

SUPERCONDUCTIVITY.

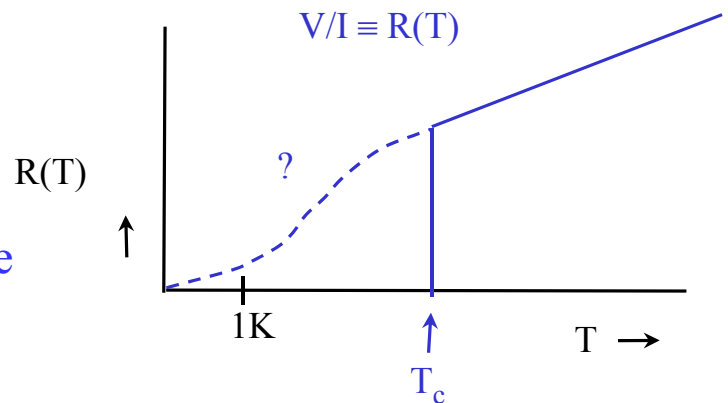
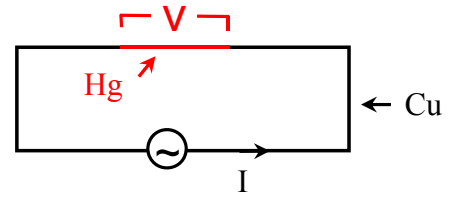
1. History:

Kamerlingh Onnes
(1911, Hg):

zero voltage drop

superconductivity \equiv absence
of resistance?

theoretical attempts 1911–1933



Meissner and
Ochsenfeld (1933):
total expulsion of flux
(**equilibrium** effect)

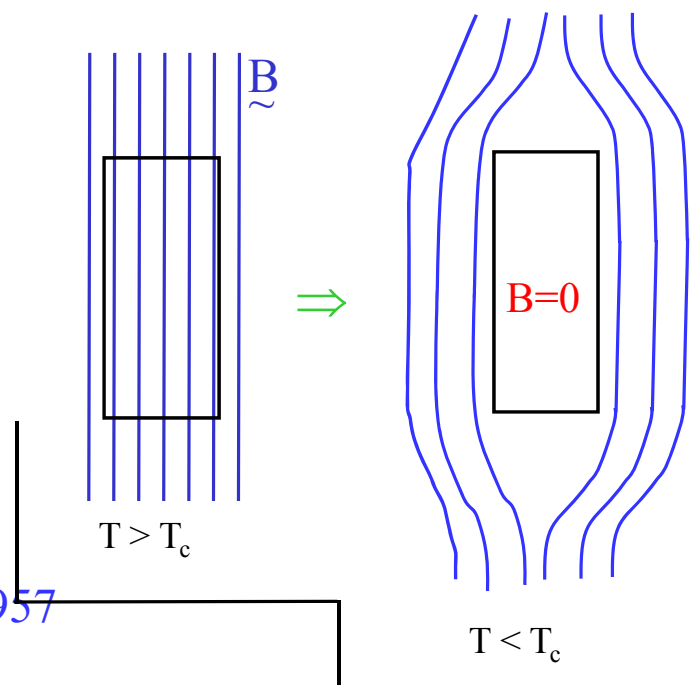
Londons 1935
isotope effect 1950
Ginzburg-Landau 1950

BCS theory (microscopic) 1957

Josephson effect 1962

[“exotic” superconductivity 1975, high-temperature (cuprate)
superconductivity 1986]

In these lectures, “superconductivity” always “classic” (BCS)



SUPERCONDUCTIVITY (cont.)

2. Phenomenology (classic, type-I)

Superconductivity sets in **abruptly** at temperature T_c (typically 1–20 K). Below T_c , superconducting state differs **qualitatively** from normal ($T > T_c$) state in 3 respects:

1. dc conductivity $\rightarrow \infty$ (e.g. persistent currents in ring)
2. magnetic flux completely expelled (**Meissner effect**)
3. Peltier coefficient $\rightarrow 0$

Occurrence:

metals, alloys, semiconductors

in metals, more towards middle of periodic table

Δ : “best” metals (Cu, Ag, Au) **not** superconductors

not sensitive to nonmagnetic impurities (e.g. many very “dirty”

alloys good superconductors with $T_c \sim 20\text{K}$), **very** sensitive to **magnetic** impurities.

