

Collective motion of polarized dipolar Fermi gases in the hydrodynamic regime

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Recently, a seminal stimulated Raman adiabatic passage (STIRAP) experiment allowed the creation of $^{40}\text{K}^{87}\text{Rb}$ molecules in the rovibrational ground state [K.-K. Ni *et al.*, *Science* **322**, 231 (2008)]. To describe such a polarized dipolar Fermi gas in the hydrodynamic regime, we work out a variational time-dependent Hartree-Fock approach. With this we calculate dynamical properties of such a system, for instance, the frequencies of the low-lying excitations and the time-of-flight expansion. We find that the dipole-dipole interaction induces anisotropic breathing oscillations in momentum space. In addition, after release from the trap, the momentum distribution becomes asymptotically isotropic, while the particle density becomes anisotropic.

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Even before the realization of Bose-Einstein condensation (BEC) with ^{52}Cr [1], much experimental and theoretical interest was dedicated to ultracold quantum gases interacting through the long-range and anisotropic dipole-dipole interaction (DDI) [2]. For bosonic dipolar particles, the starting point of the theoretical investigations was the construction of a corresponding pseudopotential by Yi and You [3]. After that, an exact solution of the Gross-Pitaevskii equation in the Thomas-Fermi regime was found for cylinder-symmetric traps [4]. Moreover, the DDI has been shown to shift the BEC critical temperature in a characteristic way in polarized systems [5] and to give rise to the Einstein-de Haas effect, when spinorial degrees of freedom are considered [6]. From the experimental point of view, time-of-flight (TOF) techniques demonstrated both the first DDI signature through small mechanical effects [7] and strong dipolar effects in quantum ferrofluids [8]. Furthermore, the shape of the trap was manipulated to stabilize a purely dipolar BEC against collapse [9].

Concerning fermionic dipolar systems, recent theoretical studies have considered interesting properties of homogeneous gases such as zero sound [10], Berezinskii-Kosterlitz-Thouless phase transition [11], and nematic phases [12]. In harmonically trapped systems, amazing predictions like anisotropic superfluidity [13], fractional quantum Hall physics [14], and Wigner crystallization [15] have been made. With respect to experimental investigations, the most promising atomic candidate is the fermionic chromium isotope ^{53}Cr [16], which has a magnetic moment of $m = 6$ Bohr magnetons. For these atoms, calculations of equilibrium properties have shown that the DDI is only a small perturbation [17,18]. However, by applying a stimulated Raman adiabatic passage (STIRAP) process, it has recently been achieved to cool and trap $^{40}\text{K}^{87}\text{Rb}$ molecules into their rovibrational ground state, where they possess an electric dipole moment of $d = 0.566$ Debye [19–22]. Due to the resulting strong DDI a considerable deformation of the momentum distribution is expected [17,18]. Once these systems would have been further cooled into the quantum degenerate regime, the main task will be to identify

unambiguously the presence of the DDI. In this respect, TOF experiments and oscillation frequency measurements represent the most fundamental diagnostic tools in the field of ultracold quantum gases. Their outcomes reveal important information on the nature of the system under investigation. They differ drastically depending on whether the system is in the collisionless (CL) regime, where collision rates are small, or in the hydrodynamic (HD) regime, where collisions take place so often that they lead to local equilibrium. To date, investigations of dynamical properties of trapped dipolar Fermi gases either have been restricted to the CL regime [23] or have excluded a deformation of the momentum distribution in the HD regime [24]. Since experiments with ultracold polar molecules are performed under strong dipolar interactions, one should expect them to lead the system into the HD regime, and thus an analysis allowing for anisotropy in the momentum distribution must be carried out. In this article, we use a variational time-dependent Hartree-Fock approach to address this question.

