

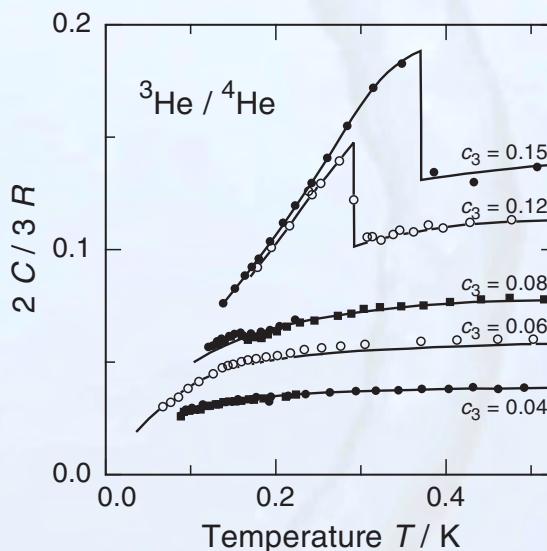
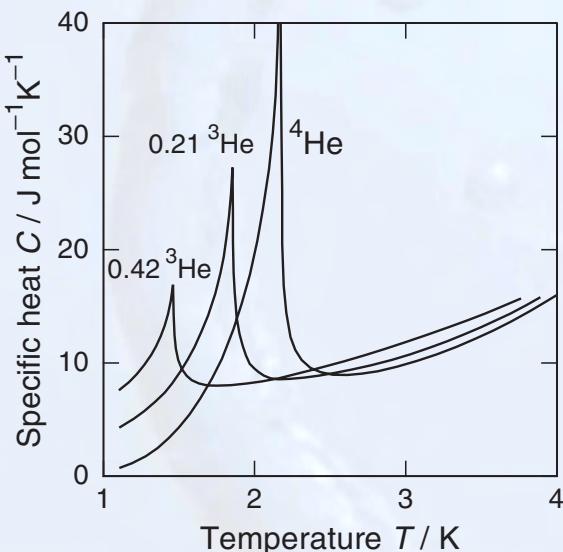
5) $^3\text{He}/^4\text{He}$ Mixtures

interesting for **technical reasons**: dilution cryostats

test for different **theories**: Fermi liquids, RGT, ...

first experiments 1947
observation of second sound 1950

5.1 Specific heat and phase diagram



- lambda transition **shifts to lower temperatures** with $c_3 = N_3/(N_3 + N_4)$

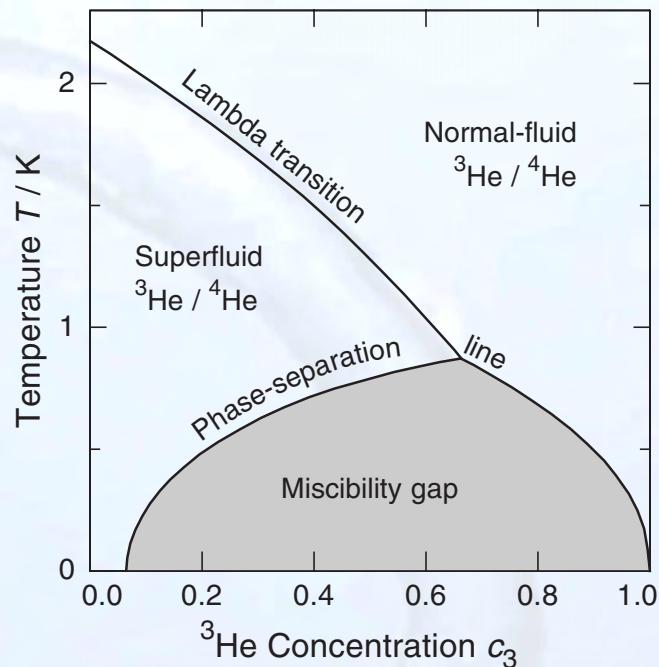
- low temperatures: **jump in specific heat**
 - 1st order phase transition
 - **de-mixing** of ^3He and ^4He

