Transport in very dilute solutions of $^3$He in superfluid $^4$He

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Motivated by a proposed experimental search for the electric dipole moment of the neutron (nEDM) utilizing neutron-$^3$He capture in a dilute solution of $^3$He in superfluid $^4$He, we derive the transport properties of dilute solutions in the regime where the $^3$He are classically distributed and rapid $^3$He-$^3$He scatterings keep the $^3$He in equilibrium. Our microscopic framework takes into account phonon-phonon, phonon-$^3$He, and $^3$He-$^3$He scatterings. We then apply these calculations to measurements by Rosenbaum et al. [J. Low Temp. Phys. 16, 131 (1974)] and by Lamoreaux et al. [Europhys. Lett. 58, 718 (2002)] of dilute solutions in the presence of a heat flow. We find satisfactory agreement of theory with the data, serving to confirm our understanding of the microscopies of the helium in the future nEDM experiment.

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I. INTRODUCTION

Dilute solutions of $^3$He in superfluid $^4$He have been an ideal testing ground for theories of quantum liquids, with past focus generally on $^3$He concentrations and temperatures for which the $^3$He forms a degenerate Fermi gas. The proposed use of ultra-dilute solutions in the search for a neutron electric dipole moment (nEDM) utilizing neutron-$^3$He capture in a dilute solution of $^3$He in superfluid $^4$He at the Oak Ridge National Laboratory Spallation Neutron Source (SNS) [1, 2], as well as earlier experiments by Rosenbaum et al. [3] and by Lamoreaux et al. [4], all require careful treatment of the transport properties of the solutions. The characteristic temperatures in all cases considered here are of order 0.5 K, at which phonons are the dominant excitation of the $^4$He. In the low temperature degenerate $^3$He regime, phonons have small effect on the $^3$He properties. However, with increasing dilution, when the $^3$He become classically distributed, the situation is reversed, and the phonons play a more and more important role. In the proposed nEDM experiment, a crucial issue is to be able to periodically sweep out the $^3$He by imposing a temperature gradient [5]; the underlying physics of the transport is scattering of phonons in the superfluid $^4$He against the $^3$He. The Lamoreaux et al. experiment, which measured the effect of a heat source on the steady state distribution of $^3$He atoms in a dilute solution, was a prototype for the effects of a “phonon wind” on the $^4$He. The Rosenbaum et al. experiment measured the thermal conductivity of dilute solutions as a function of temperature and concentration. We show below that the results of these experiments can be understood in terms of a simple thermal conductivity determined using well-established scattering amplitudes.

In the experiments of Refs. [3] and [4], the $^3$He number concentrations, $n_3 = n_3/(n_3 + n_4)$, where $n_3$ and $n_4$ are the $^3$He and $^4$He number densities, are in the range $7 \times 10^{-3}$ to $1.5 \times 10^{-2}$ in the non-degenerate regime. Here momentum carried by the phonons goes primarily into the $^3$He, which then transfer it to the walls. The physical dimensions of the experimental container are sufficiently large that the transport properties are determined locally by the microscopic scatterings of the $^3$He and the phonons. The $^3$He-$^3$He interactions are sufficiently strong that they keep the $^3$He in thermal equilibrium at rest at the local temperature $T(\vec{r})$, while phonon-phonon interactions keep the phonons in drifting local equilibrium. By contrast, in the proposed SNS experiment, phonon momentum is transferred primarily to the walls by viscous forces, with the $^3$He playing a negligible role. Furthermore, collisions of $^3$He with the phonons are responsible for establishing equilibrium in the $^3$He cloud. A common characteristic of these experiments is the effect on the dilute solution produced by a localized, static heat source. In this paper we focus on calculating transport properties in the higher concentration regime in Refs. [3] and [4], where the fact that the $^3$He-$^3$He mean free path is short greatly simplifies the transport theory. At lower $^3$He concentrations, $x \lesssim 10^{-6}$, the transport must be calculated by solving the coupled Boltzmann equations for the phonons and $^3$He, taking into account viscous forces at the boundaries; these results as well as their impact on the transport of $^3$He in the much lower concentration regime of the SNS nEDM experiment will be described elsewhere [6].

In Sec. II we examine the hydrodynamic constraints deriving from the steady state situation and from the properties of the superfluid. The basic scattering mechanisms determining the transport properties of dilute solutions—$^3$He-$^3$He, $^3$He-phonon, and phonon-phonon interactions—are described in Sec. III. In Sec. IV, we calculate the thermal conductivity (dominated by the phonons) in this situation; in this calculation we include inelastic recoil of the $^3$He in scattering against the phonons (detailed in Appendix A). We find that, contrary to the earlier treatment in Ref. [7] used by Rosenbaum...
et al., the rapid relaxation of phonons along rays of constant phonon direction dominates their distribution \[8\]. We calculate the conductivity by considering the drag force on the phonons due to their scattering against the stationary \(^{3}\)He, using the method close in spirit to that introduced earlier in a calculation of the mobility of ions in superfluid \(^{4}\)He \[9\], and later used by Bowley \[10\] in his accounting of the Lamoreaux et al. experiment. Despite a superficial similarity of the present approach to these earlier calculations, the underlying physics is different here. In Sec. V, we show how the microscopic theory satisfactorily explains the experimental findings in both the Rosenbaum et al. and the Lamoreaux et al. experiments.

