Steady-State Magneto-Optical Trap with 100-Fold Improved Phase-Space Density

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Laser cooled and trapped atoms are at the core of most ultracold quantum gas experiments [1], state-of-the-art clocks [2], and sensors based on atom interferometry [3]. Today, these devices typically operate in a time-sequential manner, with distinct phases for sample preparation and measurement. For atomic clocks a consequence is the need to bridge the dead time between measurements using a secondary frequency reference, typically a resonator. This introduces a problem known as the Dick effect [4], in which the sampling process inherent to a clock’s cyclic operation down-converts or aliases high frequency noise from the secondary reference into the signal band, thus degrading performance [5]. Recently, a new generation of atomic clocks using degenerate atoms in a three-dimensional optical lattice has been demonstrated using Sr [6]. To reach the potential of such a clock, it will be necessary to overcome the Dick effect, which can be achieved by reducing the dead time and/or by creating vastly improved secondary references. Our steady-state MOT can lead to significant advances in both directions. It approaches the high flux and low temperature requirements needed for a steady-state clock, which would completely eliminate the Dick effect. Furthermore, our MOT is created under conditions compatible with the creation of degenerate samples or an atom laser [7,8]. This would be the ideal source for a secondary frequency reference based on superradiant lasing, which is expected to outperform current references [9–13]. Our source and a future atom laser based on it might also be valuable for atomic inertial sensors [8]. Improved clocks and inertial sensors will allow tests of fundamental physics [14] or be suitable for gravitational wave astronomy [3,15–17].

Over the years, many creative approaches have honed laser cooling to produce pulsed samples of ever increasing phase-space density (PSD) [18–29]. Pulsed MOTs using $^{88}$Sr have demonstrated phase-space densities of $10^{-2}$ [30] while atoms held in dipole traps recently reached degeneracy [7,31]. Despite the exquisite performances, these techniques suffer from extremely small capture velocities. As a consequence, atoms must first be captured and precooled, and thus these techniques have only been used as part of time-varying sequences.

Several continuous high PSD MOT schemes have been demonstrated mostly based on bichromatic MOTs using alkaline earth atoms [32,33]. The most successful reached a steady-state PSD of $1.2 \times 10^{-5}$ [34]. This scheme used a MOT on a broad linewidth transition to capture atoms which then leak into a metastable state cooled by a MOT on a narrow transition. Narrow-line MOTs fed by a 2D MOT or Zeeman slower on the broad transition have been demonstrated for Yb and Er although steady-state PSDs are not measured [35,36]. Another approach is the dark SPOT MOT [37], which creates a central spatial region of reduced laser interaction. Adding a further steady-state trapping and cooling stage to a MOT can increase the PSD substantially [38–40], with a steady-state PSD of $\sim 4 \times 10^{-4}$ reached for Cr atoms at a temperature of $\sim 50 \mu$K [40].

In this Letter, we demonstrate a $^{88}$Sr MOT at a temperature of $\sim 2 \mu$K and a steady-state phase-space density of $1.3(2) \times 10^{-3}$, 2 orders of magnitude higher than reported for previous steady-state MOTs [34]. Combining our MOT with techniques such as Refs. [7,38–42] promises yet higher PSDs and potentially a steady-state BEC and atom laser. Our result is achieved by flowing atoms through a series of spatially separated cooling stages as illustrated in Fig. 1. First we use the high capture velocity of the broadbandwidth, “blue,” $^1S_0 - ^3P_1$ transition (30 MHz linewidth, 461 nm wavelength) in several stages to slow and cool atoms to mK temperatures, finishing with a 2D blue MOT. Next, we capture the atoms in a 3D MOT using the narrow linewidth, “red,” $^1S_0 - ^3P_1$ transition (7.4 kHz linewidth, 689 nm wavelength), which can reach temperatures close to
the recoil limit [30]. We operate the two MOTs in separate chambers to avoid heating of the 3D red MOT by blue photons scattered from surfaces and from fluorescing atoms in the 2D blue MOT. The transfer of atoms between chambers is ensured by two key ingredients: first, the atomic beam from the 2D blue MOT is collimated by a red optical molasses and second, the atoms are slowed in the second chamber by a hybrid slower + MOT configuration operated on the low capture velocity red transition. We show that this approach allows the red MOT to produce clouds with unprecedented phase-space densities for a steady-state apparatus. Furthermore, we show that the red MOT location is sufficiently protected to form BECs even with all the blue cooling stages operating.