Consider N spin-polarized fermionic dipoles of mass M trapped in a cylinder-symmetric harmonic potential $U_{\text{tr}}(\mathbf{x}) = M\omega_x^2(x^2 + y^2 + \lambda^2 z^2)/2$ with trap anisotropy λ at ultralow temperatures. Since the Pauli principle inhibits a contact interaction, they interact dominantly through DDI. As we assume that the fermionic cloud is polarized along the symmetry axis of the trap, the DDI potential reads $V_{\text{dd}}(\mathbf{x}) = \frac{C_{\text{dd}}}{4\pi|\mathbf{x}|^3} [1 - 3\frac{z^2}{|\mathbf{x}|^2}]$. For magnetic dipole moments m the DDI is characterized by $C_{\text{dd}} = \mu_0 m^2$, whereas for electric moments d we have $C_{\text{dd}} = 4\pi d^2$. In the following we restrict ourselves to the normal phase in the limit $T \rightarrow 0$ because the critical temperature for superfluidity is very low, depending exponentially on $a_{\text{dd}} = MC_{\text{dd}}/(4\pi\hbar^2)$ [13]. Furthermore, this limit is restricted by the HD requirement that the relaxation time τ_R is small in comparison with the time scale $1/\bar{\omega}$ defined by the average trap frequency $\bar{\omega} = (\omega_x^2 \omega_z)^{1/3}$. The necessity of an HD approach can be inferred as follows. As τ_R is not known for dipolar interactions, we estimate it by assuming the DDI to be equivalent to a contact interaction with scattering length a_{dd} . Then we use the fact that for a two-component, degenerate, normal Fermi gas with contact interaction, one has $(\bar{\omega}\tau_R)^{-1} = (N^{1/3} a_{\text{dd}} \sqrt{M\bar{\omega}/\hbar})^2 F(T/T_F)$, where $F(T/T_F)$ is of the order 0.1 in the quantum temperature regime (see,

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e.g., [25]). Thus, we expect for the one-component, dipolar gas to enter the HD regime for $N^{1/6}\epsilon_{\text{dd}} \gg 1$, with the dimensionless parameter $\epsilon_{\text{dd}} = C_{\text{dd}}(M^3\bar{\omega}/\hbar^5)^{1/2}N^{1/6}/4\pi$ measuring the strength of the DDI. In the current setup of Ref. [22] one has 4×10^4 $^{40}\text{K}^{87}\text{Rb}$ molecules with a radial trapping frequency of $\omega_x = \omega_y \approx 2\pi \times 175$ Hz. Assuming an average trap frequency of that value yields at least $\epsilon_{\text{dd}} \approx 5.3$ and $(\bar{\omega}\tau_R)^{-1} \approx 0.1 \times (N^{1/6}\epsilon_{\text{dd}})^2 \approx 96$, which drives the system into the HD regime.

In this article we work out a time-dependent Hartree-Fock approach by extremizing the action $\mathcal{A} = \int dt \langle \Psi | i\hbar \frac{\partial}{\partial t} - \hat{H} | \Psi \rangle$, where $\Psi(x_1, \dots, x_N, t) = \langle x_1, \dots, x_N | \Psi \rangle$ is a Slater determinant and \hat{H} denotes the underlying Hamilton operator. To describe the HD regime, we follow a standard procedure of nuclear physics [26] and assume that frequent particle collisions assure that all one-particle orbitals have the same local phase $\chi(x, t)$, yielding the velocity field $\mathbf{v} = \nabla\chi$. Thus, we can factorize out the phases and define a Slater determinant through $\Psi_0(x_1, \dots, x_N, t) = e^{-iM \sum_i \chi(x_i, t)/\hbar} \Psi(x_1, \dots, x_N, t)$, which contains only the moduli of the one-particle orbitals and, therefore, is invariant under time reversal. This yields a time-even one-body density matrix $\rho_0(x, x'; t) = e^{-iM[\chi(x, t) - \chi(x', t)]/\hbar} \rho(x, x'; t)$ [27]. With this the action reduces to

