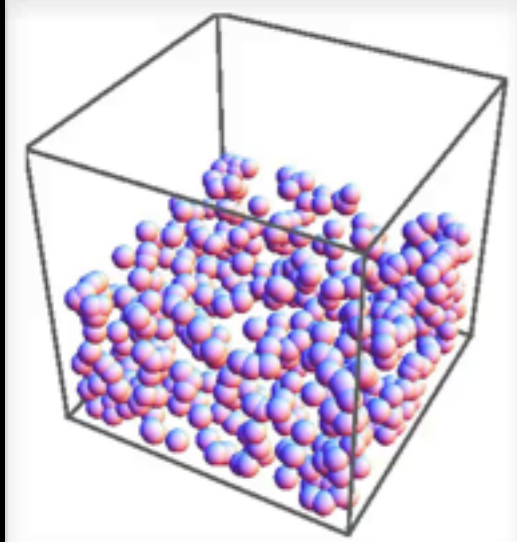
# Phase changes



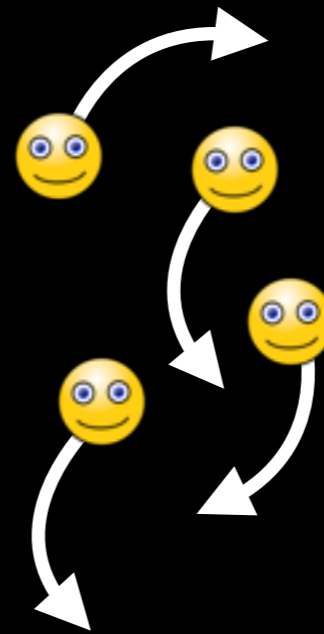
Solid



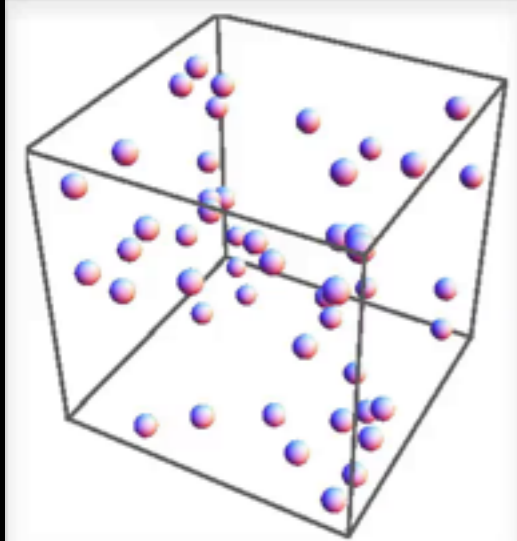
In a phase change



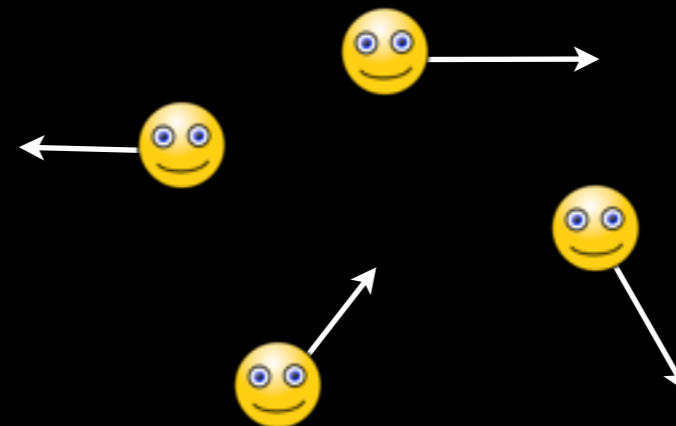
liquid



Energy is used to break molecule bonds



gas



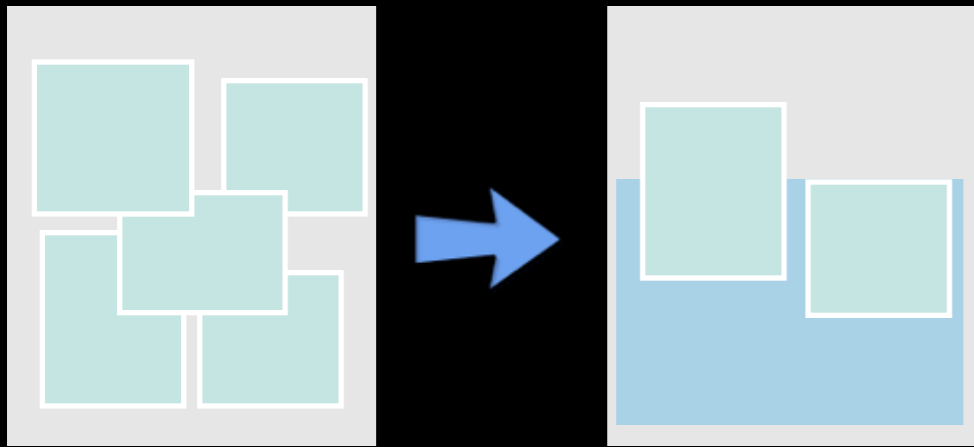
Temperature stays constant

# Phase changes

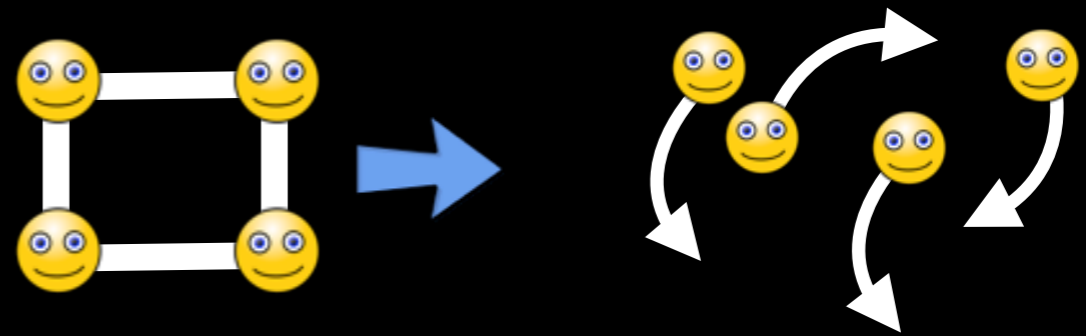
# Quiz

A glass is full of ice at  $0^{\circ}\text{C}$

After 1 hour, half the ice has melted. Is the temperature,  $T$ ...



Heat energy goes into breaking molecular bonds



(A)  $< 0^{\circ}\text{C}$

(B)  $> 0^{\circ}\text{C}$

(C)  $0^{\circ}\text{C}$

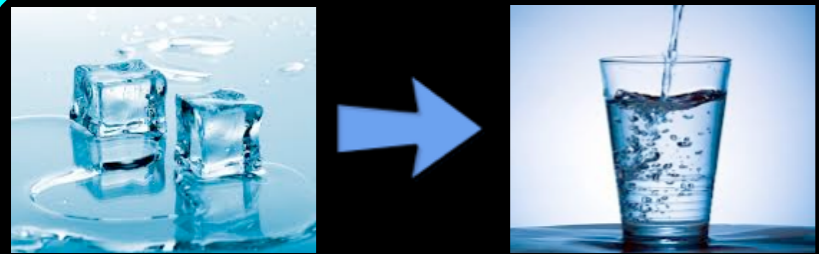
(D) Depends on the glass

while there is ice, all energy used in bond breaking.

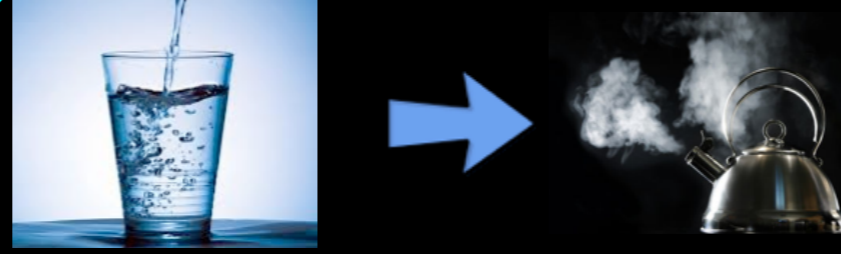
no energy raises molecule's kinetic energy (temperature)

# Phase changes

Energy / mass needed for a phase change: **heat of transformation,  $L$**   
[ J/kg ]



Solid-liquid change:  
**heat of fusion,  $L_f$**



Liquid-gas change:  
**heat of vaporization,  $L_v$**



Solid-gas change:  
**heat of sublimation**

$$Q = Lm$$

(heat of transformation)

Energy needed to change  
phase of mass,  $m$

**M**

# Phase changes



Solid-gas change:  
**heat of sublimation**

CO has a low temperature for sublimation  
(dry ice)

