Ultrasonic Attenuation in Liquid⁴ He under Pressure^{*}

Pat R. Roach, J. B. Ketterson, and M. Kuchnir Argonne National Laboratory, Argonne, Illinois 60439 (Received 6 December 1971)

In an attempt to help resolve previously observed discrepancies between the theory for ultrasonic attenuation and the data at the vapor pressure, we have extended our measurements to higher pressures. We report data on the temperature dependence of the attenuation from 0. 1 to 1.⁰ K at frequencies of 15, 45, 105, and 256 MHz for pressures between ⁰ and 24. ⁷ atm. At low pressures or temperatures, the data resemble the theoretical predictions, but the measured attenuation is about twice that predicted by most theories. At higher pressures and for temperatures above about 0. 3 K, the attenuation shows a sharp reduction below the value extrapolated from low temperatures according to previous theories. It has recently been suggested that this unexpected behavior is due to a restriction on the number of thermal phonons that are able to play a part in the attenuation process at higher temperatures and pressures.

INTRODUCTION

At very low temperatures, where there is a negligible roton population, the attenuation of sound in liquid He is thought to arise from three-phonon events. In the limit $\omega \tau_{\text{pp}} \gg 1$, where ω is the sound angular frequency and τ_{pp} is the appropriate thermalphonon relaxation time, the attenuation has generally been thought to be given by'

$$
\alpha(\omega, T) = \frac{\pi^2}{30} \frac{(u+1)^2}{\rho} \frac{k_B^4}{\hbar^3 c^6} \omega T^4
$$

$$
\times \left[\tan^{-1} (2\omega \tau_{\text{pp}}) - \tan^{-1} (3\gamma \bar{p}^2 \omega \tau_{\text{pp}}) \right], \qquad (1)
$$

where ρ is the density, $\,c\,$ is the sound velocity $u \equiv (\rho/c) (\partial c/\partial \rho)$ is the Grüneisen constant, $\bar{p} \equiv 3k_BT/c$ is the average thermal-phonon momentum, and γ is the dispersion constant defined by the relation $\epsilon = c p (1 - \gamma p^2)$, where ϵ and p are the energy and momentum, respectively, of an elementary excitation.

In previous experiments at the vapor pressure² the measured temperature dependence of the attenuation was in qualitative agreement with the $T⁴$ law but was larger than that predicted by Eq. (1) by about a factor of 2. The frequency dependence, measured in the range 12-208 MHz, was more complex; for temperatures below about 0. 3 K an approximately linear behavior was observed while for temperatures between 0. 3 and 0. 4 the frequency dependence fell below linear above 36 MHz.

Several attempts to explain the previous data have involved the assumption that γ is negative (anomalous dispersion}. Maris and Massey' pointed out that for negative γ the limit $|3\gamma \bar{p}^2 \omega \tau_{\rm m}| \gg 1$ results in twice the usually expected magnitude of the attenuation according to Eg. (1) (the arctangent functions are then additive). A lack of knowledge of tenuation according to Eq. (1) (the arctangent fu
tions are then additive). A lack of knowledge of
 τ_{pp} , however, made it unclear whether this sug-
gestion could be applied to our data. Very rece gestion could be applied to our data. Very recently, $Maris⁴ presented a calculation which assumes$

anomalous dispersion but differs from $Eq. (1)$ in that no relaxation-time approximation is made for the three-phonon collision process. Preliminary comparisons at 0. 35 K with our previous data show fairly good agreement both in magnitude and frequency dependence.

