Stable spin domains in a nondegenerate ultracold gas

S. D. Graham,¹ D. Niroomand,¹ R. J. Ragan,² and J. M. McGuirk¹ ¹Department of Physics, Simon Fraser University, Burnaby, British Columbia, Canada V5A 1S6 ²Department of Physics, University of Wisconsin–La Crosse, La Crosse, Wisconsin 54601, USA

(Received 2 March 2018; published 8 May 2018)

We study the stability of two-domain spin structures in an ultracold gas of magnetically trapped ⁸⁷Rb atoms above quantum degeneracy. Adding a small effective magnetic field gradient stabilizes the domains via coherent collective spin rotation effects, despite negligibly perturbing the potential energy relative to the thermal energy. We demonstrate that domain stabilization is accomplished through decoupling the dynamics of longitudinal magnetization, which remains in time-independent domains, from transverse magnetization, which undergoes a purely transverse spin wave trapped within the domain wall. We explore the effect of temperature and density on the steady-state domains, and compare our results to a hydrodynamic solution to a quantum Boltzmann equation.

DOI: 10.1103/PhysRevA.97.051603

Atomic Bose-Einstein condensates can form stable magnetic domains, for instance, due to spin-dependent interaction energies [1,2], yet at higher temperatures the diffusion of magnetic inhomogeneities in a weakly interacting nondegenerate gas is typically a fast process. So long as external field inhomogeneities are small enough that any differential spindependent energy is much less than the thermal energy, entropy dominates and the system rapidly relaxes to a near-uniform equilibrium. However, the addition of quantum coherence can inhibit diffusion and lead to stable, time-independent magnetic states, even above quantum degeneracy.

Macroscopic collective behavior in nondegenerate gases was observed in diffusion experiments with hydrogen [3] and helium [4] that showed that connecting two reservoirs of spin-polarized gas led to unexpected nuclear magnetic resonance signals featuring coherence collapses and longlived revivals. These results were interpreted as an instabilitydriven creation of domains [5], which were the result of collective behavior induced by coherent exchange scattering in binary collisions, known as the identical spin rotation effect (ISRE) [6,7]—similar to the Leggett-Rice effect [8,9]. Theoretical analysis suggested that the presence of a small magnetic field gradient allowed the existence of steady-state spin domains, giving rise to the surprisingly long-lived echoes [10–12]. However, direct imaging of the stable domains was elusive.

Collective effects with similar origins were seen in the hydrodynamic behavior of ultracold gases, beginning with the observation of spin waves in nondegenerate trapped Rb and K atoms [13,14]. In addition to driving precessing spin currents that lead to coherent spin oscillations, exchange-driven collective behavior also drives the collapse and revival of coherence in trapped ultracold gases [15], and can even be used to prolong coherence to extreme times [16]. The diffusion of spin inhomogeneities in ultracold gases is also strongly affected by exchange-driven collective behavior, including quantum-limited diffusion in strongly interacting Fermi gases [17–19] and significant deviations from classical diffusion in a weakly interacting Bose gas [20].

Collective spin effects are particularly striking in weakly interacting gases because of the disparity of the energy scales involved. Typical thermal energies of a nondegenerate ultracold gas are $k_{\rm B}T/h \sim 15$ kHz, while mean-field interaction energies are a thousand times less. Instead, the requirement for significant collective spin effects is that the spin rotation parameter μ be large, where $\mu = \omega_{\rm ex}\tau$, for elastic collision time τ and mean-field exchange frequency $\omega_{\rm ex} = gn/\hbar$. The coupling constant $g = 4\pi \hbar^2 a/m$, with *s*-wave scattering length *a*, mass *m*, and density *n*. When μ is large, even small potential energy inhomogeneities can drive ensemble-wide dynamics, and the addition of small external effective magnetic fields can induce dramatic effects on the diffusive dynamics of such a system.