# Phase changes

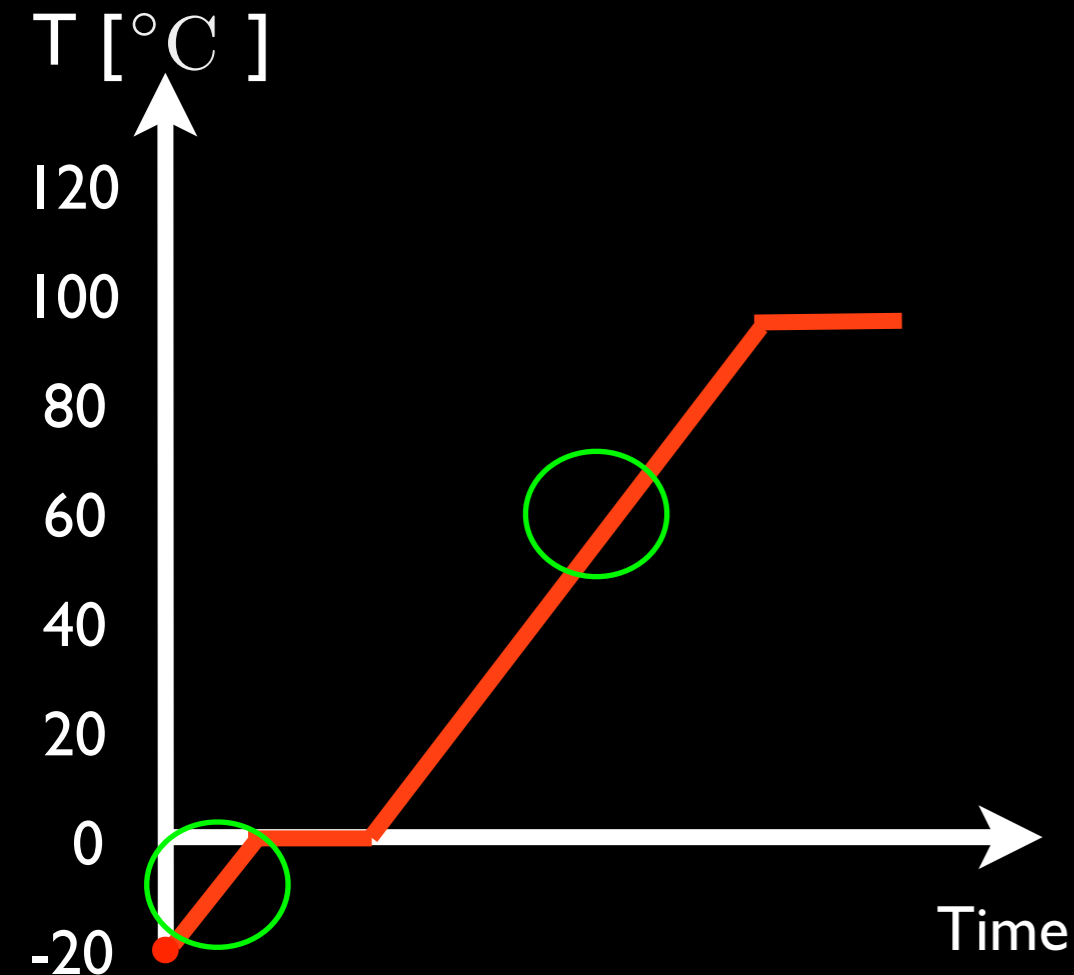
**Table 17.1** Heats of Transformation (at Atmospheric Pressure)

<b>Substance</b>	<b>Melting Point (K)</b>	<b><math>L_f</math> (kJ/kg)</b>	<b>Boiling Point (K)</b>	<b><math>L_v</math> (kJ/kg)</b>
Alcohol, ethyl	159	109	351	879
Copper	1357	205	2840	4726
Lead	601	24.7	2013	858
Mercury	234	11.3	630	296
Oxygen	54.8	13.8	90.2	213
Sulfur	388	53.6	718	306
Water	273	334	373	2257
Uranium dioxide	3120	259	3815	1533

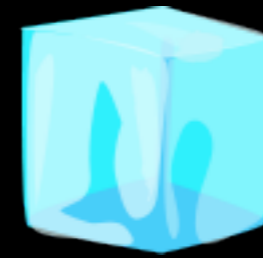


# Phase changes

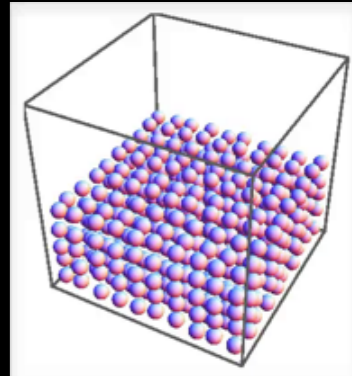
When heating ice, what happens to T?



Start at  $-20^\circ\text{C}$  .



Temperature increases to  $0^\circ\text{C}$



@  $0^\circ\text{C}$  , ice begins to melt (phase transition)



$T = \text{constant}$



When all ice is water,  $T$  increases again:

$c_{\text{water}} > c_{\text{ice}}$  water heats more slowly than ice: smaller gradient specific heat capacity

@  $100^\circ\text{C}$  , water becomes steam



$T = \text{constant}$

# Phase changes

# Quiz

How much energy does it take to melt a 45 g ice cube?

$$L_f = 334 \text{ kJ/kg}$$

(a) 22 kJ

$$Q = L_f m$$

(b) 15 kJ

$$= (334 \text{ kJ/kg})(0.045 \text{ kg})$$

(c) 0.1 kJ

$$= 15.03 \text{ kJ}$$

(d) 15480 kJ

# Phase changes

Ex.

200 g of ice at  $-10^\circ\text{C}$  are added to 1.0 kg of water at  $15^\circ\text{C}$

Is there enough ice to cool the water to  $0^\circ\text{C}$ ?

Energy removed ( $Q_1$ ):

Energy to raise T to  $0^\circ\text{C}$  + Energy to melt ice

$$m_{\text{ice}}c_{\text{ice}}\Delta T_{\text{ice}}$$

$$m_{\text{ice}}L_f$$

$$(0.20 \text{ kg})(2.05 \text{ kJ/kg} \cdot \text{K})(10 \text{ K}) + (0.20 \text{ kg})(344 \text{ kJ/kg})$$

$$Q_1 = 4.1 \text{ kJ} + 66.8 \text{ kJ} = 70.9 \text{ kJ}$$

$$Q_1 > Q_2$$

Yes!

To cool water:  $Q_2 = m_{\text{water}}c_{\text{water}}\Delta T_{\text{water}}$

$$= (1.0 \text{ kg})(4.184 \text{ kJ/kg} \cdot \text{K})(15 \text{ K}) = 62.8 \text{ kJ}$$

# Phase changes

# Quiz

How much heat must you add to vapourise (turn to steam) 2.0 g of water at 0° C?

(a) 1100 cal

(b) 1100 kcal

(c) 1200 cal

(d) 1300 cal

$$l_f = 80 \text{ cal/g}$$

$$l_v = 539 \text{ cal/g}$$

$$c = 1.0 \text{ cal/g} \cdot \text{K}$$

**watch units!**

# Phase changes

# Quiz

How much heat must you add to vapourise (turn to steam) 2.0 g of water at 0° C?

