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Citation: [Review of Scientific Instruments](#) **81**, 063103 (2010); doi: 10.1063/1.3430538

View online: <http://dx.doi.org/10.1063/1.3430538>

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^{85}Rb tunable-interaction Bose–Einstein condensate machine

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(Received 25 March 2010; accepted 26 April 2010; published online 9 June 2010)

We describe our experimental setup for creating stable Bose–Einstein condensates (BECs) of ^{85}Rb with tunable interparticle interactions. We use sympathetic cooling with ^{87}Rb in two stages, initially in a tight Ioffe–Pritchard magnetic trap and subsequently in a weak, large-volume, crossed optical dipole trap, using the 155 G Feshbach resonance to manipulate the elastic and inelastic scattering properties of the ^{85}Rb atoms. Typical ^{85}Rb condensates contain 4×10^4 atoms with a scattering length of $a = +200a_0$. Many aspects of the design presented here could be adapted to other dual-species BEC machines, including those involving degenerate Fermi–Bose mixtures. Our minimalist apparatus is well suited to experiments on dual-species and spinor Rb condensates, and has several simplifications over the ^{85}Rb BEC machine at JILA, which we discuss at the end of this article. © 2010 American Institute of Physics. [doi:10.1063/1.3430538]

I. INTRODUCTION

In recent years there have been a number of important discoveries in condensed matter physics permitted by the ability to change the interactions between ultracold bosons. In the weakly interacting regime, near-ideal Bose gases have recently been used to observe Anderson localization^{1,2} and promised to improve coherence times in atom interferometers.^{3,4} The strongly interacting regime, on the other hand, has been used to study beyond mean-field effects, including the collapse and explosion of attracting Bose–Einstein condensates (BECs) known as the “bosonova.”⁵

The majority of these experiments have used a Feshbach resonance, a magnetically tunable molecular bound state, to vary the elastic scattering properties of the Bose gas. Tuning a bound state around the atomic scattering threshold using the Zeeman effect results in the divergence of the s -wave scattering length a according to

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

where a_{bg} is the scattering length far from B_0 , the magnetic field at the resonance, and the width Δ is the difference between B_0 and the magnetic field at which $a=0$. Such resonances have been used to modify the s -wave scattering length in BECs of ^{23}Na ,⁶ ^{85}Rb ,⁵ ^{133}Cs ,⁷ ^{39}K ,⁸ ^{52}Cr ,^{9,10} and ^7Li .¹¹ Feshbach resonance physics also led to the successful creation of a molecular BEC from a Fermi gas¹² and has allowed significant advances in the field of quantum chemistry.¹³

The first BEC with tunable interactions was of ^{85}Rb , created by Wieman group⁵ at JILA in 2000. This species has a broad, low-field Feshbach resonance that allows the s -wave scattering length to be tuned over several orders of magnitude, including both positive and negative values (Fig. 1).

However, poor elastic and inelastic scattering cross sections make condensation of ^{85}Rb notoriously difficult to achieve. Indeed, until now only one group has successfully realized this goal.¹⁴

In this article we describe our setup for achieving the Bose–Einstein condensation of ^{85}Rb , based on the recent work of Papp and co-worker.^{15,16} We exploit the large inter-species collision cross section between the two isotopes¹⁷ to sympathetically cool ^{85}Rb using ^{87}Rb as a refrigerant, first in a magnetic trap and subsequently in a weak, large-volume optical dipole trap. Our machine creates tunable-interaction ^{85}Rb BECs containing 4×10^4 atoms. We have refined the approach of Papp,¹⁵ resulting in a substantially simpler setup and a shorter duty cycle. A detailed comparison of the two machines is presented at the end of this paper. Our setup and method also have similarities to the $^{39}\text{K}/^{87}\text{Rb}$ machine of Roati *et al.*,⁸ and can be proven useful in other dual-species machines, particularly those requiring sympathetic cooling and scattering length manipulation, and for investigating degenerate Fermi–Bose mixtures.

II. OVERVIEW

In our system, a cold beam containing both ^{85}Rb and ^{87}Rb atoms is produced in a two-dimensional (2D) magneto-optical trap (MOT) and is directed through the vacuum system to a retroreflected, three-dimensional (3D) MOT in the main chamber, which is evacuated to a pressure of $\sim 10^{-11}$ Torr. The atoms are pumped into their lower ground states before being loaded into a quadrupole Ioffe–Pritchard configuration (QUIC) magnetic trap. Here, the ^{87}Rb is selectively evaporated by a radiofrequency sweep which drives transitions to untrapped states, sympathetically cooling the ^{85}Rb through thermal contact. Once the temperature of the combined sample has fallen to 20 μK , the atoms are transferred to a crossed optical dipole trap and a large magnetic bias field is applied to suppress inelastic collisions in ^{85}Rb . Finally, the depth of the dipole trap is reduced, resulting in further evaporation of both species. With the appropriate

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FIG. 1. Variation in s -wave scattering length with magnetic field near the ^{85}Rb Feshbach resonance. The parameters of the resonance are $a_{bg} = -443a_0$, $B_0 = 155.0$ G, and $\Delta = 10.7$ G (Ref. 14).

magnetic field strength, the ^{85}Rb scattering length can be made positive and a stable condensate of 4×10^4 atoms is created.

