

# Realization of Bose-Einstein Condensation with Lithium-7 Atoms

by

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Submitted to the Department of Physics  
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## Abstract

Ultra-cold atoms are atoms that are at a temperature close to absolute zero, typically at the order of micro Kelvin or lower. At these low temperatures, the quantum properties of the atoms, which are usually dominated by thermal effect at room temperature ( $\approx 300K$ ), becomes important and the atoms can form interesting new states of matter including Bose-Einstein condensation (BEC) for bosons and degenerate Fermi gas for fermions.

This thesis presents our work on developing and improving the techniques of trapping and cooling an ultra-cold cloud of Lithium-7 atoms and the realization of the Bose-Einstein condensation. The techniques used in this experiment includes Zeeman slower, magneto-optical trap (MOT), gray molasses, static magnetic trap, evaporative cooling, optical dipole trap (ODT), etc.

Thesis Supervisor: Wolfgang Ketterle

Title: Professor



# Acknowledgments

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# Introduction

Predicted in 1924-25 by Satyendra Nath Bose and Albert Einstein from Bose statistics, the Bose Einstein condensate is a phase of matter at ultra-cold temperature that emerges completely because of quantum effect. It was first produced in the laboratory in 1995-96 at the University of Colorado Boulder, Massachusetts Institute of Technology and Rice University using laser cooling and evaporative cooling techniques. Since then, people have been using it to study a lot of quantum effect. Among them, one effort is to simulate complex condensed matter systems using simplified and well controlled model systems created by loading BEC into optical lattices.

In our experiment, we use the Lithium-7 atoms to create Bose-Einstein condensate. Because of the lightness and several Feshbach resonances, the Lithium-7 atom has very fast dynamics and greater tunability, making it a perfect candidate for simulating and studying the phase diagrams of certain condensed matter models. The ultimate goal of the experiment is to study the anti-ferromagnetic phase in the anisotropic Heisenberg model ( $XXZ$  model) and the work in this thesis focuses on getting a Bose-Einstein condensate using Lithium-7, one of the important steps before studying the dynamics in an optical lattice.

The presentation of this thesis is divided into two chapters. In the first chapter, I discuss the theory of our experiment, including the Bose-Einstein condensate (BEC) and the various cooling and trapping techniques we use. The second chapter describes the setup of the experiment, the alignment and optimization procedure we developed and the experimental result we have got for each steps.



# Chapter 1

## Theory

In this chapter, I am going to describe the theories behind our experiment. It is divided into three parts. The first section explains the theories of cold Bose gas and the Bose-Einstein condensation relevant to the experiment. The second section briefly describes some important properties of the Lithium-7 atom. Finally, in the third section, the cooling, trapping and state manipulation techniques used in the experiment are presented, including Zeeman slower (1.3.1), MOT(1.3.2), gray molasses (1.3.3), magnetic trap (1.3.4) and optical dipole trap (1.3.5).

### 1.1 Bose-Einstein Condensate in Harmonic Trap

Every real particles can be classified as one of the two families according to their spins, fermions which have half integer spins and bosons which have integer spins. According to quantum field theory[1, 3] the many-particles wave function of identical particles must be symmetric or anti-symmetric under particle exchange for bosons or fermions respectively. For bosonic particles, because of the symmetry of the wave function, the possibility for particles to be in the same state is greatly enhances. As a result, boson gas at ultra-low temperature forms a Bose-Einstein condensate (BEC), in which almost all of the particles are condensed to the lowest energy state. In the following, the relevant properties of BECs such as critical temperature and density distribution are described.

Since the wave function of bosons is symmetric, multiple bosons can be in the same state. From this fact, the energy distribution can be calculated for bosons,

$$f(\varepsilon) = \frac{1}{e^{\beta(\varepsilon-\mu)} - 1}$$

Since the distribution has to be positive for all energy states, in particular the  $\varepsilon = 0$  ground state, we have  $\mu \geq 0$ . For all the state except the ground state, this sets a finite upper limit on the number of atoms in each state for a fixed temperature. Therefore, if the number of atoms exceeds a certain value, all the extra atoms will go into the ground state. These atoms condensed in the ground state are called the Bose Einstein condensate.

In order to calculate the atom number in the condensate as well as the critical temperature, we can estimate the maximum atom number in the excited states (thermal atoms) with an integral,

$$N_{th} = \int_0^\infty \frac{g(\varepsilon)}{e^{\beta\varepsilon} - 1} d\varepsilon$$

where  $g(\varepsilon)$  is the energy density of states. In our experiment, we create the BEC in a harmonic optical dipole trap (see 1.3.5) for which the energy density is,

$$\begin{aligned} g(\varepsilon) &= \frac{\varepsilon^2}{2\hbar^3\omega^3} \\ N_{th} &= \frac{1}{2\hbar^3\omega^3} \int_0^\infty \frac{\varepsilon^2}{e^{\beta\varepsilon} - 1} d\varepsilon \\ &= \frac{1}{2\hbar^3\omega^3\beta^3} \int_0^\infty \frac{x^2}{e^x - 1} dx \\ &= \frac{k_B^3 T^3}{2\hbar^3\omega^3} \zeta(3)\Gamma(3) \end{aligned}$$



The critical temperature of the transition, determined by  $N_{th} = N$ ,

$$\begin{aligned} T_C &= \frac{\hbar\omega}{k_B} \sqrt[3]{\frac{2N}{\zeta(3)\Gamma(3)}} \\ &= 0.9405 \frac{\hbar\omega \sqrt[3]{N}}{k_B} \end{aligned}$$

Condensate fraction (for large  $N$ ),

$$\begin{aligned} \frac{N_0}{N} &= 1 - \frac{N_{th}}{N} \\ &= 1 - \left(\frac{T}{T_C}\right)^3 \end{aligned}$$

## 1.2 Lithium-7 Atoms

The atoms used in this experiment is the Bosonic isotope of Lithium, Lithium-7. In this section, I will present some properties of the Lithium-7 atoms that are important for the cooling and trapping techniques we use in the experiment.

