

Role of the confinement-induced effective range in the thermodynamics of a strongly correlated Fermi gas in two dimensions

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We theoretically investigate the thermodynamic properties of a strongly correlated two-dimensional Fermi gas with a confinement-induced negative effective range of interactions, which is described by a two-channel model Hamiltonian. By extending the many-body T -matrix approach by Nozières and Schmitt-Rink to the two-channel model, we calculate the equation of state in the normal phase and present several thermodynamic quantities as functions of temperature, interaction strength, and effective range. We find that there is a nontrivial dependence of thermodynamics on the effective range. In the experiment, where the effective range is set by the tight axial confinement, the contribution of the effective range becomes nonnegligible as the temperature decreases down to the degenerate temperature. We compare our finite-range results with recent measurements on the density equation of state, and show that the effective range has to be taken into account for the purpose of a quantitative understanding of the experimental data.

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I. INTRODUCTION

Recent developments in creating ultracold atomic gases in lower dimensions has opened up a new and exciting field to explore strongly correlated many-body systems [1,2]. Two-dimensional (2D) ultracold atomic Fermi gases are of interest due to the increased role of thermal and quantum fluctuations, which lead to, for example, suppressed superfluid long-range order at nonzero temperature and a quasi-ordered transition to superfluidity, the Berezinskii-Kosterlitz-Thouless (BKT) transition [3,4]. In addition to presenting a range of novel and intriguing quantum phenomena, such as scale invariance and the breathing mode anomaly [5–9] and novel topological phases [10–12], 2D Fermi gases provide an important tool in understanding confined many-body systems from diverse fields of physics, such as high-temperature superconductors [13], thin ^3He films [14], neutron stars in a nuclear pasta phase [15], and exciton-polariton condensates in a microcavity [16].

The key advantage of 2D ultracold Fermi gases is their unprecedented controllability: the interatomic interaction can be tuned continuously with Feshbach resonances to realize the crossover from a Bardeen-Cooper-Schrieffer (BCS) superfluid of weakly interacting Cooper pairs to a Bose-Einstein condensate (BEC) of tightly bound molecules [2,17], the population of spin components can be changed [18], the confining potential can be made homogeneous with a box potential [19], and nonabelian synthetic gauge field (i.e., spin-orbit coupling) can be devised [20].

With these advancements, a set of seminal experiments directly probed the universal thermodynamics of a 2D interacting Fermi gas at given interaction strengths [17,21–23], by measuring the in-trap density profile. This allows for a direct comparison between theoretical predictions and experimental data. To date, it has been found that, a single-channel model

with a single 2D s -wave scattering length works reasonably well in describing the equation of state (EoS) experimental results, both in the normal state [24–28] and below the BKT transition [29–33]. However, in the strongly correlated regime it turns out that, the theoretical EoS determined by accurate auxiliary-field quantum Monte Carlo (AFQMC) simulations [31] slightly overestimates the thermodynamics compared to the measured EoS [2,17,21], suggesting the inefficiency of the single-channel model. The necessity of using a more appropriate theoretical model was highlighted by the most recent measurements on the breathing mode [34–36], which present an interesting example of a quantum anomaly (i.e., violation of the classical scale invariance [6]). It was found that the measured breathing mode is *notably* lower than the prediction from the single-channel model [8,9]. This discrepancy cannot be fully understood by the nonzero but small temperature found in the experiments [37]. Instead, it is caused by a confinement-induced effective range of interactions, which is negative and turns out to be significant under the current experimental conditions [38]. By adopting a two-channel model to account for the effective range, both measurements on low-temperature EoS and breathing mode anomaly can now be satisfactorily explained by calculations at *zero temperature* [39].

In this work, we explore how the confinement-induced effective range changes the *finite-temperature* thermodynamic properties of a normal, strongly correlated 2D Fermi gas. To date there have been no beyond-mean-field two-channel calculations on the thermodynamics of 2D Fermi gases in the normal state, where previous calculations focused on the description of the quasi-2D model [40,41] and the zero temperature behavior [42,43]. Our purpose is two-fold. First, characterizing the thermodynamics of the normal state of strongly correlated Fermi gases may be relevant to understanding the role of many-body pairing [44], which is a precursor to

superfluidity [24]. The results presented here can be used to better understand the existing two experimental measurements of the EoS at finite temperature in the normal phase [22,23]. Through the simplest many-body T -matrix theory developed by Nozières and Schmidt-Rink (NSR) [45], we determine the thermodynamic potential of the two-channel model, and calculate how the pressure EoS, energy, and entropy change with a *negative* effective range. Second, we then consider the realistic effective range in two recent experiments [22,23] and compare our finite-range results to the experimental data on density EoS. We find that in the experiments the effect of the effective range starts to show up when the temperature is decreased down to Fermi degeneracy. To quantitatively explain the experimental density EoS at finite temperature near superfluid transition, therefore, the effective range has to be taken into account in future refined theoretical works.

We note that, for a three-dimensional (3D) interacting Fermi gas, the effect of a negative effective range has been recently discussed by Tajima [46,47], following the seminal work by Ohashi and Griffin [48], who extended the NSR approach to the two-channel model. There, the effective range is related to the width of Feshbach resonance and the 3D interacting Fermi gas may experience severe atom loss in the narrow resonance limit (i.e., at a large negative effective range) [49,50]. In our case, the confinement-induced effective range is intrinsically set by the tight axial confinement and the 2D interacting Fermi gas is always mechanically stable.

