

Direct observation of s -wave scattering of fermions crossing a Feshbach resonance

D Genkina¹, LM Aycock^{1,2}, BK Stuhl¹, H-I Lu¹ and IB Spielman¹[‡]

¹Joint Quantum Institute, National Institute of Standards and Technology, and University of Maryland, Gaithersburg, MD, 20899 USA

²Physics Department, Cornell University, Ithaca, NY 14850 USA

E-mail: spielman@umd.edu

Abstract. We directly mapped s -wave scattering across a Feshbach resonance in ^{40}K . In this experiment, we collided a pair of degenerate clouds and directly imaged the scattered atoms. Owing to their low density, few atoms scattered, even near the resonance. To optimize signal to noise, we develop techniques to interpret absorption imaging in the regime where the optical intensity changes dramatically as light traverses the cloud, and recoil induced detuning corrections are significant. We apply these techniques to our s -wave scattering data to extract an estimate for the resonant magnetic field value. These techniques are generally applicable, and can be used to observe effective p-wave scattering in the presence of spin-orbit coupling in a spin polarized Fermi gas.

Keywords: Quantum gases, Atomic physics, Something else

1. Introduction

Feshbach resonances provide a stand-by technique for tuning the interaction strength in ultracold atomic gases. Even weak interactions play a crucial role in the physics of atomic Bose-Einstein Condensates (BECs), giving rise to, for example, their characteristic Thomas-Fermi density profiles. The effects of interactions in Fermi gases, however, are harder to observe. The density of Fermi clouds is reduced by a factor of 10^3 from that of BECs, making it necessary to enhance the strength of interactions in order to observe them. Feshbach resonances alter the inter-atomic scattering length via the magnetic field. Not only do they enable detection of interactions, but they make it possible to tune interactions from attractive to repulsive, allowing for the phase transition from the BCS to BEC regime at sufficiently low temperatures [6].

A Feshbach resonance occurs when a bound molecular state of two atoms energetically approaches the free atomic state [1]. The energy of the atomic states is defined by their hyperfine state and tunable with a bias magnetic field. The Feshbach resonance can thus be approached by changing the bias field. The effect of the resonance on the scattering length between two free atoms in the appropriate hyperfine state is given by

$$a(B) = a_{bg} \left(1 - \frac{\Delta}{B - B_0} \right), \quad (1)$$

where a is the scattering length, a_{bg} is the scattering length away from the resonance, Δ is the width of the resonance, and B_0 is the field value at which the resonance occurs. Note that the scattering length tends to infinity from either side of the resonance.

To use Feshbach resonances as a tool it is necessary to have a good measurement of the parameters in the above equation. The exact value of the resonant field value B_0 is impossible to calculate analytically and must be estimated via numerical models or determined experimentally. Some experimental techniques that have been used include observation of loss due to three-body inelastic scattering, measurements of re-thermalization timescales, or measurements of anisotropic expansion, all of which infer the elastic scattering cross section from collective behavior of the cloud [3–5].

Here we present a new technique for measuring the location and width of a Feshbach resonance. We collide two clouds of a degenerate Fermi gas and directly image the resulting s -wave scattering halo as a function magnetic field strength. This allows us to directly observe the enhancement in scattering without relying on collective behavior. We measured the fraction of atoms scattered during the collision, and from this fraction deduced the resonant magnetic field and width of the resonance.

Our Fermi gases are so dilute that even with Feshbach resonance enhancement of the scattering cross section, very few atoms will scatter as a fraction of total atom number. This makes direct detection of s -wave scattering halos difficult due to detection uncertainty, which disproportionately affects low atom numbers. In order to optimize the signal to noise for low atom numbers, we entered an imaging regime that relies on simulation for an accurate interpretation of the images. We performed these simulations

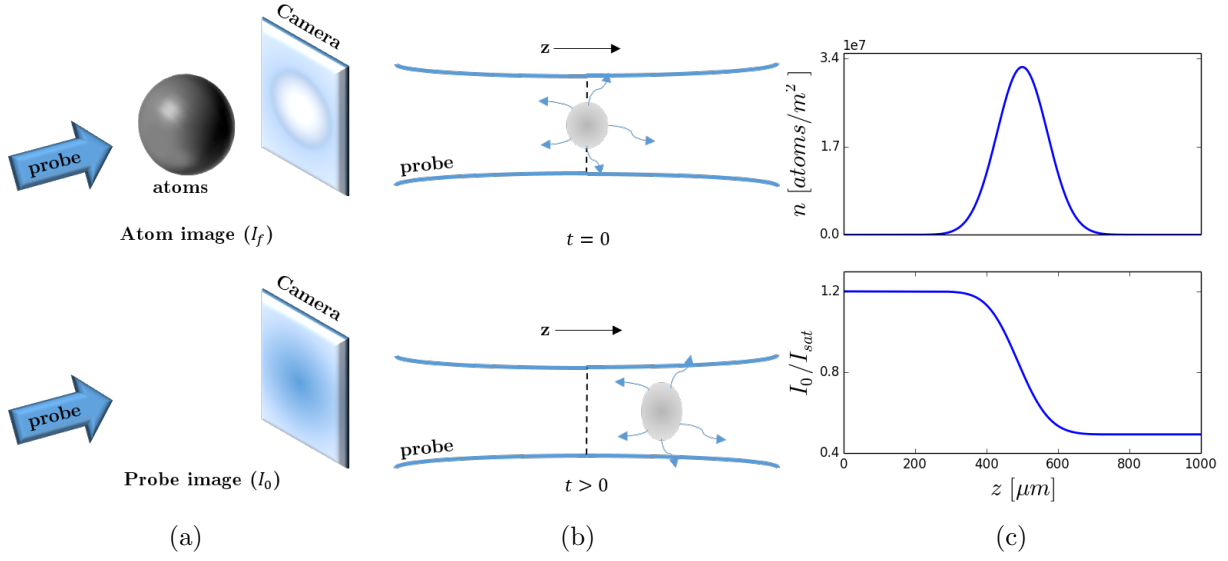


Figure 1. Absorption imaging. a. First, a near resonant probe light is shined on the atoms, and the shadow of that light is imaged on the camera. Then, the atoms are allowed to escape their trap and an image of just the probe light is taken. b. An atomic cloud illuminated by a probe light field absorbs photons from the probe and re-emits them in arbitrary directions. This process results in a net velocity of the cloud in the direction of the probe light as well as diffusive spreading in the transverse directions. c. As the probe beam travels through the atomic cloud, some of its power is absorbed, and the intensity seen by atoms further along the imaging direction z is lowered.

and used the results to extract the atom number and the scattered fraction from our images.