II. RESPONSE TO STATIC DISTURBANCES

To understand how dilute solutions respond to a localized static heat source, the physical mechanism of interest in the experiments, we first review the hydrodynamics of the dominant excitations of the \(^{4}\)He, and the momentum density or mass current density of the \(^{4}\)He is

\[
\vec{g}_4 = \rho_s \vec{v}_s + \rho_{ph} \vec{v}_{ph},
\]

where \(\vec{v}_s\) is the superfluid flow velocity, \(\vec{v}_{ph}\) is the phonon fluid flow velocity, \(\rho_s\) is the superfluid mass density, and

\[
\rho_{ph} = \frac{2\pi^2}{45} \frac{T^4}{s^5} \ll \rho_s
\]

is the \(^{4}\)He normal fluid density. Similarly, the \(^{3}\)He momentum density is

\[
\vec{g}_3 = \rho_3 \vec{v}_3,
\]

where \(\vec{v}_3\) is the \(^{3}\)He flow velocity, and \(\rho_3 = m^* n_3\), with \(m^* = m_3 + \delta m \simeq 2.34 m_3\) the effective mass \[5\]. The total mass current, \(\vec{g}\), is \(\vec{g}_4 + \vec{g}_3\). When the \(^{4}\)He mass flow vanishes, \(|v_s| = (\rho_{ph}/\rho_s) |v_{ph}| \ll |v_{ph}|\) at temperatures \(\lesssim 0.6\) K, and the only relevant flow velocities are those of the phonons and possibly the \(^{3}\)He. Then force balance in the dilute solutions implies that to linear order,

\[
\nabla P = \eta_{ph} \nabla^2 \vec{v}_{ph} + \eta_3 \nabla^2 \vec{v}_3,
\]

where \(P = P_3 + P_{ph}\) is the total pressure, with \(P_3 = n_3 T\) the \(^{3}\)He partial pressure (we generally work in units with \(\hbar\) and Boltzmann’s constant, \(k_B\), equal to unity), \(T\) is the temperature, \(P_{ph}\) is the phonon partial pressure, \(\eta_{ph}\) is the first viscosity of the normal fluid and \(\eta_3\) the first viscosity of the \(^{3}\)He. (In the situations of interest, in a steady state, \(\nabla \cdot \vec{v}_{ph}\) and \(\nabla \cdot \vec{v}_3\) both vanish.) For a container large compared with microscopic viscous mean free paths, and for \(^{3}\)He concentrations in the range of those in Refs. \[3\] and \[4\], both viscosity terms are insignificant compared with the drag forces between the \(^{3}\)He and the phonons, as we discuss in Sec. IV and thus can be neglected. (At the much lower \(^{3}\)He concentrations of the proposed SNS experiment, however, the phonon viscosity does play a significant role \[3\].) The total pressure is effectively constant throughout the system, \(\nabla P = 0\).

In addition, as one sees from the linearized superfluid acceleration equation \[11\],

\[
m_4 \frac{\partial \vec{v}_s}{\partial t} + \nabla \mu_4 = 0,
\]

the \(^{4}\)He chemical potential, \(\mu_4\), is constant in a steady state. The Gibbs-Duhem relation for the solutions, \(\nabla P = n_4 \nabla \mu_4 + n_3 \nabla \mu_3 + S \nabla T\), with \(\mu_3\) the \(^{3}\)He chemical potential, \(T\) the temperature, and \(S\) the total entropy density, together with the constancy of the pressure and the relation, \(\nabla P_3 = n_3 \nabla \mu_3 + S_3 \nabla T\) (which neglects the effects of \(^{3}\)He-phonon interactions on the thermodynamics, e.g., small terms in the total pressure of order \(P_{ph} n_3/n_4 \ll P_{ph}\), then implies that

\[
\nabla P_3 + S_{ph} \nabla T = 0,
\]

where \(S_{ph} = 4 P_{ph}/T\) is the phonon entropy density, and \(dP_{ph} = S_{ph} dT\). Equation \[10\] and the foregoing then give the simple relation between the temperature and \(^{3}\)He density gradients in the system:

\[
\nabla T = -\frac{T}{S_{ph} + n_3} \nabla n_3;
\]

in a steady state a gradient of the \(^{3}\)He density is always accompanied by a gradient of the temperature. (Note that the \(n_3\) in the denominator arises when one consistently includes a non-zero temperature gradient at every step of the calculation, unlike in earlier studies \[13\] \[14\] where the \(^{3}\)He pressure was tacitly assumed to obey \(dP_3 = T dn_3\).) A heat flux, \(\vec{Q}\), in the system is related to the temperature gradient by \(\vec{Q} = -K \nabla T\), where \(K\) is the thermal conductivity of the solution. Thus Eq. \[7\] relates the \(^{3}\)He density gradient to the heat flux by

\[
\nabla n_3 = \frac{S_{ph} + n_3}{TK} \vec{Q}.
\]