One of the most important properties for laser manipulation of cold atoms is the energy levels. As all other alkali atoms, Lithium-7 has one unpaired valence electron with a  $nS$  ground state and  $nP$  excited states, where  $n = 2$  for Lithium-7. The fine structure split the excited states into  $2^2P_{1/2}$  and  $2^2P_{3/2}$  states corresponding to the  $D1$  and  $D2$  lines that are about  $10GHz$  apart. On top of these, the hyperfine structure caused by the nucleus spin ( $I = 3/2$  for Lithium-7) further split each of these states into different levels with different total angular momentum  $F$ . the precise frequencies of these transitions measured in the experiment are shown in figure 1-1

In an external magnetic field, each  $F \neq 0$  levels will be split because of the Zeeman effect. At low field, this splitting is described by the Landé  $g$ -factor  $\Delta = g_F \mu_B m_F B_z$  as shown in table 1.1. At higher magnetic field, different hyperfine levels starts to mix and the Zeeman effect is no-longer linear. Most importantly, for  $2^2S_{1/2}$  ground

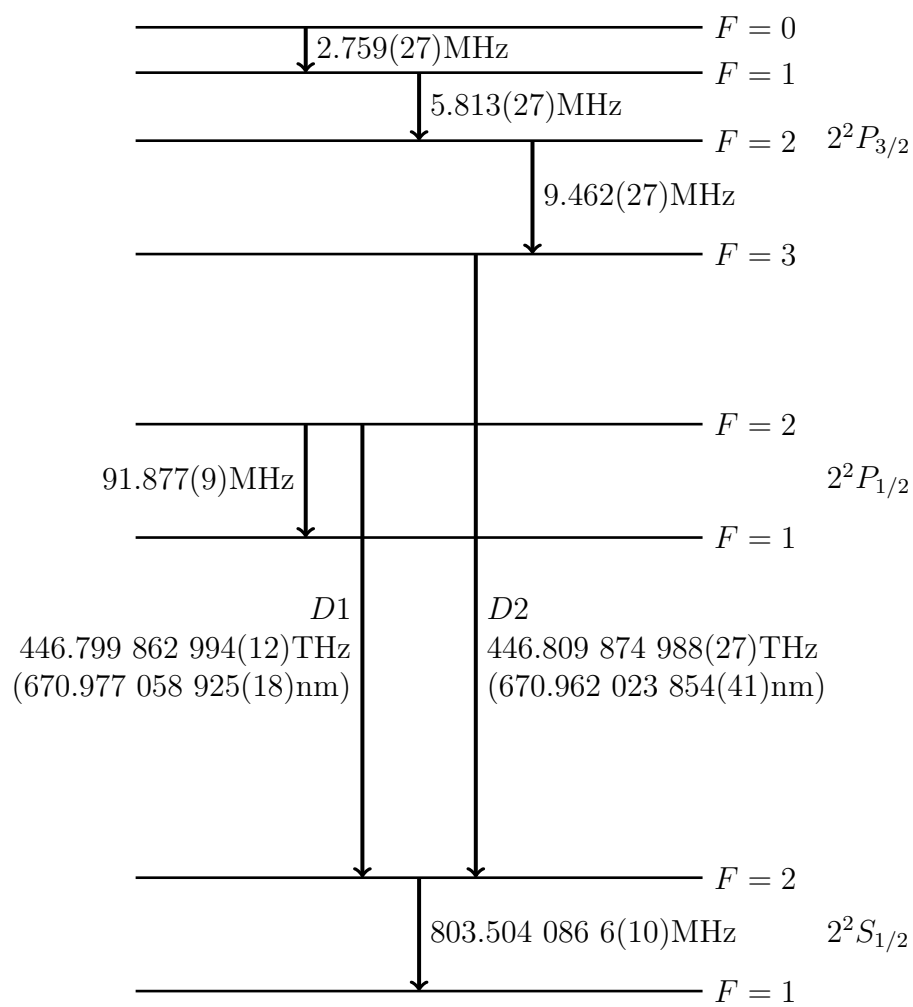


Figure 1-1: Energy levels of Lithium-7[4]

Table 1.1: Landé  $g$ -factors of Lithium-7

Fine Structure	$F$	$g$ -factor
$2^2S_{1/2}$	2	$\frac{1}{2}$
	1	$-\frac{1}{2}$
$2^2P_{1/2}$	2	$\frac{1}{6}$
	1	$-\frac{1}{6}$
$2^2P_{3/2}$	1, 2, 3	$\frac{2}{3}$

states the  $F = 1$  states have a negative slope at high magnetic field and all the  $F = 2$  states except  $m_F = -2$  has a positive slope. (Note that  $F$  and  $m_F$  are no-longer good quantum numbers at high magnetic field but there are states at high field that are adiabatically connected with these  $F$  and  $m_F$  eigenstates). The states with a positive Zeeman shift are called magnetically trappable states since they can be trapped by a static magnetic field with a minimum in the center. Only  $|2, 2\rangle$  and  $|2, 1\rangle$  are both trappable at high field and low field.

### 1.3 Cooling and Trapping Theory

The atoms used in an alkali atom experiment often come from an atom beam coming out of a oven kept above the melting temperature of the metal. The oven used in this experiment operates at 485°C producing an atom beam traveling at several hundred meters per second. In order to achieve low temperature and high density, several stages of slowing, trapping and cooling are implemented in this experiment which finally bring the atoms down to the Bose-Einstein condensate condition. In this section, I am going to talk about the theory behind these techniques we are using in our experiment.

### 1.3.1 Zeeman Slower

The atoms coming out of the oven has an average velocity determined by the Maxwell-Boltzmann distribution,

$$\bar{v} = \sqrt{\frac{k_B T}{2\pi m}}$$

We can slow down the atoms using the recoil of photon scattering by shining resonance light to the beam. However, since the atomic transition resonance is very narrow ( $\Gamma = 2\pi \cdot 5.9\text{MHz}$ ) compared to the Doppler shift ( $\Delta\nu = \nu v/c \approx 590\text{MHz}$ ), the atom will soon shift out of resonance once it slows down. There are several ways to solve this problem. One way is to modulate the laser frequency, either swiping with time (frequency chirping) or making it broadband (white light). The Zeeman slower solves the problem by changing the resonance of the atom with Zeeman effect. This is the most popular way for alkali atoms which usually have a cycling transition with linear Zeeman effect in a large range.

The Zeeman slower has a magnetic field aligned with the atom beam with a spatially variant amplitude. By shining circularly polarized light against the atom beam, the atoms in the beam are optically pumped to one of the two stretch states determined by the relative direction of the field and the light polarization. If the magnetic field is changing in a way such that the Zeeman shift of the atom follows the Doppler shift when atoms slow down, the atom beam will be always in resonance and continuously being slowed down, i.e.

$$g_F m_F \mu_B B = \frac{v}{c} \nu + \Delta$$

For constant acceleration, this implies,

$$B = \frac{\sqrt{v_0^2 - 2ax}}{g_F m_F \mu_B c} \nu + B_0$$

The maximum deceleration of the Zeeman slower is limited by the maximum scattering rate, therefore the life time of the excited state,

$$a_{max} = \frac{h\Gamma}{2\lambda}$$

where  $\Gamma$  is the linewidth of the transition.

### 1.3.2 Magneto-Optical Trap (MOT)

As in most cold atom experiment, our experiment starts with loading Zeeman slowed atom into a magneto-optical trap (MOT). MOT is a technique that uses laser and magnetic field gradient to provides both molasses cooling and confinement. In order to understand how MOT works, we will consider a one-dimensional model consisting of a spatially varying magnetic field  $B = B_x = B'x$ , two counter propagating, red-detuned, circularly polarized light with atoms sitting close to the zero of the magnetic field.