(a) 1100 cal

(b) 1100 kcal

(c) 1200 cal

(d) 1300 cal

$$Q = mc\Delta T + mL_v$$

$$= (2.0 \text{ g})(1.0 \text{ cal/g} \cdot \text{K})(100 \text{ K}) +$$

$$(2.0 \text{ g})(539 \text{ cal/g})$$

$$= 1278 \text{ cal}$$

# Phase diagrams

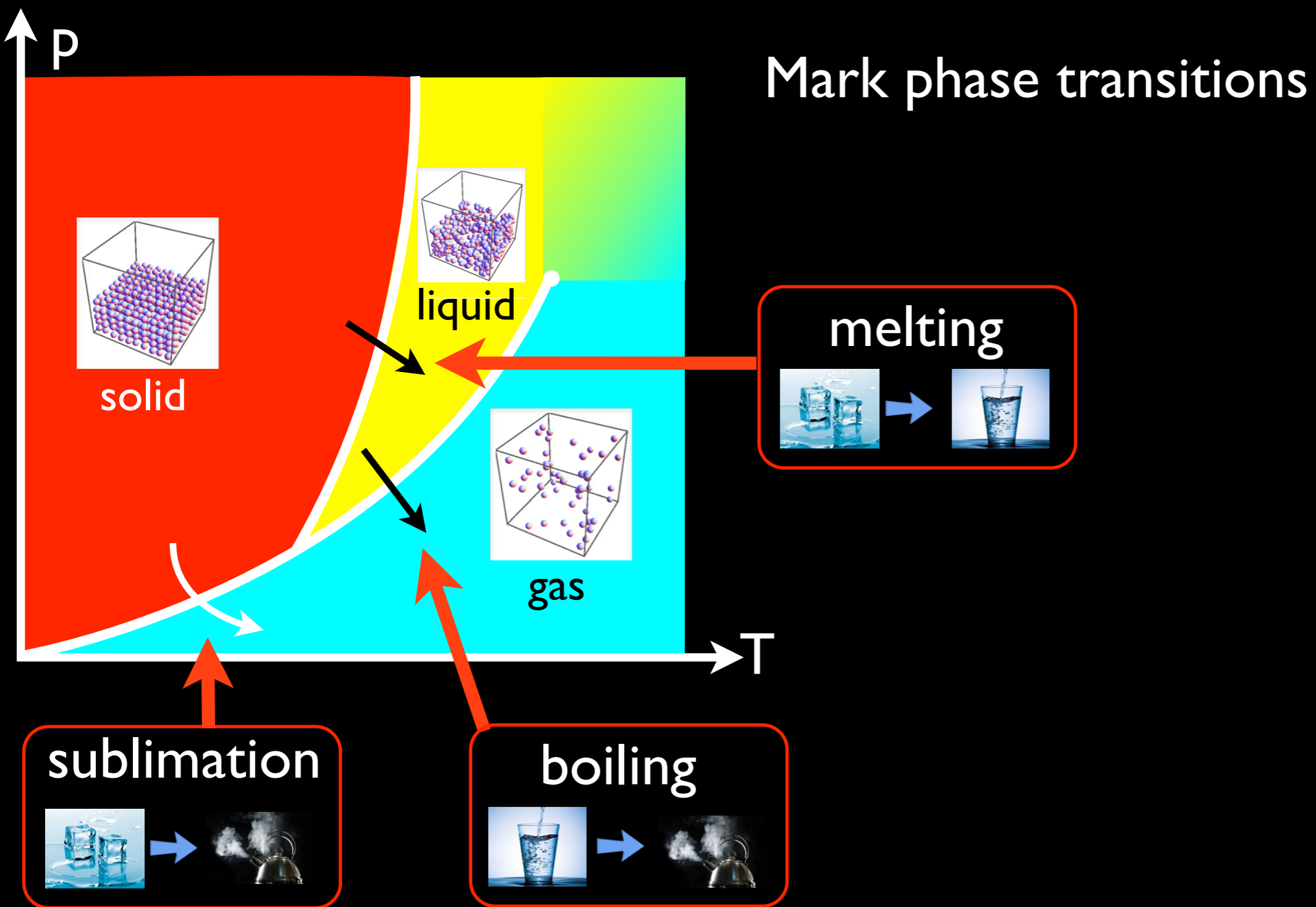
Phase changes occur at a given  $T$  ... .. and a given  $p$