The use of an optical dipole trap to confine the sample at low temperatures facilitates control of the scattering properties of ^{85}Rb using the magnetic Feshbach resonance at 155 G. However, direct loading of the optical trap from the MOT is not feasible due to large density-dependent inelastic losses in ^{85}Rb . Thus we require precooling of the sample in a magnetic potential before transfer to the dipole trap for final evaporation. In our system the QUIC trap, the dipole trap, and the Feshbach field coils are all coaxial, which facilitate transfer between the traps and control of the bias field. We image the atoms through absorption of resonant light.

III. VACUUM SYSTEM

A schematic of our vacuum system is shown in Fig. 2. The setup is simple and compact compared to many BEC machines. Two $25 \times 25 \times 80$ mm³ quartz glass cells for the 2D and 3D MOTs are flanked by a 75 l/s ion pump and a titanium sublimation pump housed in a 6 in. tube. An electrical feedthrough provides current to two 50 mg rubidium

FIG. 2. Vacuum system schematic. The 2D and 3D MOTs are collected in coaxial quartz glass cells, connected by a small impedance which maintains a pressure differential between the two sections.

TABLE I. The laser frequencies used in this experiment. Typical values for the power available after fiber coupling and the detuning relative to the specified transition are given in the columns on the right.

Purpose	Transition	Detuning (MHz)	Power (mW)
^{87}Rb			
3D MOT trapping	$F=2 \rightarrow F'=3$	-24	60
3D MOT repumping	$F=1 \rightarrow F'=2$	0	20
2D MOT trapping	$F=2 \rightarrow F'=3$	-12	150
2D MOT repumping	$F=1 \rightarrow F'=2$	0	20
Push beam	$F=2 \rightarrow F'=3$	+5	0.15
Imaging	$F=2 \rightarrow F'=3$	-6	0.4
^{85}Rb			
3D MOT trapping	$F=3 \rightarrow F'=4$	-20	15
3D MOT repumping	$F=2 \rightarrow F'=3$	0	2.5
2D MOT trapping	$F=3 \rightarrow F'=4$	-12	35
2D MOT repumping	$F=2 \rightarrow F'=3$	0	3.0
Push beam	$F=2 \rightarrow F'=3$	+29	0.03
Imaging	$F=2 \rightarrow F'=3$	0	0.4

dispensers (Alvatec AS-Rb-50-F, Althofen, Austria) mounted in the 2D MOT cell. Pressure readings are taken from the ion pump and from a cold cathode gauge (Pfeiffer IKR 270, New Hampshire, USA) mounted on one of the large tubes, and agree to within a factor of 2 in their specified operating ranges. After construction, the entire system was baked at 300 °C for 6 days.

To create a BEC, an ultrahigh vacuum (UHV) of $\lesssim 10^{-10}$ Torr is required to reduce atom loss by collisions with background vapor particles, in order to enable efficient evaporative cooling. However, to collect a large MOT a relatively high pressure is optimal—up to Rb vapor pressure of $\sim 10^{-7}$ Torr. To fulfill both these requirements, our vacuum system is partitioned into two sections by a small impedance—a stainless steel tube protruding into the 2D MOT glass cell with a 0.8 mm diameter hole. The low conductance $C = 7 \times 10^{-3}$ l/s of this tube allows the pressure in the two sections to differ by several orders of magnitude. A 2D MOT in the high vacuum section produces a beam of cold atoms, which are pushed through the impedance to feed the 3D MOT in the UHV cell. The pressure in the UHV cell is maintained at 3×10^{-11} Torr.

IV. LASER SYSTEM

The laser light used for the ^{85}Rb and ^{87}Rb MOTs and for absorption imaging of the atom clouds is derived from six homebuilt external cavity diode lasers (ECDLs) in the Littrow configuration (Table I). These use feedback from a low-efficiency holographic diffraction grating to narrow the line-width of a 780 nm laser diode (Roithner ADL-78901TX, Vienna, Austria), giving between 50 and 80 mW of output at the frequency of the rubidium D_2 line ($5^2S_{1/2} \rightarrow 5^2P_{3/2}$). The lasers are locked using saturated absorption spectroscopy and either current or Zeeman modulation,¹⁸ and double passed through acousto-optic modulators which allow the light to be shifted to the desired frequency and swiftly shuttered. The light from two of these lasers (those driving the ^{87}Rb MOTs)

is amplified by 1.5 W tapered amplifier chips (m2k laser TA-780-1500). All the light is guided to the experiment in single-mode polarization-maintaining optical fibers, which serve to isolate the experiment from the laser table and to mode clean the light. Each fiber carries two frequencies—one for each species—with orthogonal polarization.

