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### Critical ultrasonic attenuation in superfluid helium: Mixing of order-parameter and fluctuation contributions

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The mixing of the Landau-Khalatnikov (order-parameter) and fluctuation contributions to the critical ultrasonic attenuation is calculated in the  $\epsilon$  expansion. The resulting scaling function agrees with the experimental data over nine octaves.

When a system undergoes a second-order phase transition and enters a state of broken symmetry, this change is characterized by a nonvanishing average value of the order parameter  $\langle\phi\rangle\neq 0$ . A pressure wave propagating through the system produces a local modulation of  $\langle\phi\rangle$ . If the relaxation rate  $\gamma$  is comparable to the angular frequency  $\omega$  of the wave, the modulation of  $\langle\phi\rangle$  will lag behind the input pressure signal. The Landau-Khalatnikov<sup>1</sup> (LK) theory of ultrasonic attenuation is based on this hysteresis. The LK attenuation exhibits a characteristic relaxational maximum when  $\gamma=\omega$ . It vanishes at the critical point, where  $\langle\phi\rangle=0$ . The actual critical ultrasonic attenuation, however, by no means vanishes at the critical point. This discrepancy can be attributed to the effect of fluctuations. Being a mean-field theory, the LK approach neglects fluctuations. Williams and Rudnick,<sup>2</sup> in interpreting their measurements of the ultrasonic attenuation in liquid <sup>4</sup>He II, achieved success by simply superposing the LK component on top of the fluctuation component. They assumed the latter to be the same function of the reduced temperature below the  $\lambda$  point as it was above, where there is no LK contribution. This approach, however, has been found by Tozaki and Ikushima<sup>3</sup> to fail at higher frequencies. It has clearly become necessary to have a theory of critical ultrasonic attenuation which does not *ad hoc* assume a linear superposition of the order-parameter (LK) and fluctuation contributions. It is essential instead to take the fluctuations into account from the outset. The purpose of this short note is to present such a theory. Our principal result is the scaling function shown in Fig. 2,

which fits the experimental data over a frequency range that spans three orders of magnitude. From a more abstract point of view, the general significance of our work is that there is an inescapable mixing of the two components which necessarily invalidates any approach that assumes a linear superposition. This "hybridization" also occurs in the intensity of light scattering<sup>4,5</sup> and can be expected to occur in a wide range of phenomena not restricted to superfluid helium.

Recall that the origin of the ultrasonic attenuation is a lagging temperature signal<sup>6</sup> resulting from a complex frequency-dependent specific heat  $C_p(\omega, \gamma)$ . Thus the attenuation in one wavelength is

$$\alpha\lambda = (\text{const}) \text{Im} \frac{-1}{C_p(\omega, \gamma)} \quad (1)$$

For  $(\omega, \gamma) > 1$  MHz liquid helium is in the van Hove precritical regime<sup>7,8</sup> where the kinetic coefficient for order-parameter relaxation equals its noncritical background value  $B_\psi$ . Therefore the temperature dependence of the relaxation rate is given by

$$\gamma = B_\psi \kappa^2 = B_\psi \kappa_0^2 |t|^{4/3} \quad (2)$$

where  $t$  is the reduced temperature,  $\kappa^{-1}$  is the correlation length, and  $\kappa_0$  is a nonuniversal constant of the fluid that has to be determined empirically and independently above and below the  $\lambda$  point.

The specific heat that we need for Eq. (1) is related to the correlation function  $G_Q$  of the quadratic field  $Q = Q_1 + Q_2$ , where the first- and second-order components are  $Q_1 = 2\langle\phi\rangle\phi_L$  and  $Q_2 = \phi^2 = \phi_L^2 + \phi_T^2$ .

