Magnetic trapping of ytterbium and the alkaline-earth metals

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Atomic ytterbium (Yb), magnesium (Mg), calcium (Ca), and strontium (Sr) possess a simple yet versatile internal level structure and a diversity of naturally abundant fermionic and bosonic isotopes, making these systems ideal for studies of cold collisions and weakly interacting quantum degenerate gases. Unlike alkali-metal atoms, however, Yb, Mg, Ca, and Sr cannot be magnetically trapped in the ground state. We analyze a solution to this problem involving magnetic trapping in a low-lying metastable excited state and predict that significant magnetic trap populations can be obtained via continuous, in situ loading from Yb and Sr $^1S_0$ $^1P_1$ magneto-optical traps.

Magnetic trapping of ground-state neutral atoms [1] has enabled unique explorations into two-body interatomic interactions [2] and collisions [3], radio-frequency and optical spectroscopy [4], trapped atom laser-induced [5] and evaporative cooling [6], and weakly interacting quantum degenerate gases [7,8]. Atomic ytterbium (Yb) and the alkaline-earth atoms magnesium (Mg), calcium (Ca), and strontium (Sr) [hereafter collectively labeled singlet-triplet (ST) atoms] possess a diversity of naturally abundant fermionic and bosonic isotopes [9], spectrally narrow transitions that enable ultralow temperature, high spatial density laser cooling [10–12], and a simple internal level structure ideally suited to quantitative studies of cold collisions [13]. As such, ST atoms offer unique opportunities for future studies of magnetically trapped samples. Lacking the requisite ground-state magnetic substructure, however, ST atoms cannot be held in ground-state magnetic traps.

In this paper we analyze the solution to this problem proposed by Loftus et al. [14] which relies on magnetically trapping ST atoms in the low-lying $^3P_2$ ($m_j=2$) weak-field seeking metastable [15–18] excited state. We begin by showing that for each ST atom a single magnetic field supports both a $^1S_0$ $^1P_1$ transition magneto-optical trap (MOT) and a pure magnetic trap capable of holding $^1S_0$ $^1P_1$ MOT-cooled atoms. Next, we discuss three in situ techniques for loading $^3P_2$ ($m_j=2$) magnetic traps with atoms precooled in $^1S_0$ $^1P_1$ or $^1S_0$ $^3P_1$ MOTs and provide numerical estimates, based on the relevant Zeeman substate dependent decay paths [19], for the resulting magnetic trap loading rates. Finally, we discuss a zero-background technique for observing the magnetically trapped atoms. For simplicity, our discussion is confined to magnetic trapping of the most abundant, even atomic mass number ST atom isotopes. Note, however, that the procedures outlined here are applicable to the remaining isotopes, even those with nonzero nuclear spin.

Figure 1(a) shows a contour plot (calculated according to analytic expressions given by Bergman et al. [20]) of the typical anti-Helmholtz magnetic field used for a previously reported $^1S_0$ $^1P_1$ Yb MOT [14]. Coil windings lie in the centers of the four black dots and equimagnitude field contours are labeled in Gauss. The spatial extent of a magnetic trap employing this field is set by the direction-dependent radius of the maximum closed contour centered about the magnetic field null. We label the magnitude of this contour $B_M$ where in Fig. 1(a), $B_M=40$ G. In Fig. 1(b) we plot the $^3P_2$ ($m_j=2$) magnetic trap depth for each species as a function of $B_M$. The trap depth is scaled by $T_S$, the species specific $^1S_0$ $^1P_1$ Doppler-limited MOT temperature. The inset shows the trap depth normalized by $T_T$, the species specific $^1S_0$ $^3P_1$

