Steady-state magneto-optical trap of fermionic strontium on a narrow-line transition

González Escudero, R.; Chen, C.-C.; Bennetts, S.; Pasquiou, B.; Schreck, F.

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I. INTRODUCTION

Optical atomic clocks [1] are proving to be powerful tools to answer fundamental questions in physics. They enable searches for dark matter [2–4], the study of gravitational waves [5], tests of general relativity [6,7], or variations in nature’s fundamental constants [8–10]. In the applied domain they are playing an increasingly important role in geology and geodesy [11–14].

Typically, state-of-the-art optical clocks operate in a time-sequential manner, with distinct phases for sample preparation and measurement. This time-sequential mode of operation introduces noise, known as the Dick effect [15], which limits their performance [16]. The Dick effect results from the down-conversion or aliasing of high-frequency noise, present in the laser used to interrogate the atoms, down to frequencies lower than the clock’s interrogation frequency. To address it, several approaches have been followed: employing high-precision nondestructive detection techniques [17], developing novel experimental architectures to achieve high duty cycles [18], or interleaving the clock interrogations between two different atomic samples [19–21].

A new class of optical clocks based on the principle of superradiant lasing has already been demonstrated [25], in particular using strontium’s $^1S_0 - ^3P_1$ clock transition [26,27]. There are ongoing efforts to incorporate steady-state atomic sources with this new type of clock [25,28–30]. While there exist several strontium sources achieving the required flux and temperature [31,32], all of these operate using bosonic isotopes, due to their simpler atomic structure. However, superradiant lasing using the clock line of bosonic strontium has, so far, not been demonstrated.

For bosonic strontium, the clock linewidth is orders of magnitude smaller than for its fermionic counterpart, and requires high external magnetic fields for the linewidth to be broadened enough for clock interrogation [33,34]. Moreover, such fields must feature an exceptional degree of homogeneity and stability across the atomic sample for all atoms to participate in superradiant lasing [35]. Conversely, the laser cooling stages that are necessary to achieve the desired temperature and density operate best at low magnetic fields and gradients. These two adverse requirements complicate the integration of a steady-state source of bosonic strontium into an active optical clock. By contrast, a fermionic source seems better suited. Indeed, due to the nonzero nuclear magnetic moment of fermionic strontium, the clock transition is naturally much stronger than for its fermionic counterpart, and requires flux and temperature [31,32], all of these operate using bosonic isotopes, due to their simpler atomic structure. However, superradiant lasing using the clock line of bosonic strontium has, so far, not been demonstrated.

A new class of optical clocks based on the principle of superradiant lasing has been proposed as next-generation devices [22,23]. Their working principle is inherently suited to continuous operation, thus circumventing the Dick effect. Furthermore, their potential goes beyond addressing this challenge: in the future they could outperform conventional clocks in terms of stability, detection bandwidth, and, by relaxing the requirements on external high-finesse cavities, portability, and robustness [23,24].

Pulsed superradiant lasing has already been demonstrated [25], in particular using strontium’s $^1S_0 - ^3P_0$ clock transition [26,27]. There are ongoing efforts to incorporate steady-state atomic sources with this new type of clock [25,28–30]. While there exist several strontium sources achieving the required flux and temperature [31,32], all of these operate using bosonic isotopes, due to their simpler atomic structure. However, superradiant lasing using the clock line of bosonic strontium has, so far, not been demonstrated.

For bosonic strontium, the clock linewidth is orders of magnitude smaller than for its fermionic counterpart, and requires high external magnetic fields for the linewidth to be broadened enough for clock interrogation [33,34]. Moreover, such fields must feature an exceptional degree of homogeneity and stability across the atomic sample for all atoms to participate in superradiant lasing [35]. Conversely, the laser cooling stages that are necessary to achieve the desired temperature and density operate best at low magnetic fields and gradients. These two adverse requirements complicate the integration of a steady-state source of bosonic strontium into an active optical clock. By contrast, a fermionic source seems better suited. Indeed, due to the nonzero nuclear magnetic moment of fermionic strontium, the clock transition is naturally much stronger than for bosonic strontium [31,36], making it accessible without the need for the high external magnetic fields that are typically required with bosons.

Here we demonstrate a continuously loaded magneto-optical trap (MOT) of ultracold fermionic $^{87}\text{Sr}$ atoms, operated on the 7.5-kHz-wide $^3S_0 - ^3P_1$ intercombination line. This narrow-line MOT contains $8.4\times10^7$ atoms at an average temperature of $12.0(1)\mu\text{K}$, loaded with a rate of $1.31(2)\times10^7$ atoms/s, and reaches a peak density of $2.12(4)\times10^9\text{Sr/cm}^3$. This steady-state source could be used to provide the high flux of ultracold atoms necessary to refill the gain medium of a steady-state superradiant active clock [24–26].
while Secs. III–V focus on the specificities pertaining to
the fermionic isotope $^{87}$Sr. Section III describes a two-
dimensional (2D) molasses, making use of the sawtooth-wave
adiabatic passage (SWAP) technique to collimate an atomic
beam passing from one chamber to the next. Section IV de-
scribes a narrow-line Zeeman slower adapted to fermionic
strontium, thanks to optical pumping into suitable states of
the hyperfine manifold of ground-state strontium. Section V
characterizes the MOT performance and directly compares the
$^{88}$Sr and $^{87}$Sr cases. Lastly, Sec. VI discusses the application
of this atomic source to superradiant active optical clocks and
concludes.

