

# Demonstration of a quantum logic gate in a cryogenic surface-electrode ion trap

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We demonstrate quantum control techniques for a single trapped ion in a cryogenic, surface-electrode trap. A narrow optical transition of  $\text{Sr}^+$  along with the ground and first excited motional states of the harmonic trapping potential form a two-qubit system. The optical qubit transition is susceptible to magnetic field fluctuations, which we stabilize with a simple and compact method using superconducting rings. Decoherence of the motional qubit is suppressed by the cryogenic environment. AC Stark shift correction is accomplished by controlling the laser phase in the pulse sequencer, eliminating the need for an additional laser. Quantum process tomography is implemented on atomic and motional states using conditional pulse sequences. With these techniques we demonstrate a Cirac-Zoller Controlled-NOT gate in a single ion with a mean fidelity of 91(1)%.

## I. INTRODUCTION

Trapped ions are promising candidates for realizing large-scale quantum computation [1, 2]. Significant progress has been made in demonstrating the fundamental ingredients of a quantum processor, with much progress in gate fidelities [3] and multi-ion entanglement [4, 5]. In recent years there has been increasing interest in microfabricated surface-electrode traps, due to their inherent scalability [6, 7]. However, quantum gates have yet to be demonstrated in such systems. An issue with miniaturization of traps is that anomalous heating of the ion's motional state scales unfavorably with trap size [8], potentially limiting gate fidelity in traps of suitable dimensions for scalability [9]. Recently, it has been shown that by cooling to cryogenic temperatures, the heating rate can be reduced by several orders of magnitude from room-temperature values [10], thus providing one potential solution to this problem. In this work, we demonstrate a quantum gate in a microfabricated surface-electrode ion trap that is operated in a cryogenic environment, and present some control techniques developed for this experiment.

We implement a Cirac-Zoller controlled-NOT gate using qubits represented by the atomic and motional states of a single ion. The  $S \leftrightarrow D$  optical transition in  $^{88}\text{Sr}^+$  is used as one of the qubits. The motional ground state and first excited state of the ion in the harmonic trap potential forms the second qubit. The optical transition has the advantage of a long lifetime while requiring only a single laser (unlike hyperfine qubits), but the qubit is first-order Zeeman sensitive, which makes it susceptible to magnetic field noise. Taking advantage of the cryogenic environment, we stabilize the magnetic field using a pair of superconducting rings [11]. Since the  $\text{Sr}^+$  ion qubit is not an ideal two-level system, the coupling between the sideband and carrier transitions causes level shifts known as the AC Stark shift, which must be corrected. In previ-

ous work, this has been accomplished with an additional laser field with the opposite detuning to cancel out the shift [12]. Here, to reduce the experimental complexity of the additional acousto-optical modulators (AOMs) and optics required, Stark shift corrections are implemented in the experiment control scheme, by shifting reference frames as is done in NMR [13]. For read-out, the qubit encoded in the motional state of the ion normally cannot be measured directly, but conditional pulse sequences allow full state tomography of the qubit system.

The control techniques developed here may be applicable to using a single ion to probe and manipulate other systems, even though they focus on a single ion and do not necessarily imply scalability. Some such systems include coupling ions to superconducting qubits [14], micromechanical cantilevers [15], cavities [16], and wires [17]. In many of these experiments, maximizing the coupling requires proximity of the ion to a surface, and coherence of the motional state is also desired.

This paper is organized as follows. The experimental setup, including the magnetic field stabilization scheme, is described in Section II. Section III briefly discusses motional state decoherence and shows that such decoherence has an insignificant effect on the gate performance in our system. Section IV presents a theoretical model of the Stark shift correction and experimental implementation of the method. Section V describes the state preparation and measurement sequences that allowed us to implement quantum process tomography on the single-ion system. Section VI describes the realization of the CNOT gate, along with a discussion of gate performance and error sources.

## II. EXPERIMENTAL SETUP

### A. Cryogenic microfabricated trap

The microfabricated trap is a five-rod surface-electrode design identical in geometry to that described in Ref. [10]. The trap is made of niobium, and the fabrication process is similar to prior methods [10] employed for gold

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traps, and is described briefly here. A 440 nm Nb layer is grown on a sapphire substrate by sputtering. The sheet resistance is 0.3 Ohms/sq at 295 K, and the superconducting transition is at  $T_c = 9.15$  K. Trap electrodes are patterned using NR9-3000 photoresist and etched using reactive ion etching with  $\text{CF}_4 + \text{O}_2$ . The trap center is 100  $\mu\text{m}$  above the surface. For the experiment described here, the axial and two radial trap frequencies are  $2\pi \times \{1.32, 2.4, 2.7\}$  MHz respectively. Though a superconducting trap was used for the work described here, the effects of the superconducting material on trapping behavior will be described elsewhere [18].

The trap is cooled and operated in a 4 K bath cryostat described in Ref.[19]. Typical ion lifetime is on the order of several hours, limited only by the liquid helium hold time. Loading is done via photoionization of a thermal vapor.

### B. $\text{Sr}^+$ qubit and laser system

An atomic ion confined in a harmonic trapping potential can encode two qubits: one in its optical atomic transition and one in its lowest motional states. The  $^{88}\text{Sr}^+$  ion has a narrow optical transition,  $S_{1/2} \leftrightarrow D_{5/2}$  with a linewidth of  $\sim 1$  Hz. The  $m = -1/2 \leftrightarrow m = -5/2$  levels are used for the atomic qubit transition. This transition is chosen for convenience, as sideband cooling can be performed without addressing additional optical transitions. The degeneracy of the multiple Zeeman levels is lifted by applying a constant field of 4 G with external coils. To address this transition at 674 nm, a diode laser is grating-stabilized and locked to an external cavity via optical feedback [20]. It is further stabilized by locking to a high-finesse ULE cavity similar to Ref.[21]. The frequency noise, indicated by the Pound-Drever-Hall error signal as measured with a spectrum analyser, is 0.3 Hz for noise components above 1 kHz. Below 1 kHz, acoustic noise broadens the laser linewidth to 300 Hz. This laser beam propagates along the axial direction of the trap, so we ignore the radial modes of motion in sideband cooling and quantum operations. Doppler cooling is performed on the  $S_{1/2} \leftrightarrow P_{1/2}$  transition with a 422 nm diode laser. Two IR diode lasers, at 1092 nm and 1033 nm, repump the ion from the  $D_{5/2}$  and  $D_{3/2}$  states. For all measurements, the ion is initialized to the  $S_{1/2}(m = -1/2)$  state and the motional ground state via a sequence of Doppler cooling, sideband cooling, and optical pumping. Fig 1 shows the relevant levels of  $\text{Sr}^+$  for the experiment.