The \(^{3}\)He density on the right can be significant. Since

\[
T S_{ph} = s^2 \rho_{ph}
\]

one has

\[
\frac{n_3}{S_{ph}} = \frac{300 x_3}{T^2},
\]

with \(T\) measured in K; at \(T = 0.45\) K and \(x_3 = 3 \times 10^{-4}\) the ratio is unity.

We note that, in general, the response to a heat current in a steady state is a temperature gradient with the constant of proportionality being the thermal conductivity; the particle current in general depends on both gradients of concentration and temperature \[15\]. In
the present case, because of the relation between $\nabla n_3$ and $\nabla T$, Eq. (7), the $^3$He diffusion coefficient is proportional to the thermal conductivity and thus the results of Ref. 4 can also be described in terms of a diffusion coefficient. However, when viscous effects become important (for $x_3 < 10^{-6}$), Eq. (7) is no longer valid and the thermal conductivity and diffusion constant are not related in a simple way 4. In the following we will generally talk about the response to a heat current in terms of the thermal conductivity.

III. MICROSCOPIC SCATTERING PROCESSES

The microscopic processes that determine the transport properties of dilute solutions are $^3$He-$^3$He 16, phonon-phonon 8, and phonon-$^3$He 7 scatterings; the amplitudes of these processes have been well studied in earlier helium research. At low energies, the total cross section for a $^3$He atom scattering from a second $^3$He atom of opposite spin can be written as

$$\sigma_{33} = \frac{9\pi^3}{k_D^2 v_{33,0}} \left( \frac{m^*}{m_4} \right)^2 \left( \frac{m_4 s}{k_D} \right)^4 = 10.3 \text{Å}^2, \quad (11)$$

where $k_D$ is the Debye wave number, defined by $n_4 = k_D^3/6\pi^2$, and $v_{33,0} = -0.064$ measures the strength of the $^3$He-$^3$He interaction at zero momentum transfer 16. Phonon-phonon scattering rates at forward angles 8 are rapid compared with phonon-$^3$He rates. While such scatterings do not directly affect the heat current, they do play the important role of keeping the phonon momentum distribution, $n_q$, in local thermodynamic equilibrium with a drift velocity, $\bar{v}_{ph}$,

$$n_q = \frac{1}{e^{(sq - \bar{v}_{ph}/T)} - 1}. \quad (12)$$

This is the key difference between the current treatment and that in Refs. 3 and 7, where it is assumed that the phonon-$^3$He scattering dominates the phonon relaxation.

The amplitude for scattering of a phonon of wavevector $q$ against a $^3$He of wavevector $p$ to final states $q'$ and $p'$ is determined theoretically in terms of the measured excess volume, $\alpha$, of a $^3$He atom compared with that of a $^4$He atom and the $^3$He effective mass, $m^*$. The scattering amplitude is approximately 7

$$\langle p' q' | T | p q \rangle = \frac{s(qq')^{1/2}}{2n_4} (A + B \cos \theta), \quad (13)$$

where $\theta$ is the angle between $q'$ and $q$, $A = n_4 d\alpha/dn_4$ and $B = (1 + \alpha + \delta m/m_4)(m_4/m^*)(1 + \alpha - m_3/m_4).

At very low temperatures, $T \ll 1$ K, the scattering can taken to be elastic. The momentum dependent scattering rate of phonons from the $^3$He is then 6

$$\gamma_3 (q) = \frac{sq^4 x_3}{4\pi n_4} J, \quad (14)$$

where

$$J = A^2 + B^2/3 - 2AB/3; \quad (15)$$

the $q^4$ in this rate is characteristic of Rayleigh scattering. More generally, as Bowley emphasized 10, recoil of the $^3$He in scattering produces a significant correction to the effective phonon-$^3$He scattering rate 17. As we show in Appendix A, $^3$He recoil corrections to the result 14 are of relative order $T/1.36$ K.

