On the order of BEC transition in weakly interacting gases predicted by mean-field theory

L Olivares-Quiroz and V Romero-Rochin

Instituto de Física, Universidad Nacional Autónoma de México, Apartado Postal 20-364, 01000 México, DF, Mexico
E-mail: olivares@fisica.unam.mx and romero@fisica.unam.mx

Received 3 March 2010, in final form 14 July 2010
Published 4 October 2010
Online at stacks.iop.org/JPhysB/43/205302

Abstract
Predictions from Hartree–Fock (HF), Popov (P), Yukalov–Yukalova (YY) and t-matrix approximations regarding the thermodynamics from the normal to the Bose–Einstein condensation (BEC) phase in weakly interacting Bose gases are considered. By analysing the dependence of the chemical potential \( \mu \) on the temperature \( T \) and particle density \( \rho \), we show that none of them predicts a second-order phase transition as required by symmetry-breaking general considerations. In this work, we find that the isothermal compressibility \( \kappa_T \) predicted by these theories does not diverge at criticality as expected in a true second-order phase transition. Moreover, the isotherms \( \mu = \mu(\rho, T) \) typically exhibit non-single-valued behaviour in the vicinity of the BEC transition, a feature forbidden by general thermodynamic principles. This behaviour can be avoided if a first-order phase transition is appealed. The facts described above show that although these mean field approximations give correct results near zero temperature, they are endowed with thermodynamic anomalies in the vicinity of the BEC transition. We address the implications of these results in the interpretation of current experiments with ultracold trapped alkali gases.

1. Introduction

In the recent past it has been widely discussed that Bose–Einstein condensation (BEC) in interacting Bose systems, i.e. the passage from the normal to the condensed phase where all particles occupy a single-particle state, shows a spontaneous \( U(1) \) gauge symmetry breaking with the condensate fraction \( N_0/N \) playing the role of the order parameter \([1–4]\). The overwhelming task faced when trying to solve the full interacting quantum gas has motivated the search of physical approximations for the dilute and weakly interacting gas \([5, 6]\). This approximation, formally known as the theory of weakly interacting Bose gases, has become the standard tool for analysing the BEC transitions for gases confined either in a box of volume \( V \) or trapped through external potentials in optical setups.

In spite of the great success achieved, the theory of weakly interacting Bose gas still awaits a formal and complete analytical solution. In order to extract some partial results, different approximations have been developed in an upper ‘layer’ build up on top of the theory of weakly interacting gases \([2, 10–15]\). In this work, we focus our attention to four of the approximations most widely used to describe the interacting Bose gas in the dilute approximation. These are Hartree–Fock (HF), Popov (P), Yukalov–Yukalova (YY) and t-matrix approximations. As we show in this work none of these approximations exhibit BEC as a second-order phase transition. The fact that HF neither exhibits a second-order
phase transition nor displays the correct energy spectrum has been known for a while [10, 11]. Here, we show that Popov (P), t-matrix approach [2, 13] and the recently introduced Yukalov–Yukalova (YY) [12] approximations also fail to predict a second-order phase transition. In addition, we show that all of these theories display an unstable region implying that the BEC transition is at best, a first-order phase transition.

To achieve this goal we proceed as follows. We start by calculating the equation of state $\mu = \mu(T, \rho)$, that is, the dependence of the chemical potential $\mu$ on the temperature $T$ and particle density $\rho = N/V$, with $N$ being the number of particles and $V$ the volume of the system in each of the approximations considered. Equilibrium thermodynamics ensures that the isothermal compressibility $\kappa_T$ can be obtained from the equation of state $\mu = \mu(T, \rho)$ through the relationship $\kappa_T^{-1} = \rho^2(\partial \mu / \partial \rho)_T$ from which information regarding the order of the transition predicted by these theories can be inferred. Since the isotherm compressibility $\kappa_T$ is indeed a thermodynamic manifestation of the fluctuations in the particle density, the isotherms $\mu = \mu(T, \rho)$ contain fundamental information on the nature of the phase transition involved [16].