Normal state ($T > T_c$) of superconducting metal essentially a “textbook” metal described by Sommerfeld-Bloch-Landau theory

Isotope effect: for a given (elemental) metal

$$T_c \propto M^{-1/2} \quad \text{isotopic mass}$$

\Rightarrow Dynamics of nuclei (i.e. phonons) must play a role

Microscopic properties in superconducting state:

| | | | |
|-------------------------|----------|---|--|
| specific heat | C_v | } $\Rightarrow 0$ for $T \rightarrow 0$ | Relative to extrapolated N-state values |
| spin susceptibility | χ | | |
| ultrasound attenuation | α | | |
| thermal conductivity | K | | |
| nuclear relaxation rate | T^{-1} | | |



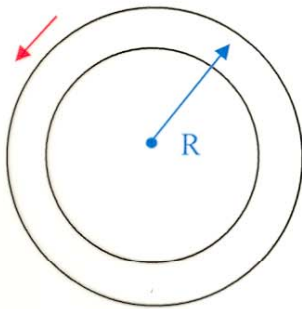
SUPERFLUIDITY IN LIQUID ^4He

| | | |
|---------------------------------------|------|--|
| ^4He liquefied: | 1908 | |
| $T < T_\lambda$ (2.17 K): | 1920 | \updownarrow ~ 20 YEARS! |
| Frictionless flow below T_λ : | 1938 | |

Modern point of view:

Define

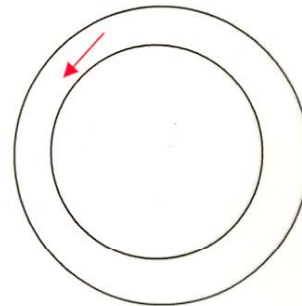
$\omega_c \equiv \hbar/mR^2 \equiv$ quantum unit of rotation ($\sim 10^{-4}$ Hz for $R \sim 1\text{cm}$)



EXPT. A
("Hess-Fairbank" effect)

walls rotate with
ang. velocity $\lesssim \omega_c$,
liquid stationary

EQUILIBRIUM
EFFECT



EXPT B
(Persistent currents)

walls at rest,
liquid rotating with
ang. velocity $\gg \omega_c$.

METASTABLE
EFFECT

BEC IN A NONINTERACTING BOSE GAS:
THE EFFECTS OF STATISTICS

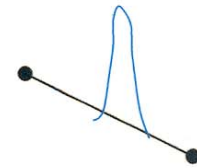
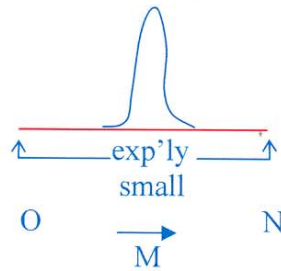
I. Qualitative argument:

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

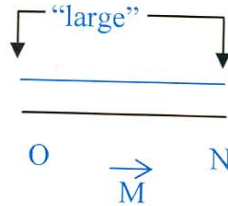
A. Objects distinguishable

(≡ coin toss):

$$P(M) = \frac{N!}{M! (N-M)!}$$



B. Objects indistinguishable (bosons):



$$P(M) = 1/N$$



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!



BEC IN A GENERAL (INTERACTING, NONEQUILIBRIUM) SYSTEM:

Can always find set of “single-particle” states $\chi_i(\mathbf{r},t)$ st. average
no. of atoms in state i is $n_i(t)$ (and $\langle a_i^\dagger a_j \rangle \equiv 0, i \neq j$)

Df of (“simple”) BEC:

one and **only** one single-particle state
 i (say $i = 0$) has $n_i(t) = O(N)$, rest $o(1)$

Then,

$N_0(t) \equiv$ “condensate number”

$\chi_0(\mathbf{r},t) \equiv$ “condensate wave function”

WHY BEC IN GENERAL CASE?