Earlier experiments in spin-polarized H and He were typically performed well into the hydrodynamic regime, where damping of small perturbations is often long, and thus spontaneously formed domain structures have time to emerge. This is not the case in the work presented here; instead, the system is initialized in a domain structure, and the stability of the state is explored. In this Rapid Communication, we report the existence of stable spin domains in a trapped nondegenerate gas of weakly interacting ⁸⁷Rb atoms. By applying a small effective magnetic field gradient, the diffusion of magnetic inhomogeneities can be dramatically slowed and even stopped. In contrast to previous results [20], which showed that the presence of a large amplitude dipolar spin wave dramatically slows longitudinal spin diffusion, this work provides experimental evidence of time-independent spin domain states that do not diffuse until spin coherence is lost.

The experimental system consists of ⁸⁷Rb atoms cooled to near degeneracy in an axisymmetric quasi-one-dimensional (1D) harmonic magnetic trap ($\omega_{z,\rho} = 2\pi \times 6.7,255$ Hz respectively). The spinor is a pseudo-spin-1/2 doublet comprising two hyperfine ground states coupled via a two-photon microwave transition at 6.8 GHz ($|1\rangle \equiv |F,m_F = 1,-1\rangle$ and $|2\rangle \equiv |2,1\rangle$). A two-domain spin structure is initialized from $|1\rangle$ using an optical masking technique, wherein a masked off-resonant laser shifts one side of the atomic distribution out of resonance for a π pulse, thereby transferring only the



FIG. 1. Spatiotemporal evolution of radially averaged M_{\parallel} for T = 650 nK and peak density $n_0 = 1.4 \times 10^{13}$ cm⁻³ at (a) G = 0, (b) 54, and (c) -19 Hz/mm. The initial domain undergoes dipole oscillations at G = 0, which increase in frequency and damping rate for G < 0. Positive gradients stabilize the initial domain-wall configuration. The domain wall drifts toward the $M_{\parallel} = 1$ domain over time, as $|2\rangle$ decays via dipolar relaxation. (d) Time evolution of the spin dipole moment $\langle zM_{\parallel} \rangle$ in different effective-field gradients. Larger G also leads to faster damping, through decoherence from the inhomogeneous applied field.

unmasked half of the distribution from $|1\rangle$ to $|2\rangle$ (see Ref. [20]). This coherent preparation maintains magnetization $|\vec{M}| = 1$ throughout the ensemble.

The effective magnetic field is created with an optical dipole potential using a frequency- and amplitude-modulated acousto-optic modulator to produce a time-averaged linear gradient in optical intensity along the trap axis [21]. A laser detuned ~3.4 GHz from the D_2 excited-state transition creates a differential energy shift that locally alters Larmor precession, analogous to a magnetic field gradient torquing real spins. The gradient size *G* is adjusted with laser intensity. In all experiments, loss from spontaneous emission is below other loss processes, such as dipolar relaxation in $|2\rangle - |2\rangle$ collisions.

Following the preparation and evolution of the spin domains, the longitudinal magnetization is measured destructively by direct measurement of the populations of each spin state, $M_{\parallel}(z,t) = N_2(z,t) - N_1(z,t)$. Earlier work showed that, with no applied external field, domain structures in a uniform differential potential still undergo trap oscillations and diffusion [Fig. 1(a)], despite a significant slowing of dynamics due to collective behavior driven by coherent spin currents [20]. Figure 1(b) shows that the addition of a small effective-field gradient (G = 54 Hz/mm) stabilizes the domains against both diffusion and trap oscillations. Classically, diffusion should occur on a timescale given by the elastic collision time τ , which at ~24 ms for the conditions shown in Fig. 1 is much too fast to explain the domain lifetimes. Nor is the domain stabilization due to any differential mechanical force. For instance, a 50 Hz/mm field gradient shifts the relative trap centers by only 0.1 μ m, less than 0.1% of the Gaussian half width of the distribution z_0 . This effect is instead driven by coherent spin interactions.

Furthermore, the sign of the effective-field gradient is greatly important. Though the transport equation [Eq. (1) below] supports steady-state solutions for both positive and negative gradients, the initial domain preparation breaks this symmetry and leads to a preferred orientation. The sign of G must oppose the initial spin gradient in the domain wall for the stable configuration to emerge. To rotate smoothly from one domain to the other as a spin traverses the domain wall, spin rotation from the external gradient must balance mean-field-induced spin torque. Otherwise, the system is initialized far from the gradient-driven stable state, and large amplitude transients consume virtually all the magnetization [Fig. 1(c)]. The domain behavior as a function of G can be summarized by dipole moment oscillations [Fig. 1(d)].