From a more general point of view, the equation of state $\mu = \mu(\rho, T)$ of an interacting quantum gas contained in a rigid vessel of volume $V$ has a fundamental relevance of itself. Such a quantity enables the prediction of the density profile $n(\vec{r})$ of the interacting gas confined by an inhomogeneous trap potential $V_{\text{ext}}(\vec{r})$ which is the main measurable property in the current experiments with ultracold alkali gases [7–9, 17]. The connection between the thermodynamics of the gas in a box of volume $V$ and the thermodynamics of the gas confined within an external potential $V_{\text{ext}}(\vec{r})$ can be achieved through the local density approximation (LDA). In such a procedure one obtains $\rho = \rho(\mu, T)$ by the inversion of the equation of state, and then replacing $\mu$ by $\mu - V_{\text{ext}}(\vec{r})$. This gives the density profile $n(\vec{r}) = \rho(\mu - V_{\text{ext}}(\vec{r}), T)$. This procedure has been shown to be exact in the appropriate thermodynamic limit of a gas confined by a trap $V_{\text{ext}}(\vec{r})$ [18–20]. As we shall carefully discuss, if the trapped gas is in the BEC phase the density profile bears the information of the isotherm $\rho(\mu, T)$ above, below and at the transition. Thus, knowledge of $\mu = \mu(\rho, T)$ also yields the equation of state for the trapped gases. Inaccurate calculations of the equation of state in the homogeneous case will be inherited by the inhomogeneous ones. One of the purposes of this paper is to suggest high-resolution measurements of the density profiles in the current experiments. These would not only settle the issue of the validity or the mean-field calculations, but would also pave the way to improve theoretical descriptions and thus better understanding of the BEC phase.

The paper is organized as follows. In section 2 a general discussion of the thermodynamics of phase transitions and the weakly interacting gas is presented. Section 3 is devoted to a brief presentation of the approximations considered in this work and the results for the equation of state $\mu = \mu(T, \rho)$ corresponding to each of them. Section 4 discusses the implications of these results within the general theory of the weakly interacting gas and in the interpretation of current experiments in ultracold alkali gases.

2. General thermodynamic considerations and the weakly interacting Bose gas

For our analysis, we consider the equation of state $\mu = \mu(T, \rho)$, that is, the dependence of the chemical potential $\mu$ on the temperature $T$ and particle density $\rho = N/V$, with $N$ being the number of particles and $V$ the volume of the sample. The extensivity property of the Helmholtz free energy $F = F(N, V, T)$ allows us to either write $F = F(\rho, T)$ or $F = N \tilde{f}(v, T)$ with $v = V/N = \rho^{-1}$ such that $\tilde{f} = \rho \tilde{f}$, and both forms carry the same physical information. One can either calculate the chemical potential as

$$\mu(T, \rho) = \left( \frac{\partial F}{\partial \rho} \right)_T,$$  \hspace{1cm} (1)

or the pressure

$$p(T, v) = -\left( \frac{\partial \tilde{f}}{\partial v} \right)_T.$$  \hspace{1cm} (2)

Therefore, $\mu = \mu(T, \rho)$ and $p = p(v, T)$ also carry the same information, and the phase diagram may be inferred from any of these forms. The laws of thermodynamics require that both of these expressions are single valued, that is, for any given values of $\rho$ and $T$, or $v$ and $T$, there must only exist a single value of $\mu$ or $p$, respectively. In addition, these formulae carry out crucial information on the order of the phase transition involved. The stability of a particular thermodynamic state is ensured if the isothermal compressibility $\kappa_T$ is positive, where $\kappa_T$ may be either calculated as

$$\kappa_T^{-1} = \rho^2 \left( \frac{\partial \mu}{\partial \rho} \right)_T,$$  \hspace{1cm} (3)

or

$$\kappa_T^{-1} = -v \left( \frac{\partial p}{\partial v} \right)_T.$$  \hspace{1cm} (4)

These equations imply that $(\partial \mu / \partial \rho)_T > 0$ and $(\partial p / \partial v)_T < 0$ for any thermodynamic equilibrium state. An exception is a critical point where the latter derivatives become zero.

The emergence of a negative region for the isothermal compressibility signals the onset of a first-order phase transition where two phases with different densities, entropies and energies coexist at the same temperature, pressure and chemical potential [16]. Let us consider here that such states have densities $\rho_1$ and $\rho_2$. The equal-areas Maxwell construction enables us to calculate the values of the thermodynamic properties of interest for both coexisting states. In order to see this, it is enough to consider the Gibbs–Duhem relationship

$$-SdT + Vdp - Nd\mu = 0,$$  \hspace{1cm} (5)