The cooling and trapping effect in a MOT can be understood separately. Let us first consider an atom in the center of the MOT moving in the  $+x$  direction. Since the light is red-detuned, the Doppler effect will shift the atom closer to the light in  $-x$  direction and farther from the light in  $+x$  direction. The atom will therefore scatters more photons from the  $-x$  light and experience an average force proportional but in the opposite direction with its velocity. Such a force will damp the motion of the atom and therefore cool down the temperature of the cloud.

In order to understand the source of the trapping force in a MOT, we can consider an atom with a  $F = 0$  ground state and  $F = 1$  excited states. As we can see in the schematic of this model (Figure 1-2), since the magnetic field changes its direction in the center, both laser beams are  $\sigma^-$  light on the side it is coming from and therefore is shifted closer to resonance by the Zeeman effect. Therefore, atoms shifted from the center will scatter more photon from this direction and on average feel a net force

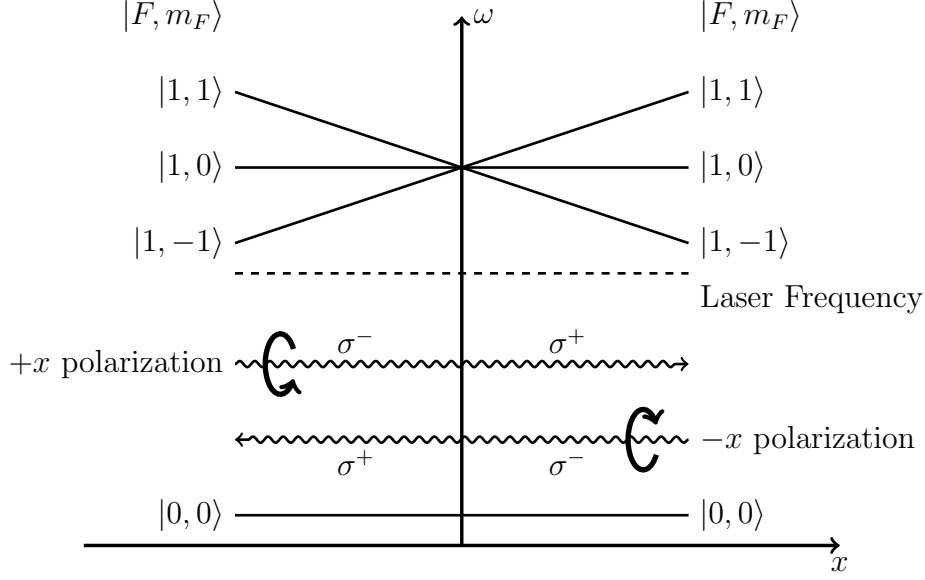


Figure 1-2: Schematic of a 1D MOT with a  $F = 0$  to  $F = 1$  transition.

from recoil pointing toward the center of the trap.

The performance of the MOT is usually limited by the random kick from scattered photons (Doppler limit on temperature) and light induced collisions (limits the density and atom number).

### 1.3.3 Gray Molasses

Since the  $2^2P_{3/2}$  excited state of Lithium-7 we use for molasses cooling in the MOT does not have resolved hyperfine structure (Figure 1-1). We cannot perform conventional polarization gradient cooling using the  $D2$  line. Instead, we implement a gray molasses on the  $D1$  transition. This scheme was first studied and demonstrated by Christophe Salomon[2] to achieve sub-Doppler temperature on some alkali isotopes that are not amenable for conventional polarization cooling ( $^7\text{Li}$ ,  $^6\text{Li}$  and  $^{40}\text{K}$ ). In this section, I am going to present a semi-quantitative explanation of this cooling scheme. See the original paper[2] for more detail discussion.

The gray molasses cooling uses three pairs of counter-propagating beams that can

cool the cloud in all three dimensions. Like what we did for MOT, we will consider a  $1 - D$  case for simplicity. The model we will use has a pair of counter-propagating beams, each has a high intensity pumper, with a large blue detuning  $\delta$  from the  $F = 2$  ( $|2\rangle$ ) to  $F' = 2$  ( $|3\rangle$ )  $D1$  transition, and a low intensity repumper, blue detuned from the  $F = 1$  ( $|1\rangle$ ) to  $F' = 2$   $D1$  transition by the same detuning  $\delta$ . Let the Rabi frequency for the pumper and the repumper be  $\Omega_2(x)$  and  $\Omega_1(x)$  respectively.

Just as any other laser cooling method, the performance of the gray molasses is determined by the balance between cooling force and heating from photon scattering. Since the two beams (pumper and repumper) satisfies the Raman resonance condition, the system has a dark state which is the superposition of  $|1\rangle$  and  $|2\rangle$ ,

$$|NC\rangle = \frac{\Omega_2|1\rangle - \Omega_1|2\rangle}{\sqrt{\Omega_1^2 + \Omega_2^2}}$$

which will be the state most of the atoms will be pumped into. (The state is dark for stationary atoms and has a non-zero but small scattering rate for moving atoms.) Since the scattering is suppressed by the presence of a dark state, the gray molasses produce little heating.

The cooling effect of the gray molasses can be understood in the dressed atom picture for the atom and the pumper beam as shown in figure 1-3, where,

$$\begin{aligned} |2'\rangle &= \frac{\delta^2}{\delta^2 + \Omega_2(x)^2}(|2\rangle - i\Omega_2(x)|3\rangle) \\ |3'\rangle &= \frac{\delta^2}{\delta^2 + \Omega_2(x)^2}(|3\rangle - i\Omega_2(x)|2\rangle) \end{aligned}$$

Since the detuning  $\delta$  is large, the mixing between  $|2\rangle$  and  $|3\rangle$  is small leaving  $|3'\rangle$  a short lifetime and  $|2'\rangle$  a longer lifetime. The two counter-propagating pumper beams create a standing wave which causes periodic AC-Stark shift in the dressed  $|2'\rangle$  and  $|3'\rangle$  states. For a repumper with the same or slightly smaller blue detuning compare to the pumper. It is the most closest to the resonance with the dressed states at the