II. APPARATUS OVERVIEW

In this section we first outline the main components of our
apparatus, by describing every laser cooling stage necessary
to produce a continuous narrow-line MOT. We then highlight
the main differences between steady-state narrow-line MOTs
of $^{88}$Sr (bosonic strontium) and $^{87}$Sr (fermionic strontium).

Similarly to our previous work on bosonic strontium [32],
we stream a beam of atoms through a sequence of spatially
separated laser cooling stages, until reaching the $\mu$K regime.
Initially, atoms coming from an oven at $\sim 550$ °C are slowed
down by a Zeeman slower operated on the 30-MHz-wide
$^1S_0 - ^1P_1$ transition. For this work, we have upgraded the power
of the Zeeman slower beam to improve our atomic flux by
more than an order of magnitude, using 500 mW diodes now
available from Nichia (Part NDB4916) [37]. The high cycling
rate of the $^1S_0 - ^1P_1$ transition is necessary to decelerate and
cool a significant part of the velocity distribution of the beam
exiting the oven. However, parasitic light resonant with this
transition, coming, for example, from uncontrolled reflec-
tions or fluorescence from the laser-cooled atoms, will be
extremely damaging to superradiantly lasing ensembles and
quantum degenerate gases [26,38,39]. It will also be harmful
to the lifetime and temperature of our narrow-line MOT. To
address these issues, we opted for a two-chamber design,
consisting of an upper and a lower chamber, separated by
$\sim 40$ cm (see Fig. 1). We use light addressing $^1S_0 - ^1P_1$ in
the upper beam but not in the lower one. Indeed, by inducing
a light shift on the $^3P_1$ excited state, the 7.48(5)-kHz-wide
$^1S_0 - ^3P_1$ transition can be locally shifted out of res-
onance. In such a way, long-lived quantum degenerate gases
can exist in the presence of laser cooling light addressing the
$^1S_0 - ^3P_1$ transition [38,39].

In the upper chamber, Zeeman-slowed atoms are gathered
in a 2D MOT using the $^1S_0 - ^1P_1$ transition (see Fig. 1). This
2D MOT produces a vertical, downward propagating atomic
beam with a radial velocity $\lesssim 0.5$ m s$^{-1}$. In order to efficiently
transfer atoms to the lower chamber through a 20-mm-long,
8-mm-diameter baffle, this atomic beam requires collimation.
In typical double-chamber experiments, this collimation can
be achieved by accelerating the atoms exiting the 2D MOT
to high velocities. However, due to the low scattering rate of
the $^1S_0 - ^3P_1$ transition, we would not be able to stay below the
capture velocity in the lower chamber. Thus, in order to
achieve a high-transfer efficiency we need to provide addi-
tional radial cooling for the beam. We achieve this goal by
employing a 2D molasses operating on the $^1S_0 - ^3P_1$ transition
and located directly below the 2D MOT.

Due to gravity, atoms accelerate and reach the center of the
lower chamber, located $\sim 40$ cm below the 2D MOT, with a
downward velocity distribution peaked at $\sim 4$ m s$^{-1}$. A con-
trappropagating laser beam slows the atoms over an $\sim 5$ cm
distance using the $^1S_0 - ^1P_1$ transition. Finally, a narrow-line
five-beam three-dimensional (3D) MOT using the same transition
captures the atoms. The vertical, upward propagating
beam serves a dual purpose, both slowing the atoms coming
from the upper chamber and acting as a MOT beam. Once in
the MOT, atoms are slow enough that gravity is sufficient to
provide the downward restoring force.

The above description applies to our previous work on $^{88}$Sr
[32] and to our new work on $^{87}$Sr described here. However, in
order to deal with the internal structure of $^{87}$Sr, our new
architecture implements two key changes. Firstly, the collimi-
gating 2D molasses needs to be upgraded using the SWAP
technique [42], to account for the magnetic sensitivity of all
$^{87}$Sr $^3P_1$ Zeeman sub-levels (see Sec. III). Secondly, optical
pumping into suitable states of the Sr $^1S_0$ hyperfine manifold
is necessary to ensure efficient slowing of falling atoms in the
lower chamber. We use additional laser light addressing the
$^1S_0 F = 9/2 - ^3P_1 F' = 9/2$ transition [43] to perform optical
pumping thus enhancing the atomic flux into the MOT (see
Secs. IV and V).

III. 2D SWAP MOLASSES

In this section, we describe how the 2D molasses using
the narrow $^1S_0 - ^3P_1$ transition collimates the atomic beam in
order for the atoms to pass from the upper chamber to the lower chamber. We first detail how we previously achieved this collimation for bosonic $^{88}\text{Sr}$, then highlight the additional challenges pertaining to fermionic $^{87}\text{Sr}$. We resolve these challenges using SWAP cooling, a technique which uses sawtooth-shaped detuning ramps of the laser cooling light. We recall two criteria for this technique to work. We then explore its limits in our implementation and show that our 2D molasses performs better using SWAP laser detuning ramps rather than inverted SWAP or symmetric ramps. Finally, we measure and model the cooling efficiency of the 2D molasses in dependence of the laser powers and ramp durations.