This paper is in two parts. In the first, we study absorption imaging in the presence of significant Doppler shift and show how we use our results to interpret data. In the second, we describe our s-wave scattering experiment and extract a measure of the location of the Feshbach resonance in ^{40}K .

2. Absorption imaging in the presence of strong recoil induced detuning

Absorption imaging is one of the two most common imaging techniques used in ultracold atomic physics, fluorescence imaging being the other. To obtain an absorption image, one shines an on or near resonant probe beam onto the atomic cloud, and captures the part of the beam that made it through the cloud onto a camera. Then, the atoms are allowed to leave the trap, and the probe light is shined directly at the camera, to calibrate the intensity of light that the atoms saw. The intensity in the second camera image at some point in time is called the initial intensity, $I_0(t)$, as it is assumed to be the intensity before interaction with atoms. The intensity in the first image at some point in time is called the final intensity, $I_f(t)$.

It is convenient to define the optical density, $\nu = -\ln \frac{\int I_f(x,y,t)dt}{\int I_0(x,y,t)dt}$. Assuming the

intensity seen by the camera during the first and second pulse is constant in time, this reduces to $\nu = -\ln \frac{I_f(x,y)}{I_0(x,y)}$. This is an observed quantity, and it is then the job of the experimentalist to relate it to the number of atoms $n = \int \rho(x,y,z) dz$, where ρ is the 3d atomic distribution, and z is the imaging axis. Another useful quantity is the optical depth, $OD = \sigma_0 \rho$, where σ_0 is the on-resonant scattering cross section. Though in general one needs a theoretical model to relate the optical density ν to the optical depth and thus the atom number, it is convenient that in the limit of zero probe intensity, these quantities coincide, $OD_0 = \nu$, as will be seen below. We call this OD_0 , because it is the simplest, in a sense "0th order" model for relating the optical density to the optical depth.

The intensity of light from the beam as a function of distance is equal to the power scattered by the atoms. Without taking the atomic recoil velocity into account, and thus assuming both the intensity at every point in the cloud to be constant in time, this is given by [7]

$$\frac{dI(z)}{dz} = -\rho\sigma_0 \frac{I(z)}{1 + I(z)/I_{sat}}, \quad (2)$$

where I_{sat} is the saturation intensity. We've suppressed the x and y dependence, or alternatively we've focused on a single pixel. This equation can be easily integrated over z to obtain

$$OD_1 = \sigma_0 n = \nu + \frac{I_0 - I_f}{I_{sat}}, \quad (3)$$

This gives us a model for the optical depth that includes a correction due to saturation intensity. In the limit of far below saturation probe intensity, it reduces to our "0th order" model, OD_0 .

Once you include the recoil velocity, the atoms become doppler shifted out of resonance and thus change the amount of probe light that will get absorbed. This detuning varies both with imaging time t and distance along the propagation direction z (Figure 2). Thus, the intensity lost to the atoms also acquires a time dependence:

$$\frac{dI(t,z)}{dz} = \sigma_0 \rho \frac{I(t,z)}{1 + \frac{4}{\Gamma^2} \delta(t,z)^2 + I(t,z)/I_{sat}}, \quad (4)$$

where Γ is the linewidth of the atomic transition, and δ is the detuning, given by

$$\delta(t,z) = \frac{v_r}{\hbar c \rho} \int_0^t \frac{dI(z,\tau)}{dz} d\tau, \quad (5)$$

where v_r is the recoil velocity. In this case, one can no longer obtain a straightforward relation between the atom number and the intensities in the two absorption images.

One can consider this equation perturbatively in time and obtain corrections to second order in imaging time, OD_2 [2]. However, the perturbative treatment breaks down after about a recoil of imaging time (Figure 3). In order to adequately correct for the recoil induced detuning of the atoms, we must simulate the process and obtain numerical predictions for I_f given a certain imaging time, atomic density, and probe intensity.

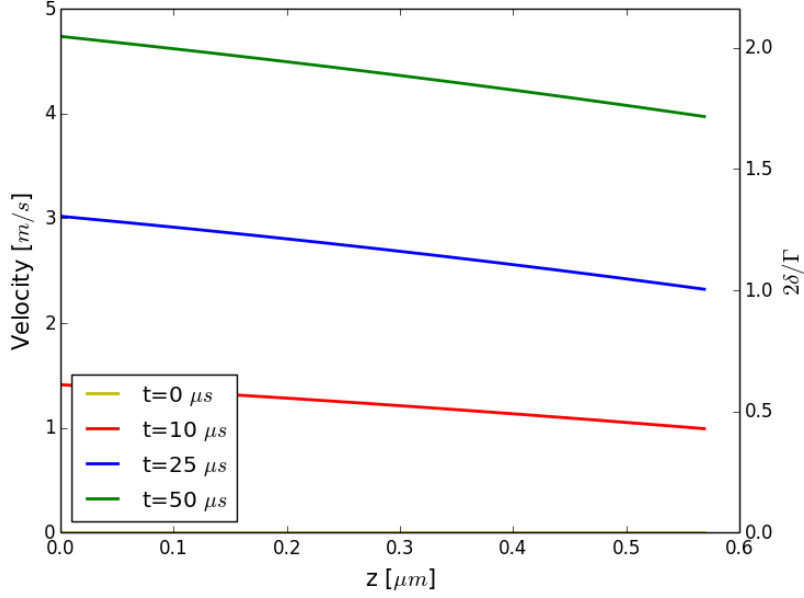


Figure 2. Distribution of generalized detuning $\Delta = \frac{2\delta}{\Gamma}$ across an atomic cloud of ^{40}K for three different imaging times, as obtained by numerical simulation.

