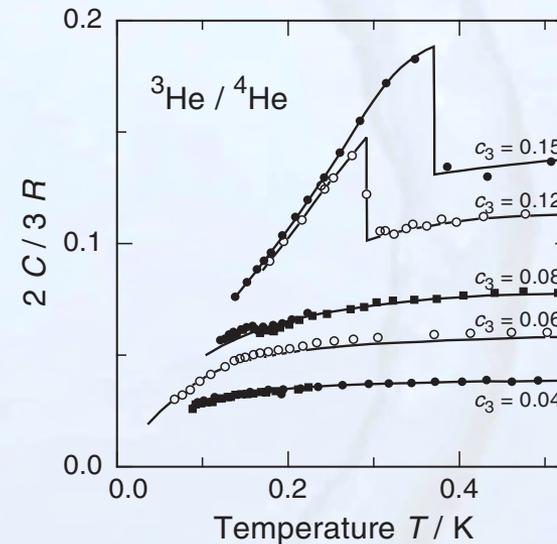
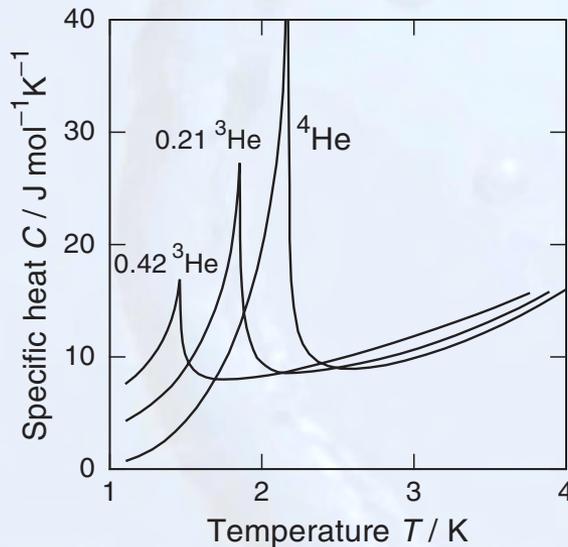




- interesting for technical reasons: dilution cryostats
- test for different theories: Fermi liquids, RGT, ...

first experiments 1947  
observation of second sound 1950

## 5.1 Specific heat and phase diagram



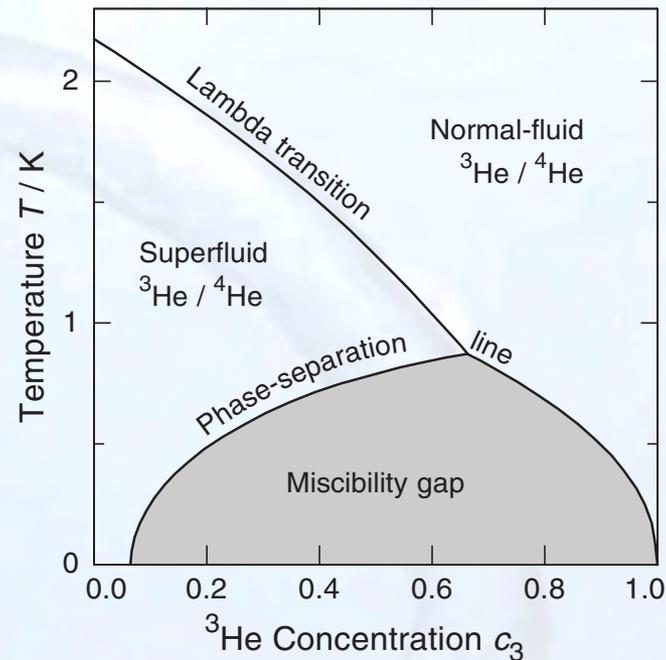
- ▶ lambda transition shifts to lower temperatures with  $c_3 = N_3/(N_3 + N_4)$

- ▶ low temperatures: jump in specific heat
  - 1<sup>st</sup> order phase transition
  - de-mixing of  $^3\text{He}$  and  $^4\text{He}$



