

Chapter 4 : FERMI SYSTEMS

Review

We studied the degenerate electron gas (or, *jellium*) in Section 3, and homework problem 6. There the discussion was limited to first order perturbation theory.

Now using QFT we'll consider a more accurate result.

12. The DEGENERATE ELECTRON GAS

Consider particles that interact by a spin-independent potential,

$$V(\vec{x}, \vec{x}')_{\lambda\lambda', \mu\mu'} = V(\vec{x} - \vec{x}') \delta_{\lambda\lambda'} \delta_{\mu\mu'}$$

$$V_0(x, x')_{\lambda\lambda', \mu\mu'} = V(\vec{x} - \vec{x}') \delta_{\lambda\lambda'} \delta_{\mu\mu'} \delta(t - t')$$

This is



Then

$$\langle \hat{V} \rangle = \frac{1}{2} \int d^3x d^3x' V(\vec{x} - \vec{x}') \langle \Phi | \psi_\alpha^\dagger(\vec{x}) \psi_\beta^\dagger(\vec{x}') \psi_\beta(\vec{x}') \psi_\alpha(\vec{x}) | \Phi \rangle$$

$$\text{Now, let } n(\vec{x}) = \psi_\alpha^\dagger(\vec{x}) \psi_\alpha(\vec{x})$$

$$\langle \hat{V} \rangle = \frac{1}{2} \int d^3x d^3x' V(\vec{x} - \vec{x}') \left\{ \langle n(\vec{x}) n(\vec{x}') \rangle - \delta^3(\vec{x} - \vec{x}') \langle n(\vec{x}) \rangle^2 \right\}$$

Or, let $\tilde{n}(\vec{x}) = n(\vec{x}) - \langle n(\vec{x}) \rangle$.

For a uniform system,

$$\langle n(\vec{x}) \rangle = n \quad (= N/V)$$

Then

$$\langle \hat{V} \rangle = \frac{1}{2} \int d^3x d^3x' V(\vec{x} - \vec{x}')$$

$$\left[\langle \tilde{n}(\vec{x}) \tilde{n}(\vec{x}') \rangle - n^2 - \delta^3(\vec{x} - \vec{x}') n \right]$$

"density correlation function"

Define a time ordered density correlation function

$$iD(x, x') = \langle \Psi_0 | T [\tilde{n}(x) \tilde{n}(x')] | \Psi_0 \rangle$$

also

$$iD_0(x, x') = \langle \Phi_0 | T [\tilde{n}_I(x) \tilde{n}_I(x')] | \Phi_0 \rangle$$

↪ interaction picture

↪ n_H : Heisenberg picture

Then

$$\langle \hat{V} \rangle = \langle \Phi_0 | \hat{V} | \Phi_0 \rangle$$

↪ This part was calculated in section 3.

$$+ \frac{1}{2} \int d^3x d^3x' V(\vec{x} - \vec{x}')$$

$$[iD(\vec{x}'t, \vec{x}t) - iD^0(\vec{x}'t, \vec{x}t)]$$

Thus the ground state energy is

$$E = \langle \Phi_0 | \hat{H} | \Phi_0 \rangle \quad \leftarrow \text{known from section 3}$$

$$+ \frac{1}{2} \int_0^1 \frac{d\lambda}{\lambda} d^3x d^3x' \lambda V(\vec{x} - \vec{x}')$$

$$[iD^{\lambda}(\vec{x}'t; \vec{x}t) - iD^0(\vec{x}'t, \vec{x}t)]$$

↪ using a trick for calculating the potential energy (Eq. 7.30)

The second term is called the "Correlation energy." In momentum space,

$$E_{\text{corr}} = \frac{V}{2} \int_0^1 \frac{d\lambda}{\lambda} \int \frac{d^4q}{(2\pi)^4} \lambda V(\vec{q}) [iD^{\lambda}(\vec{q}, \omega) - iD^0(\vec{q}, \omega)]$$

is for noninteracting particles;

By Wick's theorem

$$iD^0(x, x') = (2s+1) \overset{\text{spins}}{G^0(x, x')} G^0(x', x)$$

$$= \begin{array}{c} x \\ \curvearrowright \\ x' \end{array} \quad \left\{ \text{or, } \begin{array}{c} \uparrow \\ \curvearrowright \\ \downarrow \\ q \end{array} \right. \left. \begin{array}{c} k \\ \curvearrowright \\ k+q \end{array} \right\}$$

which is the lowest order polarization

insertion  =  +  + ...

So

$$D^0(x, x') = \hbar \Pi^0(x, x')$$

Same is true for the interacting theory,

$$D(x, x') = \hbar \Pi(x, x')$$

$$\text{Dyson's equation: } \int d^4x_1 U_0(x, x_1) \Pi(x_1, x')$$

$$= \int d^4x_1 U(x, x_1) \Pi^*(x, x')$$

sum of proper polarization
in sections.