where $S$ is the entropy. By assuming that the system is analysed along an isotherm, we can set $dT \equiv 0$ and look for the coexisting values of the density or the volume. Then, we
can impose equal chemical potential \( \mu = \mu_{\text{coex}} \) at both phases yielding
\[
\int_1^2 v \, dp = \int_1^2 d\mu = 0
\]
\[
= p_{\text{coex}}(v_2 - v_1) - \int_{v_1}^{v_2} p \, dv = 0,
\]
where \( p_{\text{coex}}, v_1 \) and \( v_2 \) are the volume values at coexistence, with \( v_1 = \rho_1^{-1} \) and \( v_2 = \rho_2^{-1} \). The value of the chemical potential is \( \mu_{\text{coex}} = \mu(T, \rho_1) = \mu(T, \rho_2) \). Equation (6) is the usual Maxwell equal-area construction in the \( p-v \) diagram. Alternatively, if one imposes equal pressures at coexistence, namely \( p = p_{\text{coex}} \), one finds from equation (5)
\[
\int_1^2 \rho \, d\mu = \int_1^2 d\rho = 0
\]
\[
= \mu_{\text{coex}}(\rho_2 - \rho_1) - \int_{\rho_1}^{\rho_2} \mu \, d\rho = 0,
\]
with \( \mu_{\text{coex}}, \rho_1 \) and \( \rho_2 \) being the corresponding values at coexistence. Equation (7) is also a Maxwell equal-areas construction but in the \( \mu-\rho \) diagram.

The onset of a second-order phase transition is featured by the divergence of the isothermal compressibility \( K_T \) and the heat capacity \( C_v \), characterized by universal exponents [16, 21, 23]. In this work, we focus our attention to the isothermal compressibility \( K_T \) whose divergence can be expressed by the vanishing derivative \( (\partial \mu/\partial \rho)_T = 0 \) or \( (\partial \rho/\partial v)_T = 0 \), at the transition. This condition demands that the isothermal \( \mu = \mu(T, \rho) \) must become ‘flat’ at the critical density \( \rho = \rho_c \). We shall explicitly show that none of the analysed approximations for the weakly interacting Bose gas show this strong requirement. It is worth recalling that an ideal Bose gas confined in a box of volume \( V \) does show a diverging compressibility at the critical temperature \( T_c \) [21, 23].

The theory of the weakly interacting Bose gas assumes that the atoms have no structure since they typically are in the same hyperfine state [9]. If a pairwise interatomic potential \( U = U(|\vec{r}_i - \vec{r}_j|) \) is assumed, then the Hamiltonian in the second quantization can be written as
\[
\hat{H} = \sum_k \epsilon_k \hat{a}_k^\dagger \hat{a}_k + \frac{1}{2V} \sum_{k,k',q} \hat{U}(q) \hat{a}_{k+q}^\dagger \hat{a}_{k'-q}^\dagger \hat{a}_{k'} \hat{a}_k,
\]
where \( k \) is the three-dimensional wavevector quantized in a box of volume \( V \), \( \hat{a}_k^\dagger \) and \( \hat{a}_k \) are the creation and annihilation operators of particles with momentum \( \hbar \vec{k}, \epsilon_k^0 = \hbar^2 k^2/(2m) \) is the one-particle kinetic energy and \( \hat{U}(q) \) is the Fourier transform of the interparticle potential \( U(r) \). The main assumption for the description of an interacting gas at low temperatures is the contact potential approximation, namely \( \hat{U}(q) \approx U_0 \) for all \( q \), with \( U_0 = 4\pi\hbar^2 a/m \) representing the strength of the interaction and \( a \) being the s-wave scattering length which, in this work, we shall assume as positive. In the \( t \)-matrix approximation also considered here, corrections to this simple potential are also included [2]. A variational ansatz completes the framework. A Bose gas is considered weakly interacting if \( \rho a^3 \ll 1 \). All the approximations considered in this work will satisfy these general requirements. In the following, we succinctly present the differences between each of them that yield to different equations of state \( \mu = \mu(\rho, T) \). We refer the reader to the original sources cited along this work for further details.

For our calculations, we use units \( \hbar = m = a = 1 \). We shall analyse three isotherms in the HF, P and YY approximations, \( k_B T = 0.01, 0.1 \) and 1.1 in dimensionless units, while only \( k_B T = 0.1 \) in TM approach. Before proceeding to the calculations it is worth stating some words on the physical regime associated with the isotherms chosen. Typical experiments on \(^{23}\)Na ultracold gases [17] show that BEC transition temperature is nearly \( T_c \approx 100 \) nK with a scattering length \( a \approx 55a_0 \), where \( a_0 \) is the Bohr radius. This yields in dimensionless units \( k_B T \approx 10^{-4} \), thus indicating that the chosen temperatures are a bit out of range, but this is not necessarily so. As we shall discuss below, while in HF and P the behaviour is qualitatively the same for all temperatures, YY does show three different qualitative behaviours corresponding approximately to those temperatures. Moreover, recent advances in the experimental achievement of ultracold atomic gases have shown that the scattering length \( a \) can be tuned by external magnetic fields to larger values near a Feshbach resonance [24–27]. Hence, if an increase of tenfold in the scattering length is achieved, then a temperature of \( T \approx 100 \) nK would give \( k_B T \approx 0.01 \) in dimensionless units which corresponds perfectly to the first isotherm considered in this work. The tunability of the scattering length \( a \) together with the fact that each time is possible to achieve condensates with a larger number of particles suggests that in the near future BEC in ultracold gases will be achieved for higher temperatures. It is important to underline that the criterion for a weakly interacting gas is fulfilled here. The gas parameters associated with the isotherms are \( \rho a^3 = 0.000 17, 0.0052 \) and 0.19, respectively. Admittedly, the last value is at the border of where one should consider the gas as weakly interacting.