5.1 Specific heat and phase diagram



Phase diagram

- tri-critical point $T = 0.87 \text{ K}$, $c_3 = 0.67$
- miscibility gap is observed



$$\text{light phase } ({}^3\text{He rich}): \quad c_4 = (1 - c_3) = a \sqrt{T^3} e^{-b/T}$$
$$a = 0.85 \text{ K}^{-3/2}$$
$$b = 0.56 \text{ K}$$

$$\text{heavy phase } ({}^4\text{He rich}): \quad c_3 = c_{3,0} (1 + \tilde{a} T^2 + \tilde{b} T^3)$$
$$c_{3,0} = 0.0648$$
$$\tilde{a} = 8.4 \text{ K}^{-2}$$
$$\tilde{b} = 9.4 \text{ K}^{-3}$$

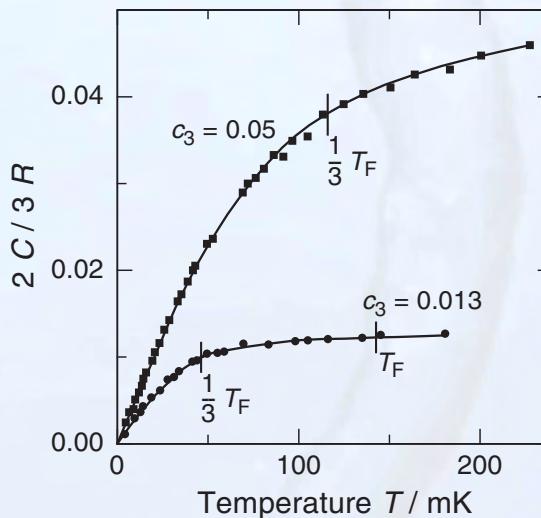
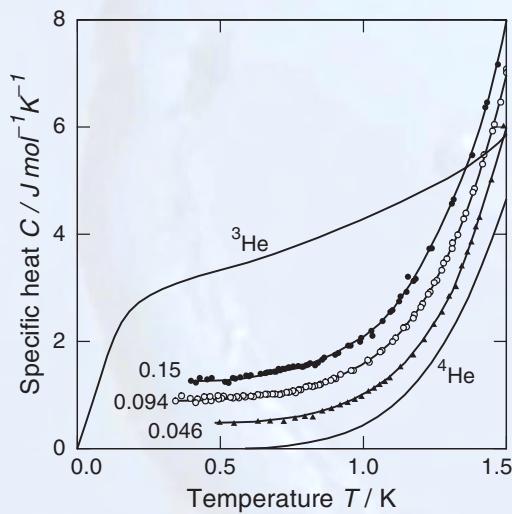


5.1 Specific heat and phase diagram

dilute solutions of ^3He in He-II ($c_3 < 0.15$, $T < 0.5$ K)→ ^4He : passive background fluid→ ^3He : “free” atoms in a quasi vacuum and effective mass $m_3^* = 2.4 m_3$

Fermi gas $T_F = \frac{\hbar^2}{2m_3^* k_B} (3\pi^2 n_3)^{2/3} \propto c_3^{2/3}$

$$C \begin{cases} \rightarrow T > T_F, \quad C \propto c_3 T^0 \quad (\hat{=} \frac{3}{2} R) & \text{high } T \\ \rightarrow T < \frac{1}{3} T_F, \quad C \propto T & \text{low } T \end{cases}$$



- ▶ T_λ depends on c_3
- ▶ pure ^3He : transition Fermi gas → Fermi liquid
- ▶ high T , dilute solution: classical gas with m^*
- ▶ low T : transition classical gas → Fermi gas
- ▶ lines correspond to theory



Finite solubility of ^3He in liquid ^4He at $T = 0$

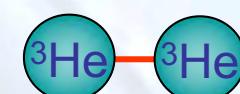
reason: difference in zero-point motion of ${}^3\text{He}$ and ${}^4\text{He}$

v. Waals interaction **identical** for ${}^3\text{He}$ - ${}^3\text{He}$ and ${}^3\text{He}$ - ${}^4\text{He}$

but: larger zero-point motion of ${}^3\text{He}$ weakens the bonding

stronger effective binding for

compared to



in equilibrium on finds

► $T = 0 \longrightarrow c_3 = 1$ for concentrated phase (pure ${}^3\text{He}$)

