A Quantum Monte Carlo Study of Static Properties of One $^3$He Atom in Superfluid $^4$He

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Abstract

The local environment and the energetic properties of one $^3$He atom solved in bulk superfluid $^4$He are studied by means of the diffusion Monte Carlo method. The chemical potential of the $^3$He impurity is calculated with a generalized reweighting method which allows for a reliable estimation of this quantity. Results for the chemical potential, radial distribution and structure functions, volume-excess parameter, and effective mass are given for several pressures and compared with available experimental data. An overall agreement with experiment is obtained except for the kinetic energy of the $^3$He atom which, in accordance with previous theoretical estimations, appears to be considerably larger than determinations from deep-inelastic neutron scattering.

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I. INTRODUCTION

Isotopic $^3$He-$^4$He mixtures have deserved theoretical and experimental interest for many years due to their unique properties. Among them one may recognize the only isotopic mixture which remains stable at a certain $^3$He concentration down to zero temperature, and the only liquid system in which the two quantum statistics, bosons ($^4$He) and fermions ($^3$He), are put together and one influences the other through the interatomic potential. As a result of this interplay, it has been observed both experimentally and theoretically, that the $^4$He superfluid fraction decreases and simultaneously the $^4$He condensate fraction increases when the $^3$He concentration increases. On the other hand, the $^3$He momentum distribution in the mixture appears largely influenced by the presence of $^4$He showing a considerably larger depletion above the Fermi momentum in comparison with pure $^3$He. Experimental information on the $^4$He condensate fraction ($n_0$) and the kinetic energy of both $^4$He and $^3$He in the mixture have been recently extracted from deep-inelastic neutron scattering. These analysis show a large enhancement of $n_0$ with respect to pure $^4$He, and $^3$He kinetic energies very similar to the ones of pure $^3$He. In contrast, all the theoretical calculations have shown only an small increment of $n_0$ when the $^3$He concentration ($x$) increases (mainly due to the change in the total density at a fixed pressure) and a $^3$He kinetic energy appreciably larger.

The maximum solubility of $^3$He in $^4$He is $x^m = 0.066$ at zero pressure and presents a maxim value of $x^m = 0.095$ at $P \approx 10$ atm. These $x$ values are sufficiently small to stimulate the theoretical interest in describing microscopically the limit of zero $^3$He concentration which, on the other hand, has also been experimentally analyzed and a number of characteristic properties are nowadays available. From a theoretical viewpoint, this limiting system has been studied considering a single $^3$He atom solved in bulk $^4$He. The most useful approach in the past has been the variational method combined with the resolution of the hypernetted chain equations (HNC) coupled or not to an Euler-Lagrange optimization procedure. The results obtained with this approach reproduces the energetic and struc-
tural properties of the system with a good accuracy but the impurity effective mass appears slightly underestimated. The application of Monte Carlo methods, both variational and \textit{ab initio}, to the impurity system in order to calculate a basic property as the chemical potential of the impurity in the bulk ($\mu_I$), has been seriously hindered by the fact that $\mu_I$ results from the difference of two energy terms of order $N$, $N$ being the number of particles. In fact, a straightforward application of the Monte Carlo method cannot estimate $\mu_I$ because the statistical fluctuations would mask it completely.

In the present calculation, the reported results for $\mu_I$ have been obtained using a particular reweighting procedure suitable for the diffusion Monte Carlo (DMC) method, which has allowed a direct calculation of $\mu_I$ with a statistical error reduced to a manageable level. Using this method, we have been able to obtain reliable results for $\mu_I$ that fit accurately the experimental data from the equilibrium up to the freezing $^4$He densities. The local environment of the impurity, reflected in the crossed radial distribution and structure functions, has been studied by means of a pure estimator to remove the bias associated to the trial wave function. We have, finally, focused our attention to the calculation of the impurity effective mass and its kinetic energy for several densities. As in previous quantum Monte Carlo (QMC) applications, the effective mass is extracted from the diffusion coefficient in imaginary time and, in spite of some uncertainties inherent to the extrapolated estimator used in this calculation, a reasonable agreement with recent experimental determinations is attained. Our results concerning the kinetic energy of the impurity, derived from the Hellmann-Feynman theorem, avoid the residual importance-sampling dependence and show values which are definitely larger than the experimental data, as pointed out previously in variational and path integral Monte Carlo (PIMC) calculations.

