Sympathetic cooling of 85Rb and 87Rb

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(Received 22 December 2000; published 10 July 2001)

We demonstrate sympathetic cooling between magnetically trapped ⁸⁵Rb and ⁸⁷Rb atoms. Up to 6×10^6 ⁸⁵Rb atoms are cooled via elastic collisions with a reservoir of 8×10^{8} ⁸⁷Rb atoms. The temperature of the binary atomic ensemble is adjusted by using radio-frequency evaporation of 87Rb. Temperatures as low as *T* $=100$ nK have been realized in the binary ensemble. Through a cross-dimensional mixing approach we have been able to measure the interspecies ⁸⁷Rb-⁸⁵Rb scattering cross section $\sigma = 1.52(34) \times 10^{-15}$ m².

DOI: 10.1103/PhysRevA.64.021402 PACS number(s): 32.80.Pj, 03.65.Nk, 03.75.Fi, 05.30.Jp

The advent of powerful cooling schemes for neutral atoms has enabled ground breaking experiments in physics. Laser cooling and evaporative cooling have proved to be the crucial ingredients in the realization of Bose-Einstein condensation in dilute atomic gases $|1|$. Laser cooling can be applied to most atomic species, as long as the required laser wavelengths are available. The conditions under which evaporative cooling can be used to chill a trapped atomic gas are more stringent. This cooling method requires a combination of high atomic density, large atom number, and favorable characteristics of the atomic collisions $[2]$. Only a very limited range of atomic species has therefore been evaporatively cooled to ultralow temperatures. Sympathetic cooling may provide a way out of this dilemma. It relies on cooling a certain atomic or even molecular species by bringing this species into thermal contact with a cold reservoir of atoms. In the ultracold regime, sympathetic cooling has been demonstrated for different spin states of one atomic species $|3,4|$ and is now being applied to boson-fermion mixtures $[5]$.

In this Rapid Communication we report on the realization of sympathetic cooling between two different bosonic isotopes, namely between the two rubidium isotopes ⁸⁷Rb and 85 Rb. An initially laser cooled sample of less than $10^{7} {}^{85}$ Rb atoms is sympathetically cooled via elastic collisions with an evaporatively cooled cloud of ⁸⁷Rb atoms. Temperatures as low as 100 nK are reached in the mixed isotope gas. We have also been able to produce $87Rb$ condensates in the presence of an ultracold thermal cloud of ⁸⁵Rb atoms.

Due to its favorable scattering properties, the $87Rb$ isotope can easily be cooled by evaporation and it is therefore ideally suited to act as a refrigerator for other atomic species. In contrast, the 85 Rb isotope is difficult to cool by evaporation since its *s*-wave scattering cross section vanishes for collision energies of $E \approx k_B \times 375$ μ K (k_B : Boltzmann's constant). Evaporative cooling of $85Rb$ has only been possible so far in the lower hyperfine ground state by exploiting a Feshbach resonance to tune the scattering length $[6]$. By mixing both isotopes we achieve almost an order of magnitude shorter cooling times than those reported before $[6]$. The cooling efficiency for the 85 Rb atoms is boosted by the interactions between the two atomic species. Since only ⁸⁷Rb atoms are selectively evaporated, the method is not sensitive to the initial number and density of the ⁸⁵Rb sample.

The crucial aspect of sympathetic cooling is the interaction between the different atomic species. We have quantitatively measured the scattering cross section for collisions between 87 Rb and 85 Rb atoms and obtain a value for the cross section that is two times larger than the scattering cross section for collisions between $87Rb$ atoms, in good agreement with theoretical predictions. Sympathetic cooling of atomic mixtures may also pave the way towards degenerate bosonboson or boson-fermion $\begin{bmatrix} 5 \end{bmatrix}$ quantum gases of different atomic species. Both systems provide a unique testing ground for the physics of interacting quantum systems with a rich variety of new physical phenomena $[7-12]$ to be explored.