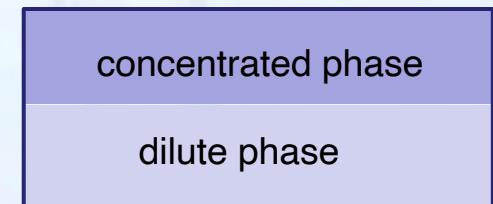
- necessary energy to bring one ${}^3\text{He}$ atom into “vacuum” $L_3(T = 0)$

$$\longrightarrow \mu_{3,c}(0,1) = \mu_3(0) = -L_3(0) = -2.473 \text{ K} \quad \text{latent heat}$$

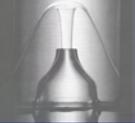
► dilute phase: $E_3 = -\mu_{3,d}(0,0)$ → binding energy

$$c_{3,d} \rightarrow 0$$

- with **increasing** concentration, the **effective** binding energy for ${}^3\text{He}$ is reduced because of the Pauli principle \longrightarrow Fermi gas: $E_F = k_B T_F(c_3)$

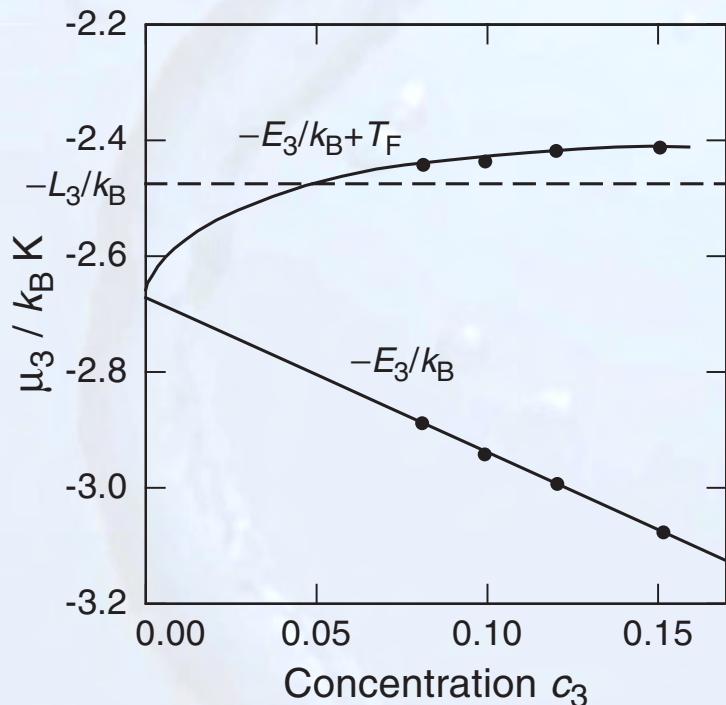


5.1 Specific heat and phase diagram

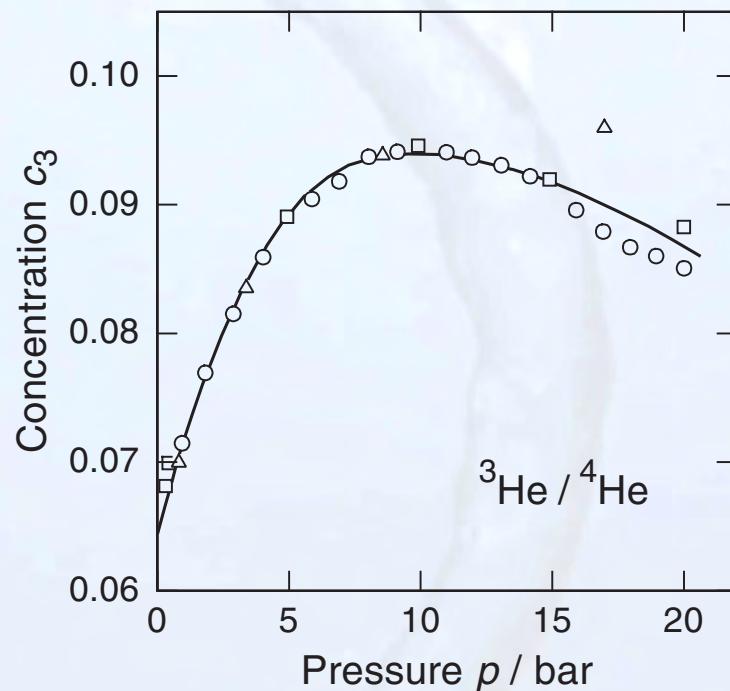


equilibrium concentration at $T = 0$

$$-L_3(0) = -E_3(0, c_3) + k_B T_F(c_3)$$



