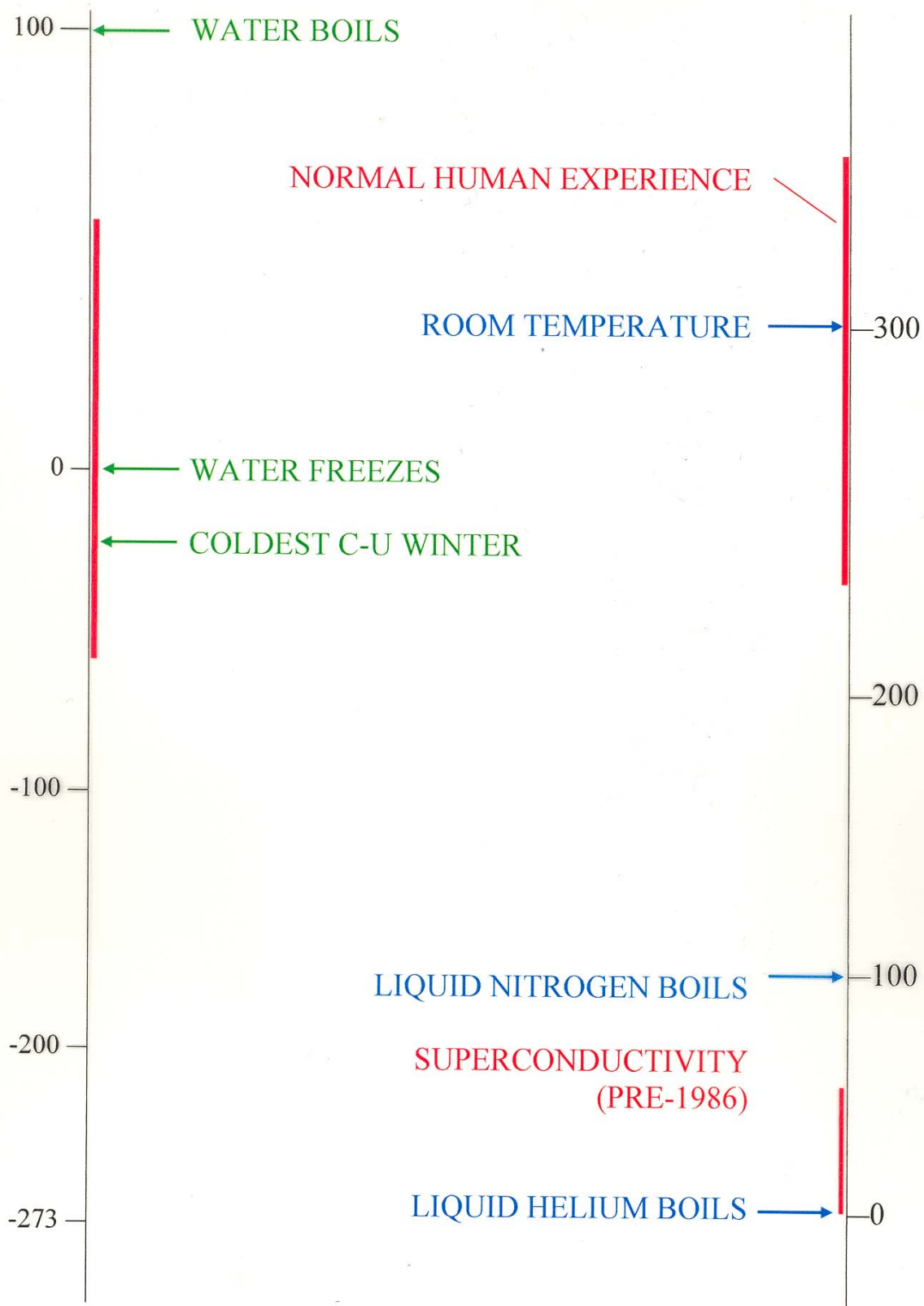


What Can We Do With a Quantum Liquid?