## Phase diagram

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



$a = 0.85 \text{ K}^{-3/2}$        $b = 0.56 \text{ K}$

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

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

$\tilde{a} = 8.4 \text{ K}^{-2}$        $\tilde{b} = 9.4 \text{ K}^{-3}$

$c_{3,0} = 0.0648$



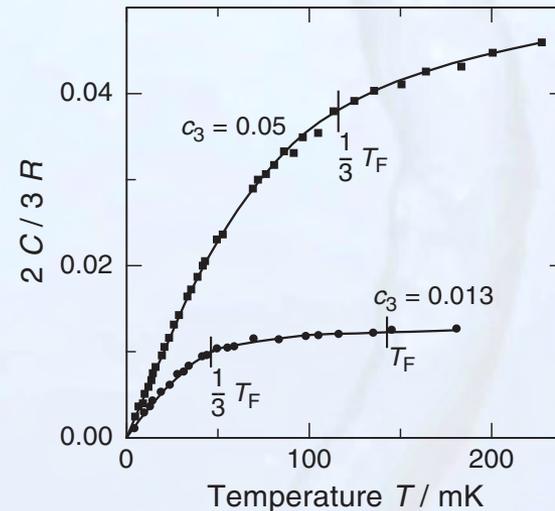
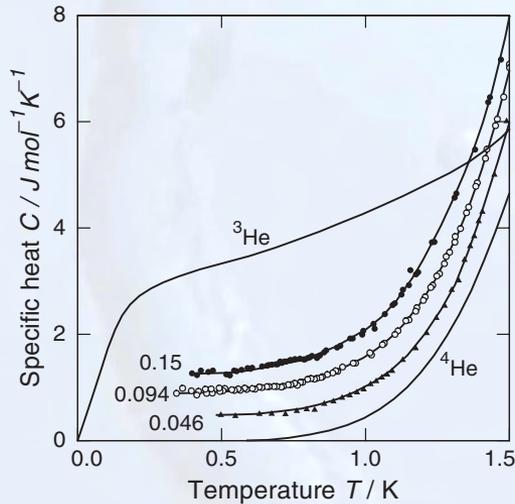
dilute solutions of  $^3\text{He}$  in He-II ( $c_3 < 0.15$ ,  $T < 0.5$  K)

→  $^4\text{He}$ : passive background fluid

→  $^3\text{He}$ : “free” atoms in a quasi vacuum and effective mass  $m_3^* = 2.4 m_3$

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

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



- ▶  $T_\lambda$  depends on  $c_3$
- ▶ pure  $^3\text{He}$ : transition Fermi gas → Fermi liquid
- ▶ high  $T$ , dilute solution: classical gas with  $m^*$
- ▶ low  $T$ : transition classical gas → Fermi gas
- ▶ lines correspond to theory



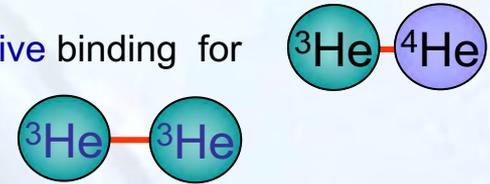
Finite solubility of  $^3\text{He}$  in liquid  $^4\text{He}$  at  $T = 0$

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

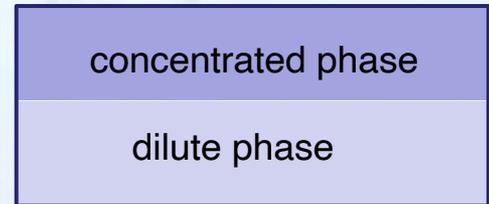


in equilibrium one finds

$$\mu_{3,d}(T, c_{3,d}) = \mu_{3,c}(T, c_{3,c})$$

dilute

concentrated



▶  $T = 0 \longrightarrow c_3 = 1$  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:  $E_3 = -\mu_{3,d}(0, 0) \longrightarrow$  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:  $E_F = k_B T_F(c_3)$