$$\mathcal{A} = -M \int dt \int d^3x \left\{ \dot{\chi}(x, t) \rho_0(x; t) + \frac{\rho_0(x; t)}{2} [\nabla\chi(x, t)]^2 \right\} - \int dt \langle \Psi_0 | \hat{H} | \Psi_0 \rangle, \quad (1)$$

where $\rho_0(x; t) = \rho_0(x, x; t)$ denotes the particle density and $\langle \Psi_0 | \hat{H} | \Psi_0 \rangle$ consists of the kinetic energy E_{kin} , the trapping potential E_{tr} , and the interaction. The latter is divided into the direct or Hartree term E_{dd}^{D} and the exchange or Fock term E_{dd}^{E} . Due to the exchange term, the ground-state energy $\langle \Psi_0 | \hat{H} | \Psi_0 \rangle$ is not a function of the particle density $\rho_0(x; t)$ alone but also contains the nondiagonal part $\rho_0(x, x'; t)$.

As it is not possible to solve analytically the resulting Euler-Lagrange equations for $\chi(x, t)$ and $\rho_0(x, x'; t)$, we propose here a variational extremization of the action. To this end, we express each energy contribution in terms of the Wigner transform of the one-body density matrix $\nu_0(\mathbf{X}, \mathbf{k}; t) = \int d^3s \rho_0(\mathbf{X} + \frac{s}{2}, \mathbf{X} - \frac{s}{2}; t) e^{-i\mathbf{k}\cdot\mathbf{s}}$. The kinetic and trapping energies are then given by

$$E_{\text{kin/tr}} = \int \frac{d^3x d^3k}{(2\pi)^3} \nu_0(\mathbf{x}, \mathbf{k}; t) \epsilon_{\text{kin/tr}}(\mathbf{x}, \mathbf{k}), \quad (2)$$

with $\epsilon_{\text{kin}} = \hbar^2 \mathbf{k}^2 / 2M$ and $\epsilon_{\text{tr}} = U_{\text{tr}}(\mathbf{x})$, respectively. The direct term, which accounts for the deformation of the particle density, and the exchange term, which is related to the momentum space deformation, read

$$E_{\text{dd}}^{\text{D}} = \int \frac{d^3x d^3k d^3x' d^3k'}{2(2\pi)^6} \nu_0(\mathbf{x}, \mathbf{k}; t) V_{\text{dd}}(\mathbf{x} - \mathbf{x}') \nu_0(\mathbf{x}', \mathbf{k}'; t),$$

$$E_{\text{dd}}^{\text{E}} = - \int \frac{d^3X d^3k d^3s d^3k'}{2(2\pi)^6} \nu_0(\mathbf{X}, \mathbf{k}; t) V_{\text{dd}}(\mathbf{s}) \nu_0(\mathbf{X}, \mathbf{k}'; t) \times e^{i\mathbf{s}\cdot(\mathbf{k}-\mathbf{k}')}.$$
(3)

At this point, we adopt the variational ansatz $\chi(x, t) = [\alpha_x(t)(x^2 + y^2) + \alpha_z(t)z^2]/2$ for the phase and $\nu_0(\mathbf{x}, \mathbf{k}; t) =$

$\Theta(1 - \frac{x^2+y^2}{\tilde{R}_x(t)^2} - \frac{z^2}{\tilde{R}_z(t)^2} - \frac{k_x^2+k_y^2}{\tilde{K}_x(t)^2} - \frac{k_z^2}{\tilde{K}_z(t)^2})$ for the Wigner phase space function, with $\Theta(\cdot)$ being the step function. We are now in the position to extremize the action (1) with respect to the time-dependent variational parameters $\alpha_i(t)$ for the phase as well as $\tilde{R}_i(t)$ and $\tilde{K}_i(t)$ for the Thomas-Fermi radii and the Fermi momenta. At first, one obtains $\alpha_i = \tilde{R}_i/R_i$, which is used to eliminate the parameters α_i from the rest of the formalism. Under conservation of the particle number,