The outline of the paper is as follows. In the next section we briefly introduce the DMC algorithm for the impurity system and present a DMC reweighting technique that permits a direct estimation of arbitrarily small differences. In Sec. III, we present the results and compare them with available experimental and theoretical data. We close in Sec. IV with the summary and final remarks.
II. THE DIFFUSION MONTE CARLO METHOD WITH REWEIGHTED CONFIGURATIONS

The DMC method allows for a very accurate description of the ground-state properties of an interacting $N$-body system. In the DMC formulation the imaginary-time Schrödinger equation for the function $f(R, t) = \Psi_T(R)\Phi(R, t)$,

$$-\frac{\partial f(R, t)}{\partial t} = \sum_{i=1}^{N} -D_i \left[ \nabla_i^2 f(R, t) - \nabla_i \cdot (F_i(R) f(R, t)) \right] + (E_L(R) - E) f(R, t) , \quad (1)$$

is turned into a stochastic process which provides a sample of configuration points $R$ (walkers) and weights $w(R)$ in a $3N$-dimensional space, whose probability distribution is given by $f(R, t)$. $\Psi_T(R)$ is a time-independent trial wave function that acts as an importance-sampling function, and $\Phi(R, t)$ is the exact wave function of the system. In this form, the Schrödinger equation appears as a diffusion-like differential equation with a diffusion, drift and branching terms corresponding to the first, second and third terms of the rhs of Eq. (1), respectively. In Eq. (1), $E_L(R) = \Psi_T(R)^{-1}H\Psi_T(R)$ is the local energy, $F_i(R) = 2\Psi_T(R)^{-1}\nabla_i\Psi_T(R)$ is the quantum drift force, and $D_i = \hbar^2/(2m_i)$ acts as the free-diffusion constant of the $i$ particle. At sufficiently long imaginary times the probability density evolves to a stationary solution given by $\Phi_0(R)\Psi_T(R)$, $\Phi_0(R)$ being the ground-state wave function from which the exact ground-state energy is obtained as the average of the local energy $E_L(R)$.

Let us now turn to the implementation of the reweighting method. In the DMC algorithm, the distribution probability of the walkers is modified in every single operation. Consider in particular the stochastic process originated by the diffusion term, which is a random gaussian displacement $R \rightarrow R'$. The new weight and distribution probability are $w'(R') = w(R)$ and $p'(R') = \int e^{-\frac{(R'-R)^2}{4D\Delta t}} p(R) \, dR$, respectively. In this stochastic process we can make use again of importance sampling in order to perform a modified diffusion random displacement. In this case, if the transition probability of going from $R$ to $R'$ following a modified diffusion process is $G(R' - R)$, the new distribution probability is given by
\[ p'(R') = \int G(R' - R) \, p(R) \, dR . \]  

(2)

The statistical sample of walkers provides unchanged averaged values if one uses accordingly a new weight given by

\[ w'(R') = \frac{e^{-\frac{(R' - R)^2}{4D\Delta t}}}{G(R' - R)} \, w(R) . \]  

(3)

This means that a system can be studied using a variety of diffusion random laws, although the efficiency of the method will be related to the magnitude of the changes. In general, the modification has to be small enough so that the same configuration space is sampled.

The reweighting method is specially useful in the calculation of differences between two almost identical systems. Performing independent samplings for both systems generates a global uncorrelated noise that prevents a direct measure of the difference. Assume, however, that given a common starting configuration point, a single drift process brings both walkers to new positions \( R_1 \) and \( R_2 \) which are very close (in particular, separated a distance much smaller than the typical size of a random gaussian displacement). The configuration region attainable after a combined drift and diffusion process is the same, and the transition probabilities \( G_1(R' - R_1) \) and \( G_2(R' - R_2) \) are almost equal. Equations (2) and (3) may be used then to change \( G_1(R' - R_1) \) into \( G_2(R' - R_2) \). Therefore, there is no need of taking averages using two independent walkers for the two systems, and it may be highly preferable to use correlated walkers, in the sense of carrying a single random walk to obtain statistical values for both systems. Furthermore, notice that this technique may be applied to modify the diffusion process of the whole walker, i.e., all the particles of the system, or only a subset of it.

The generalized reweighting method is an appropriate tool for studying the quantum liquid in which we are now interested. It is composed by \( N - 1 \) \( ^4 \)He particles and one \( ^3 \)He atom (I) enclosed in a simulation box with periodic boundary conditions. The Hamiltonian of the system is

\[ H = -D_4 \sum_{i=1}^{N-1} \nabla_i^2 - D_I \nabla_I^2 + \sum_{i<j}^{N} V(r_{ij}) , \]  

(4)
and the trial wave function $\Psi_T(R)$ has been chosen to be of Jastrow type

$$\Psi_T(R) = \exp \left( \sum_{i<j}^N u(r_{ij}) \right) \tag{5}$$

without distinguishing between the $(4, I)$ and $(4, 4)$ pairs of particles. This simplification in the wave function, known as average correlation approximation (ACA), has been used in several variational calculations, obtaining a quite good description of the impurity properties. In the DMC method the trial wave function acts only as a guiding wave function for the walkers driving them to regions where $\Phi_0(R)$ is expected to be large and thus a particular choice, as the ACA one in the present case, does not bias the expected value for the ground-state energy. On the other hand, the sign problem that would emerge in a simulation of a finite $^3$He concentration in $^4$He does not appear here and an exact energy for the system, apart from statistical uncertainties, can be safely obtained.