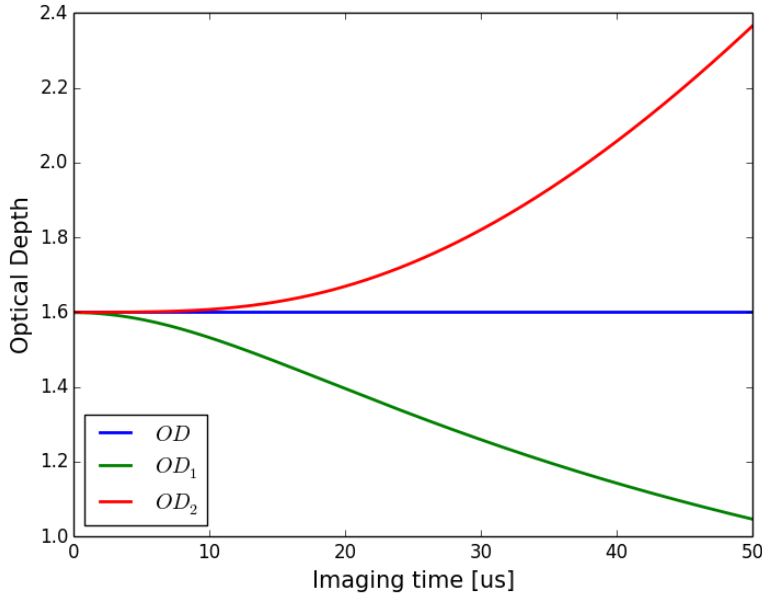


Figure 3. Using time dependent I_f values obtained from recoil detuning corrected simulation of on-resonant imaging of ^{40}K atoms, this graph shows the optical depths each model would deduce from such images. The true optical depth is given at 1.6. OD_1 is the high probe intensity corrected optical depth given by 3. OD_2 is the model that includes high probe intensity corrections and imaging time corrections from expanding 4,5 to second order in imaging time [2]. Note that the value obtained using the second order expansion in time starts to differ significantly from the true value after about 15us. The probe intensity is $0.8 I_{sat}$.

In the following, we describe two versions of this simulation. First, we take the simplistic approach that the on-axis distribution of atoms does not change dramatically during the imaging time and can be treated statically. We test this approach in known limits and then use then check the validity of the static assumption. It turns out that, for realistic input parameters, this assumption is grossly incorrect. So, we take a slightly more sophisticated approach and allow the atoms to move within the cloud during the imaging time. This allows us to simulate the phase space evolution of atoms subjected to probe light. However, in the end we find that numerical differences in predicted OD only vary on the 0.05% level between the two models.

2.1. Stationary atom model

In order to solve eq 4,5, we start with an input 1-d distribution of atom densities $\rho(z)$, usually gaussian in shape. We divide the cloud into spatial bins (the bin size was chosen such that decreasing the bin size further produced less than a 0.001% difference in the result). Each bin carries a certain number of atoms. In this approximation, we keep the number of atoms in each bin constant. Then, we input a probe intensity, and propagate the intensity bin by bin according to 4, such that each bin only sees the probe intensity that was not absorbed in the previous bins. Then, we update the average velocity of the atoms in each bin according to 5 for our time step size dt . Then, we propagate the probe intensity through each bin again, starting with the same intensity at the first bin but this time taking into account the average recoil induced detuning of each bin in calculating the intensity absorbed. We continue this two-step process until we reach the desired imaging time.

We sum the intensity that made it through the entire cloud at each time step $\int_{t=0}^{t_f} I_f(\tau) d\tau$, the observable that is actually detected on the first absorption image. From this we can obtain a simulated optical density, $\nu_s = -\ln \frac{\int I_f(\tau) d\tau}{I_0 t_f}$. This optical density depends on the initial probe intensity and the input atom number, or optical depth. Thus, we can take our two absorption images, which give us a value for the optical density as well as I_0 , and infer what optical depth would have to be put into the simulation to obtain the observed optical density. We call this inferred optical depth OD_{corr} , the optical depth corrected for recoil induced detuning effects.

To check the validity of our simulation, we can take it to certain limits in which the problem becomes analytically solvable. In the limit that the probe intensity is much weaker than saturation, $I_0 \ll I_{sat}$, the atoms will not absorb enough light to significantly detune from resonance. We can then neglect time dependence and 4 reduces to

$$\frac{dI(z)}{dz} = -\rho\sigma_0 I(z), \quad (6)$$

from which we recover the simple

$$\sigma_0 n = OD_0 = \nu. \quad (7)$$

In the limit that the probe intensity is much stronger than saturation, $I_0 \gg I_{sat}$, even far detuned atoms will absorb light at their maximum, allowing us to again neglect the