A. Statistics

B. Interactions (“Fock” term):

e.g. if $V(\mathbf{r}) = V_0 \delta(\mathbf{r})$:

$$N \text{ atoms in } \chi_0(\mathbf{r},t): \langle V \rangle(t) = \frac{1}{2} V_0 N^2 \cdot \int |\chi_0(\mathbf{r},t)|^4 d\mathbf{r}$$

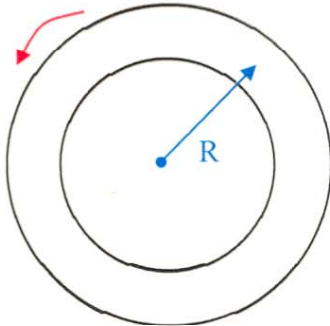
$$N_1 \text{ in } \chi_1, N_2 \text{ in } \chi_2: \langle V \rangle(t) = 2V_0 N_1 N_2 \int |\chi_1(\mathbf{r},t)|^2 |\chi_2(\mathbf{r},t)|^2 d\mathbf{r}$$

\Rightarrow if $V_0 > 0$, advantageous to have all in one state

what if $V_0 < 0$?



EXPLANATION OF HESS-FAIRBANK EFFECT IN TERMS OF BEC:



Walls rotating with ang. velocity

$$\omega \lesssim \omega_c \leftarrow \equiv \hbar/m R^2$$

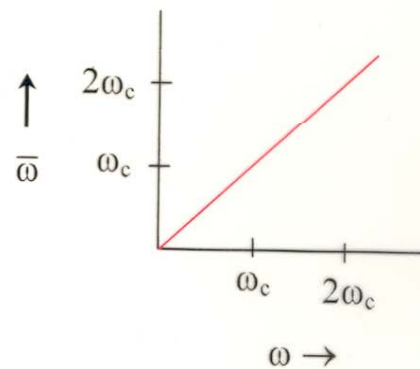
What does liquid do?

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

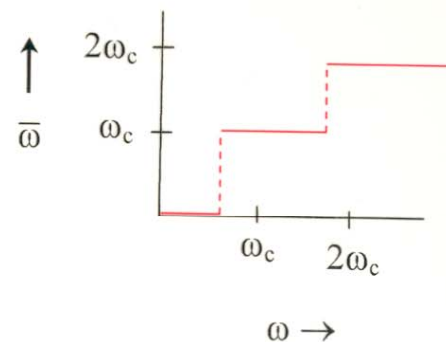
\uparrow : Single-atom states must obey
quantization condition: $\omega = n\omega_c$ ($\ell = n\hbar$)

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 \rightarrow must have **same**
value of n (n_0) $\Rightarrow \bar{\omega} \cong n_0 \omega_c$



INTERACTIONS
"OPTIONAL"

^4He : PERSISTENT CURRENTS

Initially, after walls stopped,

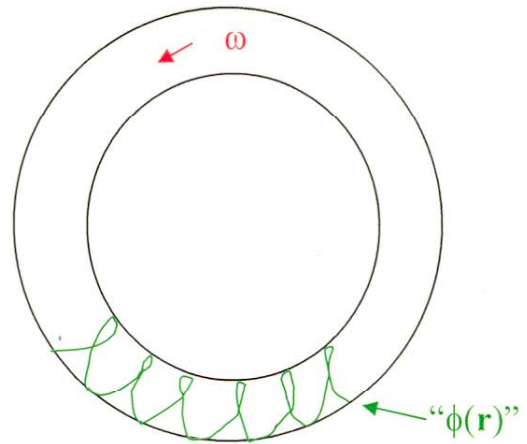
$$\langle L \rangle = N_0 \ell_0 \hbar, \quad \ell_0 \gg 1 \quad (\bar{\omega} \gg \omega_c)$$

But groundstate has $\langle L \rangle = 0$. ($\omega = 0$)

Why no relaxation?

$$\chi_0(\mathbf{r}) = |\chi_0(\mathbf{r})| \exp i \phi(\mathbf{r})$$

↑
condensate w.f.



$$\text{Df: "winding no." } n \equiv \oint \frac{\nabla \phi \cdot d\mathbf{l}}{2\pi}$$

Initially, $n = \ell_0$: eq^m state has $n = 0$.