- Anthony J. Leggett
- University of Illinois at
Urbana-Champaign

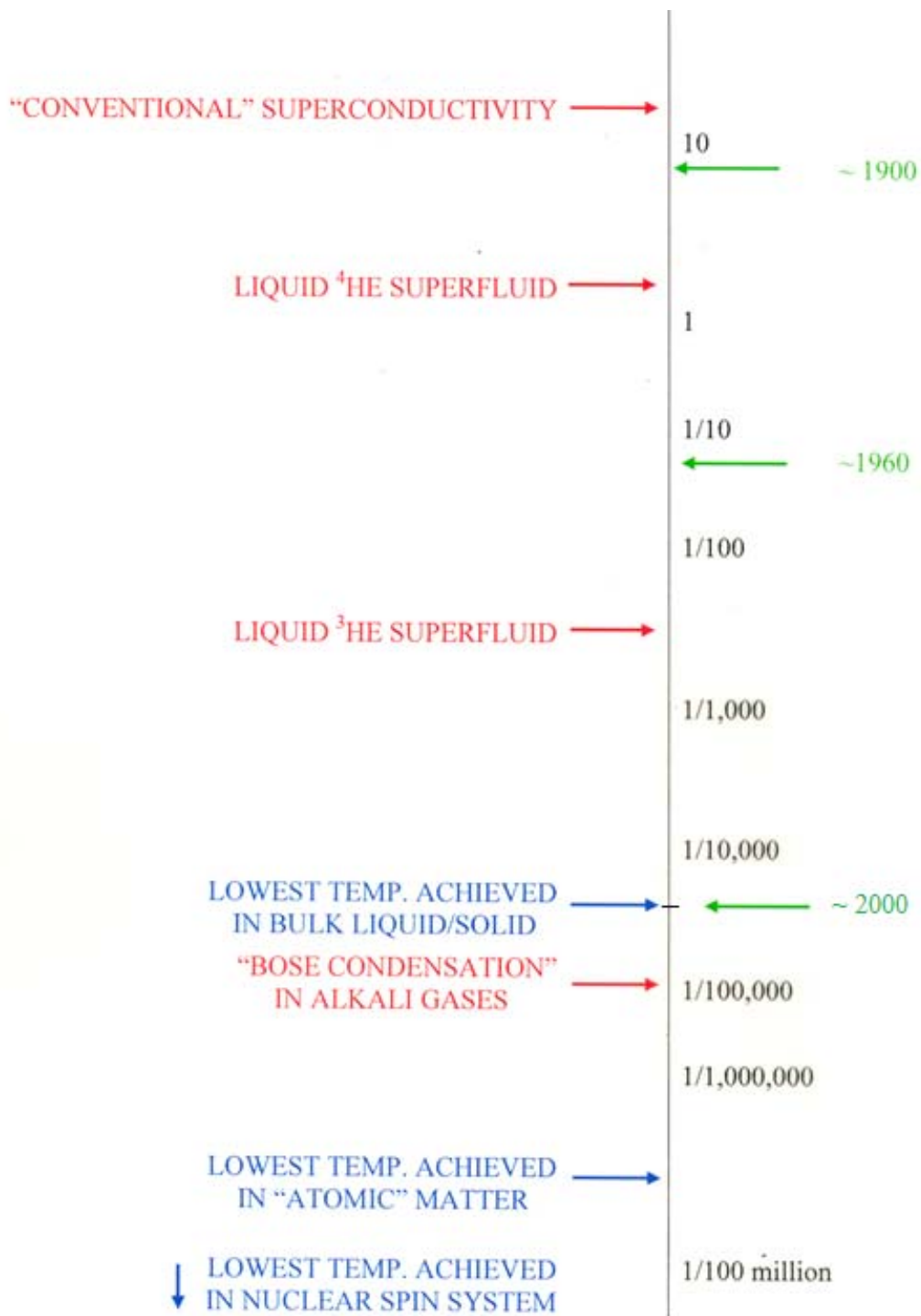


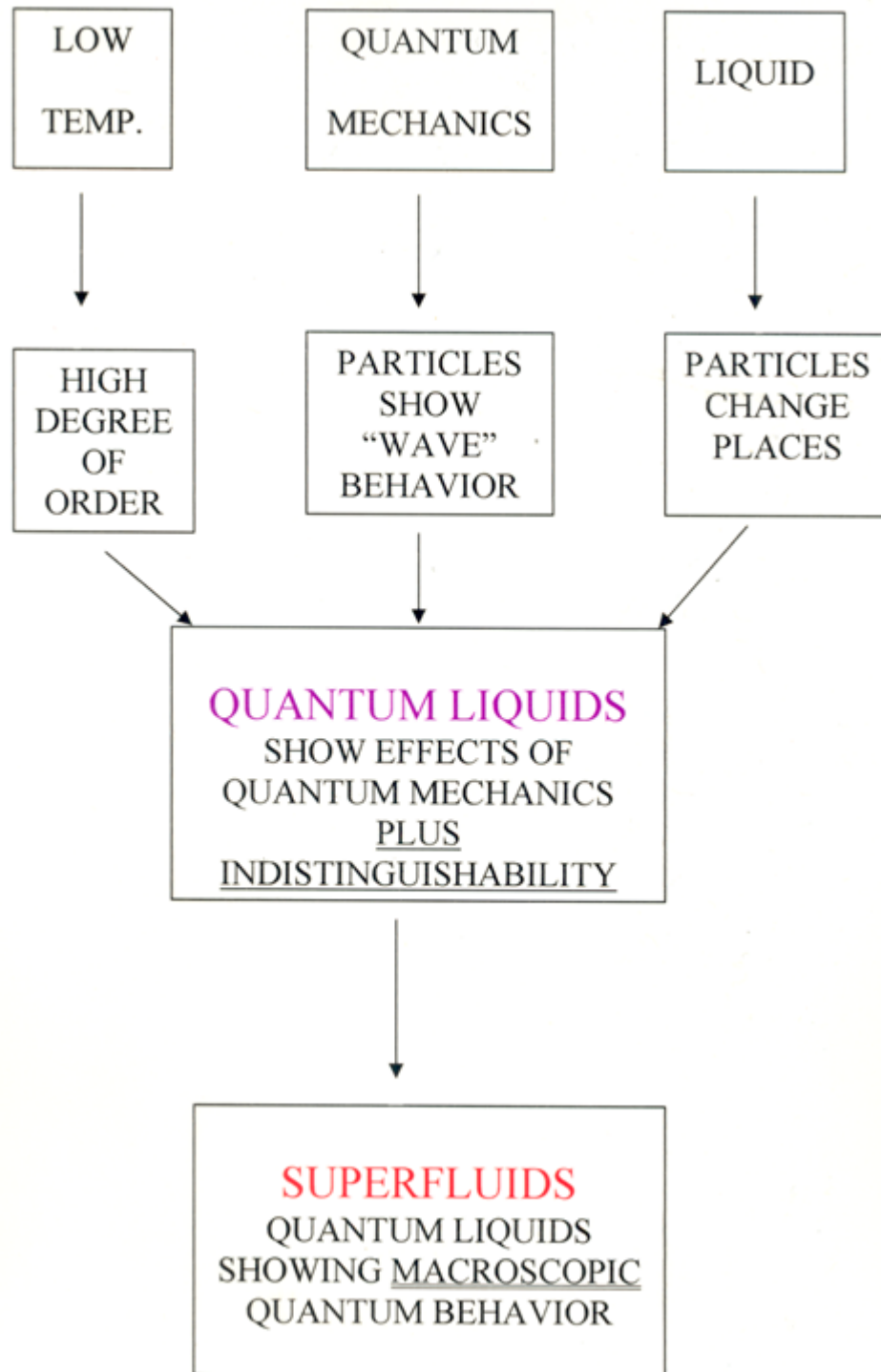


SUPERCONDUCTIVITY
(1997)



“LOGARITHMIC” TEMPERATURE SCALE
 (EACH INTERVAL CORRESPONDS TO A FACTOR OF 10)





TEMPERATURE, ORDER and DISORDER

HIGH TEMPERATURE

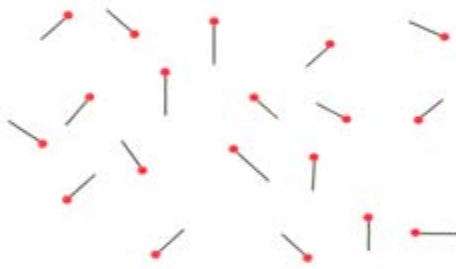


LIQUID

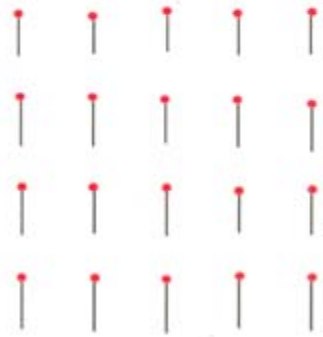
LOW TEMPERATURE



SOLID



PARAMAGNETIC



FERROMAGNETIC



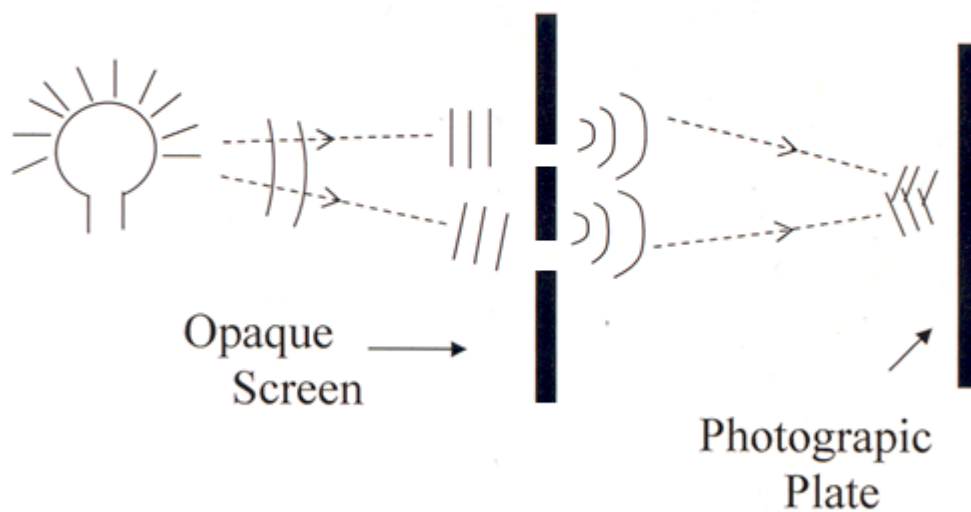
DISORDERED ALLOY



ORDERED ALLOY



PARTICLES AS WAVES



Quantitative particle-wave relation
("de Broglie relation"):

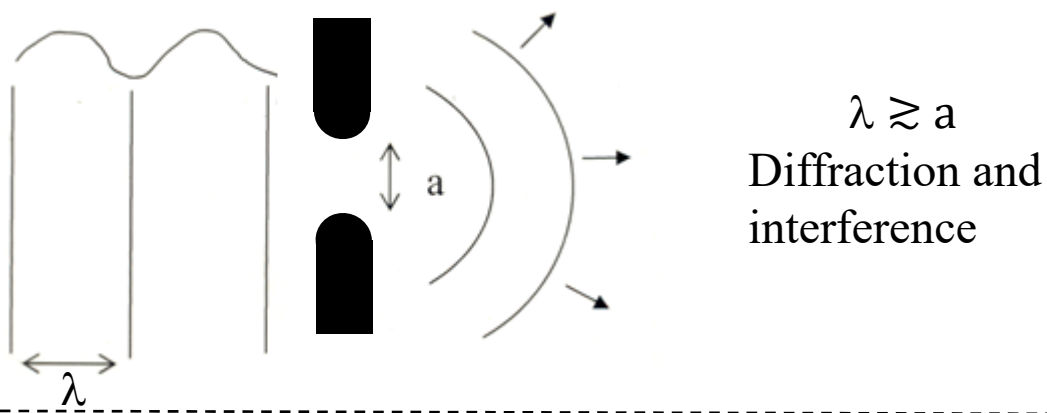
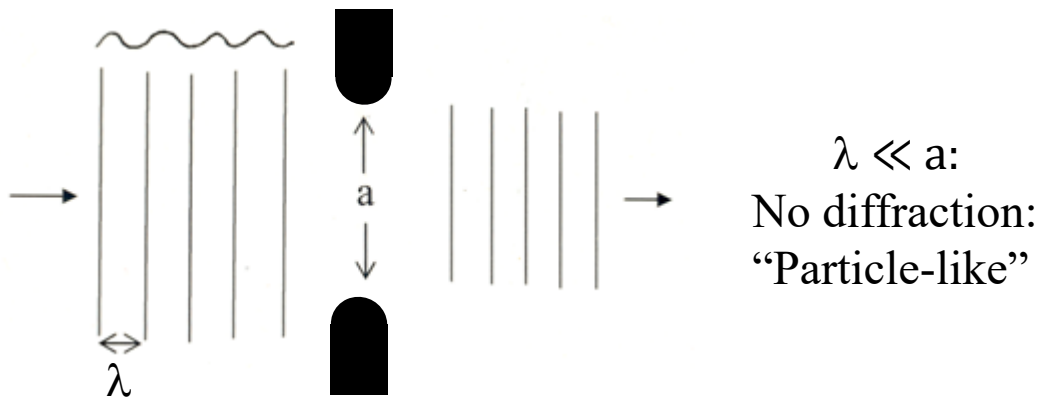
$$\lambda = \frac{h}{mv}$$

wavelength (wave property) \leftarrow λ
 Planck's constant, $\sim 6 \times 10^{-34}$ joule secs \leftarrow h
 mass \leftarrow m
 velocity \leftarrow v

(particle properties)



When does a “wave” behave like a “particle”?



since $\lambda = h/mv$ (De Broglie) to get $\lambda \gtrsim a$ need

$$v \lesssim h/ma: \text{ but } \frac{1}{2}mv^2 \sim k_B T$$

so to see “wave” effects need

$$T \lesssim h^2/2mk_B a^2$$

Boltzmann’s constant, $\sim 10^{-23}$ joules/degree

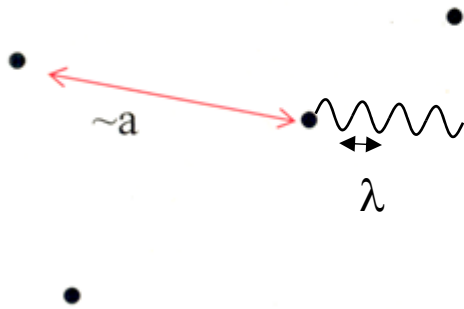
In a gas/liquid/solid, take “slit width” $a \sim$ interparticle spacing

\Rightarrow to get “wavelike” behavior, need (for atoms)

$$T \lesssim 20^\circ\text{K}/(\text{atomic number})$$

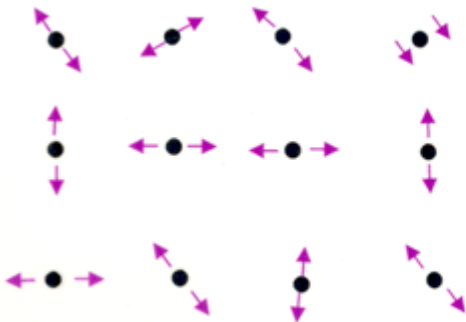
I (electrons show “wavelike” behavior for all T in liquid/solid phase)

Why “Quantum Liquids”?