In the experiment, the two rubidium isotopes are first collected in a vapor cell magneto-optical trap (MOT) with large beams (apertured to a diameter of 40 mm). The main cooling light for both isotopes is derived from a master-oscillator power amplifier (MOPA) diode laser system. Up to 100 mW of this laser power is available after an optical fiber at the MOT. The MOT loading process is divided into two stages. During the first 15 s the MOPA system exclusively provides cooling light on the $|F=2\rangle \rightarrow |F=3\rangle$ ⁸⁷Rb transition (*F* denotes the total angular momentum of the atom). Then 10% of the MOPA laser power is used to additionally collect ⁸⁵Rb atoms over the next 900 ms. This is achieved by injecting another laser running on the $|F=3\rangle \rightarrow |F=4\rangle$ ⁸⁵Rb transition into the amplifier $[13]$. The laser frequencies are red detuned by 18 MHz with respect to the $|F=2\rangle \rightarrow |F=3\rangle$ ⁸⁷Rb transition and by 7 MHz with respect to the $|F=3\rangle$ \rightarrow $|F=4\rangle$ ⁸⁵Rb transition. After an 8-ms optical molasses phase the isotopes are optically pumped into the weak-fieldseeking states $|F=2,m_F=2\rangle$ (⁸⁷Rb) and $|F=3,m_F=3\rangle$ $(85Rb)$, where m_F denotes the magnetic sublevel. A magnetic quadrupole potential is then rapidly turned on in order to confine the atoms. Using a series of computer activated magnet coils, the quadrupole potential is moved over 33 cm into an UHV glass cell $[14]$, where the magnetic quadrupole potential is converted into a Ioffe-type potential using the quadrupole-Ioffe configuration trap mechanism [15]. The trapping frequencies of the final magnetic trap are $\omega_z = 2\pi$ \times 24 Hz in the axial and ω_0 =2 π ×219 Hz in the radial direction.

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FIG. 1. Potentials of the trapped atomic states $|F=2,m_F=2\rangle$ of ⁸⁷Rb and $|F=3,m_F=3\rangle$ of ⁸⁵Rb (thick line) and their neighboring magnetic sublevels $|F=2,m_F=1\rangle$ (⁸⁷Rb) (dashed line) and $|F=3,m_F=2\rangle$ (⁸⁵Rb) (thin line). In a mixed cloud at thermal equilibrium the radio frequency, indicated by the arrows, predominantly evaporates 87 Rb atoms, since the energy at which the 85 Rb atoms are removed is higher than the energy at which ⁸⁷Rb atoms are removed. Both species experience the same confining potential and overlap completely.

Approximately 8×10^{8} ⁸⁷Rb and 3×10^{6} ⁸⁵Rb atoms are present in the final magnetic trap. The two atomic species are initially prepared in states of equal magnetic moment, so that they experience the same confining potential. However, the different Landé g_F factors of the two hyperfine states (g_F $=1/2$ for ⁸⁷Rb and $g_F=1/3$ for ⁸⁵Rb) lead to an unequal splitting between neighboring magnetic sublevels. For the same magnetic field, the energy splitting due to the linear Zeeman effect between neighboring magnetic sublevels in the $|F=2\rangle$ ⁸⁷Rb manifold is 50% larger than in the $|F=3\rangle$ $85Rb$ manifold (see Fig. 1). Radio-frequency evaporation therefore predominantly removes hot ${}^{87}Rb$ atoms. The ${}^{85}Rb$ atoms remain almost unaffected as long as they thermalize with the cooler ⁸⁷Rb atoms on a time scale that is faster than that given by the radio-frequency sweep. Adjusting the time scale of the evaporation process to the ⁸⁵Rb thermalizaton time therefore allows us to be highly isotope selective in the removal of hot atoms. A perfectly isotope selective evaporation could be realized by using microwave photons tuned to the ⁸⁷Rb hyperfine energy splitting.