The path of atoms through the setup begins with a high-flux atomic beam source adapted from our previous design [43–45]. In brief, this source is composed of an oven similar to Ref. [46], followed by a transverse cooling stage and a Zeeman slower, both using laser cooling on the broad-linewidth blue transition. A 2D blue MOT [47–49], whose nonconfining axis is oriented in the direction of gravity, is located approximately 5 cm after the exit of the Zeeman slower [see Fig. 1(a)]. This MOT has a loading rate of $2.66(16) \times 10^9$ $^{88}$Sr atoms/s (measured by absorption imaging) and cools atoms to about 1 mK in the radial ($xz$) plane. To prevent atoms from escaping upward, a pair of downward propagating blue “plug” beams are placed symmetrically to each other at an $8^\circ$ angle from the $y$ axis, increasing the flux by around 30%.

The high phase-space density MOT must be operated on the narrow-linewidth red transition while being protected against photons from the broad-linewidth blue transition [32]. In order to ensure such protection, we position the red MOT in a separate chamber, 41 cm below the 2D blue MOT. The two chambers are separated and baffled by a set of four stacked 1-inch absorptive neutral density filters (Thorlabs NE60A). These filters are separated by 20 mm and have an 8 mm diameter center hole allowing atoms to pass.

Upon exiting the 2D blue MOT, atoms have a radial velocity of about 0.5 m/s, an average downward velocity of 3 m/s [45], and they fall accelerated by gravity towards the bottom chamber. Radial expansion during the 185 mm drop to the bottom of the baffle section would give a transfer efficiency less than 2%. Accelerating the atoms downward is out of the question, since we can only use the low capture velocity red transition in the bottom chamber. Instead, we radially cool atoms emerging from the 2D MOT using a 2D molasses operating on the red $^1S_0 \rightarrow ^3P_1 \pi$ transition, which is insensitive to the spatially varying magnetic field. To compensate the Doppler broadening of the atomic transition, which is more than 100 times the 7.4 kHz natural linewidth, it is necessary to modulate the frequency of the molasses laser beams forming a frequency comb spanning from 30 to 750 kHz to the red of the transition, with 25 kHz spacing [44]. This technique is used on all our red-transition laser beams. To prevent the two red $^1S_0 \rightarrow ^3P_1 \sigma$ transitions towards the $m_J = \pm 1$ states from hindering the radial slowing process, we produce a Zeeman shift of about 3 MHz by applying a bias magnetic field of 1.4 G in the vertical direction. Such a small field doesn’t disturb the operation of the 2D blue MOT. This molasses reaches steady state after a few 10 ms, easily provided by four horizontal molasses beams with $1/e^2$ diameters of 45.6 mm along the $y$ axis.

After a 41 cm fall from the top to the bottom chamber, the atomic beam has a measured downward velocity distribution peaked at 4 m/s. Our protection scheme necessitates slowing and capturing these falling atoms using only the red transition. However, the small scattering rate on this line allows a maximum deceleration of only $\sim 16g$, where $g$ is earth acceleration. To overcome this extreme limitation we implement a hybrid slower + MOT.

Our first hybrid setup configuration labeled “Red MOT I” [see Fig. 1(a)] uses a magnetic quadrupole field centered 23 cm directly below the bottom baffle between chambers. The gradients are $(0.55, 0.35, 0.23)$ G/cm in the ($x, y, z$) directions. Horizontally propagating laser beams in the $x, z$ axes are placed in a MOT configuration and provide radial cooling and confinement. On the $y$ axis, a single, upward propagating beam is used, with circular polarization as needed for the MOT. Because of the weakness of the

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**FIG. 1.** (a) Schematic of our setup, showing the main cooling stages, and the position of the three red MOT configurations (see text). (b) Electronic level scheme of strontium, with the blue and red transitions used for laser cooling. (c) In situ absorption picture of a steady-state $^{88}$Sr MOT with a PSD of $1.3(2) \times 10^{-3}$ [Red MOT II(b) configuration].
transition this upward propagating beam and gravity are sufficient to confine atoms in the vertical direction without a downward propagating MOT beam [50]. The vertical beam is directed slightly to the side of the baffle onto the lowest neutral density filter, to prevent it from affecting the cooling processes in the top chamber. This beam is converging to exert a restoring force towards the beam center during the slowing process [51].

Upon reaching the second chamber, atoms enter the region illuminated by the circularly polarized upward-propagating beam, whose frequency is set to the red of the \( ^1S_0 - ^3P_1 \) transitions. We now describe the slowing process using an upward pointing quantization axis; see Fig. 2(a). The Doppler shift \( \delta_{\text{Doppler}}(v) \) of atoms with a downward velocity \( v \) brings them into resonance with the \( \sigma^+ \) transition from \( ^1S_0 \) to the state \( m_f = +1 \) of \( ^3P_1 \). As radiation pressure slows atoms down, \( \delta_{\text{Doppler}} \) diminishes, which is partially compensated by the spatial variation of the magnetic field, following the principle of a Zeeman slower. The narrow linewidth of the red transition does not allow for a slowing process robust against magnetic and laser intensity fluctuations, so we modulate the laser to allow for a slowing process robust against magnetic and laser intensity fluctuations, so we modulate the laser frequency \( \nu_L \) with a span of \( \Delta \nu_L = 4.05 \text{ MHz} \), as used for example in “white light” slowing [52]. If atoms are successfully slowed and reach the region below the quadrupole field center, \( \delta_{\text{Doppler}} \) is small and atoms are resonant with the laser as in a standard broadband narrow-line MOT [30]. Note that for the experimental configurations used the angle between the local magnetic field and the beam direction can be big, leading to significant additional absorption on the \( x \) transition during the slowing process.