### A. SWAP molasses needed for fermionic Sr

The collimation molasses is located just below the broad-line 2D MOT that uses the $^1S_0 - ^3P_1$ transition. It exploits the much lower Doppler temperature limit of the narrow-line $^1S_0 - ^3P_1$ transition to further reduce the radial velocity spread of the downward propagating atomic beam (see Fig. 1). As atoms fall with a velocity of $\sim 3 \text{ m s}^{-1}$, radial cooling with this 2D molasses reaches steady state in at most 10 ms. To achieve such a long interaction time, the molasses beams are elliptical, with a $1/e^2$ diameter of 45.6 mm along the vertical axis and 18.2 mm along the horizontal axis.

The molasses needs to address the atoms efficiently, despite a Doppler broadening about 100 times larger than the transition’s natural width $\gamma = 2\pi \times 7.5 \text{ kHz}$. For bosonic strontium we therefore broaden the molasses beams' frequency spectrum, forming a frequency comb spanning from 30 to 750 kHz to the red of the transition with a 25 kHz spacing [32]. An additional challenge arises from the presence of a residual 2D quadrupole magnetic field with a gradient of 2 G cm$^{-1}$ at the location of the molasses. This field originates from the 2D quadrupole field of the nearby 2D MOT and it cannot be completely removed. This residual field leads to Zeeman broadening of the magnetic $^1S_0 - ^3P_1$ transitions and position dependent light scattering forces. This challenge is overcome by addressing only the $|J = 0, m_J = 0\rangle - |J' = 1, m_J = 0\rangle$, $\pi$, nonmagnetic transition of $^{88}\text{Sr}$. To this end, we add a vertical bias field of 1.2 G to Zeeman shift the other two, $\sigma^\pm$, transitions away from resonance, as the solution previously used in Ref. [32]. At the location of the atomic beam, the average field amplitude is 2.4 G, creating a Zeeman shift of about 5 MHz, a significant part of the field being along the $z$ axis (see Fig. 1). The field orientation is not fully vertical because the atomic beam is displaced sideways with respect to the zero of the 2D quadrupole field, resulting in an additional component along the $z$ axis.

Due to the hyperfine structure of $^{87}\text{Sr}$, nonmagnetic transitions from $^1S_0$ to $^3P_1$ do not exist [see Fig. 2(a)]. Therefore, the method just described no longer provides efficient transfer of atoms to the lower chamber. To overcome this difficulty we use SWAP cooling [42,44–48], a technique that uses two rapid adiabatic passages to transfer atoms to a long-lived excited state and back to the ground state. In this work these states are $^3P_1$ and $^1S_0$, respectively. SWAP cooling exploits the time ordering of the adiabatic passages, transferring photons between two counterpropagating laser beams, and ensuring that the recoil momenta associated with photon absorption and stimulated emission are both oriented against the atoms’ direction of travel (see Fig. 3). SWAP cooling has the advantage of robustly addressing a wide range of effective detunings, while increasing deceleration. Therefore it can be used on atomic samples whose Doppler broadening is much wider than the $^1S_0 - ^3P_1$ transition’s width, and it can address several

![FIG. 2. (a) $^{87}\text{Sr}$ energy levels relevant for our experiment. The arrows indicate all transitions necessary to obtain our steady-state narrow-line MOT. The hyperfine shifts of the $^3P_1$ manifold are referenced to the single $^3P_1$ state of $^{88}\text{Sr}$. (b) Magnetic sublevels of the $^3P_1 F' = 9/2$ state for an atom at rest in the presence of a field of 2.4 G. Also shown is a schematic of the detuning sweep (red lines) used by the 2D SWAP molasses to collimate the atoms propagating toward the lower chamber. (c) Schematic of the symmetric sweep used to collimate bosonic atoms [32]. The ramp parameters shown are the sweep duration $\Delta t$, the laser start detuning $\Delta_1$, and stop detuning $\Delta_2$.](image)

![FIG. 3. SWAP working principle. The plain black trace describes the laser frequency sweep in the laboratory frame. In the reference frame of the atom, the detuning of each of the two counterpropagating beams (blue and red traces) is Doppler shifted by $\hat{k} \cdot \mathbf{v}$. This shift introduces an effective time ordering, ensuring that the momenta transferred to the atom are oriented against its direction of travel.](image)
transitions with distinct differential Zeeman shifts in a single sweep.

B. SWAP cooling criteria

For SWAP cooling to be effective, two main criteria must be fulfilled [42,44]. The first is the adiabaticity of the rapid adiabatic passage from one internal state to the other, which is required for both absorption and stimulated emission. To be adiabatic, the square of the Rabi frequency of the laser light $\Omega^2$ needs to be high compared to the laser detuning sweep rate $\alpha$, i.e., $\kappa = \Omega^2/\alpha \gg 1$. Here $\alpha = \Delta \omega/\Delta t$, $\Delta \omega$ is the full detuning range spanned, and $\Delta t$ is the duration of a single sweep. Indeed, for a Landau-Zener type crossing the probability of transfer is given by $P_{LZ} = 1 - \exp(-\pi \kappa / 2)$ [44,49].