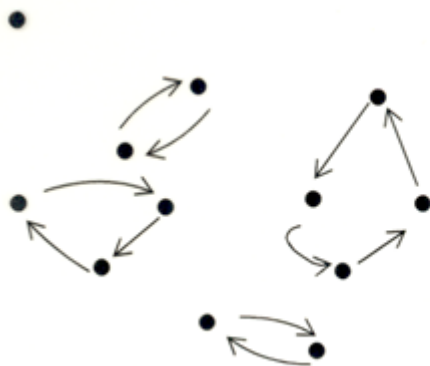
Gas: (usually)

$\lambda \ll a$
so no “wave”
(quantum) effects



Solid at low T:

$\lambda \gtrsim a$ but atoms
don't change places



Liquid at low T:

$\lambda \gtrsim a$ and
atoms change places

need: $T \lesssim 20^\circ \text{K}/(\text{atomic no.})$ and liquid!

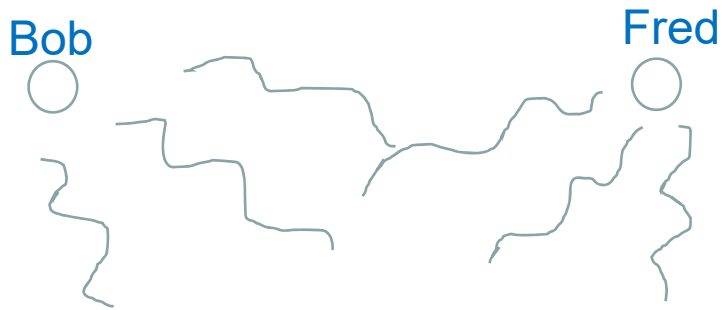
Atoms: helium (and ultracold atomic gases)

Electrons: all liquid/solid metals

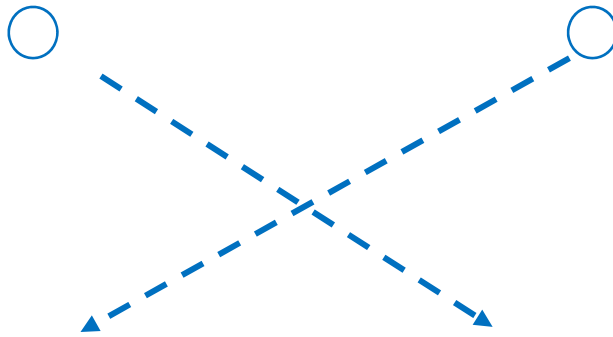


Indistinguishability of elementary particles

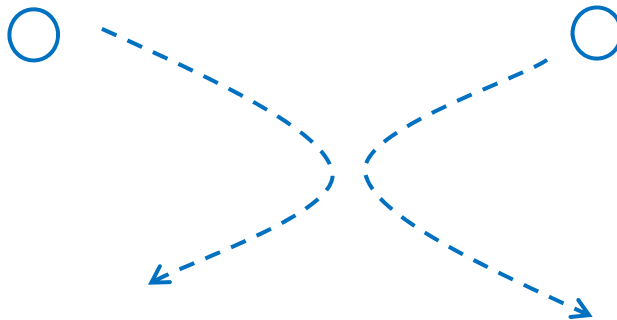
Because particles behave like waves, impossible to “tag” them.



○ ← Which is Bob and → ○
which is Fred?



or



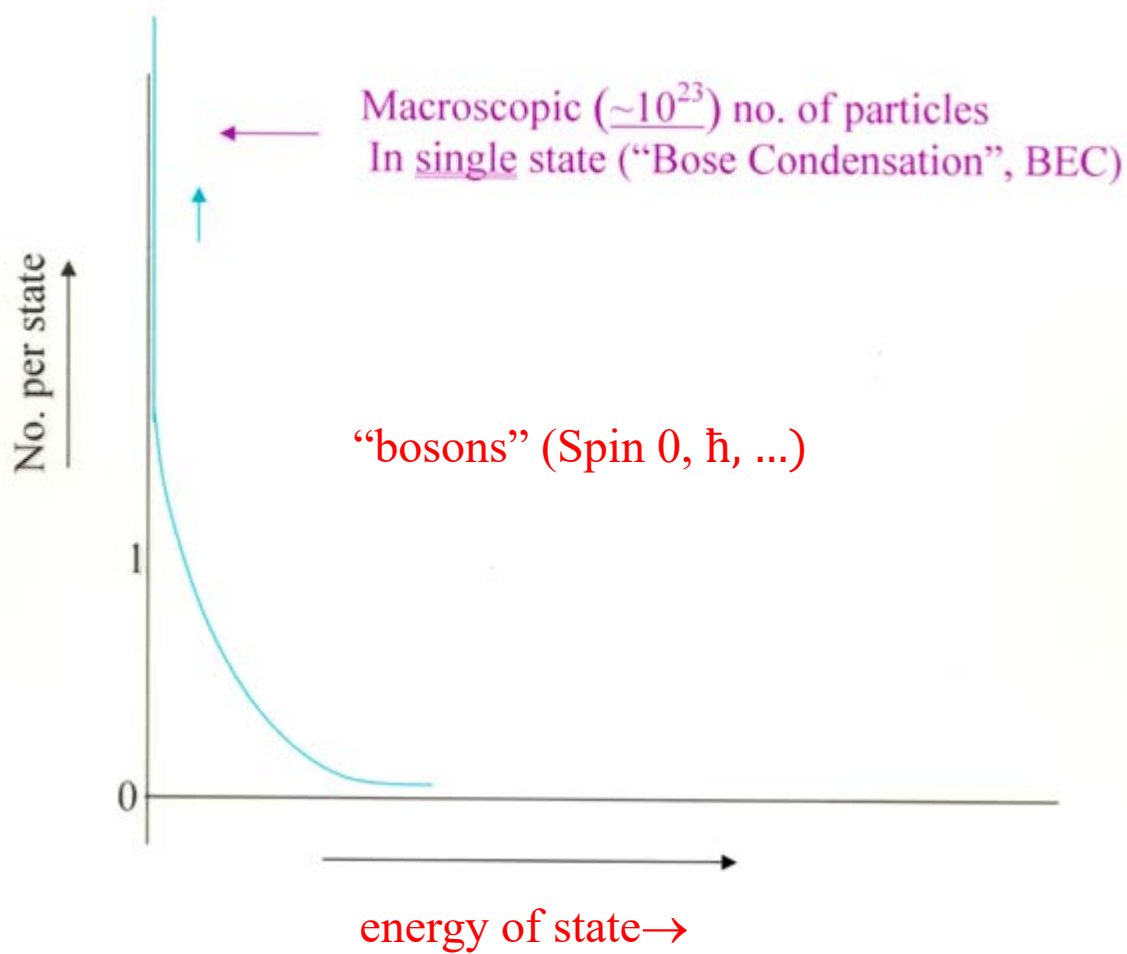
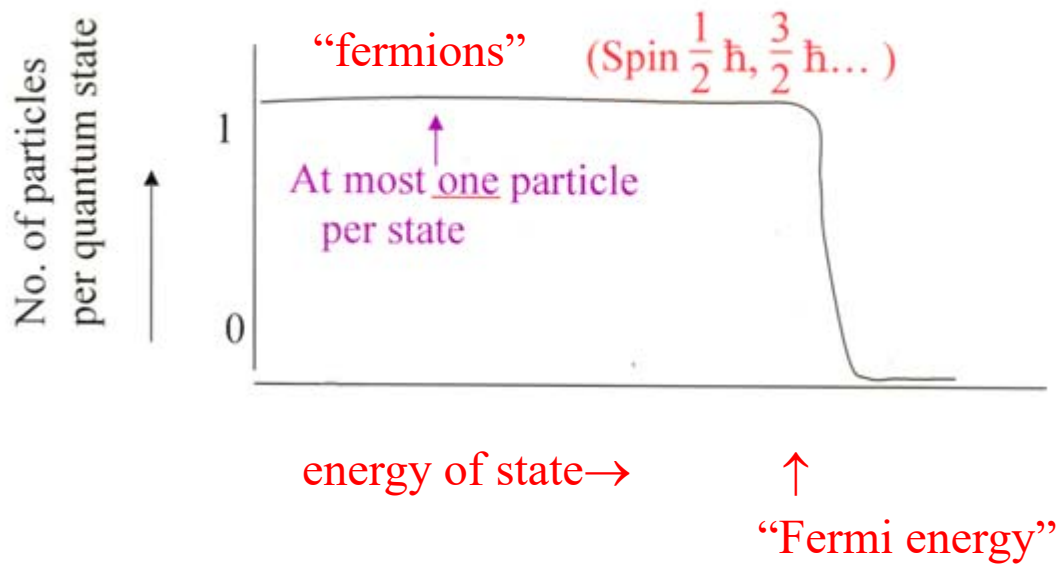
?