We now estimate the importance of 3-3 versus phonon scattering in bringing the $^3$He into equilibrium. The mean free path of a $^3$He scattering on unpolarized $^3$He is

$$\ell_{33} = \frac{2}{n_3 \sigma_{33}} = \frac{8.66 \times 10^{-8}}{x_3} \text{cm}; \quad (16)$$

the factor $n_3/2$ is the density of opposite spin $^3$He. Similarly, the mean free path for scattering of $^3$He of momentum $p$ on phonons is given approximately by $p^2/m\Gamma$, where

$$\Gamma = \frac{1}{n_3} \int \frac{d^6 q}{(2\pi)^3} q^3 (q) n_q^0 \left( 1 + n_q^0 \right) \approx \frac{270\pi J}{s^2} \left( \frac{S_{ph}}{n_4} \right)^2 T^3 $$

$$\sim T^9, \quad (17)$$

in the limit $p \gg q$. Here $n_q^0 = (e^{sq/T} - 1)^{-1}$ is the equilibrium phonon distribution function. Replacing $p^2$ by $3m^2 T$, an appropriate thermal average, we find

$$\ell_{3ph} = \sqrt{\frac{3}{2J}} \left( \frac{S_{ph}}{n_4} \right) \frac{m^*/s^2}{T_3/2} = 0.077 \frac{(0.45 K)}{T}^{15/2} \text{cm.} \quad (18)$$

Comparing the mean free paths 16 and 18, we find

$$\frac{\ell_{3ph}}{\ell_{33}} = 0.89 \times 10^6 x_3 \frac{(0.45 K)}{T}^{15/2}. \quad (19)$$

For $T = 0.45$ K and $x_3 = 10^{-6}$, $\ell_{3ph} \approx \ell_{33}$, while for $T = 0.65$ K and $x_3 = 3 \times 10^{-4}$, $\ell_{3ph}/\ell_{33} \approx 16.9$. Thus for $T \sim 0.5$ K, and $x_3 \sim 10^{-5}$ a good first approximation is to assume that scattering of $^3$He by $^3$He atoms is fast compared with scattering of $^3$He by phonons. In fact, as will be shown in detail in 6, we may take the momentum distribution of the $^3$He to be simply that of a classical gas in equilibrium and at rest,

$$f_p^0 = e^{-(p^2/2m^* - \mu_3)/T}. \quad (20)$$

IV. THERMAL CONDUCTIVITY

The thermal conductivity of the dilute solutions can in general be calculated by solving the coupled phonon and
\[3^3\text{He} \] Boltzmann equations to determine the steady state momentum distributions of the phonons and the \(3^3\text{He}\), and, from these distributions, the heat and particle currents. However, here, where the \(3^3\text{He}\) are approximately stationary and in equilibrium, the phonon thermal conductivity can be found simply, by calculating the rate at which the phonons lose momentum by scattering against \(3^3\text{He}\) [8, 10].

In a steady state the net force density on the phonons drifting at velocity \(\vec{v}_{ph}\) is balanced by the phonon pressure gradient, \(\nabla P_{ph} = S_{ph} \nabla T\). Microscopically then,

\[
\nabla P_{ph} = -\int \frac{d^3q}{(2\pi)^3} \int \frac{d^3q'}{(2\pi)^3} \int d^3p \frac{d^3p}{(2\pi)^3} \times q|\langle T \rangle|2\pi\delta(sq + p^2/2m^* - sq' - p'^2/2m^*)
\times \left[n_p n_q (1 + n_{q'} - f_p', n_{q'} (1 + n_q))\right],
\]

(21)

where \(\vec{q}' - \vec{q} = \vec{q} - \vec{q}' \equiv \vec{k}, \langle T \rangle = \langle p|q|T|pq \rangle\), \(n_q\) is the phonon distribution, and the factor of 2 is from the \(3^3\text{He}\) spin sum. In terms of the \(3^3\text{He}\) structure function,

\[
S_3(k, \omega) = 2 \int \frac{d^3p}{(2\pi)^3} \delta(\omega + p^2/2m^* - p'^2/2m^*) f_p',
\]

\[= n_3 \left(\frac{m^*}{2\pi k^2 T}\right)^{1/2} e^{-m^*(\omega - k^2/2m^*)^2/2k^2 T}, \]

(22)

which obeys \(S_3(k, -\omega) = e^{-\omega/T} S_3(k, \omega)\), we can rewrite Eq. (21) as,

\[
\nabla P_{ph} = -\int \frac{d^3q}{(2\pi)^3} \int \frac{d^3q'}{(2\pi)^3} \int d\omega \times q|\langle T \rangle|^2 \pi\delta(s - sq + sq') S_3(k, \omega)
\times \left[n_q (1 + n_{q'}) - n_{q'} (1 + n_q) e^{-\omega/T}\right].
\]

(23)