The number and temperature of the atoms during the evaporation process are measured using absorption imaging after an adjustable ballistic expansion time. In a single run of the experiment we are able to detect both atomic species by using the following procedure. We first apply a $100 - \mu s$ pulse of light tuned to the ⁸⁵Rb $|F=3\rangle \rightarrow |F=4\rangle$ transition and 2 ms later we apply a 100- μ s pulse of light tuned to the ⁸⁷Rb $|F=2\rangle \rightarrow |F=3\rangle$ transition. During 18 s of radio-frequency evaporation we are able to cool the mixed atomic sample to temperatures well below 1 μ K. The number of atoms of both isotopes are displayed in Fig. 2 for different temperatures below 60 μ K along the evaporation trajectory. While cooling the sample to a temperature of 2 μ K, the number of ⁸⁷Rb atoms has decreased by roughly three orders of magnitude, whereas the number of ⁸⁵Rb atoms has decreased only by a factor of 3. We have furthermore observed that all ${}^{85}Rb$ atoms are removed in the evaporation process if there are no

FIG. 2. Number of atoms versus temperature along the evaporation trajectory of the mixed species ensemble. The filled circles denote the number of 87Rb atoms, while the open circles denote the number of ⁸⁵Rb atoms.

 $87Rb$ atoms present in the magnetic trap. This is caused by the fact that thermalization of the pure $85Rb$ cloud is suppressed by the vanishing *s*-wave collision cross section of ⁸⁵Rb for collision energies of $E \approx k_B \times 375$ μ K [16]. Therefore all atoms are removed during evaporation. This property is responsible for the severe difficulties associated with evaporatively cooling ${}^{85}Rb$ [6]. In a cloud of ${}^{85}Rb$ atoms adiabatic compression results in a dramatic decrease of the elastic collision rate, as the scattering cross section decreases by orders of magnitude with increasing temperature $[16]$. For our case the initial 85Rb intraspecies collisions are completely negligible and we therefore conclude that interspecies collisions play a crucial role in thermalizing the $85Rb$ cloud of atoms at higher temperatures during the evaporation process. Indeed, Burke *et al.* [16] predict an almost constant and large interspecies collision cross section between the $87Rb-85Rb$ isotopes for temperatures up to 1 mK. For temperatures below 1 μ K the evaporative cooling process becomes inefficient, as can be seen from the increased slopes of the $\log N$ vs $\log T$ graph of Fig. 2. This can partly be attributed to the increasing number of inelastic events predicted for collisions of atoms in the $|F=2,m_F=2\rangle$ (⁸⁷Rb) and $|F=3,m_F=3\rangle$ (⁸⁵Rb) states and also to the fact that the number of atoms in our 87 Rb cooling medium are approaching the number of 85 Rb atoms so that we cannot sustain a completely isotope selective evaporation. Furthermore, the triplet scattering length in $85Rb$ has a large and negative value, and allows only condensates with a very small number below 100 atoms to be produced. Presently, such a low number of atoms cannot be detected unambiguously in our setup. Nevertheless, we have been able to produce $87Rb$ condensates of up to $10⁵$ atoms coexisting with a thermal cloud of ⁸⁵Rb atoms.