V. MAGNETO-OPTICAL TRAPS

In a typical experimental run, 10^{10} ^{87}Rb atoms and 10^8 ^{85}Rb atoms are collected in the 3D MOT. A low power push beam slightly detuned from the cooling transition drives atoms from the 2D MOT through the impedance into the UHV section of the vacuum system, where they are captured in the 3D MOT. Each MOT uses retroreflected beams apertured to 20 mm in diameter. The trapping lasers for the 3D MOT are detuned 24 and 20 MHz to the red of the ^{87}Rb $|F=2\rangle \rightarrow |F'=3\rangle$ transition and the ^{85}Rb $|F=3\rangle \rightarrow |F'=4\rangle$ transitions, respectively. The 2D MOT trapping lasers are each tuned 12 MHz to the red of these transitions. To prevent loss of atoms through off-resonant transitions to the lower ground states, which occur roughly once in 10^4 absorption events, repumping light resonant with the ^{87}Rb $|F=1\rangle \rightarrow |F'=2\rangle$ transition and the ^{85}Rb $|F=2\rangle \rightarrow |F'=3\rangle$ transition is mixed in with the trapping light in both MOTs.

We have also experimented with using a single ECDL locked to the ^{87}Rb cooling transition to drive both 2D MOTs by passing the light through an electro-optic modulator (EOM) in a microwave cavity driven at 1.1265 GHz—corresponding to the frequency difference between the cooling transitions for each isotope. Up to 20% of the total power could thus be transferred into a sideband resonant with the ^{85}Rb cooling transition (a further 20% in the upper sideband was not resonant with either species). This light was then amplified by a 1.5 W tapered amplifier chip; 250 mW was available at the experiment after fiber coupling. Although this setup was easier to align (since the 2D MOTs were automatically coincident) and allowed us to produce larger ^{85}Rb MOTs, we found that thermal and mechanical instabilities in the EOM caused large run-to-run atom number fluctuations.

Using either approach, the flux from the 2D MOT is sufficient to fill the 3D MOT within 5 s. After a 20 ms polarization-gradient cooling stage, the MOT repumping lasers are switched off for 1 ms to allow both species to be optically pumped into their lower ground states (^{87}Rb $|F=1\rangle$ and ^{85}Rb $|F=2\rangle$) in preparation for sympathetic cooling.

VI. MAGNETIC TRAP

Once the MOT light has been switched off, the low-field seeking states ^{85}Rb $|F=2, m_F=-2\rangle$ and ^{87}Rb $|F=1, m_F=-1\rangle$ are captured in the quadrupole magnetic field produced by the MOT coils and then magnetically transported over a distance of 40 mm to a second quadrupole trap with the aid of a rectangular transfer coil (see Fig. 3). In this process, the current in the transfer coil is ramped up to push the cloud toward the Ioffe–Pritchard trap coils, then the MOT and transfer coils are turned off as the current in the

FIG. 3. Magnetic trap coils mounted around the UHV cell. Not shown are the rf coil, which produces the oscillating magnetic field used for evaporating atoms from the magnetic trap and the imaging coil, which defines a quantization axis to ensure maximum absorption of the probe light during imaging.

quadrupole coils ramps up. This quadrupole trap is then converted into a harmonic Ioffe–Pritchard potential using the QUIC configuration (QUIC).¹⁹

The QUIC trap coils were designed using a full numerical optimization of the trapping frequencies at a given power. The calculation was performed taking into account the various geometric constraints of the experiment and provided the optimal size and position of the coils, the diameter of the wire to be used for the windings, and the current to flow in each coil. The quadrupole coils each consist of 137 turns of enameled copper wire with diameter of 1.35 mm and run at 25 A, dissipating a total power of 270 W. The Ioffe coil has 33 turns of 0.9 mm wire and also runs at 25 A, dissipating 40 W.

The trap coils are cooled by a recirculating solid-state chiller, which pumps water at 13 °C through holes in the aluminum trap mount. To maximize the cooling efficiency, the coils were wound directly onto the mount using a high thermal conductivity epoxy (Cotronics Duralco 132, New York, USA). Before winding the Ioffe coil, we mixed fine (filed) copper dust with the epoxy in roughly a 1:2 ratio to improve its thermal conductivity ($k_{\text{Cu}}=380$ W/m K compared to $k_{\text{epoxy}}=6$ W/m K). This had no detrimental effect on the properties of the epoxy.

Since we do not use the magnetic trap to study condensates, there is no stringent requirement on the stability of the bias field, thus we can use separate power supplies to drive the quadrupole and Ioffe coils. This facilitates small adjustments of the bias field. Fast switch-off of the trap coils is essential for absorption imaging of the atoms in the magnetic trap and when transferring samples to the dipole trap. We are able to achieve a switching time of 200 μs using a solid-state relay with a diode/resistor shunt around the coils. Radial slots cut into the aluminum coil mount also help by minimizing eddy currents during switch-off.