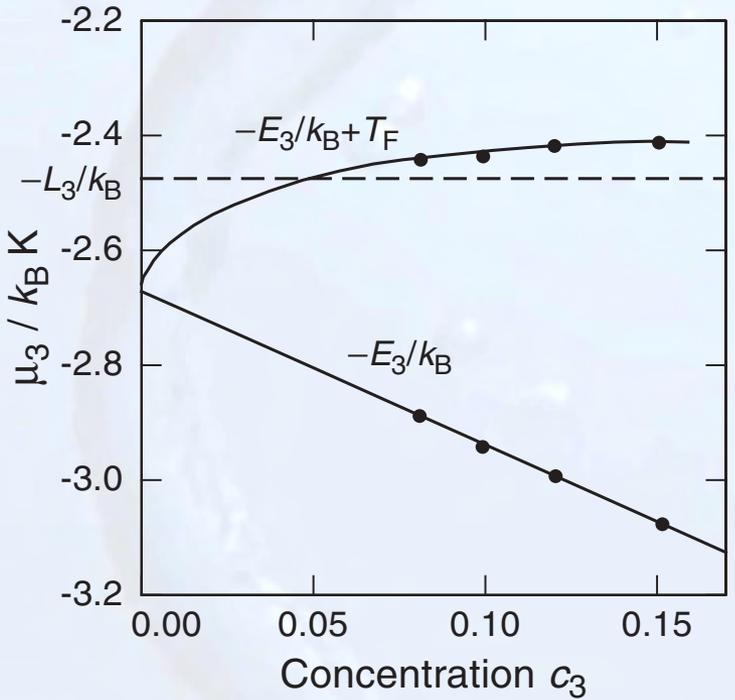


# 5.1 Specific heat and phase diagram

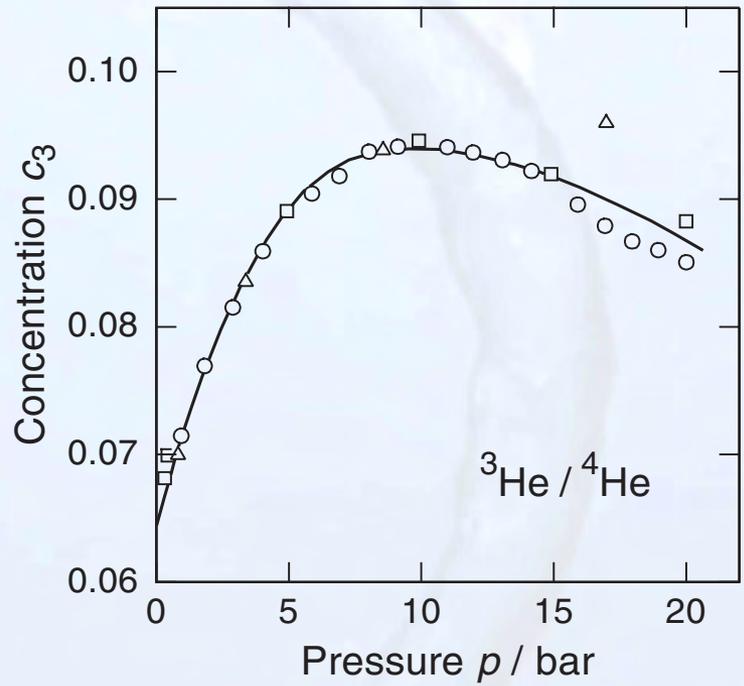


equilibrium concentration at  $T = 0$

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



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

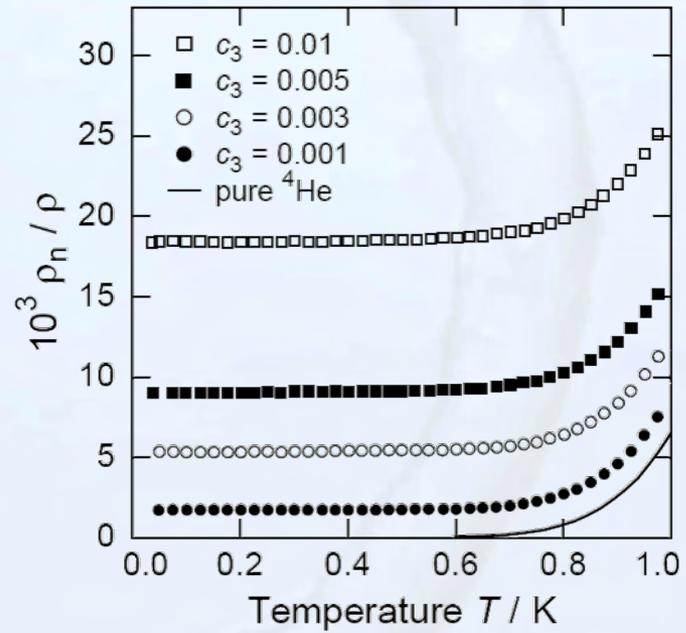
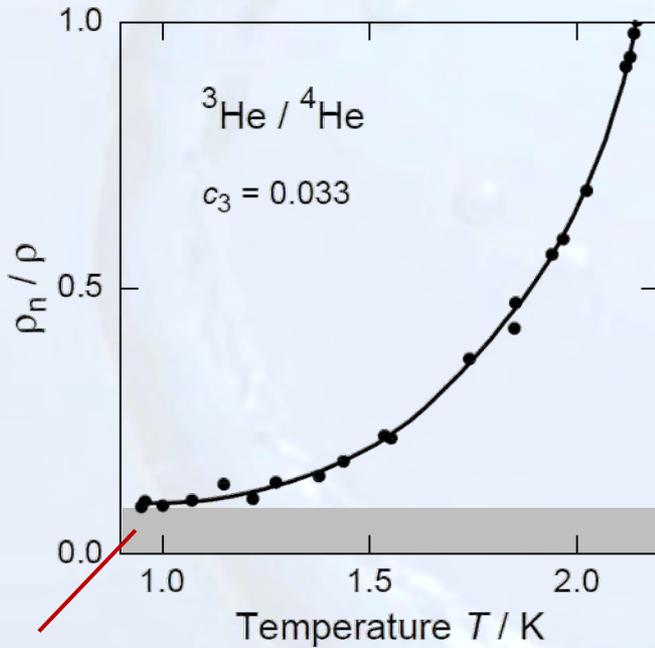