To change n , must depress $|\chi_0(\mathbf{r})|$ to zero somewhere!

(a) Electron in atom:

Schrödinger eqn. **linear** \Rightarrow nodes cost no extra energy, e.g.

$$\psi(t) = a(t) \psi_p + b(t) \psi_s \quad \begin{cases} t \rightarrow -\infty: a=1, b=0 \\ t \rightarrow +\infty: a=0, b=1 \end{cases}$$

$$\langle E \rangle(t) = |a(t)|^2 E_p + |b(t)|^2 E_s = \text{monotonically decreasing}$$

(b) BEC (^4He):

$$\text{Extra term in energy: } \langle V \rangle = V_0 \int |\chi_0(\mathbf{r}, t)|^4 d\mathbf{r} \quad \begin{matrix} \nearrow > 0 \\ \text{red arrow} \end{matrix}$$

\Rightarrow energy **NOT** monotonically decreasing!

(REPULSIVE) INTERACTIONS ESSENTIAL!



CORRESPONDENCE BETWEEN SUPERFLUIDITY AND SUPERCONDUCTIVITY

- a) Persistent currents in ^4He in annular geometry \Leftrightarrow persistent currents in superconducting ring.
- b) What is superconducting analog of Hess-Fairbank effect?
- i. Behavior of superconductor under rotation (“London moment”)
 - ii. less obviously: behavior in magnetic field

Neutral system observed
from **rotating** frame

$$\begin{aligned}\hat{H} &= \hat{H}_0 - \underline{\omega} \cdot \hat{\underline{L}} \\ &\equiv \frac{1}{2m} \sum_i (p_i - m \underline{\omega} \times \underline{r}_i)^2 + \sum_i V(r_i) \\ &+ \frac{1}{2} \sum_{ij} U(r_i - r_j) \\ &- \frac{1}{2} \sum_i m (\underline{\omega} \times \underline{r}_i)^2 \leftarrow \text{centrifugal term,} \\ &\quad \text{affects only meniscus}\end{aligned}$$

So:

neutral system observed in
container rotating with
velocity $\underline{\omega}$ viewed from
rotating frame

\Leftrightarrow

charged system in magnetic
field \underline{B} , viewed from **lab**
frame

(with "scaling" $\underline{B} \Leftrightarrow \frac{e}{2m} \underline{\omega}$)

\Downarrow

HF effect: (part of) system at
rest in lab. frame \Rightarrow moving
in rotating frame

\Leftrightarrow

(part of) system **moving in**
lab frame (diamagnetism)

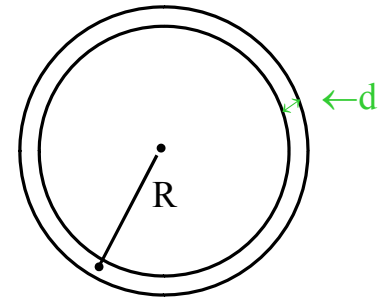


SUPERFLUID-SUPERCONDUCTOR CORRESPONDENCE (cont.)

Quantitative correspondence (T=0): consider in each case “thin” ring (d≪R) then

$$\underline{v} \equiv \underline{\omega} \times \underline{r} \cong \underline{\omega} \times \underline{R} \equiv R \underline{\omega} \times \hat{n} \neq f(r) \quad \text{and (prima facie!)} \quad \underline{A}(\underline{r}) \equiv \frac{1}{2} \underline{B} \times \underline{r} = \frac{1}{2} R \underline{B} \times \hat{n} \neq f(r)$$