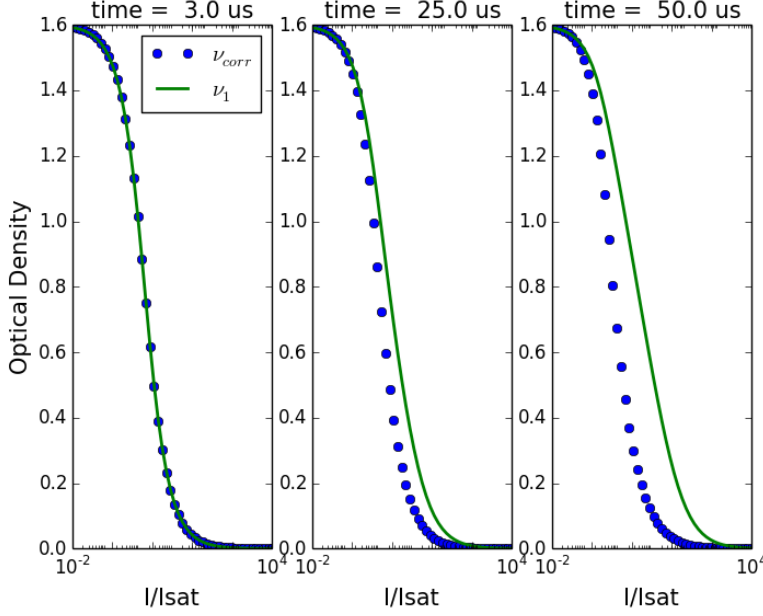


Figure 4. Optical densities as a function of probe intensity as predicted by the simulation and by 3, for three different imaging times. The predictions agree in both the high and low intensity limits, and differ for probe intensities comparable to the saturation intensity. The difference is enhanced with increased imaging time.

time dependence and reduce 4 to

$$\frac{dI(z)}{dz} = -\rho\sigma_0 I_{sat}, \quad (8)$$

which integrates out to

$$\sigma_0 n = \frac{I_0 - I_f}{I_{sat}}. \quad (9)$$

We recognize 7 and 9 as the two terms in the expression for OD_1 , 3. Thus, in both limits OD_{corr} should coincide with OD_1 . Or equivalently, if the optical depth is an input parameter, in those limits both 3 and our simulation should produce the same prediction for optical density. Indeed, they coincide as seen in figure 4.

Encouraged by the reasonable behavior of our simulation, we can now use it to see if the stationary atom assumption is self consistent, ie if the distance travelled by the atoms in one bin during the imaging time is less than the bin size. However, as can be seen from figure 8, it turns out that not only do the atoms travel more than the bin size, but they also travel more than the size of the whole cloud, and the back of the cloud travels less than the beginning for long enough imaging times. Thus, the atomic distribution as a function of position changes dramatically during the imaging time, and our stationary assumption is completely invalid.

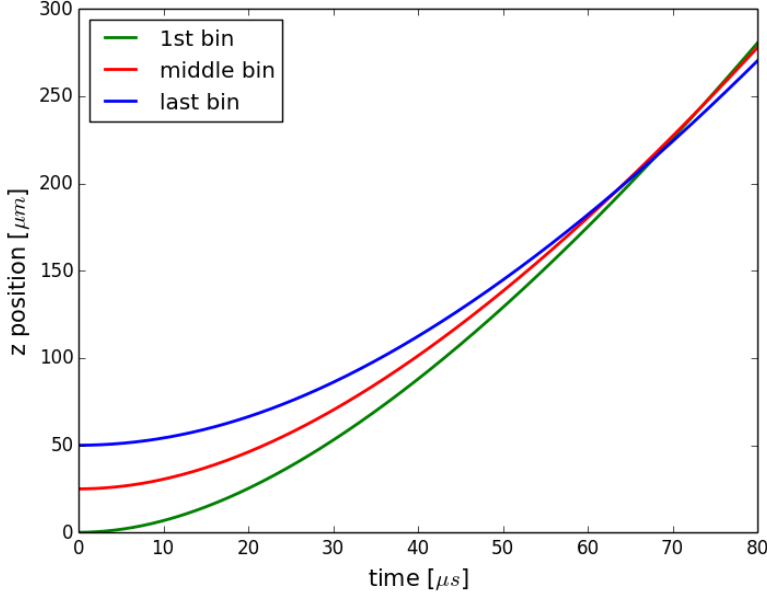


Figure 5. Position of atoms as a function of imaging time for atoms in the first, middle, and last bins of the simulation. The probe intensity here is $1.2I_{sat}$, and the optical depth is 1.6.

2.2. Traveling atom model

To account for the changing atomic distribution during the imaging pulse, we shift our framework slightly from spatial bins of equal size with varying atoms number to superatoms. That is, we divide the initial atomic distribution into clumps of a set number of atoms, called superatoms, with initial positions that reflect the atomic distribution. The simulation then proceeds as before, propagating the probe light through one superatom at a time. But now, after the probe has been propagated through and the velocities of the superatoms are updated, the positions of the superatoms are advanced accordingly before the next time step is taken. Thus, we can simulate an evolving atomic distribution and track the resultant detunings.

First, we should check the velocity behavior of our superatoms against calculable limits. One such limit is that of a single superatom. In this case, the intensity seen by the superatom is constant at the full probe intensity, and the only time dependence in the problem is in the velocity, and thus detuning, of the superatom. 5 can then be re-written as

$$\frac{d\delta(t)}{dt} = \frac{v_r}{\hbar c \rho} \frac{dI}{dz}(t) = \frac{v_r \sigma_0}{\hbar c} \frac{I_0}{1 + \frac{4}{\Gamma^2} \delta(t)^2 + I_0/I_{sat}}, \quad (10)$$

or in dimensionless form,

$$\frac{d\Delta(t)}{dt} = kv_r \frac{\tilde{I}}{1 + \Delta^2 + \tilde{I}}, \quad (11)$$

where $\Delta = 2\delta/\Gamma$, $\tilde{I} = I_0/I_{sat}$ and k is the photonic wavevector. 11 can be integrated