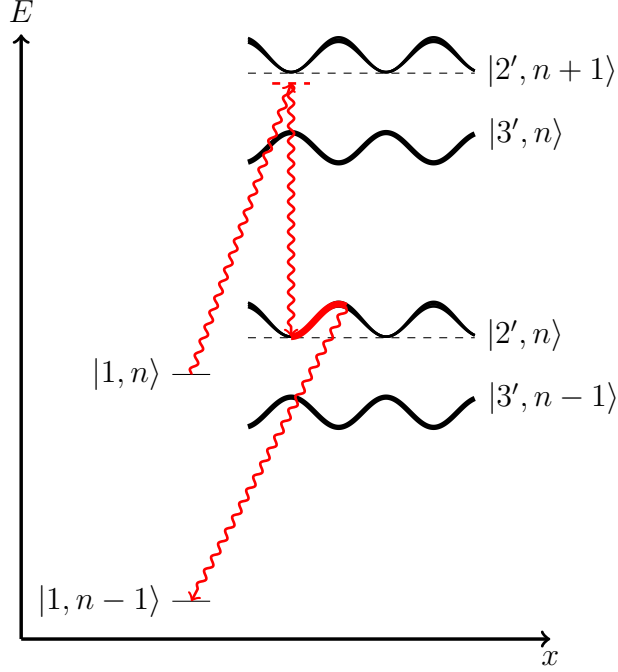


Figure 1-3: Schematic of a 1D Gray Molasses.[2]

nodes of the standing wave where the mixing is minimized. Therefore, the repumper will most likely pumps atoms into the  $|2'\rangle$  state (through  $|3'\rangle$  as shown in figure 1-3) at the node of the standing wave and the atoms will most likely exit the state at the anti-node of the standing wave where the lifetime of  $|2'\rangle$  is the shortest because of the maximum mixing. Since potential energy for atoms in  $|2'\rangle$  is minimized at the nodes and maximized at the anti-nodes the atoms will on average feel a damping force by climbing up the hill in  $|2'\rangle$ . The height of the hill, therefore the cooling force, is proportional to the laser intensity. Combined with the low scattering rate due to the dark state, the gray molasses can be used to quickly cool the cloud after MOT to sub-Doppler temperature.

#### 1.3.4 Evaporation in Static Magnetic Trap

The density and temperature of a laser cooled cloud is usually limited by the scattering of photons. Therefore, in order to reach quantum degeneracy, one usually need to use a way that can cool the atoms without shining resonant light onto them. The



way we use in our experiment is evaporative cooling in a magnetic trap and an optical dipole trap. In this section, I will describe the theory of static magnetic quadrupole trap and radio frequency evaporation and the theory of evaporation in an optical dipole trap will be discussed in the next section.

The static magnetic trap, as the name suggested, is created using static magnetic field and the Zeeman effect of the atoms. Assuming a monotonic the Zeeman effect, the atoms will be trapped at an extremum of the field. Since the magnetic field in the vacuum satisfies the source-less Maxwell equation, we have,

$$\begin{aligned}
0 &= \nabla \times (\nabla \times B) \\
&= \nabla(\nabla \cdot B) - \nabla^2 B \\
&= -\nabla^2 B \\
\nabla^2(B^2) &= 2\nabla(B \cdot \nabla B) \\
&= 2(\nabla B)^2 + 2B \cdot \nabla^2 B \\
&= 2(\nabla B)^2 \\
&\geq 0
\end{aligned}$$

Therefore, the only local extremum that is possible for a static magnetic field in the vacuum is a local minimum and only states with a positive Zeeman shift (low field seeking) can be trapped magnetically.

The static magnetic field with a local minimum in the center can be implemented in many different ways. The configuration used in this experiment is a quadrupole field created by a set of Helmholtz coils carrying opposite current, which provides the tighter confinement compare to other configurations. Closed to the center of the

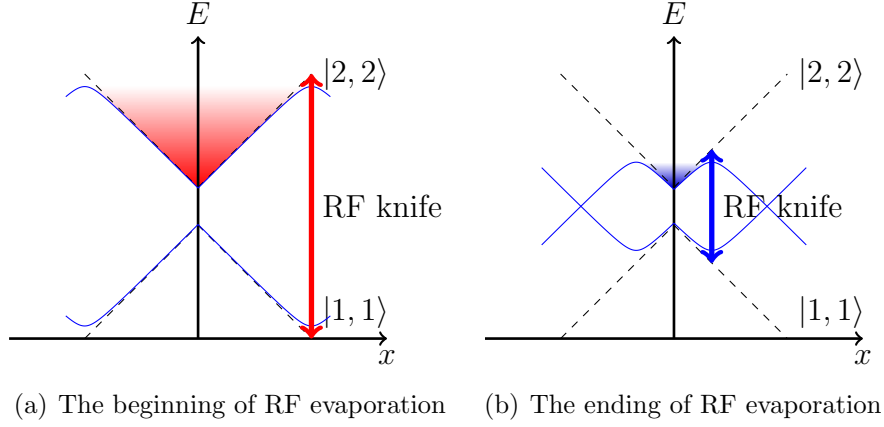


Figure 1-4: RF evaporation in magnetic trap. The temperature goes down and the phase space density goes up during evaporation.

trap, the magnetic field can be expressed as,

$$B_x = B'x$$

$$B_y = B'y$$

$$B_z = -2B'z$$

Since the magnetic field is 0 at the center of the trap where all the Zeeman levels are degenerate, spin-flip of atoms (Majorana loss) occurs in the center of a quadrupole trap. This problem is fixed in the experiment by plugging the hole with a repulsive blue detuned laser beam.

After trapping the atoms with the magnetic field, we use radio frequency induced spin-flip to evaporate the cloud. As shown in figure 1-4, the  $|2, 2\rangle$  has a positive Zeeman effect whereas the  $|1, 1\rangle$  state has a negative Zeeman shift. After all the atoms are trapped in the  $|2, 2\rangle$  state, a RF field that is blue detuned to the zero field resonance is applied which mixes the two states and turns the potential curve around at the resonance field. As a result, the depth of the trap is lowered and atoms which has higher energy can now escape from the trap. If the RF knife is placed much higher than the thermal energy  $k_B T$ , each leaving atoms will carry away energy a lot higher

than the averaged value leaving a lower average energy, therefore lower temperature, for the atoms remaining in the trap. After the cloud gets colder, less and less atoms will have the energy to pass the RF knife, at which point the RF frequency can be sweep down to accelerate the evaporation. In order for the evaporative cooling to work efficiently, a high elastic collision rate (high rate for creating atoms with high energy) is also necessary.

### 1.3.5 Evaporation in Optical Dipole Trap

Although RF evaporation in the magnetic trap is very convenient and efficient, it does have some limitations,

1. The atoms have to be in a hyperfine state.
2. Magnetic field cannot be freely changed.
3. Hard to manipulate with RF field (due to the field gradient across the cloud.)

Moreover, for Lithium-7, the trappable  $|2, 2\rangle$  state has a negative scattering length ( $-27a_0$ ) making it impossible to create a stable Bose-Einstein condensate with a large atom number. In order to solve the problem, we switch to an optical dipole trap and do the final evaporation there.

The optical dipole trap (ODT) is created using the negative AC Stark shift of a red detuned laser beam. Since the AC Stark effect is proportional to  $\delta^{-1}$  (where  $\delta$  is the detuning) and the scattering (heating) is proportional to  $\delta^{-2}$ , high power beams with huge detuning ( $\approx 100THz$ ) is often used in order to introduce the least amount of heating for the same trapping potential.