The Ring Approximation

(FW say this is sometimes called the "random phase approximation")

$$\text{wavy line} = \text{bare} + \text{ring} + \text{ring} + \dots$$

$$E_{\text{ring}} = \sum_{n=2}^{\infty} E_{\text{ring}, n}$$

$$= \frac{i\hbar V}{2} \int_0^1 \frac{d\lambda}{\lambda} \int \frac{d^4 q}{(2\pi)^4} \sum_{n=2}^{\infty} [\lambda U_0(q) \Pi^0(q)]^n$$

$$= \dots \int \frac{d^4 q}{(2\pi)^4} \frac{(\lambda U_0 \Pi^0)^2}{1 - \lambda U_0 \Pi^0}$$

$$= \dots \int \frac{d^4 q}{(2\pi)^4} \lambda U_0 \Pi^0 \underbrace{U_{\text{RING}}(q)}_{\text{defines the effective interaction}}$$

λU_0 = bare interaction

U_{RING} = effective interaction

So, we have then ten leading contributions to the correlation energy

$$E_{\text{RING}} = \text{diagram 1} + \text{diagram 2} + \text{diagram 3} + \dots$$

Results for $\Pi^0(\vec{q}, q_0)$ when $v = \frac{\hbar q_0}{2(\hbar^2 k_F^2/2m)}$

$$\begin{aligned} \text{Re } \Pi^0(\vec{q}, q_0) &= \frac{2mk_F}{\hbar^2} \frac{1}{4\pi^2} \left\{ -1 + \right. \\ &\quad \left. + \frac{1}{2q} \left[1 - \left(\frac{v}{q} - \frac{q}{2} \right)^2 \right] \ln \left| \frac{1 + (v/q - q/2)}{1 - (v/q - q/2)} \right| \right. \\ &\quad \left. - \frac{1}{2q} \left[1 - \left(\frac{v}{q} + \frac{q}{2} \right)^2 \right] \ln \left| \frac{1 + (v/q + q/2)}{1 - (v/q + q/2)} \right| \right\} \end{aligned}$$

and $\text{Im } \Pi^0(\vec{q}, q_0)$ is more complicated.

The ring approximation for the ~~the~~ correlated energy is

$$\begin{aligned}
 E_{\text{Ring}} &= \frac{i\hbar V}{2} \int \frac{d^4 q}{(2\pi)^4} \int_0^1 \frac{d\lambda}{\lambda} \frac{\left[\lambda U_0(\mathbf{q}) \Pi^0(\mathbf{q}) \right]^2}{\left[1 - \lambda U_0(\mathbf{q}) \Pi^0(\mathbf{q}) \right]} \\
 &= \frac{-i\hbar V}{2} \int \frac{d^4 q}{(2\pi)^4} \left\{ \ln \left[1 - U_0(\mathbf{q}) \Pi^0(\mathbf{q}) \right] + U_0(\mathbf{q}) \Pi^0(\mathbf{q}) \right\} \\
 &= \frac{N e^2}{2 a_0} E_{\text{RWC}} \text{ for Vee Coulomb interaction.}
 \end{aligned}$$

$\frac{e^2}{2a_0} = \text{the Rydberg energy.}$

$$E_{\text{RING}} = \frac{2}{\pi^2} (1 - \ln 2) \ln r_s + \text{constant}$$

in the limit $r_s \rightarrow 0$.

From Feynman, Statistical Mechanics,
page 264

$$\frac{E}{N} = \frac{2.22}{r_s^2} - \frac{0.916}{r_s} + 0.0622 \ln r_s - 0.096$$

$+ O(r_s) \quad \text{Rydbergs}$

r_s = average spacing in unit of Bohr radius

$$r_s = r_0 / a_B$$

Homework Problem #16.

12e. The effective interaction

$$U_r(q, 0) = \frac{4\pi e^2}{q^2 + (4\alpha r_s / \pi) k_F^2 g(q/k_F)}$$

$$g(x) = \frac{1}{2} - \frac{1}{2x} \left(1 - \frac{1}{4}x^2\right) \ln \left| \frac{1 - \frac{1}{2}x}{1 + \frac{1}{2}x} \right|$$

$$U_r(q, 0) \underset{r_s \rightarrow 0}{\approx} \frac{4\pi e^2}{q^2 + (4\alpha r_s / \pi) k_F^2}$$

$$V_r(x) \approx e^2 e^{-q_{TF} x} x^{-1}$$

The e^2/r Coulomb potential is screened, with screening length $1/q_{TF}$, by the other charges (positive and negative) in the neighborhood of the electrons.