A pulse sequencer [22] consisting of a field-programmable gate array (FPGA, OpalKelly XEM3010-1000) and direct digital synthesis boards controls the phases, amplitudes, and lengths of laser pulses. Switching of the beam and setting of the desired frequency and phase shift are accomplished using AOMs on the 674, 422, and 1033 nm lasers. Phase coherent switching is implemented by computing the expected phase at time  $t$ , referenced to a fixed point in the past, for a given fre-

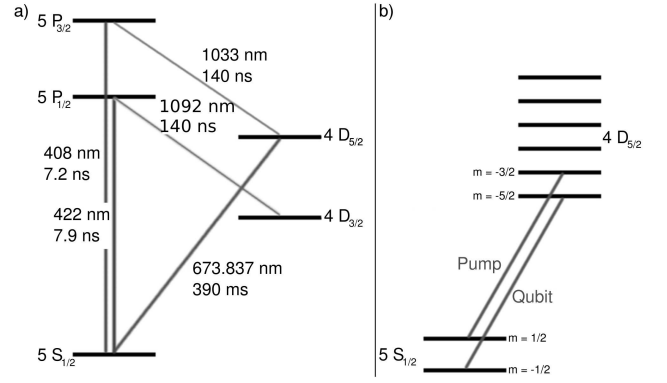


FIG. 1: (a)  $^{88}\text{Sr}^+$  level diagram. The 422 nm and 1091 nm transitions are used for Doppler cooling and detection. The 673.837 nm transition couples the qubit levels. (b) Details of the qubit states with Zeeman levels explicitly drawn. The “pump” transition is used to pump the ion out of the  $S_{1/2}(m = 1/2)$  state during initialization.

quency  $f$  using  $\phi_0(t) = f \times t \pmod{2\pi}$ . Then, after a frequency switch at time  $T$ , the absolute phase of the waveform is adjusted to equal  $\phi_0(T) + \phi$ , where  $\phi$  is any desired phase. This process allows for frequency switching while maintaining phase information throughout any arbitrary pulse sequence.

### C. Magnetic field stabilization

When the optical qubit is encoded in a pair of levels which are first-order sensitive to magnetic fields, field fluctuations on the timescale of gate operations will decrease gate fidelity. One way of passively stabilizing the field is using a mu-metal shield, which is expensive and inconvenient for optical access, and also mainly effective for low-frequency noise. Active stabilization of magnetic field using a flux gate sensor and coils have been implemented in another experiment [23], at the cost of higher complexity.

Superconducting solenoids have been employed for passively stabilizing ambient magnetic field fluctuations in NMR experiments, with field suppression by a factor of 156 [11]. A similar method for ion traps which would permit good optical access is desired. In the NMR implementation, the field needs to be stabilized over a region 1 cm in length, whereas in an ion trap the region of interest is much smaller. Our method uses the same principle of superconductive shielding, but the small region and requirement for optical access suggest a more compact approach.

We stabilize the magnetic field by employing the persistent current in two superconducting rings, placed closely adjacent to the ion trap chip. This is a very compact and experimentally convenient arrangement, with high passive field stability and little barrier to optical access. Below the trap is a 1 cm  $\times$  1 cm square Nb plate

with a 1.5 mm diameter hole, located 0.5 mm below the trap center. Above the trap is a 50 cm square plate with a 11 mm diameter hole, located 7 mm above the trap center (Fig.2a). Both rings are 0.5 mm thick. This geometry was chosen to optimize the field suppression at the trap location using the method to calculate magnetic fields in superconducting rings described in Ref.[24].

With a single trapped ion we measured the field suppression by applying a constant field with external coils, cooling the trap and Nb rings to below  $T_c$ , and reducing the field while measuring the  $S \leftrightarrow D$  transition frequency. The magnetic field is calculated from the Zeeman splitting between the  $m = -1/2 \leftrightarrow m = -5/2$  transition and the  $m = +1/2 \leftrightarrow m = -3/2$  transition. A 50-fold reduction in field sensitivity was observed (Fig.2), in agreement with the numerical calculation. To determine the effectiveness of the noise suppression on coherence of the atomic qubit, we measured the decay of Ramsey fringes as a function of the separation of the Ramsey  $\pi/2$  rotations on the carrier transition. Such a measurement also includes effects caused by laser linewidth and the drift in laser-ion distance. We found that reducing the magnetic field noise by a factor of 50 did not improve the coherence time by more than a factor of 2, suggesting that magnetic field noise is no longer a dominant source of decoherence.

Greater reduction can be obtained by optimizing the geometry further, e.g. by decreasing the distance between the plates to 4mm, but is not implemented due to physical constraints in the apparatus. This method only stabilizes the magnetic field along the axis of the superconducting rings, but since the 4 G bias field defining the quantization axis is applied in the same direction, field noise in the x- or y- direction only contributes quadratically to the change in the total field [11].

### III. MOTIONAL STATE COHERENCE

The Cirac-Zoller CNOT gate employs superpositions of ion motional states as intermediate states during the gate, and thus is sensitive to motional decoherence. In particular, a high ion heating rate will reduce the gate fidelity. An upper bound on the maximum heating rate tolerable  $\dot{n}_{\max}$  can be given by considering the total time  $T_{\text{gate}}$  required for the pulse sequence implementing the CNOT gate, together with a design goal for the gate error probability  $p_{\text{gate}}$  desired. Assuming that a single quantum of change due to heating will cause a gate error, then  $\dot{n}_{\max} < T_{\text{gate}}/p_{\text{gate}}$ . For  $T_{\text{gate}} \sim 230 \mu\text{s}$  (for our experiment), a heating rate of  $\dot{n}_{\max} < 40$  quanta/s is needed to get  $p_{\text{gate}} \sim 0.01$ .