The second criterion concerns the time an atom must spend in the excited state in order to perform a complete cycle of absorption and stimulated emission. This time must be smaller than the natural decay time by spontaneous emission $1/\gamma$. If this criterion is not met, spontaneous emission will occur before the stimulated emission, reducing the cooling efficiency. Moreover, if the atom absorbs a second photon by adiabatic passage from the copropagating beam, momentum will be imparted in the wrong direction. This second absorption and stimulated emission results in net momentum gain or loss. A positive slope (SW AP ramps), such as the one shown in Fig. 2(b), results in deceleration and cooling, while a negative slope (inverted SW AP ramps) results in acceleration and heating [44,47]. In Fig. 3 we compare the atom number loaded into the MOT (see Sec. V) for molasses following SWAP, inverted SWAP, and symmetric ramps, with varying sweep frequency $1/\Delta t$. For all explored $\Delta t$, the flux of loaded atoms is the highest using SWAP ramps, while inverted SWAP ramps perform the worst. This difference is the most pronounced at $\kappa = 1$, corresponding to long durations, $\Delta t \gg e^{-\kappa}$. For $\kappa \gg 1$, corresponding to long durations, the adiabaticity parameter decreases to $\kappa \ll 1$, the probability of stimulated emission or absorption becomes negligible. This reduces the cooling efficiency and effectively breaks the time ordering. In this limit the cooling efficiencies of SWAP and inverted SWAP ramps converge and result in similar fluxes. For $\kappa \gg 1$, corresponding to long durations, the adiabaticity criterion is fulfilled, but the second criterion is broken, as $\tau_r \propto 1/|\alpha| \propto \Delta t$. For $\tau_r \gg 1/\gamma$, the cooling reduces to two independent stimulated absorption events each followed by spontaneous emission: one from the counterpropagating beam and one from the copropagating beam. In this situation the average slowing force approaches zero for all three ramp types.

C. SWAP and inverted SWAP working regimes

For SWAP cooling, the sign of the sawtooth ramp slope determines whether the time ordering associated with absorption and stimulated emission results in net momentum gain or loss. A positive slope (SWAP ramps), such as the one shown in Fig. 3, results in deceleration and cooling, while a negative slope (inverted SWAP ramps) results in acceleration and heating [44,47]. In Fig. 4 we compare the atom number loaded into the MOT (see Sec. V) for molasses following SWAP, inverted SWAP, and symmetric ramps, with varying sweep frequency $1/\Delta t$. For all explored $\Delta t$, the flux of loaded atoms

\[ P_{LZ} = 1 - \exp(-\pi \kappa / 2) \]

is the highest using SWAP ramps, while inverted SWAP ramps perform the worst. This difference is the most pronounced at $1/\Delta t \approx 15$ kHz, corresponding to an adiabaticity criterion $\kappa \sim 1.5$, while in the limits of short or long sweep durations the fluxes become similar. This comparison of the three types of ramp shows that the time ordering of the molasses beams’ action matters and that the adiabaticity of the emission and absorption of the rapid adiabatic passages plays a significant role here.

The results of Fig. 4 can be explained by considering the two SWAP criteria previously given. Indeed, for short durations the adiabaticity parameter decreases to $\kappa \ll 1$ and the probability of stimulated emission or absorption becomes negligible. This reduces the cooling efficiency and effectively breaks the time ordering. In this limit the cooling efficiencies of SWAP and inverted SWAP ramps converge and result in similar fluxes. For $\kappa \gg 1$, corresponding to long durations, the adiabaticity criterion is fulfilled, but the second criterion is broken, as $\tau_r \propto 1/|\alpha| \propto \Delta t$. For $\tau_r \gg 1/\gamma$, the cooling reduces to two independent stimulated absorption events each followed by spontaneous emission: one from the counterpropagating beam and one from the copropagating beam. In this situation the average slowing force approaches zero for all three ramp types.
falling atoms in the lower chamber. For this purpose, we beam powers with varying sweep frequency $1/\Delta t$. The powers stated in the legend refer to the average laser power for each beam. Each data point represents the average of ten independent measurements and the error bars show the standard deviation of these measurements. The vertical dashed lines highlight the location of $\kappa = \Omega^2/\alpha = 1.5$. (b) Numerical estimations of the molasses’ average restoring force exerted on an atom exiting the 2D MOT, with the same parameters as in (a). The curves represent thermal averages over a 1D Maxwell-Boltzmann distribution at 1 mK for two regimes: one integrated only over the low-velocity class $|v| < 0.25 \text{ m s}^{-1}$ (solid line) and the other integrated over the high-velocity class $|v| > 0.25 \text{ m s}^{-1}$ (dashed lines). The given Rabi frequencies result from spatial averages of the intensity across a 1/e$^2$ beam waist.

D. Modeling SWAP molasses

To explore the power dependence of the 2D SWAP molasses, we measure the relative transfer efficiency to the lower chamber for different laser beam powers and sweep durations. These results are shown in Fig. 5(a).