Evaporation in an ODT is done simply by lowering the potential (beam power). Since a positive and big scattering length is necessary for efficient evaporation to BEC, we spin-flip the atoms into  $|1, 1\rangle$  and evaporate close to a Feshbach resonance with an appropriate scattering length.



# Chapter 2

## Experimental Setup and Results

In this chapter, I will describe our experimental setup and results. It includes a brief discussion of our optical tables setup (2.1 and 2.2), the implementation, optimization and performance of each steps (2.3, 2.4, 2.5, 2.6 and 2.7) and finally some basic characteristic of our BEC (??).

### 2.1 Laser System

The laser table is where we prepare all the light resonance with the Lithium-7 transition ( $\approx 671nm$ ). As discussed in the previous chapter, there are four distinct lines that we need in the experiment, the combination of  $D1$ ,  $D2$  with  $F1$ ,  $F2$ . At a certain time, we need either  $D1$  or  $D2$  light with the probability of using both  $F1$  and  $F2$  at the same time and the laser table is designed to do just this.

As shown in figure 2-1, since the separation between the  $D1$  and  $D2$  line ( $\approx 10GHz$ ) is larger than the range of common optical frequency shifters (e.g. AOMs), the  $D1$  and  $D2$  light are created separately using two diode lasers (TA pro and DL pro in figure 2-1). Both of these lasers are actively locked to the appropriate atomic transitions using saturated absorption spectroscopy to better than 2MHz. For the  $D2$  path, part of the light ( $\approx 80mW$ ) is red shifted and used as the Zeeman slower light ( $ZSL$  in the figure) and the rest goes to the  $D2$ ,  $D1$  selecting switch, which uses two

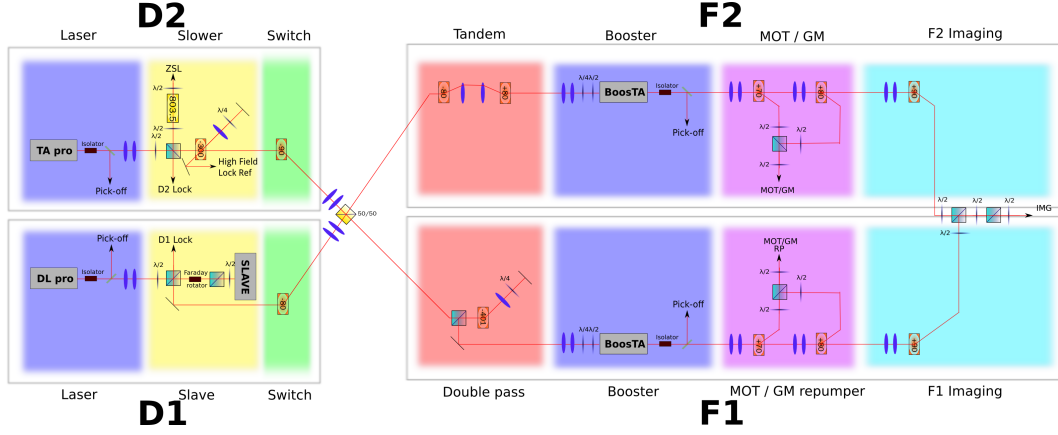


Figure 2-1: Schematic of the laser table design

AOMs (acousto-optical modulator) and a 50-50 cube to feed both the  $F1$  and  $F2$  path with either  $D1$  and  $D2$  light. For the  $D1$  path, a slave diode is used to amplify the light before going to the switch.

In the  $F1$ ,  $F2$  path, a tandem and a double pass are used to continuously shift the frequency of the laser (the double pass is also used to shift the light from  $F2$  to  $F1$ ). After that, the light in each path is amplified using tapered amplifier and then go through a polarization switch consists of two AOMs and a polarization beam splitter which can but the light into polarization maintaining fibers (MOT/GM and MOT/GM RP) with two orthogonal linear polarization's (the use of these beams is further described in section 2.2.2). At the end of the chain, two more AOMs are used to control the light we put into the imaging fiber so that we can image on any of the four transitions. Each fibers also has a mechanical shutter that is used to block any possible leaking light that may cause heating.



Figure 2-2: Top View of the Vacuum Chamber (Top/Bottom MOT/Gray Molasses beams not included, not to scale).

## 2.2 Vacuum Chamber and Main Coils Configuration

In this section, I am going to describe some important setup on the machine table where our vacuum chamber sits on.

### 2.2.1 Optical Access of the Vacuum Chamber

The windows and connections except the top and bottom window on our main vacuum chamber is shown in figure 2-2. There are in total 6 MOT/Gray Molasses beams, 2 ODT beams, 3 Imaging beams, a plug and a Zeeman slower beam going into the chamber. When multiple colors need to be sent in through the same window, they are combined outside the vacuum chamber using appropriate dichroic.



Figure 2-3: MOT/Gray Molasses Cage.

### 2.2.2 MOT-Gray Molasses Cage

The delivery of the MOT and gray molasses beam from the two in-coupler on the laser table to the six output on the machine table is done with a polarization maintaining evanescent wave fiber splitter which takes the light from the two input fibers and split them equally into the six output fibers. However, the MOT beams and the gray molasses beams have different requirement in beam sizes. On one hand, the MOT needs a bigger beam size for a bigger capture volume. On the other hand, as discussed earlier, for the gray molasses to work effectively, high intensity, therefore smaller beam size, is required. In order to have a different size for the two beams coming out of the same fiber, we send in the two beams with orthogonal polarization's (figure 2-1) and add polarization dependent beam expanders after the fiber output to shape the two beams differently. This beam shaping is done with a cage system shown in figure 2-3. The upper beam path is used for expanding the MOT beam whereas the lower path shapes the gray molasses beam to a smaller size. Each of the four lenses can slide in the cage, which are used to tweak the size and divergence of the beams. The alignment between beams is done with the two bottom mirrors and the two half wave-plates between the two polarization beam splitter cubes are used to balance the intensities.