The final magnetic potential has trapping frequencies of $\omega_z=2\pi\times 16$ Hz axially and $\omega_\rho=2\pi\times 156$ Hz radially for ^{87}Rb atoms in the $|F=1, m_F=-1\rangle$ hyperfine state, and a bias field of 3.7 G. Typically, 3×10^9 ^{87}Rb atoms and 1×10^7

^{85}Rb atoms are present at 200 μK in the QUIC trap before evaporation. The lifetime of the atoms in the magnetic trap is measured to be 20 s, resulting in a background loss of roughly 50% during the 15 s of rf evaporation. This loss is due to collisions with background gas particles, predominantly rubidium vapor from the 2D MOT chamber. The lifetime can be increased by lowering the pressure in the 2D MOT cell; however this reduces both the rate at which the 3D MOT fills and the final number of atoms collected. The pressure is controlled by the current flowing through the dispensers, and is chosen to maximize the number of atoms present at the end of the evaporation.

Radiofrequency forced evaporation is used to cool the samples in the magnetic trap to microKelvin temperatures. An oscillating rf field induces magnetic dipole transitions between trapped and untrapped spin states to expel the most energetic atoms, and those remaining rethermalize via elastic collisions. We use an arbitrary waveform generator (Agilent 33250A, California, USA) and a 4 W rf amplifier (Mini-Circuits TIA-1000-1R8) to drive a two-loop coil of radius 12 mm placed against the glass cell. The function generator produces a logarithmic rf sweep from 50 to ~ 3 MHz over 15 s, with the final frequency determining the temperature of the sample. In the absence of ^{85}Rb atoms, we are able to produce pure ^{87}Rb $|F=1, m_F=-1\rangle$ condensates containing 2×10^6 atoms in the QUIC trap after 15 s of evaporation.

VII. CONTROL AND IMAGING

Our machine is controlled by a desktop computer running National Instruments LABVIEW 7, Texas, USA. The computer is equipped with two eight-channel 16-bit analog output cards (NI PCI-6733), giving variable voltage outputs of -10 to 10 V, and a 32-channel digital card (NI PCI-6533) with 5 V outputs. A computer clock signal updates the state of each channel, providing timing control accurate to <100 ns with a resolution of 100 μs . During a long run, the memory required to store the state of all analog channels at each update can exceed the capacity of the card's buffer. To overcome this difficulty, two of the digital channels are reserved for triggering analog updates only when a channel value is altered. Since there are often long periods during a run when no updates are required (e.g., during MOT loading), this substantially reduces the amount of information that must be stored in the buffer.

Absorption imaging is our primary diagnostic tool for ultracold atom clouds. A 100 μs pulse of repumping light is applied first to pump the atoms out of the lower ground state, immediately followed by a 100 μs pulse of light on the cooling transition. The imaging light is circularly polarized and a small magnetic bias field is applied along the direction of the probe light to ensure that the imaging transition is closed. The shadow left by the atoms on the imaging laser is captured with an IEEE1394 charge-coupled device camera (Point Grey Research Dragonfly, British Columbia, Canada) and acquired using CORIANDER for LINUX (Ref. 20) on a separate computer. The images are processed using MATLAB to determine the number, spatial distribution, and temperature of either species at the end of a run.

FIG. 4. Atom number as a function of temperature during rf-induced evaporative precooling in the magnetic trap. The evaporation trajectories of ^{85}Rb (filled circles) and ^{87}Rb (open circles) are shown, as well as the single-species trajectory for ^{87}Rb (squares).

VIII. SYMPATHETIC COOLING

Sympathetic precooling in the magnetic trap is critical to achieving ^{85}Rb BEC in our setup. This is because achieving a sufficiently high phase space density in an optical trap loaded directly from a MOT would either involve very high atomic densities, for which density-dependent inelastic losses would prevent efficient evaporation or would require a prohibitively large amount of laser power. Precooling in a magnetic trap allows us to create cold samples with phase space densities on the order of 10^{-3} , which can then be loaded into a shallow, large-volume optical dipole trap for final evaporation to quantum degeneracy.

With both species in the QUIC trap, rf evaporation predominantly removes ^{87}Rb $|F=1, m_F=-1\rangle$ atoms, cooling both species as long as they remain in thermal contact. The reason for the isotope selectivity is twofold: First, since the Landé factor g_F has a larger magnitude for the ^{87}Rb ground state ($|g_F|=1/2$) than for ^{85}Rb ($|g_F|=1/3$), the ^{87}Rb evaporation surface at a given radiofrequency is closer to the center of the trap than for ^{85}Rb ; second, the ^{85}Rb $|F=2, m_F=-2\rangle$ cloud is more tightly confined in the magnetic trap (i.e., smaller at a given temperature) due to its larger $g_F m_F$ factor. The interspecies ^{87}Rb – ^{85}Rb collision cross section is large in the temperature range of interest (below 200 μK),²¹ and the two clouds remain spatially overlapped until the temperature drops below ~ 1 μK , where the difference in the gravitational sag for each species becomes comparable to the size of the clouds. The present limit to this precooling stage is therefore only the number of ^{87}Rb atoms initially present in the magnetic trap: Although we collect enough to create pure ^{87}Rb condensates, the addition of ^{85}Rb places a heat load on the sample which prevents us from reaching submicroKelvin temperatures in ^{85}Rb .