We obtain this data by collecting fluorescence from the falling atoms in the lower chamber. For this purpose, we illuminate the passing atoms with light addressing the fast cycling $^1S_0 - ^3P_1$ transition, with no other beam present in the lower chamber. This detection method provides a more direct measure of the atomic flux arriving into the lower chamber than what is given by the MOT atom number.

For each chosen power we observe the existence of an optimal sweep frequency close to $1/\Delta t_{opt} \sim 10 \text{ kHz}$, that shows only a small dependence with the laser beam power. The location of this optimum does not quite correspond to the value of the adiabaticity criterion $\kappa$, indicated in Fig. 5(a) by vertical dashed lines. This mismatch might arise from the laser beams’ finite interaction time with the atoms. Indeed, in order to pass through the 8-mm-diameter baffle between the two vacuum chambers, the radial velocity must be reduced below $v < v_{\text{min}} \approx 0.07 \text{ m s}^{-1}$. However, atoms exiting the 2D MOT have a radial velocity up to $0.5 \text{ m s}^{-1}$. Successful passage to the lower chamber therefore requires that the 2D molasses removes about 50 recoil momenta per beam pair. Assuming a momentum transfer of one photon recoil per sweep, achieving this deceleration within the finite interaction time with the laser beams would require a minimum sweep frequency of $1/\Delta t \sim 3 \text{ kHz}$. The measured optimum sweep frequency is three times slower, indicating that on average only 0.3 photons might be transferred per sweep, corresponding to a restoring force of $\sim 3\hbar \text{kHz/ms}$.

To deepen our insight into SWAP molasses cooling, we theoretically estimate the average restoring force exerted on the atoms exiting the 2D MOT [see Fig. 5(b)]. We limit our discussion to the average force exerted by the molasses, as reproducing the data shown in Fig 5(a) would require introducing free parameters into our model that we do not have access to. Such parameters include the atomic distribution exiting the 2D MOT, the transit time and the maximum transfer velocity of the atoms. Moreover, including them in the model would not provide deeper insight into the molasses cooling process itself.

The results of Fig. 5(b) are obtained by expanding on the theory developed by Bartolotta et al. [44] to take into account the internal structure of $^{87}\text{Sr}$. In particular, we numerically integrate the optical Bloch equations for the sweep durations and Rabi frequencies shown in Fig. 5(b) using the finite set of states,

$$\{|m_F, p + n\hbar k\}, \{m_F, p + n\hbar k\},$$

where $n \in \{2, 1, 0, -1, -2\}$, $p$ defines the starting momentum used in the calculation, and we have used shorthand notation for a particle with momentum $p + n\hbar k$ in the $^1S_0 F = 9/2 m_F$ and $^3P_1 F' = 9/2 m_F$ states, respectively.

We solve these equations stepping through several starting momenta, $p/\hbar k \in \{70, 69.5, 69, \ldots, 0\}$ and one sweep cycle of duration $\Delta t$ to obtain a momentum dependent force $F(p) = (\text{tr}(\rho(\Delta t) - \text{tr}(\rho F(p)/\Delta t))/\Delta t$. Here $\rho$ is the atomic momentum operator, $\rho_{p, m_F} = \sum_{m_F} P_{m_F, p} |m_F, p\rangle \langle m_F, p|$ is the initial state used in the calculation, and $\rho_{p, p'}$ is the final state after the single sweep.

To take into account optical pumping, the initial distribution over internal states $m_F$ described by $P_{m_F, p}$ in $\rho_{p}$ is obtained by an iterative process. Starting from a fully depolarized state with momentum $70\hbar k$, i.e., $P_{m_F, 70\hbar k} = 1/10$, the final occupation of the internal states of each momentum step is used as an input occupation for the following step, i.e., $P_{m_F, p'} = \sum_{m_F} \langle m_F, p + n\hbar k | \rho_{p, p'} | m_F, p + n\hbar k \rangle$, with $p' = 0.5\hbar k$ corresponding to our choice of discretization. We checked that the result of the calculation depends weakly on this somewhat arbitrary choice and that the shape of the curves shown in Fig. 5(b) remains similar when using a grid with $p - p' = (0.5, 1.2)\hbar k$.

The curves shown in Fig. 5(b) average the force obtained over a 1D Maxwell-Boltzmann distribution split into two velocity classes, $|v| < v_c$ and $|v| > v_c$. Atoms in the low-velocity class are rapidly deexcited via stimulated emission, whereas atoms in the high-velocity class remain unaddressed long enough to spontaneously emit a photon. Here we choose to distinguish these two classes for $1/\Delta t = 15 \text{ kHz}$, which gives $v_c = 0.25 \text{ m s}^{-1}$ using the second SWAP cooling criterion (see Sec. III B).

Our theoretical results confirm that for the high-velocity atoms, $v > v_c$, the cooling mostly reduces to a single absorption by adiabatic passage from the counterpropagating beam, followed by spontaneous emission. Low-velocity atoms $v < v_c$ have an effective detuning dominated by the Zeeman
shifts. Since our sweep range is red detuned from the unperturbed transition, it leaves the magnetic sublevels with $m_F > 3/2$ unaddressed. However, our simulations show that optical pumping is optimal for a sweep duration corresponding to \( \sim 10 \) kHz. This optical pumping enhances the population of the states that are efficiently addressed by the molasses beams, particularly those with low \(|m_F|\) values. Failing to take the effect of optical pumping into account shifts the optima shown in Fig. 5(b) to significantly lower sweep duration.