3. BEC transition within four mean-field theories

In this section, we analyse four different approximations that describe the weakly interacting Bose gas. These schemes have provided accurate descriptions at and near zero temperature. Given this success their use has been extended to finite temperatures near criticality. As suggested before, BEC can be attained either at fixed density lowering the temperature or at fixed temperature increasing the density. In this work, we shall make use of the latter approach and thus BEC transition will be accomplished in \( \mu - \rho \) space at the fixed temperature \( T \) by varying the total density. In this case, the transition occurs at a critical density \( \rho_c \), the gas being normal for \( \rho < \rho_c \), while BEC sets in for \( \rho \geq \rho_c \).

3.1. Hartree–Fock

The Hartree–Fock approximation is a self-consistent approach in which the state of the \( N \)-particle system is expressed in terms of effective one-particle states, yielding a gas of non-interacting excitations whose energy spectrum depends self-consistently on its density and the actual interparticle
interaction [10, 11]. Then, one can construct the grand potential as \( \Omega = \langle H \rangle - TS + \mu(N) \), in terms of variational occupation numbers \( f_k \), with

\[
\langle H \rangle \approx \frac{N_0}{2V} U_0 + \sum_{k \neq 0} \left( e_k^0 + 2\rho U_0 \right) f_k - \frac{U_0}{V} \sum_{k \neq 0} f_k f_{-k} \tag{9}
\]

and

\[
\langle N \rangle \approx N_0 + \sum_{k \neq 0} f_k, \tag{10}
\]

\( N_0 \) being the number of particles in the condensate with \( k = 0 \). Furthermore, one assumes that the entropy is given as that of an ideal Bose gas, but in terms of the effective one-particle occupation states \( f_k \), this is \( S = k \sum_k [(1 + f_k) \ln(1 + f_k) - f_k \ln f_k] \). At zero temperature, HF is considered an adequate approximation being equivalent to the Gross–Pitaevskii approach. However, at low but finite temperatures it predicts an excitation spectrum with a gap \( \epsilon_k \), as seen from equation (9) contrary to the expected gapless spectrum [28], that the P and YY approximations do take into account. However, in the normal gas region, that is, for densities below and at the transition, HF is an acceptable approximation.

By looking for the values of \( f_k \) that minimize \( \Omega \) at a constant temperature, one is led to the thermodynamic equations in the HF approximation. One finds that for densities \( \rho < \rho_c \), the gas is in the normal phase with [10, 11]

\[
\rho = \frac{1}{\lambda_T^3} g_{3/2} \left[ \beta (\mu - 2 \rho U_0) \right], \tag{11}
\]

On the other hand, for \( \rho \geq \rho_c \) BEC sets in, and it is found that

\[
\rho = \rho_0 + \frac{1}{\lambda_T^3} g_{3/2} (\beta \rho_0 U_0) \quad \text{with} \quad \mu = (2 \rho - \rho_0) U_0, \tag{12}
\]

where \( \rho \) and \( \rho_0 \) are the total and condensate densities. In the above equations \( \lambda_T = h/\sqrt{2\pi mk_BT} \), and we used the Bose integral

\[
g_n(\alpha) = \frac{1}{\Gamma(n)} \int_0^\infty x^{n-1} e^{-x} e^{-\alpha x} dx, \tag{13}
\]

with \( \beta = 1/k_BT \).

The onset of BEC is assumed to occur at \( \rho_0 = 0 \) in equation (12), namely at \( \mu_c = 2\rho_0 U_0 \). This defines a relation between \( T \) and \( \rho_c \), which is the same as in the ideal gas,

\[
\rho_c = \frac{1}{\lambda_T^3} g_{3/2}(0), \tag{14}
\]

Equations (11) and (12) yield \( \mu = \mu(\rho, T) \) as a continuous function of \( \rho \) at BEC. As we now show, this continuity does not guarantee the existence of a second-order phase transition.