The system is described theoretically by considering a quasi-one-dimensional trapped gas with a spin-density distribution function $\vec{m}(z, p, t)$. The evolution of the magnetization in the presence of weak interactions is described by a 1D Boltzmann equation,

$$\partial_t \vec{m} + \partial_0 \vec{m} - \frac{1}{\hbar} (U_{\text{diff}} \hat{z} + g \vec{M}) \times \vec{m} = \partial_t \vec{m}|_{\text{coll}}, \qquad (1)$$

where $\partial_0 = \frac{p}{m_{\rm Rb}} \partial_z - m_{\rm Rb} \omega_z^2 z \partial_p$, for mass $m_{\rm Rb}$ and radially averaged collisional relaxation rate $\partial_t \vec{m}|_{\rm coll}$ [22]. $\vec{M}(z,t) = \int \vec{m} dp/2\pi \hbar$ is the spatial component of the spin distribution, and is the observable in these experiments. $U_{\rm diff}$ represents any differential energies experienced by the spin components and acts as an effective externally applied magnetic field.

Analysis of the kinetic equation in the presence of a linear effective magnetic field gradient $U_{\text{diff}}/\hbar = Gz$ indeed reveals the existence of steady-state solutions featuring two oppositely oriented longitudinal spin domains $(M_{\parallel} = \pm 1)$ joined by a narrow domain wall [10–12]. In the collisionless limit, Fomin derived a relation between the applied field gradient *G* and the equilibrium size of the domain wall λ_{eq} in an open normal Fermi liquid [10]. Following these predictions, Ref. [23] obtained a similar expression for a trapped gas in the hydrodynamic limit,

$$G_{\text{hydro}} = \frac{\omega_z}{z_0} \frac{1}{\mu M} \left(\frac{\pi/2}{1.1\lambda_{\text{eq}}/z_0} \right)^3 \omega_z \tau, \qquad (2)$$

which scales as T/n. To provide the spin rotation required to maintain stable domains, the domain wall must be nearly fully polarized—that is, a helical domain wall where the spin smoothly rotates from one longitudinal orientation to another and $|\vec{M}| \simeq 1$ —so that $\mu M \gg 1$.

To test these predictions, we study the dynamics of the domain wall, which provides insight into the behavior of the steady-state domains. We characterize the domain



FIG. 2. (a) Normalized distribution of atoms in $|1\rangle [N_1(z), \mathbf{\nabla}]$ and $|2\rangle [N_2(z), \mathbf{A}]$ at T = 650 nK and $n_0 = 2.8 \times 10^{13}$ cm⁻³. A Gaussian is fit to the sum (•) to extract temperature. The difference $M_{\parallel} = N_2 - N_1$ (**II**) is fit with the same Gaussian multiplied by tanh z/λ to determine domain-wall width λ . (b) Time dependence of λ immediately after application of *G* (given in Hz/mm), exhibiting linear relaxation at short times. (c) Initial domain-wall relaxation rate λ for three densities as a function of effective-field gradient *G*. Uncertainties represent a statistical uncertainty in fitting λ due to shot-to-shot temperature and density fluctuations. A linear fit is performed to determine the gradient G_0 that stabilizes the initial domain wall λ_0 .

wall with a fit to a phenomenological model, $M_{\parallel}(z,t) = \exp(-z^2/2z_0^2) \tanh z/\lambda(t)$, giving an initial domain-wall width $\lambda_0 \simeq 73(3) \mu m$ [Fig. 2(a)]. Changes in λ due to the effective-field gradient allow determination of the relationship between *G* and λ_{eq} . When the applied gradient is small, the stable domain-wall width λ_{eq} is large—much larger than λ_0 —and the domain wall relaxes until it matches λ_{eq} for that *G*. Conversely, if *G* is large, then λ shrinks. However, if *G* is chosen so that $\lambda_{eq} = \lambda_0$, no domain-wall dynamics is observed, until eventually, at long times, dephasing from the effective-field inhomogeneity removes enough coherence so that $\mu M < 1$ and classical diffusion dominates.

