



Basic idea of Fritz London:

dissipation-less motion

 $\longleftrightarrow$ 

macroscopic wave function

Einstein 1924 Bose 1925 London 1938

#### a) Ideal Bose gas

non-interacting Bose gas (rough approximation for liquid He)

let's consider: 1 cm<sup>3</sup> cube of liquid <sup>4</sup>He  $riangle 10^{22}$  atoms with mass m

eigenstates for free particles in a cube:

$$E_n=rac{\hbar^2}{2m}\left(rac{\pi}{L}
ight)^2n^2$$
 with  $n^2=n_x^2+n_y^2+n_z^2$ 

$$T=0$$
 all atoms are in the ground state  $E_{111}$  trivial!

But at finite temperatures?



consider energy difference between ground state and first excited state

$$\Delta E/k_{\rm B} = (E_{211} - E_{111})/k_{\rm B} \approx 2 \times 10^{-14} \,\rm K$$



if Boltzmann statistics would hold - no condensate at 1 K!!!

however, Bose-Einstein distribution is relevant here

$$f(E,T)=rac{1}{{
m e}^{(E-\mu)/k_{
m B}T}-1}$$
 chemical potential  $\ \mu=rac{\partial F}{\partial N}$ 

what we know: 
$$\mu < E_{111}$$
  $\longrightarrow$  otherwise, negative occupation

$$u \neq 0$$

 $\mu \neq 0$  since particle number conserved



## Occupation of ground state $E_{111} = 0$

$$f(0,T) = rac{1}{\mathrm{e}^{-\mu/k_{\mathrm{B}}T}-1}$$
 — occupation depends critically on  $\mu$ 

$$f(0,T o 0) o \infty \quad \text{if} \quad \mu o 0 \quad \text{faster than} \quad T o 0 \qquad \left[ \begin{array}{c} \frac{1}{e^0-1} o \infty \end{array} \right]$$

What is the temperature dependence of  $\mu(T)$ ?

for this let us consider a real, but non-interacting gas

$$\mu = -k_{\rm B}T \ln \left(\frac{V_{\rm A}}{V_{\rm Q}}\right)$$
 quantum volume  $V_{\rm Q} = \left(\frac{h}{\sqrt{2\pi m k_{\rm B}T}}\right)^3 = \lambda_{\rm B}^3$  thermal de Broglie wavelength 
$$\lambda_{\rm B}^3 = (8.7~{\rm \AA})^3 ~{\rm at}~1~{\rm K}$$
 
$$V_{\rm A} = V/N = (3.8~{\rm \AA})^3 ~{\rm in}~{\rm comparison}$$



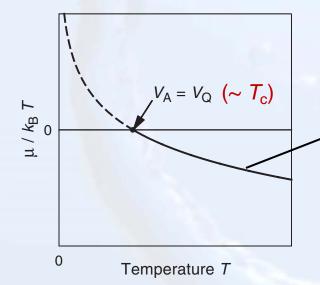
for 
$$T \to 0$$
  $\longrightarrow$   $V_{\rm Q} \to \infty$ 



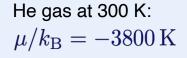
at sufficiently low temperatures  $\,V_{
m A} = V_{
m Q}\,$ 

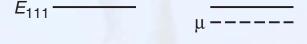
$$\ln\left(\frac{V_{\rm A}}{V_{\rm O}}\right) \to 0 \longrightarrow \mu \to 0$$





classical regime 
$$\mu$$
 is negative





$$T > 0$$
  $T \longrightarrow 0$ 

igoplus this means  $|\mu|$  becomes smaller than  $\Delta E/k_{
m B}=(E_{211}-E_{111})$  at finite T



Calculation of  $\mu$ : how large is  $\mu$  at 1K? (revers argument)

for 
$$T \to 0$$
  $\longrightarrow$   $f_{111} \to N$ 

$$\lim_{T \to 0} f(0,T) = N_0(T) = \lim_{T \to 0} \left( \frac{1}{e^{-\mu/k_{\rm B}T} - 1} \right)$$

