## Scale Invariance of a Spherical Unitary Fermi Gas

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A unitary Fermi gas in an isotropic harmonic trap is predicted to show scale and conformal symmetry that have important consequences in its thermodynamic and dynamical properties. By experimentally realizing a unitary Fermi gas in an isotropic harmonic trap, we demonstrate its universal expansion dynamics along each direction and at different temperatures. We show that as a consequence of SO(2,1) symmetry, the measured release energy is equal to that of the trapping energy. We further observe the breathing mode with an oscillation frequency twice the trapping frequency and a small damping rate, providing the evidence of SO(2,1) symmetry. In addition, away from resonance when scale invariance is broken, we determine the effective exponent  $\gamma$  that relates the chemical potential and average density along the BEC-BCS crossover, which qualitatively agrees with the mean field predictions. This Letter opens the possibility of studying nonequilibrium dynamics in a conformal invariant system in the future.

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Strongly interacting Fermi gases are created by tuning the interaction strength between atoms of different spin states via Feshbach resonance [1,2]. The unitary Fermi gas, realized when the *s*-wave scattering length is tuned to infinity, is of special interest in various geometries including harmonic [3–7] and box traps [8–11]. It is not only strongly correlated but also an example of a scale-invariant quantum many-body system. One of the basic tools used to explore the properties of unitary Fermi gas is the expansion dynamics [12] and much insight has been obtained about the role of interactions [13–15].

The strongly interacting Fermi gas at finite temperature is described by a hydrodynamic theory [see Eq. (1)], where the transport behaviors are determined by viscosities [16– 18]. At unitarity, the bulk viscosity  $\zeta_B$  vanishes, and the friction force arise from shear viscosity  $\eta$ . For a unitary Fermi gas in an anisotropic trap studied previously, the conformal symmetry is broken and the shear viscosity plays a dominant role, which allowed its extraction from expansion dynamics [19–21]. On the other hand, for a spherical unitary Fermi gas, the transverse relative motion of the atomic cloud is absent ( $\sigma_{ii} = 0$ ), and consequently the effect of the shear viscosity can be neglected. The system without viscosity contribution would have scale invariance of the mean square cloud radius  $\langle x^2 \rangle \rightarrow \lambda^2 \langle x^2 \rangle$ under the transformation  $x \rightarrow \lambda x$ , being connected to a noninteracting gas. Contrary to the anisotropic system, the spherical unitary Fermi gas has a hidden SO(2,1) symmetry [22,23] with Hamiltonian and raising and lowering operators composing three parts of the SO(2,1) Lie algebra, which predicts the exact relations between trapping potential energy and total energy and the breathing mode with an oscillation frequency twice the trapping frequency. However, the preparation and exploration of the universal properties of a spherical unitary Fermi gas are yet to be demonstrated experimentally.

In this Letter, we produce a spherical Fermi gas in an optical dipole trap (ODT) and explore scale invariant behaviors in strongly interacting regimes. By tuning the interaction strength to unitarity, the expansion of the system shows the scale invariance along each direction and at different temperatures, which is absent in an anisotropic system. We find that the trapping potential energy equals to the half of the total energy, verifying the virial theorem at unitarity. Furthermore, we observe the breathing mode with an oscillation frequency twice the trapping frequency and a small damping rate, providing the evidence of SO(2,1)symmetry [22]. In addition, we explore expansion dynamics away from unitarity when scale invariance is broken, and measure the effective exponent  $\gamma$  of  $\mu(n) \propto n^{\gamma}$  where  $\mu$ is the chemical potential and *n* is the average density. To the best of our knowledge, this is the first experiment on the 3D ultracold quantum gases with SO(2, 1) symmetry.

