

Universal three-body physics for fermionic dipoles

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A study of the universal physics for three oriented fermionic dipoles in the hyperspherical adiabatic representation predicts a single long-lived three-dipole state, which exists in only one three-body symmetry, should form near a two-dipole resonance. Our analysis reveals the spatial configuration of the universal state, and the scaling of its binding energy and lifetime with the strength of the dipolar interaction. In addition, three-body recombination of fermionic dipoles is found to be important even at ultracold energies. An additional finding is that an effective long-range repulsion arises between a dipole and a dipolar dimer that is tunable via dipolar interactions.

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The recent advances in producing ground-state ultracold dipolar molecules [1] have sparked great interest in the novel phases that could be experimentally accessible [2]. With the ability to control the dipolar interaction by an external electric field [3], studies of dipolar molecules could stimulate the discovery of intriguing phenomena across a broad range of disciplines, from condensed matter physics to ultracold chemistry. In particular, the effects of the long-range anisotropic dipolar interaction on ultracold physics has already been studied for three bosonic dipoles [4, 5]. A recent development has shown [5] that the Efimov effect [6] is more favorable for three bosonic dipoles although the dipolar interaction has ingredients that could conceivably have destroyed it.

Stimulated by their ability to mimic a variety of exotic physical systems from superfluids to neutron stars, quantum degenerate fermi gases with nondipolar interactions have been studied extensively in recent years [7]. In contrast to bosonic gases whose lifetime is mainly determined by three-body recombination [8], fermionic gases exhibit extraordinary stability against three-body inelastic collisions except when the two-body interaction is tuned near a p -wave resonance [9, 10]. Moreover, three-body physics for identical fermions is expected to be nonuniversal, i.e., short-range-dependent [9, 11]. Consequently, three-body physics for identical fermions have to some extent been regarded as less exciting and/or difficult to observe. Nevertheless, the influence of dipolar interactions on the universal behavior of bosons triggers new interest in the study of universality for fermionic dipoles, and it is becoming increasingly important to understand such physics in order to guide ongoing experiments on ground-state fermionic dipolar molecules [3].

In this Letter, the behavior of three identical fermionic dipoles is demonstrated to be universal, thus differing fundamentally from the nondipolar cases. In particular, near a dipole-dipole resonance (i.e., where the two-dipole binding energy $E_{2d} \rightarrow 0$), a single long-lived, universal three-dipole state is formed. Its size and lifetime both grow with stronger dipolar interactions. The stability of the fermionic dipolar gases is assessed through numerical calculations of the rate coefficient K_3 for three-dipole

recombination, $D + D + D \rightarrow D_2 + D$, which is the dominant expected loss mechanism for nonreactive ground-state dipolar molecules [12]. In fact, we have found that K_3 grows rapidly with the strength of the dipolar interactions, and it is strongly enhanced near a dipole-dipole resonance. This study also shows evidence for a long-range, dipolar interaction-dependent repulsion that suppresses $D_2 + D$ relaxation collisions when D_2 is deeply-bound.

Consider the three-dipole Schrödinger equation in hyperspherical coordinates (in atomic units):

$$\left(-\frac{1}{2\mu} \frac{\partial^2}{\partial R^2} + H_{ad}\right) \psi = E\psi. \quad (1)$$

where $\psi = R^{5/2}\Psi$ is the scaled three-body wavefunction, $\mu = m/\sqrt{3}$ is the three-body reduced mass for identical particles with mass m . The adiabatic H_{ad} is

$$H_{ad} = \frac{\Lambda^2(\Omega)}{2\mu R^2} + \frac{15}{8\mu R^2} + V, \quad (2)$$

where Λ^2 is the usual grand angular momentum operator [13], and Ω are the hyperangles representing positions of the three particles at fixed hyperradius R . The potential energy $V = v(\vec{r}_{12}) + v(\vec{r}_{23}) + v(\vec{r}_{23})$ is a pairwise sum of two-dipole interactions:

$$v(\vec{r}) = V_0 \text{sech}^2(r/r_0) + \frac{2d_\ell}{m} \frac{1 - 3(\hat{z} \cdot \hat{r})^2}{r^3} f(r). \quad (3)$$

The sech^2 term is isotropic and short-range, and it encapsulates the complicated interactions between two polar molecules. The second term with a short-range cutoff $f(r)$ is the interaction between two dipoles aligned in the \hat{z} direction. The dipole length is defined in terms of the electric dipole moment d_m as $d_\ell = md_m^2/2$. The universality of our calculations is tested by varying both d_ℓ and V_0 .