Our paper is set out as follows. In Sec. II, we introduce the two-channel model Hamiltonian and renormalize the relevant parameters by solving the two-body scattering problem. In Sec. III, we briefly outline the many-body effective field theory of the model Hamiltonian and show how to calculate the thermodynamic observables. In Sec. IV, we discuss the thermodynamic properties of the 2D interacting Fermi gas, as functions of the effective range, temperature, and interaction strength. Finally in Sec. V, we summarize our results. For simplicity we set $\hbar = 1$ throughout.

II. HAMILTONIAN AND TWO-PARTICLE SCATTERING

We begin our calculation of the thermodynamic properties by considering a two-component Fermi gas in the normal state, described by the two-channel Hamiltonian [46–48,51,52]

$$\begin{aligned} \mathcal{H} = & \sum_{\mathbf{k}\sigma} \xi_{\mathbf{k}} c_{\mathbf{k}\sigma}^\dagger c_{\mathbf{k}\sigma} + \sum_{\mathbf{q}} (\epsilon_{\mathbf{q}}/2 + \nu - 2\mu) b_{\mathbf{q}}^\dagger b_{\mathbf{q}} \\ & + g \sum_{kq} (b_{\mathbf{q}} c_{\mathbf{q}/2+\mathbf{k}\uparrow}^\dagger c_{\mathbf{q}/2-\mathbf{k}\downarrow}^\dagger + \text{H.c.}), \end{aligned} \quad (1)$$

where H.c. is the Hermitian conjugate, $c_{\mathbf{k}\sigma}$ are the annihilation operators of fermionic atoms with spin $\sigma = \uparrow, \downarrow$ and mass M in the open channel, and $b_{\mathbf{q}}$ the annihilation operators of bosonic molecules in the closed channel. The kinetic energy of atoms measured from the chemical potential μ is $\xi_{\mathbf{k}} = \epsilon_{\mathbf{k}} - \mu$, where $\epsilon_{\mathbf{k}} = \mathbf{k}^2/(2M)$. The threshold energy or detuning of the diatomic molecules is ν and the Feshbach coupling between atoms and molecules is given by g .

To find the many-body properties we require the bare two-body scattering parameters to be rewritten in terms of

measurable or renormalizable scattering parameters. To relate the detuning ν and the channel coupling g to physical observables, we consider the two-body T -matrix in a vacuum ($E^+ \equiv k^2/M + i0^+$) [38,51],

$$T_{2B}^{-1}(E^+) = U_{\text{eff}}^{-1} + \sum_{\mathbf{p}} \frac{1}{2\epsilon_{\mathbf{p}} - E^+}, \quad (2)$$

where the effective interaction in the presence of the channel coupling is given by

$$U_{\text{eff}}(E^+) = \frac{g^2}{E^+ - \nu}. \quad (3)$$

Taking a large momentum cutoff $\Lambda \rightarrow \infty$, we write the two-body T -matrix as

$$T_{2B}^{-1}(E^+) = \frac{k^2/M - \nu}{g^2} + \frac{M}{4\pi} \left(\ln \left[\frac{\Lambda^2}{k^2} - 1 \right] + i\pi \right). \quad (4)$$

Alternatively, we may rewrite it in the form

$$T_{2B}(E^+) = \frac{m}{4\pi} (-2 \ln[ka_s] - R_s k^2 + i\pi), \quad (5)$$

where we wrote the detuning and Feshbach coupling in terms of the 2D scattering length a_s and effective range R_s [52],

$$a_s = \frac{1}{\Lambda} e^{\frac{2\pi\nu}{gM}}, \quad R_s = -\frac{4\pi}{M^2} \frac{1}{g^2}. \quad (6)$$

As can be seen from Eq. (6), we recover the single-channel model in the broad resonance limit when $g \rightarrow \infty$. We may remove the cutoff Λ by considering the pole of the two-body T -matrix $T_{2B}(E)$, $E = E_B$. We find that

$$\nu = E_B + g^2 \sum_{\mathbf{k}} \frac{1}{2\epsilon_{\mathbf{k}} - E_B}. \quad (7)$$

The binding energy can be set by $\epsilon_B = -E_B = \kappa^2/M$, where $k = i\kappa$ corresponds to the pole of the two-body T -matrix.

III. MANY-BODY T MATRIX

We consider the strong-coupling effects and pairing fluctuations by applying the normal-state NSR approach [45], which has been widely used in different context and has also been extended to superfluid phase [30,53–55]. For completeness, here we briefly go through the derivation of the thermodynamic potential using the functional path-integral formalism. All the thermodynamic properties can be obtained through the thermodynamic potential, $\Omega = -k_B T \ln \mathcal{Z}$, where the partition function \mathcal{Z} for the two-channel Hamiltonian is given by

$$\mathcal{Z} = \int \mathcal{D}(c, c^\dagger) \mathcal{D}(b, b^\dagger) e^{-S[c, c^\dagger, b, b^\dagger]}, \quad (8)$$

for atomic fields ($c_{\mathbf{k}\sigma}, c_{\mathbf{k}\sigma}^\dagger$) and molecular fields ($b_{\mathbf{q}}, b_{\mathbf{q}}^\dagger$), and the action at the inverse temperature $\beta = 1/(k_B T)$ is given by

$$\mathcal{S} = \int_0^\beta d\tau \left[\sum_{\mathbf{k}\sigma} c_{\mathbf{k}\sigma}^\dagger(\tau) c_{\mathbf{k}\sigma}(\tau) + \sum_{\mathbf{q}} b_{\mathbf{q}}^\dagger(\tau) \partial_\tau b_{\mathbf{q}}(\tau) + \mathcal{H}(\tau) \right]. \quad (9)$$