We measured the heating rate of the trap at the operating secular frequency of  $2\pi \times 1.32$  MHz. The number of motional quanta is measured by probing the blue and red sidebands of the  $S \leftrightarrow D$  transition using the shelving technique, and comparing the ratio of shelving probability on each sideband [25]. The heating rate is determined

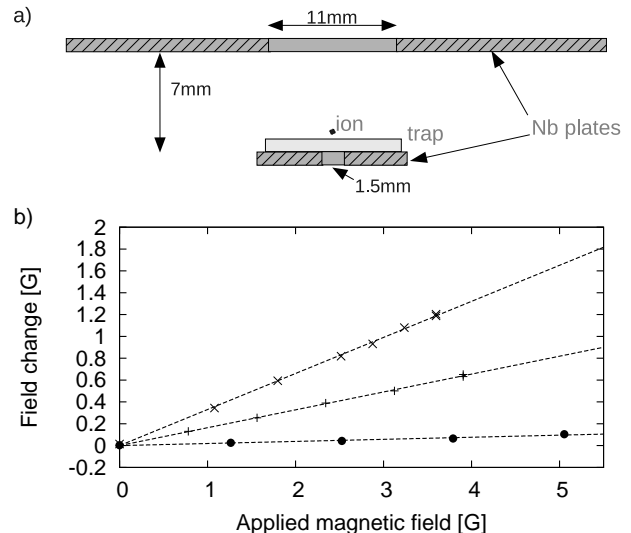


FIG. 2: (a) Two superconducting discs, one below and one above the trapped ion, stabilize the magnetic field in the  $\hat{z}$  direction. (Not to scale.) (b) Magnetic field fluctuation suppression due to the top disc only ( $\times$ ), the bottom disc only ( $+$ ) and both discs ( $\bullet$ ). When both discs are used, field changes are suppressed 50-fold.

by varying the delay before readout and comparing the number of quanta vs delay time. The lowest heating rate measured in this trap was  $2.1(3)$  q/s.

Fig.3a shows Rabi flops on the blue sideband after the ion is initialized to the motional ground state with average number of quanta  $\bar{n} < 0.01$ . The fitted initial contrast is  $97.6(3)\%$  and the frequency is  $46.7$  kHz. Motional state coherence is demonstrated by performing Ramsey spectroscopy on the blue sideband (Fig.3b). The coherence time  $T_2^*$  is  $622(37) \mu\text{s}$ . This is comparable to the coherence time of  $660(12) \mu\text{s}$  of the atomic qubit as measured by the same method on the carrier transition.

### IV. STARK SHIFT CORRECTION

When a two-level atom encoding a qubit is driven off-resonance, such as on a sideband transition, it excites the carrier transition and creates an AC-Stark shift. In a real ion with multiple levels, additional complication comes from other transitions that contribute shifts which are independent of the laser detuning. In the past, correction for the Stark shift has been done by using an additional laser detuned to the opposite sideband transition to cancel the shift [12]. In qubits addressed by a Raman transition, this can also be accomplished by changing the power ratio of the Raman pulses [26]. In this work, the Stark shift correction is done by calculating the shift and accounting for it in the pulse sequencer, following an example in NMR [13]. Here, we develop a systematic model of the light shifts seen by a single trapped ion.

The Stark shift is traditionally a phase shift caused

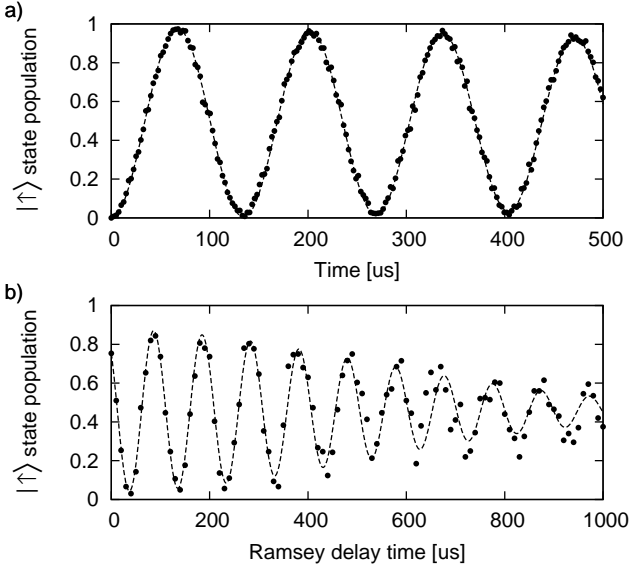


FIG. 3: (a) Rabi oscillations on the blue sideband. The fitted initial contrast is 97.6(3)% and the frequency is 46.7 kHz. (b) Ramsey spectroscopy on the blue sideband. The fitted Gaussian envelope of the decay has time constant  $T_2^* = 622(37) \mu\text{s}$ .

by a small change in the transition frequency due to level shifts. In reality it is a unitary transform involving more than just a change of energy levels. Below, we also take this into consideration as a “generalized” Stark shift. The model presented here is adapted from well-known methods in NMR and included for pedagogical reasons. Section IV A outlines a general method for switching between different reference frames, and Section IV B identifies the reference frames useful for discussing the single ion in the context of quantum control. The generalized Stark shift correction operation is then derived as a result of switching between these frames. In Section IV C we apply this method to our single ion system and describe how to calculate the appropriate Stark shift correction for any gate in an arbitrary gate sequence. Section IV D describes the measurement of the AC Stark shift and results of the Stark shift correction.

### A. Frames of reference

The Hamiltonian for the system in the laboratory reference frame is often inconvenient to use, both for calculations and for modeling the system as it is used for a quantum computation. Specifically, the simplest way to exponentiate a Hamiltonian and obtain the time evolution operator is in a frame of reference where the Hamiltonian is time-independent. In contrast, in the laboratory frame, the electric field of the laser varies with time. Also, because a two-level system such as an atom evolves with time in the lab frame, it is convenient to define a frame of reference which follows this evolution, so that a stationary definition of a qubit can be obtained. This

is accomplished experimentally by keeping a clock which accurately follows the evolution of the qubit in the absence of any excitation.