Figure 1 shows solutions to equations (11) and (12) for the three isotherms, \( k_B T = 0.01, 0.1 \) and 1.1. The behaviour of \( \mu \) versus \( \rho \) for all temperatures is essentially the same. In the normal gas region, \( \rho < \rho_c \), the derivative \( (\partial \mu/\partial \rho)T \rightarrow 2U_0 \) as \( \rho \rightarrow \rho_c \). This straightforwardly predicts that the isothermal compressibility does not diverge at BEC. The behaviour of \( \mu \) versus \( \rho \) corresponding to the BEC region, i.e. the solution to equation (12), shows an anomalous thermodynamic behaviour: this solution, in principle only valid for densities \( \rho \geq \rho_c \), intrudes into the normal region yielding a multiple valued chemical potential as a function of density. Thermodynamics forbids this multiple valuedness. However, as done originally in [14] and [15], such a behaviour can be avoided by invoking a first-order phase transition, joining two phases with different densities by means of an equal-areas Maxwell construction, as shown in figure 1 for the isotherm \( k_B T = 1.1 \). As discussed above, equal-areas Maxwell construction ensures that the two phases with different densities have the same temperature, pressure and chemical potential, namely that the phases coexist. This is the signature of a first-order phase transition, and one finds that the conditions for a second-order phase transition, as discussed before, are never met. As described previously, the fact of having the co-existence of two states with different densities implies two different entropies, namely a latent heat, and two values of the energy. In the same fashion, a multiple-valued chemical potential also gives rise to an unacceptable multiple-valued pressure of the system.

### 3.2. Popov

The Popov approximation is considered correct at zero temperatures, being essentially the same as the Bogoliubov approximation [28]. This approximation, as shown below, yields a gapless excitation spectrum [2, 3, 11], linear in the excitation momentum as \( k \rightarrow 0 \) and gives rise to the phenomenon of superfluidity, just as in \( ^4\text{He} \) [6]. Bragg spectroscopy in an ultracold \(^{87}\text{Rb} \) gas [29] has shown that the excitation spectrum is clearly given by the Bogoliubov expression, thus providing validity to this approximation at very low temperatures and, incidentally, confirming the superfluid nature of these gases at the BEC state.
The number-conserving Hamiltonian of the system at this level of approximation is given by [11]

$$\hat{H} \approx \frac{N E_0^2}{2V} + \sum_k \left( \epsilon_k^0 + \frac{2N U_0}{V} \right) a_k^\dagger a_k - \frac{U_0}{V} \sum_{k,q \neq 0} f_k f_q$$

$$+ \frac{N_0 U_0}{2V} \sum_{k \neq 0} (a_k^\dagger a_{-k} + a_k a_{-k}). \quad (15)$$

It looks very similar to the Hartree–Fock version except for the last term which represents the annihilation of two particles into the condensate and the creation of two particles from the condensate. For a finite-temperature calculation of the thermodynamics once again one deals with the problem in the grand canonical ensemble. After performing the usual Bogoliubov transformation [11], the Popov procedure follows a similar line as HF and the system emerges as a kind of ideal gas with elementary excitations whose excitation spectrum is

$$\epsilon_k = \sqrt{\frac{h^2 k^2}{2m} + 2 \rho U_0 - \mu} - \rho^2 U_0^2. \quad (16)$$

At a fixed temperature $T$ and for densities below the critical one $\rho_c$, the equation in the P approximation for the density $\rho$ as a function of temperature $T$ and chemical potential $\mu$ turns out to be the same as in HF, equation (11), but for densities $\rho \geq \rho_c$, it is found that the density of the fluid is

$$\rho = \rho_0 + \frac{2}{\pi^{1/2} N_0 \sqrt{T}} \int_0^\infty \frac{x^{1/2} dx}{\sqrt{x^2 + 2 \beta \rho_0 U_0 x}} \left[ \coth \left( \frac{m c^2 x}{2 k_B T} \right) - 1 \right], \quad (17)$$

while the chemical potential is again given by $\mu = (2 \rho - \rho_0) U_0$ as in HF. The transition occurs when the condensate density vanishes, $\rho_0 = 0$, and the critical density $\rho_c$ is given by condition (14). The solution $\mu(\rho, T)$ for $\rho \geq \rho_c$ becomes continuous at $\rho_c$ with the solution to equation (11). Figure 2 shows three isotherms $\mu$ versus $\rho$ at $k_B T = 0.01, 0.1$ and $1.1$, from the numerical solution of Popov equations (11) and (17). $\mu_c$ and $\rho_c$ are the values at BEC. The gaseousness parameters at criticality are $\rho_c a^3 = 0.00017, 0.0052$ and $0.19$, respectively.