Using the Hubbard-Stratonovich transformation to decouple the Feshbach coupling term and integrating out the atomic fields, we obtain an effective action for Cooper pairs and molecules. By further truncating the perturbative expansion over bosonic fields (for both pairs and molecules) at the Gaussian fluctuation level, we arrive at the NSR thermodynamic potential [48]

$$\Omega = \Omega_F + \Omega_B - \sum_{\mathbf{q}, i\nu_n} \ln[1 + g^2 D_0(\mathbf{q}, i\nu_n) \Pi(\mathbf{q}, i\nu_n)], \quad (10)$$

where $\nu_n = 2n\pi/\beta$ are the bosonic Matsubara frequencies, $\Omega_F = 2 \sum_{\mathbf{k}} \ln(1 + e^{-\beta \epsilon_{\mathbf{k}}})$ and $\Omega_B = \sum_{\mathbf{q}} \ln(1 - e^{-\beta \epsilon_{\mathbf{q}}^B})$ are the free fermionic and bosonic thermodynamic potentials for atoms and molecules, respectively. We defined the pair correlation function

$$\Pi(\mathbf{q}, i\nu_n) = \sum_{\mathbf{k}} \frac{1 - f(\xi_{\frac{\mathbf{q}}{2} - \mathbf{k}}) - f(\xi_{\frac{\mathbf{q}}{2} + \mathbf{k}})}{2\epsilon_{\mathbf{k}} - 2\mu + \epsilon_{\mathbf{q}}/2 - i\nu_n}, \quad (11)$$

and $D_0(\mathbf{q}, i\nu_n) = 1/(i\nu_n - \epsilon_{\mathbf{q}}^B)$ is the Green's function of a free molecular boson with dispersion $\epsilon_{\mathbf{q}}^B = \epsilon_{\mathbf{q}}/2 - \nu + 2\mu$. The thermodynamic potential can be rewritten into the following form:

$$\Omega = \Omega_F - \sum_{\mathbf{q}, i\nu_n} \ln[-\Gamma^{-1}(\mathbf{q}, i\nu_n)], \quad (12)$$

where we introduced the vertex function

$$\Gamma^{-1}(\mathbf{q}, i\nu_n) = U_{\text{eff}}^{-1}(\mathbf{q}, i\nu_n) + \Pi(\mathbf{q}, i\nu_n), \quad (13)$$

and the in-medium effective interaction $U_{\text{eff}}(\mathbf{q}, i\nu_n) \equiv g^2 D_0(\mathbf{q}, i\nu_n)$, which can be explicitly written as

$$\frac{1}{U_{\text{eff}}(\mathbf{q}, i\nu_n)} = - \sum_{\mathbf{k}} \frac{1}{2\epsilon_{\mathbf{k}} + \epsilon_B} - \frac{M^2 R_s}{4\pi} \left(i\nu_n - \frac{\epsilon_{\mathbf{q}}}{2} + 2\mu + \epsilon_B \right). \quad (14)$$

Using Eq. (14) we cancel off the divergent parts in $\Pi(\mathbf{q}, i\nu_n)$. Analytically continuing the vertex function, $i\nu_n \rightarrow \omega + i0^+$, we calculate the thermodynamic potential

$$\Omega = \Omega_F - \frac{1}{\pi} \sum_{\mathbf{q}} \int_{-\infty}^{\infty} \frac{d\omega}{e^{\beta\omega} - 1} \delta(\mathbf{q}, \omega), \quad (15)$$

where $\delta(\mathbf{q}, \omega) \equiv -\text{Im} \ln[-\Gamma^{-1}(\mathbf{q}, \omega + i0^+)]$ is the phase of the vertex function. The number equation $nV = N = -d\Omega/d\mu$, where N is the total number of atoms and molecules and V is the area (or the volume in 2D), is solved to yield the chemical potential μ at a given set of reduced temperature T/T_F , binding energy ϵ_B/ϵ_F , and effective range $k_F^2 R_s$. Here, we define $k_F = (2\pi n)^{1/2}$, $E_F = k_F^2/(2M)$, and $T_F = E_F/k_B$.

We note that within the many-body T -matrix framework, we cannot calculate the superfluid transition temperature, i.e., the BKT transition temperature. This is due to the fact that the Thouless criterion in two dimensions becomes inapplicable as a direct consequence of Hohenberg's theorem and the loss of long-range order due to quantum fluctuations [13,56]. The self-consistent calculation of the chemical potential and Thouless criterion always leads to a *zero* critical temperature.

To consider the BKT transition the superfluid density needs to be calculated for the two-channel model in the superfluid phase, following the work in Ref. [33]. Such a scheme is beyond the scope of this work.

To calculate the thermodynamic properties of the entropy and energy, we define the dimensionless pressure equation of state f_p ,

$$\frac{P\lambda_T^2}{k_B T} \equiv f_p \left(x = \frac{T}{T_F}, y = \frac{\epsilon_B}{\epsilon_F}, z = k_F^2 R_s \right), \quad (16)$$

where the thermal wavelength is $\lambda_T = \sqrt{2\pi/(mk_B T)}$. All the other thermodynamic observables can then be calculated as derivatives from the dimensionless pressure equation of state, Eq. (16). The entropy is given by

$$S = - \left(\frac{\partial \Omega}{\partial T} \right)_{\mu} \equiv N k_B f_s, \quad (17)$$

where the dimensionless entropy takes the form

$$f_s = 2f_p - \frac{\tilde{\mu}}{x} f_{p_x} - \frac{k_{BY}}{x} f_{p_y}, \quad (18)$$

and $f_{p_x} \equiv \partial f_p / \partial x$, and $\tilde{\mu} = \mu/\epsilon_F$. The energy is found through $E = -TS + \Omega + \mu N \equiv N \epsilon_F f_E$, and we obtain

$$f_E = x f_s + x \frac{f_p}{f_{p_x}} + \tilde{\mu}. \quad (19)$$

To close this section, it is worth emphasizing that the many-body T -matrix theory does not take into account the interactions between the underlying characteristic particles. It works well either in the tight-binding limit, where the binding energy $\epsilon_B \gg \epsilon_F$ and molecules are well-formed, or in the noninteracting limit, where the binding energy is exponentially small. In the BEC-BCS crossover regime of most interest, such as $\epsilon_B \sim [0.01 - 1.0]\epsilon_F$, the interactions between performed molecules are strong, and the many-body T -matrix theory provides only a qualitative description.