The expansion of strongly interacting Fermi gases is described by the hydrodynamic theory [14,19,23]

$$\frac{d^2}{dt^2} \frac{m\langle x_i^2 \rangle}{2} = \left\langle x_i \frac{\partial U}{\partial x_i} \right\rangle_0 + \frac{1}{N} \int d^3 r [\Delta p - \Delta p_0] \\ - \frac{1}{N} \int d^3 r (\eta \sigma_{ii} + \zeta_B \sigma'), \tag{1}$$

where  $\langle x_i^2 \rangle$  represents the mean square cloud radius along the *i*th axis (i = x, y, z), *U* is the trapping potential, *t* is the expansion time, the subscript (<sub>0</sub>) denotes the initial condition in the trap at t = 0, and  $\Delta p = p - (2/3)\varepsilon$  is the scale-invariance breaking pressure, where  $\varepsilon$  is the energy density [27]. The last term on the right describes the friction forces arising from shear viscosity  $\eta$  and bulk viscosity  $\zeta_B$ . Here,  $\sigma_{ii} = 2\dot{b}_i/b_i - (2/3)\sum_j \dot{b}_j/b_j$  represents the transverse relative motion and  $\sigma' = \sum_i \dot{b}_i/b_i$  for the dilation process, where  $b_i$  denotes the expansion scale factor. In the unitary regime, both  $\Delta p$  and  $\zeta_B$  vanish [19,28–30]. The value of  $\sigma_{ii}$  depends on the geometry or symmetry of the atomic cloud. Only for a spherical gas, the relative motion is absent with  $\sigma_{ii} = 0$ , and in this case, we obtain the expansion behavior

$$\langle x_i^2 \rangle = \langle x_i^2 \rangle_0 + \frac{t^2}{m} \left\langle x_i \frac{\partial U}{\partial x_i} \right\rangle_0.$$
(2)

Equation (2) shows the ballistic expansion analogous to a noninteracting ideal gas, and the interaction is included in the *in situ* atomic cloud size  $\langle x_i^2 \rangle_0$ .

The scale-invariant expansion along each direction can be tested by determining the value

$$\tau_i^2(t) = \frac{\langle x_i^2 \rangle_t - \langle x_i^2 \rangle_0}{\langle x_i^2 \rangle_0 \omega^2},\tag{3}$$

where  $\omega_x = \omega_y = \omega_z = \omega$  is the trapping frequency. According to Eq. (2),  $\tau^2(t) = t^2$ . The ideas of using ballistic expansion and  $\tau^2(t)$  as measure of scale invariance were suggested and demonstrated experimentally in an anisotropic Fermi gas [14], where only sum of mean square radii along three axes shows the scale invariance.

We prepare a spherical Fermi gas based on our previous works [31,32]. We initially prepare a <sup>6</sup>Li atomic degenerate Fermi gas with two spin states  $|F = 1/2, m_F = \pm 1/2\rangle$  in an elongated ODT and at the Feshbach-resonance magnetic field of 834 G. The experimental setup of the isotropic trap is schematically displayed in Fig. 1(a), where some special techniques are applied. First, a magnetic field with a gradient  $B'_z = 1.05$  G/cm is applied along z axis to simultaneously compensate the gravity force of the two spin states. This is valid for <sup>6</sup>Li atoms because the hyperfine interaction is much smaller than the Zeeman shift at the applied magnetic field. Second, two elliptic optical beams with a cross-sectional aspect ratio of  $\sqrt{2}$ , propagating perpendicularly in the horizontal plane, form the isotropic trap. Under these conditions, the trapping frequency can be varied by adjusting the optical power (see Supplemental Material [23] for details). We transfer the unitary Fermi gas to the isotropic trap with an efficiency of more than 90%. After performing the evaporative cooling in the isotropic trap, we slowly increase the optical power to 3.8 W in about 75 ms.



FIG. 1. Production of a spherical Fermi gas and scale-invariant expansion along each direction. (a) Schematics of the experimental setup. Two elliptic optical beams, propagating along xand y axes, respectively, form the isotropic trap. The ratio of the beam width along z direction to that in the horizontal plane is  $\sqrt{2}$ . A pair of anti-Helmholtz coils produce a gradient magnetic field along vertical direction to compensate the gravity. Another pair of Helmholtz coils (not shown) produces a homogeneous magnetic field along vertical direction to tune the interactions. (b) Values of  $\tau_i^2(t)$  versus the expansion time t along different directions (i = x, y, z, y'). The inset shows typical atomic images taken in the vertical direction, where the expansion time is 0.5, 0.8, 1.1, 1.4, and 1.7 ms. y' direction is between x and y axes. The error bar is the standard deviation of several measurements. The black solid curve represents the scale theory  $\tau^2(t) = t^2$ .  $T/T_{\rm F} = 0.36(3)$ , where  $T_{\rm F}$  is the Fermi temperature of the noninteracting Fermi gas.