Equation (1) can be routinely solved, either using the adiabatic representation [13] or else through slow variable discretization (SVD) [14]. Our study shows it is advantageous to use the adiabatic scheme to treat the asymptotic behavior of three dipoles in scattering calculations, while

SVD efficiently handles sharp avoided crossings at small distances. In both cases, the biggest challenge is to solve the adiabatic eigenvalue equation:

$$H_{ad}\Phi_{\nu}^{\Pi,M}(R;\Omega) = U_{\nu}(R)\Phi_{\nu}^{\Pi,M}(R;\Omega), \quad (4)$$

where $U_{\nu}(R)$ is the adiabatic potential for channel ν , M is the space-fixed frame projection of the total angular momentum J , and $\Phi_{\nu}^{\Pi,M}(R;\Omega)$ is the corresponding eigenfunction. For interacting oriented dipoles, the major difficulty in solving Eq. (4) is that the total orbital angular momentum J is not conserved. The method of Ref. [5] is implemented to solve Eq. (4). Briefly, $\Phi_{\nu}^{\Pi,M}$ is expanded in terms of the Wigner D functions [see Eq. (4) in Ref. [5]], truncated at $J_{\max} = 13$. Tests of this truncation confirm that it yields adequate convergence.

The nature of three-dipole states near a dipole-dipole resonance emerges from study of the adiabatic potentials for $M^{\Pi} = 0^{+}$. Note that the even and the odd J values decouple in this symmetry. This symmetry with only odd J is of particular interest because it includes the $J^{\Pi} = 1^{+}$, the least repulsive partial wave for three noninteracting identical fermions [16]. One would also expect the binding of three dipoles is most likely to occur at $m_{2d} = 0$ dipole-dipole resonance, where m_{2d} is the projection of two-dipole angular momentum along the field direction. However, $m_{2d} = 0$ is not allowed for $M^{\Pi} = 0^{+}$ symmetry, so the three-dipole system is considered instead near the $m_{2d} = 1$ resonances. Figure 1 shows the typical behavior of the adiabatic potentials. Asymptotically, the three-body continuum potentials behave in the same manner as the nondipolar interaction case, but for dipole plus dipolar dimer channels, each threshold has a family of potential curves whose centrifugal barriers carry different effective angular momenta due to the J -coupling.

Irregardless of the complicated topology of the adiabatic potentials shown in Fig. 1, near a dipole-dipole resonance it is possible to trace a diabatic potential well in the channel closest to the three-body continuum. It cuts through multiple sharp avoided crossings with deeply-bound channels, as is visible in the inset of Fig. 1. Remarkably, all these potentials for different d_{ℓ} and for different short-range potentials fall on top of each other after a proper scaling. A key implication of the scaling of the universal potentials is that the hyperradius of the repulsive barrier increases with d_{ℓ} , which suppresses decay of the three-dipole state and increases its lifetime as the dipolar interaction is increased. To quantify the number of three-dipole states and their energies in the universal potential wells, we have solved Eq. (1) by using SVD. The energy E_{3d} and the width Γ of the three-body states are revealed by adding complex absorbing potentials in the lower decaying channels. Interestingly, exactly one quasi-bound state is supported by each of the universal potentials. Table I lists E_{3d} and Γ for a few values of d_{ℓ} . The energy E_{3d} shown in Table I suggests