IV. RESULTS AND DISCUSSIONS

A. Pressure equation of state

To begin our analysis of the thermodynamic properties of the system, we plot the pressure equation of state (EoS) in Fig. 1, which is normalized with the pressure of an ideal Fermi gas at the same temperature, $P_0 = -2\pi\lambda_T^{-4} \text{Li}_2(-e^{\beta\mu})$, where $\text{Li}_s(z)$ is the polylogarithm function. We show the pressure EoS at temperatures (a) $T/T_F = 0.5$, (b) $T/T_F = 1$, (c) and $T/T_F = 2$, for a range of interaction strengths, from $\epsilon_B/\epsilon_F = 0.01$ to $\epsilon_B/\epsilon_F = 2$. We find for decreasing temperature and increasing binding energy the normalized pressure increases, indicating that the system is becoming more strongly interacting. As the effective range decreases, the normalized pressure increases for all temperatures and interaction strengths. This general trend is anticipated, as the system becomes less correlated as the number of molecules increases. We can calculate the bosonic thermodynamic potential in the infinite negative effective range limit, $R_s \rightarrow -\infty$, where the thermodynamic potential reduces to the mean-field theory of a mixture of

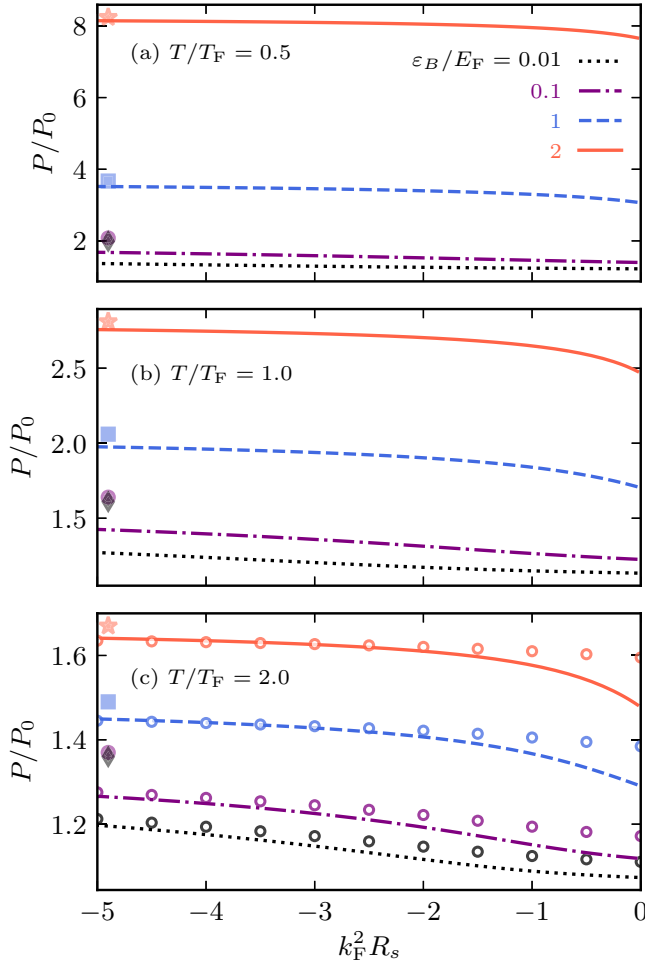


FIG. 1. Pressure EoS, normalized by the ideal pressure P_0 at the same temperature, is plotted as a function of the effective range for interaction strengths $\varepsilon_B/\varepsilon_F = 0.01$ (black-dotted), $\varepsilon_B/\varepsilon_F = 0.1$ (purple-dot dashed), $\varepsilon_B/\varepsilon_F = 1$ (blue-dashed), and $\varepsilon_B/\varepsilon_F = 2$ (red-solid), at temperatures (a) $T/T_F = 0.5$, (b) $T/T_F = 1.0$, and (c) $T/T_F = 2.0$. At high temperature in (c), the second-order virial expansion results for each interaction strength are shown by circles. The results of the infinite negative effective range are given by the symbols, where the star is for $\varepsilon_B/\varepsilon_F = 2$, square for $\varepsilon_B/\varepsilon_F = 1$, circle for $\varepsilon_B/\varepsilon_F = 0.1$, and diamond for $\varepsilon_B/\varepsilon_F = 0.01$.

atoms and molecules, $\Omega = \Omega_F + \Omega_B$ and

$$\Omega_B = -2 \sum_{\mathbf{q}} \ln[1 - e^{-\beta(\varepsilon_{\mathbf{q}}/2 - 2\mu - \varepsilon_B)}]. \quad (20)$$

The symbols in Fig. 1 are the limiting values for each binding energy calculated from solving the mean-field number equation, where the star is for $\varepsilon_B/\varepsilon_F = 2$, square for $\varepsilon_B/\varepsilon_F = 1$, circle for $\varepsilon_B/\varepsilon_F = 0.1$, and diamond for $\varepsilon_B/\varepsilon_F = 0.01$. We see that the pressure EoS is approaching the infinite negative effective range limit values for large binding energies and all temperatures, due to there being more molecule formation on the tight-binding side. However, in the BEC-BCS crossover regime (i.e., for $\varepsilon_B/\varepsilon_F = 0.1$ or 0.01), the system is approaching the limit rather slowly.