The present investigation was initiated in order to test how well the theories work at elevated pressure. A preliminary report of this work has appeared previously. '

EXPERIMENTAL TECHNIQUE

The sound cell with which these data were taken was attached to the copper mixing chamber of a 3 He- 4 He dilution refrigerator which regularly cools below 20 mK. Temperatures were determined by measuring the susceptibility of single crystals of cerium magnesium nitrate which were thermally anchored to the sound cell via coil foil. To ensure thermal contact between the liquid 4 He being measured and the copper walls of the sound cell to which the magnetic thermometer was attached, the sound cell had added to it a chamber filled with sintered copper in contact with the sample liquid. The sound was generated and detected by separate 15-MHz unloaded x -cut quartz transducers. The transducers were separated by a 2. 02-cm-long hollow quartz spacer. The phase and amplitude of the received sound signal can be compared with a signal coming from the oscillator in order to determine both changes in the time delay through the liquid (due to changes in the sound velocity) and changes in the attenuation through the liquid. Details of the sound cell and the ultrasonic comparator have been described previously.⁶

EXPERIMENTAL RESULTS

Figures 1-4 show our ultrasonic-attenuation measurements at frequencies of 15, 45, 105, and 256 MHz, respectively. At each frequency a number

 $\overline{5}$

2205

FIG. 1. Temperature dependence of the measured attenuation for a frequency of 15 MHz at various values of pressure. The solid lines are smooth curves through the data.

of attenuation-vs-temperature isobars are shown for pressures up to 24. 7 atm. The data at 105 MHz have been reported previously.⁵ Earlier measurements^{$7,8$} of the attenuation under pressure have shown the general decrease in attenuation with increasing pressure that we have observed. However, these previous measurements did not extend above 15 MHz in frequency and did not show the features to be discussed for our data. We have compared our data with the theory of Khalatnikov and Chernikova⁹ using the data of Van den Meijdenberg. Taconis, and De Bruyn Ouboter¹⁰ for the pressure dependence of the roton parameters and using our previous ultrasonic measurements¹¹ for the pressure dependence of ρ , c , and u . Figure 5 shows curves of the theory at 105 MHz and various pressures corresponding to some of the experimental curves. These curves are representative of the theory for the entire frequency and pressure range of our experiment. They show a well-defined $T⁴$ temperature dependence at low temperatures and a peak in the attenuation at around 1 K. Our

data, however, show this behavior only at low pressures and even then the measured attenuation is consistently larger than that predicted by theory. . At higher pressures the data show a sharp departure from theory; in the vicinity of 0. ⁵ K the higher pressure data show a shoulder followed by a rapid increase of the attenuation with temperature until the vicinity of the maximum is reached at about 1 K. At the highest pressure (24. 7 atm) only the very rapid rise below the maximum is seen; neither the $T⁴$ dependence nor the shoulder could be resolved in the present experiment.

Tables I-IV give most of our data in numerical form. The accuracy with which the attenuation is given in the tables $(\pm 0.005 \text{ dB/cm})$ is only meaningful at the lower values of attenuation. At attenuations above 10 dB/cm , signal-to-noise problems limited our accuracy to several percent in the attenuation measurements. Because the data are presented on logarithmic scales, it is very important that the zero of attenuation be accurately known. In the present experiment this was empirically de-

FIG. 2. Temperature dependence of the measured attenuation for a frequency of 45 MHz at various values of the pressure. The solid lines are smooth curves through the data.

FIG. 3. Temperature dependence of the measured attenuation for a frequency of 105 MHz at various values of the pressure. The solid lines are smooth curves through the data. IO.O

termined for each of the curves shown by finding that value which made the low-temperature part of the curve most nearly linear (on a log-log plot) and/or consistent with the curves for other pressures. This seldom involved a correction of more than 0.01 dB/cm to the data. The accuracy quoted in the tables for the temperature $(±0.1$ mK) is at best useful only as a measure of the relative temperatures of the points. The absolute accuracy of our temperature calibration is estimated to be a few percent while our temperature resolution is everywhere better than 1% . Our pressures were measured on a fused quartz bourdon gauge. 12 Although they are quoted to only \pm 0.1 atm, in fact the pressures were held to ± 0.001 atm during the taking of data at any of the given pressures.