Once more, nevertheless, for a given temperature $T$ and in the normal region $\rho < \rho_c$, the relevant equations are those of HF, equation (11). For densities above criticality, $\rho \geq \rho_c$, a new set of equations are given for the value of the density $\rho$ and the anomalous average $\sigma_1$, namely

$$\rho = \rho_0 + \frac{1}{3 \pi^2 \hbar} \left( \frac{m c}{\hbar} \right)^3 \int_0^\infty \left[ \coth \left( \frac{m c^2 x}{2 k_B T} \right) - 1 \right] dx, \quad (19)$$

$$\sigma_1 = \left( \frac{m c}{\hbar} \right)^3 \left[ \left( \frac{\rho_0 U_0}{\pi^2 m c^2} \right)^{1/2} - \frac{1}{2 \pi^{1/2}} \right] \times \int_0^\infty \frac{\sqrt{1 + x^2} - 1}{\sqrt{1 + x^2}} \left[ \coth \left( \frac{m c^2 x}{2 k_B T} \right) - 1 \right] dx, \quad (20)$$

where the speed of the elementary excitations $c$ is given in terms of the condensate density $\rho_0$ and $\sigma_1$ by $m c^2 = (\rho_0 + \sigma_1) U_0$. In this approximation, the chemical potential is given by [12]

$$\rho \mu = \rho_0 (2 \rho - \rho_0 + \sigma_1) U_0 + (\rho - \rho_0) (2 \rho - \rho_0 - \sigma_1) U_0. \quad (21)$$

3.3. Yukalov–Yukalova

This recent scheme is based on the inclusion of anomalous averages $\sigma_1 = \langle \hat{\psi}_1(\vec{r}) \hat{\psi}_1(\vec{r}) \rangle$ into the Hamiltonian. According to its authors the anomalous averages should not be neglected since their contribution is of the same order as that of the condensate density [12]. The operator $\hat{\psi}_1(\vec{r})$ is given in terms of the creation operators $a_k$ as

$$\hat{\psi}_1(\vec{r}) = \sum_{k \neq 0} \phi_k(\vec{r}) a_k. \quad (18)$$

It is discussed in [12] that these anomalous averages are not properly considered in the Popov approximation and therefore their inclusion provides a better description for the BEC transition. In order to obtain the thermodynamics of the system, the YY approximation considers explicitly the symmetry breaking of the state by the appearance of the condensate and implements the so-called representative ensemble calculation where one Legendre multiplier is associated with the condensate particles $\rho_0$ and a different one to the non-condensate fraction $\rho - \rho_0$. This takes into account the correlations of uncondensed particles and allows for a clear distinction of the superfluid and condensate fractions at finite temperatures. At zero temperature, this approximation also agrees with that of Bogoliubov [28] as expected. The details of the calculation can be consulted in the original works of Yukalov and Yukalova [12].

Figure 2. Isotherms $\mu$ versus $\rho$ at $k_B T = 0.01, 0.1$ and $1.1$, from the numerical solution of Popov equations (11) and (17). $\mu_c$ and $\rho_c$ are the values at BEC. The gaseousness parameters at criticality are $\rho_c a^3 = 0.00017, 0.0052$ and $0.19$, respectively.
In this case BEC sets in when both $\sigma_1$ and $\rho_0$ vanish. This yields $\mu_\epsilon = 2\rho_c U_0$ at the transition with $\rho_c$ determined by condition (14). As in HF and P, the solution $\mu(\rho, T)$ to (19)–(21) is continuous with the solution of (11) at BEC. This approximation is much richer than the two previous ones. The behaviour of the isotherms $\mu$ versus $\rho$ shows three different regimes depending on the temperature, see figure 3. At very low temperatures, i.e. $k_B T = 0.01$, there are no multiple-valued solutions, neither an unstable region. Thus, there is no need to appeal to a first-order phase transition. However, despite the fact that the transition appears continuous, it cannot be considered as a bona-fide second-order one, since neither regime shows a divergent compressibility at BEC; rather, the compressibility becomes nearly zero, that is, the gas becomes incompressible at BEC. This property is completely unexpected and it is contrary to the standard results in weakly interacting systems.

while for $\rho \gtrsim \rho_c$

$$\rho = \rho_0 + \frac{1}{\pi^{1/2} \beta k_B T} \int_{0}^{\infty} \left[ \frac{1 + \beta \Delta}{E_x} \coth \frac{E_x}{2} - 1 \right] x^{1/2} \mathrm{d}x. \quad (23)$$