Evidently, for this property to be important, **must be able to change places**



Result of indistinguishability:

“QUANTUM STATISTICS”

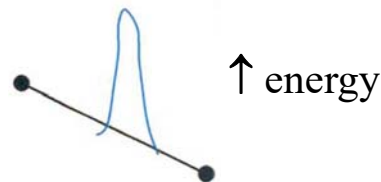
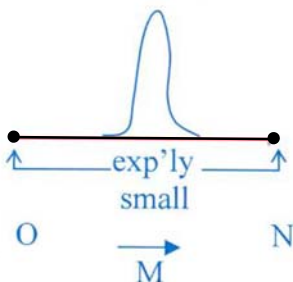


WHY BEC?

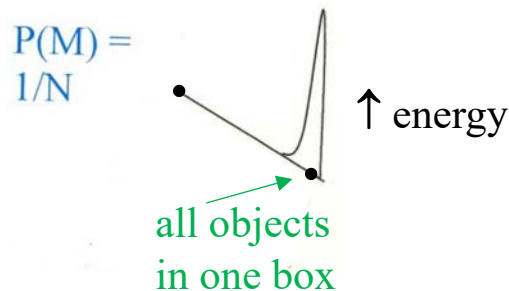
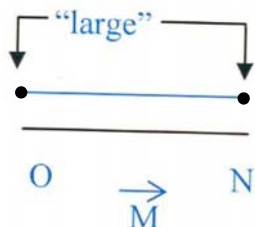
I. Qualitative argument:

Distribute N objects between 2 boxes: what is probability $P(M)$ of finding M in one box?

A. Objects distinguishable
 (\equiv coin toss):
 $P(M) = \frac{N!}{M! N-M!}$



B. Objects indistinguishable
 (bosons):



II. Quantitative argt. (Einstein, 1925):

$$n_i(T) = [\exp(\epsilon_i - \mu(T)/k_B T - 1)]^{-1}$$

chemical potential, ≤ 0

$$\mu(T) \text{ fixed by: } \sum_i n_i(T; \mu(T)) = N \leftarrow \text{total no. of particles}$$

$T \rightarrow \infty \Rightarrow \mu \rightarrow -\infty$; $T \downarrow \Rightarrow \mu \uparrow$. But what if

$$\sum_i [\exp(\epsilon_i/k_B T) - 1]^{-1} < N?$$

Einstein: Macroscopic no. of particles occupy lowest ($\epsilon=0$) state!

Condition for this to happen: roughly, $T \lesssim h^2/2mk_B a^2$ (as above)





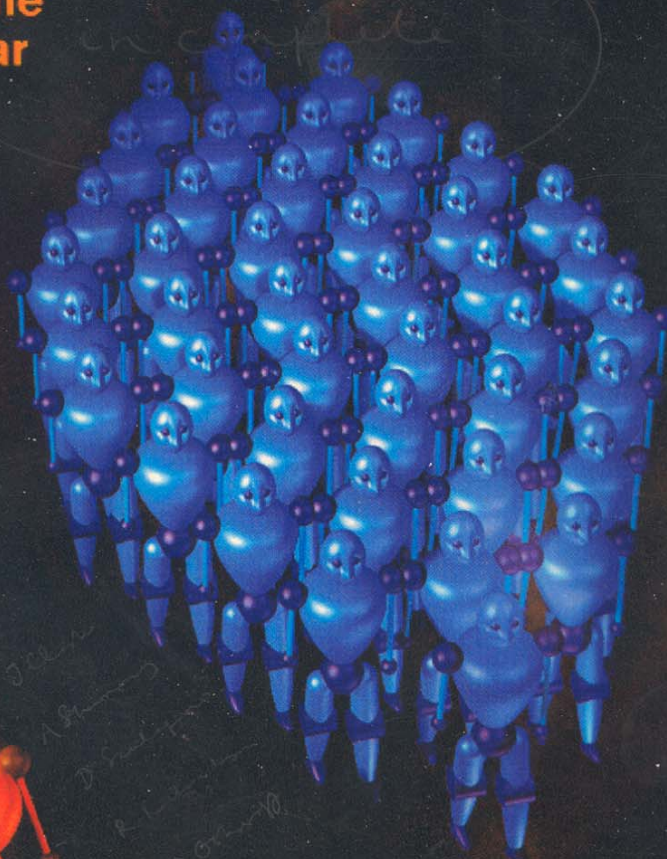
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ASSOCIATION FOR THE
ADVANCEMENT OF
SCIENCE