If  $p \downarrow \rightarrow T_{\text{phase}} \downarrow$



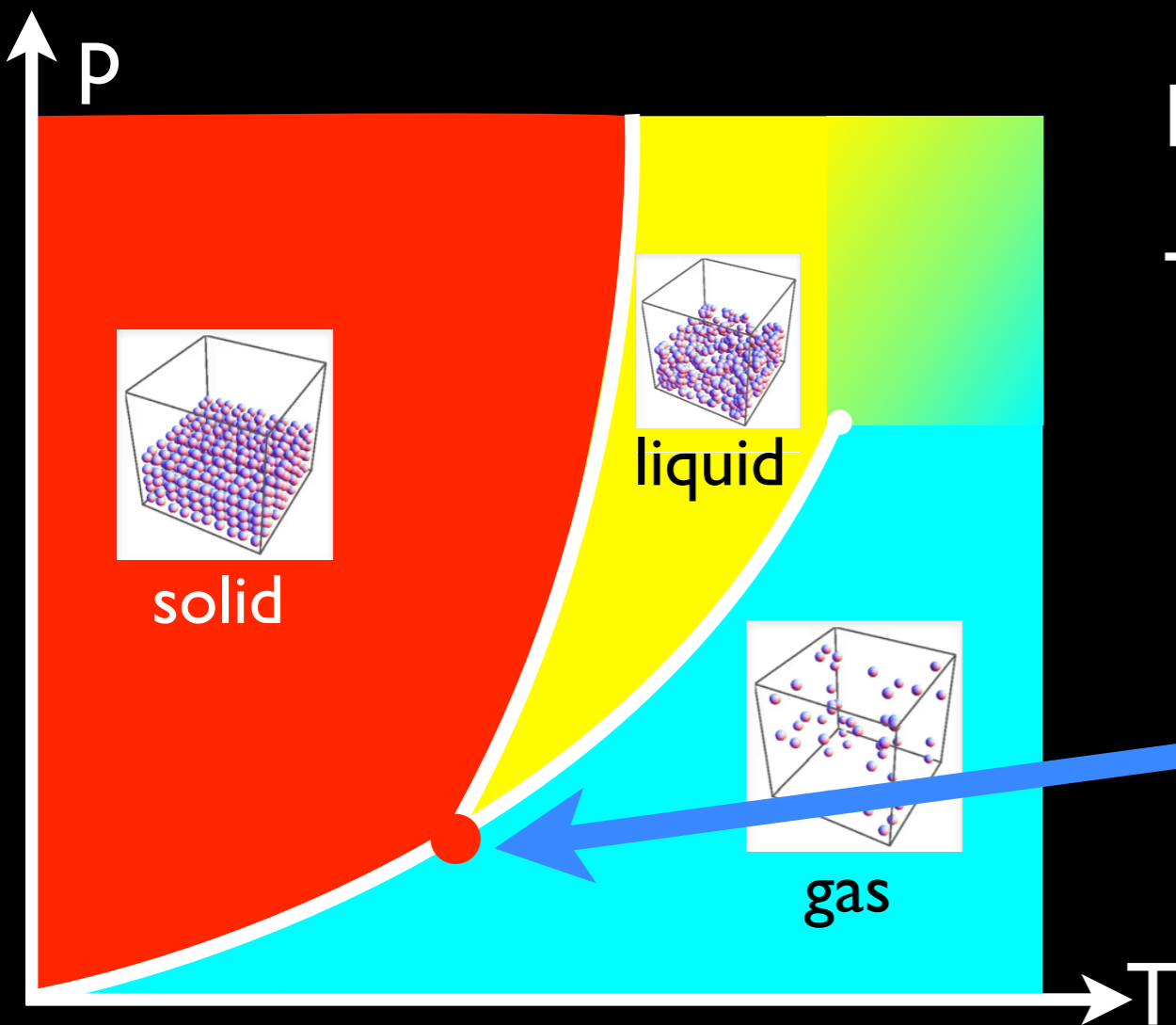
# Phase diagrams

Phase diagram: plot of pressure vs. temperature



# Phase diagrams

Phase diagram: plot of pressure vs. temperature



Mark phase transitions

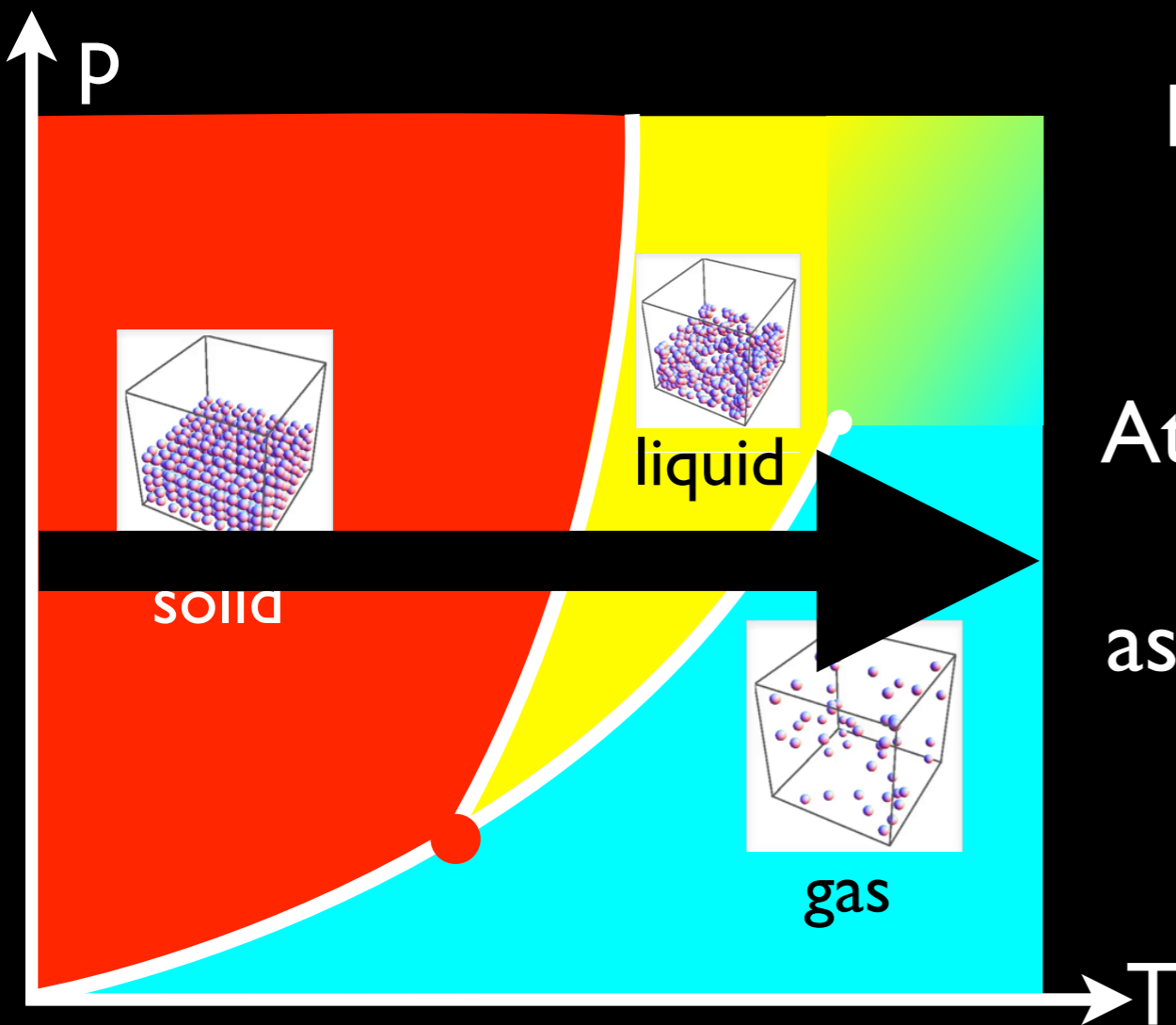
Triple point: solid, liquid, gas all exist





# Phase diagrams

Phase diagram: plot of pressure vs. temperature



Mark phase transitions

At atmospheric (normal) p,

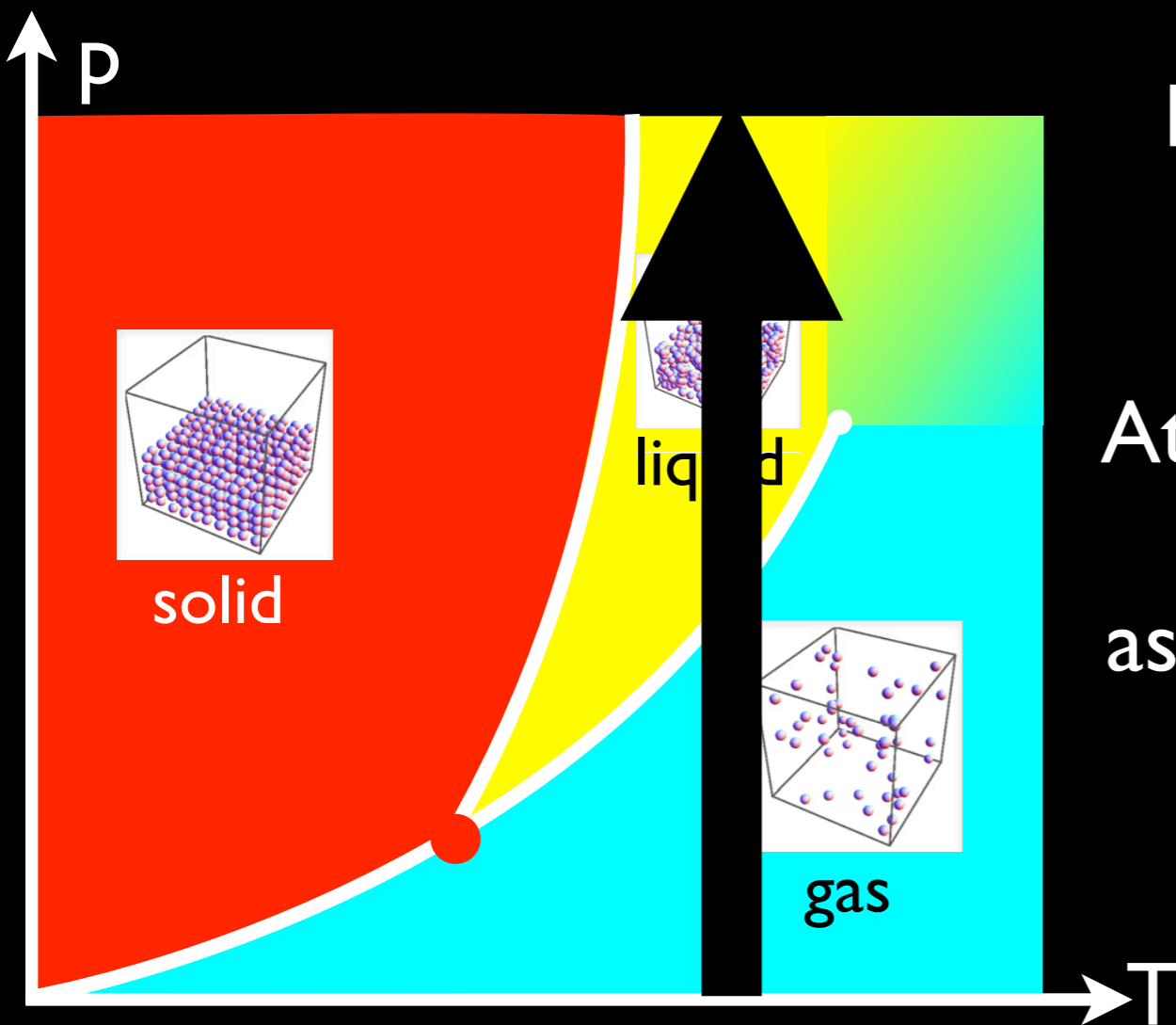
as  $T \uparrow$ : solid  $\rightarrow$  liquid  $\rightarrow$  gas phase change

Note: phase change is not fast!