Mathematically, such frames of reference are constructed and used in the following method. Let  $H_0$  be the Hamiltonian of the full system in the laboratory reference frame. Let  $|\psi\rangle$  be a state in this rest frame. Suppose we want to follow the system in a different frame of reference, defined by another Hamiltonian,  $H_{\text{frame}}$ . Often, this Hamiltonian will be constructed from terms in the system Hamiltonian. We may define a state vector in this new frame in terms of the state vector in the rest frame, as (letting  $\hbar = 1$ ):

$$|\phi(t)\rangle = e^{+iH_{\text{frame}}t}|\psi(t)\rangle. \quad (1)$$

In terms of this new state vector, the Schrodinger equation for the system is

$$i\partial_t|\phi(t)\rangle = [e^{+iH_{\text{frame}}t}H_0e^{-iH_{\text{frame}}t} - H_{\text{frame}}]|\phi(t)\rangle. \quad (2)$$

This is the Schrödinger equation for a system evolving under a different Hamiltonian,

$$V = e^{+iH_{\text{frame}}t}H_0e^{-iH_{\text{frame}}t} - H_{\text{frame}} \quad (3)$$

such that for time  $t = t_2 - t_1$ , the state (in the rotating frame) is given by

$$|\phi(t_2)\rangle = e^{-iVt}|\phi(t_1)\rangle. \quad (4)$$

Thus, in the laboratory frame, the state is

$$|\psi(t_2)\rangle = e^{-iH_{\text{frame}}t_1}e^{-iH_{\text{frame}}t}e^{-iVt}e^{iH_{\text{frame}}t_1}|\psi(t_1)\rangle. \quad (5)$$

Note that when all the Hamiltonians commute, this expression simplifies to give the normal expected evolution. This equation shows that converting between frames involves not just conjugation by  $e^{-iH_{\text{frame}}t_1}$ , but also computation of  $e^{-iH_{\text{frame}}t}e^{-iVt}$ .

### B. Stark shift on carrier: simple free ion model

There are several useful frames of reference to describe the two-level atom model. Consider a single ion at a fixed position in free space, interacting with a single mode laser. Let this be described by the laboratory reference frame Hamiltonian (with rotating wave approximation)

$$H_0 = \omega_0\sigma_z + \Omega\sigma_x \cos\omega t + \Omega\sigma_y \sin\omega t \quad (6)$$

where  $\omega_0$  is the optical transition frequency,  $\sigma_x, \sigma_y, \sigma_z$  are spin-1/2 operators corresponding to the Pauli matrices with eigenvalues  $\pm 1/2$ , and  $\Omega$  is the Rabi frequency.

Let the laser be applied at frequency  $\omega = \omega_0 + \delta$ , such that we may define

$$H_L = \omega\sigma_z \quad (7)$$

as a convenient frame of reference. In the frame of the laser, the Hamiltonian is

$$V_L = -\delta\sigma_z + \Omega\sigma_x. \quad (8)$$

The frame of reference we wish to use for quantum computation (QC frame) is defined by the Hamiltonian

$$H_{QC} = \omega_0\sigma_z. \quad (9)$$

Thus, if we define a state in this frame as

$$|\gamma(t)\rangle = e^{+iH_{QC}t}|\psi(t)\rangle \quad (10)$$

where  $|\psi(t)\rangle$  is the state in the lab frame, then we find that

$$|\gamma(t)\rangle = e^{i\delta\sigma_z t} e^{-i(\delta\sigma_z + \Omega\sigma_x)t} |\gamma(0)\rangle \quad (11)$$

assuming that  $|\gamma(0)\rangle = |\psi(0)\rangle$ .

The generalized Stark shift correction operation which needs to be applied is thus  $R^\dagger$ , where

$$R = e^{i\delta\sigma_z t} e^{-i(\delta\sigma_z + \Omega\sigma_x)t}. \quad (12)$$

This is an operator that rotates about an axis

$$\hat{n} = \frac{\hat{z} + \frac{\Omega}{\delta} \hat{x}}{\sqrt{1 + \left(\frac{\Omega}{\delta}\right)^2}}. \quad (13)$$

When the detuning is very large compared with the Rabi frequency, the maximum rotation about the  $\hat{x}$  axis, which corresponds to a population change, can be bounded by  $\Omega^2/\delta^2$  for a  $\pi$ -rotation about  $\hat{n}$ . For our experimental parameters (Sec VIA), this is less than 1%. Therefore the Stark shift is traditionally approximated as a rotation about the  $\hat{z}$  axis,  $R_z(\theta) = e^{i\theta\sigma_z}$ . We can compute what this operation and the rotation angle  $\theta$  would be by looking for the  $R_z$  closest to  $R$ . The angle of rotation of the operator  $e^{-i(\delta\sigma_z + \Omega\sigma_x)t}$  is  $\sqrt{\delta^2 + \Omega^2}t$ , while the angle of rotation of the operator  $e^{i\delta\sigma_z t}$  is  $\delta t$ . Thus, if one ignored the axes of rotation and treated the first operator as if it were also a rotation about  $\hat{z}$ , then the Stark shift correction would be a rotation by angle

$$\left[\delta - \sqrt{\delta^2 + \Omega^2}\right] t \quad (14)$$

about  $\hat{z}$ .

### C. Stark shift corrections for arbitrary gate sequences

We now examine a real experimental situation with a multi-level ion. To verify our proposed Stark shift correction and later to consider the effect of error sources on gate fidelity, we simulated gate operations by modelling the action of lasers on the full system Hamiltonian in the space formed by  $\{|D\rangle, |S\rangle\} \otimes \{|0\rangle, |1\rangle, |2\rangle\}$ .

Exact simulation of the action of the lasers on the computational space requires the use of both laser and QC frames. The full Hamiltonian is time-independent in the laser frame, suggesting that gate operations should be computed in that frame. The states used in quantum computation are defined in the QC frame, where they are stationary without the interaction applied. Simulation of a gate sequence will therefore require frequent switching between the frames, for which we define the operator  $U_{\text{lqc}}(t)$ .

Computation is performed by moving to the laser frame, exponentiating  $V_L$ , and moving back to QC frame. For example, a gate performed by applying a laser pulse of detuning  $\delta$ , phase  $\phi$ , starting at time  $t_0$  for time  $t$ , can be computed in QC frame to be

$$U(\delta, \phi, t, t_0) = U_{\text{lqc}}(t + t_0) e^{-iV_L t/\hbar} U_{\text{lqc}}(t_0)^\dagger \quad (15)$$

where  $V_L$  and  $U_{\text{lqc}}$  depend on laser detuning, phase and Rabi frequency as well as the trap parameters. Note that each laser detuning and trap frequency defines a separate laser frame. The operator  $U_{\text{lqc}}$  moves between the unique QC frame and one of the infinite number of laser frames.