From these simulations we obtain the maximum average restoring force $F_{\text{max}} = 3.8 \hbar k/\text{ms}$ provided by the SWAP molasses. This relatively small force is enough to collimate most atoms such that they pass through the 8-mm-diameter hole between chambers, and is consistent with the rough estimation $\sim 3 \hbar k/\text{ms}$ we gave previously.

To finalize this section, we summarize the SWAP molasses cooling parameters used in the remainder of this work. The optical powers are 14.2 mW for one pair of beams and 15.2 mW for the other. Their polarization is linear and perpendicular to the $y$ axis. The sweep duration is $\Delta t = 71 \mu s$, while the molasses detuning is modulated between $\Delta_1 = -2\pi \times 2 \text{ MHz}$ and $\Delta_2 = +2\pi \times 0.2 \text{ MHz}$. Based on our experimental explorations, we find this set of parameters to provide optimal flux to our narrow-line 3D MOT.

IV. LASER SLOWING STAGE

Atoms falling through the 2D SWAP molasses region can be captured in the lower chamber by a narrow-line MOT operating on the $^1S_0 \rightarrow ^3P_1$ transition (see Fig. 1). Because of the limited deceleration achievable with the $^1S_0 \rightarrow ^3P_1$ transition, we slow the atoms before they reach the MOT. In the following, we first describe the operation of this slowing stage for bosonic isotopes. Then we highlight the additional challenges for the fermionic isotope. We finish this section by describing the frequency ramps used to efficiently slow the atoms.

Accelerated by gravity, atoms reach the entrance of the lower chamber with a downward velocity distribution peaked at $\sim 4 \text{ m/s}$, which exceeds the MOT’s capture velocity. Therefore we employ a counterpropagating laser beam addressing the $^1S_0 \rightarrow ^3P_1$ transition to begin slowing the atoms before they reach the MOT, which is located $\sim 40 \text{ cm}$ below the 2D SWAP molasses. This beam is circularly polarized to address the $\sigma^+$ transition and its frequency is red detuned from the unshifted resonance. For the laser slowing stage, the magnetic field is dominated by the quadrupole field used to generate the narrow-line MOT.

For bosonic strontium, the slowing stage operates as a Zeeman slower and its working principle is sketched in Fig. 6(a). As atoms decelerate, the change in Doppler shift $\delta_{\text{Doppler}}$ is partially compensated, thanks to the vertical magnetic field gradient produced by the MOT, $dB/dy = 0.32 \text{ G cm}^{-1}$.

The gradient is mostly linear up to 10 cm away from the MOT quadrupole center, beyond which the magnetic field amplitude decreases. Due to the high initial atom velocity the Zeeman shift alone is insufficient to fully address the atoms, so we also modulate the laser detuning with a span of $\Delta_{\text{Doppler}} = 2\pi \times 4.05 \text{ MHz}$, thus forming a “white light” slower [50].

The choice of laser detuning $\Delta_2$, together with the maximum magnetic field experienced by the falling atoms $B_{\text{max}}$, defines a capture velocity for the slowing stage, $v_{\text{capture}} = |\Delta_2 (B_{\text{max}}) - \Delta_1|/k$ (see Fig. 6). For bosonic strontium $v_{\text{capture}} = 5.3 \text{ m/s}$, faster than the velocity at which the atoms fall of $\sim 4 \text{ m/s}$. For fermionic strontium, the hyperfine structure complicates the slowing stage [see Fig. 6(b)].

All from the hyperfine transitions (see Fig. 2) only the $^1S_0 F = 9/2 \rightarrow ^3P_1 F' = 11/2$ “cooling” transition can have high enough capture velocity to slow the atoms while being compatible with the MOT configuration described in Sec. V. Therefore we use this transition in order to slow atoms entering the lower chamber. Note that the condition $v_{\text{capture}} > 4 \text{ m/s}$ can only be fulfilled for atoms in the $m_F > 3/2$ states, where $\vec{y}$ defines the quantization axis. Without optical pumping, fast atoms entering the lower chamber in other states will be lost. This loss mechanism can be particularly relevant as our simulations show that the collimating 2D molasses in the upper chamber results in a mixture of several $m_F$ states, predominantly with low $|m_F|$ values.

To optically pump ground-state atoms to the $m_F > 3/2$ Zeeman sublevels, we use dual-frequency laser light for the vertical slower beam, addressing two different hyperfine manifolds. First, we use the cooling transition, because of its high capture velocity. Secondly, we use the $^1S_0 F = 9/2 \rightarrow ^3P_1 F' = 9/2$ “pumping” transition to optically pump atoms that would otherwise remain unaddressed. For
As described above, optical pumping is important for the operation of the slower. We enhance it by using the dual-frequency configuration for the slower beam, thereby improving the loading efficiency by a factor 2.2(3) compared to when addressing only the cooling transition. Note that only using the pumping transition does not enable us to load any atoms in the MOT.