pressure dependence



- ▶ calculation of $E_3(0, c_3)$ is not trivial
- Bardeen, Baym, Pines model

- ▶ maximum at 8.7 bar
- ▶ concentration $c_3 = 0.096$

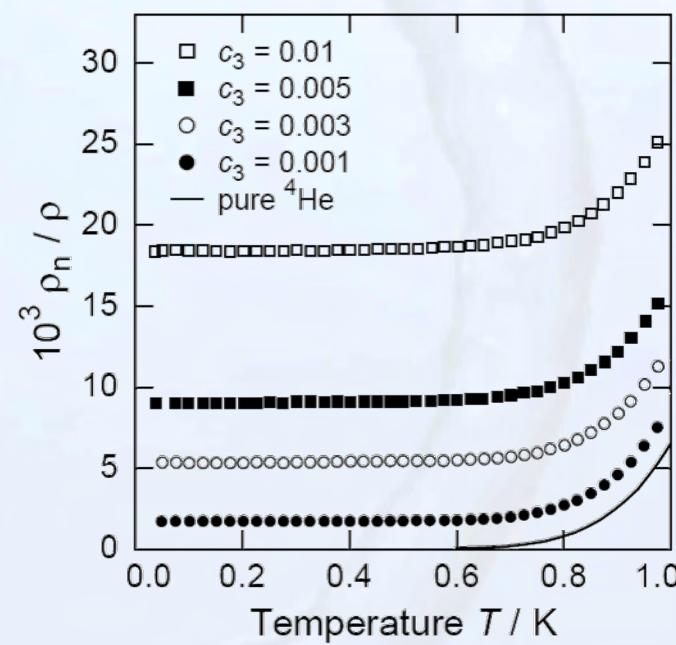
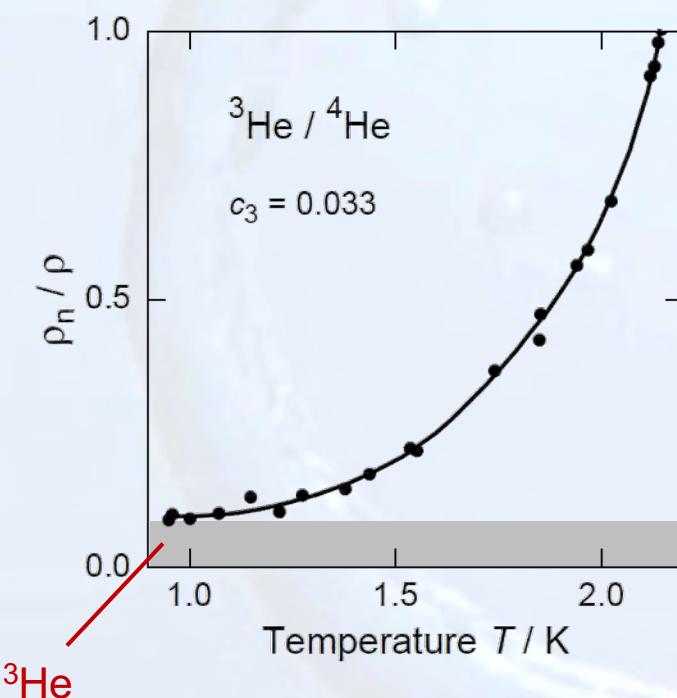
5.2 Normalfluid Component

determination of ρ_n → Andronikasvili-type experiment

$$\rho_n = \rho_{n,4} + \rho \frac{m_3^*}{m_4} c_3 \quad \text{const}$$

pure He-II

const



→ $\rho_n(T \rightarrow 0) = \text{const} \propto c_3$



5.2 Normalfluid Component

Osmotic pressure

► ${}^4\text{He}$ flows to solution to thin the ${}^3\text{He}$ concentration

► ${}^3\text{He}$ is blocked

→ osmotic pressure

van't Hoff law ($T \gg T_F$, classical regime)