We use this domain-wall dynamics to study the steady-state configuration, because one cannot wait until λ has stabilized to measure λ_{eq} as a function of *G*. Significant transients are observed during relaxation to the steady state, and damping of those transients removes magnetization, as does decoherence driven by the inhomogeneous effective field. Thus, by the time equilibration occurs, $|\vec{M}| \ll 1$, which in turn changes λ_{eq} . Instead, we focus on times much shorter than these damping timescales and determine which gradient G_0 stabilizes λ_0 in a fully polarized system.

Figure 2(b) shows the initial behavior of $\lambda(t)$ for several gradients, where again $\lambda(t)$ comes from phenomenological fits to $M_{\parallel}(z,t)$. At short times, $\lambda(t)$ is approximately linear, and the rate of relaxation λ depends on the difference between λ_0 and λ_{eq} . Figure 2(c) displays λ vs *G*; the horizontal intercept



FIG. 3. Stabilizing gradient G_0 as a function of peak density and temperature for $\lambda_0 = 73 \ \mu$ m. Points at $n_0 = 1.4 \times 10^{13} \ \text{cm}^{-3}$ have been offset slightly for clarity. The data span the crossover from collisionless to hydrodynamic regimes, as indicated by the Knudsen number for $\lambda_0 = 73 \ \mu$ m. The data are described by G_{hydro} at high density for T = 650 nK (dashed) and 425 nK (dashed-dotted). Dotted lines show G_{hydro} for $\lambda_{\text{eq}} = \ell_{3\text{D}}$ (Kn = 1), demarking the region where the hydrodynamic model's prediction of a steady-state solution for some λ_0 is expected to be valid.

gives the gradient G_0 that produces the stable domain solution where $\lambda_{eq} = \lambda_0$ and $\dot{\lambda} = 0$.

Density and temperature play important roles in the relationship between λ_{eq} and G_0 (Fig. 3). At high density, the system nears the hydrodynamic regime and transport is inhibited; thus, smaller G_0 is needed to counteract spin gradients. Figure 3 shows the Knudsen number, $\text{Kn} = \ell/\lambda_0$, calculated at the domain-wall center using the mean free path ℓ and $\lambda_0 =$ 73 μ m. As Kn decreases to 1, though still in the crossover region between collisionless and hydrodynamic behavior, the 1/n dependence of G_{hydro} agrees well with measured values of G_0 . Since the atom cloud is collisionally thick at high n_0 and atoms are more localized, we use a radially averaged density in G_{hydro} , instead of an ensemble average. We note the λ_0^{-3} dependence produces a 15% uncertainty in G_{hydro} .

Furthermore, when temperature is lowered at high density, G_0 decreases. The quantitative agreement with G_{hydro} is better at T = 650 nK than at 425 nK, as assumptions contained within the model become more strained at lower temperature namely, that $\lambda_0 \ll z_0$ and $\omega_z \tau \ll 1$. The former ensures slow, or unbounded, diffusion, while the latter is necessary for a large enough collision rate to ensure local equilibrium in the spin current. However, since $\mu = 4.9-3.5$ for T = 425-850 nK, the approximation $\mu M \gg 1$ strengthens at lower T as collisions become more quantum in nature. Colder, smaller clouds do give larger uncertainties as the signal-to-noise ratio (SNR) drops; additionally, the highest density point at T = 425 nK has $T/T_c = 1.08$, and shot-to-shot fluctuations may occasionally include small condensates, which are inclined to phase separate to minimize energy in even small field gradients.