$$\tilde{R}_x^2 \tilde{R}_z^2 \tilde{K}_x^2 \tilde{K}_z^2 = 1, \quad (4)$$

the equations of motion for the Thomas-Fermi radii read

$$\frac{1}{\omega_x^2} \frac{d^2 \tilde{R}_x}{dt^2} = -\tilde{R}_x + \frac{2\tilde{K}_x^2 + \tilde{K}_z^2}{3\tilde{R}_x} + \epsilon_{\text{dd}} A(\tilde{R}_x, \tilde{R}_z, \tilde{K}_x, \tilde{K}_z), \quad (5)$$

$$\frac{1}{\omega_z^2} \frac{d^2 \tilde{R}_z}{dt^2} = -\tilde{R}_z + \frac{2\tilde{K}_x^2 + \tilde{K}_z^2}{3\tilde{R}_z} + \epsilon_{\text{dd}} B(\tilde{R}_x, \tilde{R}_z, \tilde{K}_x, \tilde{K}_z). \quad (6)$$

Here we use a tilde to represent the quantity expressed in units of the noninteracting Thomas-Fermi radius $R_i^{(0)} = \sqrt{2E_F/M\omega_i^2}$ and the Fermi momentum $K_F = \sqrt{2E_F/\hbar^2}$ with the Fermi energy $E_F = (6N)^{1/3}\hbar\bar{\omega}$. The auxiliary functions are defined according to

$$A = -\frac{c_d}{\tilde{R}_x^3 \tilde{R}_z} \left[1 - \frac{3\tilde{R}_x^2 \lambda^2 f_s(\tilde{R}_x \lambda / \tilde{R}_z)}{2(\tilde{R}_z^2 - \tilde{R}_x^2 \lambda^2)} - f_s\left(\frac{\tilde{K}_z}{\tilde{K}_x}\right) \right],$$

$$B = -\frac{c_d}{\tilde{R}_x^2 \tilde{R}_z^2} \left[-2 + \frac{3\tilde{R}_z^2 f_s(\tilde{R}_x \lambda / \tilde{R}_z)}{(\tilde{R}_z^2 - \tilde{R}_x^2 \lambda^2)} - f_s\left(\frac{\tilde{K}_z}{\tilde{K}_x}\right) \right],$$

with the numerical constant $c_d = \frac{2^{38/3}}{3^{23}/6 \cdot 5 \cdot 7 \cdot \pi^2} \approx 0.2791$. Furthermore, the anisotropy function

$$f_s(x) \equiv \begin{cases} \frac{2x^2+1}{1-x^2} - \frac{3x^2 \tanh^{-1} \sqrt{1-x^2}}{(1-x^2)^{3/2}}, & x \neq 1, \\ 0, & x = 1, \end{cases} \quad (7)$$

decreases monotonically from 1 at $x = 0$ to -2 at $x = \infty$, passing through zero at $x = 1$ [4,5]. In addition, the variational parameters are restricted to obey

$$\tilde{K}_z^2 - \tilde{K}_x^2 = \epsilon_{\text{dd}} C(\tilde{R}_x, \tilde{R}_z, \tilde{K}_x, \tilde{K}_z), \quad (8)$$

with $C = \frac{3c_d}{\tilde{R}_x^2 \tilde{R}_z} [1 - \frac{(2\tilde{K}_x^2 + \tilde{K}_z^2) f_s(\tilde{K}_z/\tilde{K}_x)}{2(\tilde{K}_z^2 - \tilde{K}_x^2)}]$. This equation can be traced back to the exchange term and shows explicitly that a nonzero ϵ_{dd} implies a deformed momentum distribution $\tilde{K}_z \neq \tilde{K}_x$ for finite \tilde{R}_x, \tilde{R}_z as first pointed out in Ref. [17].