Crossing lines takes a lot of energy

# Phase diagrams

Phase diagram: plot of pressure vs. temperature



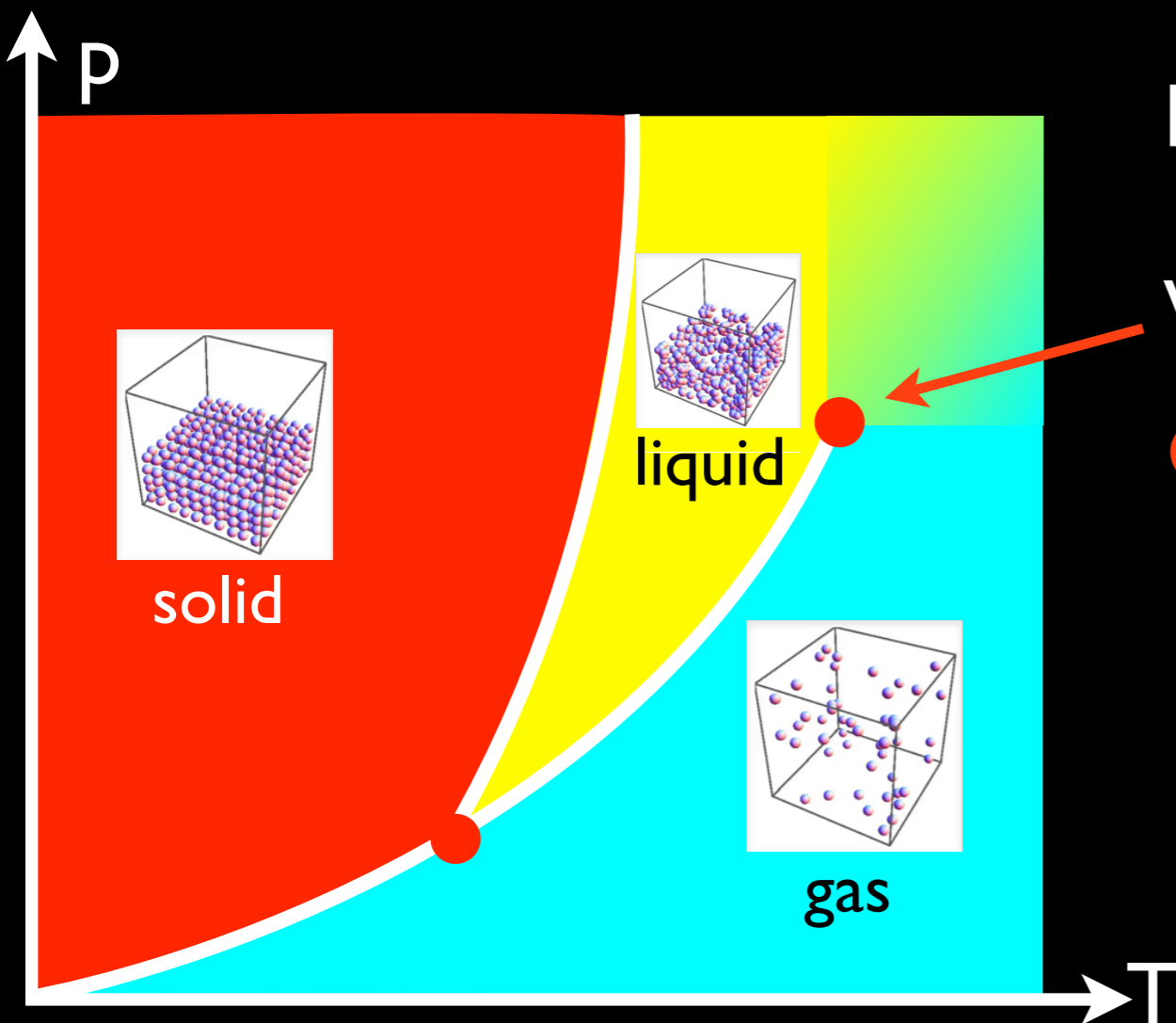
Mark phase transitions

At constant T

as  $p \uparrow$ : phase change without heat input

# Phase diagrams

Phase diagram: plot of pressure vs. temperature



Mark phase transitions

What happens here?

**Critical point :**

Sudden phase change disappears

No clear boundary between liquid and gas

gas changes gradually into liquid

# Phase diagrams

# Quiz

Everest Base Camp is at 5365 m.

Water will boil here at:

(a)  $100^{\circ}\text{C}$

(b)  $> 100^{\circ}\text{C}$

(c)  $< 100^{\circ}\text{C}$

(d)  $< 0^{\circ}\text{C}$



# Thermal Expansion

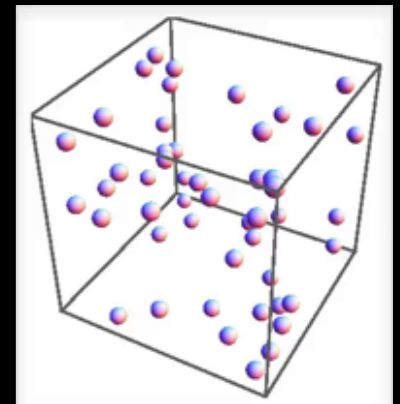
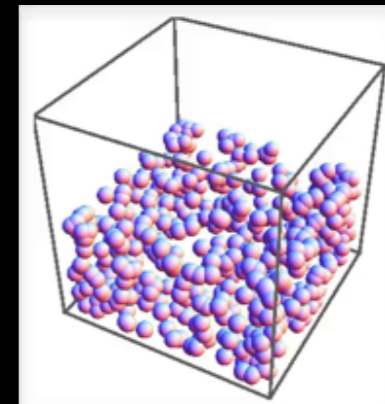
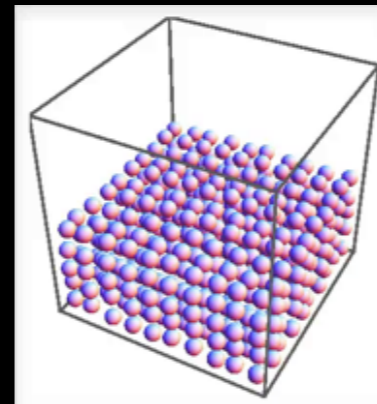
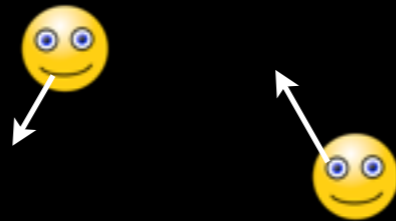
Gas:  $pV = nRT \rightarrow V \propto T$

Also true for liquid and solid but...

molecules closer



than in a gas.



➔ harder to expand and compress

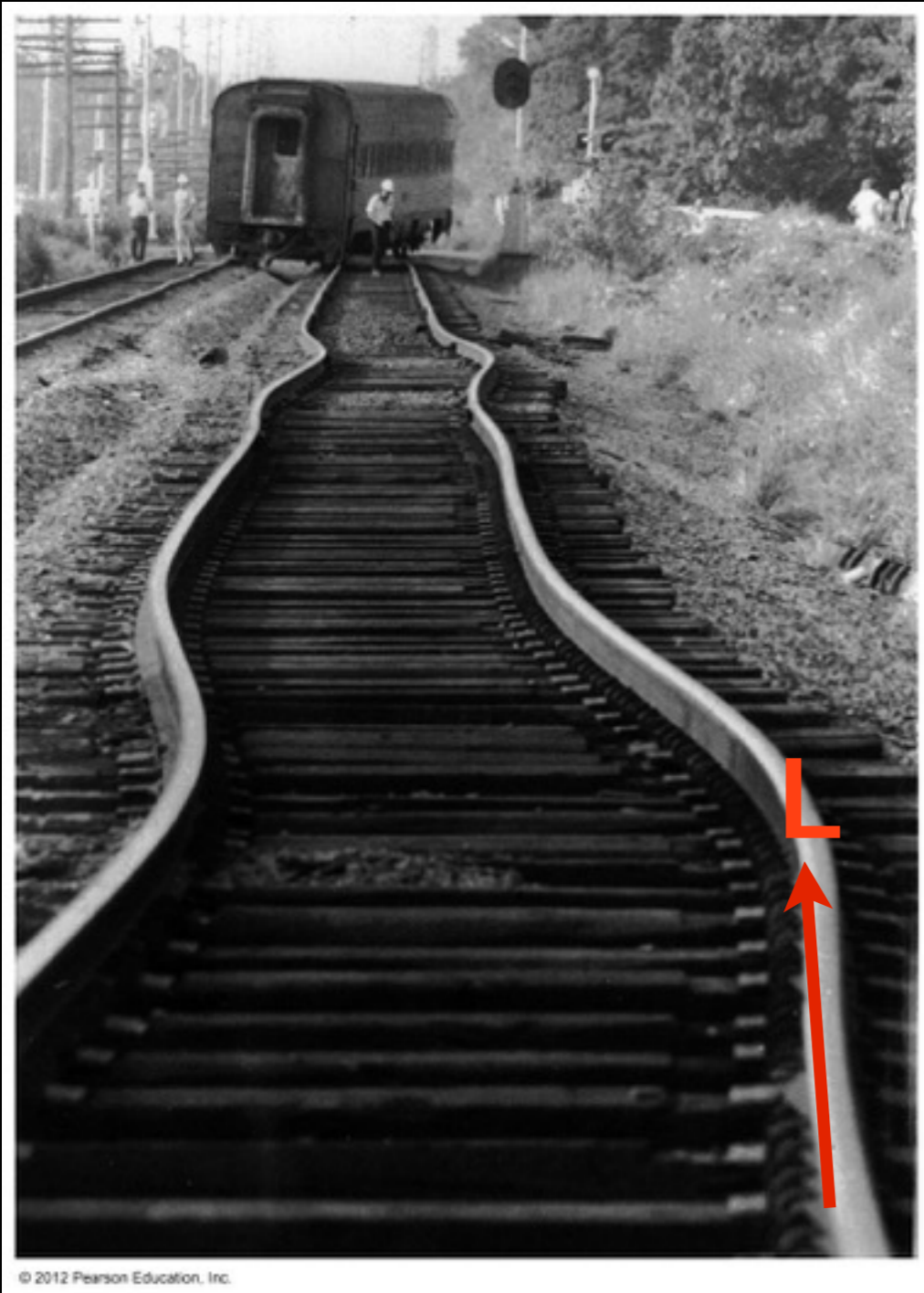


Define how much solid/liquid expands:

**coefficient of volume expansion**  $\beta = \frac{\Delta V/V}{\Delta T}$

(assumes  $\beta$  not dependent on T and constant p )

# Thermal Expansion



If object is long and thin...