At low density, the hydrodynamic approximation breaks down, and G_0 deviates from G_{hydro} . The ultimate limits of the hydrodynamic model are shown by the dotted lines in Fig. 3, where G_{hydro} is evaluated for the largest λ_{eq} that



FIG. 4. (a) Initial transverse spin magnitude M_{\perp} as measured from Ramsey fringe amplitudes. Also shown are M_{\parallel} and $|\vec{M}|$. Minimal decoherence is expected during the short preparation sequence. Any discrepancies in $|\vec{M}| \simeq \sqrt{M_{\parallel}^2 + M_{\perp}^2}$ are attributed to noise-degrading Ramsey fringe fits. Solid lines are fits to a Gaussian $(|\vec{M}|)$, and the same Gaussian multiplied by sech z/λ (M_{\perp}) and tanh z/λ (M_{\parallel}). (b) Spatiotemporal evolution of the orientation of $\vec{M}_{\perp}(z,t)$, from the shaded region in (a). (c) The time evolution of ϕ exhibits dipolar spin-wave oscillations, highlighted for two locations in the domain wall [dotted lines in (b)].

satisfies Kn = 1 for a given n_0 and T, i.e., $\lambda_{eq} = \ell_{3D}$. Here, ℓ_{3D} uses an ensemble-averaged density that is more appropriate in the low-density limit, instead of the radially averaged Kn shown on the Fig. 3 axis. The hydrodynamic model predicts that steady-state domains exist for the parameter space to the right of the dotted lines, but does not guarantee the existence of any stable domains to the left. In fact, at low densities the measured value of the stabilizing gradient corresponds roughly to $\lambda_{eq} = \ell_{3D}$. A linearized moment-method analysis of Eq. (1) in the collisionless limit suggests that steady-state solutions do indeed exist for large domain walls ($\lambda_0 > z_0$) at small gradients, but it is not clear that narrow domain-wall preparations such as initialized here can be stabilized in the collisionless regime.

Low density complicates the experimental technique of using domain-wall relaxation to find G_0 . The second-lowest density in Fig. 3, $n_0 = 0.84 \times 10^{13}$ cm⁻³, shows unambiguous signatures of steady-state domains and behavior consistent with higher values of n_0 , but the G_0 found for $n_0 =$ 0.55×10^{13} cm⁻³ may not truly represent a stable domain solution for $\lambda_0 = 73 \ \mu$ m. Large transients are observed here, where the initial spin current lies further from its equilibrium value, and rapid dephasing in the collisionless regime renders analysis of λ less reliable in determining G_0 . It is likely that the measurement of G_0 at this density is tainted by transients and also represents an average value for 0 < M < 1 due to dephasing. Indeed, analysis of λ for very short times (t < 8 ms) and small gradients (G < 40 Hz/mm) suggests that G_0 should be several times higher; however, measurements at higher G reveal only rapid dephasing and even faster domain-wall relaxation rates. Thus it appears that steady-state solutions may not exist for narrow domain walls in the collisionless limit.

Lastly, an important component to understanding the stabilization of domains is the behavior of the transverse magnetization $\vec{M}_{\perp} = M_{\perp} e^{i\phi}$. We study the dynamics of \vec{M}_{\perp} for the conditions in Fig. 1(b) by applying a $\pi/2$ pulse after a variable delay time to produce Ramsey fringes [20]. The amplitude and phase of Ramsey fringes give the magnitude M_{\perp} and phase angle ϕ , respectively. There are several striking features in the behavior of M_{\perp} (Fig. 4). First, M_{\perp} undergoes phase oscillations, i.e., a dipolar transverse spin wave. This represents a decoupling in dynamics between a purely transverse spin wave and the stable longitudinal spin domains, as opposed to the coupled transverse-longitudinal spin dynamics observed in Ref. [20] at G = 0. The frequency of this spin wave approaches ω_z , consistent with strongly driven, highly nonlinear dipolar spin waves [21]. Second, stable longitudinal domains preclude any diffusion of transverse spin as well, and the transverse spin wave remains trapped within the domain wall.

A confined, purely transverse spin wave is necessary to support the steady-state domains. The precessing transverse spin current provides the spin rotation as atoms move across the domain wall from $M_{\parallel} = 1$ to -1. Though microscopic spins remain coupled via the ISRE, the macroscopic magnetization decouples completely. By contrast, in the absence of an effective-field gradient, the longitudinal magnetization gradient induces spin currents that determine the transverse phase gradient, which would be stable with no field gradient. However, longitudinal domains are unstable at G = 0 and undergo dipole oscillations, dragging the transverse phase with them and leading to a rapid collapse and revival of M_{\perp} [20].