$\phi_{L,T}$  are the longitudinal and transverse fluctuations of the order parameter, respectively. The correlation function of  $Q$  between times 1 and 2 is consequently composed of three terms,

$$G_Q(12) = \langle Q(1)Q(2) \rangle = G_{11} + 2G_{12} + G_{22} \quad (3)$$

where

$$\begin{aligned} G_{11} &= \langle Q_1(1)Q_1(1) \rangle = 4\langle \phi \rangle^2 \langle \phi_L(1)\phi_L(2) \rangle \\ &\equiv 4\langle \phi \rangle^2 G(12) \quad (4a) \end{aligned}$$

The mixing of the two contributions to  $Q$  is clearly exhibited by the cross term

$$\begin{aligned} 2G_{12} &= \langle Q_1(1)Q_2(2) \rangle + \langle Q_2(1)Q_1(2) \rangle \\ &= 2\langle \phi \rangle [\langle \phi_L(1)\phi^2(2) \rangle + \langle \phi^2(1)\phi_L(2) \rangle] \quad (4b) \end{aligned}$$

The purely fluctuational component is

$$G_{22} = \langle Q_2(1)Q_2(2) \rangle_c = \langle \phi^2(1)\phi^2(2) \rangle_c \quad (4c)$$

(The subscript on the above equilibrium average indicates the connected part.) We now work with the Fourier transform and convert all correlation functions to response functions. In the broken symmetry state there is a three-point vertex which produces the mixing and interaction effects illustrated by the graphs of Fig. 1. We treat these effects in the  $\epsilon$  expansion. Graph (a) involves only the noninteracting response function  $\gamma g_0$  corresponding to correlation function

$$g_0 = \frac{1}{\gamma - i\omega} \quad (5)$$

The zero-order part of Eq. (4a) becomes therefore the response function

$$g_{11}^0 = 4\langle \phi \rangle^2 g_0 = C_0 \gamma g_0 = C_0 \frac{\gamma}{\gamma - i\omega} \quad (6)$$

of the LK form. The proportionality of  $\langle \phi \rangle^2$  to  $\gamma$ , which is valid to lowest order in the  $\epsilon$  expansion, is expressed by the constant  $C_0$ . As expected, Eq. (6) vanishes at the  $\lambda$  point, where  $\gamma = 0$ . In the thermodynamic limit  $\omega \rightarrow 0$  it adds the constant  $C_0$  to the specific heat for all  $\gamma > 0$  (i.e., for all temperatures below the  $\lambda$  point). This is the finite-frequency form

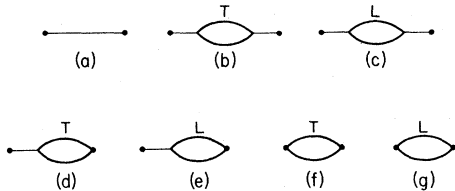


FIG. 1. Low-order contributions to the specific heat below the  $\lambda$  point.

of the familiar mean-field step-function contribution of the order parameter to the specific heat.

The interaction corrections to Eq. (4a) corresponding to graphs (b) and (c) become

$$\Delta g_{11} = 4\langle \phi \rangle^2 \Delta g \propto (\gamma g_0)^2 L_{T,L} \quad (7)$$

where  $L_{T,L}$  represent the "bubbles" of graphs (f) and (g). Each "leg" of a graph corresponds to a factor of  $\gamma g_0$ . Thus Eq. (4b) becomes

$$g_{12} \propto \gamma g_0 L_{T,L} \quad (8)$$

Paying due attention to numerical factors, we find

$$g_Q^T(\omega, \gamma) = (1 - \gamma g_0)^2 \Delta L_T + \text{Re} L_T(-i\gamma, \gamma) \quad (9a)$$

and

$$g_Q^L(\omega, \gamma) = (1 - 3\gamma g_0)^2 \Delta L_L + L_L(-i\gamma, \gamma) \quad (9b)$$

for the total contribution to the specific heat from all of the low-order graphs of Fig. 1 involving transverse fluctuations and longitudinal fluctuations, respectively. We have made a subtraction at the pole  $\omega = -i\gamma$  as this is, by definition, the renormalized value.