Figure 4 shows the number of atoms as a function of temperature for each species during the magnetic trap precooling. The single-species ^{87}Rb evaporation is also shown.

The isotope selectivity of the evaporation is clearly demonstrated by the fact that no loss of ^{85}Rb is detected during cooling from 200 to 20 μK , corresponding to an increase in phase space density of over three orders of magnitude. The sympathetic cooling trajectory of ^{85}Rb begins to roll off at around 20 μK , as the number of each species present becomes comparable. This is to be expected since beyond this point the density of ^{85}Rb at the evaporation surface will not necessarily be lower than that of ^{87}Rb and so the rf begins to remove both species from the trap. With a larger ^{87}Rb reservoir, it may be possible to continue the sympathetic cooling further. However, the inelastic scattering properties of ^{85}Rb are known to prevent the containment of high density samples at low magnetic fields—it is expected that sympathetic cooling would no longer be effective below ~ 5 μK , depending on the density of the sample.¹⁵

Under optimal experimental conditions, we can create samples of 8×10^6 ^{85}Rb atoms at 10 μK in the QUIC trap, with a phase space density of 6×10^{-4} . When making BEC, we typically begin with a smaller sample of ^{85}Rb in order to have more ^{87}Rb remaining in the trap and cease rf evaporation at 20 μK with 6×10^6 ^{85}Rb atoms and 4×10^7 ^{87}Rb atoms available for transfer to the optical dipole trap.

IX. OPTICAL DIPOLE TRAP

In order to create and study a ^{85}Rb BEC with more than ~ 100 atoms, it is necessary for the atoms to be in a magnetic field at which the s -wave scattering length a is positive. The most convenient point at which this occurs is on the high-field side of the 155 G Feshbach resonance. However, it is difficult to make a tightly confining magnetic trap with such a large bias field B_0 since for the standard Ioffe–Pritchard-type traps the radial trapping frequency falls as $\omega_p \propto 1/\sqrt{B_0}$. Furthermore, many of the experiments one would like to perform on a ^{85}Rb condensate involve changing the scattering length via the magnetic field, and the relationship between B_0 and ω_p means that bias field adjustments would couple into changes in radial confinement, which could cause undesirable excitations in the sample. For this reason, we confine the atoms in a far-detuned crossed-beam optical dipole trap for the final stage of the experiment and use a set of dedicated “Feshbach” coils to generate a homogeneous magnetic field for manipulation of the scattering length. The light for the dipole trap is sourced from a 20 W Er-doped fiber laser (SPI Lasers redPOWER compact, Southampton, UK) operating at 1090 nm. The laser beam has a diameter of 5 mm, an M^2 value of <1.1 , and an emission linewidth of 2 nm full width at half maximum.

The optical setup for the dipole trap (Fig. 5) is remarkably simple. The unpolarized output from the fiber laser is split evenly on a 1 in. polarizing beam splitter. Each beam then passes through a 2 in. planoconvex lens with focal length $f=100$ cm. The axial beam is directed into the end of the glass cell through the center of the Ioffe coil, parallel to the long axis of the QUIC trap. The cross beam also passes through the cell in the horizontal plane (see Fig. 5), making an angle of 75° with the axial beam.²² The short coherence length (<1 mm) of the laser prevents standing wave pat-

FIG. 5. (a) Schematic of the optical setup for the dipole trap. A and B are the paths taken by the axial and cross beam, respectively, which cross in the cell at an angle of 75° . (b) The position of the dipole beams and Feshbach coils in relation to the magnetic trap.

terns forming in the overlap region, eliminating the need for the frequency of the cross beam to be shifted using an acousto-optic modulator, which would reduce the power available for trapping.

The intensity of the dipole trap during an experimental run is controlled by a 0–10 V analog input on the laser controller. Without any active stabilization, we are able to evaporatively cool in the optical trap over 7 s and hold condensates for up to 20 s. We measure the relative rms intensity fluctuations in the dipole trap laser to be 2×10^{-4} in a 1 Hz bandwidth around the mean trapping frequency $\bar{\omega} = \frac{2\pi}{\times 43 \text{ Hz}}$, giving a relative intensity noise of $-37 \text{ dBc}/\sqrt{\text{Hz}}$ at $\bar{\omega}$.