In conclusion, we have demonstrated the stabilization of spin domains in a nondegenerate gas using optically induced effective magnetic field gradients. The gradients allow domain lifetimes more than 30 times the classical diffusion time. This effect is driven by quantum symmetry in microscopic atomatom interactions, and leads to decoupling of the dynamics of the longitudinal spin domains and a trapped transverse spin wave. Our experiments show good agreement with a hydrodynamic approximation at high density and suggest limitations to steady-state domains in the collisionless limit.

This project was supported by the Natural Sciences and Engineering Research Council of Canada (NSERC).

S.D.G. and D.N. contributed equally to this work.

- D. S. Hall, M. R. Matthews, J. R. Ensher, C. E. Wieman, and E. A. Cornell, Phys. Rev. Lett. 81, 1539 (1998).
- [3] B. R. Johnson, J. S. Denker, N. Bigelow, L. P. Lévy, J. H. Freed, and D. M. Lee, Phys. Rev. Lett. 52, 1508 (1984).
- [2] J. Stenger, S. Inouye, D. M. Stamper-Kurn, H.-J. Miesner, A. P. Chikkatur, and W. Ketterle, Nature (London) 396, 345 (1998).
- [4] G. Nunes, C. Jin, D. L. Hawthorne, A. M. Putnam, and D. M. Lee, Phys. Rev. B 46, 9082 (1992).
- [5] B. Castaing, Physica B 126, 212 (1984).

- [6] E. P. Bashkin, Pis'ma Zh. Eksp. Teor. Fiz. 33, 11 (1981)[JETP Lett. 33, 8 (1981)].
- [7] C. Lhuillier and F. Laloë, J. Phys. 43, 197 (1982).
- [8] A. Leggett and M. Rice, Phys. Rev. Lett. 20, 586 (1968).
- [9] J. N. Fuchs, D. M. Gangardt, and F. Laloë, Eur. Phys. J. D 25, 57 (2003).
- [10] V. V. Dmitriev and I. A. Fomin, Pis'ma Zh. Eksp. Teor. Fiz. 59, 353 (1994) [JETP Lett. 59, 378 (1994)].
- [11] R. J. Ragan and D. M. Schwarz, J. Low Temp. Phys. 109, 775 (1997).
- [12] A. E. Meyerovich, J. H. Naish, J. R. Owers-Bradley, and A. Stepaniants, Low Temp. Phys. 23, 411 (1997).
- [13] J. M. McGuirk, H. J. Lewandowski, D. M. Harber, T. Nikuni, J. E. Williams, and E. A. Cornell, Phys. Rev. Lett. 89, 090402 (2002).
- [14] X. Du, L. Luo, B. Clancy, and J. E. Thomas, Phys. Rev. Lett. 101, 150401 (2008).
- [15] J. M. McGuirk and L. F. Zajiczek, Phys. Rev. A 83, 013625 (2011).

- [16] C. Deutsch, F. Ramirez-Martinez, C. Lacroûte, F. Reinhard, T. Schneider, J. N. Fuchs, F. Piéchon, F. Laloë, J. Reichel, and P. Rosenbusch, Phys. Rev. Lett. 105, 020401 (2010).
- [17] A. Sommer, M. Ku, G. Roati, and M. W. Zwierlein, Nature (London) 472, 201 (2011).
- [18] M. Koschorreck, D. Pertot, E. Vogt, and M. Köhl, Nat. Phys. 9, 405 (2013).
- [19] S. Trotzky, S. Beattie, C. Luciuk, S. Smale, A. B. Bardon, T. Enss, E. Taylor, S. Zhang, and J. H. Thywissen, Phys. Rev. Lett. 114, 015301 (2015).
- [20] D. Niroomand, S. D. Graham, and J. M. McGuirk, Phys. Rev. Lett. 115, 075302 (2015).
- [21] J. M. McGuirk and L. F. Zajiczek, New J. Phys. 12, 103020 (2010).
- [22] T. Nikuni, J. E. Williams, and C. W. Clark, Phys. Rev. A 66, 043411 (2002).
- [23] R. Ragan and J. Baggett, J. Low Temp. Phys. 134, 369 (2004).