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FIG. 1. (a) Contour plot of the anti-Helmholtz magnetic field used for a previously reported $^1S_0$ $^1P_1$ Yb MOT [14]. Coil windings lie in the centers of the four black dots and equimagnitude field contours are labeled in Gauss. (b) $^3P_2$ ($m_j=2$) magnetic trap depth as a function of $B_M$, the maximum effective trap field [~40 G in Fig. 1(a)]. The trap depth is scaled by $T_S$, the species specific $^1S_0$ $^1P_1$ Doppler-limited MOT temperature. (c) $^3P_2$ ($m_j=2$) magnetic trap depth versus the trap size, r, with a fixed axial field gradient of 60 G/cm [the axial gradient in (a)]. Here, r is measured relative to the magnetic field null point. Insets to (b) and (c) show the $^3P_2$ ($m_j=2$) magnetic trap depth normalized by $T_T$, the species specific $^1S_0$ $^1P_1$ Doppler limited temperatures for Yb and Sr. Note that a field gradient of ~10 G/cm (~5 G/cm) is required to suspend Yb (Sr) atoms against gravity.
Depicts the 3MOT precooling followed by population transfer to the 3N state. As demonstrated elsewhere, type II or type III transfer employs 3S0 − 1P1 MOT precooling followed by population transfer to the 3P2 (mJ = 2) state via optical pumping. As discussed above, all three transfer schemes are applicable to Yb, Ca, and Sr while, due to the details of the Mg internal level structure, Mg requires the use of type II or type III transfer. Note that in all cases approximately equal populations are loaded into the 3P2 (mJ = 2) and 3P2 (mJ = 1) weak-field seeking states. As demonstrated elsewhere [8], however, the resulting spin-mixed sample could be purified by ejecting 3P2 (mJ = 1) atoms from the magnetic trap with a sweep-frequency rf field. The remainder of our discussion, therefore, will focus on atoms loaded into the more tightly confined 3P2 (mJ = 2) state. In the following, we write the continuous or quasi-continuous type I and type II transfer rates, RM, as R = f(Ns/τs) = Rsf where f < 1 is an effective 1P1 → 3P2 (mJ = 2) or 3S1 → 3P2 (mJ = 2) cascade decay factor, Ns (τs) is the population (lifetime) of a 1S0 − 1P1 MOT, and Ns is a 1S0 − 1P1 MOT loading rate. Type III loading, which involves transferring population from a 1S0 − 1P1 MOT to the 3P2 (mJ = 2) state in a pulsed manner, is characterized by f′Nf where f′ < 1 is a 3P2 (mJ = 2) transfer fraction and Nf is the steady-state population of a 1S0 − 3P1 MOT. Note that f and f′ depend on spontaneous decay rates and Zeeman substate dependent weighting factors. For the former, we use either measured or theoretically predicted values (see Fig. 3 and Refs. [13,16,17,24–32]) while the latter are calculated according to the Wigner-Eckhart theorem [33].

In type I and type II transfer [Figs. 2(a) and 2(b), respectively], the atoms are initially loaded into 1S0 − 1P1 MOTs. Type I transfer then consists of allowing the atoms to radiatively cascade from the 1P1 state to the 3P2 (mJ = 2) state while they are held in the 1S0 − 1P1 MOT. Note that in this case τs < τf/1000 for all four ST atoms where τf is the time required to slow atoms from the MOT capture velocity to T5. Thus, the atoms are generally cooled to T5 before the radiative cascade from the 1P1 state to the 3P2 (mJ = 2) state occurs. For Yb and Sr, this transfer scheme is relatively efficient (see Table I) since cascade radiative decay from the 1P1 state which terminates in the 3P2,0 states (3P2 state is

![Figure 2](image-url)  
**FIG. 2.** Simplified energy-level diagrams showing (a) type I, (b) type II, and (c) type III transfer processes and a zero-background technique for observing atoms magnetically trapped in the 3P2 (mJ = 2) state. In (a), transitions and states labeled with dotted-dashed lines apply only to Yb. In (b), Pump I (Pump II and Pump III) is used for incoherent (semicoherent) transfer. For all three, the detection (probe) transitions are denoted by dashed (dotted) lines.