As atoms fall in the lower chamber they can fall out of resonance and be lost if they are not efficiently slowed [see Fig. 6(a), black dashed line]. To increase the available force, we ramp the detuning of both transitions addressed by the slower beam using an asymmetric sawtooth, as in Fig. 3. Operating the slower beam in this manner can yield a force up to twice the typical radiation pressure force [46]. Here, because atoms are always counterpropagating toward a single beam, ramps with positive or negative slope yield the same flux. Using an asymmetric sawtooth frequency sweep instead of a symmetric one serves only to suppress stimulated emission into the slower beam. Using the asymmetric frequency sweep we enhanced our loading efficiency by a factor of ~1.5 compared to using only the symmetric sweeps. Here the sweep time has been independently optimized for each sweep configuration and we have used the laser detunings and powers shown in Table I (YMOT beam).

V. STEADY-STATE NARROW-LINE MOT

Atoms that are successfully slowly accumulated in a narrow-line 3D MOT operating on the $^1S_0 - ^3P_1$ transition. In this section we describe how this MOT is set up and characterize its performance. In particular, we will focus on the comparison of the flux and lifetime between MOTs of $^{88}$Sr (bosonic strontium) and $^{97}$Sr (fermionic strontium).

The MOT confinement is provided by five beams: four in the horizontal plane and one propagating upward. The downward restoring force is provided by gravity. The upward propagating beam is also used as a slower beam (see Sec. IV). The properties of these beams are given in Table I, while the field gradients used for the MOT are $\{0.55, 0.32, 0.23\} \text{ G cm}^{-1}$ along the $\{x, y, z\}$ axes. The magnetic field gradient is produced by three different coil pairs in anti-Helmholz configuration. The choice of using three different coil pairs to generate the magnetic gradient is taken for experimental convenience. Indeed, a single coil pair oriented along the $x, y,$ or $z$ axes could also be used.

In order to characterize the performance of this MOT, we measure the flux, temperature, steady-state atom number, and lifetime. We then compare the MOT for fermionic strontium with one for bosons produced with the same apparatus. The results are summarized in Table II. In order to characterize the MOT we take absorption images using the $^1S_0 - ^1P_1$ transition (see inset of Fig. 7).

To record the loading of the MOT, we repeat such measurements for different loading times after switching the laser beams on [see Fig. 7(a)]. As the flux of atoms entering the lower chamber is constant, we fit these data with the function $N(t) = N_\infty (1 - e^{-t/t_{\text{load}}})$, where $N_\infty$ is the steady-state atom number, and $t_{\text{load}}$ is the $1/e$ loading time. We obtain from the fit $t_{\text{load}} = 6.4(1)\text{s}$ and $N_\infty = 8.41(4) \times 10^7$ atoms.

TABLE I. Properties of the MOT beams shown in Fig. 1. For each beam, we give the power of the cooling $||$ pumping beam, $1/e^2$ beam diameter $D$, and laser detuning sweep parameters: start detuning $\Delta_1$, stop detuning $\Delta_2$, and repetition rate $1/\Delta t$. Cooling and pumping beams have the same polarization, the same frequency modulation, and are combined within the same optical fibers before being sent to the atoms. The vertical MOT beam is focused $\sim 22\text{ cm}$ above the quadrupole center and we give the diameter of the beam $\sim 22\text{ cm}$ below it. The high power for the vertical beam is required to achieve high flux in the experiment, while the remaining parameters offer more flexibility and have been tuned to maximize the flux in the experiment.
TABLE II. Characteristics of the steady-state MOT, for both $^{87}$Sr and $^{88}$Sr isotopes (see also description in the main text). The steady-state MOT extents $\sigma_x$, $\sigma_y$, and $\sigma_z$ are Gaussian widths extracted by fitting a 2D Gaussian function to the images of the MOT. All uncertainties are given as the standard deviation from the fitted data.

<table>
<thead>
<tr>
<th></th>
<th>$^{87}$Sr MOT</th>
<th>$^{88}$Sr MOT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flux $\Phi$ (atoms/s)</td>
<td>$(1.31(2) \times 10^7)$</td>
<td>$(2.29(6) \times 10^6)$</td>
</tr>
<tr>
<td>$T_1$ (µK)</td>
<td>14.2(2)</td>
<td>12.6(1)</td>
</tr>
<tr>
<td>$T_2$ (µK)</td>
<td>7.5(2)</td>
<td>9(1)</td>
</tr>
<tr>
<td>$T_{\text{average}}$ (µK)</td>
<td>12.0(1)</td>
<td>11.4(4)</td>
</tr>
<tr>
<td>Width $\sigma_x$ (mm)</td>
<td>0.689(8)</td>
<td>0.632(8)</td>
</tr>
<tr>
<td>Width $\sigma_y$ (mm)</td>
<td>1.91(2)</td>
<td>2.82(2)</td>
</tr>
<tr>
<td>Atom number $N_\infty$</td>
<td>$(8.41(4) \times 10^7)$</td>
<td>$(100.8(9) \times 10^7)$</td>
</tr>
<tr>
<td>Peak density (atoms/cm$^3$)</td>
<td>$(2.12(4) \times 10^9)$</td>
<td>$(1.28(2) \times 10^{10}$)</td>
</tr>
<tr>
<td>1/e loading time $\tau_{\text{load}}$ (s)</td>
<td>6.4(1)</td>
<td>4.4(1)</td>
</tr>
<tr>
<td>1/e lifetime $\tau_{\text{life}}$ (s)</td>
<td>4.84(3)</td>
<td>4.6(1)</td>
</tr>
</tbody>
</table>