The lens in the axial (cross) beam is placed 98 cm (106 cm) from the atoms so that the beam radius at the cloud is approximately 150 μm (200 μm). Although the cloud is

FIG. 6. Splitting of the cloud during the ramp up of the bias field, with the current in the Feshbach coils at 0%, 50%, and 100%. The upper panel shows contour lines of the magnetic plus gravitational potential, with the axial dipole trapping beam overlaid (not to scale). The lower panel shows the shape of the total potential seen by the atoms in the y direction (along the dashed lines on the contour plots). The formation of a double well splits the trapped cloud in two, and those atoms not confined by the optical trap are lost as the magnetic trap sags under gravity and eventually reaches the edge of the glass cell.

not precisely at the waist of either beam, the long Rayleigh range $z_R > 5$ cm and the resulting weak trapping force along each beam (equivalent to < 1 Hz) means that the trap profile in each direction is dominated by the perpendicular beam. At full power, with around 9 W in each beam, the trap is approximately 30 μK deep—corresponding to roughly half of the effective depth of the magnetic trap at the final frequency of the rf evaporation. The radial trapping frequency is $\omega_\rho = 2\pi \times 110$ Hz.

Following the sympathetic precooling stage, the dipole beams are suddenly (within < 1 ms) superimposed onto the trapped cloud. If the magnetic field is switched off, all of the atoms remain trapped radially, but spread out along the axial beam since the cross beam is not large enough in this direction to contain the entire cloud, which is about 2 mm long. For this reason, we leave the QUIC trap coils running at 50% to create a magnetic field curvature corresponding to approximately 11 Hz along the axis. After further evaporation, the cloud is small enough to fit into the cross beam and this field can be switched off. The radial confinement of the magnetic trap is removed automatically once the Feshbach magnetic field is applied, as described below.

X. FESHBACH MAGNETIC FIELD

Manipulating the scattering properties of ^{85}Rb using the 155 G Feshbach resonance is crucial in the final stages of cooling toward BEC, for two reasons: First, a BEC of significant size will not be stable far from the Feshbach resonance due to the negative background s -wave scattering length $a_{bg} = -443a_0$; second, as the cloud cools and its density increases, inelastic losses become increasingly severe. Although our precooling stage is at present limited by the number of refrigerant ^{87}Rb atoms in the magnetic trap, it has been shown that inelastic collisions preclude efficient cool-

ing of ^{85}Rb below about 5 μK at low field.¹⁵ Both of these issues may be overcome by tuning the scattering cross sections using the Feshbach resonance.

The coils we use to apply the Feshbach magnetic field are wound from the same wire as the quadrupole coils and are mounted in the same water-cooled aluminum block for mechanical and thermal stability. Each coil consists of 55 turns and runs at 13–15 A to produce a field in the range of 150–170 G. The coils are in Helmholtz configuration, producing a field parallel to that of the QUIC trap, and are centered on the QUIC trap minimum, which is 5 mm from the center of the quadrupole coils. This ensures optimal field uniformity over the trapped cloud: The field gradient at the atoms is well below 1 mG/cm and the cloud at 20 μK is only 2 mm in length.

Immediately after the optical dipole trap is switched on, the current in the Feshbach coils is ramped up over 500 ms to increase the magnetic bias field to 168 G. Since the radial curvature of the QUIC trap (which is now running at 50%) is inversely related to the bias field, this relaxes the radial confinement of the magnetic trap to the point where it is insufficient to support the atom cloud against gravity. Once at the full bias field, therefore, the atoms are held in a kind of hybrid trap, with the radial confinement provided purely by the axial dipole trapping beam and the axial confinement provided by the magnetic field curvature of the weakened QUIC trap.

An unfortunate side effect of this process is the splitting of the cloud by the resulting double-well potential as the bias field is ramped up. This effect is illustrated in Fig. 6. Initially, the atoms sit at the minimum of the potential created by the magnetic field and gravity (contour plot) and it is to this position that the optical dipole trap is aligned. However, as the bias field increases, the radial magnetic curvature decreases as described above and so the minimum of the mag-

netic plus gravitational potential moves downward (the gravitational sag is proportional to g/ω_p^2). This creates two distinct minima in the total potential seen by the atoms and as they separate, the cloud is split.

In addition to those lost by the splitting mechanism described above, some atoms leave as the bias field is ramped up because the dipole trap is shallower than the magnetic trap at the end of the rf evaporation. This can itself be considered evaporation since it is only the most energetic atoms that will spill over the edge of the trap. The combination of these two effects causes a loss of about 50% in number for each species, but keeps the phase space density of the sample constant during the transfer from QUIC trap to optical trap. We begin the final evaporation with 3×10^6 ^{85}Rb atoms and 2×10^7 ^{87}Rb atoms in the dipole trap at a temperature of 5 μK and a phase space density of 10^{-3} .

Before attempting evaporation to BEC in the optical trap, it is important to know the magnetic field produced by the Feshbach coils precisely. We calibrate the field by addressing the atoms with radiofrequency to drive transitions between the m_F sublevels. At the fields of interest (>100 G), the nonlinear contribution to the Zeeman splitting is large compared to the frequency width of the cloud so that the transitions are well separated from each other and only two states are coupled by the rf.