determination of  $\varrho_n \longrightarrow$  Andronikasvili-type experiment

15 mica sheets  
4 cm diameter  
190  $\mu\text{m}$  spacing

$$\varrho_n = \varrho_{n,4} + \varrho \frac{m_3^*}{m_4} c_3$$

pure He-II const



${}^3\text{He}$

$\longrightarrow \varrho_n(T \rightarrow 0) = \text{const} \propto c_3$



## 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)

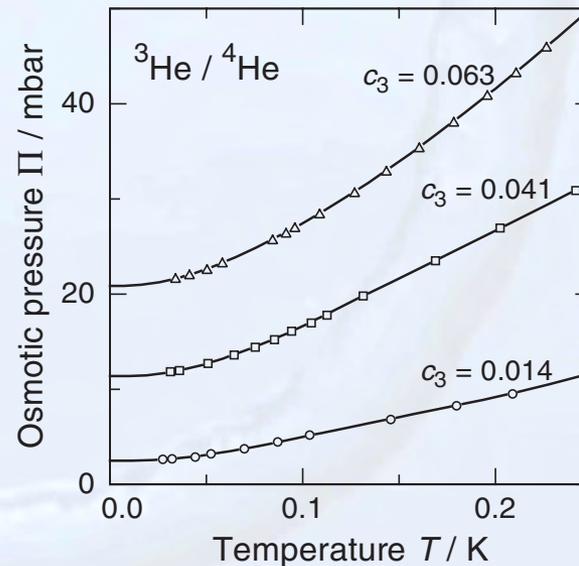
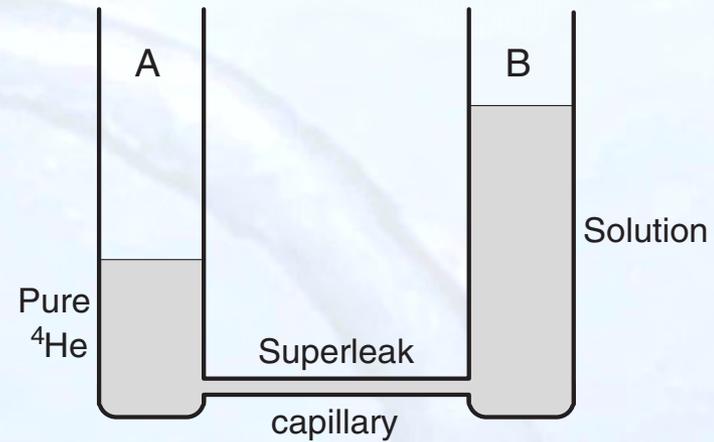
$$\Pi = n_3 k_B T \propto c_3 T$$