The temperature is adjusted by controlling the optical power of the evaporative cooling. The atom number is  $N = 2.9(3) \times 10^4$  and the spin polarization is less than 6%. The trapping frequencies are  $(\omega_x, \omega_y, \omega_z) = 2\pi \times [1234(6), 1165(11), 1204(3)]$  Hz, which are nearly the same along three axes [23].

We switch off the isotropic ODT and measure the cloud width versus the expansion time t at the magnetic field B = 834 G. Two laser beams with a frequency difference of 76 MHz, propagating along vertical and horizonal directions, respectively, are applied to detect two spin states. Typical atomic images during the expansion are shown in the inset of Fig. 1(b), indicating an isotropic expansion in direct contradiction to an elongated Fermi gas [2]. We use a fringe-removal algorithm [23,33] to reduce the imaging noise. The cloud radius  $\langle x_i^2 \rangle_t$  is obtained by fitting a Gaussian distribution to the atomic density profile.



FIG. 2. Scale-invariant expansion at different temperatures. (a) Atomic cloud size  $\langle x^2(t) \rangle$  versus the expansion time *t* in the unitary regime  $(a \to \infty)$ .  $T/T_F = 0.36(3)$  (black), 0.62(2) (red), and 0.81(2) (blue), respectively. For comparison, the noninteracting Fermi gas  $(a \to 0)$  is also shown with  $T/T_F = 0.91(3)$  (green). The solid curves represent the calculations of Eq. (2). The error bar is the standard deviation of several measurements. (b) Values of  $\tau^2(t)$  versus the expansion time *t* at different temperatures. The solid black curve denotes the scale theory  $\tau^2(t) = t^2$ .

In the unitary regime, the temperature is determined by analyzing the atomic density distribution, and the cloud radius in the trap  $\langle x_i^2 \rangle_0$  can be theoretically calculated [3,23,31]. Values of  $\tau^2(t)$  are calculated according to Eq. (3). As shown in Fig. 1(b), the expansion behaviors along different directions all obey the scale theory  $\tau^2(t) = t^2$ , which indicates the absence of the effect of viscosity. This scale-invariant expansion along each direction is unique for a spherical Fermi gas.

In Fig. 2, we measure the atomic expansion at different temperatures. Only expansion along the *x* axis is displayed for simplicity. Because of the finite-temperature effect, the atomic cloud size  $\langle x^2(t) \rangle$  shows an obvious difference, as shown in Fig. 2(a). System at a higher temperature has a larger *in situ* cloud radius  $\langle x_i^2 \rangle_0$ , leading to a faster expansion, which agrees well with the theoretical prediction of Eq. (2). While values of  $\tau^2(t)$  at different temperatures are consistent, all obeying the scale theory  $\tau^2(t) = t^2$  [see Fig. 2(b)]. For comparison, the expansion behavior of a noninteracting Fermi gas  $(a \to 0)$ , where *a* is the *s*-wave scattering length) is also shown. The Fermi gas in the unitary regime  $(a \to \infty)$  has the same scaled expansion behavior with that of the noninteracting Fermi gas.



FIG. 3. Verifying the virial theorem for a unitary Fermi gas. The trapping potential U shows a linear dependence on the release energy  $E_{\rm rel}$ . The error bar of the trapping potential comes from the uncertainty of the trapping frequency. The error bar of the release energy from the standard deviation of several measurements is smaller than the mask size. Both energies are normalized by the Fermi energy  $E_{\rm F} = (3N)^{1/3} \hbar \omega$ , where N is the atom number and  $\omega$  is the trapping frequency. The solid line represents the relation  $U = E_{\rm rel}$ .

As predicted by SO(2,1) symmetry, the total energy should be twice the trapping potential energy [22]. This energy relation, which is called the virial theorem, also could be derived based on the equation of state and verified by measuring the trapping potential energy [13]. Here, we verify the virial theorem using the expansion method. The total energy of the trapped gas is the sum of trapping potential energy U, kinetic energy  $E_{kin}$ , and interaction energy  $E_{\text{int}}, E_{\text{tot}} = U + E_{\text{kin}} + E_{\text{int}}$ . After switching off the trapping potential (U = 0), the release energy  $E_{rel} = E_{kin} + E_{rel}$  $E_{\text{int}}$  remains constant during the expansion [34–36] and will be completely converted to the kinetic energy in the longtime expansion. So we only need to demonstrate the relation  $U = E_{\rm rel}$ . By fitting the slope of atomic cloud radius respect to the expansion time, we obtain the release energy  $E_{\rm rel} = (3/2)mv_x^2$ , where  $v_x$  is the expansion velocity along x axis. We can also determine the trapping potential energy  $U = (3/2)m\omega^2 \langle x^2 \rangle_0$ , which varies with atomic temperature. The experimental results are shown in Fig. 3, where the atomic temperature changes across the Fermi degeneracy. The virial theorem is valid over a wide range of energies.