 $E_{111} = 0$ , ground state

$$\approx \lim_{T\to 0} \left(\frac{1}{1-\mu/(k_{\rm B}T)+\ldots-1}\right) \approx -\frac{k_{\rm B}T}{\mu}$$

close to 
$$T=0$$

$$\rightarrow \text{ at } T = 1 \text{ K} \longrightarrow \mu/k_{\text{B}} \approx 10^{-22} \text{ K}$$



Calculation of  $\,N_0\,$  and  $\,N_{
m e}$  :

number of particles in excited states

$$\sum_i f(E_i,T) = N = N_0(T) + N_{\rm e}(T)$$
 
$$= N_0(T) + \int\limits_0^\infty D(E)\,f(E,T)\,{\rm d}E$$
 density of states for free particles without  $D(0)$ 

density of states for free particles  $\,E_k \propto k^2$ 

$$D(E) = \frac{V(2m)^{3/2} \sqrt{E}}{4\pi^2 \hbar^3}$$

with  $E/k_{\rm B}T=x$  and  $|\mu|\ll \Delta E$   $\longrightarrow$   $\exp[(E-\mu)/k_{\rm B}T]\approx \exp(E/k_{\rm B}T)$ 

$$N = N_0 + \frac{V}{4\pi^2} \left(\frac{2m}{\hbar^2}\right)^{3/2} (k_B T)^{3/2} \int_0^\infty \frac{\sqrt{x}}{e^x - 1} dx$$

$$\Gamma(5/2) \times \zeta(5/2) \approx 1.783$$





with 
$$V_{\mathrm{Q}} = \left(\frac{h}{\sqrt{2\pi m k_{\mathrm{B}} T}}\right)^{3} = \lambda_{\mathrm{B}}^{3}$$

$$N \approx N_0 + 2.6 \, \frac{V}{V_{\rm Q}}$$

$$N_0 = N - 2.6 \frac{V}{V_{\rm Q}}$$

Interpretation

as long as  $2.6 \frac{V}{V_{\rm O}} \ll 10^{22}$ , which means that the de Broglie wavelength is

significantly larger as an atom condensation



factor 
$$\sqrt[3]{2.6} = 1.37$$



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 $NV_{\mathsf{A}}$ 

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- $lacksquare T=0 \longrightarrow N_0=N$  trivial!
- $ightharpoonup 0 < T < T_c \longrightarrow N_0$  still macroscopically large!
- $ightharpoonup N_{
  m e} ext{ } ext$

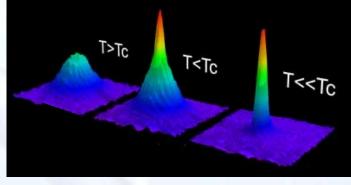
comment:

 $\lambda_{B}^{3}$  must not be as large as the vessel as proposed by London



#### What is the value of the condensation temperature?

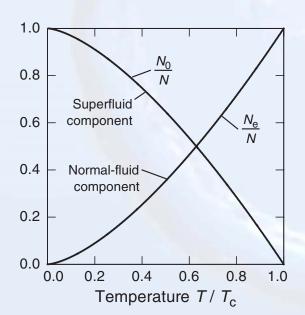
$$\left. egin{aligned} N_{0}(T_{
m c}) &= 0 \ N_{
m e}(T_{
m c}) &= N \end{aligned} 
ight. \qquad T_{
m c} = rac{2\pi\hbar^{2}}{k_{
m B}m} \left(rac{N}{2.6V}
ight)^{2/3} \ \end{array}$$



Bose Einstein condensate of atomic gas

gas  $T_{
m c}pprox 0.5\,{
m K}$  , but boiling point is at  $4.2\,{
m K}$ 

liquid  $\,T_{
m c} = 3.1\,{
m K}$  , works well in comparison to  $\,T_{\lambda} = 2.17\,{
m K}$ 



$$\frac{N_{\rm e}}{N} = \left(\frac{T}{T_{\rm c}}\right)^{3/2}$$

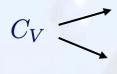
the condensation of a normal gas in real space corresponds to the Bose-Einstein condensation in momentum space, which means all atoms have the same wave vector and are strongly correlated.