The steady-state MOT is operated with less detuning and without stirring beam, are addressed at a distance $y_0 = h\Delta I/(g_m\mu_B dB/dy)$ from the MOT quadrupole magnetic field center, where $\mu_B$ is the Bohr magneton. For $m_F = +9/2$, $|y_0| \approx 14$ mm below the center, which is much bigger than the vertical Gaussian width $\sigma_y$ of the MOT sample. This large distance relative to the MOT size implies that atoms always lie well below the quadrupole field center. Here they are constantly addressed by the vertical beam and optically pumped mainly into the $F=1$ ground state, $\mu_B F = 11/2$ excited state. Indeed, atoms within such MOTs can be optically pumped into states that are not trapped if they travel to the opposite side of the magnetic field center. Typically these losses are avoided by adding a second set of laser beams addressing the pumping transition. This transition generates ‘‘stirring,’’ a fast randomization of the population over all the $m_F$ states [52]. We apply this solution by using the pumping light described in Sec. IV with the parameters reported in Table I. Figure 7(b) compares the MOT’s decay with and without pumping beams. Surprisingly, we found that the presence of the pumping light does not affect the $^{87}$Sr MOT lifetime. While the flux increases by a factor of 2.2(3) the lifetime remains the same.

This behavior can be explained by the small gradients of the MOT quadrupole field and large detuning of the MOT beams, combined with the sample’s low temperature. Atoms slowed by the vertical beam and captured in the MOT are addressed at a distance $y_0 = h\Delta I/(g_m\mu_B dB/dy)$ from the MOT quadrupole magnetic field center, where $\mu_B$ is the Bohr magneton. For $m_F = +9/2$, $|y_0| \approx 14$ mm below the center, which is much bigger than the vertical Gaussian width $\sigma_y$ of the MOT sample. This large distance relative to the MOT size implies that atoms always lie well below the quadrupole field center. Here they are constantly addressed by the vertical beam and optically pumped mainly into the $m_F = +9/2$ state. Therefore the losses observed in other $^{87}$Sr MOTs that are operated with less detuning and without stirring beam, are not present here. The convenient polarization mechanism at play here, resulting from weak optical forces, gravity, and the large detuning of the MOT beams, has already been observed in MOTs of other elements such as dysprosium and erbium [53–55]. We note that, even if present, the pumping light is
unlikely to have a strong effect on the MOT because it is strongly detuned at its location (see Fig. 6). The role of the pumping light is to increase the atom flux \( \Phi \) from the slower stage.

VI. DISCUSSION AND CONCLUSION

Let us now compare the performance of our \(^{87}\text{Sr} \) MOT with the requirements for a superradiant active optical clock operating on the mHz-wide clock transition [24–26]. Such a clock is a bad-cavity laser, which uses as a gain medium a cloud of ultracold atoms within the mode of a high-finesse optical cavity. After being optically pumped to an excited state, these atoms can emit into the cavity mode, with a rate of emission enhanced via collective, superradiant emission. To sustain the emission of this laser, the excited population must be renewed. Atoms could be renewed by pumping them back into the excited state, which leads to unfavorable heating, or by bringing new excited state atoms into the superradiantly lasing ensemble. This translates into requirements for a stream of new incoming atoms needed for allowing the laser’s continuous-wave operation. For a laser operating on the \(^{1}\text{S}_0 \rightarrow ^{3}\text{P}_1 \) transition of strontium, we estimate that, without optical pumping, the flux of new atoms must be superior to \( 10^6 \) \(^{87}\text{Sr} \) atoms/s. Moreover, MOT atoms must be transported and loaded into the cavity mode, located somewhere outside of the MOT region, where they cannot be addressed by laser cooling light. Such transport could be achieved by a moving optical lattice [56]. For MOT atoms to be efficiently loaded into a lattice of reasonable depth, their temperature should be lower than \( \sim 10 \) \( \mu \)K. Our steady-state fermionic MOT satisfies these requirements and is an excellent candidate for feeding atoms into a superradiant laser system. A lower temperature or higher density can be achieved by using less light intensity within MOT beams, to the detriment of the atomic flux. A higher flux can be straightforwardly achieved by operating at a higher oven temperature and/or by using “repumper” beams to plug losses on the \(^{1}\text{S}_0 \rightarrow ^{1}\text{P}_1 \) cooling cycle [57].

To conclude, we have demonstrated a steady-state magneto-optical trap operating on the narrow intercombination line of strontium that works with the fermionic isotope \(^{87}\text{Sr} \), despite the additional challenges arising from its hyperfine structure. Two key ingredients were necessary: (1) the use of 2D SWAP molasses to collimate the atomic beam before atoms pass from the upper vacuum chamber to the next; and (2) a dual-frequency narrow-line Zeeman slower and MOT capable of slowing and capturing the incoming atoms. This MOT is well suited to provide atoms for the gain medium of a continuous-wave superradiant laser operating on strontium’s mHz-wide clock transition.

Note added: The raw data and the analysis tools used in this research can be found in Ref. [58].

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