From the energetic viewpoint, the more fundamental quantity in the study of the $^3$He impurity in $^4$He is the impurity chemical potential or binding energy

$$\mu_I = \langle H(N+I) \rangle_{N+I} - \langle H(N) \rangle_N, \tag{6}$$

both energy estimations being evaluated at fixed volume $\Omega = N/\rho$. If the total number of particles is also conserved, and therefore one $^4$He atom is substituted by the $^3$He impurity, $\mu_I$ is given by

$$\mu_I = \mu_4 + \left( \langle H((N-1)+I) \rangle_{(N-1)+I} - \langle H(N) \rangle_N \right). \tag{7}$$

We have chosen the second option in which the difference between the two energy estimations is much less density dependent than in Eq. (3), and moreover because it is more convenient if a correlated estimation of the difference is intended. The pure $^4$He chemical potential $\mu_4$ entering in Eq. (7) has been determined in a previous work using also the DMC method with a nice agreement with experimental data.

The drawback of an ab initio MC estimation of $\mu_I$, that has precluded such a calculation for years, is that an independent calculation of $\langle H((N-1)+I) \rangle_{(N-1)+I}$ and $\langle H(N) \rangle_N$ followed by its difference, produces a result completely hidden by the statistical error. In order
to overcome this serious problem, we have directly sampled the difference by means of the reweighting method above introduced. Our purpose was to perform two correlated DMC runs, one of bulk $^4$He and the other with one $^3$He impurity. In this case, Eq. (2) has allowed us to use the same environment for both the $^3$He atom in the impurity system and the equivalent $^4$He atom in the pure liquid. In fact, the drift of the surrounding $N - 1$ particles in Eq. (2) is almost insensitive to the mass of the impurity, i.e., the resulting positions in the impurity system ($\mathbf{R}'_{N-1}$) and in the pure phase ($\mathbf{R}^4_{N-1}$) are very close. If one decides to change the diffusion process of the environment in the pure system with

$$G(\mathbf{R}'_{N-1} - \mathbf{R}^4_{N-1}) \equiv \exp \left( -\frac{(\mathbf{R}'_{N-1} - \mathbf{R}^I_{N-1})^2}{4D_4\Delta t} \right),$$

then the distribution probability of the environment, after the double process of drift and diffusion, is the same for the two systems. In this form, the statistical fluctuations coming from regions far from the impurity and its corresponding $^4$He atom cancel exactly, and the remaining signal corresponds only to their local environment making feasible a direct estimation of $\mu_I$.

In addition to the impurity chemical potential $\mu_I$, the knowledge of other properties as the crossed radial distribution function $g^{(4,I)}(r)$, the impurity effective mass and its kinetic energy are also relevant in a microscopic characterization of the $^3$He impurity. Expectation values of operators $\mathcal{O}$ that do not commute with the Hamiltonian $H$ are however biased because the probability density is $\Psi_T(\mathbf{R})\Phi_0(\mathbf{R})$ and not $|\Phi_0(\mathbf{R})|^2$. Thus, the natural expectation values, called mixed estimators ($m$), have to be corrected in order to reduce or eliminate this systematic source of error. In the extrapolation methods\[21\] this correction is approximated by

$$\langle \mathcal{O} \rangle_{el} = 2 \langle \mathcal{O} \rangle_m - \langle \mathcal{O} \rangle_v ,$$

or

$$\langle \mathcal{O} \rangle_{eq} = \frac{\langle \mathcal{O} \rangle^2_m}{\langle \mathcal{O} \rangle^2_v} ,$$

(10)
\[ \langle O \rangle_v \] being a variational Monte Carlo (VMC) estimation. Both \( \langle O \rangle_{el} \) and \( \langle O \rangle_{eq} \) are accurate to first order in \( \delta \Psi \), with \( \Psi_T = \Phi_0 + \delta \Psi \), but in general it is not enough to completely eliminate the influence of \( \Psi_T \) in \( \langle O \rangle \). In order to go beyond this approximation, we have used for the expectation values of coordinate operators the pure estimators following the methodology of Ref. [13] based on the future-walking strategy [22]. As proved in pure \( ^4 \)He [13], the pure estimator removes all the dependence on \( \Psi_T \) providing results as exact as the ones for the total energy.