As a comparison in the high-temperature regime, we consider the virial expansion up to second order [26,57–60],

which is, in principle, exact for large-enough temperatures. The dimensionless pressure equation of state is given by

$$f_p = \int_0^\infty dt \ln[1 + ze^{-t}] + \Delta b_2 z^2 + \dots, \quad (21)$$

where Δb_2 is the second-order virial coefficient. By using the elegant Beth-Uhlenbeck relation [57], we obtain

$$\Delta b_2 = e^{\beta\varepsilon_B} - \int_0^\infty \frac{dk}{k} e^{-2k^2} \frac{2k^2 R_s + 2}{(\ln[\frac{2k^2}{\varepsilon_B}] + k^2 R_s)^2 + \pi^2}, \quad (22)$$

which in the zero-range limit reduces to the known results [22,25]. As shown in Fig. 1(c) at high temperature $T = 2T_F$, we find that as the effective range decreases, the virial expansion provides an excellent description of the pressure EoS, even for the largest binding energy. This is quite different from the zero-range limit. At $k_F^2 R_s = 0$, where the single-channel model is applicable, the virial expansion systematically overestimates the pressure EoS. The improved applicability of virial expansion is again due to the weaker correlation at larger effective range which means a smaller g and hence weaker interaction effect.

B. Entropy and energy

Having discussed the pressure EoS, we now consider the entropy per particle $S/(Nk_B)$, as shown in the upper panels of Fig. 2 as a function of the negative effective range at three different temperatures, (a) $T/T_F = 0.5$, (b) $T/T_F = 1.0$, and (c) $T/T_F = 2.0$, for a range of binding energies. We also plot the infinite negative effective range limit for each interaction and temperature, as shown by the symbols. We find that the entropy has a consistent behavior for all interaction strengths and temperatures: the entropy increases as the effective range decreases. In particular, at low temperature as the negative effective range decreases, we see for large binding energies (i.e., $\varepsilon_B/\varepsilon_F = 1, 2$) the entropy increases rapidly. The entropy is quickly converging to the infinite-range limit for low temperature. For high temperatures at the BEC-BCS crossover regime, as shown in (c) the residual interaction at large effective range seems still nonnegligible and the entropy is not fully approaching the infinite limit.

In the lower panels of Fig. 2, we plot the energy per particle, $E/(N\varepsilon_F)$, with the two-body contribution from pairs (where we assumed that there are $N/2$ pairs each with energy $-\varepsilon_B$) subtracted. We see that the energy is decreasing with decreasing temperature for all interaction strengths. For the weakest binding energy, the energy decreases as the effective range decreases, for all temperatures. However, for larger binding energies ($\varepsilon_B/\varepsilon_F = 1, 2$), the energy increases at low temperature with decreasing effective range; while at higher temperatures the energy decreases. The different effective-range dependence of the energy at low and high temperatures may be understood from the many-body pairing. In the low-temperature regime for large binding energy, the many-body pairing is important and the system becomes rigid with respect to the change of the effective range. Hence, the energy slightly increases, similar to the pressure EoS. In contrast, at high temperature the many-body pairing is less significant and