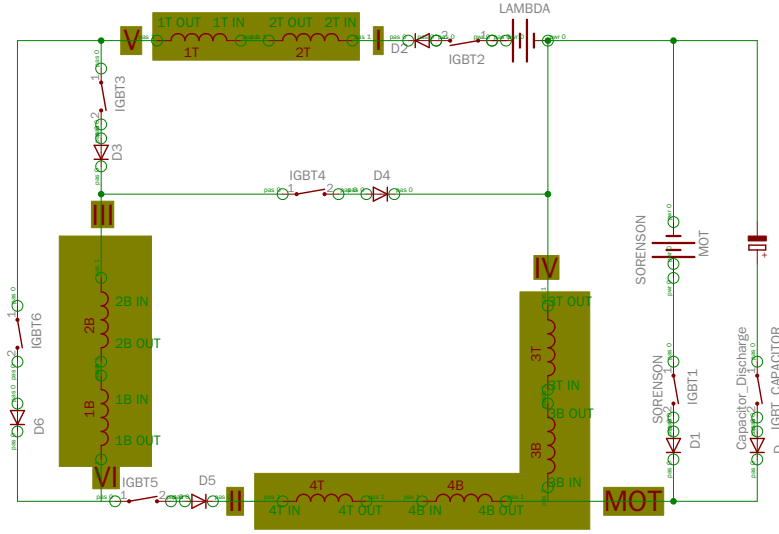


Figure 2-4: Main Coils Control Circuit

### 2.2.3 Main Coils Configuration

In our experiment, a lot of different field configurations are used for different purpose. All of these configurations have to be implemented using our main coil which has five independent layers closed to the Helmholtz condition on each sides with sub millisecond switching time between them. A modified H-bridge (figure 2-4) was build with IGBTs in order to accomplish these goals using the minimum amount of independent layers, IGBTs and power supplies. The list of possible configurations are shown in table 2.1.

## 2.3 Magneto-Optical Trap (MOT) and Compressed-MOT

The MOT loading is optimized by changing the parameters of the lasers to maximize the number of atoms in the MOT after 6 seconds of loading. The optimum settings we have found are listed in table 2.2 and the performance of our MOT can be found

Closed switches(IGBTs)	Usage of the power supply	
	Lambda	Sorenson
1, 3, 5	-	Quadrupole field with layer 3 for MOT and ODT clean up (2.7)
2, 3, 5	Quadrupole field with layer 1, 2, 3 and 4 for magnetic trap	-
1, 2, 4, 6	Homogeneous Feshbach resonance field with layer 1 and 2	Field gradient on top of the bias field with layer 3 for tilt and levitation.

Table 2.1: List of coil configurations

MOT power	Repumper power	MOT detuning	Repumper detuning	MOT Current
16mW	7mW	-38.5mW	-28mW	57A

Table 2.2: MOT parameters

in table 2.3.

Since our MOT is mainly optimized for loading rate, it is not necessarily optimized for density and cooling. Therefore, we added a compress-MOT (CMOT) step after the MOT in order to increase the density and decrease the temperature. In the CMOT step, we ramp the MOT frequency closer to resonance in 4.5ms and decrease the intensity of the repumper. As a result, the cloud is compressed by radiation pressure and pumped to the  $F1$  state. The temperature of the cloud is also decreased because of the decrease in scattering. Figure 2-5 shows the F pumping effect in the CMOT and the performance of the CMOT is listed in table 2.3.

## 2.4 Gray Molasses

The gray molasses step is used to cool the CMOT cloud further before transferring to the magnetic trap for evaporative cooling. Since the gray molasses beams have a smaller size ( $d \approx 5\text{mm}$ ), the beams need to be carefully aligned with the cloud

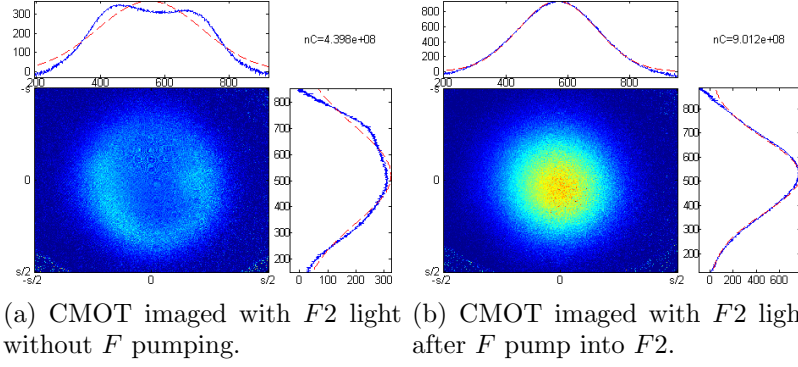


Figure 2-5: Image of the CMOT. Atoms are pumped into the  $F1$  states.

Step	Atom Numbers	Temperature(K)	Density( $\text{cm}^{-3}$ )
Oven	-	760	-
Zeeman Slower	-	0.5	-
MOT	$2 \cdot 10^{10}$	$1.5 \cdot 10^{-3}$	$1 \cdot 10^{11}$
CMOT	$2 \cdot 10^{10}$	$1 \cdot 10^{-3}$	$2 \cdot 10^{11}$
Gray Molasses	$1 \cdot 10^{10}$	$1 \cdot 10^{-4}$	$2 \cdot 10^{11}$

Table 2.3: Laser cooling performance.

which has a similar diameter. In our experiment, it is done putting  $F2$  light in each gray molasses beam individually and imaging the overlap between the beam and the CMOT cloud directly. Figure 2-6(a) shows the CMOT atoms after pumped into  $F2$  state imaging with  $F2$  light. After we pump the cloud with  $F2$  light from one of the gray molasses beam path, the image looks like figure 2-6(b), in which the missing part is pumped back into  $F1$  by the gray molasses beam and is invisible on the image. We then maximum the overlap between the beam and the cloud by minimize the atoms left in the image. Figure 2-6(c) shows the same image after the gray molasses beam is properly aligned. After we repeated this on all the gray molasses beams, we took time of flight images while scanning the frequencies of the light. As shown in figure 2-7, a clear decrease in temperature (a decrease in time of flight size) can be seen just as what we expected from the theory 1.3.3 and the experimental result from other groups[2].