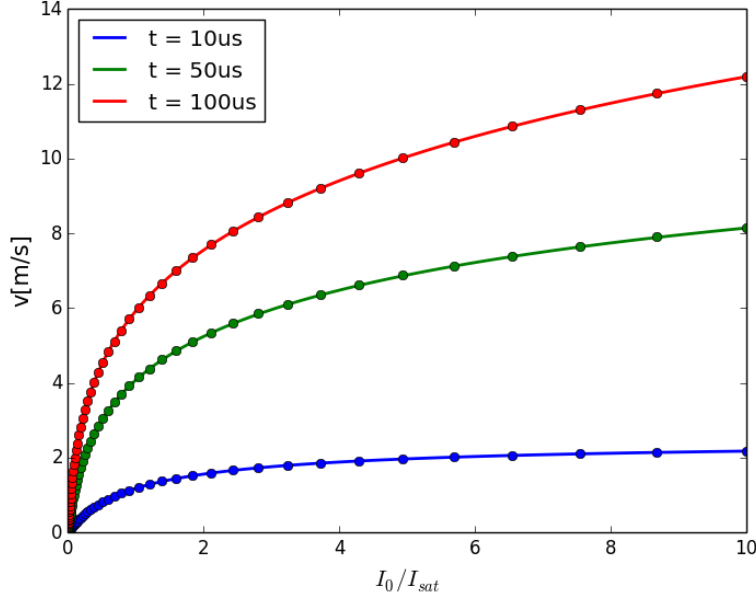


Figure 6. The velocity of a single superatom as a function of probe intensity for various imaging times. The dots represent simulation data, while the lines represent analytical solutions.

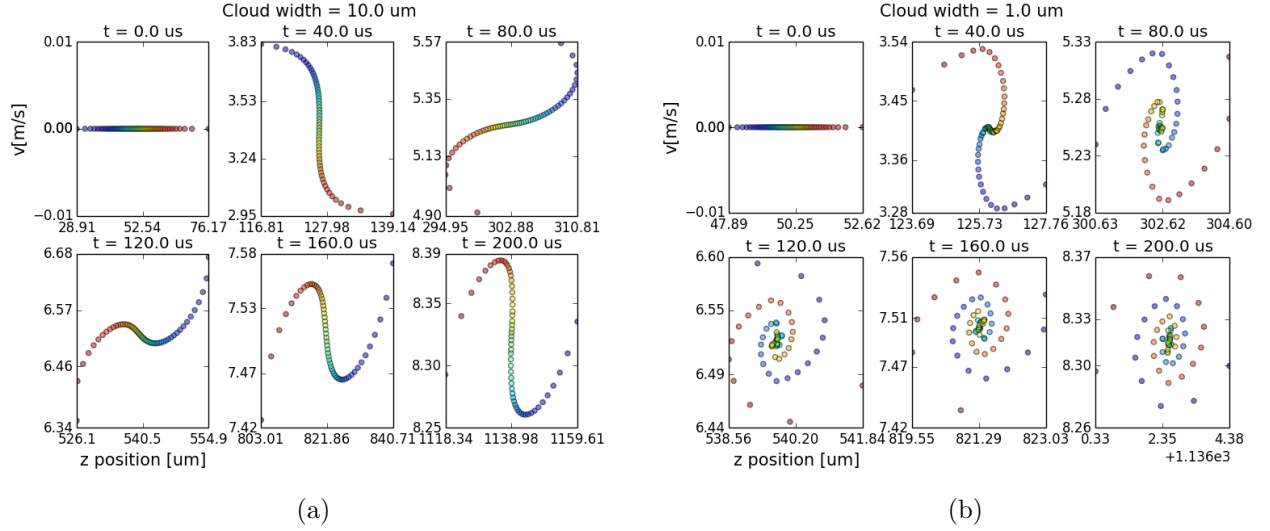


Figure 7. Phase space evolution of an atomic cloud exposed to probe light of $1.2I_{sat}$. The optical depth is 1.6, and the initial cloud is a gaussian with width a. $10\mu m$ and b. $1\mu m$

and solved exactly with one real root. We can compare that result to our simulation, and it agrees as seen in figure 6.

We are now in a position to study the time dependance of the cloud evolution. This can be visualized as a phase space evolution of superatoms, as seen in figure 7. We see that the cloud shape is actually strongly distorted in the imaging time.

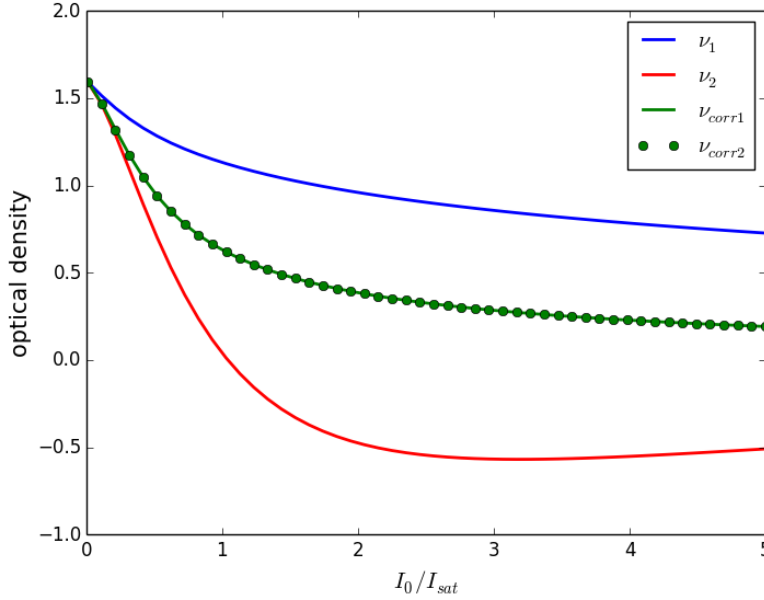


Figure 8. Predictions for optical density as a function of probe intensity for an imaging time $t = 50\mu s$. Note that the two versions of simulated optical density, ν_{corr1} and ν_{corr2} are virtually indistinguishable.

Having performed both the stationary and traveling atom simulations, we can compare the optical densities predicted by each, which we call ν_{corr1} and ν_{corr2} respectively. We find that, despite the significant changes in atomic distribution during the imaging time, the predicted optical densities are so slightly changed by including this effect that it is practically undetectable. In fact, the difference $|\nu_{corr1} - \nu_{corr2}|/\nu_{corr1} \leq 0.05$. Thus, for the purposes of deducing the optical density from experimental optical depths, simply using a stationary model is sufficient. Furthermore, the actual atomic distribution $\rho(z)$ is largely irrelevant, and the only observable is the total atom number $n = \int \rho(z)dz$.