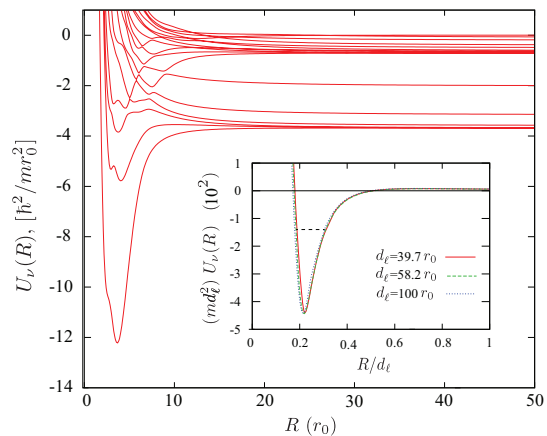


FIG. 1: (color online) A typical set of adiabatic hyper-spherical potentials $U_{\nu}(R)$ for three fermionic dipoles with $d_{\ell}/r_0 = 58.2$ and $E_{2d} \rightarrow 0$. The inset shows rescaled, diabatic potentials exhibiting universal behavior for a few values of d_{ℓ} at a dipole-dipole resonance. The horizontal dashed line in the diabatic potential wells indicates the position of the universal three-dipole states.

$E_{2d} \rightarrow 0$	$d_{\ell} (r_0)$	$md_{\ell}^2 E_{3d}$	$md_{\ell}^2 \Gamma$
	39.7	171	42
	58.2	135	43
	100	139	17
$\Theta \approx 16^{\circ}$	$b_l \approx 0.26d_{\ell}$	$b_s \approx 0.14d_{\ell}$	$\Delta \approx 15^{\circ}$

TABLE I: The upper rows: the energies E_{3d} and the widths Γ of the universal three-dipole states for different values of d_{ℓ} . The bottom row: the angle Θ between the three-dipole triangle and field direction, the long and short bond length b_l and b_s , and the smaller bond angle Δ from the most probable configuration of the three-dipole states.

a universal trend, but the width Γ exhibits less universal behavior, presumably due to the nonuniversal couplings to deeply-bound channels. Nevertheless, the $1/d_{\ell}^2$ suppression in the width implies an increased lifetime of the universal three-dipole states as d_{ℓ} increases. Our numerical study also shows that the universal three-dipole states slowly become unbound or deeply-bound as the system shifts away from a dipole-dipole resonance. In a broad region near a dipole-dipole resonance, the properties of the three-dipole states are essentially unchanged.

Importantly, the interactions of the aligned dipoles break the overall rotational symmetry, reflecting that the three-dipole states have a preferential orientation in space. The geometry of these universal states is analyzed by calculating the spatial distribution of the dipoles using the three-body wavefunctions. Specifically, the probability density for finding a dipole at \vec{r} , is

$$\rho(\vec{r}) = \frac{1}{3} \langle \Psi | \sum_i \delta(\vec{r} - \vec{r}_i) | \Psi \rangle, \quad (5)$$

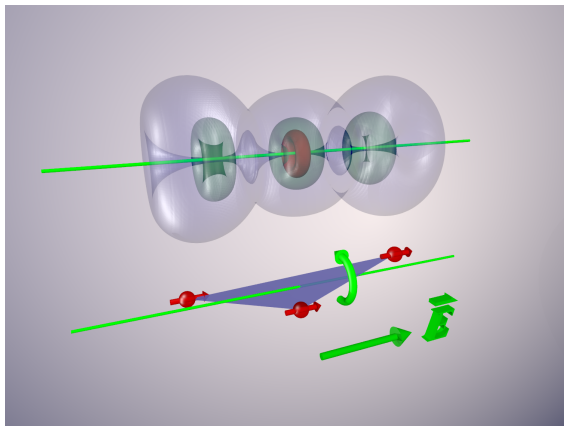


FIG. 2: (color online) The upper part: the isosurfaces of the one-dipole probability density $\rho(\vec{r})$ for the universal three-dipole states, centered in the space-fixed center of mass frame. By normalizing the peak density to be unity, the density of the following surfaces: the red (innermost), three green (middle), and the light blue (outermost) are 0.8, 0.3 and 0.015, respectively. This crankshaft-resembling structure is a figure of rotation about the field axis. The lower part: the most probable orientation and geometry of the triangle formed by three dipoles. The green axes show the direction of the external electric field.