Expansion / contraction greatest in 1D

$$\alpha = \frac{\Delta L / L}{\Delta T}$$

$$\beta = 3\alpha$$



only important to solids

(liquids / gases can change shape)

# Thermal Expansion

# Quiz

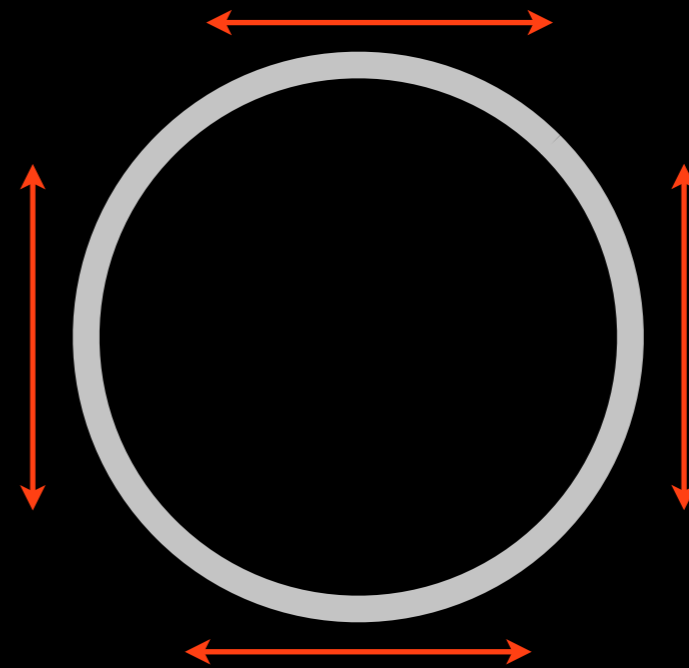
You heat up an iron ring.

The hole in the middle...

(a) larger?

(b) smaller?

(c) does not change?



all dimensions expand equally

# Thermal Expansion

**Table 17.2** Expansion Coefficients\*

Solids	$\alpha$ ( $\text{K}^{-1}$ )	Liquids and Gases	$\beta$ ( $\text{K}^{-1}$ )
Aluminum	$24 \times 10^{-6}$	Air	$3.7 \times 10^{-3}$
Brass	$19 \times 10^{-6}$	Alcohol, ethyl	$75 \times 10^{-5}$
Copper	$17 \times 10^{-6}$	Gasoline	$95 \times 10^{-5}$
Glass (Pyrex)	$3.2 \times 10^{-6}$	Mercury	$18 \times 10^{-5}$
Ice	$51 \times 10^{-6}$	Water, 1°C	$-4.8 \times 10^{-5}$
Invar <sup>†</sup>	$0.9 \times 10^{-6}$	Water, 20°C	$20 \times 10^{-5}$
Steel	$12 \times 10^{-6}$	Water, 50°C	$50 \times 10^{-5}$

\*At approximately room temperature unless noted.

<sup>†</sup>Invar, consisting of 64% iron and 36% nickel, is an alloy designed to minimize thermal expansion.

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64 % iron, 36% nickel: designed for small thermal expansion

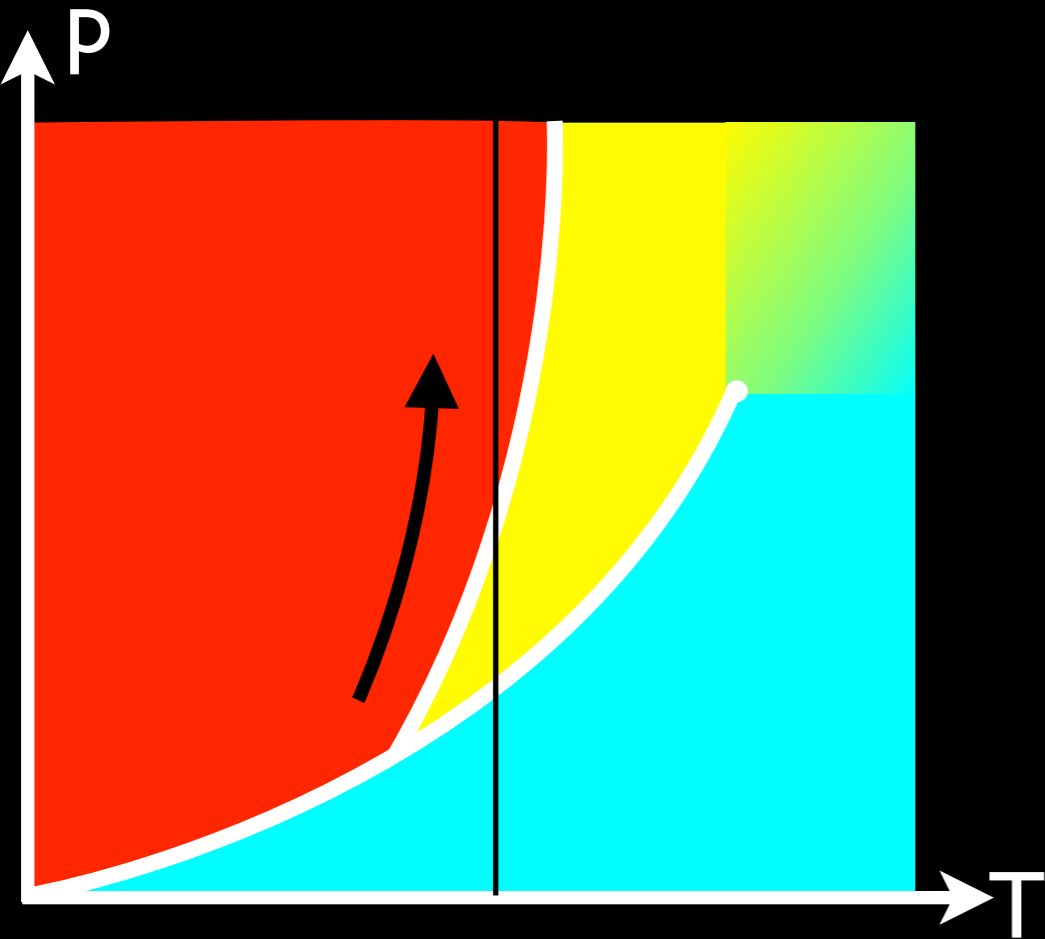
... negative?



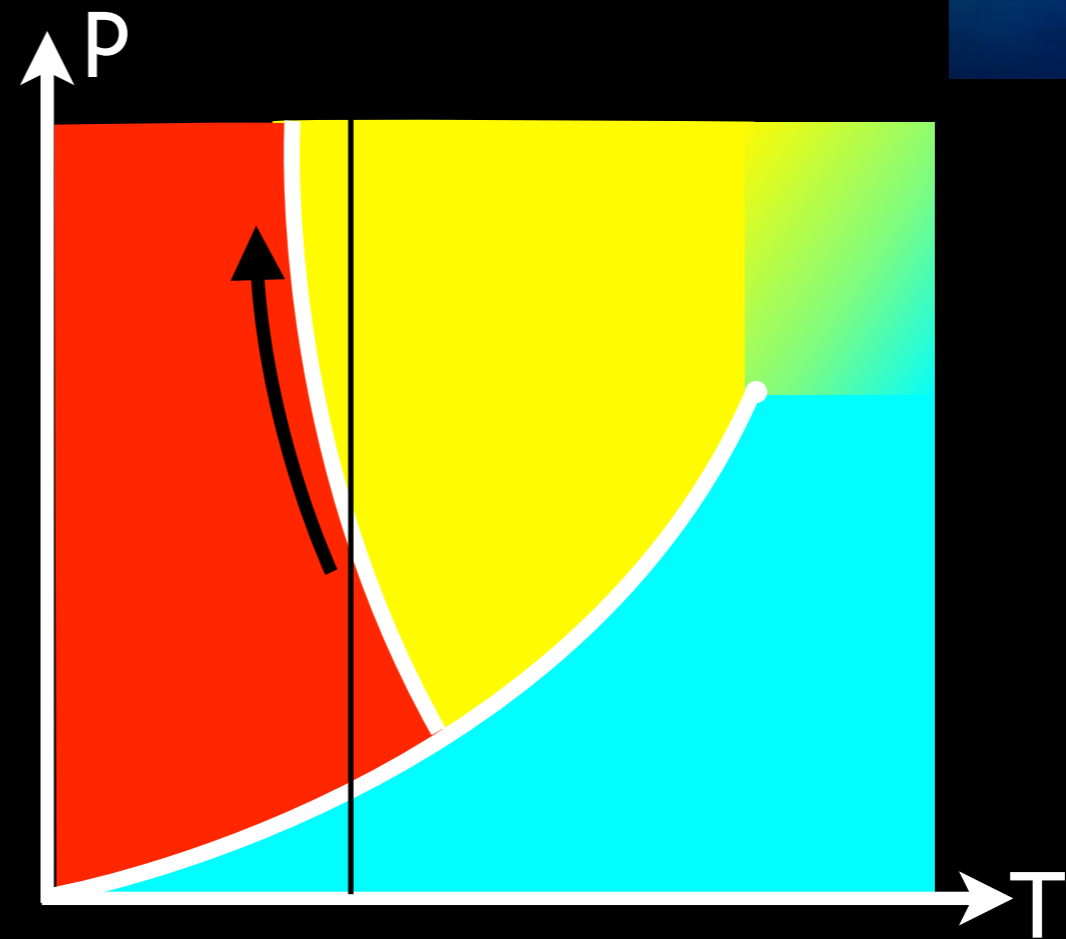
# Thermal Expansion



Water ... is strange



normal



water

At constant  $T$  ...