2.3. Calibration of saturation intensity

Now we come to the question of how to interpret the actual output of an imaging camera. Each camera pixel outputs an integer number of "counts", proportional to the total energy it absorbed during a pulse. The precise proportionality constant of these counts to the intensity as seen by the atoms depends on many factors, such as the quantum efficiency of the camera, the photoelectric conversion factor of the camera, and the polarization of the probe light.

The most accurate way to determine this factor is through a direct experiment. In the limit where the system is adequately described by 7, only the ratio of the initial and final intensities matters, and thus this proportionality constant is irrelevant. In all other regimes, however, the ratio of the initial and final intensities to the saturation

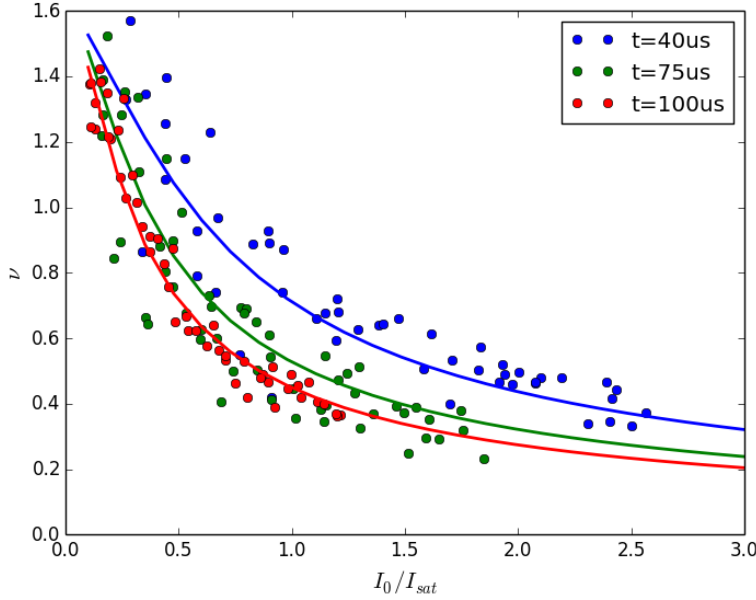


Figure 9. The optical density as a function of probe intensity for three imaging times. The dots represent experimental data and the lines represent the best fit of simulated data. The optimal fit parameters pictured are optical depth of 1.62 and saturation intensity of 29 counts/us.

intensity also comes into play, making the proportionality constant significant. One way to approach this is to calibrate the saturation intensity in terms of camera counts per unit time.

In order to calibrate the saturation intensity in camera counts per unit time, we take absorption images of a cloud of ^{40}K atoms at three different imaging times, 40us, 100us, and 200us, and at varying probe intensities. We pick a small square in the center of the cloud, where the atomic density is approximately uniform, and average the initial and final intensities of each pixel in the square. Thus, for each image we obtain one I_0 and one I_f , in units of counts per microsecond, or equivalently one optical density and one I_0/I_{sat} . We then do a least squares fit of ν_{corr} , our simulated optical density, to the data. The two fit parameters are the optical depth at the center of the cloud, representing the actual atom number, and the value of I_{sat} in counts per microsecond. As seen in figure 9, the model produces a good fit to the experimental data.

2.4. Signal to noise optimization

Now that we have the ability to interpret absorption images at larger imaging times, we can choose an imaging time based on optimal signal to noise ratio. Here, we consider only the photon shot noise that results in camera count noise, with the uncertainty proportional to $\sqrt{N_p}$, where N_p is the photon number. We then propagate this uncertainty through our correction scheme to deduce the final uncertainty in the

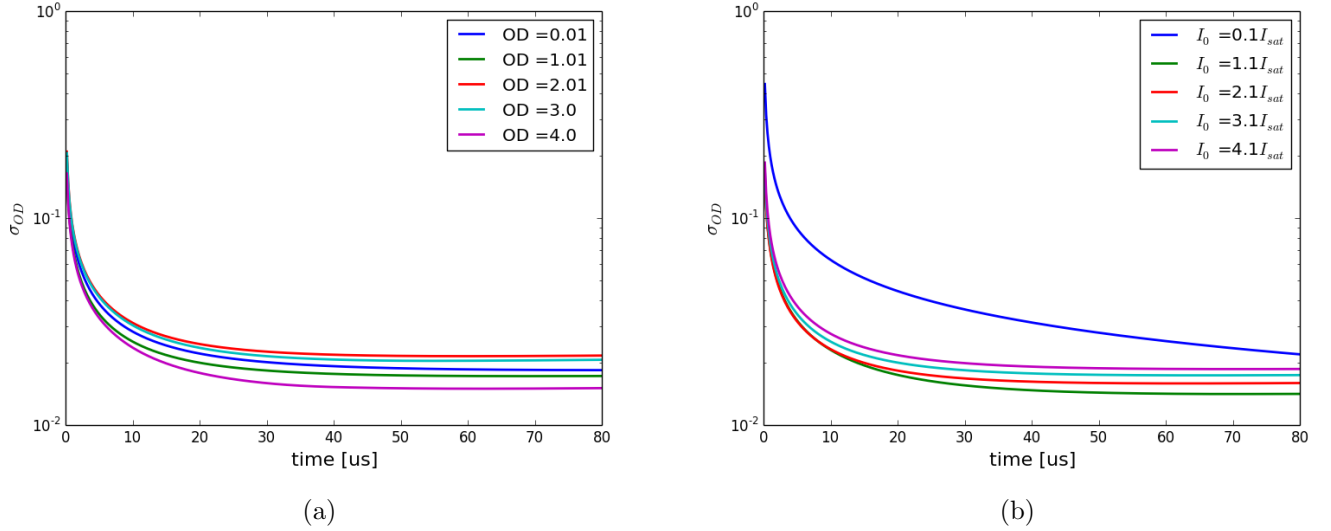


Figure 10. Dependence of detection uncertainty of optical depth on imaging time, after correcting for recoil induced detuning. a. Uncertainty at fixed probe intensity $I_0/I_{sat} = 2.6$ for a range of optical depths and b. fixed $OD = 1.5$ for a range of probe intensities.

calculated optical depth.