Let  $U_\phi(\phi) = e^{-i\phi}$  be a phase shift on the D states:

$$U_\phi(\phi) = \begin{pmatrix} e^{i\phi} & 0 \\ 0 & 1 \end{pmatrix} \otimes I_3, \quad (16)$$

where  $I_n$  is the identity matrix of size  $n \times n$ . Here the  $2 \times 2$  matrix acts on  $\{|D\rangle, |S\rangle\}$  and  $I_3$  acts on the motional states. Experimentally this operator is equivalent to shifting the laser phase by  $\phi$ . In a sequence of gates, applying such a phase rotation implies shifting the laser phases of all subsequent gates.

In a multi-level atom, there are other transitions which are off-resonantly coupled to the laser and contribute to additional phase shifts that are detuning-independent. In our modeling of the  $\text{Sr}^+$  computation presented here, we include the  $S_{1/2} \leftrightarrow P_{1/2}$ ,  $S_{1/2} \leftrightarrow P_{3/2}$ , and  $D_{1/2} \leftrightarrow P_{3/2}$  transitions. The matrix elements of all these transitions which determine the resulting shift can be calculated as in Ref.[27].

For gates performed on the carrier transition, since the duration of carrier gates are shorter than sideband gates by the Lamb-Dicke factor  $\eta$  ( $= 0.06$  for our case), carrier gates take only a small fraction of the total time in a typical gate sequence ( $\sim 2\%$  in the CNOT pulse sequence). Thus we ignore off-resonant coupling to the motional sidebands and coupling to far-off-resonant transitions. For gates on the sideband transitions, consider an interaction with laser detuning  $\delta \gg \Omega$ , carrier Rabi frequency  $\Omega$  and phase  $\phi$  applied for time  $t$  starting at time  $t_0$ . There are three separate phase shifts which need to be cancelled:

- Stark shift of the ground and excited state can be removed by rotating the phase of the  $|e\rangle$  state by  $\phi_s = -(\delta - \sqrt{\delta^2 + \Omega^2})t$ , equivalent to applying  $e^{-i\sigma_z \phi_s}$ , following the offending gate. Z rotations

can be performed by changing the phases of all subsequent laser pulses by  $\phi_s$ .

- Resonant excitation of sidebands is applied at a frequency Stark-shifted due to the carrier. The laser frame corresponding to that frequency will rotate with respect to the unshifted states at a rate proportional to the Stark shift. To bring the laser frame and unshifted states in phase, the laser phase has to be shifted by  $\phi_f = (\delta - \sqrt{\delta^2 + \Omega^2}) t_0$ . Such phase shift is equivalent to applying  $e^{-i\sigma_z \phi_f}$  before the gate, and  $e^{i\sigma_z \phi_f}$  after.
- Off-resonant phase shifts account for approximately 10% of the total Stark shift in  $\text{Sr}^+$ . Let  $\Delta_o$  be a constant factor to account for these off-resonant phase shifts.

Define the carrier gate  $U_c$ , sideband gate  $U_m$ , and phase correction  $\Delta$  as follows. Along with the gate time  $t$  and gate starting time  $t_0$ , these variables contain all the information relevant to calculating the required Stark shift correction.

$$U_c = U \quad (17)$$

$$U_m = U_\phi(-\Delta(t + t_0)) U U_\phi(\Delta t_0) \quad (18)$$

$$\Delta = \left[ \delta - \sqrt{\delta^2 + \Omega^2} \right] + \Delta_o \quad (19)$$

From the definition of  $U$  and properties of the exponential function, it can be shown that the phase correction on the  $n$ -th gate  $U^n$  in an arbitrary gate sequence is

$$U^n = U_\phi \left( -\sum_{i=1}^n \Delta^i t^i \right) U \left( \delta, \phi + \Delta^n t_0^n - \sum_{i=1}^{n-1} \Delta^i t^i, t^n, t_0^n \right). \quad (20)$$

This phase correction consists of the appropriate correction for that particular gate plus a global phase, the sum of all phase corrections applied to previous gates. In our pulse sequencer [22], the global time and global phase are kept as internal registers, and are used to calculate the appropriate phase correction every time the qubit laser phase is set during a pulse sequence.

#### D. Results

Ramsey spectroscopy on the blue sideband can be used to characterize the effectiveness of the Stark shift correction. Using the methods described in Ref.[12], we measure the AC Stark shift for various detunings and compare to the theoretical model. Fig.4b shows the typical oscillation in the shelving probability  $P(D)$  when a pulse detuned from the  $S \leftrightarrow D$  transition (Stark pulse) of varying duration is applied. The AC Stark shift is given by the oscillation frequency. This shift is measured for several values of detuning, and is shown in Fig.4c along with a one-parameter fit to  $A/x + b$  where the fixed parameter is  $A = \Omega^2/(2\omega_{\text{sec}})$ , the detuning-dependent shift

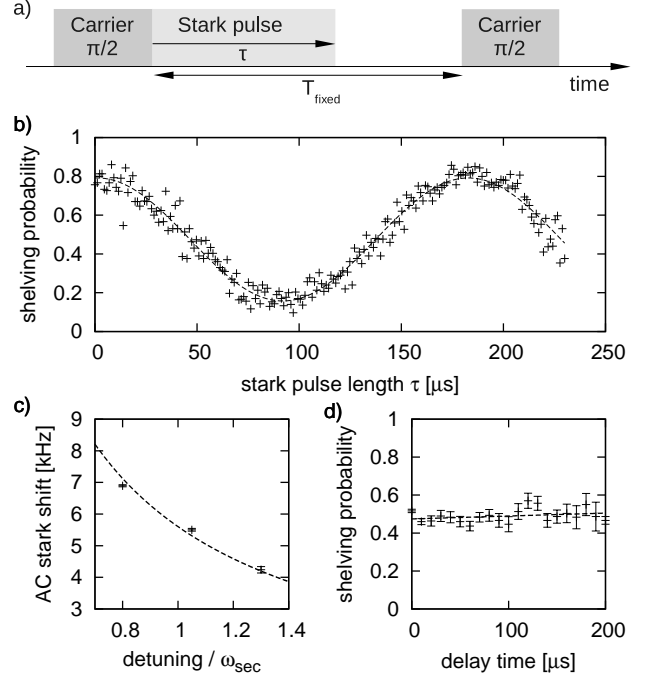


FIG. 4: (a) Pulse sequence used to measure AC Stark shift, from Ref.[12]. The delay between two carrier  $\pi/2$  pulses is  $T_{\text{fixed}} = 230 \mu\text{s}$  while the length  $\tau$  of the stark pulse is varied. (b) Typical measurement with the pulse sequence in (a). The AC Stark shift is given by the oscillation frequency of the shelving probability. Here, the fixed laser detuning is  $1.05\omega_{\text{sec}}$  which gives an AC stark shift of  $2\pi \times 5.50(4) \text{ kHz}$ . (c) Measured AC Stark shift as a function of detuning, fitted to the Stark shift model with the detuning-independent shift as a free parameter. (d) Ramsey spectroscopy on the sideband demonstrating compensation of Stark shift.

to first order, and  $b$  is the detuning-independent Stark shift caused by farther off-resonant transitions. The fitted offset is  $b = -2\pi \times 0.5(1) \text{ kHz}$ , in agreement with  $-2\pi \times 0.50 \text{ kHz}$  predicted by theory.