Equations (4)–(6) and (8) govern the static as well as the dynamic properties of a polarized dipolar Fermi gas in the HD regime and represent the main result of this article. They determine the temporal evolution of both the spatial and the momentum distribution of a dipolar Fermi gas, which are directly experimentally accessible via TOF techniques. The static solutions agree precisely with the ones obtained in Refs. [17] and [18]. In Fig. 1 we present our findings for the spatial aspect ratio as a function of the dipolar strength ϵ_{dd} . The characteristic feature is that a minimal value of λ is required for stabilizing a system with a given ϵ_{dd} . Thus, for future experiments with $^{40}\text{K}^{87}\text{Rb}$ molecules in the quantum

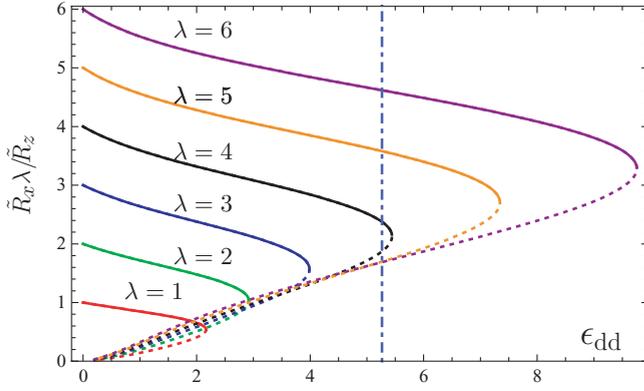


FIG. 1. (Color online) Spatial aspect ratio for different trap anisotropies λ ; the upper (solid) branches correspond to the local minimum of the mean-field energy, and the lower (dotted) branches to the maximum. Note that the critical value $\epsilon_{\text{dd}}^{\text{crit}}$, where two branches meet, decreases more slowly for lower values of λ . The dashed vertical line marks the estimated value of the interaction strength for $^{40}\text{K}^{87}\text{Rb}$ molecules, $\epsilon_{\text{dd}} \approx 5.3$.

degenerate regime, one should choose the anisotropy λ to be larger than the minimal value $\lambda_{\text{min}} \approx 3.89$ to render the system stable against collapse. Amazingly, the minimum value of λ supporting stability decreases slowly, and samples with $\lambda = 0.05$ are stable if $\epsilon_{\text{dd}} \lesssim 1.6$.

Having summarized the most important aspects of the static solutions, we turn now to their dynamical properties. In a cylinder-symmetric system the mono- and quadrupole low-lying oscillation modes couple to each other. To obtain the frequency of these modes in the HD regime, we expand the radii and momenta around their respective equilibrium values according to $\tilde{R}_i = \bar{R}_i(0) + \eta_i e^{i\Omega t}$, $\tilde{K}_i = \bar{K}_i(0) + \zeta_i e^{i\Omega t}$, where η_i (ζ_i) denotes a small oscillation amplitude in the i th direction in real (momentum) space and Ω represents the oscillation frequency. Inserting these into the equations of motion, (4)–(6) and (8), a linearization initially yields, for the ratio of the momentum amplitudes,

$$\frac{\zeta_x}{\zeta_z} = \frac{\tilde{K}_x \tilde{K}_x^2 + \tilde{K}_z^2 - \epsilon_{\text{dd}} \tilde{K}_z \partial C / \partial \tilde{K}_z}{\tilde{K}_z 2\tilde{K}_z^2 - \epsilon_{\text{dd}} \tilde{K}_z \partial C / \partial \tilde{K}_z}, \quad (9)$$

where all terms are evaluated at equilibrium. This quantity is plotted against ϵ_{dd} for $\lambda = 5$ in the lower (red) curve in Fig. 2 and is compared to the corresponding equilibrium momentum aspect ratio [upper (blue) curve]. Setting $C = 0$, that is, removing the exchange term, one has $\zeta_x = \zeta_z$, whereas for nonzero C , the ratio ζ_x/ζ_z decreases monotonically from 1 to about 0.28 in the interval $0 < \epsilon_{\text{dd}} < \epsilon_{\text{dd}}^{\text{crit}} \approx 7.34$. This shows that the exchange term induces characteristic *anisotropic breathing oscillations in momentum space*, which can be regarded as a trademark sign of the DDI in fermionic quantum gases.