Derivative operators as the kinetic energy cannot be evaluated with the pure estimator, and the extrapolation methods generate more unreliable results than in the case of \( O(R) \). In a pure phase it is not a severe problem because the kinetic energy can be calculated through the difference \( E/N - V/N \), \( V/N \) being the pure estimation of the potential energy. That it is not obviously possible in the impurity system because the total energy includes the kinetic energy of the medium and the one of the \( ^3 \)He impurity. To overcome this difficulty and go to an unbiased estimation of the \( ^3 \)He kinetic energy one can invoke the Hellmann-Feynman theorem [23]. It states that

\[
\langle T_I \rangle = D_I \frac{\partial E}{\partial D_I}, \tag{11}
\]

\( E \) being the exact ground-state energy. We have then evaluated \( T_I \) discretizing the derivative \( \partial E/\partial D_I \) and computing the difference in the total energy (with \( \Delta D_I/D_I = 0.1 - 10\% \)) by means of the generalized reweighting method.

\[ \text{III. RESULTS} \]

The microscopic properties of a \( ^3 \)He atom immersed in bulk \( ^4 \)He have been investigated putting it in a simulation box with \( N - 1 \) \( ^4 \)He atoms in such a way that the volume is \( \Omega = N/\rho \), with \( \rho \) the input density. In all the simulations \( N = 108 \) particles have been used and the time step and population bias have been analyzed in order to remove any systematic error. We have also verified that for \( N \gtrsim 100 \) the finite-system size introduces an error which
is smaller than the statistical noise, indicating that the influence of the replicas of the $^3$He impurity implied by the use of periodic boundary conditions is negligible. The interatomic interaction, which does not distinguish between the two isotopes, is the HFD-B(HE) Aziz potential $^{24}$ which has proved its high accuracy in a DMC calculation of the equation of state of superfluid $^4$He at zero temperature. $^{19,20}$ Concerning the trial wave function (5), the two-body factor proposed in Ref. $^{25}$ with the parameters optimized for pure $^4$He $^{19}$ has been considered.

We present the results of our calculations starting with a microscopic analysis of the local environment of the $^3$He impurity in the medium. This information is mainly contained in the crossed two-body radial distribution function $g^{(4,I)}(r)$. In Fig. 1, mixed (short-dashed line) and pure (solid line) estimations of $g^{(4,I)}(r)$ at densities 0.365, 0.401 and 0.424 $σ^{-3}$ ($σ = 2.556$ Å) are reported. In all the three densities the pure or exact results appear shifted to the right with respect to the mixed estimations pointing to a larger hole that is absolutely absent in the trial wave function. On the other hand, the height of the main peak in the pure $g^{(4,I)}(r)$ is slightly reduced at positive pressures and remains unchanged at the equilibrium density. In Fig. 2, the evolution of $g^{(4,I)}(r)$ with density is compared with the one shown by the pure $^4$He distribution function $g^{(4,4)}(r)$. Both functions show an increase of the localization when the density increases as well as a shift of the main peak to shorter interparticle distances. At a given density, the height of the main peak of $g^{(4,I)}(r)$ is smaller than the one of $g^{(4,4)}(r)$ and, what is more relevant, it appears localized to the right of the main peak of $g^{(4,4)}(r)$ pointing manifestly to the existence of an excluded-volume region due to the smaller mass of the $^3$He atom. The size of the excluded volume decreases when the density increases as one qualitatively can see comparing $g^{(4,4)}(r)$ and $g^{(4,I)}(r)$ at equilibrium and at the highest density plotted in Fig. 2.

Additional information on the local environment of the impurity is contained in the crossed static structure factor $S^{(4,I)}(k)$,

$$S^{(4,I)}(k) = \left\langle \Phi_0 \left| e^{i\mathbf{k} \cdot \mathbf{r}_I} \sum_{i=1}^{N-1} e^{-i\mathbf{k} \cdot \mathbf{r}_i} \right| \Phi_0 \right\rangle, \quad (12)$$
which corresponds to the Fourier transform of $g^{(4,I)}(r)$

$$S^{(4,I)}(k) = \rho \int dr \ e^{ik \cdot r} (g^{(4,I)}(r) - 1),$$  \hspace{1cm} (13)

$\rho$ being the density of pure $^4$He. From the above definition it is easy to check that the value of $S^{(4,I)}(k)$ at the origin is

$$S^{(4,I)}(0+) = -(1 + \alpha)$$  \hspace{1cm} (14)

with $\alpha = v / v_4$ the quotient between the molar volume of the impurity system ($v$) and that of pure $^4$He ($v_4$). In Fig. 3, $S^{(4,I)}(k)$ is plotted in comparison with $S^{(4,4)}(k) - 1$ at the $^4$He equilibrium density, $S^{(4,4)}(k)$ being the pure $^4$He static structure factor

$$S^{(4,4)}(k) = 1 + \rho \int dr \ e^{ik \cdot r} (g^{(4,4)}(r) - 1).$$  \hspace{1cm} (15)

The function $S^{(4,I)}(k)$ shown in the figure has been obtained Fourier transforming $g^{(4,I)}(r)$ for values $k > 1 \ \text{Å}^{-1}$ and by a direct calculation of Eq. (12) for $k \leq 1 \ \text{Å}^{-1}$. The main peak of $S^{(4,I)}(k)$ appears slightly depressed with respect to the one of $S^{(4,4)}(k) - 1$ reflecting the same feature observed in the comparison of the radial distribution functions. Nevertheless, the largest differences between the two static structure functions are at low $k$ values ($k \lesssim 1 \ \text{Å}^{-1}$).