A particularly satisfactory feature of Eq. (9a) is that, although  $\Delta L_T$  does not have a thermodynamic limit, Eq. (9a) does. This is because of the vanishing of the prefactor of  $\Delta L_T$  as  $\omega \rightarrow 0$ . For this reason we do not have to sum a series of bubbles, as is often done in this kind of problem. It would be natural to represent superfluid helium, which has one longitudinal and one transverse mode, by an equal mixture of Eqs. (9a) and (9b). We, however, follow a different approach which adheres more closely to the  $\epsilon$  expansion. We take as the hallmark of the superfluid not that it has just one transverse component of the order parameter, but rather that its critical specific-heat exponent vanishes (or very nearly so). It is well known that to describe this situation to first order in the  $\epsilon$  expansion we need three transverse components. We therefore argue that the correct weighting of Eqs. (9a) and (9b) is  $\frac{3}{4}$  and  $\frac{1}{4}$ , respectively.

At this point a computational convenience enters. We have found that the results are not significantly changed (error of a few percent) by using 100% of Eq. (9a). Therefore, from now on, for the sake of simplicity, we drop Eq. (9b) and use only Eq. (9a). A further simplification, justified by a detailed study to be published elsewhere, is to neglect the oscillatory nature of the intermediate lines in graph (f). Thus, ignoring second sound effects and treating the lines as relaxation only, we obtain  $L_T(\omega, \gamma) = L_\lambda(\omega)$ , independent of  $\gamma$ , where  $L_\lambda(\omega)$  is the  $\lambda$ -point specific heat. The subtraction is consequently

$$\Delta L_T = L_\lambda(\omega) - \text{Re} L_\lambda(-i\gamma) = \ln \frac{\gamma}{-i\omega} \quad (10)$$

Denoting  $\text{Re} L_\lambda(\omega)$  by  $L_{1\lambda}(\omega)$  and writing  $g_Q^T(\omega, \gamma)$  as  $L(\omega, \gamma)$  we find from substituting Eq. (10) into

Eq. (9a)

$$L(\omega, \gamma) = \frac{C_0 \gamma}{\gamma - i\omega} + \frac{\omega^2}{(\omega + i\gamma)^2} \ln \frac{\gamma}{-i\omega} + \ln \frac{\omega}{\gamma}$$

$$+ L_{1\lambda}(\omega) = \frac{C_0}{1 - i\Omega} - \frac{\Omega^2}{(\Omega + i)^2} \ln(-i\Omega)$$

$$+ \ln \Omega + L_{1\lambda}(\omega) \quad (11)$$

where we have introduced the scaled frequency<sup>9</sup>  $\Omega = \omega/\gamma$ . With  $L = L_1 + iL_2$ , the imaginary part gives the scaling function

$$F(\Omega) = \frac{2}{\pi} L_2(\Omega)$$

$$= \frac{2}{\pi} \frac{C_0 \Omega}{1 + \Omega^2} + \Omega^2 \frac{\Omega^2 - 1}{(\Omega^2 + 1)^2}$$

$$+ \frac{4}{\pi} \frac{\Omega^3}{(\Omega^2 + 1)^2} \ln \Omega \quad (12a)$$

which is shown by the solid curve in Fig. 2(b). The dashed curve shows the first term of Eq. (12a) alone, the LK contribution. The real part of  $L$  does not scale, being a function of two variables

$$L_1(\omega, \gamma) = \frac{C_0}{1 + \Omega^2} + \frac{\pi}{2} \frac{\Omega^3}{(\Omega^2 + 1)^2}$$

$$- \Omega^2 \frac{\Omega^2 - 1}{(\Omega^2 + 1)^2} \ln \Omega + \ln \Omega + L_{1\lambda}(\omega) \quad (12b)$$