In order to characterize the interactions between the two atomic species we have measured the ${}^{87}Rb-{}^{85}Rb$ interspecies scattering cross section through a cross-dimensional mixing approach [18]. At temperatures of 6.6 μ K, samples of atoms are prepared with 3.3×10^5 ⁸⁵Rb atoms and 1.5×10^5 ⁸⁷Rb atoms. The mixed atomic ensemble is then heated parametrically by modulating the radial trapping frequency at twice its resonance frequency for 50 ms. Energy is thereby added to the radial axes of the atomic ensemble. This leads to a nonequilibrium situation with a radial temperature of the cloud larger than the axial temperature of the cloud. In this nonequilibrium state the cloud of atoms has a smaller aspect ratio $\Delta z/\Delta \rho$ than in the equilibrium situation, where $\Delta z/\Delta \rho = \omega_o/\omega_z$. Here $\Delta \rho$ and Δz are the radial and axial extensions of the trapped cloud of atoms. The subsequent relaxation of the aspect ratio of the trapped ensemble of ${}^{87}Rb$ atoms towards the equilibrium value due to elastic collisions is then monitored versus time. In order to verify that this relaxation of the aspect ratio is in fact due to elastic collisions and not due to anharmonic mixing between the motion of the atoms along the three trapping axes, we have verified that the relaxation rate scales with the density of our trapped ensembles. We have also checked that the change in density of both atomic species due to loss of atoms or heating rates is below 2% for a thermalization time of 1 s. Furthermore, the total energy of the atoms remains constant within our measurement time. We obtain the interspecies scattering cross section by comparing the time constant of the relaxation process of $87Rb$ in the presence of $85Rb$ with the time constant of the relaxation of a pure cloud of ⁸⁷Rb.

The relaxation rate Γ_{87}^{mix} for the ⁸⁷Rb component of the mixed atomic gas can be described by

$$
\Gamma_{87}^{mix} = \alpha^{-1} \Gamma_{87}^{el} + \tilde{\alpha}^{-1} \Gamma_{87-85}^{el}
$$

=
$$
\alpha^{-1} \bar{n}_{87} \sigma_{87} \bar{\upsilon}_{87} + \tilde{\alpha}^{-1} \bar{n}_{85} \sigma_{87-85} \bar{\upsilon}_{87-85}.
$$
 (1)

Here Γ^{el} denotes the elastic collision rate, α and $\tilde{\alpha}$ are constants relating the elastic collision rate to the crossdimensional mixing time, \overline{n} is the effective density of the atoms $\overline{n} = \int n^2(\mathbf{r})d^3r/\int n(\mathbf{r})d^3r$, σ is the elastic collision cross section, and \overline{v} is the average relative velocity of the atoms. The respective indices 87, 85, and 87–85 indicate whether values for the single atomic species or the mixed species have to be inserted into the formula. The average relative velocity between 87 Rb atoms is given by \bar{v}_{87} $=4\sqrt{k_B T/\pi m_{87}}$, whereas the average velocity between ⁸⁷Rb and 85 Rb atoms can be expressed by \bar{v}_{87-85} $= \sqrt{8k_B T/\pi} \sqrt{(m_{87}+m_{85})/(m_{87}m_{85})}$ [17].

Through Monte Carlo simulations of the collision process and an analytical model, the constant α for intraspecies collisions has been found to be $\alpha \approx 2.7$ [18,19]. For the interspecies ⁸⁷Rb⁻⁸⁵Rb collisions we expect $\tilde{\alpha}$ to be slightly smaller, as the ⁸⁵Rb cloud is almost thermalized already after a short time due to its large scattering cross section. Therefore energy is more efficiently redistributed for collisions of $87Rb$ atoms with these thermalized $85Rb$ atoms. Using an analytical model based on Boltzmann's equation and a multispecies Monte Carlo simulation of the thermalization process, we find that $\tilde{\alpha} \approx \alpha/1.2(1)$ for our measurement parameters. In the analysis we have also assumed dominant *s*-wave scattering for ⁸⁵Rb-⁸⁷Rb interspecies and ⁸⁷Rb intraspecies collisions.

FIG. 3. Relaxation of the aspect ratio of an ensemble of $87Rb$ atoms after parametric heating of the radial axes of the trapped atom cloud. (a) Closed circles denote the data for a pure $87Rb$ ensemble of 1.5×10^5 atoms, while the open circles are the datapoints for the ⁸⁷Rb ensemble in the presence of 3.3×10^5 ⁸⁵Rb atoms. (b) Close-up of the ${}^{87}Rb$ data in the presence of ${}^{85}Rb$. The solid and broken lines are fits to the data assuming an exponential decay of the aspect ratio towards the equilibrium value ω_p / ω_z . Each datapoint is an average over eight measurements and the error bars display the corresponding statistical errors. The temperature of the ensemble during the measurement was $T \approx 7$ μ K. The 1/*e* time constant for the relaxation of the pure ⁸⁷Rb ensemble is $1/\Gamma_{87}$ $=1160(100)$ ms. For the ⁸⁷Rb ensemble in the presence of ⁸⁵Rb it is $1/\Gamma_{87}^{mix}$ = 190(20) ms.