The effectiveness of AC Stark shift compensation were evaluated by performing Ramsey spectroscopy on the sideband and varying the delay time between the two pulses, with both Ramsey pulses shifted by  $\pi/2$ . In the absence of uncorrected Stark shifts, the expected  $P(D)$  is  $1/2$  for all delay times. Fig.4d shows the result of such a measurement. Here, the secular frequency is determined by taking a spectrum and fitting to the sideband, then the Ramsey sequence is performed. The parameter  $\phi_s$  is fixed in the experiment control hardware while  $\Delta_o$  is tuned such that  $P(D)$  is maintained near  $1/2$ . From the slope of Fig.4d we estimate the residual Stark shift to be  $2\pi \times 24(20) \text{ Hz}$ .

## V. QUANTUM PROCESS TOMOGRAPHY ON A SINGLE ION

With two qubits encoded in a single ion and methods of coupling and controlling these states, a Cirac-Zoller CNOT gate can be implemented [28]. The CNOT gate is universal in that all quantum operations can be decomposed into single-qubit operations and the CNOT gate, and is thus of interest for implementing quantum information processing in ion traps. To evaluate the performance of such a gate, we prepare the system in a set of basis states that spans the Hilbert space and perform a set of measurements that completely specifies the resulting state (state tomography). Quantum process tomography (QPT) is performed on the two qubits to construct the process matrix, allowing a full characterization of the gate. Section V A gives a brief summary of state tomography using conditional measurements. Section V B describes a minimal set of available measurements and operations in this two-qubit single-ion system necessary for QPT. Section V C and V D lists the pulse sequences for preparing all basis states and measuring the outcome. Section V E briefly describes the construction of the process matrix which fully characterizes the gate from these measurements.

### A. Two qubit state tomography for one ion

State tomography on the single-ion system of atomic and motional qubits requires a non-trivial set of operations, since a single qubit rotation on the motional qubit cannot be realized directly except by first swapping it with the internal state, performing the desired gate, then swapping back. The swap operation is complicated since the most straightforward physical operations, red and blue sideband pulses, generally take the system out of the computational space, and into higher order motional states such as  $|2\rangle$  [29]. For the CNOT gate, a set of composite pulse sequences can keep the system in the computational space. But if the goal is measurement of the two-qubit state space rather than the realization of a coherent operation, an alternative approach can be employed. A sequence of measurements, with the second conditioned on the results of the first, can suffice to allow full state tomography on the two-qubit atomic+motional state space. This is an extension of the single-ion tomography technique described in [30].

The conditional measurement sequence is as follows. First we apply an optional  $\pi$  pulse on the carrier transition; then the internal atomic state is measured by fluorescence detection. When this measurement scatters photons, it provides information about the internal state only and the motional state information is lost. When this first measurement does not scatter photons, a  $\pi$  pulse is applied on the blue sideband transition, which allows measurement of the population in the state pairs  $\{|S0\rangle, |S1\rangle\}$  or  $\{|D0\rangle, |D1\rangle\}$ , depending on whether the

initial carrier  $\pi$  pulse was applied or not. Two measurements, with and without the carrier pulse, are sufficient to determine the population in all 4 states.

### B. Process tomography: operator definitions

The state tomographic measurements just described measure state population only, which are the diagonal elements of the full density matrix. Relative phases between qubit states, which determine coherence properties of the state, are also needed in order to perform complete process tomography. The phases can be obtained by appropriate rotations of the qubits prior to measurement. Here we define the measurement and rotation operators for the sections following.

The single available measurement is the usual fluorescence detection, which is a projective measurement into the  $|S\rangle$  state, denoted  $P_S$ . Let  $P_D$  denote a projection into the  $|D\rangle$  state. The matrices for  $P_S$ ,  $P_D$  in the basis  $|D0\rangle, |D1\rangle, |D2\rangle, |S0\rangle, |S1\rangle, |S2\rangle$  are:

$$P_S = \begin{pmatrix} 0 & 0 \\ 0 & 1 \end{pmatrix} \otimes I_3, \quad P_D = I_6 - P_S \quad (21)$$

The available unitary operations are:

- $R_x(\theta)$ ,  $R_y(\theta)$ : Single qubit (carrier) rotations on the  $\{|S\rangle, |D\rangle\}$  qubit.
- $R_x^+(\theta)$ ,  $R_y^+(\theta)$ : Blue sideband rotations, connecting  $\{|S0\rangle, |D1\rangle\}$  and  $\{|S1\rangle, |D2\rangle\}$  (neglecting higher order vibrational modes).  $\theta$  is the rotation angle on the  $\{|S0\rangle, |D1\rangle\}$  manifold.
- Red sideband rotations can be defined similarly, but are actually not necessary for constructing a complete measurement set.

Explicitly, these rotation matrices are defined as follows:

$$\begin{aligned} R_x(\theta) &= \exp(-i\theta(\sigma_x \otimes I_3)) \\ R_y(\theta) &= \exp(-i\theta(\sigma_y \otimes I_3)) \\ R_x^+(\theta) &= \exp(\theta(\sigma_+ \otimes a^\dagger - \sigma_- \otimes a)/2) \\ R_y^+(\theta) &= \exp(-i\theta(\sigma_+ \otimes a^\dagger + \sigma_- \otimes a)/2) \end{aligned} \quad (22)$$

where  $a^\dagger$ ,  $a$  are the creation and annihilation operators in the Jaynes-Cummings Hamiltonian.