$T \ll T_F$ , degenerate Fermi gas

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

depends on  $c_3$

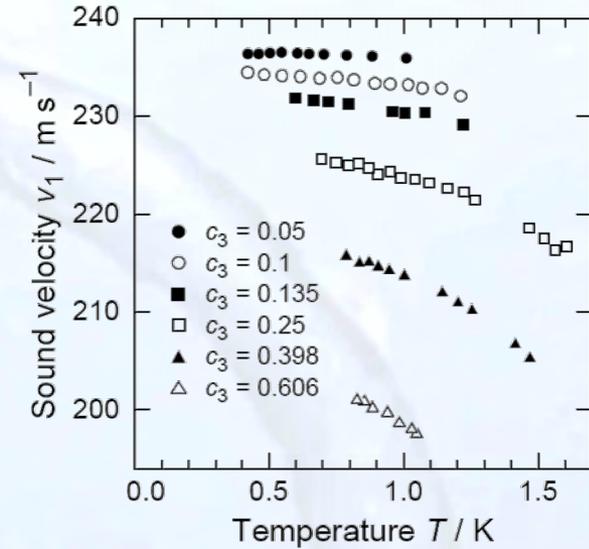
→ transition from FG to classical gas





## First sound

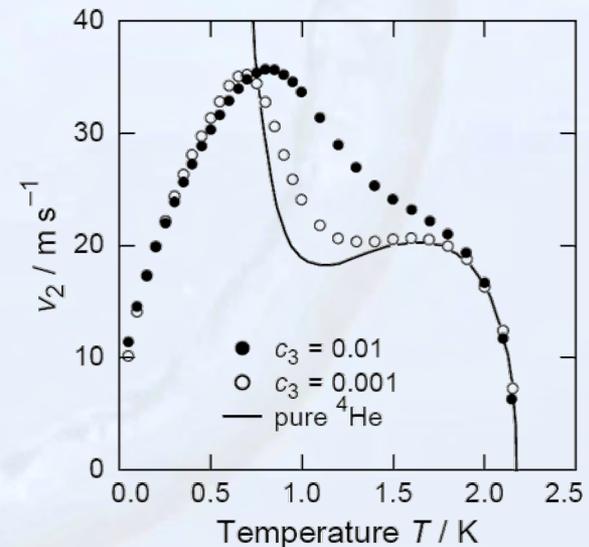
$$v_1^2 = \left( \frac{\partial p}{\partial \rho} \right)_{S, c_3} \left[ 1 + \frac{\rho_s}{\rho_n} \left( \frac{\partial \rho}{\partial c_3} \frac{c_3}{\rho} \right)^2 \right]$$



## Second sound

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

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





Thermal transport (rather complex)

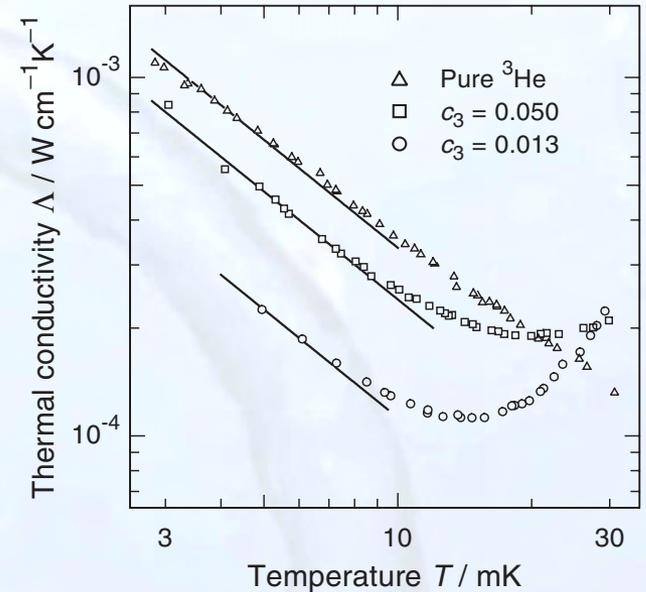
high Temp.:  $\varrho_n$  flow leads to  $^3\text{He}$  concentration gradient

- $^3\text{He}$  atoms diffuse back
- $^3\text{He}$  form scattering centers for  $\varrho_n$
- 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  $\longrightarrow$   $^4\text{He}$ ,  $^3\text{He}$ , and dilute  $^3\text{He}$

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



acoustic mismatch hinders cooling

new initiative:

$\longrightarrow$  cooling by melting of  $^4\text{He}$  crystal

$\longrightarrow$  lowest temperature so far  $90 \mu\text{K}$