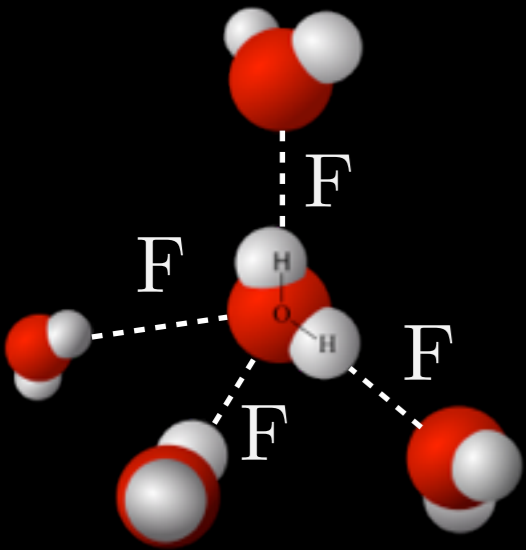
Solid will melt if  $P$  decrease



Solid will melt if  $P$  increased



# Thermal Expansion



Water is  $\text{H}_2\text{O}$

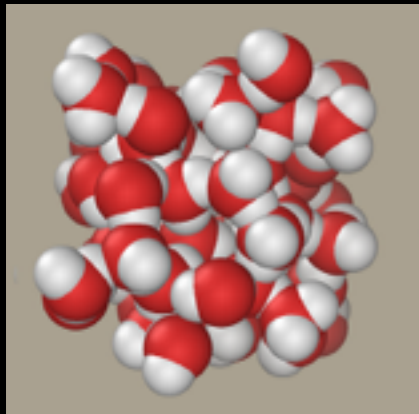


2 hydrogen atoms ●

1 oxygen atom ●

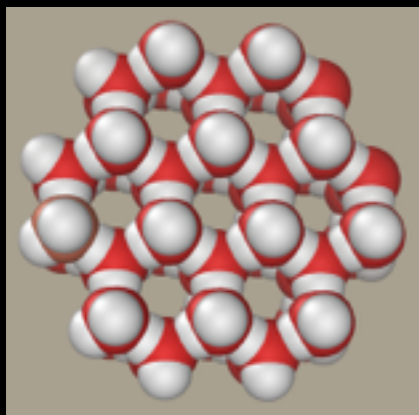
force between molecules (with H)

*'Hydrogen bonding'*



In water, *hydrogen bonding* moves molecules close

➔ dense structure



In ice, fixed structure moves molecules apart

➔ less dense

> 4°C high K moves molecules apart again.

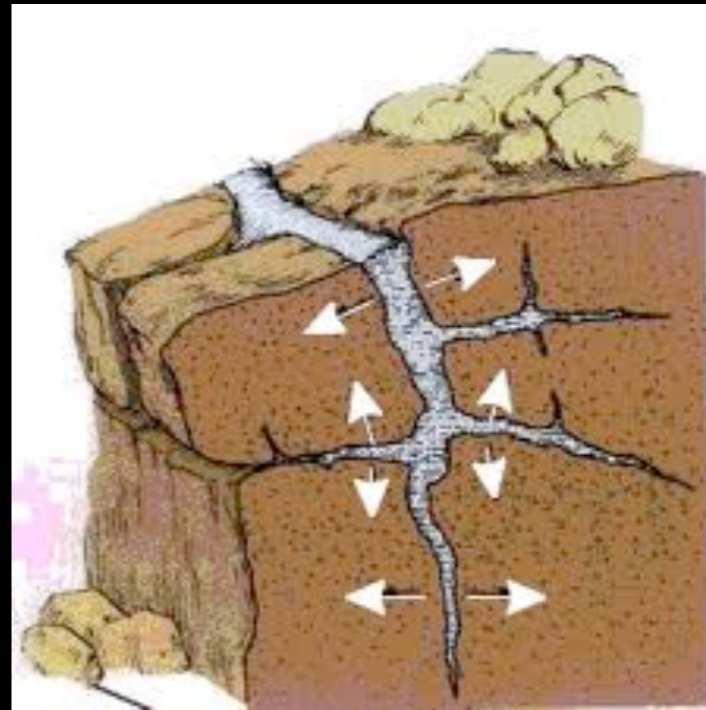
➔ densest water at 4°C

# Thermal Expansion



Good for sea life!

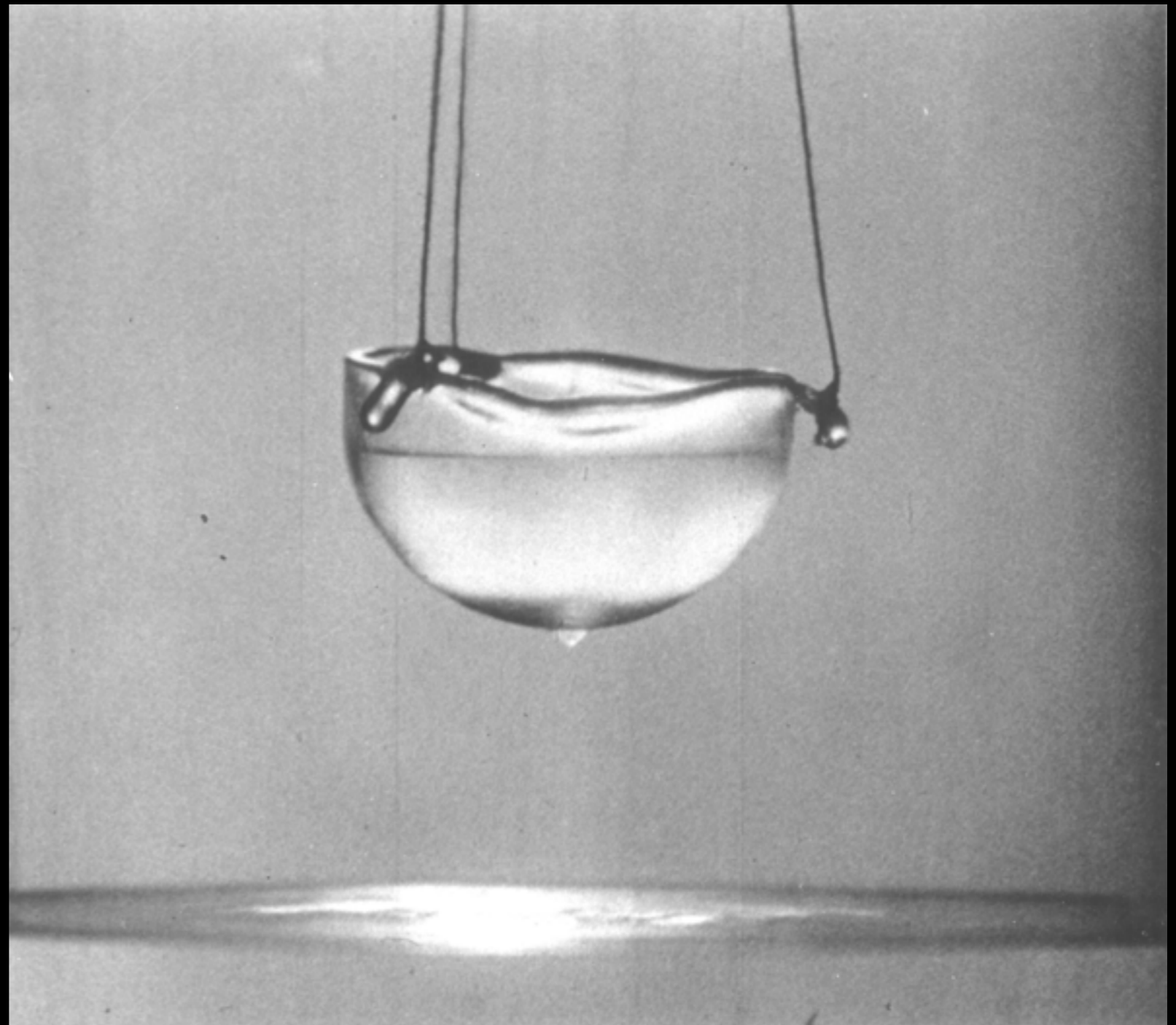
In winter, ice forms only on surface



Bad for roads!

