THE SARFACE SYSTEM OF DILUTION SOLUTION OF HE3 IN HE4 TREATED AS A SYSTEM OF TWO DIMEUSIONAL IDEAL FERMI GAS

用二度空間佛米氣體之模型來處理 He³ 與 He⁴ 稀 溶液的表面系統

Dilute solutions of He³ in He⁴ have been the subject of a good deal of theoretical and experimental research in the last several years and are reasonably well understood. They were first treated theoretically by Landau and Pomeranchuk. (1),(2) They considered the change in the liquid He 4 due to the presence of He 3 impurity atoms in terms of elementary excitations called He³ quasiparticles which have the energy spectrum

$$E = -E_s + \frac{p^2}{2m^*} \tag{1}$$

Where -E₃ is the binding energy of a single He³ atom in the ground state in He⁴, m* is the effective mass of the quasiparticle and p is its momentum. At temperatures below 0.5-0.6k when phonon and roton excitations are negligible, Edwards, et al (3) and Anderson, et al (4) showed in their heat capacity measurement experiment that these dilute system can be treated as a system of ideal Fermi gas of He³ quasiparticles if one allowed for the effective mass varies with concentration*. The number density of He³ quasiparticles equals to that of He³ impurity atoms. A natural extension of ones thought along this line is that the surface of dilute He^3 - He^4 solutions might behave in a similar way to a two dimensional ideal Fermi gas of surface He³ quasiparticles.

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Andreev⁽⁵⁾ treated the surface system of dilute He³ - He⁴ mixtures as a system of two dimensional ideal gas of surface He³ quasiparticles with energy spectrum

$$E = -E_{s} - \varepsilon_{o} + \frac{p^{2}}{2M}$$
 (2)

where $-\varepsilon_0$ is the difference of the ground state energy of the surface state and the He³ quasiparticle in the bulk. M can be considered as the effective mass of the surface He³ - quasiparticle. Starting from equation (2), he calculated the surface tension σ of the system in the Boltzmann region to be

$$\sigma = \sigma_4 - x \frac{\hbar \rho}{m_4} \frac{M}{m^*} \left(\frac{2\pi k_B T}{m^*} \right)^{\frac{1}{2}} e^{\frac{\epsilon_0}{k_B T}}$$
(3)

Where σ_4 is the surface tension of the pure solvent He⁴, x is the He³ concentration, m_4 is the mass of He⁴ atom, ρ is the density of liquid He⁴, k_B is the Boltzmann constant, T is the temperature and \hbar is Planck constant divided by 2π . He compared his results with Eselson's (6) experimental results and later Zinoveva (7) et al did an experiment checked with Andreev's predication. The results indicated that two dimensional ideal gas is a good working model for the surface system. This work is to extend Andreev's calculation to the degenerate region. By comparing the result with experimental data in the lower temperature region, one should be able to determine whether the two dimensional Fermi gas is a good model or not.

The average occupation number $< n_{\vec{p}} >$ of a Fermi gas is

$$\langle n_{\vec{p}} \rangle = \frac{1}{\eta^{-1} e \times p\beta \epsilon_{\vec{p}} + 1}$$
 (4)

Where $\epsilon_{\overline{p}}$ is the energy in the one particle state with momentum \overline{p} , $\eta=e^{\beta\mu'}=e^{\beta(\mu_3+\epsilon_0)}$, $\beta=\frac{1}{k_BT}$, μ_s is the He³ chemical potential in the bulk of He³ - He⁴ mixtures. At equilibrium

 μ_3 is equal to the He³ surface chemical potential. For the surface system, the excess number of He³ particles per unit area of the surface n , the surface adsorption, is then

$$n_{s} = \frac{2}{h^{2}} \int_{0}^{\infty} \frac{\eta e \times p \left(-\beta \frac{p^{2}}{2M}\right)}{1 + \eta e \times p \left(-\beta \frac{p^{2}}{2M}\right)} \cdot 2\pi p dp$$
let $x = \beta \frac{p^{2}}{2M}$, $1 + \eta e \times p \left(-\beta \frac{p^{2}}{2M}\right)$
then
$$n_{s} = \frac{4\pi M}{\beta h^{2}} \int_{0}^{\infty} \frac{\eta e^{-x}}{1 + \eta e^{-x}} dx = \frac{-M}{\pi \beta h^{2}} \ln \left(1 + \eta e^{-x}\right) \Big|_{0}^{\infty}$$

$$\therefore n_{s} = \frac{M k_{B}T}{\pi h^{2}} \ln \left(1 + e \times p \left(\frac{\mu_{s} + \epsilon_{0}}{k_{B}T}\right)\right)$$
(5)