There are a number of ways in which transitions between m_F states can be observed. With either species, the substates can be spatially separated by applying a magnetic field gradient during the expansion, and then imaged separately. Alternatively, using ^{85}Rb the trapped $|F=1, m_F=-1\rangle$ state can be coupled to the untrapped $|F=1, m_F=0\rangle$ state. These atoms no longer feel the axial magnetic confinement and leave the trap along the axial beam, resulting in observable loss from the cloud. Interestingly, we also observe loss when coupling the $|F=2, m_F=-2\rangle$ and $|F=2, m_F=-1\rangle$ states in ^{85}Rb , despite the fact that both are low-field seeking and should remain in the trapping region. We attribute this to inelastic collisions in the $|m_F=-1\rangle$ state, which occur at a rate almost 6 times higher than in the $|m_F=-2\rangle$ state.²³

Once the frequency which drives transitions between neighboring Zeeman states is known, the Breit-Rabi equation²⁴ can be used to calculate the field at the center of the cloud. Using this technique we are able to calibrate the Feshbach bias field to within 50 mG.

XI. BOSE-EINSTEIN CONDENSATION

After single-species precooling in the QUIC trap, we are able to produce pure ^{87}Rb $|F=1, m_F=-1\rangle$ condensates containing 2×10^6 atoms in the QUIC trap after 7 s of evaporation in the optical dipole trap. The strength of the Feshbach bias field is not important in this case, and using Landau-Zener sweeps to affect rapid adiabatic passage between the Zeeman states we can create and study the behavior of spinor ^{87}Rb $|F=1\rangle$ condensates as a function of magnetic field.

Inelastic losses in ^{85}Rb $|F=2, m_F=-2\rangle$ are minimized at a magnetic field of 168 G.^{23,25} With the Feshbach coils set to this field, we proceed to evaporate the sample further by reducing the power in the dipole trap. This lowers the depth

FIG. 7. (Color online) (a) Absorption images taken after 20 ms of ballistic expansion showing the formation of a ^{85}Rb BEC as the depth of the crossed dipole trap is reduced. The elongation of the cloud along the tight trapping direction demonstrates the reversal of aspect ratio characteristic of BEC. (b) Optical depth profiles along the z direction, with fits showing the transition from a thermal Gaussian distribution, to a bimodal distribution and to a pure condensate.

of the trap and allows the most energetic atoms to spill out. With the axial magnetic field curvature still present, the evaporation is only 2D (atoms leave the trap only radially). At 168 G the s -wave scattering length of ^{85}Rb is near zero so this cooling is again sympathetic—although the isotope selectivity is less pronounced since both species see the same trapping potential.

After 5 s of evaporation, the cloud is small enough to be contained entirely within the cross beam of the dipole trap and the QUIC trap coils are switched off. At this point, the Feshbach field is slowly ramped from the inelastic minimum at 168 G to near 163 G, where the ^{85}Rb scattering length is $a=+200a_0$, suitable for the formation of a stable condensate. With a further 2 s of evaporation at this field, we observe the Bose-Einstein condensation of ^{85}Rb (Fig. 7). The final trapping frequencies of the crossed dipole trap are $\omega_{x,y,z}=2\pi \times (50, 57, 28)$ Hz. By varying the relative number of ^{85}Rb and ^{87}Rb initially loaded in the MOTs, we can optimize for maximum final phase space density in ^{85}Rb and choose whether to have a ^{87}Rb BEC of up to 10^6 atoms coexisting in the trap.

We typically reach the phase transition with 3×10^5 ^{85}Rb atoms and can create nearly pure condensates containing 4×10^4 atoms with a peak density of $2 \times 10^{13} \text{ cm}^{-3}$. The presence of a BEC is clearly indicated by the emergence of a bimodal density distribution and by the reversal of the aspect ratio of the cloud during expansion. The lifetime of the condensate in the crossed dipole trap is around 3 s.

FIG. 8. (Color online) Phase separation of ^{85}Rb and ^{87}Rb condensates in the immiscible regime. At $a_{85}=+200a_0$, the interspecies repulsion exceeds the intraspecies repulsion (miscibility parameter $\mu < 1$) and the clouds separate, producing a small, crescent-shaped ^{85}Rb condensate at the edge of a larger ^{87}Rb cloud.