In spite of the impossibility of calculating $S^{(4,\alpha)}(k)$ below a certain $k_{\text{min}}$, imposed by the use of a finite-size simulation box and periodic boundary conditions, if a linear extrapolation to $k = 0$ is carried out one obtains $S^{(4,4)}(0) - 1 \simeq -1$ and $S^{(4,I)}(0) \simeq -1.3$. If the latter is compared with Eq. (14), it results $\alpha = 0.3$ to be compared with the experimental value $\alpha^{\text{expt}} = 0.284$. The volume-excess parameter $\alpha$ decreases with pressure but this feature may be hardly observed in the limiting behaviour of $S^{(4,I)}(k)$ at different densities (Fig. 4).

One of the most relevant magnitudes in the study of the impurity system is the binding energy of the $^3$He atom in the medium or, otherwise, the chemical potential of the impurity $\mu_I$. In Table I, we report DMC results of the pure $^4$He chemical potential $\mu_4$ and $\mu_I$ at three densities which correspond to the pressures also contained in the table. The results for the pressure and $\mu_4$ reproduce the experimental data with high accuracy as pointed out.
in Refs. 19,20. Also, in the present case, one gets a nice agreement between the calculated
\( \mu_I \) and the experimental data,\[1\] the statistical uncertainties in the values of \( \mu_I \) being less
than 10%. A more exhaustive comparison between theoretical and experimental values for
\( \mu_I \) is displayed in Fig. 5. In the figure, two additional results are plotted: one at a pressure
higher than 20 atm, and another located at a density smaller than the equilibrium one (\( \rho_0 \))
which corresponds to a negative pressure of -6 atm. The solid line is a polynomial fit to the
DMC results and has to be compared with the available experimental data of Ref. [1], also
reported in the figure. As one can see, the agreement between theory and experiment is
excellent and a minimum in \( \mu_I(\rho) \) is not observed in this region. In fact, if a minimum exists
it is located at lower densities, even below the spinodal density of \( ^4\text{He} \) (\( \rho_s = 0.264 \sigma^{-3} \)).\[20\]

It is worth noticing that \( ^3\text{He} \) energetically prefers to remain in the surface of liquid \( ^4\text{He} \)
forming an Andreev state rather than penetrate in the bulk.\[21,22\] We have verified\[28\] that if
the \( ^3\text{He} \) impurity is replaced by a \( \text{H}_2 \) molecule there is a minimum of \( \mu_I \) at a density below
\( \rho_0 \) that nearly coincides with the local density of the preferred location of \( \text{H}_2 \) in \( ^4\text{He} \) clusters
obtained in a DMC calculation of Barnett and Whaley.\[29\]

In ACA the chemical potential of the impurity is given by\[8,30\]

\[
\mu_I^\text{ACA} = \mu_4 + \left( \frac{m_4}{m_I} - 1 \right) T_4 ,
\]

i. e., it can be calculated from the knowledge of properties of the pure liquid. This approx-
imation provides upper bounds (see Table I) which, using DMC values for \( \mu_4 \) and \( T_4 \), come
close to the DMC and experimental values.

The volume-excess parameter \( \alpha \) may be obtained from the knowledge of \( \mu_I(\rho) \), or equi-
valently \( \mu_I(P) \), through the thermodynamic relation

\[
\alpha = \rho \frac{\partial \mu_I}{\partial P} - 1 .
\]