where $\delta(\vec{r})$ is the Dirac delta function, \vec{r}_i is the position vector for the i 'th dipole. Figure 2 shows the isosurfaces of $\rho(\vec{r})$. The three-dipole states have a strongly preferential spatial configuration, as indicated by the triangle in the lower part of the figure. The isosurfaces can be viewed as the revolution of the triangle along the field axis. It is also interesting to note that the configuration of the three-dipole states are universal, only their overall size scales linearly with d_ℓ .

The configuration of a three-dipole state shown in Fig. 2 can be understood in terms of the anisotropic nature of the dipolar interactions and the symmetry of the state. The three dipoles tend to align in a line where the dipolar interactions in Eq. (3) are the most attractive, but this is forbidden by the odd partial wave symmetry. The state therefore has the configuration of an obtuse, isosceles triangle making a small angle with the field direction. The lower part of Table I lists the angle between the plane of the triangle and the field direction, the short and long bond lengths, and the smaller bond angle determined from the most probable and also the dominant configuration of the three-dipole states.

The possibility of universal three-dipole states for other symmetries have been explored, in particular the three-dipole system for $M^\Pi = 0^-$ symmetry with odd J 's, which includes the less repulsive partial wave contribution $J^\Pi = 1^-$ [16]. However, no universal three-dipole states have been found near both $m_{2d} = 0$ and $m_{2d} = 1$ dipole-dipole resonances. The situation is similar for $M^\Pi = 1^-$ and 1^+ symmetries, where the adiabatic po-

tentials are found to be more repulsive. The potentials for higher M near higher m_{2d} resonances are expected to be more repulsive, suggesting the conclusion that the $M^\Pi = 0^+$ three-dipole states presented above are the only class of universal states for identical fermions.

The universal three-dipole states affect scattering processes. As d_ℓ increases, a resonance in three-dipole recombination is expected whenever a three-dipole state passes through the three-body breakup threshold before a new two-dipole state becomes bound. To determine the stability of nonreactive fermionic dipolar gases, however, it is more important to know the general scaling of three-dipole recombination with d_ℓ and the behavior near a dipole-dipole resonance. The threshold behavior of the recombination rate K_3 for three fermionic dipoles can be obtained from a Wigner law analysis [16] of the $M^\Pi = 0^+$, J -odd low-energy-dominant contribution, giving

$$K_3 = \frac{C_3}{\mu} k^4, \quad (6)$$

where $k = \sqrt{2\mu E}$ is the three-body wavenumber.

For short-range, nondipolar interactions, the coefficient C_3 in recombination rate is expected to scale with the scattering volume [9, 10] and the effective range [10]. For dipolar interactions, however, the scattering physics is fundamentally different even in the two-body level [17]. In particular, we have numerically identified a distinct low-energy expansion of the p -wave phase shift δ for two fermionic dipoles. The real part of δ can be expanded as

$$\text{Re}[\delta(k_2)] \approx -ak_2 - bk_2^2 - Vk_2^3 \quad (k_2 \rightarrow 0), \quad (7)$$

where k_2 is the relative two-dipole wavenumber. This expansion has a linear term in k_2 implying a nonvanishing scattering length a [17] as expected for r^{-3} interaction and also a quadratic dependence on k_2 , both of which are absent for nondipolar interactions. But similar to the nondipolar interaction case, the scattering volume V goes through a pole across a dipole-dipole resonance.