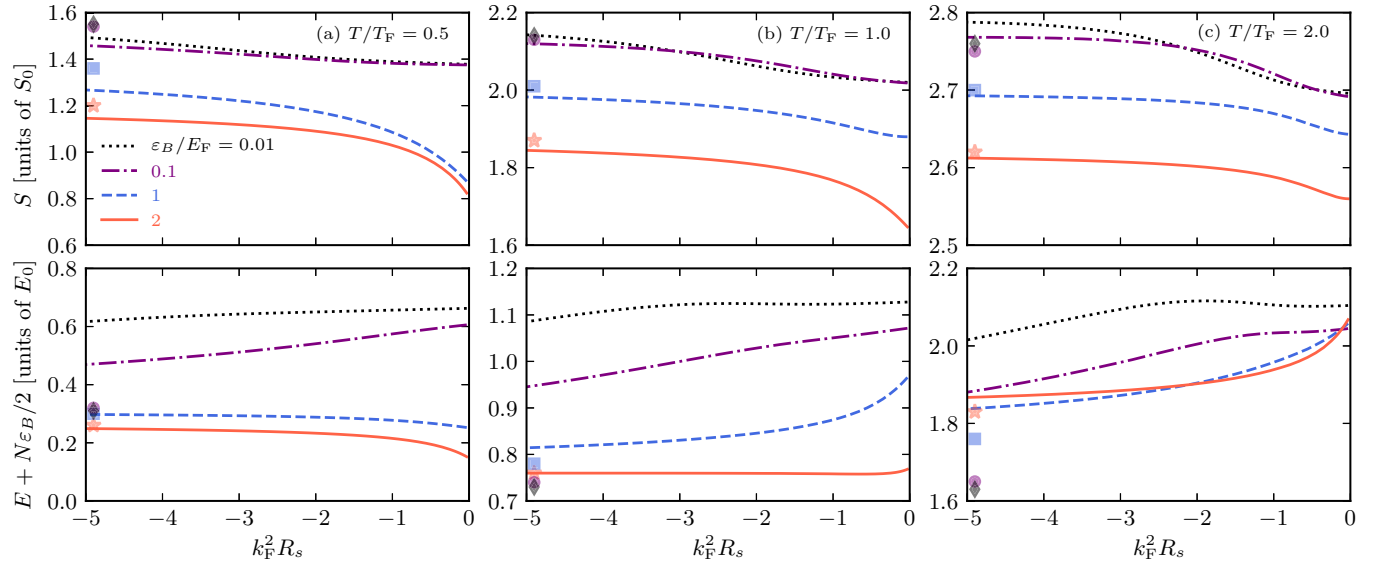


FIG. 2. Upper panels: Entropy normalized in units of $S_0 = k_B N$ as a function of the effective range at different interaction strengths $\epsilon_B/\epsilon_F = 0.01$ (black-dotted), $\epsilon_B/\epsilon_F = 0.1$ (purple-dot-dashed), $\epsilon_B/\epsilon_F = 1$ (blue-dashed), and $\epsilon_B/\epsilon_F = 2$ (red-solid). The columns (a), (b), and (c) are for temperatures $T/T_F = 0.5$, $T/T_F = 1$, and $T/T_F = 2$, respectively. Lower panels: The effective range dependence of the energy, in units of $E_0 = N\epsilon_F$, with the same convention as in the upper panels. The results of the infinite negative effective range are given by the symbols, where the star is for $\epsilon_B/\epsilon_F = 2$, square for $\epsilon_B/\epsilon_F = 1$, circle for $\epsilon_B/\epsilon_F = 0.1$, and diamond for $\epsilon_B/\epsilon_F = 0.01$.

the system experiences a character change from atoms to molecules with increasing effective range. The energy then decreases, roughly following the picture of a noninteracting Bose and Fermi mixture.

To see the interaction effects on the thermodynamic properties, we show in Fig. 3 the dimensionless entropy (upper panels) and energy (lower panels) as a function of the binding energy ϵ_B/ϵ_F , for a range of negative effective ranges from $k_F^2 R_s = 0$ to -3 at temperatures (a) $T/T_F = 0.3$, (b) $T/T_F =$

0.5, and (c) $T/T_F = 1.0$. For comparison, we show also the ideal gas limits of Fermi and Bose gases for the entropy and energy. For the ideal Fermi gas limit, we use the following formulas:

$$\frac{S^F}{Nk_B} = \frac{2\text{Li}_2(-e^{\mu/\epsilon_F})}{\text{Li}_1(-e^{\mu/\epsilon_F})} - \frac{\mu}{\epsilon_F}, \quad (23)$$

$$\frac{E^F}{N\epsilon_F} = -\left(\frac{T}{T_F}\right)^2 \text{Li}_2(-e^{\mu/\epsilon_F}), \quad (24)$$

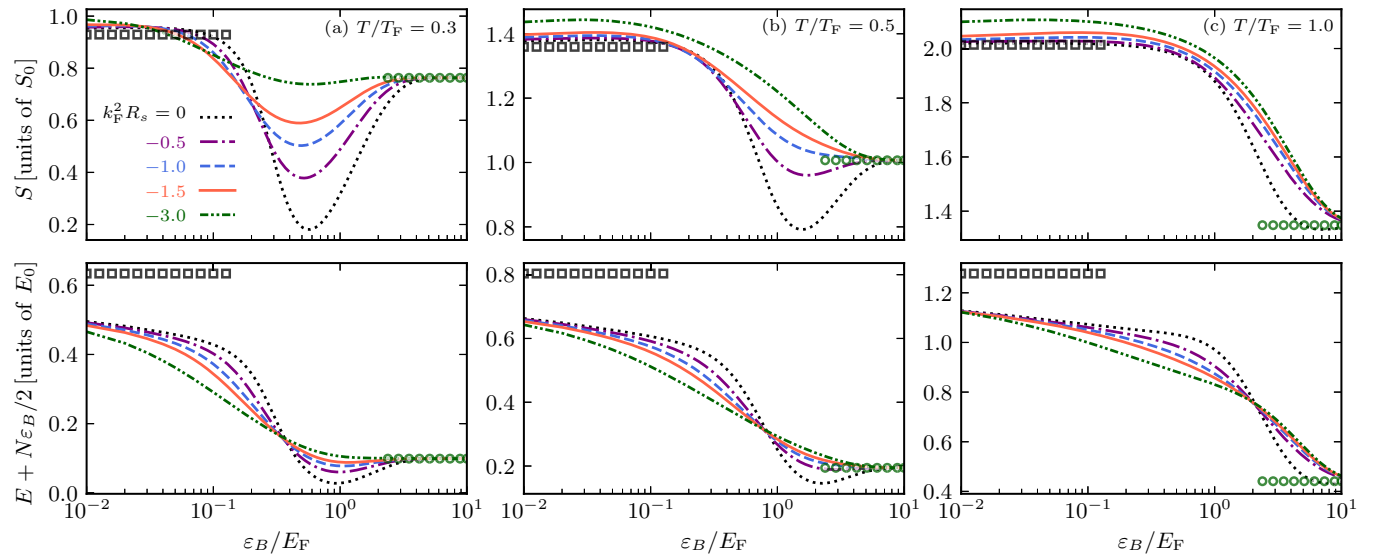


FIG. 3. Upper panels: Entropy is plotted in units of $S_0 = Nk_B$ as a function of the interaction strength at different negative effective ranges: $k_F^2 R_s = 0$ (black dotted), $k_F^2 R_s = -0.5$ (purple dot-dashed), $k_F^2 R_s = -1$ (blue dashed), $k_F^2 R_s = -1.5$ (red solid), and $k_F^2 R_s = -3$ (green dot-dot-dashed), for temperatures (a) $T/T_F = 0.3$, (b) $T/T_F = 0.5$, and (c) $T/T_F = 1.0$. The square and circular symbols are the ideal Fermi and Bose gas limits (see text), respectively. Lower panels: Energy is plotted in units of $E_0 = N\epsilon_F$. The same convention was taken as in the upper panels.

with a noninteracting chemical potential μ determined by using the number equation. To find the entropy and energy for noninteracting Bose gases S^B and E^B , we assume a gas of $N/2$ noninteracting molecules with mass $2M$ and similarly solve a molecular chemical potential.