## b) Interacting Bose "gas" (He)

specific heat



ideal Bose gas

experimental results

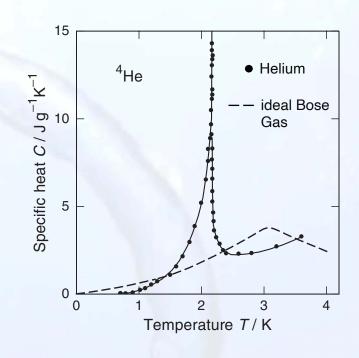
interactions are important



depletion of the ground state

collective excitation

(has first been proposed by Bogoliubov 1947)

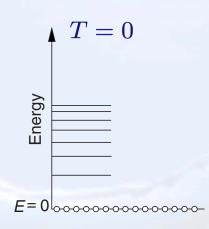


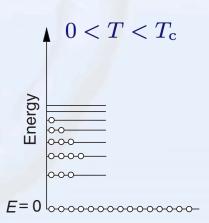
ideal Bose gas

$$T = 0$$
,  $N_0 = N$ 

$$T \neq 0$$
,  $N_0 \leq N$ 

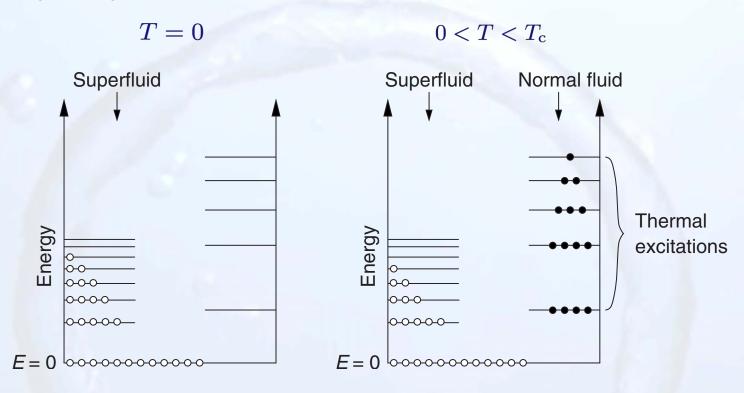
excited atoms







#### interacting Bose gas



T=0,  $N_0 \le N$ : significant number of atoms are not in the ground state

 $T \neq 0$ ,  $N_0 < N$ : in addition, collective excitations, nature of excitations changes



#### Experimental determination of the condensate

there is no direct way to measure the condensate fraction:  $N_0/N = n_0$ 

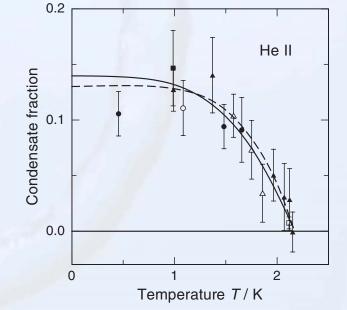
- $\omega$
- a) neutron scattering: measuring the dynamic structure factor  $S(Q,\omega) = n(p)$  via inelastic neutron scattering

momentum distribution

b) X-ray scattering: pair correlation function g(r) at transition to superfluid state becomes broader because of the condensation in momentum space

$$g(r) - 1 = (1 - n_0)^2 \left[g^*(r) - 1
ight]$$
  $g(r)$  above

c) surface tension: complicated but possible



condensate fraction for  $T \rightarrow 0$  just 13 %

Qs is not equal with condensate fraction