Our scaling laws for three-dipole recombination are based on a calculation of the three-dipole scattering matrix using the eigenchannel R -matrix propagation method [18]. First, SVD solution of Eq. (1) obtains the R -matrix out to a distance that includes all the sharp-avoided crossings. The R -matrix is then propagated farther in the adiabatic representation [13] in the asymptotic region until K_3 is fully converged. K_3 has been calculated for $V > 0$ near a dipole-dipole resonance, where the dipolar dimer is weakly bound. At ultracold energies, our calculation shows that only the lowest three-body continuum channel is important for recombination, and K_3 is dominated by recombination into the weakly-bound dipole plus dipolar dimer channel having the lowest effective angular momentum barrier asymptotically. Figure 3 shows the scaling coefficient C_3 near $m_{2d} = 1$

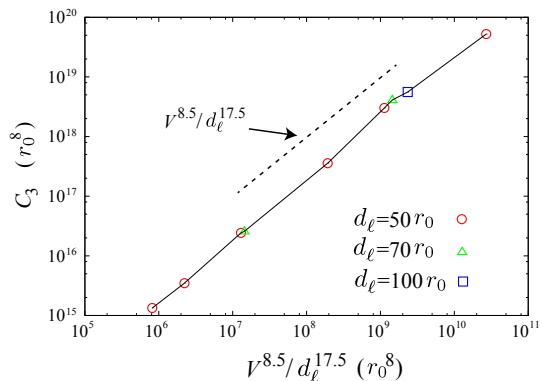


FIG. 3: (color online) The dependence of the recombination scaling coefficient C_3 on the two-body scattering volume V and the dipole length d_ℓ near a dipole-dipole resonance. The lines connecting the numerical points just guide the eye.

dipole-dipole resonances for a few d_l with different short-range interactions. The dipole-dipole resonances with higher m_{2d} are not of current interest because of their extremely narrow widths [17]. Dimensional analysis has suggested, and our numerical results in Fig. 3 confirm, that a universal scaling of C_3 exists

$$C_3 = \lambda V^{17/2} d_\ell^{-35/2}, \quad (8)$$

where dimensionless $\lambda \approx 2 \times 10^9$. This scaling is stronger with V than for nondipolar interactions [9, 10]. Although Eq. (8) also seems to imply a strong suppression of three-dipole recombination with d_ℓ , instead a general scaling of $C_3 \propto d_\ell^8$ is expected away from a dipole-dipole resonance, due to the d_ℓ^3 scaling of the background value of V [19]. This rapid growth of C_3 with d_ℓ may be unfortunate for experiments that require stable fermionic dipolar gas in three dimensions, since three-dipole recombination can be dramatic even in the ultracold threshold regime ($kd_\ell \ll 1$). On the other hand, scenarios exist where three-body recombination has a positive role, as a means to produce weakly-bound dipolar dimers that are stable against collision with a free dipole, in close analogy to the technique used in ultracold spin-mixed fermions with nondipolar interactions [20].

Finally, consider the behavior of deeply-bound dipole plus dipolar dimer channels and their impact on the stability of a mixed gas containing both dipoles and dipolar dimers. Similar to the bosonic dipole case [5], these channels have a centrifugal barrier characterized by a non-integer effective angular momentum quantum number, and it grows with increasing d_ℓ . Although our numerical study shows that this d_ℓ -dependence is nonuniversal, the effective angular momentum barrier can nevertheless “protect” a mixture of dipoles and dipolar dimers from collisional decay, and it appears to provide an alternative way to tune interactions in a mixed-species gas.

In summary, a class of universal physics has been identified for three fermionic dipoles. For scattering volumes

near a dipole-dipole resonance, one universal three-dipole state exists at an energy near the three-dipole breakup threshold. This state has a universal geometry and orientation in the external field, with a variable size dependent on d_l . Moreover, its long lifetime can be beneficial for experimental observation and manipulation. Our study also shows a universal trend of the three-dipole recombination rate, which increases rapidly near each dipole-dipole resonance, and grows rapidly with stronger dipolar interactions. And the effective repulsion between a dipole and a dipolar dimer can be tuned to control collisions in mixed quantum gases.

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