Since \(n_{q'} (1 + n_q) = n_q (1 + n_{q'}) e^{\omega/\pi k})\), the final line in Eq. (23) to first order in \(v_{ph}\) is \(n_3^0 (1 + n_{q'}) (\vec{k} \cdot \vec{v}_{ph})/T\). Finally, symmetrizing the integrand in \(q\) and \(q'\), and carrying out the angular averages, we have, in agreement with Bowley,

\[
\nabla P_{ph} = -\frac{\vec{v}_{ph}}{6T} \int \frac{d^3q}{(2\pi)^3} \int \frac{d^3q'}{(2\pi)^3} \int d\omega \times 2\pi\delta(s - sq + sq') S_3(k, \omega)
\times \left[n_q (1 + n_{q'}) - n_{q'} (1 + n_q) e^{-\omega/T}\right].
\]

(24)

Neglecting \(3^3\text{He}\) recoil is equivalent to setting \(S_3(k, \omega) = n_q \delta(\omega)\). In this case,

\[
\nabla P_{ph} = -\frac{\vec{v}_{ph}}{6T} \int \frac{d^3q}{(2\pi)^3} \int \frac{d^3q'}{(2\pi)^3} n_q^0 (1 + n_{q'}^0) \left|\langle T \rangle\right|^2 \times 2\pi\delta(s - sq')
\times \left[n_q (1 + n_{q'}) - n_{q'} (1 + n_q) e^{-\omega/T}\right].
\]

(25)

The phonon heat current density,

\[\dot{Q}_{ph} = s^2 \int \frac{d^3q}{(2\pi)^3} \dot{q} n_q = T S_{ph} \dot{v}_{ph}, \]

(26)

in steady state determines the phonon thermal conductivity, \(\dot{Q}_{ph} = -K_{ph} \nabla T\), and since \(\nabla P_{ph} = S_{ph} \nabla T\), we find the simple result [18]

\[K_{ph} = \frac{s \ell_{visc}}{s \ell_{ph}}, \]

(27)

where \(\ell_{ph} = 3T^2 S_{ph}/s n_3 \Gamma\) defines an effective mean free path for phonons scattering on \(3^3\text{He}\).

In Appendix A we include \(3^3\text{He}\) recoil to leading order in \(T/m^* s\), which we find increases the thermal conductivity by \(25\sim 33\%\) in the range \(T = 0.45\sim 0.65\) K. Figure 1 shows the phonon thermal conductivity computed from Eq. (A13), as well as the approximate thermal conductivity, Eq. (27), together with the measurements of Ref [3]. We use here the parameters \(A = -1.2 \pm 0.2\) [14, 20], and \(B = 0.70 \pm 0.03\) [19, 21], which lead to \(J = 2.2\pm 0.6\). The largest uncertainty is in \(A\), owing to a systematic difference between the measurements [14, 20] of the pressure dependence of the density of dilute solutions.

We return now to justify neglecting viscous stresses in the pressure equation [14]. The drag force per unit volume of the \(3\text{He}\) on the phonons is \(\sim \rho_{ph}s/\ell_{ph}\). We assume laminar flow in a circular pipe of radius \(R\), for which \(\nabla^2 \dot{v}_{ph} = -8(\dot{v}_{ph})/R^2\) is a constant throughout [22], thus overestimating the viscous force (density) on the phonons by neglecting the drag against the \(3^3\text{He}\). The phonon viscosity is \(\sim \rho_{ph}s\ell_{visc}/5\), where \(\ell_{visc}\) is the phonon mean
VI. APPLICATION TO EXPERIMENT

To further illustrate the physics, we now determine the temperature and concentration distributions for two geometries (Refs. [3] and [2]) of interest. Conservation of energy implies

$$\nabla \cdot \vec{Q} = -\nabla \cdot (K_{ph} \nabla T) = -C \nabla \left( \frac{\nabla T}{P_3} \right) = 0, \quad (32)$$

where, from Eq. (27), \( C = n_3^2 \beta_0 / 90 \pi J \). For simplicity, recoil effects are neglected in writing the equations here; however, they are taken into account in all numerical calculations reported below. We eliminate \( P_3 \) using the solution

$$P_3 + \frac{1}{4} S_{ph} T = P, \quad (33)$$

of Eq. (6), where the constant \( P \) is the total pressure of the excitations. Equation (32) then reduces to a partial differential equation in \( T \) alone (even when including the recoil corrections), which we solve using the finite element code FlexPDE [24].

Given \( T(\vec{r}) \), we determine \( n_3(\vec{r}) \) from Eq. (33), and determine the constant \( P \) by fixing the total number of \(^3\)He atoms in the system.

We begin by applying this theoretical description to the experiment of Rosenbaum et al. [3], where the thermal conductivity of mixtures in the concentration range \( 1.1 \times 10^{-4} \leq x_3 \leq 1.3 \times 10^{-2} \) were measured at temperatures \( 0.084 \leq T \leq 0.65 \) K. They determined the thermal conductivity by measuring the temperature difference over a 5 cm length of pipe, 0.26 cm in diameter, in the presence of a localized heat source. The pipe was connected to small reservoirs at either end containing the thermometers, the coupling to the dilution refrigerator and the heater.