DISCUSSION

In the time since the original presentation⁵ of parts of this data, several features of the data have been more or less successfully accounted for theoretically.

Klein and Wehner^{13, 14} have considered the pres-

sure dependence of our new data in the low-temperature T^4 region. They do not account for the magnitude of the data but they observe that the pressure dependence of the attenuation is not completely explained by the pressure dependence of u , ρ , and c ; they conclude that, in addition, a negative value of γ combined with its pressure dependence as measured by Phillips et al. ¹⁵ is necessa dence as measured by Phillips ${\it et \ al.}^{\text{15}}$ is necessar to explain the data. Since Maris⁴ has had some success in accounting for the magnitude of our zero-pressure data, it seems reasonable to anticipate that his calculations will yield equally good agreement with our high-pressure data.

Jäckle and $Kehr^{16}$ have suggested an explanation for the shoulder in our data. They assume that the phonon spectrum is initially curved upwards as in Fig. 6 (where the effect is exaggerated). For this situation there will be a momentum q_a above which thermal phonons cannot absorb an ultrasonic phonon by the three-phonon process. This momentum is the point on the spectrum where the group velocity v of a thermal phonon is equal to the velocity c of

FIG. 4. Temperature dependence of. the measured attenuation for a frequency of 256 MHz at various values of the pressure. The solid lines are smooth curves through the data.

FIG. 5. Temperature dependence of the attenuation as calculated from the theory of Khalatnikov and Chernikova for 105 MHz and for pressures corresponding to some of the experimental values.

an ultrasonic phonon $(q \approx 0)$. When a temperature is reached where a significant fraction of the thermal phonons have momenta above q_c , then the attenuation due to the three-phonon process will begin to be suppressed. An additional assumption is made that q_c decreases considerably with pres-

FIG. 6. Possible form of anomalous dispersion (greatly exaggerated) at two different pressures. q_c is the momentum above which thermal phonons cannot absorb an acoustic phonon by the three-phonon process.

| Temp. | | Attenuation |
|--------|-------------|-------------|
| (mK) | | (dB/cm) |
| | | |
| | $P=0.0$ atm | |
| | А | |
| | | |
| 148.4 | | 0.050 |
| 180.2 | | 0.125 |
| 209.8 | | 0.240 |
| 245.7 | | 0.530 |
| 282.0 | | 1.000 |
| 347.4 | | 2.275 |
| 401.1 | | 3.635 |
| 448.6 | | 4.975 |
| 523.5 | | 7.580 |
| | | |
| | B | |
| | | |
| 127.5 | | 0.030 |
| 140.5 | | 0.050 |
| 155.1 | | 0.068 |
| 170.4 | | |
| | | 0.100 |
| 185.0 | | 0.146 |
| 204.8 | | 0.220 |
| 223.4 | | 0.336 |
| | | |
| | $P=1.1$ atm | |
| | А | |
| | | |
| 132.4 | | 0.025 |
| 141.1 | | 0.030 |
| 151.7 | | 0.045 |
| 162.1 | | 0.065 |
| 184.9 | | 0.120 |
| 205.8 | | 0.195 |
| 223.7 | | 0.270 |
| 249.2 | | 0.435 |
| 276.0 | | 0.705 |
| | | |
| | В | |
| | | |
| 227.4 | | 0.270 |
| 271.6 | | 0.630 |
| 331.0 | | 1.430 |
| 394.2 | | 2.610 |
| 499.2 | | 5.095 |
| 595.8 | | 7.950 |
| 771.8 | | 15.170 |
| 809.8 | | 17.450 |
| 833.3 | | 18.835 |
| 860.3 | | 20.010 |
| 869.0 | | 20.195 |
| 869.0 | | 20.310 |
| | | |
| 889.1 | | 20.760 |
| 898.4 | | 21.010 |
| 905.5 | | 21.010 |
| 910.2 | | 21.110 |
| 922.4 | | 21.160 |
| 955.6 | | 21.260 |
| 980.1 | | 21.360 |
| 1026.7 | | 21.710 |
| 1134.8 | | 19.060 |
| 1259.1 | | 12.260 |

attenuation in liquid He under pressure at 14.92 MHz.