In the above equations, $\Delta$ is determined by an additional quantity

$$\alpha = \frac{1}{\pi^{1/2} \beta k_B T} \int_{0}^{\infty} \left[ \frac{1}{E_x} \coth \frac{E_x}{2} - \frac{1}{x} \right] x^{1/2} \mathrm{d}x, \quad (24)$$

where for $\rho < \rho_c, E_x = x - \beta \Delta$ and for $\rho \gtrsim \rho_c, E_x = \sqrt{x^2 + 2\beta \Delta x}$. These equations must be solved self-consistently with the corresponding equation for the chemical potential

$$\mu = \Delta + \frac{2\rho U_0}{(1 + \alpha U_0)}. \quad (25)$$

BEC occurs when $\Delta \to 0$ and $\alpha \to \infty$ from both sides. This yields $\mu = 0$ at the transition, as in the ideal gas, and the transition density is again given by condition (14).

Figure 4 shows the isotherm $k_B T = 0.1$, with $\rho_c a^3 = 0.0052$. Given the fact that this theory goes beyond mean-field by considering the renormalization of the coupling parameter, it is somewhat surprising that its predictions, in a way, fare less satisfactorily than HF and P. It is seen that in the normal region $\rho < \rho_c$ the solution is unstable near BEC, i.e. $(\partial \mu/\partial \rho)_T < 0$. In the BEC side, $\rho \geq \rho_c$, the situation is more worrisome, since it appears that $(\partial \mu/\partial \rho)_T$ becomes extremely large, probably diverging at the transition. This indicates that the compressibility becomes nearly zero, that is, the gas becomes incompressible at BEC. This is completely opposite to a critical behaviour where large density fluctuations are due to the large compressibility of the gas [22].
4. Final remarks and perspectives

In this work, we considered four of the most used mean-field theories to describe the onset of the BEC phase in weakly interacting gases. We calculated the equation of state \( \mu = \mu(\rho, T) \) corresponding to each of these theories at three different temperature regimes (with exception of the \( t \)-matrix theory which has calculated only at one temperature) satisfying the requirements of dilute gas in the contact potential approximation. The central conclusion is that none of the descriptions considered display a true second-order phase transition at BEC as expected by general symmetry breaking considerations. In addition, and excepting the Yukalov–Yukalova approximation at \( k_B T = 0.01 \) and 0.1, all the approximations considered exhibit unphysical behaviour in the vicinity of BEC. This anomalous property restricts severely its applicability at finite temperatures near the transition.

The interest in the equation of state \( \mu = \mu(\rho, T) \) goes beyond the order of the phase transition involved. As discussed in the introduction, it provides a fundamental link to the determination of the local density profile \( n(\rho) \) in gases confined by inhomogeneous magnetic or optical traps which is one of the main measurable quantities in the current experiments with ultracold alkali gases [7, 8, 30]. It is worthwhile to point out that a single profile at a temperature below BEC in the trapped gas bears information of the homogeneous one for densities \( \rho \) below, at and above the critical one \( \rho_c \).

Using the local density approximation (LDA) we have calculated density profiles for an isotropic 3D harmonic trap, \( V_{\text{ext}}(\rho) = \frac{1}{2} m \omega^2 r^2 \), using the isotherms \( \mu = \mu(\rho, T) \) obtained from the analysed theories. In this case the thermodynamic limit corresponds to \( N \to \infty, \omega \to 0 \) but \( N \omega^3 \) = constant. It is found that the predicted density profiles for HP and P are multiple valued for any temperature, while for YY this occurs only for relatively high temperatures. As an illustration of these multiple valuedness we show in figure 5 the YY profile obtained for \( k_B T = 1.1 \) by applying LDA to the corresponding case in figure 3. We recall that LDA, within the HF and P approximations, has been widely used to calculate thermodynamic properties of trapped gases, see e.g. [9, 10, 31–33], and the issue of multiple valuedness has not been brought up. This is perhaps due to the fact that when performing numerical calculations, the multiple-valued region being very small can be ‘short circuited’, inadvertently or not. In that case, the resulting profile is thus continuous and, apparently, the ensuing thermodynamic properties of the trapped gas are quite insensitive to this correction. We expect that the results provided in this work will bring again the attention to this matter. An accurate determination of density profiles would help to settle this issue and certainly it would provide solid guidelines to a better formulation of the theory of the weakly interacting gas.