In the analysis in Ref. [3], the effect of the variable \(^3\)He concentration in the pipe, and its attendant effect on the thermal conductivity, was not taken into account. We therefore determined the temperature and concentration distributions in the system by adjusting the heater power and minimum (dilution refrigerator) temperature to match the reported average temperature and the temperature difference implied by the reported thermal conductivity. We note that the \(^3\)He thermal conductivity is negligibly small owing to \( \ell_{visc} \ll \ell_{ph} \), Eq. (19).

Because the phonons push the \(^3\)He into the cold reservoir, the average concentration in the pipe is substantially lower than the concentration including the reservoirs. We show the results of our calculations in Fig. 11 for representative measurements [25] of Ref. [3]. The open circles are the thermal conductivities as reported, plotted at the average overall concentration; the solid circles are plotted at the average concentrations calculated for the pipe. Because the dimensions of the reservoirs are not given in Ref. [3], there is some uncertainty in the calculated result. The error bars shown in Fig. 11 represent changes of about \( \pm 25\% \) from the reservoir volumes estimated from the schematic in their Fig. 1. Given
FIG. 3: (Color online) Contribution of the thermoelectric transport term relative to the effective diffusion constant, $D_L$, extracted in Ref. lamoreaux. The temperatures are 0.45 K (dotted), 0.55 K (dashed) and 0.65 K (solid).

the uncertainties related to the geometry and the details of how the thermal conductivity was extracted, there is good agreement between this calculation and the measurement. As previously noted in Sec. III, the theoretical analysis in Ref. [3] incorrectly assumed that the primary phonon relaxation was due to $^3$He-phonon scattering, rather than phonon-phonon scattering along rays.

We now analyze the experiment of Lamoreaux et al. [4], which measures the $^3$He density response to a localized heat source by a novel technique. In the experiment, a dilute solution with concentration in the range $7 \times 10^{-5}$ to $3 \times 10^{-4}$ is contained in a cylindrical cell roughly 5 cm in diameter and 5 cm long, cooled at one end by a dilution refrigerator to a temperature $T$ in the range $0.45 - 0.95$ K [26]. A temperature gradient is created by a resistive heater generating 7–15 mW, located roughly midway between the ends and near the cylinder wall (see Fig. 4 below). The resulting spatial distribution of the $^3$He density is probed by a well-collimated neutron beam of diameter $\sim 0.25$ cm going through the cell. A fraction of the neutrons are captured via the reaction $n + ^3$He $\rightarrow p + t + 764$ keV. The XUV scintillation light from protons and tritons in the liquid $^4$He is converted to visible light and detected by a photomultiplier tube which views the solution through a window at the other end of the cell. The yield of scintillation light, measured as a function of cell temperature, initial $^3$He concentration, and heater power, is used to determine the thermally induced change in the $^3$He distribution. The heat flow again produces non-uniform temperature and $^3$He concentration distributions. Because the concentration and temperature gradients are directly proportional, Eq. (34), the concentration gradients implied by the measured scintillation yields here simply encode the same type of temperature difference information measured by Rosenbaum et al. We note that the proportionality constant involves $S_{ph} + n_3$, rather than just $S_{ph}$.

The response of the $^3$He density to the heat source is given here by Eq. (35). Combining this result with $\bar{Q} =$

Reference [4], which did not take into account the temperature gradient induced by the heat flow, interpreted the result for the $^3$He density gradient in terms of a simple diffusion constant, $D_L$, in the form,

$$\nabla n_3 = \frac{n_3}{D_L} \bar{v}_{ph}.$$
FIG. 5. (Color online) Representative results of the finite element calculation, including a function of the quantity \( \xi_L \) (column) density along the neutron beam as a function of the quantity \( \xi_L = T^4 (P/S_{ph}) \) where \( P \) is the heater power, along with data from Ref. [4] (circles, 0.45 K; diamonds, 0.55 K and squares, 0.65 K). The curves are for a refrigerator (and barrel) temperature of 0.45 K; diamonds, 0.55 K and squares, 0.65 K. The curves are plotted for the average temperature (greater than the boundary temperature) along the neutron beam path. For example, at the actual value \( T^4 P/S_{ph} = 2.5 \), the average \( \xi_L \) is shifted upward by \( \Delta \xi_L \approx 0.05 K^7 \text{cm}^3/\text{s} \) for \( T = 0.45 \text{ K} \), and by \( \Delta \xi_L \approx 0.5 K^7 \text{cm}^3/\text{s} \) for \( T = 0.65 \text{ K} \).