Next we will study the breathing mode, demonstrating SO(2,1) symmetry of the system at unitarity. To excite the breathing mode, we sinusoidally modulate the optical field at twice the trapping frequency for four periods, making the atomic cloud sizes oscillate in phase along three axes. After different holding times *t* in the trap, the atomic cloud is imaged with a time of flight of 1 ms. The breathing mode oscillation is defined as

$$A_B = \frac{\sum_i \langle x_i^2 \rangle(t)}{\langle \sum_i \langle x_i^2 \rangle(t) \rangle} - 1, \qquad (i = x, y, z), \qquad (4)$$



FIG. 4. Breathing oscillation as a function of the holding time *t* in the trap. Experimental data are fitted with a damped sinusoidal function (solid curve). The oscillation frequency is  $\omega_B = 2\pi \times 1142(2)$  Hz  $\approx 2\omega_0$  and the damping rate is  $\Gamma_B = 38(8)$  s<sup>-1</sup>. Error bar is the standard deviation of several measurements. Here, the temperature is  $T/T_F = 0.29(1)$  and the mean trapping frequency is  $\omega_0 = 2\pi \times 583$  Hz.

where  $\langle x_i^2 \rangle(t)$  is the mean square cloud radius and  $\langle \sum_i \langle x_i^2 \rangle(t) \rangle$  is the average value of all data. As shown in Fig. 4, the oscillation is fitted with a damped sinusoidal function  $A_B = \delta A_B \sin(\omega_B t + \phi) \exp(-\Gamma_B t)$ . Here, the mean trapping frequency along three axes is  $\omega_0 = 2\pi \times 583$  Hz. The oscillation frequency is  $\omega_B = 2\pi \times 1142(2)$  Hz  $\approx 2\omega_0$  and the damping rate is  $\Gamma_B = 38(8)$  s<sup>-1</sup>. The normalized damping rate to the oscillation frequency is very small, i.e.,  $\Gamma_B / \omega_B \approx 0.005(1)$ . Observation of the breathing mode with an oscillation frequency twice the trapping frequency and a small damping rate provides a direct evidence of SO(2,1) symmetry [22]. The SO(2,1) symmetry in a 2D quantum gas has also been demonstrated in a similar way [37–39].

For comparison, we measure the breathing mode away from the unitarity. On the BEC side  $(1/k_Fa = 0.57)$ , the oscillation frequency is  $\omega_B \approx 2.08\omega_0$  and the normalized damping rate is  $\Gamma_B/\omega_B \approx 0.009(1)$ . On the BCS side  $(1/k_Fa = -0.84)$ , the oscillation frequency is  $\omega_B \approx 1.94\omega_0$ and the normalized damping rate is  $\Gamma_B/\omega_B \approx 0.006(1)$ . Away from the unitarity, the ratio of the oscillation frequency to the trapping frequency is not equal to 2 and the damping rate increases more or less. The difference from the unitarity is large on the BEC side and small on the BCS side, which is similar to the measurements of the free expansion (see Fig. 5).

Away from the resonance with  $\Delta p \neq 0$ , the scale invariance will be broken. We assume a power-law dependence of the chemical potential,  $\mu(n) = n^{\gamma}$ , where *n* is the average atomic density and  $\gamma$  is the effective exponent [40–42]. By imposing that the total energy variation vanishes to first order, one gets the energy relation in the BEC-BCS crossover [41],  $3\gamma E_{rel} = 2U$ . Considering that the bulk viscosity is negligibly small [14] and the effect of the shear viscosity in a spherical system is zero, we obtain the expansion scale factors [40]