Although the mean-field theories here described do not predict a second-order phase transition, their behaviour near \( T = 0 \) remains correct and valuable. With the exception of HF, the others correctly incorporate the gapless elementary excitation spectrum, originally predicted by Bogoliubov [28]. The results presented here suggest strongly that at temperatures near the transition, additional properties of the interatomic potential are needed to fully capture the thermodynamics of the system. Nevertheless, it is important to stress that exact calculations of the thermodynamics of quantum systems based on complete solutions of the Hamiltonian (8) are beyond present capabilities. We believe that in the construction of a new approximation or an improvement of those, the calculation of the equation of state \( \mu = \mu(\rho, T) \) should be a useful tool to determine the predicted order of the transition, and therefore, of the correct thermodynamic behaviour of the ultracold gas in the BEC state.

Acknowledgments

We acknowledge support from grants PAPIIT-UNAM IN114308 and IN116110. LOQ gratefully acknowledges support from CONACyT and Universidad Autonoma de la Ciudad de Mexico.

References

[1] Lieb E H, Seiringer R and Yngvason J 2005 Phys. Rev. Lett. 94 080401
[2] Shi H and Griffin A 1998 Phys. Rep. 304 1
[3] Andersen J O 2004 Rev. Mod. Phys. 76 599
[4] Yukalov V I 2007 Laser Phys. Lett. 4 632
[5] Lifshitz E M and Pitaevskii L P 1980 Statistical Physics II (London: Pergamon)
[6] Fetter A L and Walecka J D 2003 Quantum Theory of Many-Particle Systems (New York: Dover)
[7] Andersen M H, Ensher J R, Matthews M R, Wieman C E and Cornell E A 1995 Science 269 198
[8] Davis K B, Mewes M O, Andrews M R, vanDruten N J, Durfee D S, Kurn D M and Ketterle W 1995 Phys. Rev. Lett. 75 3969
[9] Dalfovo F, Giorgini S, Pitaevskii L P and Stringari S 1999 Rev. Mod. Phys. 71 463
[10] Goldman V V, Silvera I F and Leggett A J 1981 Phys. Rev. B 24 2870
[11] Pethick C J and Smith H 2002 Bose–Einstein Condensation in Dilute Gases (Cambridge: Cambridge University Press)
[12] Yukalov V I and Yukalova E P 2006 Phys. Rev. A 74 063623
   Yukalov V I and Yukalova E P 2007 Phys. Rev. A 76 013602
[13] Bijlsma M and Stoof H T C 1997 Phys. Rev. A 55 498
[14] Huang K, Yang C N and Luttinger J M 1957 Phys. Rev. 105 776
[15] Reatto L and Straley J P 1969 Phys. Rev. 183 321
[16] Callen H B 1960 Thermodynamics (New York: Wiley)
[17] Magalhães K M F, Muniz S R, Henn E A L, Silva R R, Marcassa L G and Bagnato V S 2005 Laser Phys. Lett. 2 214
[18] Garrod C and Simmons C 1972 J. Math. Phys. 13 1168
[19] Marchioro C and Presutti E 1973 Commun. Math. Phys. 29 265
[20] Sandoval-Figueroa N and Romero-Rochin V 2008 Phys. Rev. E 78 061129
[21] Landau L D and Lifshitz E M 1980 Statistical Physics I (London: Pergamon)
[22] Ma S-K 1976 Modern Theory of Critical Phenomena (London: Benjamin)
[23] Huang K 1987 Statistical Mechanics (New York: Wiley)
[24] Inouye S, Andrews M R, Stenger J, Miesner H-J, Stamper-Kurn D M and Ketterle W 1998 Nature 392 15
[25] Roberts J L, Claussen N R, Burke J F Jr, Greene C H, Cornell E A and Wieman C E 1998 Phys. Rev. Lett. 81 5109
[26] Streeker K E, Partridge G B, Truscott A G and Hulet R G 2002 Nature 417 150
[27] D’Errico C, Zaccanti M, Fattori M, Roati G, Inguscio M, Modugno G and Simoni A 2007 New J. Phys. 9 223
[28] Bogoliubov N N 1947 J. Phys. (Moscow) 11 23
[29] Steinhauer J, Ozser R, Katz N and Davidson N 2002 Phys. Rev. Lett. 88 120407
[30] Silva R R, Henn E A L, Magalhães K M F, Marcassa L G, Romero-Rochin V and Bagnato V S 2006 Laser Phys. 16 687
[31] Giorgini S, Pitaevskii L P and Stringari S 1996 Phys. Rev. A 54 R4633
[32] Minguzzi A, Conti S and Tosi M P 1997 J. Phys.: Condens. Matter 9 L23
[33] Romero-Rochin V 2005 Phys. Rev. Lett. 94 130601