From Eq. (34) we see that

\[
D_L = \frac{n_3}{S_{ph} + n_3 S_{ph}} \frac{K}{S_{ph}}. \quad (36)
\]

Equivalently,

\[
D_L = D_3 - \frac{T}{S_{ph} + n_3} D_T = \frac{S_{ph}}{S_{ph} + n_3} D_3. \quad (37)
\]

The \( D_T \) term in this equation makes a significant contribution at higher concentrations (see Fig. 3); as noted, \( n_3 \sim S_{ph} \) for a large range of concentrations in the experiment of Ref. [4]. This effective \( D_L \) is not a simple diffusion constant, owing to the presence of the temperature gradient. We note that while the temperature dependence of the result for \( D_3 \) without \( ^3\text{He} \) recoil falls with temperature as \( 1/T^7 \), the temperature dependence of the effective \( D_L \) differs, owing both to recoil effects and the \( 1/T^3 \) dependence of \( n_3/S_{ph} \). The relative similarity of \( D_1 \) and \( D_3 \) also depends on the \( ^3\text{He} \) thermal conductivity being negligible.

An example of the \( ^3\text{He} \) distribution resulting from the finite element calculation for the Lamoreaux et al. experiment is shown in Fig. 4 results of the relative integrated (column) \( ^3\text{He} \) densities along the neutron beam are shown in Fig. 5 together with the data in Fig. 4 of [4]. To illustrate the effect of other variables in the problem, we also show in Fig. 5 the results for \( T = 0.65 \text{ K} \) and for the two concentrations used in the experiment. In this calculation we take the refrigerator end and the barrel of the cell to be fixed at \( T = 0.45 \) and 0.65 K as indicated; the opposite end of the cell is insulated. We note that the conductivity of the aluminum barrel is more than an order of magnitude larger than that of the phonons at these temperatures [27]. The relative column density calculated with an insulator in place of the aluminum barrel (not shown) falls well below the data. In this simulation we neglect any possibility of convective flow in the \( ^3\text{He} \).

The effect of the non-zero size of the neutron beam is shown in Fig. 5 together with the data in Fig. 4 of [4]. The relative similarity of \( D_1 \) and \( D_3 \) is shown in Fig. 5 the results for \( T = 0.65 \text{ K} \) and for the two concentrations used in the experiment. In this simulation we calculate contributions to the thermal conductivity due to the finite \( ^3\text{He} \) mass. We

VII. SUMMARY

We have laid out the basic transport theory of dilute solutions of \( ^3\text{He} \) in superfluid \( ^4\text{He} \) in the regime where scattering among the \( ^3\text{He} \) is the primary mechanism keeping the \( ^3\text{He} \) in thermal equilibrium, and phonons are the significant excitations of the \( ^4\text{He} \). We find that in the range of \( ^3\text{He} \) concentrations in the Rosenbaum et al. experiments, \( 7 \times 10^{-5} \leq x_3 \leq 1.5 \times 10^{-3} \), heat transport is dominated by phonons. The physical response of the system is not simple diffusion, described only by a \( ^3\text{He} \) diffusion constant, \( D_3 \), since the temperature gradients are important. The experiments can be characterized simply by a phonon thermal conductivity. This thermal conductivity, calculated in a microscopic framework, satisfactorily reproduces the measurements, indicating that the well-tested theory of \( ^3\text{He} \)-phonon scattering in dilute solutions of \( ^3\text{He} \) in superfluid \( ^4\text{He} \) is consistent with these experiments as well.

Acknowledgments

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Appendix A: Recoil corrections

In this Appendix we calculate contributions to the thermal conductivity due to the finite \( ^3\text{He} \) mass. We
The integral in Eq. (A1) is proportional to the inverse of the thermal conductivity. In the limit \( m^* \to \infty \) the expression in square brackets reduces to \( \delta (sq - sq') \). For finite \( m^* \) there are two effects. First, the structure factor, and therefore also the scattering rate, is reduced in magnitude because of the nonzero momentum transfer, \( \vec{k} = \vec{q} - \vec{q}' \), as is shown by the final Gaussian factor. Second, as the first Gaussian factor indicates, there is an energy transfer \( sq - sq' \) which is of order \( (Tk^2/m^*)^{1/2} \sim (T/m^*s^2)^{1/2}T \).

The contributions to the scattering amplitude for nonzero energy transfer and for nonzero velocity of the \(^3\text{He}\) atoms have not been investigated in detail, although the basic processes were discussed in full in Ref. [7]. Here we use Eq. (13) for the scattering amplitude. With prefactors omitted, the quantity to be calculated is thus

\[
\lim_{T \to 0} (TK)/TK \simeq 1 - \frac{1}{8m^*T} \langle k^4 \rangle / \langle k^2 \rangle, \tag{A3}
\]

where

\[
\langle \ldots \rangle = \int_0^\infty dq \int_{-1}^1 d\cos \theta \frac{(A + B \cos \theta)^2}{4 \sinh^2(sq/2T)} \langle \ldots \rangle. \tag{A4}
\]