sure. For pressures below 10 atm, q_c must be above the range of thermal phonons $(c_1q_{c1} > 3k_BT)$, while at the highest pressures, q_c must be within this range $(c_2q_{c2} \leq 3k_BT)$ so that the attenuation is suppressed. Jäckle and Kehr give the attenuation for the restricted three-phonon process as

$$
\alpha(\omega, p, T) = A \omega T^4 F(z) / F(\infty) , \qquad (2)
$$

where $z = cq_c/(k_B T)$ and $F(z) = \int_0^z dx x^4 f(x) [1 + f(x)];$ $f(x)$ is the Bose function. At low temperatures $(k_BT \ll cq_c)$ this becomes $\alpha = A\omega T^4$, which is equivalent to Eq. (1). For high temperatures $(k_B T \gg c q_c)$ the three-phonon process is restricted, and one obtains

$$
\alpha = \frac{1}{3}A\left(cq_c/k_B\right)^3 \omega T \tag{3}
$$

Many of our curves do show such a linear temperature dependence in the vicinity of the shoulder. Jäckle and Kehr also treat the rapid rise of the attenuation above the shoulder. They consider the effect of thermal-phonon lifetime on the threephonon process in the region above q_c where the process is not favorable because of an energy deficit. In the temperature region above 0. 6 K, where the number of rotons is increasing very rapidly, they find that the attenuation, although strongly depressed, is rising very steeply with temperature. Figure 7 shows their fit to our data at a pressure of 16.4 atm for frequencies of 15 and 105 MHz. The fitted curve for 105 MHz also includes a contribution from roton viscosity. Although the fit

TABLE II. Temperature dependence of the sound

TABLE III. Temperature dependence of the sound attenuation in liquid 4 He under pressure at 105.05 $\frac{M}{2}$

FIG. 7. Theory of Jäckle and Kehr (Ref. 16). Attenuation at 16.4-atm pressure and for frequencies of 15 (lower curves) and 105 MHz (upper curves). Solid points: experimental measurements. Dashed lines: basic threephonon result $\alpha \omega T^4$. Dash-dotted lines: restricted three-phonon process. Full line for 15 MHz: sum of restricted three-phonon and lifetime-induced attenuation. Full line for 105 MHz: the same, plus contribution of the roton viscosity.

TABLE IV. Temperature dependence of the sound attenuation in liquid 4He under pressure at 256. 22 MHz.

but also from the calculation by Maris⁴ of the magnitude and frequency dependence of our zero-pressure data and from the explanation by Jäckle of the shoulder in our data, isthat all these theories require an assumption of anomalous dispersion to account for our data. This has to be interpreted as very

*Based on work performed under the auspices of the U. S. Atomic Energy Commission.

¹C. J. Pethick and D. ter Haar, Physica 32, 1905 (1966); see also, S. G. Eckstein, Y. Eckstein, J. B. (1966); see also, S. G. Eckstein, Y. Eckstein, J. 1
Ketterson, and J. H. Vignos, in *Physical Acoustics*
edited by W. P. Mason and R. N. Thurston (Academ
Naw York, 1970), Val. 6, and the references earts edited by W. P. Mason and R. N. Thurston (Academic New York, 1970), Vol. 6, and the references contained therein.

²B. M. Abraham, Y. Eckstein, J. B. Ketterson, M. Kuchnir, and J. H. Vignos, Phys. Rev. 181, ³⁴⁷ (1969).

 3 H. J. Maris and W. E. Massey, Phys. Rev. Letters 25, 220 (1970); also, S. Havlin, thesis (Tel-Aviv University) (unpublished).