As seen in figure 10, around $40\mu s$ for reasonable parameters the uncertainty levels out and there are no longer appreciable benefits by extending the imaging time. The advantage of going to a $40\mu s$ as opposed to $10\mu s$ where the uncorrected model is appropriate is about a factor of 1.6 reduction in measurement uncertainty.

3. S-wave scattering experiment

In this section we describe our experiment with scattering two counter-propagating clouds of ^{40}K atoms and observe the resulting s-wave halo of scattered atoms. We vary the magnetic field around the Feshbach resonance during this experiment, and from the change in scattering deduce a location of the magnetic fields resonance of 202.47 Gauss and a width of 8.98 Gauss.

3.1. Experimental procedure

We load our cold Fermi gas of ^{40}K atoms into a crossed dipole trap. We then use adiabatic rapid passage to bring the gas into an equal superposition of $|9/2, -9/2\rangle$ and $|9/2, -7/2\rangle$, the states whose Feshbach resonance we are trying to study.

We then use a large pair of coils in Helmholtz configuration to bring the magnetic field close to the Feshbach resonance, but not close enough for significant collisions to occur. We hold the atoms there to allow the eddy currents in the large coils to settle, and then use a smaller set of Helmholtz coils to hop the field to the desired value. We take two sets of data: one coming from below the resonance, where we hop the large

coils to a lower field value than the resonance and use the small coils to hop up to the desired field, and one coming from below the resonance, where we hop the large coils to a field value above the resonance and then use the small coils to hop the field down to the desired value. This allows us to correct for losses due to molecule formation and three-body recombination [1].

We then use a double pulse sequence [8] of a near resonant 1-d retro-reflected optical lattice to transfer most of the atoms into the first two excited states of the lattice, giving them $\pm 2k_r$ of momentum, where k_r is the recoil momentum of the lattice. Since the initial Fermi gas has a wide momentum spread (in contrast to a BEC, which has a very narrow momentum spread), and the lattice pulsing is a momentum dependent process, not all the atoms are successfully transferred into the 1st excited band of the lattice. We optimized our pulse times to minimize the number of atoms that remain in the lowest, zero momentum, band of the lattice, but were not able to eliminate them completely. The optimized pulse times are $23\mu s$ for the first square pulse, $13\mu s$ off interval, and $12\mu s$ for the second square pulse.

We release the atoms from the trap and allow 1ms for the two opposite momentum states within the cloud to pass through each other, scattering on the way. For the data taken coming from below the Feshbach resonance, we then simply ramp down the field and image the atoms. For the data taken coming from above the Feshbach resonance, we then ramp the field back up through the resonance to recover any molecules that were created when the field was ramped from the attractive to the repulsive side of the resonance, and then quickly ramp the field back down and image the atoms. We use an imaging pulse of $40\mu s$.

3.2. Methods

We first correct each image for recoil induced detuning using the simulation described above. The simulation gives us a look-up table of observed optical density as a function of atom column density and probe intensity. From our two absorption images we observe the probe intensity and optical density for each pixel. We then use the simulation look-up table in an inverted fashion to deduce the column density or, equivalently, the optical depth. An example of the effect of this correction procedure is shown in figure 11

To improve the signal and compensate for our shot to shot number fluctuations, we take 15 equivalent images for each data point and, after correcting, add them together. We then perform a standard inverse Abel transform to extract the radial atom number dependence from the two dimensional projection we see in our images. The inverse Abel transform assumes a cylindrical symmetry, which is present in our case, with the axis of symmetry defined by the 1-D lattice.

We then extract the number of atoms that went through a single scattering event N_{scat} , as a fraction of the total atom number N_{tot} , for each bias magnetic field value. We unscattered number by counting the number of atoms in the two unscattered clouds. We obtain the number of atoms that underwent single scattering by counting the number

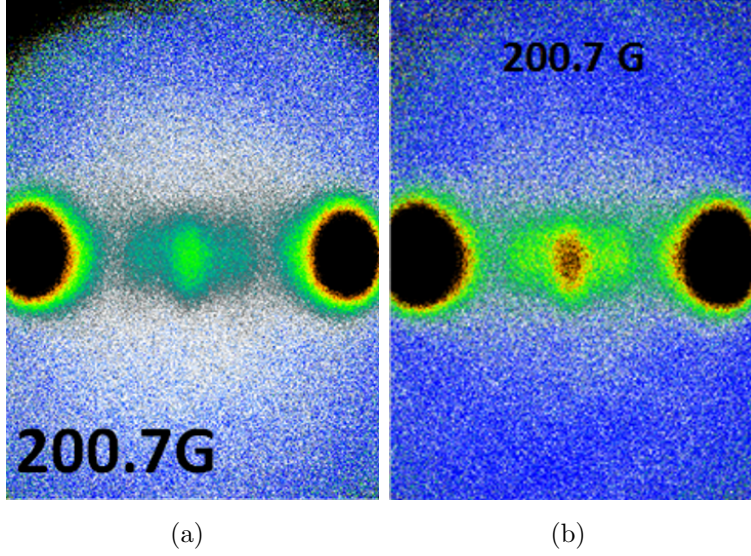


Figure 11. An example of our absorption image after 6.784ms time of flight. The 1-D lattice is along the horizontal direction at the center of our images. The two large clouds on the left and right are the atoms in the $\pm 2k_r$ momentum orders that passed through each other unscattered. The smaller cloud in the center is the atoms that remained in the lowest band of the lattice after pulsing, and thus obtained no momentum. The thin spread of atoms around these clouds is the atoms that underwent scattering. a. Raw optical density b. Optical density of the same image after correcting for recoil induced detuning by comparing to simulation

of atoms outside the Fermi radius of the unscattered clouds, but inside the arc created by rotating the Fermi radius around the original center of the cloud. This method excludes atoms that underwent multiple scatterings, as those atoms are not bound by momentum conservation to have travelled the same distance from the center as the unscattered clouds in time of flight. The total atom number is the sum of those two.