SCIENCE

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VOL. 270 • PAGES 1893-2064

\$7.00

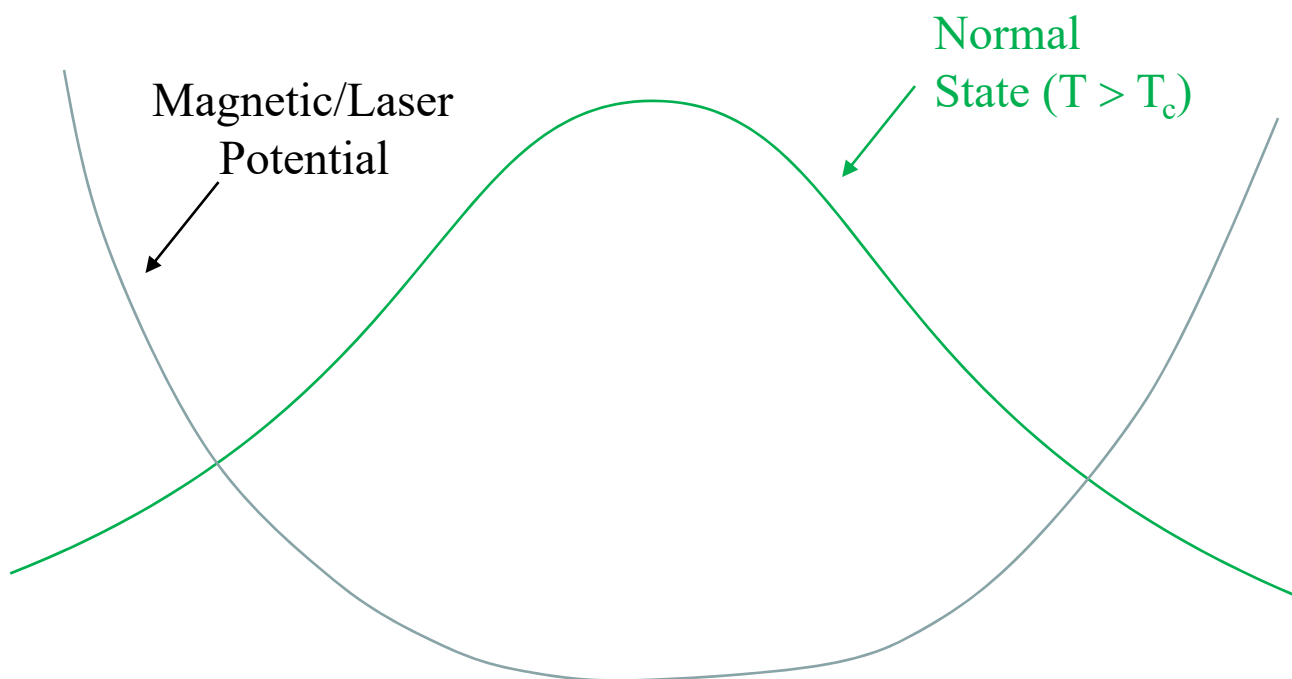
Molecule
of the
Year

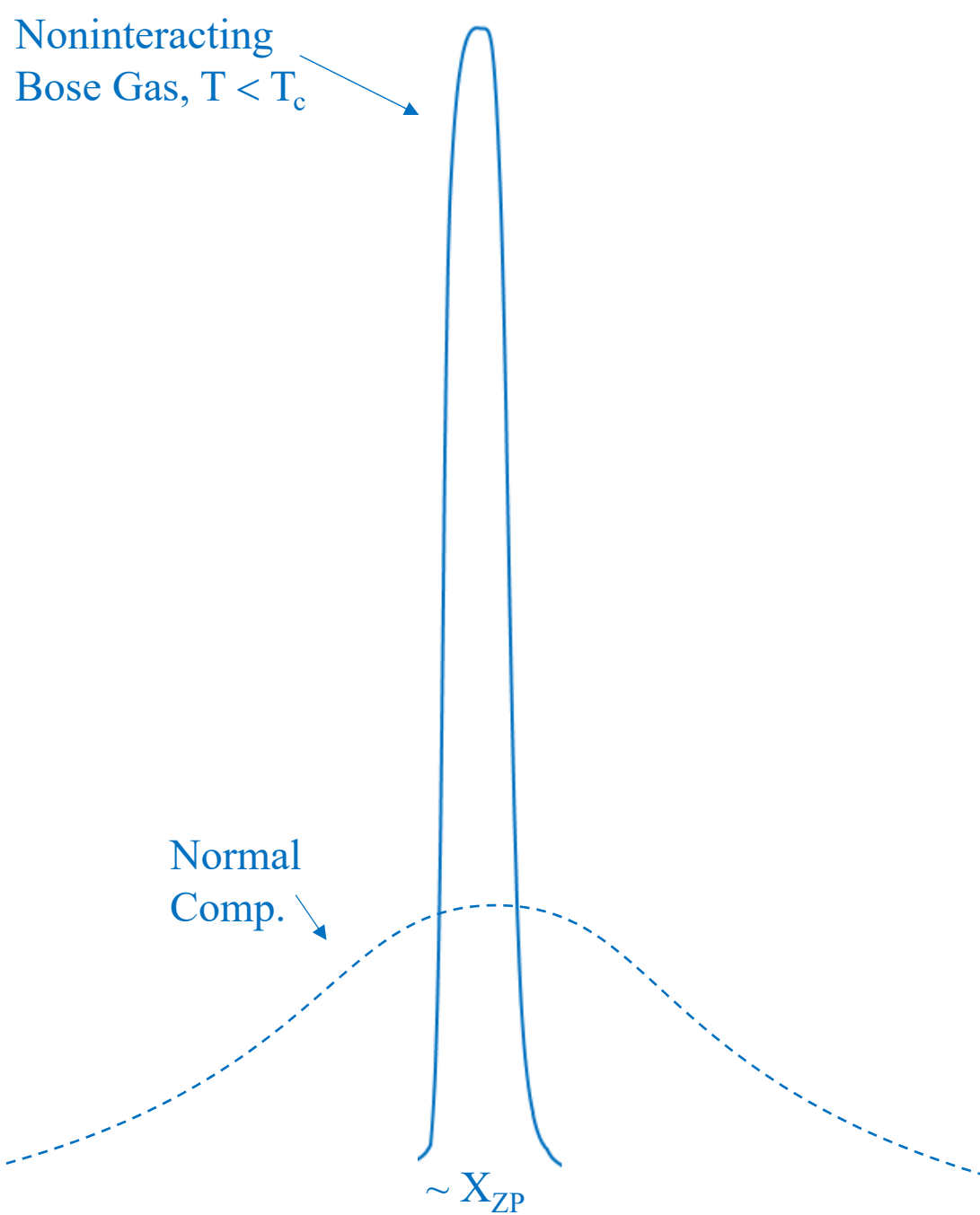


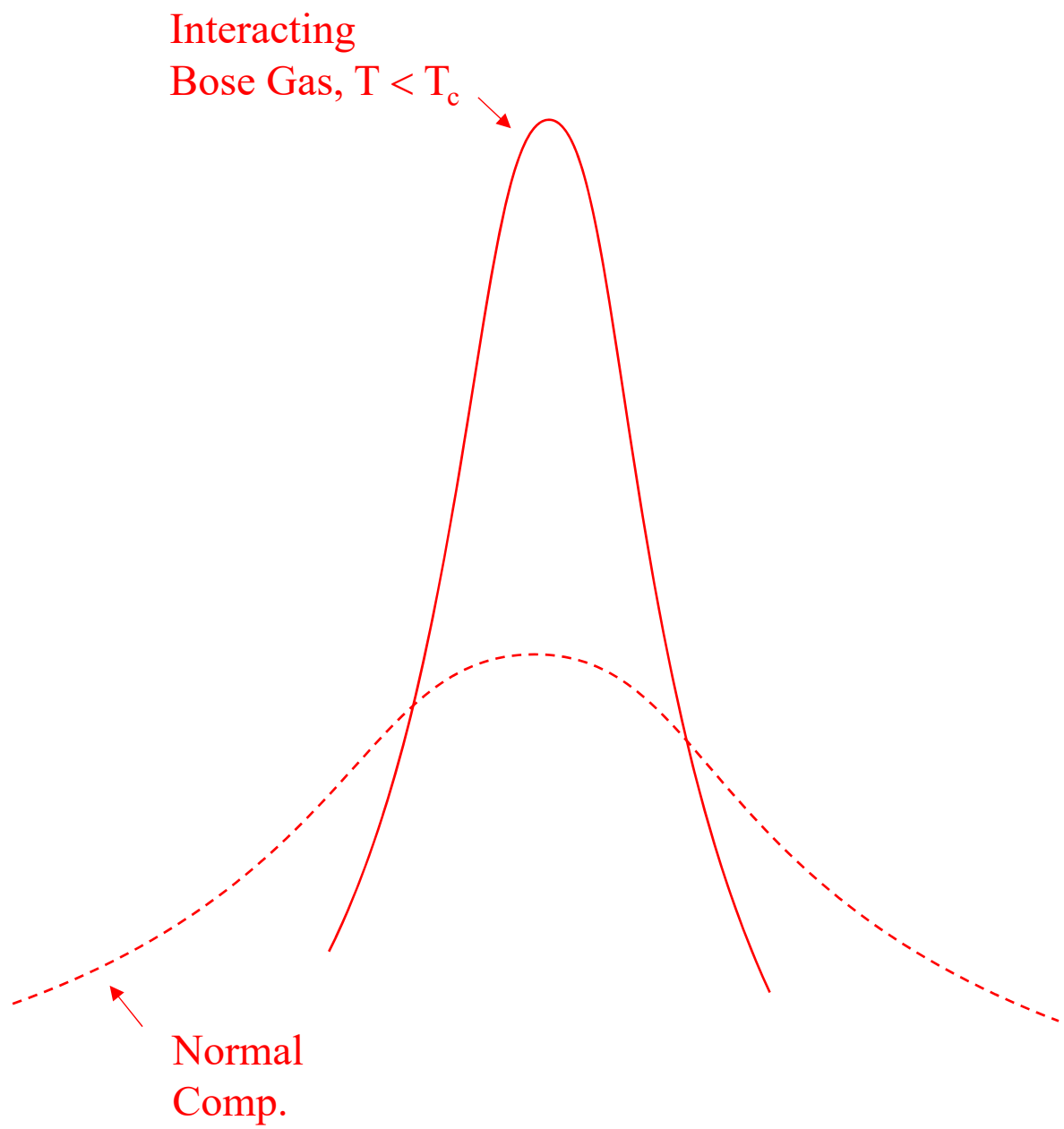
the
Bose-Einstein
Condensate

HOW TO SEE BEC OCCURRING?

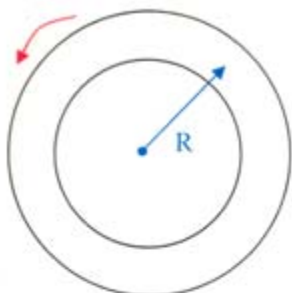
↑
LITERALLY







“NO-ROTATION” EFFECT IN LIQUID ⁴HE



Walls rotating with ang. velocity

$\omega \lesssim \omega_c \Leftrightarrow \equiv \hbar/m R^2$ (typically ~ 1 revolution/hour!)
 What does liquid do?

General principle: Average ang. velocity of atoms ($\bar{\omega}$) as close as possible to ω

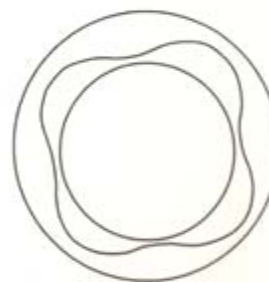
↑ : Single-atom states must obey quantization condition:

$$n\lambda = 2\pi R \quad + \text{d.B. } \lambda = h/mv$$

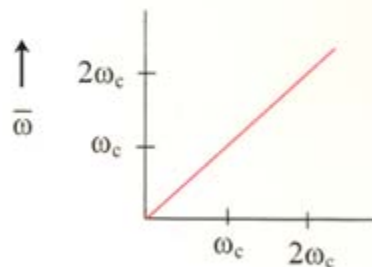
$$\Rightarrow L = mvR = n\hbar \quad (\hbar \equiv h/2\pi)$$

↙ ang. momentum

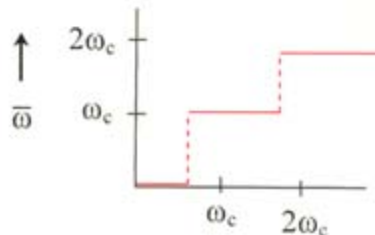
$$\Rightarrow L/mR^2 \equiv \bar{\omega} = n\omega_c$$



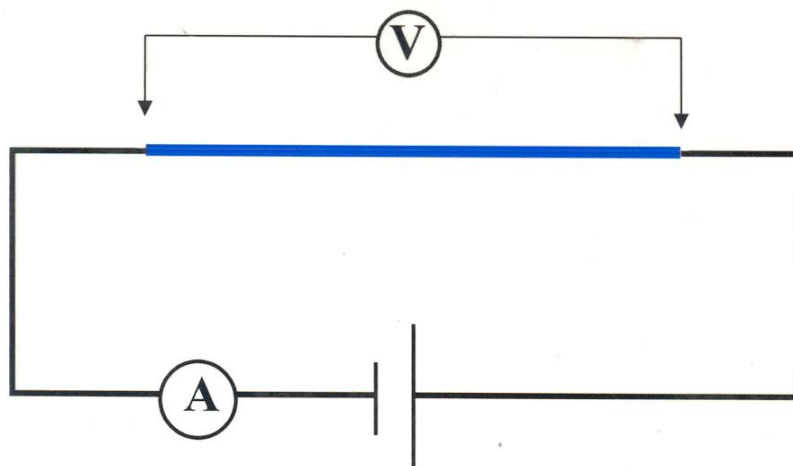
- A. “Normal” (non-BEC) system: many different single-particle states occupied (typical value of $n \sim (kT/\hbar\omega_c)^{1/2} \sim 10^7$)
 \Rightarrow to get $\bar{\omega} = \omega$, just shift atoms slightly between states.