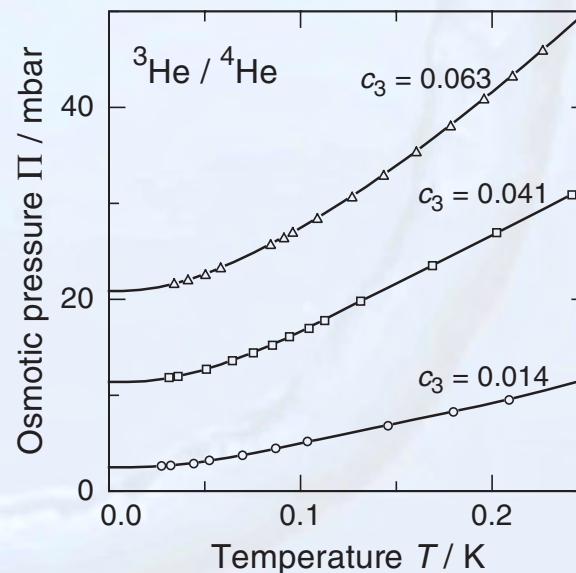
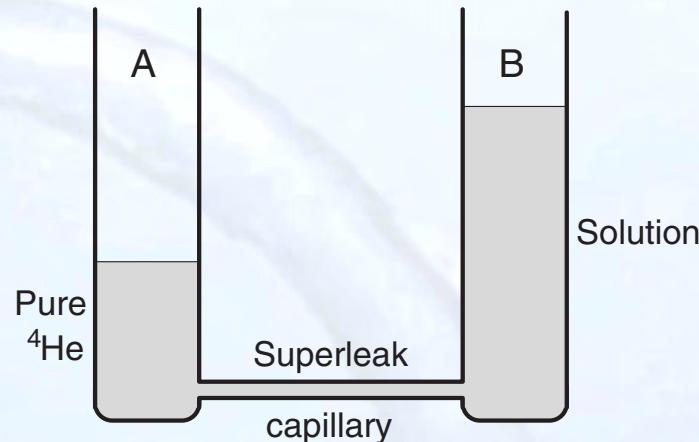
$$\Pi = n_3 k_B T \propto c_3 T$$