When creating dual-species condensates, we also observe phase separation of the two isotopes due to the large interspecies scattering length $a_{85-87}=+213a_0$.²¹ At fields where the interspecies repulsion is greater than the intraspecies repulsion, the two fluids will be immiscible and the condensates will spatially separate. Because the ^{85}Rb scattering length may be changed using the Feshbach resonance, a dual-species ^{85}Rb – ^{87}Rb BEC exhibits tunable miscibility, an effect first observed by Papp *et al.*²⁶ The miscibility parameter μ is defined as

$$\mu = \frac{a_{85}a_{87}}{a_{85-87}^2}. \quad (2)$$

For the case of ^{85}Rb and ^{87}Rb , when $a_{85} > 460a_0$, $\mu > 1$ and the condensates are miscible. In our experiment, ^{85}Rb condensates are created with a scattering length $a=+200a_0$, corresponding to the immiscible regime when both species coexist in the trap. Figure 8 shows the density distribution of a partly condensed ^{85}Rb cloud in the presence of a large ^{87}Rb condensate. The thermal ^{85}Rb component is spread symmetrically over the trapping volume, while the condensed fraction is repelled by the ^{87}Rb condensate and forms a crescent-shaped cloud at the edge of the trap.²⁷

XII. COMPARISON

The first machine to make a ^{85}Rb BEC with tunable interactions was built at JILA, Colorado in 2000.⁵ It utilized careful and lengthy direct evaporation of ^{85}Rb in a weak magnetic trap with a large and variable bias field, and produced condensates containing up to 16 000 atoms. In 2006, the same group completed a new machine for making ^{85}Rb BECs by sympathetic cooling with ^{87}Rb , producing larger condensates (up to 80 000 atoms) with a shorter duty cycle.¹⁵ It is on this latest work that our method is largely based. However, although the procedure is similar, our setup has several important simplifications and a significantly shorter duty cycle.

Our vacuum system is more compact than that used in the work of Papp,¹⁵ at the expense of a reduced background lifetime due to the science cell being in direct line of sight of the 2D MOT. However, this is compensated by speedier

evaporation. The number of atoms collected in the 3D MOT is several times larger in our machine, but we do not perform any spin polarization, resulting in a similar number of atoms being loaded into the magnetic trap. We would expect to be able to increase the efficiency of our sympathetic cooling by spin polarizing the ^{87}Rb , which could boost the atom number by up to a factor of 3 (after optical pumping, the ^{87}Rb atoms would be distributed between the three Zeeman substates of the $|F=1\rangle$ manifold). This should allow higher phase space densities to be achieved prior to loading the optical trap.

Although our apparatus requires a 2D MOT in addition to the 3D MOT, it enables a very simple single-coil magnetic transfer from the MOT to the Ioffe–Pritchard trap. This is compared to the set of ten coils and mechanical translation stage in the JILA machine, creating a large set of parameters to be optimized to ensure efficient transport with minimal heating. The sympathetic cooling in the magnetic trap follows a qualitatively similar trajectory in the two machines, including the efficiency roll-off as the number of each species becomes comparable. Two notable differences are the initial temperature—200 μK here compared to 700 μK at JILA, leading to a higher final phase space density in our machine—and the length of the rf evaporation stage; 15 s in this work compared to 44 s.

The setup for our optical trap is greatly simplified compared to that at JILA as it does not require active stabilization, which also means that we can utilize the full power of our fiber laser. Also, our much larger beam waists (150–200 μm compared to 46 μm) remove the need for high quality, diffraction-limited optics. Despite lower trapping frequencies (our initial ω_ρ is an order of magnitude lower than in Ref. 15), we evaporate to BEC in 7 s, compared to 11.5 s in the JILA machine. An entire run of the experiment takes only 30 s. Although we can create similar-sized ^{87}Rb condensates of around 10^6 atoms, our ^{85}Rb BECs are about half the size of the largest created in the JILA machine, and we find the optimum scattering length for making BEC to be $a=+200a_0$, rather than $a=60a_0$ as found by Papp *et al.* Importantly, our final samples are confined purely optically, without an axial field curvature as in Ref. 15, making possible experiments with spinor ^{85}Rb BECs.

XIII. CONCLUSION AND OUTLOOK

We have presented details of our machine for creating BECs of ^{85}Rb with tunable interactions, which has only been achieved by one other group. We employ sympathetic cooling with ^{87}Rb , first in a magnetic trap and then in a crossed optical dipole trap, to make ^{85}Rb BECs containing 4×10^4 atoms. Our apparatus has several important simplifications compared to the ^{85}Rb BEC machine at JILA. Although our ^{85}Rb BECs are currently smaller than those created in that machine, we have thus far seen increases in the number of atoms in the Ioffe–Pritchard trap translate directly into larger condensates, although inelastic collisions in ^{85}Rb will impose a limit on this increase. The machine presented here confines ^{85}Rb BECs in a purely optical potential and is also

able to produce large ^{87}Rb condensates, giving us the flexibility to perform a variety of experiments involving dual-species and spinor BECs.

ACKNOWLEDGMENTS

The authors gratefully acknowledge B. Buchler for experimental assistance, G. Dennis and J. Hope for help with numerical simulations, and N. Hinchey, S. Grieves, P. Tant, P. McNamara, and N. Devlin for technical assistance. This work was supported by the Australian Research Council Centre of Excellence for Quantum-Atom Optics.

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