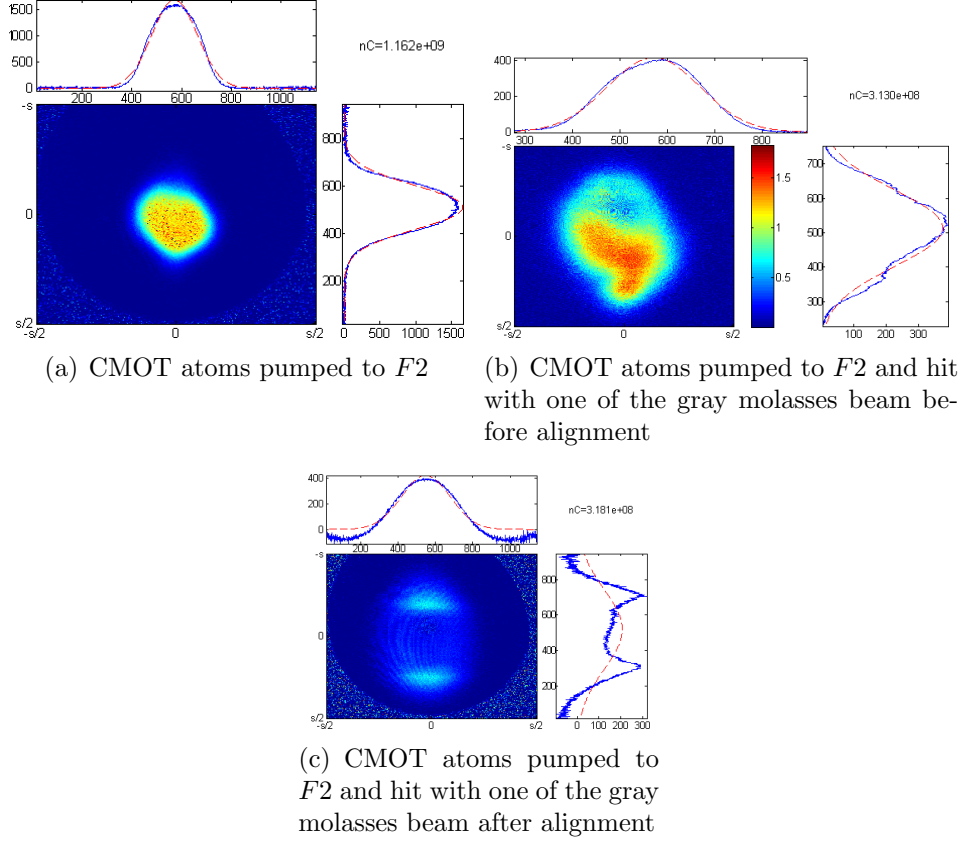


Figure 2-6: Images used to align the gray molasses beams.

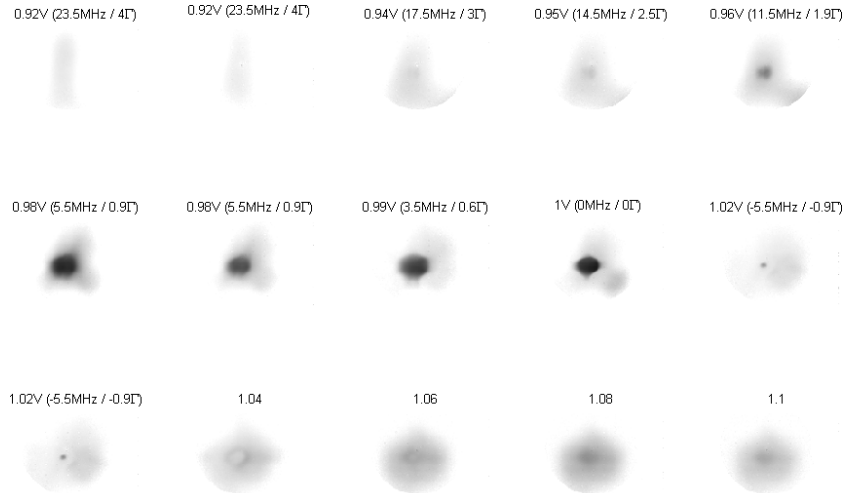


Figure 2-7: Time of flight image for gray molasses with different relative detuning (in MHz) between the pumper and repumper. (The voltages in the image is the control voltage we use to tweak the frequency.)

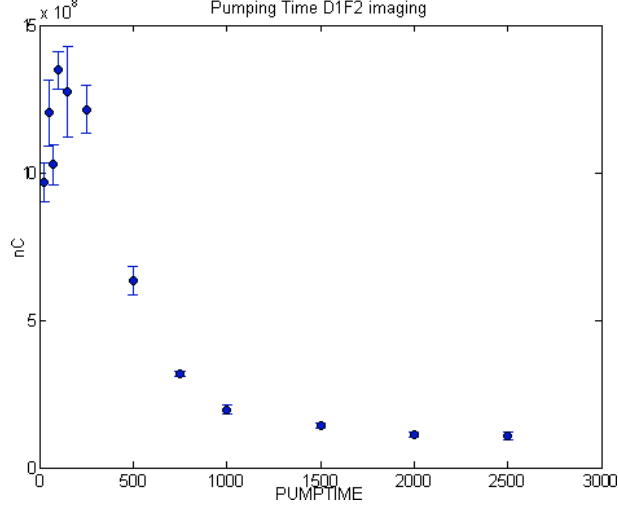


Figure 2-8:

## 2.5 Dark State Pumping

At the end of Laser cooling, the atoms are distributed in different hyperfine levels not all of which are trappable. In order to transfer to and evaporate in the magnetic trap, all the atoms have to be in a single trappable state. Making use of the dark state in the  $D1$  line with  $\sigma^+$  light, we use the dark state pumping to transfer the atoms into the  $|2, 2\rangle$  state without significant heating. In order to minimize re-scattering of the pumping light, the pumping light is blue detuned by  $\delta_{F1} = 20\text{MHz}$  and  $\delta_{F2} = 34\text{MHz}$ . Figure 2-8 shows the atoms after different pumping time imaged with  $D1F2 \sigma^+$  light. The initial increase in atom number shows atoms being pumped into  $F2$  and the slower decay is when they are going into the  $|2, 2\rangle$  state invisible to the imaging light.

## 2.6 Magnetic Trap

When the atoms are in the magnetic trap, the center of the trap need to be plugged in order to avoid Majorana loss. This is done with a 10W 532nm green laser beam, focused to  $20\mu\text{m}$  at the center of the trap. In order to keep it aligned for different trap gradient, we zero the transverse magnetic field by measuring the center position

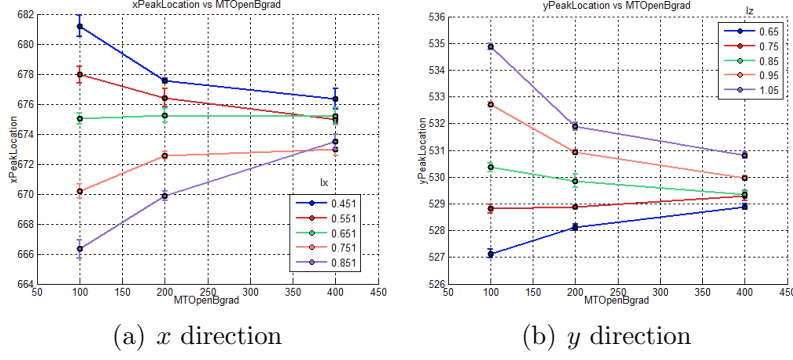


Figure 2-9: Precise Field Zeroing in Magnetic trap

of a small cloud in the magnetic trap with different gradients and different bias field. Figure 2-9 shows the result of this measurement. We pick the bias field with the smallest displacement at different gradient, for which the center of the trap moves by smaller than one pixel ( $20\mu m$ ). Figure 2-10 shows the number of atom after the magnetic trap, the saturation of the atom number with 6W of plug power shows that we have successfully suppressed the Majorana loss. Due to the high inelastic collision rate, low elastic collision rate of Lithium-7 and the atom number fluctuation in our experiment, it is hard to optimize the RF evaporation purely experimentally. Instead, the evaporation in our experiment is optimized using numeric simulation. After 2.5s of RF evaporation, we are left with  $6 \cdot 10^7$  atoms with a temperature of  $4\mu K$  and a density of  $10^{12} cm^{-3}$ .