$$\ddot{b}_i - (\omega^2/b_i)\Gamma^{-\gamma} = 0, \tag{5}$$



FIG. 5. Scale-invariance breaking in the BEC-BCS crossover. (a) Values of  $\tau^2(t)$  versus the expansion time *t* at different interactions. The expansion behaviors are represented for BEC ( $1/k_Fa = 1.1$ , red squares), unitary ( $1/k_Fa = 0$ , black circles), and BCS ( $1/k_Fa = -0.68$ , blue diamonds) regions, respectively. The solid curves denote the calculations of Eq. (6) with experimentally obtained  $\gamma$ . In the unitary regime,  $T/T_F = 0.21(2)$ . (b) Values of  $\gamma$  in the BEC-BCS crossover. The solid curve depicts the mean field calculation at zero temperature as in Ref. [40]. The dashed line denotes  $\gamma = 1$  valid deeply in the BEC regime, and the dot-dashed line denotes  $\gamma = 2/3$  valid at unitarity and deeply in the BCS regime.

where  $\Gamma = b_i^3$ . Equation (5) is a decoupled equation for each direction, where  $b_i$  can be calculated if one knows the value of  $\gamma$ . In the unitary regime with  $\gamma = 2/3$ , Eq. (5) has an analytical solution the same as Eq. (2). In the BEC-BCS crossover,  $\tau^2(t)$  can be calculated as

$$\tau^{2}(t) = \frac{b_{i}^{2} - 1}{\omega^{2}}.$$
 (6)

To measure  $\gamma$  at different interactions, we adiabatically ramp the magnetic field from 834 G to the desired value in 300 ms. As the power-law dependence of the chemical potential is valid at zero temperature, we perform the experiment at a low temperature  $T/T_{\rm F} = 0.21(2)$ . The value of  $\gamma$  at zero temperature [40] is initially input into Eq. (5) to calculate  $b_x(t)$ . Then we can obtain the cloud radius in the trap from the expansion data,  $\langle x^2 \rangle_0 =$  $\langle x(t)^2 \rangle_t / b(t)_x^2$ , to determine the trapping potential energy U. We also measure the release energy  $E_{\rm rel}$  from the long-time expansion. Using the energy relation, we obtain a new value of  $\gamma$  and input it into Eq. (5) for the next iterative calculation. We repeat the calculation until  $|\gamma_{i+1} - \gamma_i|/\gamma_i \le 10^{-5}$ , where *i* denotes the number of iterations. The obtained  $\gamma$  in the BEC-BCS crossover is shown in Fig. 5(b). In the unitary regime,  $\gamma \approx 2/3$ . On the BEC side,  $\gamma$  increases toward the molecule condensate limit with  $\gamma = 1$ . On the BCS side,  $\gamma$  decreases to some extent. The experimental measurements have the same variation trend with the mean field calculation at zero temperature [40]. The value of  $\gamma$  in the unitary regime does not change with temperature. But due to the finite-temperature effect,  $\gamma$ is smaller than the zero-temperature calculation on the BEC side, and on the BCS side it is larger. This can be reasonably understood that, as temperature increases,  $\gamma$ will change toward the thermal gas with  $\gamma = 2/3$ .  $\gamma$  could also be obtained by measuring the equation of state [43,44] and collective-mode oscillation [45].

With obtained  $\gamma$ , we can determine the cloud radius  $\langle x^2 \rangle_0 = \gamma E_{\rm rel}/m\omega^2$ . Three expansion behaviors in the BEC-BCS crossover are shown in Fig. 5(a), displaying the obvious deviation from that in the unitary regime  $(1/k_{\rm F}a = 0)$  when the interaction strength is tuned away from the resonance. The expansion is fast on the BCS side  $(1/k_{\rm F}a = -0.68)$  and slow on the BEC side  $(1/k_{\rm F}a = 1.1)$ , which can be well calculated with Eq. (6).

In conclusion, we observe the unique feature of the scale invariance induced by the coexistence of the spherical symmetry and unitary interaction, and demonstrate SO(2,1) symmetry of the system by observing the breathing mode. The virial theorem for the unitary Fermi gas has been verified. We also measure the effective exponent  $\gamma$  in the equation of state along the BEC-BCS crossover. The spherical unitary Fermi gas provides the platform to study geometrized quantum dynamics with SU(1,1) symmetry [46,47], nonequilibrium dynamics in the presence of conformal symmetry [48–50], and the bulk viscosity in the BEC-BCS crossover [51–53].

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