The values for \( \alpha \) so obtained are reported in Table I in comparison with the experimental
data of Ref. [1]. The agreement between \( \alpha \) and \( \alpha^\text{expt} \) is very good at zero and intermediate
pressures and even at high pressure where the error bar is somewhat larger.
Microscopic quantities which are also significant in the present study are the kinetic energy of the $^3$He atom and the mean potential energy $^3$He-$^4$He ($V_I$). In Table II, results for the kinetic and potential energies for the two helium isotopes are reported at several densities. All of them correspond to pure estimations. In both systems, pure liquid $^4$He and liquid $^4$He with one $^3$He impurity, the potential energies may be obtained using the same method that has been used for the radial distribution functions because they are coordinate operators. The pure $^4$He kinetic energy simply results from the difference $E/N - V/N$ but that is not the case for $T_I$ in the impurity system due to the coexistence of the two isotopes. Therefore, the kinetic energy of the impurity has been calculated using the Hellmann-Feynman theorem as commented in Sect. II. The ACA estimation of the partial energies of the impurity is $T_I^{\text{ACA}} = m_4/m_I T_4$ and $V_I^{\text{ACA}} = V_4$, the values of $T_I^{\text{ACA}}$ being explicitly given in Table II to be compared with the exact results. In going from $T_4$ to $T_I$ one can see that the largest change is due to the difference in the mass of the two isotopes, the only effect contained in $T_I^{\text{ACA}}$, and the correction due to different correlations, i.e., $T_I^{\text{ACA}} - T_I$, is in all cases less than 10%. This small correction is also observed by comparing $V_4$ and $V_I$. In the range of densities here analyzed, it is observed that $V_I$ is always smaller than $V_4$ (in absolute value) whereas the difference $T_I^{\text{ACA}} - T_I$ is not monotonous: at $P \geq 0$, $T_I^{\text{ACA}} > T_I$ but $T_I^{\text{ACA}} < T_I$ at a density $0.328 \sigma^{-3}$ ($P = -6$ atm). This striking behaviour can be better understood looking at the differences between $g_{\text{ACA}}^{(4,I)}(r) = g^{(4,4)}(r)$ and $g^{(4,I)}(r)$ at each density. In the region of positive and zero pressures the main peak of $g^{(4,I)}(r)$ is ever shifted to the right with respect to the one of $g^{(4,4)}(r)$ and with a smaller localization (Fig. 2). The environment of the impurity may then be made equivalent to a pure $^4$He liquid at a reduced density. The reduced density $\rho_r$ of the equivalent system at positive pressure can be obtained by looking for the density of pure $^4$He at which $V_I$ and $g^{(4,I)}(r)$ do correspond. If the density $\rho_r$ is then used to estimate the kinetic energy of the impurity, $T_I(\rho_r) = m_4/m_I T_4(\rho_r)$, the results for $T_I$ are the same than the ones reported in Table II. This supplies an additional test to our pure computation of $T_I$ using the Hellmann-Feynman theorem. In the case of the equilibrium density ($\rho_0 = 0.365 \sigma^{-3}$) $\rho_r = 0.358 \sigma^{-3}$. At $\rho_r$, we have performed an explicit
calculation of $^4\text{He}$ and have verified that $g^{\text{(4,4)}}(r)$ is very much the same that $g^{\text{(4,I)}}(r)$ at $\rho_0$. On the other hand, at the lowest density reported in Table II ($\rho = 0.328 \sigma^{-3}$, $P = -6$ atm) the equivalent system does not exist because the shift of the main peak of $g^{\text{(4,4)}}(r)$ with respect to the one of $g^{\text{(4-4)}}(r)$ disappears and only a small delocalization remains.

There is only a previous *ab initio* calculation of $T_I$ at the $^4\text{He}$ equilibrium density using PIMC and extrapolating to zero temperature. Our present result for $T_I$, which is more accurate than our preliminary result of Ref. 31, is appreciably larger than the value reported in Ref. 12, $T_I = 17.1(1)$ K. As a kind of closure test of our results we have calculated the mass dependence of $T_I$ in order to estimate the chemical potential of the $^3\text{He}$ impurity through the relation

$$\mu_I = \mu_4 + \int_{m_4}^{m_I} dm \frac{T_I(m)}{m}.$$  \hspace{1cm} (18)

In Fig. 6, results for $T_I$ using different masses for the impurity are displayed in comparison with the ACA prediction (dashed line). For simplicity, the kinetic energies $T_I$ correspond in this case to mixed estimations, since at $\rho_0$ and for the aforementioned trial wave function the mixed and pure results coincide for both $m_I = m_4$ and $m_I = m_3$. The PIMC result for $m_I = m_3$ is also shown as an open circle. In the ACA case, if $T_I$ in Eq. (18) is replaced by $T_I^{\text{ACA}}$ one recovers the ACA expression for $\mu_I$ (14) and the corresponding result reported in Table I, $\mu_I^{\text{ACA}} = -2.58$ K. The solid line in Fig. 6 corresponds to a fit $T_I(m) = am + b/m$, and when integrated in Eq. (18) one obtains $\mu_I = -2.70(10)$ K which is consistent with both the experimental value and our direct estimation contained in Table I. As a supplementary result, it is predicted a linear departure from the ACA prediction with the impurity mass as is clearly manifested in Fig. 7, where the function $T_I^{\text{ACA}}(m) - T_I(m)$ is shown. Finally, it is worth mentioning that our results confirm and even enlarge the discrepancies between deep-inelastic neutron scattering determinations of the $^3\text{He}$ kinetic energy in liquid $^3\text{He}^{-^4\text{He}}$ mixtures (13) ($T_3 = 11 \pm 3$ K at $P = 0$ and $x = N_3/N = 0.10$) and all the theoretical predictions. One of the reasons that may explain this disturbing difference is the importance of the high-energy tails in the dynamic structure function which
largely influence the second energy-weighted sum rule from which the kinetic energy is extracted.