### C. State preparation

For every measurement sequence, the ion is initialized to the state  $\Psi_0 \equiv |S0\rangle$ . The sequences of operations listed in Table I generates 16 basis states which span the complete Hilbert space of  $|D0\rangle, |D1\rangle, |S0\rangle, |S1\rangle$ .

$\Psi(i)$	Operations applied to $\Psi_0$	State
$\Psi(1)$	$R_y(-\pi)$	$ D0\rangle$
$\Psi(2)$	$R_x^+(-\pi)$	$ D1\rangle$
$\Psi(3)$	$I$	$ S0\rangle$
$\Psi(4)$	$R_y(\pi)R_x^+(\pi)$	$ S1\rangle$
$\Psi(5)$	$R_x^+(\pi)R_y(-\pi/2)$	$( D0\rangle +  D1\rangle)/\sqrt{2}$
$\Psi(6)$	$R_y^+(-\pi)R_y(-\pi/2)$	$( D0\rangle + i D1\rangle)/\sqrt{2}$
$\Psi(7)$	$R_y(-\pi/2)$	$( D0\rangle +  S0\rangle)/\sqrt{2}$
$\Psi(8)$	$R_x(\pi/2)$	$( D0\rangle + i S0\rangle)/\sqrt{2}$
$\Psi(9)$	$R_y(-\pi)R_x^+(-\pi/2)$	$( D0\rangle +  S1\rangle)/\sqrt{2}$
$\Psi(10)$	$R_y(-\pi)R_x^+(\pi/2)$	$( D0\rangle + i S1\rangle)/\sqrt{2}$
$\Psi(11)$	$R_x^+(\pi/2)$	$( D1\rangle +  S0\rangle)/\sqrt{2}$
$\Psi(12)$	$R_y^+(\pi/2)$	$( D1\rangle + i S0\rangle)/\sqrt{2}$
$\Psi(13)$	$R_y(\pi/2)R_x^+(\pi)$	$( D1\rangle +  S1\rangle)/\sqrt{2}$
$\Psi(14)$	$R_x(-\pi/2)R_x^+(-\pi)$	$( D1\rangle + i S1\rangle)/\sqrt{2}$
$\Psi(15)$	$R_y(-\pi)R_x^+(-\pi)R_y(\pi/2)$	$( S0\rangle +  S1\rangle)/\sqrt{2}$
$\Psi(16)$	$R_y(-\pi)R_y^+(\pi)R_y(\pi/2)$	$( S0\rangle + i S1\rangle)/\sqrt{2}$

TABLE I: State preparation operations.

#### D. Complete basis of measurements

The following is a procedure for performing complete state tomography of the two-qubit  $\{|S\rangle, |D\rangle\} \otimes \{|0\rangle, |1\rangle\}$  Hilbert space of a single ion, using the measurements and operations in Sec V B. This is a generalization of the method used to measure just the diagonal elements of the density matrix. There are two kinds of measurements used; we call them MeasU and MeasUV.

**MeasU** involves performing a unitary operation  $U$  to the input state, then projecting into the  $|S\rangle$  subspace,  $P_S$ . This is described by the measurement operator

$$\text{MeasU}(U) = U^\dagger P_S U \quad (23)$$

Typically,  $U$  will be a rotation in the  $\{|S\rangle, |D\rangle\}$  subspace, implemented by a carrier transition pulse.

**MeasUV** involves first performing a unitary operation  $U$  to the input state and making a measurement to detect fluorescence, which is equivalent to projecting to the  $|S\rangle$  subspace. Since  $|D\rangle$  is long-lived, this projection leaves the  $\{|D0\rangle, |D1\rangle, \dots\}$  subspace undisturbed, but motional state information is lost if the ion is in state  $|S, n\rangle$ . If no fluorescence is detected, the post-measurement state is  $P_D \rho P_D$ . Conditioned on the first measurement returning  $|D\rangle$  (no fluorescence), a unitary transform  $V$  is performed, and finally another into the  $|S\rangle$  subspace,  $P_S$ . If the first measurement returns fluorescence, the measurement sequence stops, in which case only information about the atomic state is obtained. MeasUV is described by the measurement operator

$$\text{MeasUV}(U, V) = U^\dagger P_D V^\dagger P_S V P_D U \quad (24)$$

Typically,  $U$  will be a rotation in the  $\{|S\rangle, |D\rangle\}$  subspace, while  $V$  will be one or more rotations on the carrier and the red or blue sideband.

The measurements listed in Table II provide a complete basis of observables from which the full density matrix  $\rho$  can be reconstructed, assuming that  $\rho$  is initially in only the two-qubit computational subspace. These measurement observables are linearly independent.

$M_j$	Measurement functions
$M_1$	$\text{MeasU}(I)$
$M_2$	$\text{MeasUV}(I, R_y^+(\pi))$
$M_3$	$\text{MeasUV}(R_y(\pi), R_y^+(\pi))$
$M_4$	$\text{MeasU}(R_y(\pi/2))$
$M_5$	$\text{MeasU}(R_x(\pi/2))$
$M_6$	$\text{MeasUV}(I, R_y(\pi/2)R_y^+(\pi/2))$
$M_7$	$\text{MeasUV}(R_y(\pi), R_y(\pi/2)R_y^+(\pi/2))$
$M_8$	$\text{MeasUV}(R_y(\pi/2), R_y(\pi/2)R_y^+(\pi/2))$
$M_9$	$\text{MeasUV}(R_y(\pi/2), R_x(\pi/2)R_y^+(\pi/2))$
$M_{10}$	$\text{MeasUV}(I, R_x(\pi/2)R_y^+(\pi/2))$
$M_{11}$	$\text{MeasUV}(R_x(\pi), R_x(\pi/2)R_y^+(\pi/2))$
$M_{12}$	$\text{MeasUV}(R_x(\pi/2), R_x(\pi/2)R_y^+(\pi/2))$
$M_{13}$	$\text{MeasUV}(R_x(\pi/2), R_y(\pi/2)R_y^+(\pi/2))$
$M_{14}$	$\text{MeasUV}(R_y(\pi/2), R_y^+(\pi/2))$
$M_{15}$	$\text{MeasUV}(R_x(\pi/2), R_y^+(\pi/2))$

TABLE II: State measurement functions.