Neutral case (T=0): as viewed from rotating frame,



mass
current
density $\rightarrow \underline{J} = -n m \underline{v}$

↑
particle density

So in charged case $\underline{J}_{el} = -\frac{ne^2}{m} \underline{A}$

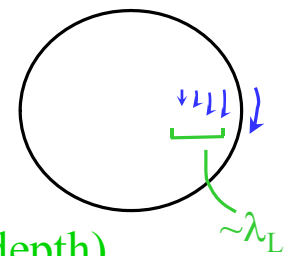
↑ electric current density ↓ vector potential

On a “sufficiently coarse-grained” scale, can interpret this as a **local** relation between \underline{J}_{el} and \underline{A} :

$$\underline{J}_{el}(\underline{r}) = -\frac{ne^2}{m} \underline{A}(\underline{r}) \quad \text{(London equation)}$$

But, in a bulk geometry, $\underline{A}(\underline{r})$ must be determined self-consistently from Maxwell’s equations, which in time-independent case $\Rightarrow \nabla^2 \underline{A}(\underline{r}) = -\underline{J}_{el}(\underline{r}) / \epsilon_0$. Hence,

$$\left. \begin{aligned} \nabla^2 \underline{A}(\underline{r}) &= \lambda_L^{-2} \underline{A}(\underline{r}) \\ \Rightarrow \nabla^2 \underline{B}(\underline{r}) &= \lambda_L^{-2} \underline{B}(\underline{r}) \end{aligned} \right\} \begin{aligned} \lambda_L &\equiv \left(\frac{m\epsilon_0}{ne^2} \right)^{-1/2} \\ &\equiv c / \omega_p \end{aligned}$$



and $\underline{B}(\underline{r})$ falls off (London penetration depth)

exponentially in bulk superconductor over distance $\sim \lambda_L$
 \Rightarrow Meissner effect.

- ⚠: 1) In multiply connected superconductor, London equation must be generalized (but $\underline{B}(\underline{r})$ still falls off exponentially)
- 2) London equation not quantitatively valid in type-I superconductors (\because not “sufficiently coarse-grained”)



HOW TO ADAPT (QUALITATIVE) IDEAS RE BEC TO

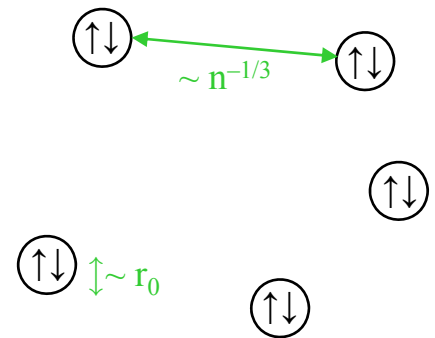
SUPERCONDUCTIVITY? (⚠: non-historical!)

Obvious problem in taking over idea of BEC directly: electrons in superconductor are fermions! So,

$$\rho_1(\underline{r}, \underline{r}') = \sum_i n_i \chi_i^*(\underline{r}) \chi_i(\underline{r}'), \quad n_i \leq 1 \quad \text{(Pauli principle)}$$

⇒ no BEC in literal sense. However: consider hypothetical dilute gas of diatomic molecules composed of 2 fermions (for simplicity with $\ell=S=0$), with $nr_0^3 \ll 1$

density molecular radius



(Ex: hypothetical gaseous D₂)

It is highly plausible that in the limit $nr_0^3 \rightarrow 0$, this system will behave just like a dilute gas of bosons (with spin 0).

Moreover, while the details of the molecule-molecule interactions depend on the short-range part of the potential, at least in the limit of “large” molecules there are strong arguments* that it should be repulsive. Thus the model is exactly that discussed above, and in the limit $T \rightarrow 0$ we expect

BEC OF DI-FERMIONIC MOLECULES

What does the many-body wave function of such a system look like? Answer:

$$\Psi(\underline{r}_1 \sigma_1 \underline{r}_2 \sigma_2 \dots \underline{r}_N \sigma_N) = \mathcal{A} \overbrace{\varphi(\underline{r}_1 \underline{r}_2 \sigma_1 \sigma_2) \varphi(\underline{r}_3 \underline{r}_4 \sigma_3 \sigma_4) \dots \varphi(\underline{r}_{N-1} \underline{r}_N \sigma_{N-1} \sigma_N)}^{\text{antisymmetrizer}} \quad \text{BEC!}$$

$$\varphi(\underline{r}_1 \underline{r}_2 \sigma_1 \sigma_2) = \frac{1}{\sqrt{2}} (\uparrow_1 \downarrow_2 - \downarrow_1 \uparrow_2) \varphi(|\underline{r}_1 - \underline{r}_2|)$$

spin singlet s-wave molecular state



*Petrov et al., PRL 93, 090404(2004): $a_{BB} = 0.6 a_{FF} (> 0)$.