From the well-known thermodynamic formula (8)

$$n_s = -(\frac{\partial \sigma}{\partial \mu_s})_T$$

equation (5) can be written in the following form

$$\frac{\pi \hbar^2}{Mk_BT} d\sigma = -\ell n (1 + e^{\beta \mu'}) d\mu'$$
 (6)

Integrate equation (6), notice that as the He^3 concentration x approaches to zero the He 3 chemical potential μ_s and hence μ' approaches to $-\infty$, we obtain

$$=\sigma_4 - \frac{M(k_B T)^2}{\pi h^2} \int_{-\infty}^{\xi} \ell n (1 + e^{\xi}) d\xi$$
 (7)

Where
$$\dot{\xi} = \beta \mu'$$
 Let $\eta = e^{\xi}$ we have

$$\sigma = \sigma_4 - \frac{M(k_B T)^2}{\pi h^2} \int_0^{\eta} \frac{\ln(1+\eta)}{\eta} d\eta$$
 (8)

In the Boltzmann region, when temperature $T\gg T_{\rm F\,s}$, where $T_{\rm F\,s}=\frac{\pi\hbar^2}{Mk_{\rm B}}n_{\rm s}$ is the Fermi temperature of the two dimensional Fermi gas, it can easily be shown, from equation (5) that $\eta \ll 1$. Under this condition equation (8) can be integrated by series ex-

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pansion. Neglecting the higher order terms.

$$\sigma = \sigma_{\bullet} - \frac{M(k_B T)^2}{\pi \hbar^2} e^{\beta (\mu_B + \epsilon_0)}$$
(9)

The He 3 chemical potential μ_s of the bulk liquid in the Boltzmann region is $^{(9)}$

$$\mu_{3} = -k_{B}T \ln \left(\frac{2m_{4}}{xp} \left(\frac{m^{*}k_{B}T}{2\pi\hbar^{2}}\right)^{\frac{3}{2}}\right)$$
 (10)

combining equation (9) and equation (10), we obtain Andreev's result in equation (3).

In the degenerate region when $T < T_{\rm F\, s}$ then $\eta < 1$, equation (8) can be integrated as follows.

$$\sigma = \sigma_{4} - \frac{M(k_{B}T)^{2}}{\pi \hbar^{2}} \left\{ \int_{0}^{1} \frac{\ln(1+\eta)}{\eta} d\eta + \int_{0}^{\eta} \frac{\ln(1+\eta)}{\eta} d\eta \right\}$$

$$\therefore \sigma = \sigma_{4} - \frac{M(k_{B}T)^{2}}{\pi \hbar^{2}} \left\{ \frac{\pi^{2}}{12} + \left(\frac{(\ln \eta)^{2}}{2} - \frac{1}{\eta} + \frac{1}{2^{2}\eta^{2}} - \frac{1}{3^{2}\eta^{3}} + \dots \right) \right\}$$

$$(-1)^{n} \frac{1}{n^{2}\eta^{n}} + \dots \right\}^{\eta}$$

$$\therefore \sigma = \sigma_{4} - \frac{M(k_{B}T)^{2}}{\pi \hbar^{2}} \left\{ \frac{\pi^{2}}{6} + \frac{(\ln \eta)^{2}}{2} - \frac{1}{\eta} + \frac{1}{2^{2}\eta^{2}} - \frac{1}{3^{2}\eta^{3}} + \dots \right\}$$

$$(-1)^{n} \frac{1}{n^{2}\eta^{n}} + \dots \right\}$$

$$\therefore \sigma = \sigma_{4} - \frac{M}{2\pi \hbar^{2}} (\mu_{3} + \varepsilon_{0})^{2} \left\{ 1 + \frac{\pi^{2}}{3\xi^{2}} - \frac{2}{\xi^{2}} e^{-\xi} + \dots (-1)^{n} \frac{2}{n^{2}\xi^{2}} e^{-n\xi} + \dots \right\}$$

$$(11)$$

In the highly degenerate region, one can keep only the first few leading terms in the bracket. With the dilution refrigerator available commercially which can operate in the milidegree range continuously, it is not a difficult task to measure the surface tension in the degenate region. $(T \leq 0.2^{\circ} k$ for most dilute concentation). By comparing with the experimental data one can determine the

validity of this model and determine the two parameters M and ε_0 in equation (2). The presence of weak interactions between the surface quasiparticles will show up on the dependence of M on the concentration x. The same result can be obtained by explaining the surface tension σ as the surface pressure (10) of the two dimensional Fermi gas and using the relation between the pressure and the grand partition function of the system.

* Experiment showed that the value of M* depends on the He³ concentration. This indicates that a weak interaction exists between He³ quasiparticles. (11)(12)

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