We numerically model this hybrid setup by evolving classical atomic trajectories, first in an idealized 1D geometry with a linear vertical magnetic field gradient and a uniform circularly polarized vertical beam, then using a Monte Carlo approach in a realistic 3D geometry including all beams and details of the magnetic fields [45].

The behavior of a falling atom can be obtained by analyzing the deceleration it experiences in dependence of time when dropped into the slower + MOT region with various starting velocities. The idealized 1D results shown in Fig. 2(b) are qualitatively confirmed by the more realistic 3D model results shown in Fig. 2(c). With these simulations we estimate a maximum capture velocity of around 6 m/s for the hybrid slower + MOT setup, which is compatible with the measured velocity distribution of the atomic beam produced by the 2D blue MOT [45].

The characteristics of the hybrid Red MOT I are summarized in Table I. The loading rate gives an estimated transfer efficiency of 19% between the two chambers. Unfortunately, the high power broadband beams needed for the hybrid setup limit the PSD to \( 2.8(1.2) \times 10^{-9} \), low compared to what can be achieved with a pure red MOT geometry in a time-varying sequence [30]. Moreover, with direct line-of-sight to a bright fluorescing blue MOT, this configuration does not provide protection against heating from blue photons sufficient to produce a steady-state BEC [45].

For these reasons, we implement a second MOT configuration, “Red MOT II(a),” located 3 cm in the \( z \) direction away from the Red MOT I position [see Fig. 1(a)]. This is achieved by adding another 5-beam geometry MOT (4 beams along \( x \) and 1 beam along \( y \)), whose beams make a smooth connection with the Red MOT I beams, and by displacing the center of the quadrupole field to the intersection of these new beams. Along the \( z \) axis the two additional beams are implemented as an 8 mm diameter low-intensity core within hollow 48 mm diameter Red
applying a 3 s evaporative cooling sequence, we produce an optical dipole trap in a time sequential manner and horizontal axes, respectively. By loading this MOT into MOT II(a) where they are confined and further cooled. The Red MOT I beams, then continuously pushed to the Red second chamber are decelerated by the hybrid slower plus MOT I beams. In this configuration, atoms entering the effective scattering rate is just enough to hold atoms against gravity, while in the surrounding region an effective atomic flux is suddenly stopped is 1.95(6) s, which is 2%, an order of magnitude lower than for Red MOT II(a). The transfer efficiency from the top chamber to this MOT is 2%, an order of magnitude lower than for Red MOT I. Simulations suggest significant losses may be attributed first to increased bouncing of atoms in the hybrid slower when the \( \pi \) transition is dominant, and second to atom trajectories intersecting with mirrors inside the vacuum chamber.

We also produce a Red MOT II(a) using the much less abundant \(^{88}\text{Sr}\) isotope, which is particularly suited to produce quantum degenerate gases, owing to its favorable scattering properties. This \(^{88}\text{Sr}\) MOT contains up to 9.0(2) \( \times 10^6 \) atoms at temperatures of 1.5(3) \( \mu \)K and 3.4(1.1) \( \mu \)K in the vertical and horizontal axes, respectively. By loading this MOT into an optical dipole trap in a time sequential manner and applying a 3 s evaporative cooling sequence, we produce \(^{88}\text{Sr}\) BECs of 3.0(2) \( \times 10^5 \) atoms.

The protection provided by the dual chamber and baffle system against photons from the broad-linewidth blue transition is determined by comparing the BEC lifetime at the location of the Red MOT II(a) with and without the 2D blue MOT operating. The atomic flux is disabled by turning off the red transition beams during these measurements. The BEC lifetime in the dipole trap is 3.0(4) s without blue light and 2.7(2) s with blue light. Lifetimes were strongly limited by one-body collisions due to poor vacuum quality. These measurements confirm significant protection from blue photons thanks to our two-chamber design, making our system suitable for experiments aimed at developing a steady-state source of degenerate quantum gas.