HOW TO ADAPT BEC IDEAS...? (cont.)

2-body problem

Consider the behavior of two isolated fermions with some interatomic potential $V(|r_1 - r_2|)$ whose strength (and/or “shape”) can be varied, in a $\underline{K} = 0, S = \ell = 0$ state



As the potential is varied, its effects on the low-energy behavior are uniquely parametrized by the quantity a_s (**s-wave scattering length**), or more conveniently by a_s^{-1} :

For a strongly attractive potential $a_s^{-1} \rightarrow +\infty$: fermions form **tightly bound molecule** (radius \sim range of potential (w)).

As potential is weakened, a_s^{-1} decreases and eventually becomes negative.

For $a_s^{-1} > 0$ but $\ll w^{-1}$, fermions form **weakly bound molecule** (radius = $a_s \gg w^{-1}$), with binding energy $\varepsilon = -\hbar^2 / ma_s^2$.

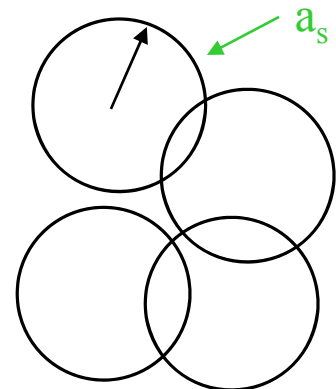
For $a_s^{-1} \rightarrow 0$ (“**unitarity**”) the energy of the molecular bound state $\rightarrow 0$, and for $a_s^{-1} < 0$ **no molecular state is possible**.

Now, back to the many-body problem: What happens if starting from a dilute BEC of di-fermionic molecules, we gradually weaken the inter-fermion attraction (while keeping $n = \text{const.}$)?

When the (2-body) s-wave scattering length a_s becomes $\sim n^{-1/3}$, “molecules” start to overlap \Rightarrow cannot neglect effects of Pauli principle. Equivalently,

$$|\varepsilon_F| \sim \frac{n^{2/3} \hbar^2}{m} \sim \frac{\hbar^2}{ma_s^2} \sim |E|.$$

When $a_s \gtrsim n^{-1/3}$, do “molecules” unbind?



DO MOLECULES UNBIND? (cont.)

BCS (1957): (A sort of) “molecules” persist for arbitrarily **weak attraction**, i.e. even for a_s -ve when no 2-body state is bound.

Unifying formalism (Yang, 1962):

General many-body pure state wave function:

$$\Psi_N^S(t) \equiv \Psi_S(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2 \dots \underline{r}_N\sigma_N : t)$$

2-body density matrix ρ_2 defined by

$$\rho_2(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2 : \underline{r}'_1\sigma'_1, \underline{r}'_2\sigma'_2) \equiv N(N-1) \sum_S p_S \sum_{\sigma_3\sigma_4\dots\sigma_N} \int d\underline{r}_3 d\underline{r}_4 \dots d\underline{r}_N$$

$$\Psi_S^*(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2, \underbrace{\underline{r}_3\sigma_3 \dots \underline{r}_N\sigma_N}_{\text{summed/integrated over}} : t) \cdot \Psi_S(\underbrace{\underline{r}'_1\sigma'_1, \underline{r}'_2\sigma'_2}_{\text{prob. of status}}, \underbrace{\underline{r}_3\sigma_3 \dots \underline{r}_N\sigma_N}_{\text{summed/integrated over}} : t)$$

$$\equiv \langle \psi_{\sigma_1}^\dagger(\underline{r}_1 t) \psi_{\sigma_2}^\dagger(\underline{r}_2 t) \psi_{\sigma'_2}(\underline{r}'_2 t) \psi_{\sigma'_1}(\underline{r}'_1 t) \rangle$$