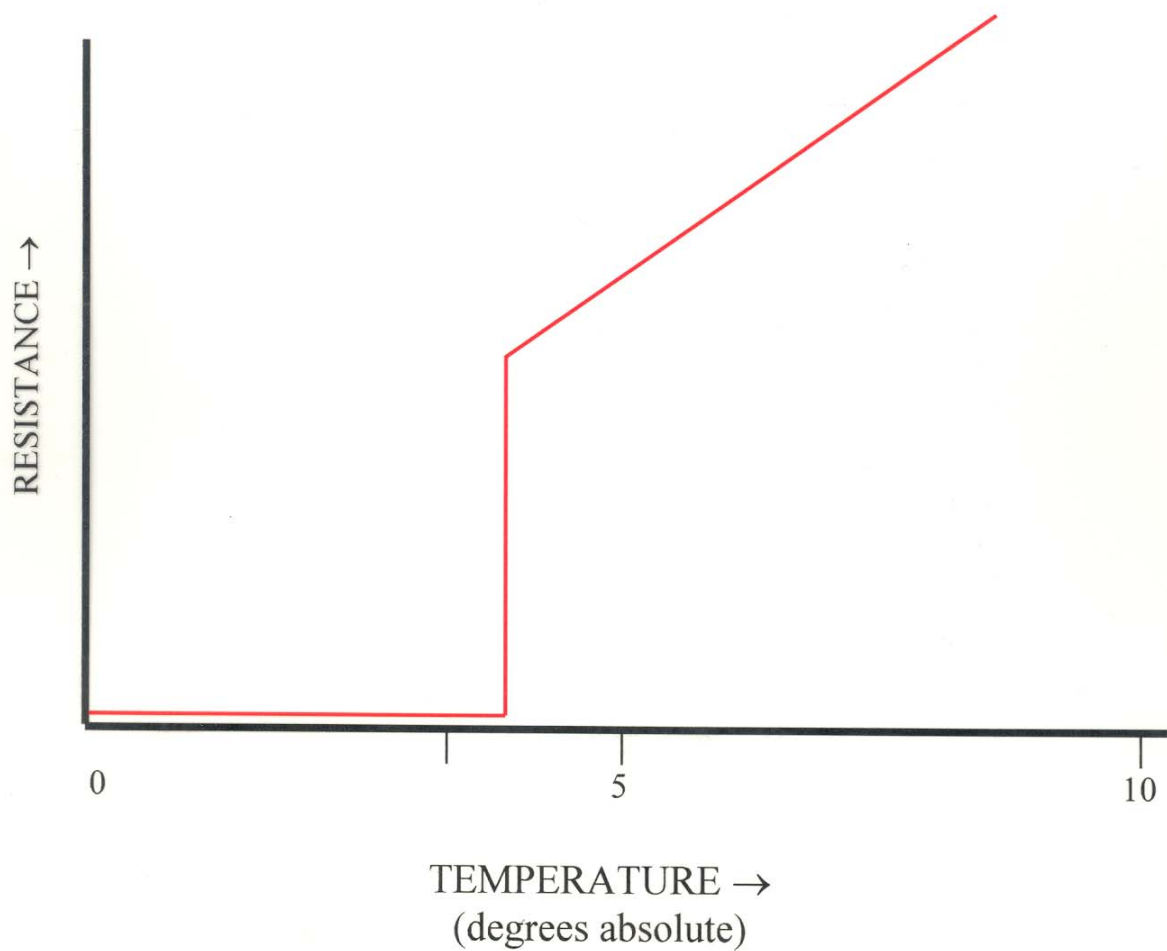
- B. BEC system ($T \ll T_c$): (almost) all atoms in condensate must have same value of n . (n_0) $\Rightarrow \bar{\omega} \cong n_0 \omega_c$