Finally, “Red MOT II(b)” is a configuration designed to optimize phase-space density at the expense of transfer efficiency. This MOT reaches a steady-state PSD of 1.3(2) \( \times 10^{-3} \) for \(^{88}\text{Sr}\), which is 2 orders of magnitude higher than demonstrated in previous steady-state MOTs [34] [see Fig. 1(c) and Table I]. Red MOT II(b) is the same as Red MOT II(a) except that we reduce the bandwidth of the MOT II beams to cover only −40 to −200 kHz detuning, resulting in three improvements. First, the smaller detuning of −40 kHz compresses the MOT, second, the reduced bandwidth reduces the total beam intensity required, and third, by ending the spectrum well before the photoassociation line located at −435 kHz detuning [53], we reduce losses due to molecule formation. The transfer efficiency from the top chamber to this MOT is 0.2%, an order of magnitude lower than for Red MOT II(a). The 1/e lifetime of Red MOT II(b) measured after the atomic flux is suddenly stopped is 1.95(6) s, which is significantly smaller than the 4.53(6) s lifetime of Red MOT I. This reduction is due to two-body light-assisted collisions, which ultimately limits the maximum density achievable [53]. Both these losses and the need to compensate for gravity set the limits on the PSD achievable with this MOT configuration.

### Table I. Characteristics of the three Red MOT configurations. All uncertainties stated in this letter are taken as ±2\( \sigma \) from the fitted data.

<table>
<thead>
<tr>
<th></th>
<th>Red MOT I</th>
<th>Red MOT II(a)</th>
<th>Red MOT II(b)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flux [^{88}\text{Sr}/s]</td>
<td>5.1(7) ( \times 10^8 )</td>
<td>5.3(4) ( \times 10^7 )</td>
<td>5.3(9) ( \times 10^6 )</td>
</tr>
<tr>
<td>Temperature ( x ) ( \mu )K</td>
<td>3.7(3)</td>
<td>1.9(1)</td>
<td>1.9(1)</td>
</tr>
<tr>
<td>Temperature ( y ) ( \mu )K</td>
<td>20(5)</td>
<td>1.9(1)</td>
<td>1.9(1)</td>
</tr>
<tr>
<td>Temperature ( z ) ( \mu )K</td>
<td>26(7)</td>
<td>2.8(2)</td>
<td>1.9(1)</td>
</tr>
<tr>
<td>Width ( \sigma_x ) ( \mu )m</td>
<td>385(4)</td>
<td>352(3)</td>
<td>150(2)</td>
</tr>
<tr>
<td>Width ( \sigma_y ) ( \mu )m</td>
<td>725(61)</td>
<td>192(3)</td>
<td>88(1)</td>
</tr>
<tr>
<td>Width ( \sigma_z ) ( \mu )m</td>
<td>2086(41)</td>
<td>528(3)</td>
<td>247(3)</td>
</tr>
<tr>
<td>Atom number [^{88}\text{Sr}]</td>
<td>2.54(10) ( \times 10^6 )</td>
<td>1.71(5) ( \times 10^6 )</td>
<td>2.5(1) ( \times 10^6 )</td>
</tr>
<tr>
<td>Peak density [^{88}\text{Sr}/\text{cm}^3]</td>
<td>5.1(7) ( \times 10^{10} )</td>
<td>2.8(2) ( \times 10^{11} )</td>
<td>4.8(4) ( \times 10^{11} )</td>
</tr>
<tr>
<td>Peak PSD</td>
<td>2.8(1.2) ( \times 10^{-6} )</td>
<td>4(1) ( \times 10^{-4} )</td>
<td>1.3(2) ( \times 10^{-3} )</td>
</tr>
<tr>
<td>1/e lifetime</td>
<td>4.53(6) s</td>
<td>2.8(2) s</td>
<td>1.95(6) s</td>
</tr>
</tbody>
</table>

\(^{88}\text{Sr}\) at the location of Red MOT I no imaging system for the \( x \) axis was available so Red MOT I density and PSD calculations assume that temperature and width are the same as \( z \) along \( x \).
To summarize, we have demonstrated the operation of a MOT with a PSD of \(1.3(2) \times 10^{-3}\) in the steady-state regime. This result requires the use of broad- and narrow-linewidth transitions to simultaneously achieve both high PSD and high capture efficiency. We use a dual chamber architecture to protect the MOT from heating by broad-linewidth transition photons, and efficient transfer between chambers is achieved using a hybrid slower + MOT configuration using only cooling on the narrow-linewidth transition. Although Sr is ideally suited to this architecture, our approach is broadly applicable to alkaline-earth metals, lanthanides, and any other species with a strong transition to precool atoms and a weak transition supporting the operation of a MOT with a high PSD \([54,55]\). Finally, we have shown that our design is compatible with the generation of quantum degenerate gases in the presence of a laser cooled influx. The use of this high-PSD source of matter, combined with a protection mechanism such as that demonstrated in Ref. \([7]\), should allow the creation of a steady-state Bose-Einstein condensate and ultimately an atom laser with uninterrupted phase coherent output.

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S. B. and C.-C. C. contributed equally to this work.

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