$\hat{\rho}_2$ is Hermitian \Rightarrow can be diagonalized:

$$\rho_2(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2 : \underline{r}'_1\sigma'_1, \underline{r}'_2\sigma'_2 : t) = \sum_i n_i(t) \chi_i^*(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2 : t) \chi_i(\underline{r}'_1\sigma'_1, \underline{r}'_2\sigma'_2 : t)$$

Theorem (Yang): All $n_i \leq N$.

Ansatz: (In thermal equilibrium at $T <$ some T_c): For arbitrarily weak attraction, \exists **one and only one eigenvalue** $\sim N$, all others ~ 1 . (“ODLRO”). Call it N_0 , and corresponding χ χ_0

BEC limit ($a_s^{-1} \rightarrow +\infty$): $N_0 = N, \chi_0 =$ molecular wave function

BCS limit ($a_s^{-1} \rightarrow -\infty$): $N_0 \ll N$ (but $\sim N$), χ_0 “molecular-like” but radius $\gg n^{-1/3}$.

intermediate case (“unitarity”) ?? (“BEC-BCS crossover” in Fermi alkali gases)



QUALITATIVE ARGUMENT FOR MAIN PHENOMENA OF SUPERCONDUCTIVITY FROM YANG HYPOTHESIS

Bose case (recap):

$$\rho_1(\underline{r}_1, \underline{r}' : t) = \sum n_i(t) \chi_i^*(\underline{r}, t) \chi_i(\underline{r}' t)$$

in thermal equilibrium, \exists , **one** eigenvalue (N_0) $\sim N$ (“BEC”), with associated eigenfunction $\chi_0(\underline{r})$. Define order parameter

$$\Psi(\underline{r}) \equiv \sqrt{N_0} \chi_0(\underline{r}) \qquad \Psi(\underline{r}) \equiv |\Psi(\underline{r})| \exp i \varphi(\underline{r})$$

Then: a) Hess-Fairbank effect follows from BEC alone.

b) stability of supercurrents follows from BEC **plus** repulsive interactions, i.e. term in energy $\sim b |\Psi(\underline{r})|^4$, $b > 0$.

Fermi case:

$$\rho_2(r_1\sigma_1, r_2\sigma_2 : r'_1\sigma'_1, r'_2\sigma'_2 : t) = \sum_i n_i(t) \chi_i^*(r_1\sigma_1, r_2\sigma_2 : t) \chi_i(r'_1\sigma'_1, r'_2\sigma'_2 : t)$$

Assumption: in thermal equilibrium \exists **one** eigenvalue (N_0) $\sim N$, with associated eigenfunction $\chi_0(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2)$.

Write

$$\chi_0(\underline{r}_1\sigma_1, \underline{r}_2\sigma_2) \equiv \chi_0(\underline{r}_1 - \underline{r}_2, \sigma_1\sigma_2 : \underline{r})$$

relative
COM
coordinate
coordinate

and fix $(\underline{r}_1 - \underline{r}_2, \sigma_1\sigma_2)$ at some “standard” values (e.g. for s-wave, $(\underline{r}_1 - \underline{r}_2 = 0, \sigma_1 = -\sigma_2 = \uparrow)$). Then $\chi_0 \equiv \chi_0(\underline{r})$, and can define similarly to Bose case an order parameter

$$\Psi(\underline{r}) \equiv \sqrt{N_0} \chi_0(\underline{r}) \qquad \Psi(\underline{r}) \equiv |\Psi(\underline{r})| \exp i \varphi(\underline{r})$$

So, arguments go through similarly to Bose case, **provided** \exists a term in energy of form $\sim b |\Psi(\underline{r})|^4$, $b > 0$.

(Note: $\Psi(\underline{r})$ is essentially the order parameter introduced in the phenomenological theory of Ginzburg and Landau, without an appreciation of its microscopic meaning.)