Superconductivity

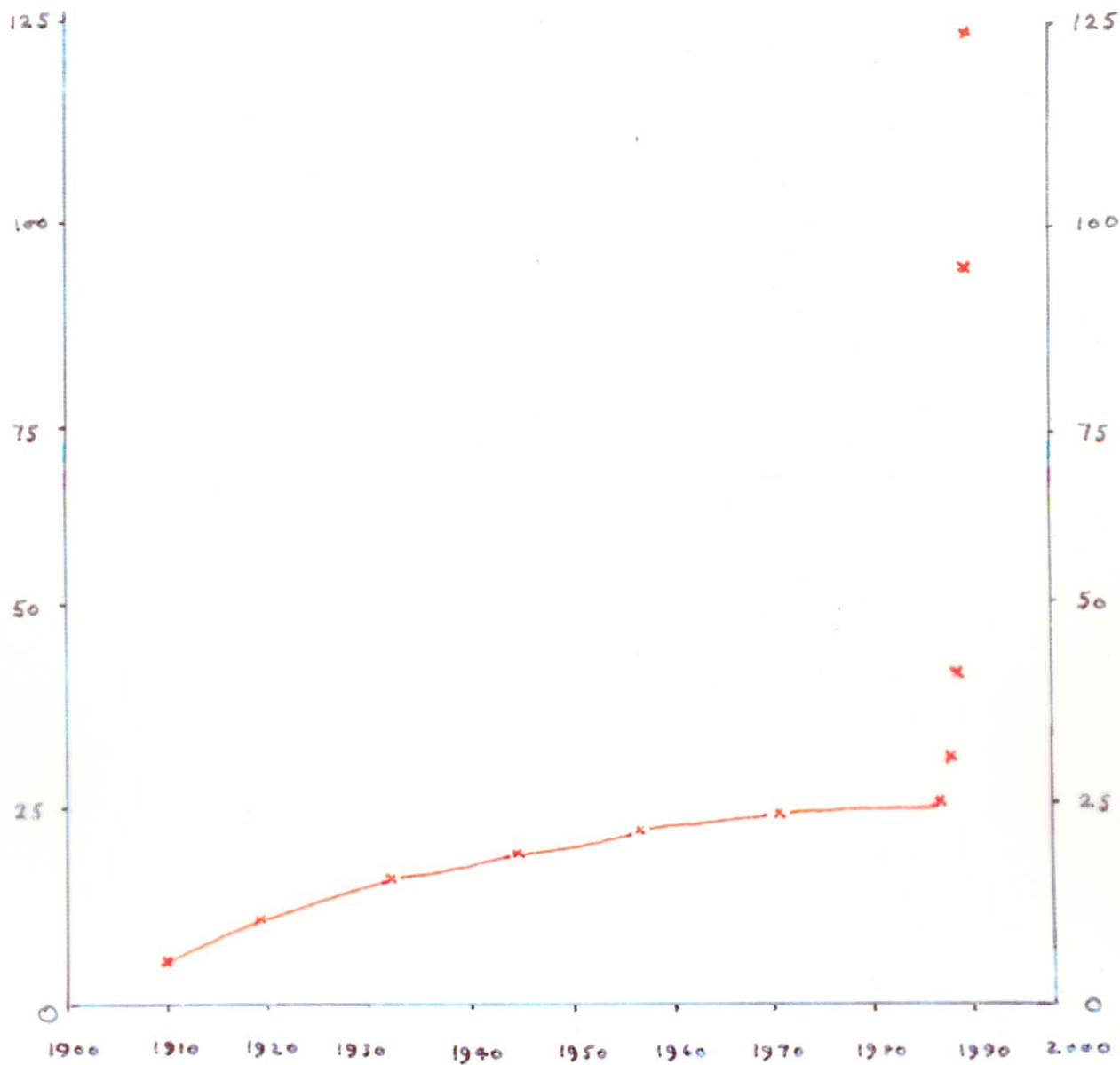


resistance of **—** = V/A = voltage/current



●
2014

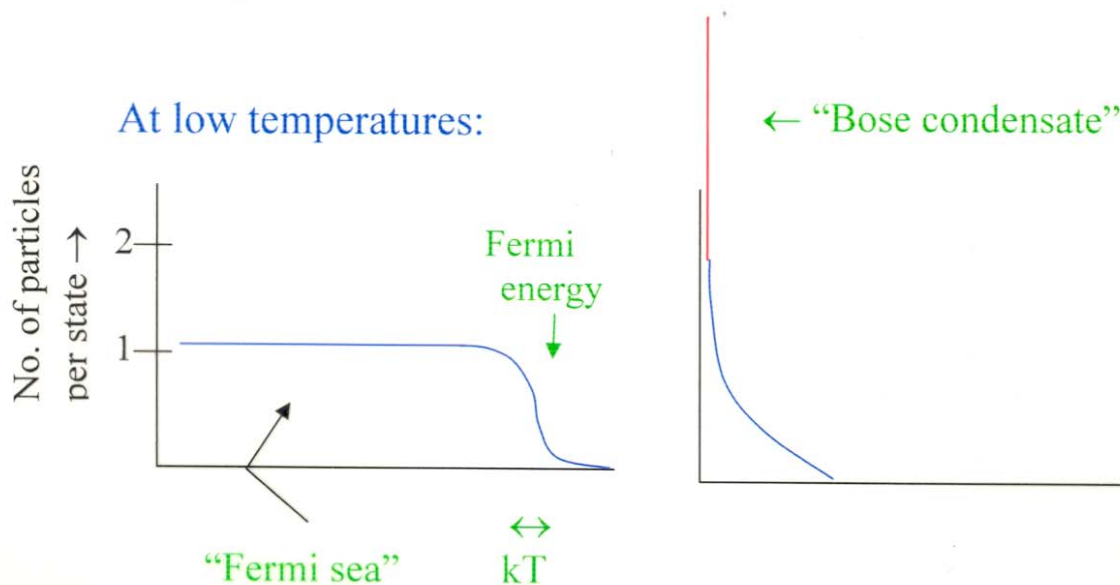
HISTORY OF THE HIGHEST TEMPERATURE
("T_c") AT WHICH SUPERCONDUCTIVITY KNOWN



PHYSICS OF SUPERCONDUCTIVITY

“Spin” of elementary particles = $\frac{n}{2} \hbar$

0, 1, 2, ... bosons
 $\frac{1}{2}, \frac{3}{2}, \frac{5}{2}, \dots$ fermions



Electrons in metals: spin $\frac{1}{2} \Rightarrow$ fermions

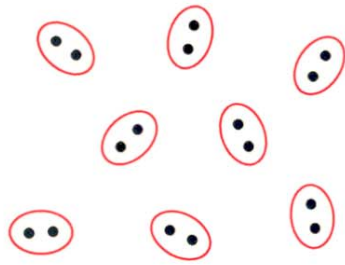
But a compound object consisting of an **even** no.

of fermions has spin 0, 1, 2 ... \Rightarrow boson.

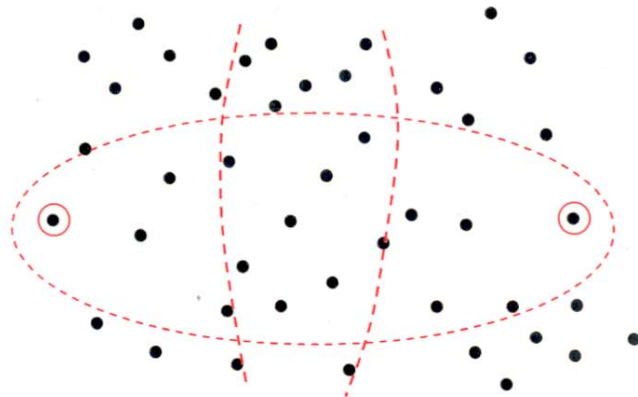
(Ex: $2p + 2n + 2c = {}^4\text{He}$ atom)

\Rightarrow can undergo Bose condensation





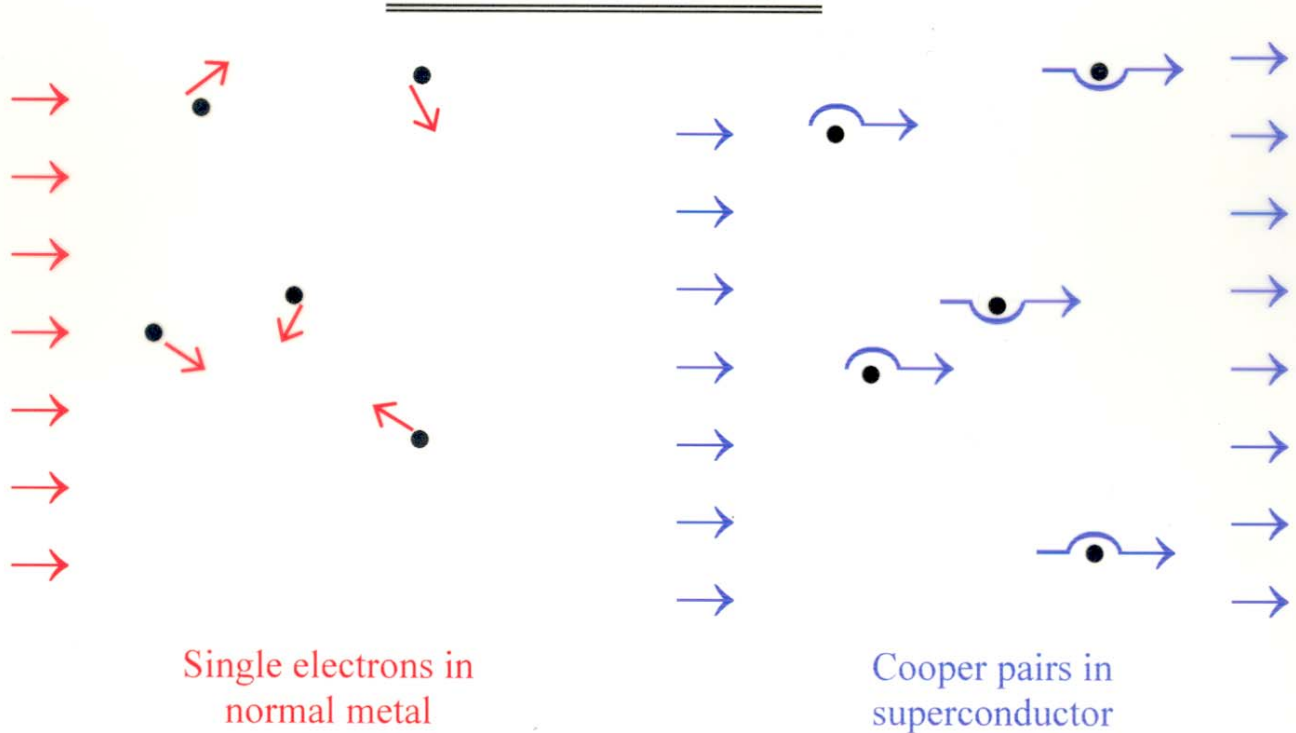
“di-electronic molecules”



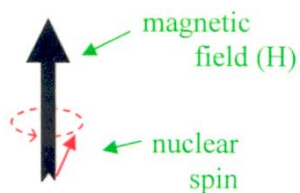
Cooper Pairs

In simplest (“BCS”) theory, Cooper pairs, once formed, must automatically undergo Bose condensation!

⇒ must all do exactly the same thing at the same time (also in nonequilibrium situation)

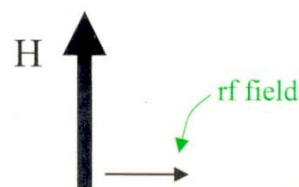


NUCLEAR MAGNETIC RESONANCE

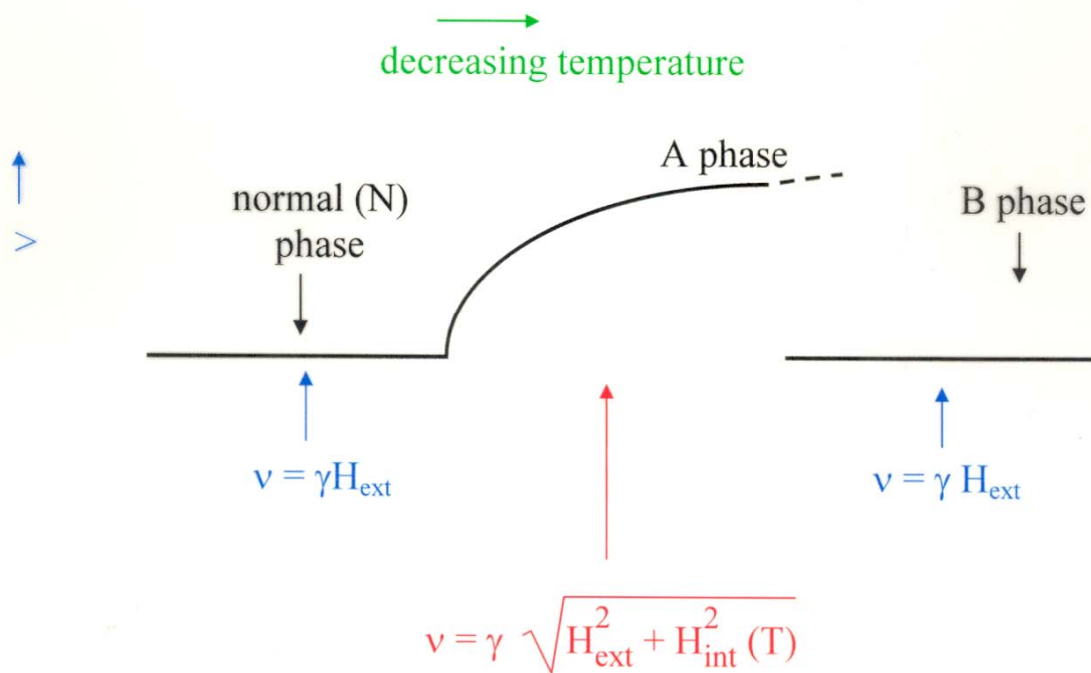


Rate of "precession"
 $\nu = \gamma H$
 "gyromagnetic ratio"

γ is known, (in ^3He , ~ 3 kHz/gauss)
 so, rate of precession (ν) measures magn. field (H)
 To measure ν , apply
 oscillating (r.f.) field $\perp H$:
 field is strongly absorbed when its frequency is ν .