We close this section with the results obtained for the impurity effective mass $m^*_I$, which has been recently measured with great accuracy in $^3\text{He}-^4\text{He}$ mixtures and also microscopically analyzed using correlated basis function (CBF) theory. The $^3\text{He}$ effective mass plays a relevant role in the study of $^3\text{He}-^4\text{He}$ mixtures characterizing the $^3\text{He}$ excitations at low momenta. In a DMC calculation, the impurity effective mass can be obtained from the diffusion coefficient of the impurity in imaginary time:

$$\frac{m_I}{m^*_I} = \lim_{\tau \to \infty} \frac{|\mathbf{r}_I(\tau) - \mathbf{r}_I(0)|^2}{6D_I \tau}, \quad (19)$$

with $D_I = \hbar^2 / (2m_I)$ the free-diffusion constant of the impurity. In Fig. 8, extrapolated estimations of $m_I/m^*_I$ are reported at densities 0.365, 0.401, and 0.424 $\sigma^{-3}$. The impurity effective mass is extracted from a linear fit to the flat asymptotic regime of that function (19) which, as the figure shows, is acquired at relatively short diffusion times. The results so obtained are reported in Table III in comparison with the experimental determinations from Refs. 14,15 and the recent CBF calculation of Krotscheck et al. Obviously, the experimental values are not direct measures but extrapolations to zero $^3\text{He}$ concentration ($x$) of determinations in $^3\text{He}-^4\text{He}$ mixtures. As pointed out by Krotscheck et al. a linear extrapolation, primarily used in the experimental works, is not satisfactory because the Fermi-liquid contributions are the most relevant in the $^3\text{He}$-concentration dependence of $m^*_I$ and these terms introduce fractional powers of $x$ in the analytical model for $m^*_I(x)$. The experimental values reported in Table III have been obtained using this more accurate extrapolation. Within the statistical errors of the DMC results, an overall agreement between our calculation and experiments is attained, with somehow a significant difference at the highest density due in part to the use of the extrapolated estimation (19). On the other hand, the CBF results of Ref. 14 come close to the DMC and experimental results but seem to be slightly smaller at the densities here reported. Another CBF calculation, due to Fabrocini et al. reported several years ago a result of $m^*_I = 2.2$ at the equilibrium density in better
agreement with the present DMC results.

IV. SUMMARY AND CONCLUSIONS

We have analyzed in this paper the most important magnitudes which characterize the static properties of a single $^3$He atom embedded in bulk superfluid $^4$He. The difficulties of an efficient calculation of the binding energy of the impurity in the medium, one of the main objectives of the present work, had prevented in the past the application of \textit{ab initio} Monte Carlo methods to this problem. In order to overcome these difficulties, it has been proved that the use of reweighting techniques can be readily extended to diffusion Monte Carlo algorithms. This generalized reweighting method has provided reliable results for $\mu_I$ which are in excellent agreement with experimental data.

The local environment of the $^3$He atom has been explored through the calculation of the crossed radial distribution and static structure functions for a wide range of densities. The use of pure estimators for these quantities removes the uncontrolled bias, remanent in the approximate extrapolation methods, and shows clear evidence of an excluded volume region surrounding the $^3$He impurity. The low $k$ behaviour of $S^{(4,1)}(k)$ also points to the expected value related to the volume-excess parameter $\alpha$, but a precise value for $\alpha$ cannot be estimated due to the absence of data for $k \leq k_{\text{min}} = 2\pi/L$, with $L$ the side of the simulation box. Nevertheless, an independent and more precise estimation of $\alpha$, through the pressure dependence of the chemical potential of the impurity, produces results which compare favorably with experimental data.

Special attention has been devoted to an accurate estimation of the partial energies, potential and kinetic, of the impurity. The usual forward walking methodology does not apply for derivative operators, and for this reason, we have used the Hellmann-Feynman theorem combined with the generalized reweighting method to calculate the $^3$He kinetic energy. The results for $T_I$ obtained with this method show smaller differences with the ACA values than a previous PIMC estimate, with a difference $T_I^{ACA} - T_I$ which increases
linearly with the mass of the isotopic impurity. Our results confirm the gap between all the theoretical results for $T_I$ and the much smaller $^3$He kinetic energies derived from the neutron scattering data of Refs. 34.

A final concern of the present work is the calculation of the $^3$He effective mass through its diffusion coefficient in imaginary time. The results obtained show a good agreement with recent experimental data that slightly worsens at high pressure due probably to uncertainties in the MC extrapolation method used in the estimation of $m^*_I$. A natural extension to the present work would be the calculation of the excitation energy of the $^3$He impurity in liquid $^4$He, which in the limit $q \to 0$ is given by $\frac{\hbar^2 q^2}{2m^*_I}$, and therefore will provide another method to estimate the impurity effective mass. In such a calculation, one can use DMC combined with the fixed-node and released-node methods, that we have already employed in the study of the phonon-roton spectrum in superfluid $^4$He. Work in this direction is in progress.