The behavior of the entropy as a function of the binding energy is nontrivial. We find that for $k_F^2 R_s = 0$, as the interaction is increased from the weakly to strongly attractive regimes, the entropy has a local minimum at $\varepsilon_B/\varepsilon_F \simeq 0.6$ for temperature $T/T_F = 0.3$, $\varepsilon_B/\varepsilon_F \simeq 1.05$ for temperature $T/T_F = 0.5$, and $\varepsilon_B/\varepsilon_F \simeq 5$ for temperature $T/T_F = 1.0$. This minimum may be understood as the position where pair formation is the strongest, and at low temperature where there is a defined Fermi surface, i.e., $\mu > 0$ for $\varepsilon_B/\varepsilon_F \lesssim 1$, this pairing is dominated by many-body pairing. As the negative effective range decreases, we see this clear minimum in the crossover interaction regime become shallower. This is due to the fact that the system more readily forms bound molecules as the negative effective range decreases. For $T/T_F = 0.3$ the minimum shifts to weaker binding energies with decreasing effective range, and for higher temperatures the minimum shifts to larger binding energies and disappears for $k_F^2 R_s < -1$, consistent with the naive picture that high temperature dampens the formation of many-body pairs. In the weakly attractive regime the entropy quickly approaches the noninteracting 2D Fermi gas limit and in the strongly attractive regime the entropy approaches the limit of $N/2$ noninteracting molecules with mass $2M$.

The behavior of the energy is qualitatively the same as the entropy. We see that there exists a local minimum that shifts to larger binding energy as the temperature increases. We find also that the minimum in the crossover interaction regime becomes much weaker as the negative effective range decreases. Unlike the entropy, the energy is slowly approaching the noninteracting 2D Fermi gas in the weakly attractive regime. However, in the strongly attractive regime the energy quickly approaches the limit of $N/2$ noninteracting molecules.

C. Comparison to experiment

We now compare our two-channel results to the experimental data on density EoS, with a realistic confinement-induced effective range. For this purpose, we match the the low-energy expansion of $T_{2B}(E^+ \equiv k^2/M + i0^+)$ to the quasi-2D scattering amplitude $f_{Q2D}(k)/M$, which describes the two-particle scattering within the ground-state manifold under a tight axial confinement with frequency ω_z [61]:

$$f_{Q2D}(k \rightarrow 0) = \frac{4\pi}{\sqrt{2\pi} a_z/a_{3D} + \varpi(k^2 a_z^2/2)}, \quad (25)$$

where $a_z \equiv \sqrt{1/(M\omega_z)}$ is the harmonic oscillator length and the function $\varpi(x)$ has the low-energy expansion, $\varpi(x \rightarrow 0) \simeq -\ln(2\pi x/\mathcal{B}) + 2x \ln 2 + i\pi$, with $\mathcal{B} \simeq 0.9049$.

By setting $T_{2B}(E^+) = f_{Q2D}(k)/M$, we obtain the well-known result [61]

$$a_s = a_z \frac{\pi}{\mathcal{B}} \exp\left(-\sqrt{\frac{\pi}{2}} \frac{a_z}{a_{3D}}\right), \quad (26)$$

in the zero-energy limit $k \rightarrow 0$. The determination of the effective range R_s is also straightforward. We require that

the two-body T -matrix $T_{2B}(E^+)$ and the quasi-2D scattering amplitude share the same pole or the same binding energy ε_B [39]. For the former, it is readily seen from Eq. (5) that the binding energy $\varepsilon_B = \kappa^2/M$ is related to the effective range R_s by

$$R_s = \frac{2 \ln(\kappa a_s)}{\kappa^2}. \quad (27)$$

The binding energy can also be obtained from the quasi-2D scattering amplitude by solving the equation [61]

$$\frac{a_z}{a_{3D}} = \mathcal{F}\left(\frac{\varepsilon_B}{\omega_z}\right), \quad (28)$$

where

$$\mathcal{F}(x) = \int_0^\infty \frac{du}{\sqrt{4\pi u^3}} \left(1 - \frac{e^{-xu}}{\sqrt{(1 - e^{-2u})/(2u)}}\right). \quad (29)$$

For a given a_z/a_{3D} , we can solve Eq. (28) for $\varepsilon_B = \kappa^2/M$, and then use Eq. (27) to calculate R_s . In this way, we can determine the dimensionless ratio R_s/a_s^2 as a function of a_z/a_{3D} [39].

Experimentally, the 2D density EoS is measured at a given magnetic field (i.e., a_{3D}) and temperature through the density profile of a cloud of N interacting fermions. By applying the local density approximation, the local density n is calibrated as a function of the normalized local chemical potential $\beta\mu$ [22,23]. This gives rise to the *homogeneous* density EoS $n(\mu)$ at dimensionless interaction parameters $\beta\varepsilon_B$ and R_s/a_s^2 , allowing for a direct comparison with the theoretical EoS.