Assuming Gaussian distributions for the density of the atoms, one can find a simple expression for the interspecies scattering cross section σ_{87-85} . By comparing the relaxation rate of ⁸⁷Rb in the mixed isotope ensemble to the relaxation rate of a pure ⁸⁷Rb ensemble at the same temperature and with the same number of ${}^{87}Rb$ atoms, one obtains

$$
\sigma_{87-85} = \sqrt{2} \left\{ \frac{\Gamma_{87}^{mix}}{\Gamma_{87}} - 1 \right\} \left(1 + \frac{m_{87}}{m_{85}} \right)^{-1/2} \frac{N_{87}}{N_{85}} \frac{\tilde{\alpha}}{\alpha} \sigma_{87}.
$$
 (2)

Using the experimentally measured relaxation times from Fig. 3 with $N_{85}/N_{87}=2.2(2)$, we obtain $\sigma_{87-85}=1.92(40)$ $\times \sigma_{87}$.

The single species ⁸⁷Rb cross section σ_{87} for atoms in the $|F=2,m_F=2\rangle$ state can be expressed through the ⁸⁷Rb trip-

let scattering length a_T^{87} as $\sigma_{87} = 8 \pi (a_T^{87})^2$. This triplet scattering length is well known through photoassociation spectroscopy, collision cross-section measurements and the location of the 85 Rb Feshbach resonance [20]. Since the rubidium isotopes are distinguishable from each other, the interspecies collision cross section is related to the interspecies triplet scattering length $a_{T_a}^{87-85}$ by $\sigma_{87-85} = 4 \pi (a_T^{87-85})^2$. Using a recent value for $a_T^{87} = 106(4) a_0$ (*a*₀ being the Bohr radius) [20,21], we obtain $\sigma_{87-85} = 1.52(34) \times 10^{-15}$ m² and $|a_T^{87-85}| = 208(23)a_0$ for the absolute value of the interspecies triplet scattering length. This is in good agreement with recent theoretical predictions of a_T^{87-85} = 213(7)*a*₀ (theory) $\lceil 21 \rceil$.

In the 85 Rb and 87 Rb mixture the atom-atom interactions can be tuned via Feshbach resonances. The system is therefore ideally suited to study the behavior of two species Bose-Einstein condensates. In ⁸⁵Rb a Feshbach resonance in the $|F=2,m_F=-2\rangle$ state [6] allows one to tune the scattering

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length between 85 Rb atoms. In addition, there is a theoretical prediction for an interspecies Feshbach resonance in collisions between ⁸⁷Rb atoms in the $|F=1, m_F=-1\rangle$ state and ⁸⁵Rb atoms in the $|F=2,m_F=-2\rangle$ state that should allow one to adjust the interspecies scattering length $[16]$.

In conclusion, we have demonstrated sympathetic cooling between 87 Rb and 85 Rb atoms in a magnetic trap. The severe difficulties in evaporatively cooling $85Rb$ due to the zero *s*-wave scattering cross section of ⁸⁵Rb at collision energies of $E \approx k_B \times 375$ μ K have been circumvented by exploiting interspecies collisions between ${}^{87}Rb$ and ${}^{85}Rb$ atoms. We have furthermore been able to measure the collision cross section between different atomic species at ultralow temperatures.

We would like to thank Anton Scheich for experimental assistance during the construction of the experiment. We also acknowledge support by the Deutsche Forschungsgemeinschaft.

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