![Figure 3](image-url)  
**FIG. 3.** Simplified energy-level diagrams for (a) Yb and (b) Sr. Electric dipole and quadrupole and magnetic dipole and quadrupole transitions are represented by solid, dashed, dotted, and dotted-dashed lines, respectively. States with a superscript o have odd parity, transition wavelengths are given in vacuum [24,25], and numbers in parentheses give transition probabilities (in s−1) [16–18, 27–31]. The inset in (a) shows an expanded view of the Yb 1P1 → 3D2,3 → 3P2,0 decay channel.
the dominant Yb (Sr) $^{1}S_{0}-^{1}P_{1}$ MOT loss mechanism [10,11,23] and intermediate states decay quickly to the $^{3}P_{2}$ state [see Figs. 3(a) and 3(b)]. Using the $N_g$ and $\tau_g$ values reported for Sr in Ref. [11], for example, gives $N_g \sim 10^9$ atoms/s. Moreover, in Yb the transfer efficiency can be increased approximately an order of magnitude by optically pumping population from the $^{3}P_{0}$ state to the $^{3}P_{2}$ ($m_j=2$) through excitation of the $^{3}P_{0}-^{3}S_{1}$ transition [see Fig. 3(a)]. As shown in Table I, type I transfer is similarly efficient for Ca. Note, however, that cold Ca atoms lost from a $^{1}S_{0}-^{1}P_{1}$ MOT via radiative decay spend ~10 ms in the intermediate $^{1}D_{2}$ state [Ca Einstein $A(^{1}D_{2}-^{3}P_{2}) = 92.9$ s$^{-1}$ [28]], a process that increases the $^{3}P_{2}$ ($m_j=2$) cloud size [using the conservative force experienced by $^{1}D_{2}$ state atoms in the Fig. 1(a) field, the atoms travel ~1 mm prior to $^{3}D_{2}-^{3}P_{2}$ decay] and results in additional heating [approximately a factor of 2 along the Fig. 1(a) trap axial direction]. Finally, note that type I transfer cannot be used for Mg since in this case the $^{1}S_{0}-^{1}P_{1}$ transition is radiatively closed [34].

In type II transfer, these limitations are overcome by optically pumping $^{1}S_{0}-^{1}P_{1}$ MOT precooled atoms to the $^{3}P_{2}$ ($m_j=2$) state via radiative decay from the $^{3}S_{1}$ state, a process that can be accomplished two ways. First, exciting the $^{1}P_{1}-^{3}S_{1}$ transition with a Rabi frequency, $\Omega \gg 1/\tau_p$, where $\tau_p$ is the $^{3}S_{1}$ state lifetime, equalizes the $^{1}P_{1}$ and $^{3}S_{1}$ state populations and fills the $^{3}P_{2}$ ($m_j=2$) state with a 1/e time of $\sim \tau_p$. This incoherent population transfer is continued until the $^{1}S_{0}-^{1}P_{1}$ MOT is emptied, can be repeated once every $\tau_s$ (note $\tau_s \gg \tau_p$), and assuming equal population distributions in the $^{1}P_{1}$ and $^{3}S_{1}$ magnetic substates, yields $f \sim 0.1$ for all four ST atom species. Alternatively, once every $\tau_s$, the $^{1}S_{0}-^{1}P_{1}$ MOT optical fields are extinguished and temporally separated, circularly polarized $\pi$ pulses excite the stepwise $^{1}S_{0}-^{3}P_{1}$ ($m_j=1$) $\rightarrow$ $^{3}S_{1}$ ($m_j=1$) transition. Subsequently, radiative decay from the $^{3}S_{1}$ ($m_j=1$) state loads the atoms into the $^{3}P_{2}$ ($m_j=2$) state. This semicoherent process enhances the type II transfer efficiency by roughly a factor of 3 and due to the spectral width of the $^{1}S_{0}-^{3}P_{1}$ transition, enables additional one-dimensional cooling [10,35]. Moreover, the resulting magnetic trap spin mixture favors atoms in the $^{3}P_{2}$ ($m_j=2$) state over atoms in the $^{3}P_{2}$ ($m_j=1$) state by a factor of 2.