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TABLE I. Chemical potential of pure liquid \(^4\)He (\(\mu_4\)), chemical potential of the \(^3\)He impurity (\(\mu_I\)), and excess-volume parameter \(\alpha\) at several densities. The experimental data is from Ref. 1.

| \(\rho (\sigma^{-3})\) | \(P(\text{atm})\) | \(\mu_4 (\text{K})\) | \(\mu_I^\text{ACA} (\text{K})\) | \(\mu_I (\text{K})\) | \(\mu_I^\text{expt} (\text{K})\) | \(\alpha\) | \(\alpha^\text{expt}\) |
|------------------------|-----------------|-------------------|-------------------|-------------------|-------------------|--------|---------|
| 0.365                  | 0.              | -7.27(1)          | -2.58             | -2.79(25)         | -2.785            | 0.284(10) | 0.284  |
| 0.401                  | 10.67           | -3.89(1)          | 1.59              | 1.38(30)          | 1.42              | 0.200(10) | 0.199  |
| 0.424                  | 20.42           | -0.97(2)          | 5.10              | 4.73(35)          | 4.83              | 0.176(20) | 0.165  |

TABLE II. Kinetic and potential energies of the pure liquid \(^4\)He and of the \(^3\)He impurity immersed in bulk \(^4\)He. All the energies are in K.

| \(\rho (\sigma^{-3})\) | \(T_4\)     | \(V_4\)       | \(T_I^\text{ACA}\) | \(T_I\)     | \(V_I\)     |
|------------------------|------------|--------------|-------------------|------------|------------|
| 0.328                  | 11.99(8)   | -19.14(6)    | 15.91(8)          | 17.0(6)    | -18.2(5)   |
| 0.365                  | 14.32(5)   | -21.59(5)    | 19.00(7)          | 18.4(5)    | -21.1(5)   |
| 0.401                  | 16.73(9)   | -23.88(9)    | 22.20(12)         | 20.5(5)    | -22.6(5)   |
| 0.424                  | 18.57(8)   | -25.45(8)    | 24.64(11)         | 23.4(8)    | -24.7(5)   |

TABLE III. \(^3\)He impurity effective mass at several densities. The CBF results are from Ref. 10.

| \(\rho (\sigma^{-3})\) | \(m_I^*\)   | \(m_I^{*\text{expt}}\) (Ref. 14) | \(m_I^{*\text{expt}}\) (Ref. 15) | \(m_I^{*\text{CBF}}\) |
|------------------------|-------------|----------------------------------|----------------------------------|------------------------|
| 0.365                  | 2.20(5)     | 2.18                             | 2.15                             | 2.09                   |
| 0.401                  | 2.36(8)     | 2.44                             | 2.39                             | 2.34                   |
| 0.424                  | 2.72(10)    | 2.64                             | 2.62                             | 2.55                   |
FIG. 1. Mixed (dashed line) and pure (solid line) estimations of $g^{(4,1)}(r)$ at densities $0.365 \, \sigma^{-3}$, $0.401 \, \sigma^{-3}$, and $0.424 \, \sigma^{-3}$, from bottom to top. A vertical shift has been introduced at $0.401 \, \sigma^{-3}$ and $0.424 \, \sigma^{-3}$ to better visualize their differences.
FIG. 2. Pure liquid $^4$He (dashed line) and impurity-medium (solid line) two-body radial distribution functions at densities 0.365 $\sigma^{-3}$, 0.401 $\sigma^{-3}$, and 0.424 $\sigma^{-3}$, from bottom to top. A vertical shift has been introduced at 0.401 $\sigma^{-3}$ and 0.424 $\sigma^{-3}$ to better visualize their differences.
FIG. 3. Pure liquid $^4$He (dashed line) and impurity-medium (solid line) static structure factor at the $^4$He equilibrium density $\rho_0 = 0.365 \, \text{Å}^{-3}$. We have plotted $S^{(4,4)}(k) - 1$ for a better comparison.
FIG. 4. Impurity-medium static structure factor at densities 0.365 $\sigma^{-3}$ (solid line), 0.401 $\sigma^{-3}$ (dashed line), and 0.424 (dotted line) $\sigma^{-3}$. 
FIG. 5. Chemical potential of the $^3$He impurity as a function of the density (full circles). The solid line is a polynomial fit to the DMC results. The open circles are experimental data from Ref. 1.
FIG. 6. Kinetic energy of the impurity as a function of its mass (full circles and solid line). The dashed line corresponds to the ACA prediction. The open circle is the PIMC result from Ref. 12.
FIG. 7. Difference between the ACA prediction and the real value for the kinetic energy of the impurity as a function of its mass (full circles and solid line). The open circle is the PIMC result from Ref. 12.
FIG. 8. The inverse of the impurity effective mass from the long-time behaviour of its diffusion coefficient. The solid, dashed and dotted lines correspond to densities $0.365 \sigma^{-3}$, $0.401 \sigma^{-3}$, and $0.424 \sigma^{-3}$, respectively.