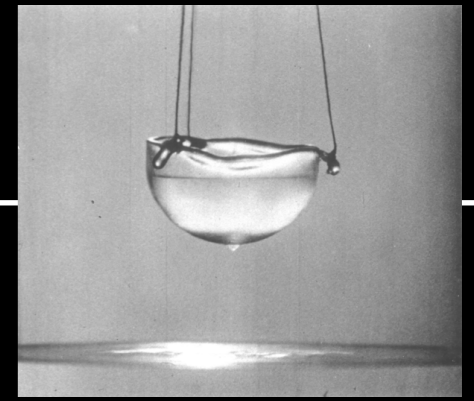


## Superfluids



# Superfluids

Helium-4 ( ${}^4\text{He}$ ) facts:

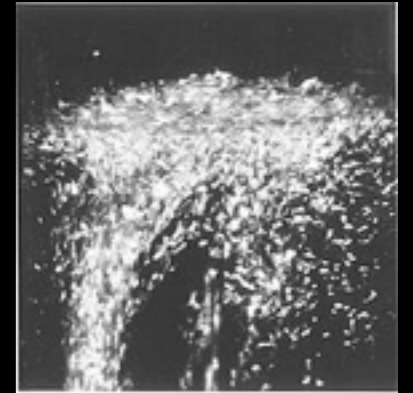


Liquid-gas change:  
(Boiling point)



$-269^\circ\text{C}$

4 K



Very difficult to make helium-4 a liquid by cooling!

(Helium-4 is only a solid at very high pressures)

Below 2.17 K, helium-4 goes through a new phase change to become a **superfluid**

# Superfluids

---

When do molecules (in theory) stop moving?

(a)  $101^{\circ}\text{C}$

(b)  $0^{\circ}\text{C}$

(c)  $0\text{ K}$

(d) They can never stop moving

# Superfluids

---

How close to 0 K have scientists ever reached?

(a)  $1 \times 10^{-9}$  K

(b)  $1 \times 10^{-6}$  C

(c) 0 K

(d)  $1 \times 10^{-6}$  K

# Superfluids

What happens when the helium becomes a superfluid?

- (a) the liquid bubbles
- (b) the liquid becomes a gas
- (c) the liquid becomes very still
- (d) the liquid sublimates





# Superfluids

The range of velocities in a superfluid are...



$v_{\min}$

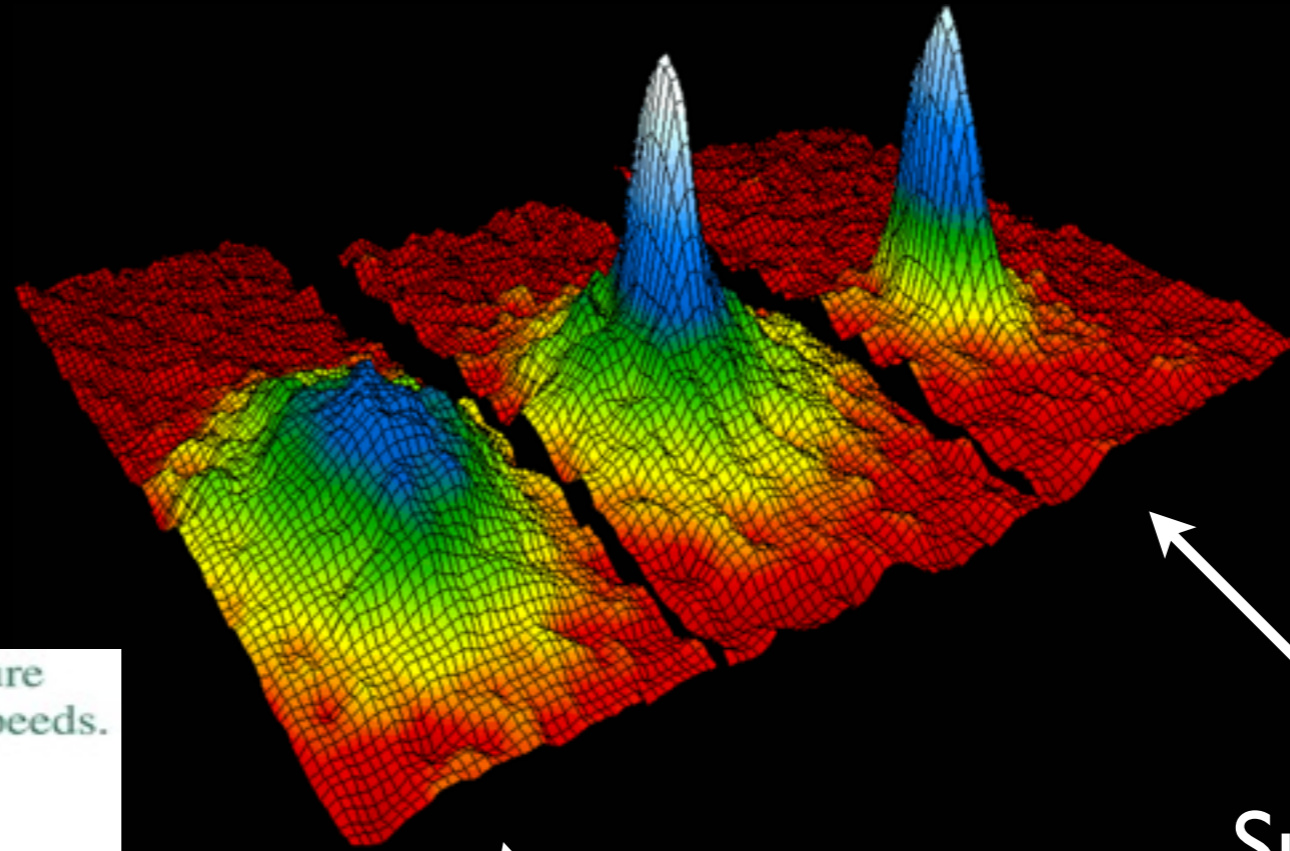


$v_{\max}$

- (a) much larger than normal fluid
- (b) same as normal fluid
- (c) much smaller than normal fluid
- (d) (almost) infinity

# Superfluids

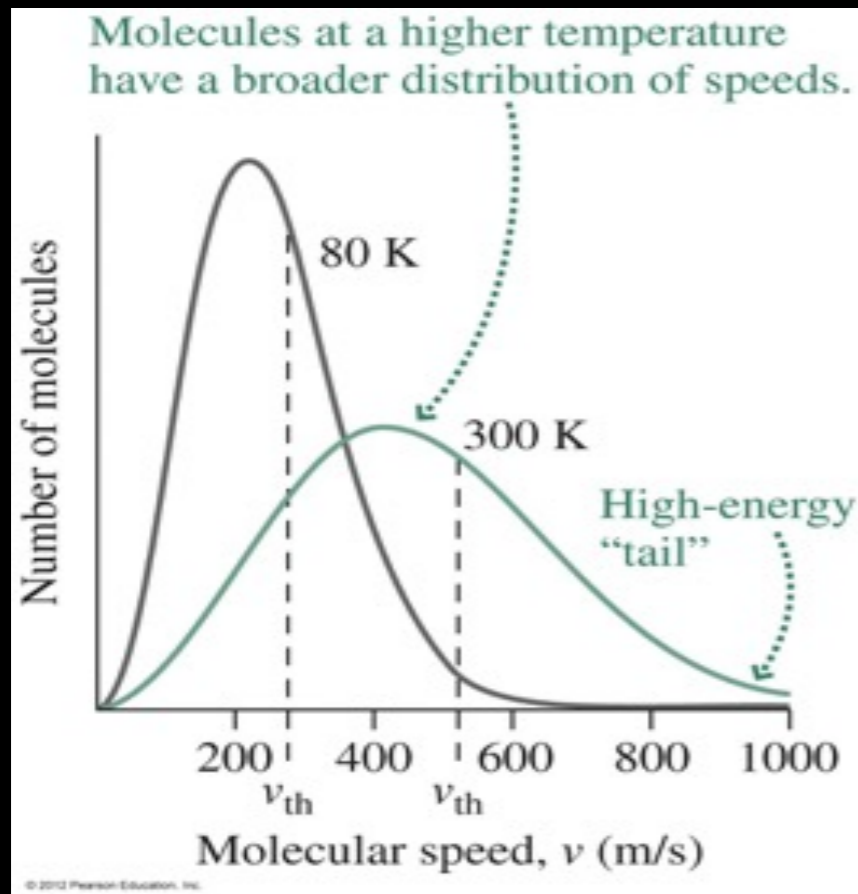
Velocity distribution:



Superfluid

Same T everywhere in fluid

VERY small velocity range



normal

Maxwell-Boltzmann distribution

# Superfluids

Why does the superfluid leak out of the bucket?



(a) the cold cracked the plug in the bucket

(b) Quantum mechanics changes the fluid properties

(c) it becomes a gas and can escape the plug

(d) no one knows

# Superfluids

A superfluid has no **viscosity**

↑  
friction for fluids

Can move through tiny holes in containers

Also, spread out and climb up walls