 4 H. J. Maris, Phys. Rev. Letters 28 , 277 (1972). 5 Pat R. Roach, J. B. Ketterson, and M. Kuchnir, Phys. Rev. Letters 25, 1002 (1970).

 6 B. M. Abraham, Y. Eckstein, J. B. Ketterson, and J. H. Vignos, Cryogenics 9, ²⁷⁴ (1969).

 7 W. M. Whitney, Phys. Rev. 105, 38 (1957).

 8K . Dransfeld, J. A. Newell, and J. Wilks, Proc. Boy. Soc. (London) A243, 500 (1958).

 9 I. M. Khalatnikov and D. M. Chernikova, Zh. Eksperim.

strong evidence for anomalous dispersion. Whether this dispersion should be described by a quadratic or a cubic term in the energy spectrum remains to be determined; more detailed theoretical calculations using both descriptions would be very desirable.

i Teor. Fiz. 49, 1957 (1965); 50, 411 (1966) [Sov. Phys. JETP 22, 1336 (1966); 23, 274 (1966)].

 10 C. J. N. Van den Meijdenberg, K. W. Taconis, and R. De Bruyn Ouboter, Physica 27, 197 (1961).

B. M.. Abraham, Y. Eckstein, J. B. Ketterson, M. Kuchnir, and Pat R. Roach, Phys. Rev. A $\frac{1}{1}$, 250 (1970).

¹²Texas Instruments Inc. , Houston, Tex.

 13 R. Klein and R. K. Wehner, Phys. Rev. Letters 23 , 1372 (1969); in Proceedings of the Twelfth International Conference on Low-Temperature Physics, Kyoto, Japan, 1970, edited by Eizo Kanda (Academic Press of Japan, Tokyo, 1971), p. 91.

 14 R. Klein and R. K. Wehner, Helv. Phys. Acta 44, 550 (1971).

 $15N$. E. Phillips, C. G. Waterfield, and J. K. Hoffer, Phys. Bev. Letters 25, 1260 (1970).

 16 J. Jäckle and K. W. Kehr, Phys. Rev. Letters 27, 654 (1971).

 ^{17}E . Feenberg, Phys. Rev. Letters 26, 301 (1971).

 18 A. Molinari and T. Regge, Phys. Rev. Letters 26, 1531 (1971).

 19 G. Barucchi, G. Ponzano, and T. Regge (unpublished).

PHYSICAL REVIEW A VOLUME 5, NUMBER 5 MAY 1972

Liquid-Helium Configuration around a Metastable Excited Helium Atom †

J. P. Hansen* and E. L. Pollock

Laboratory of Atomic and Solid State Physics, Cornell University, Ithaca, New York 14850 (Received 22 November 1971)

The pair distribution for normal He atoms in the vicinity of a metastable $2³S$ He atom is calculated in the zero-temperature limit, from first principles, using a variational Jastrow wave function and the Percus- Yevick integral equations for fluid mixtures. From this pair function, the "bubble radius, " coordination number, and energy per metastable atom are calculated.

Spectroscopic studies of the infrared emission and absorption spectra of neutral localized excitations, created in liquid helium by electron beams with an energy of the order of 10^2 keV, have identified these as metastable excited states of He atoms and $He₂$ molecules shifted only very slightly in energy from their free atomic or molecular values. $1-3$ For the atomic states, Hickman and Lane, by extending the calculations of Jortner $et al.^4$ for an excess electron in liquid helium to include the interaction of a helium core with both the excited electron and the liquid, have shown these observations to be consistent with the existence of a cavity surrounding the excited atom.⁵

The purpose of this paper is to study these "bubble states" from first principles, for the special case of metastable $2³S$ atoms (hereafter symbolized by He*) which have been observed at concentrations greater than 10^{12} atoms $\rm cm^{-3}$ by the Rice University group.^{1,2} The calculation presented here could be extended to other excitations if the pair interaction between such excitations and normal He atoms were known.

The Hamiltonian for a system of N_1 normal and