## 2.7 BEC in Optical Dipole Trap

After evaporation in the magnetic trap, we transfer the atoms into a optical dipole trap (ODT) created with two 15W 1064nm laser beams. The transfer need to be adiabatic in order to minimize heating and maximize transfer efficiency. After exploring different transfer schemes, the best method we have found is to turn on the ODT in full power before evaporating in the magnetic trap and ramp down the magnetic trap after evaporation in 200ms. In this way, we can accumulate  $2 \cdot 10^7$  atoms in the ODT at a temperature of  $20\mu K$ .

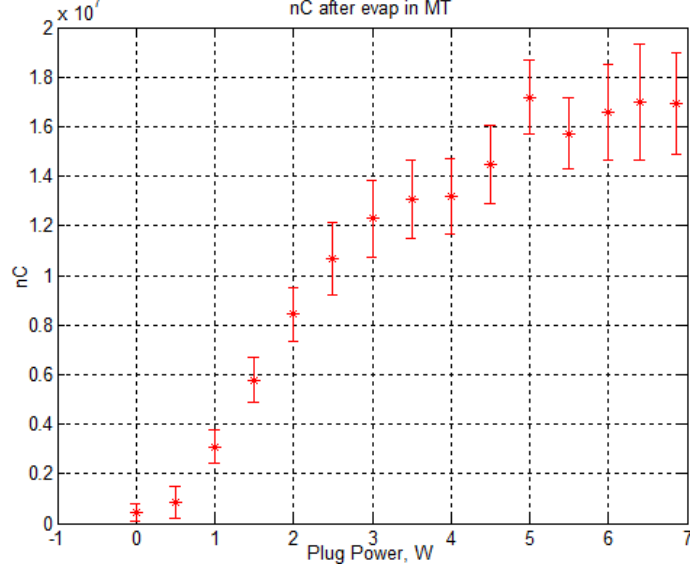


Figure 2-10: Saturation of the Plug Laser Power.

We put the atoms from  $|2, 2\rangle$  state to  $|1, 1\rangle$  state using a Landau-Zener sweep with RF frequency. Figure 2-11 shows the atom number in  $|2, 2\rangle$  versus starting frequency of a 1MHz wide RF scan in a magnetic field about 1G. We hit the resonance at 806.25MHz and the transfer efficiency is better than 80%.

The evaporation in the ODT is aided by a Feshbach resonance. We measure the resonance using the increase in three-body loss rate. Figure 2-12 shows the atom number after a certain hold time in the ODT with different Feshbach field. The resonance occurs at 285A. By comparing the resonance point with know data for the scattering length around the resonance, we do our evaporation using a Feshbach current of  $\approx 260$ A corresponding to a reasonably large scattering length ( $\approx 100a_0$ ).

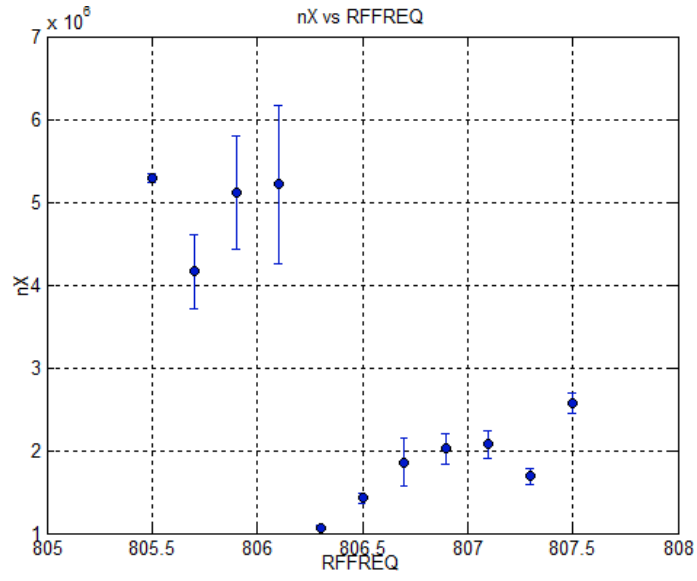


Figure 2-11: Landau-Zener Sweep.

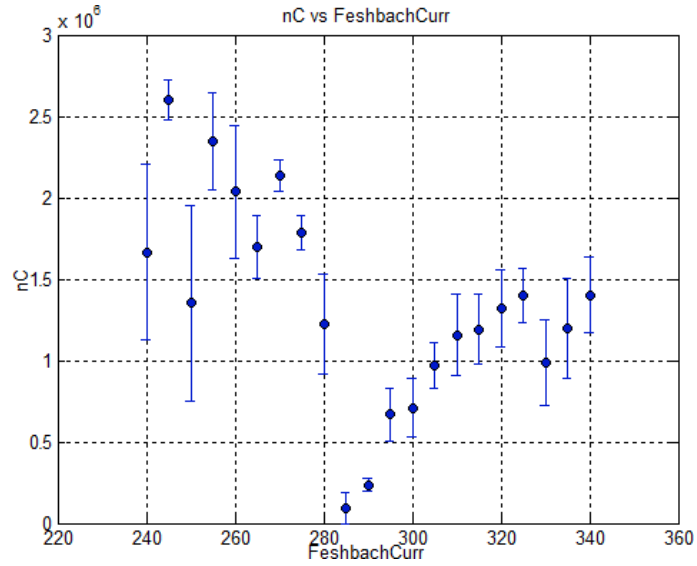


Figure 2-12: Feshbach Resonance in  $|1, 1\rangle$  state.



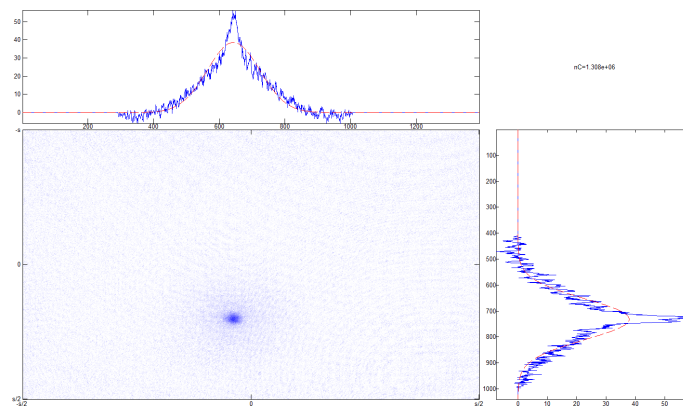


Figure 2-13: BEC with thermal wing.



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