NMR IN LIQUID ^3He BELOW 3mK:

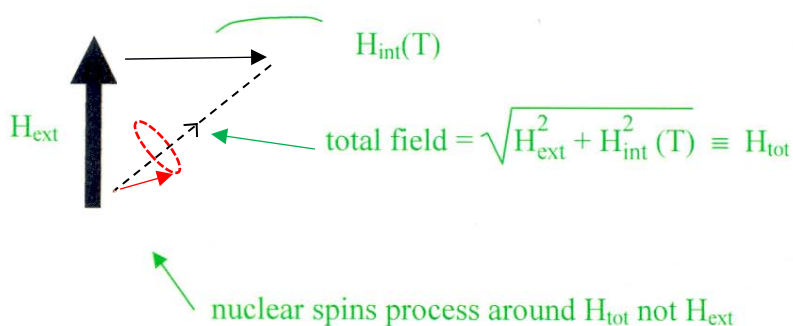


THE ^3HE NMR PUZZLE (cont.)

In A phase, precession freq. ν is larger than value (γH_{ext}) in N phase, and given by expression of form

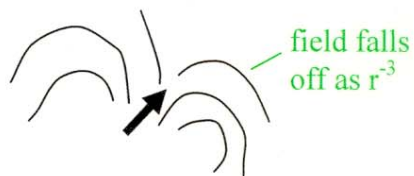
$$\nu = \gamma \sqrt{H_{\text{ext}}^2 + H_{\text{int}}^2} \text{ (T)}$$

Simplest interpretation:



Problem:

Only possible origin of $H_{\text{int}}(T)$ is other nuclear spins



Max. value of field of one nuclear spin on another
(at distance of closest approach of atoms) < 1 gauss.

But, experimental value of $H_{\text{int}}(T)$ is ~ 30 gauss!

**FIRST EVIDENCE FOR BREAKDOWN
OF QUANTUM MECHANICS?**



RESULT OF MORE SOPHISTICATED APPROACH:

A. Simple classical argument too naive.
(no “transverse” internal field)

B. Nevertheless, indeed predict formula

$$\nu = \gamma \sqrt{H_{\text{ext}}^2 + H_o^2(T)}$$

where $H_o^2(T)$ is proportional to average value of nuclear dipole interaction energy $E_{\text{dip}}(T)$.

Experimental value of $H_o(T) \rightarrow E_{\text{dip}}(T) \sim 10^{-3}$ ergs/cm³

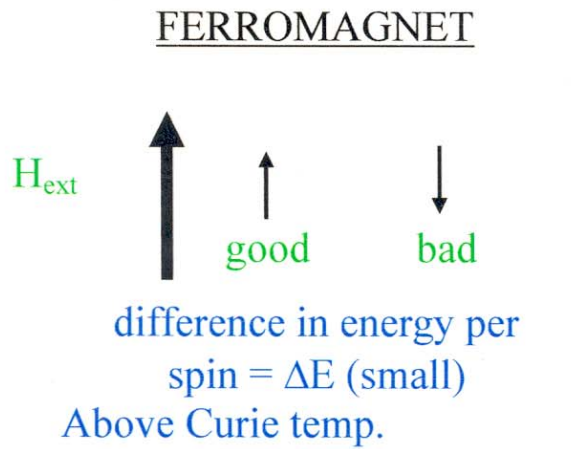
Why is this a problem?



- energy difference (ΔE) between “good” and “bad” orientations $< 10^{-7}$ K per pair.
- thermal energy (E_{th}) ($= k_B T$) $\sim 10^{-3}$ K.
 \Rightarrow preference for “good” orientation over “bad”
 only $\sim \Delta E/E_{\text{th}} < 10^{-4}$
 \Rightarrow resulting value of $E_{\text{dip}}(T)$ **much too small to fit experiment.**
 Need preference for “good” over “bad” ~ 1 not $\sim \Delta E/E_{\text{th}}!$

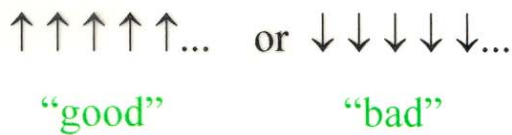
SPONTANEOUSLY BROKEN SPIN-ORBIT SYMMETRY:

the analogy with ferromagnetism



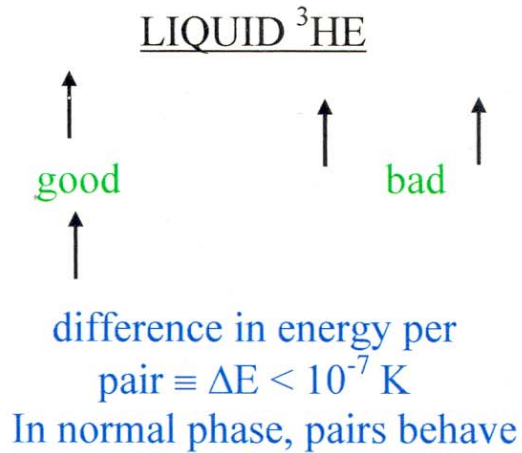
(“paramagnetic” phase), spins behave independently \Rightarrow thermal energy E_{th} competes with $\Delta E \Rightarrow$ polarization only $\sim \Delta E/E_{eth} \ll 1$

Below T_c (“ferromagnetic” phase): strong (exchange) forces **constrain all spins to lie parallel:**



$$E_{good} - E_{bad} \sim N\Delta E \gg E_{th}$$

\Rightarrow polarization ~ 1



independently $\Rightarrow E_{th}$ competes with $\Delta E \Rightarrow$ “polarization” (pref. for good orientation over bad) only $\sim \Delta E/E_{th} \ll 1$.

In A phase, **assume:** strong (kinetic-energy, VDW) forces **constrain all pairs to behave in same way** \Rightarrow either all “good” or all “bad”

$$E_{good} - E_{bad} \sim N \Delta E \gg E_{th} \quad \sim 10^{23}!$$

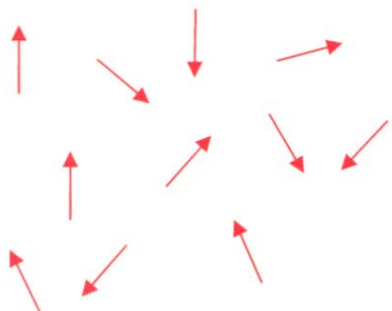
\Rightarrow polarization can be ~ 1



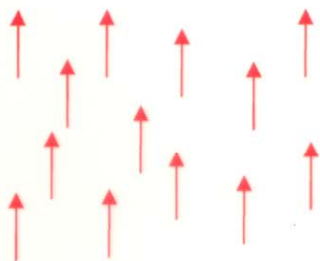
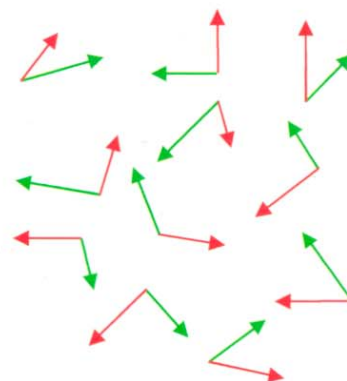
SBSOS: ORDERING MAY BE SUBTLE

FERROMAGNET

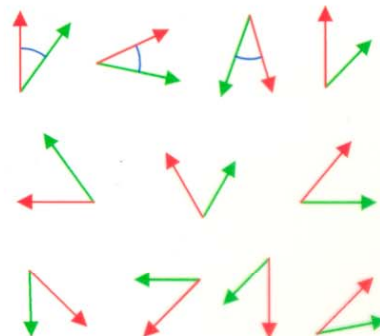
LIQUID ³HE



⇐ NORMAL PHASE ⇒



⇐ ORDERED PHASE ⇒



(= total spin of pair
 = relative orbital ang. momentum)

$\langle \tilde{S} \rangle \neq 0$

$\langle \tilde{S} \rangle = \langle \tilde{L} \rangle = 0$
 but $\langle \tilde{L} \times \tilde{S} \rangle \neq 0!$



Amplification of ultra-weak effects by BEC (cf NMR):

Example: P- (but not T-) violating effects of neutral current part of weak interaction:

For single elementary particle, any EDM \underline{d} must be of form

$$\underline{d} = \text{const. } \underline{J} \quad \leftarrow \text{violates T as well as P.}$$

But for ${}^3\text{He} - \text{B}$, can form

$$d \sim \text{const. } \underline{L} \times \underline{S} \sim \text{const. } \hat{\omega}$$



violates P but not T.

Effect is tiny for single pair, but since all pairs have same value of

$\underline{L} \times \underline{S}$, is multiplied by factor of $\sim 10^{23} \Rightarrow$

macroscopic P-violating effect?