Figure 4(a) plots the density EoS as a function of dimensionless chemical potential at $\beta\varepsilon_B = 1.2$ and $R_s/a_s^2 \simeq -0.2$, normalized by the ideal gas density at the same chemical potential, i.e., $n_0 = -\lambda_T^{-2} \text{Li}_1(-e^{\beta\mu})$. Here, the interaction parameters $\beta\varepsilon_B = 1.2$ and $R_s/a_s^2 \simeq -0.2$ are taken according to the recent density EoS measurement in Ref. [23]. We plot the single-channel prediction (red-dotted line, by artificially setting $R_s = 0$), the two-channel prediction (blue dot-dashed line), and the experimental data (squares). We see that including the *realistic* negative effective range quantitatively improves the comparison to experiment in the high-temperature regime, down to $\beta\mu = -0.25$ (which corresponds to a temperature $T = 0.7T_F$), although as $\beta\mu$ approaches $-\infty$, the effective range has no significant contribution due to the vanishingly small density and hence $k_F^2 R_s \sim 0$, as we would expect. Typically, the normal-state NSR theory breaks down at low temperature, as indicated by a divergent density EoS. The NSR calculation of the two-channel model breaks down at a higher temperature than the single-channel NSR, as a result of the fact that the chemical potential is approaching the two-body bound state of the closed-channel molecules.

In Fig. 4(b) we compare the theoretical predictions on density EoS to the experimental results of Ref. [22] at $\beta\varepsilon_B = 0.47$ and $R_s/a_s^2 \simeq -0.03$. In this case, we are on the BCS side with a larger 2D scattering length a_s , so the effect of the effective range may become relatively weaker. Although the inclusion of the effective range does significantly shift the density EoS towards the low-temperature regime (i.e., $\beta\varepsilon_B \sim 1$), it does not approach the experimental results fully.

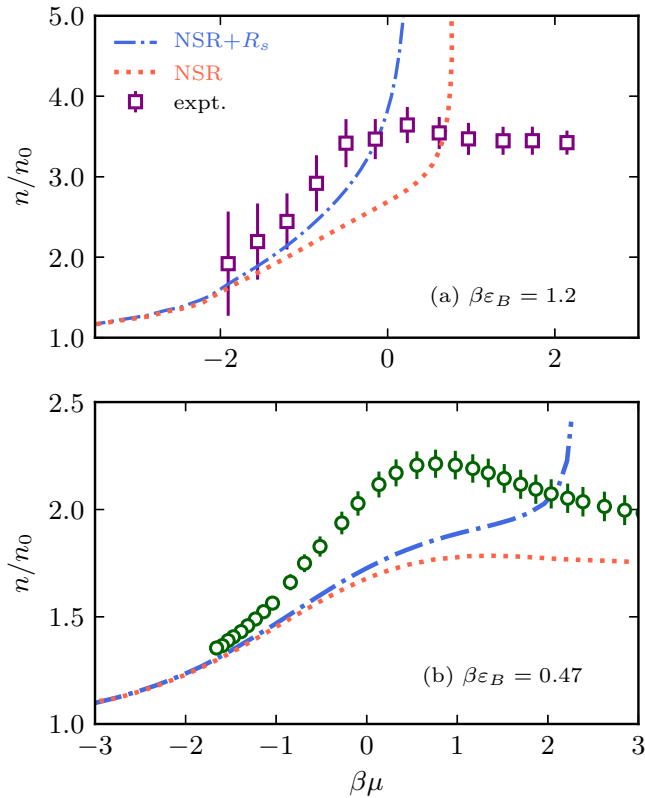


FIG. 4. (a) Density EoS, normalized by $n_0 = -\lambda_T^{-2} \text{Li}_1(-e^{\beta\mu})$, as a function of the dimensionless chemical potential $\beta\mu$ and interaction strength $\beta\epsilon_B = 1.2$, for the single-channel model (red-dotted, $R_s = 0$), the two-channel model (blue dot-dashed, $R_s \neq 0$), and the experimental results from Ref. [23]. (b) Density EoS at $\beta\epsilon_B = 0.47$, with the experimental results from Ref. [22].

This is anticipated, as the simple many-body T -matrix theory such as NSR is known to underestimate the pair correlations close to the BKT transition [28].

Although our NSR theory with realistic effective ranges cannot provide a *quantitative* explanation of the experimental

data, the message obtained from the comparison of the two cases (i.e., with and without the effective range) is clear: one needs to take into account the confinement-induced effective range in future theoretical studies of an interacting 2D Fermi gas at finite temperature. We expect the self-consistent two-channel calculation, as done in 3D by Ref. [51], would improve the numerical calculation and comparison to experiment. Although there is no *definite* reason to choose a diagrammatic scheme, the self-consistent calculation will include interactions between dressed fermionic and bosonic Green's function that are important in reduced dimensions. This calculation will be considered in future work.

V. CONCLUSION

In summary, we theoretically investigated the thermodynamics of a strongly correlated Fermi gas confined to two dimensions with a negative confinement-induced effective range. Within a two-channel model, including pairing fluctuations beyond the mean-field level we discussed the negative effective range corrections to the thermodynamic properties. Using the recent experimental data [22,23] as a benchmark, we find that density equation of state improves in our two-channel model, compared to the widely used single-channel model.

We show that as the effective range decreases, the entropy increases, and the system eventually approaches a molecular Bose-Einstein condensate with finite bound state energy, thus making it more difficult for free fermions to form pairs. In future works the many-body pairing should be carefully examined. The appearance of 2D many-body pairing at high temperature in the crossover regime remains an interesting and controversial topic to be explored [44,62–64].

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