Lastly, in type III transfer, the atoms are precooled in $^{1}S_{0}-^{1}P_{1}$ MOTs loaded with either a $^{3}S_{0}-^{3}P_{1}$ MOT [10,11] or a Zeeman slowed atomic beam [12]. The MOT optical and magnetic fields are then turned off, polarized $\pi$ pulses load atoms into the $^{3}P_{2}$ ($m_j=2$) state by exciting the stepwise $^{1}S_{0}-^{3}P_{1}$ ($m_j=1$) $\rightarrow$ $^{3}S_{1}$ ($m_j=1$) transition, and the magnetic trap is turned on around the optically prepared atoms. For a given $^{1}S_{0}-^{3}P_{1}$ MOT population, $N_T$, the resulting magnetic trap population is $f^* N_T$ where here, $f^* \sim 0.3$. Although the most technically challenging of all three transfer techniques, this approach has the benefit of producing ultracold ($\sim 3$ μK in the case of Sr [36]), large population ($\sim 10^7$ atoms [10–12]) samples and is an attractive extension of the intercombination line laser cooling currently being pursued in Sr [10,11] and Yb [12]. Note, however, that type III transfer cannot be applied directly to Mg or Ca since in these cases, $F_3^* \sim F_5^*/10$ where $F_T \sim F_3^*$ is the $^{1}S_{0}-^{3}P_{1}$ optical (gravitational) force and thus $^{1}S_{0}-^{3}P_{1}$ MOTs cannot suspend Mg or Ca against gravity. For these two species, however, magnetic trapping of ultracold samples could be achieved by substituting the quenched narrow-line laser cooling technique recently developed by Curtis et al. [37] for the $^{1}S_{0}-^{3}P_{1}$ MOT stage. Finally, we point out that due to the reduced sample temperatures achieved with type III transfer, Majorana floss will likely be a significant source of trap loss if the quadrupole magnetic field geometry assumed here is employed. This problem, however, can easily be overcome by switching in either rotating bias [38] or Ioffe [39] fields during the magnetic trapping stage.

For all three transfer schemes, zero-background detection of the magnetically trapped atoms can be realized via optical excitation of the $^{3}P_{2}-^{3}S_{1}$ transition followed by selective observation of radiative decay from the $^{3}S_{1}$ state to either the $^{3}P_{1}$ state or the $^{3}P_{0}$ state. Several factors, however, must be considered when performing these measurements. First, in contrast to traditional ground-state magnetically trapped alkali metal samples, the probability that a given atom will radiatively decay to untrapped states ($^{3}P_{1}$ and $^{3}P_{0}$) following excitation of the $^{3}P_{2}-^{3}S_{1}$ transition is approximately 1 (see Fig. 3 and Ref. [32]). Consequently, detecting the magnetically trapped atoms requires efficiently collecting the $^{3}S_{1}-^{3}P_{1,0}$ fluorescence. Additionally, although the $^{3}S_{1}-^{3}P_{2}$ transition is spectrally separated from the $^{3}S_{1}-^{3}P_{1}$ and $^{3}S_{1}-^{3}P_{0}$ transitions by $\Delta \lambda_{p}\sim 10$ nm in Yb and Sr, for Mg and Ca, $\Delta \lambda_{p}\sim 1$ nm [24,25], requiring, for zero-background measurements, a spectrometer or an equivalent spectrally selective device. Note, however, that for sufficiently high spatial densities it may be possible to use the absorptive [7] or dispersive light scattering [40] techniques commonly employed to characterize magnetically trapped alkali metals.

Studies of laser cooled Yb and alkaline earth atoms have provided important insights into light-assisted collisions, radiation-pressure induced MOT density limits, and all-optical pathways to quantum degeneracy. We have presented and analyzed several novel strategies for efficiently loading-

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<th>Loading scheme</th>
<th>Species</th>
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<tr>
<td>Type I</td>
<td>$^{174}$Yb</td>
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<tr>
<td></td>
<td>$^{40}$Ca</td>
<td>0.2</td>
</tr>
<tr>
<td></td>
<td>$^{88}$Sr</td>
<td>0.2</td>
</tr>
<tr>
<td>Type II</td>
<td>$^{174}$Yb</td>
<td>0.3</td>
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<tr>
<td></td>
<td>$^{24}$Mg</td>
<td>3</td>
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<td>$^{40}$Ca</td>
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these species into magnetic traps and demonstrated that significant trap populations can be obtained using $^{1}S_0 - ^1P_1$ MOT magnetic fields and currently achieved $^{1}S_0 - ^1P_1$ MOT populations. Our procedures may thus enable the uniquely simple yet versatile properties of Yb and the alkaline earths to be exploited in the ultracold, photon-free regime available in magnetic traps where evaporative cooling and ultimately, observations of quantum degeneracy in these species may be possible.

**Note added in proof.** Recently, refined values for the Mg, Ca, and Sr $^3P_2$ state lifetimes have become availbale. See Ref. [41].

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[36] Using the 400 nK $^{1}S_{0} \rightarrow ^{3}P_{1}$ MOT temperature reported in Ref. [11] and including loading induced photon recoil heating.