$T \ll T_F$, degenerate Fermi gas

$$\Pi = \frac{2}{5} n_3 k_B T_F \propto c_3^{5/3} = \text{const}$$

depends on c_3

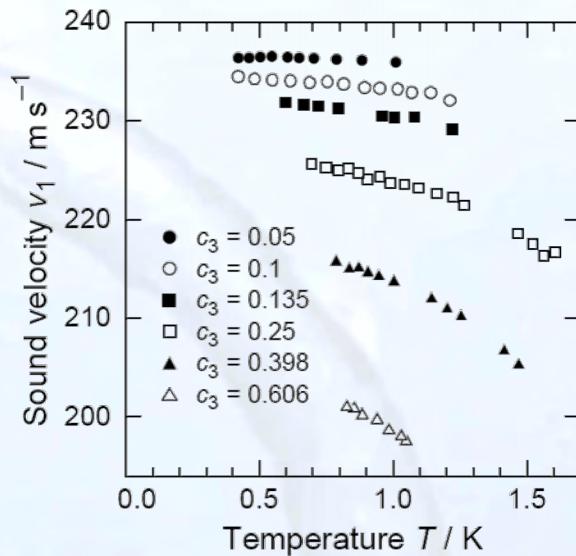
→ transition from FG to classical gas



5.3 Sound Propagation

First sound

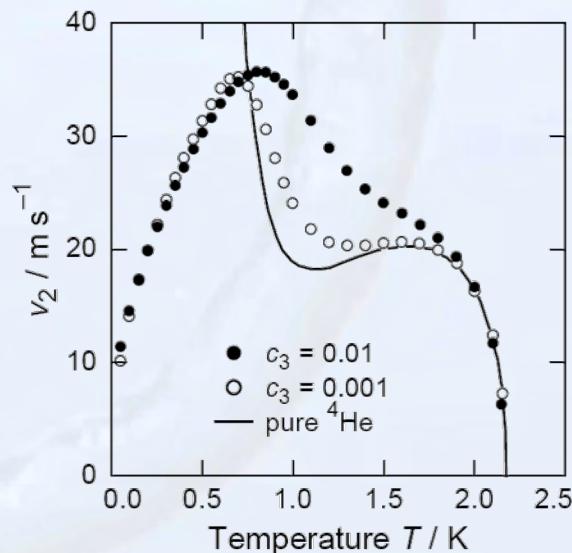
$$v_1^2 = \left(\frac{\partial p}{\partial \varrho} \right)_{S,c_3} \left[1 + \frac{\varrho_s}{\varrho_n} \left(\frac{\partial \varrho}{\partial c_3} \frac{c_3}{\varrho} \right)^2 \right]$$



Second sound

$$v_2^2 = \frac{\varrho_s}{\varrho_n} \left[\bar{S} \left(\frac{\partial T}{\partial S} \right)_{\varrho,c_3} + c_3^2 \frac{\partial(\mu_3 - \mu_4)}{\partial c_3} \right] \left[1 + \frac{\varrho_s}{\varrho_n} \left(\frac{\partial \varrho}{\partial c_3} \frac{c_3}{\varrho} \right)^2 \right]^{-1}$$

$\bar{S} = S_{4,0} - \frac{k_B}{m_4} [c_3 + \ln(1 - c_3)] + \frac{k_B}{m_3} c_3$





5.4 Transport Properties

Thermal transport (rather complex)

high Temp.: ϱ_n flow leads to ${}^3\text{He}$ concentration gradient

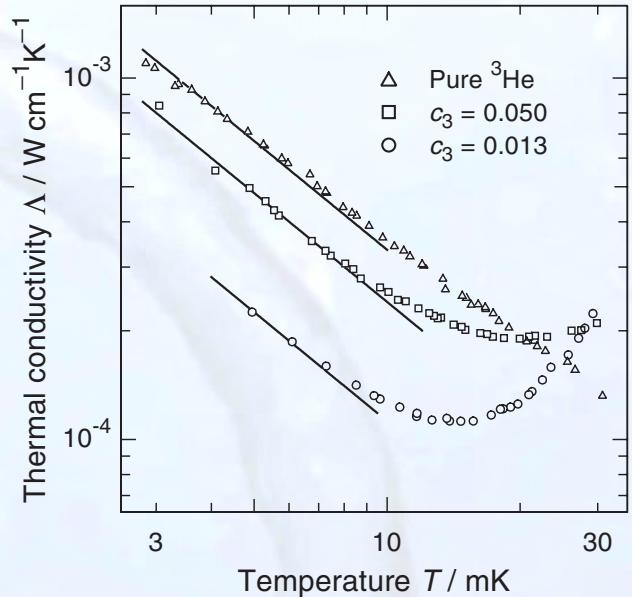
- ${}^3\text{He}$ atoms diffuse back
- ${}^3\text{He}$ form scattering centers for ϱ_n
- reduction of heat transport

low Temp.: ${}^3\text{He}$ atoms from FG

$$\Lambda = \frac{1}{3} C v_F \ell \propto \frac{c_3}{T}$$

$\ell = v_F \tau \propto (T_F/T)^2$

$$v_F = (\hbar/m_3^*)(3\pi^2 n_3)^{1/3},$$
$$C \propto T/T_F$$
$$m_3^* = (1 + F_1/3)m_3$$





very interesting: 3 superfluid phases in the same container → ${}^4\text{He}$, ${}^3\text{He}$, and dilute ${}^3\text{He}$

Problem: ${}^3\text{He}/{}^4\text{He}$ mixtures are hard to cool to below $200 \mu\text{K}$ because of Kapitza resistance



acoustic mismatch hinders cooling

new initiative:

- cooling by melting of ${}^4\text{He}$ crystal
- lowest temperature so far $90 \mu\text{K}$