Eliminating the momentum amplitudes ζ_i yields a reduced linear homogeneous system for the spatial amplitudes η_i . Demanding nontrivial solutions, we obtain an explicit but lengthy result for the monopole- (quadrupole-) oscillation frequency Ω_+ (Ω_-), which depends, via the equilibrium values of the Thomas-Fermi radii and the Fermi momenta, on the trap anharmonicity λ and the dipolar strength ϵ_{dd} . In

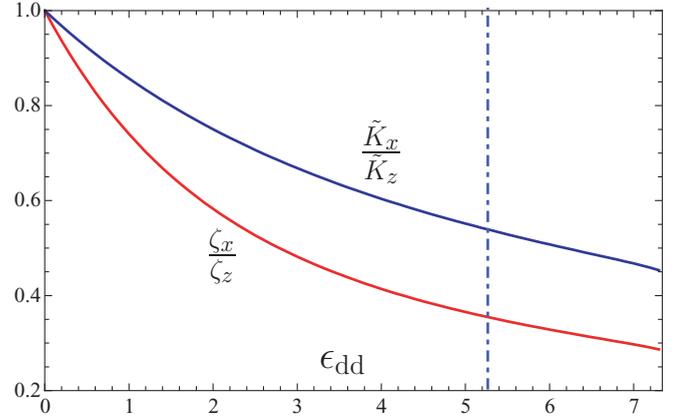


FIG. 2. (Color online) The lower (red) curve shows the ratio of the amplitudes ζ_x/ζ_z as a function of ϵ_{dd} for $\lambda = 5$. For comparison, the equilibrium aspect ratio in momentum space against ϵ_{dd} for $\lambda = 5$ is depicted by the upper (blue) curve.

the special case of an ideal Fermi gas, that is, $\epsilon_{\text{dd}} = 0$, the oscillation frequencies Ω_{\pm} reduce to the correct noninteracting values $\Omega_{\pm}^{(0)2} = \omega_x^2(5 + 4\lambda^2 \pm \sqrt{25 - 32\lambda^2 + 16\lambda^4})/3$, which were first obtained for $\lambda = 1$ in Ref. [28] and for $\lambda \neq 1$ in Ref. [29]. Figure 3 shows the oscillation frequencies of the monopole (blue) and quadrupole (red) modes plotted against ϵ_{dd} for $\lambda = 5$. As ϵ_{dd} becomes larger, we find that the monopole frequency increases and that the quadrupole frequency decreases, vanishing at $\epsilon_{\text{dd}}^{\text{crit}} \approx 7.34$, the same value for which the system becomes unstable (see Fig. 1). The inset in Fig. 3 shows how the frequencies depend on the anisotropy λ for $\epsilon_{\text{dd}} = 0.5$ (dashed) and $\epsilon_{\text{dd}} = 1.0$ (solid). It turns out that the quadrupole frequencies are larger than in the noninteracting case for $\lambda < 1$ and smaller for $\lambda > 1$, while the contrary is true