The thermodynamic limit of Eq. (12b) is  $L_1(0, \gamma) = C_0 + L_{1\lambda}(\gamma)$ . Although the  $\epsilon$  expansion gives a value for  $C_0$  of roughly the right size, it is obviously better to determine  $C_0$  empirically from the thermodynamic data. Having determined in this way  $C_0 = 5.4$  as well as  $\kappa_0 = 1.5 \times 10^8 \text{ cm}^{-1}$ , we have been able to account very well for the ultrasonic data on the basis of Eq. (1), which becomes  $\alpha\lambda \propto -\text{Im}L^{-1} = L_2/|L|^2$ . Thus, neglecting the very slight temperature dependence of the sound velocity, we find the ratio of the attenuation below the  $\lambda$  point to that at the  $\lambda$  point to be

$$\frac{\alpha}{\alpha_\lambda} = r^{-1} F(\Omega) \quad (13)$$

where  $r = |L|^2/|L|_\lambda^2$ . Because the factor  $r^{-1}$  does not scale, the product depends on the frequency of the measurement. At higher frequencies,  $|L|^2$  is a relatively strong function of  $\Omega$  and tends to cancel out the maximum of  $F(\Omega)$ . For this reason the 1-GHz data of Commins and Rudnick<sup>10</sup> as well as the 1.7-GHz data of Lambert, Legros, and Salin<sup>11</sup> hardly exhibit any maximum in  $\alpha/\alpha_\lambda$ , as seen in Fig. 2(a) which exhibits the raw data. This contrast with the pronounced maximum in the 3-MHz data of Williams and Rudnick.<sup>2</sup> The effect of frequency on the shape

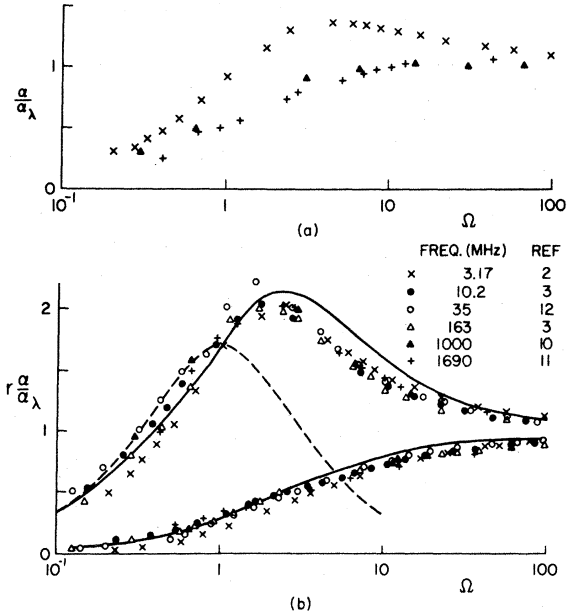


FIG. 2. (a) Unprocessed experimental data showing dependence of shape on frequency and absence of scaling. (b) Scaling functions  $F(\Omega)$  from Eq. (12a) and processed experimental attenuation data  $r\alpha/\alpha_\lambda$ , where  $r = |L|^2/|L_\lambda|^2$ . Inclusion of  $r$  restores scaling. The dashed curve shows the Landau-Khalatnikov contribution. The lower curve shows the scaling above the lambda point (Ref. 6).

of  $\alpha/\alpha_\lambda$  is also clearly evident in the data of Tozaki and Ikushima,<sup>3</sup> and has been remarked upon by them.

In order to reveal the underlying regularity in the apparently disparate data described above, it is essential to subject them to a "processing." This is simply the removal of the offending nonscaling factor  $r^{-1}$  from the right-hand side of Eq. (13) by transferring it to the left. The "processed" values  $r\alpha/\alpha_\lambda$  for the various runs referred to above, as well as for the 35-MHz run of Roe, Meyer, and Ikushima<sup>12</sup> for their 0.7% mixture, are shown in Fig. 2(b). It will be noted that not only do the data thereby coalesce into a single scaling function, but also that this function is in satisfactory agreement with the predicted scaling function of Eq. (12a).

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