We then use our data to deduce the resonant field value, the width of the resonance, etc. To do this, we must fit our data to a model. From [1], we have the expression:

$$a(B) = a_{bg} \left(1 - \frac{\Delta}{B - B_0} \right), \quad (12)$$

where a is the scattering length, a_{bg} is the scattering length away from the resonance, Δ is the width of the resonance, and B_0 is the field value at which the resonance occurs. Since we are in the low energy regime, the scattering cross-section is given by $\sigma = 4\pi a^2$.

One way to think about the scattering cross-section σ is that the probability P_{scat} that a single particle will get scattered when incident on a cloud of atoms with a surface density of $\frac{N}{A}$ is given by $P_{scat} = \sigma \frac{N}{A}$. If this experiment is repeated 1000 times, the number of times the atom will get scattered is expected to be $N_{scat} = 1000\sigma \frac{N}{A}$. In our case, half the initial cloud, with atoms number $N_{tot}/2$, is incident on the other half of the initial cloud, again with $N_{tot}/2$ atoms. Thus, the number of scattered atoms should be given by $N_{scat} = \frac{N_{tot}}{2} \sigma \frac{N_{tot}}{2} = \sigma \frac{N_{tot}^2}{4A}$, where A is the cross-sectional area of the cloud. Assuming A is constant for all our data, we can absorb the factor of $4A$ into our

definition of a_{bg} , along with the 4π , to obtain

$$\frac{N_{scat}}{N_{tot}^2} = \tilde{a}_{bg}^2 \left(1 - \frac{\Delta}{B - B_0} \right)^2. \quad (13)$$

Unfortunately, we cannot fit to this equation directly - camera readout noise is skewed towards the positive, creating an asymmetry in detected shot noise for very low and zero atom number pixels. Thus, there is a background number of what looks like scattered atoms in our treatment, giving rise to a background offset o in our data. Thus, the equation we actually fit to is

$$\frac{N_{scat}}{N_{tot}^2} = \tilde{a}_{bg}^2 \left(1 - \frac{\Delta}{B - B_0} \right)^2 + o, \quad (14)$$

where our fit parameters are \tilde{a}_{bg} , Δ , B_0 , and o .

3.3. Results

Our final data is presented in figure 12. The red line is a best fit of the model given in eq.14. The fit parameters we extract are $\tilde{a}_{bg} = 1.48e - 3$, $\Delta = 8.98$ Gauss, $B_0 = 202.47$ Gauss, and $o = 9.00e - 5$. The error bars on the fitted data are obtained solely from photon shot noise of both absorption images propagated through our analysis. The accepted values for the ^{40}K s-wave Feshbach resonance for the $|9/2, -9/2\rangle$ and $|9/2, -7/2\rangle$ states are $B_0 = 202.1 \pm 0.07$ and $\Delta = 7.8 \pm 0.6$.

4. Conclusion

We studied the effects of recoil-induced detuning effects on absorption images and found an optimal imaging time of $40\mu s$ for ^{40}K atoms for noise minimization after corrections. We use these results to observe s-wave scattering halos of the Fermi gas around the Feshbach resonance and directly verify the resonance location and width. Our analysis can be used in any absorption imaging applications where signal to noise minimization is critical. Experimental techniques developed for our s-wave scattering experiment can be readily extended to a Raman-coupled single component gas version, in which we hope to detect synthetic p-wave interactions.

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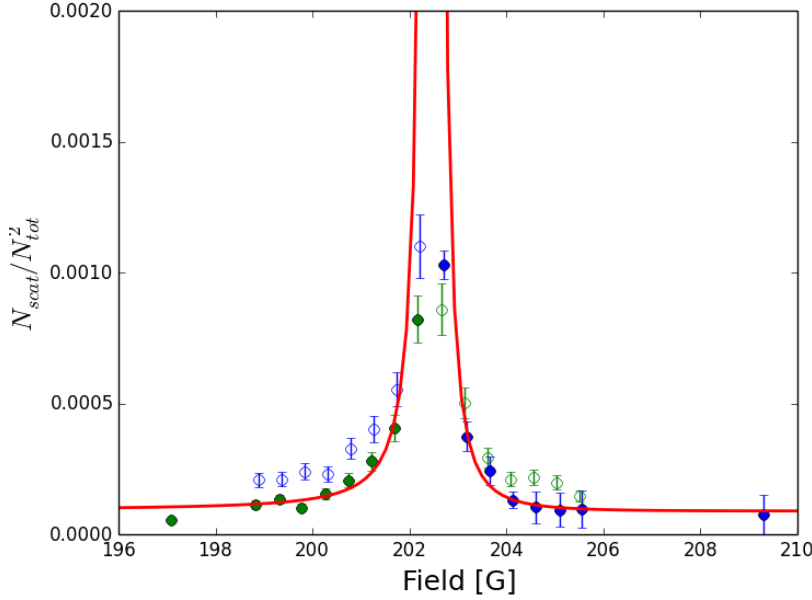


Figure 12. Scattered fraction over the total atom number as a function of bias field. Green dots represent data taken coming from below the resonance, and blue dots represent the data taken coming from above the resonance. Red is the line of best fit, where data coming from above the resonance was used above the resonance and data coming from below the resonance was used below the resonance to create the fit. The four points in the middle were not used in the fit at all, as they are in the regime where the scattering length is large enough for the atoms to behave hydrodynamically and the assumption $\sigma\rho \ll 1$, where ρ is the atom number per unit area, is no longer valid.

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