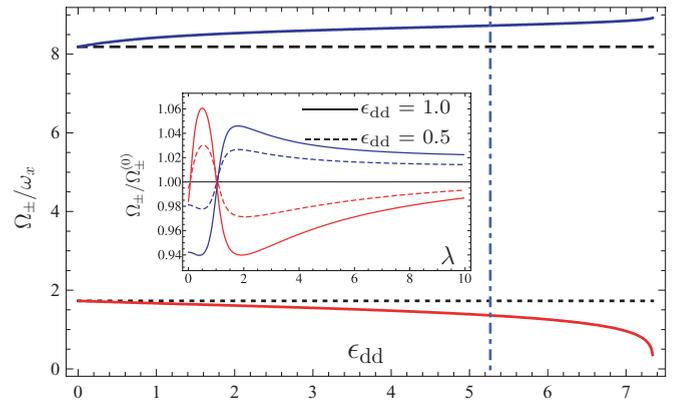


FIG. 3. (Color online) Excitation frequencies for $\lambda = 5$ as functions of the DDI strength ϵ_{dd} . The upper solid (blue) and lower solid (red) curves represent the monopole (Ω_+) and quadrupole (Ω_-) frequencies. The dashed (dotted) horizontal line represents the monopole (quadrupole) frequency of the noninteracting gas from Ref. [29]. Inset: Monopole-oscillation (blue) and quadrupole-oscillation (red) frequencies of the dipolar Fermi gas normalized by the noninteracting values from Ref. [29] against the trap aspect ratio λ for different values of the dipolar strength ϵ_{dd} . The dashed (solid) curves are for $\epsilon_{\text{dd}} = 0.5$ ($\epsilon_{\text{dd}} = 1.0$).

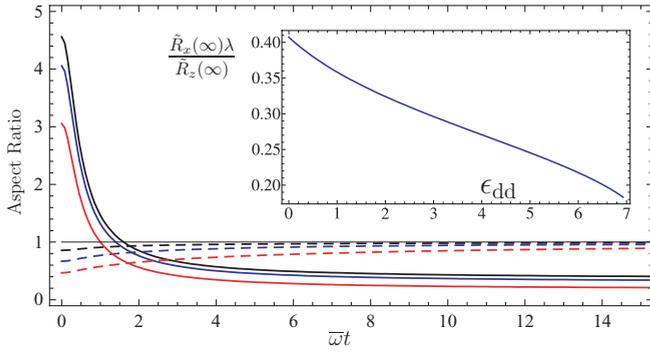


FIG. 4. (Color online) Cloud aspect ratio in TOF expansion for $\lambda = 5$ with $\epsilon_{dd} = 1, 3,$ and 7 (solid curves, top to bottom). The dashed curves depict the corresponding momentum aspect ratios. Inset: Asymptotic cloud aspect ratio against ϵ_{dd} .

for the monopole modes. This behavior agrees qualitatively with dipolar BECs [4].

It remains to study the TOF expansion of a dipolar Fermi gas. This is done by numerically solving Eqs. (4)–(6) and (8), while removing the trap frequencies. The results are presented in Fig. 4, where the spatial and momentum aspect ratios are plotted as functions of time in units of $\bar{\omega}^{-1}$ at $\lambda = 5$ for different ϵ_{dd} . The characteristic of the hydrodynamic regime

is that the asymptotic value of the aspect ratio in real space depends on ϵ_{dd} , while local equilibrium renders the momentum distribution asymptotically isotropic. We can estimate the validity of these results if we assume the previous HD criterion to be valid also during the expansion. Since the equations of motion imply $d^2\tilde{R}_i/dt^2 = 0$ for large times, yielding $(\tilde{R}_x^2\tilde{R}_z)^{1/3} \sim \bar{\omega}t$, one obtains a HD expansion provided that $(\bar{\omega}t)^2 \cdot \bar{\omega}\tau_R \ll 1$. For KRB molecules, the expansion is HD only for small times $\bar{\omega}t \ll 10$, whereas for molecules like LiCs, with $d \approx 5.5$ Debye, the expansion is HD for $\bar{\omega}t \ll 1000$.

In the present article we have investigated both low-lying oscillation frequencies and TOF expansion data for a polarized dipolar Fermi gas through a hydrodynamic approach. Our findings have revealed different fingerprints of a strong DDI. We have estimated the validity of our results and found strong evidence for hydrodynamic behavior also in the absence of superfluidity. The prospects for observing normal dipolar hydrodynamics in the quantum degenerate regime are enhanced by tight traps and the recently obtained large dipole moments.

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