In these integrals we may replace \( k^2 \) by its value \( 2q^2(1 - \cos \theta) \) for zero energy transfer. The integrals over \( q \) and \( \theta \) decouple and one finds

\[
TK \simeq \left( 1 + \frac{T}{4m^*s^2} \frac{\tilde{J}}{I_6} \right) \lim_{T \to 0} (TK), \tag{A5}
\]

where

\[
\tilde{J} = \int_{-1}^1 \frac{d\cos \theta}{2} (A + B \cos \theta)^2 (1 - \cos \theta)^2 = \frac{4}{3} A^2 - \frac{4}{3} AB + \frac{8}{15} B^2 \tag{A6}
\]

and

\[
I_n = \int_0^\infty dq \frac{x^n}{4 \sinh^2(x/2)} = n! \zeta(n), \tag{A7}
\]

where \( \zeta(n) \) is the Riemann zeta function of order \( n \). Therefore the thermal conductivity is given by

\[
K \simeq \left( 1 + \frac{25\pi^2}{11} \frac{\tilde{J}}{m^*s^2} \right) \frac{1}{T} \lim_{T \to 0} (TK). \tag{A8}
\]

We now calculate the leading correction to the thermal conductivity due to the energy transfer, which is found by neglecting the term \( k^2/8m^*T \) in the exponent in Eq. (A2). The Gaussian in the energy difference has a width small compared with \( T \), so we adopt a procedure similar to that used in making the Sommerfeld expansion for low temperature Fermi systems, where the derivative of the Fermi function approaches a delta function. In an integral of the form,

\[
G(x) = \int_0^\infty dy g(y) e^{-(x-y)^2/2\Delta^2}, \tag{A9}
\]
where $\Delta$ is a constant, and the function $g(y)$ varies slowly on the scale $\Delta$, one finds on expanding $g$ in a Taylor series about $y = x$, that

$$G(x) = g(x) + \frac{\Delta^2}{2} g''(x) + \ldots.$$  \hspace{1cm} (A10)

When this result is applied to the $q'$ integral in Eq. (A2) with the final Gaussian omitted, one finds

$$\int_0^\infty dq \int_0^\infty dq' \int_{-1}^1 d\cos \theta \left( \frac{m^* s^2}{2\pi T k^2} \right)^{1/2} \frac{k^2 (qq')^3 (A + B \cos \theta)^2 e^{-m^* s^2(q-q')^2/2k^2 T}}{4 \sinh^2(sq/2T)} \approx \frac{2q^3}{4 \sinh^2(sq/2T)} + \frac{T}{m^* s^2} \int_0^\infty dq \int_{-1}^1 d\cos \theta \frac{k^2 q^3 k^2 (A + B \cos \theta)^2}{2 \sinh(sq/2T)} \left( \frac{\partial^2}{\partial q'^2} \frac{k^2 q'^3}{2 \sinh(sq'/2T)} \right)_{q' = q},$$

which, when expressed in terms of the variables $x = sq/T$, $y = sq'/T$, and $z = \cos \theta$, is proportional to

$$\int_0^\infty dz \frac{2x^3}{\sinh(x/2)} + \frac{T}{m^* s^2} \int_0^\infty dz \frac{x^3}{2 \sinh(x/2)} \int_{-1}^1 \frac{dz}{2} (1 - z)(A + Bz)^2 \left( \frac{\partial^2}{\partial y^2} \frac{\kappa^2 y^3}{2 \sinh(y/2)} \right)_{y = x},$$

where $\kappa^2 = x^2 + y^2 - 2xyz$. The second derivative is

$$\frac{x^3}{\sinh(x/2)} \left[ 1 + (1 - z) \left( 12 - \frac{1}{4} x^2 - 4x \coth(x/2) + \frac{1}{2} x^2 \coth(x/2)^2 \right) \right].$$

On evaluating the integrals, one finds that the effect of energy transfer produces contributions to the integral \(\text{A2}\) having the form

$$1 + \left( 1 - \frac{25\pi^2 - 198}{33} \frac{\hat{J}}{J} \right) \frac{T}{m^* s^2}. \hspace{1cm} (A14)$$

The numerical factor $25\pi^2 - 198/33$ is approximately 1.477 and thus one sees that the effects of nonzero energy transfer are much less important than those due to nonzero momentum transfer. When both contributions to the thermal conductivity are included, one finds on inserting Eq. (27) for the low temperature limiting behavior,

$$K \simeq \frac{n_4 s^2}{90 \pi x_3 JT} \left( 1 + \frac{T}{m^* s^2} \left[ \frac{100\pi^2 - 198 \hat{J}}{33} - 1 \right] \right). \hspace{1cm} (A15)$$

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One should not be alarmed by the apparent divergence as $x_3 \to 0$. When $x_3$ becomes sufficiently small, the $^3$He no longer provide the mechanism for absorbing momentum from the phonons, and rather the phonon thermal conductivity becomes limited by viscous transfer of momentum from the phonons to the container walls [18].

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