The relationship between measurements and the density matrix can be expressed by a matrix  $A$  with elements

$$A_{ij} = M_j(\Psi(i)). \quad (25)$$

The full density matrix  $\rho$  can be reconstructed as:

$$\rho = \sum_{ij} m_j A_{ij}^{-1} |\Psi_i\rangle \langle \Psi_i| \quad (26)$$

where  $m_j$  is the result of measurement  $M_j$ .

#### E. Constructing the process matrix

A quantum gate including all error sources can be represented by the operation  $\mathcal{E}(\rho)$ , which can be written in the operator sum representation as

$$\mathcal{E}(\rho) = \sum_{mn} E_m \rho E_n^\dagger \chi_{mn} \quad (27)$$

where  $\rho$  is the input state and  $E_i$  is a basis of the set of operators on the state space. The process matrix  $\chi_{mn}$  contains the full gate information. For two qubits, the state space is spanned by 16 basis states, and  $16^2$  elements define the  $\chi$ -matrix, though it only has  $16 \times 15$  independent degrees of freedom due to normalization. This is reflected by the fact that only 15 measurements are needed. The  $\chi$ -matrix can be obtained by inverting the above relation. To avoid unphysical results (namely a non-positive-semidefinite  $\rho$ ,  $\text{Tr}(\rho^2) > 1$ ) caused by statistical quantum



error in the experiment, a maximum-likelihood estimation algorithm [31] is employed to determine the physical operation  $\mathcal{E}$  which most likely generated the measured data. An alternate, iterative algorithm is presented in Ref.[32].

## VI. SINGLE-ION CNOT GATE

The CNOT gate is implemented with the pulse sequence described in Ref.[33]. The optical transition is the control qubit, and the motional ground and first excited states are used as the target qubit. In the product basis  $\{|D0\rangle, |D1\rangle, |S0\rangle, |S1\rangle\}$ , the unitary matrix implemented is

$$U = \begin{pmatrix} 1 & 0 & 0 & 0 \\ 0 & -1 & 0 & 0 \\ 0 & 0 & 0 & 1 \\ 0 & 0 & -1 & 0 \end{pmatrix} \quad (28)$$

$$= \frac{1}{2}(-iY \otimes Z + Z \otimes I + Z \otimes Z + iY \otimes Z).$$

This differs from the ideal CNOT matrix by only single-qubit phase shifts. Section VIA describes the achieved gate fidelities and Section VIB discusses the major known error sources that compromises gate fidelity.

### A. Gate performance

Quantum process tomography was carried out to evaluate the performance of various gates on the two qubits of a single ion. The ion in its motional and atomic ground state is initialized to one of the 16 basis states in Table I. Then the gate is applied, and the output state is determined by making all of the measurements listed in Table II. The longest duration of the full measurement sequence (excluding the gate) is 610  $\mu$ s, and a single CNOT gate takes 230  $\mu$ s. These durations are determined by the Rabi frequency on the carrier  $\Omega = 2\pi \times 125$  kHz, on the sideband  $\Omega_{\text{BSB}} = 2\pi \times 7.7$  kHz, and secular frequency  $\omega_{\text{sec}} = 2\pi \times 1.32$  MHz. The resulting  $\chi$ -matrix for the CNOT gate is shown in Fig.5.

We evaluate the performance of the identity gate (all preparation and measurement sequences performed with no gate in between), the single CNOT gate, and two concatenated CNOT gates (CNOTx2). The results are shown in Table III. The process fidelity is defined as  $F_p = \text{Tr}(\chi_{\text{id}}\chi_{\text{exp}})$ , where  $\chi_{\text{id}}$  is the ideal  $\chi$ -matrix calculated with the ideal unitary operation  $U$ , and  $\chi_{\text{exp}}$  is experimentally obtained using maximum-likelihood estimation. We also define the mean fidelity,  $F_{\text{mean}}$ , based on the overlap between the expected and measured density matrices,  $\text{Tr}(\rho_{\text{id}}\rho_{\text{exp}})$ , averaged over all prepared and measured basis states, similar to Ref.[34]. Error bars on  $F_p$  are calculated from quantum projection noise using Monte Carlo methods [35]. The large error bars on  $F_{\text{mean}}$

are due to certain measured basis states consistently having a higher or lower overlap with the ideal states. In general, states that involve multiple pulses to create entanglement are more susceptible to error and therefore have a lower fidelity than states that are closer to pure states. The pulse sequence for some states essentially performs a CNOT gate to create and remove entanglement; thus imperfect state preparation and measurement contributes significantly to the overall infidelity. Using the data for 0, 1, and 2 gates, we can estimate the fidelity of a single CNOT gate normalized with respect to the overall fidelity of the state preparation and measurement steps. Assuming that the fidelity of the  $n$ -th gate is  $F_p^n = F_i(F_g)^n$ , where  $F_i$  is the preparation and measurement fidelity, the fitted fidelity per gate  $F_g$  is 95%.

Gate	$F_p$ (%)	$F_{\text{mean}}$ (%)
ID	90(1)	94(3)
CNOT	85(1)	91(5)
CNOTx2	81(1)	89(6)

TABLE III: Measured gate fidelities for the identity gate, the single CNOT gate, and two concatenated CNOT gates.

### B. Error sources

A number of possible error sources and their contributions to the process fidelity of the single CNOT gate are listed in Table IV. To estimate and understand error sources, we simulated the full system evolution in the (2 atomic state)  $\times$  (3 motional state) manifold using the exact Hamiltonian, including Stark shift and tomographic measurements. The magnitude of each source is measured independently and then added to the simulated pulse sequence. Laser frequency fluctuation is assumed to be the primary cause of decoherence and is measured by observing the decay of Ramsey fringes on the carrier transition. The frequency fluctuation is simulated as a random variable on the laser frequency which grows in amplitude over time, and accounted for via Monte Carlo techniques. Laser intensity fluctuations are measured directly with a photodiode. On short time scales comparable to the length of the gate, the fluctuations are  $\sim 0.1\%$  peak-to-peak; on longer time scales, up to 1% drifts are observed. Both of these effects are accounted for in the simulation. Off-resonant excitations are automatically included in the model of the full Hamiltonian. The effect can be removed from the simulation if decoherence is not included and the simulated pulses are of arbitrarily long lengths, equivalent to reducing the laser intensity. The resulting  $\chi$ -matrix and fidelity agrees well with the measured results, indicating that the observed fidelity is well-understood in terms of technical limitations.

Off-resonant excitations, caused by the square pulse shape used to address all transitions, is expected to be the largest source of error, as previous work has found [36].

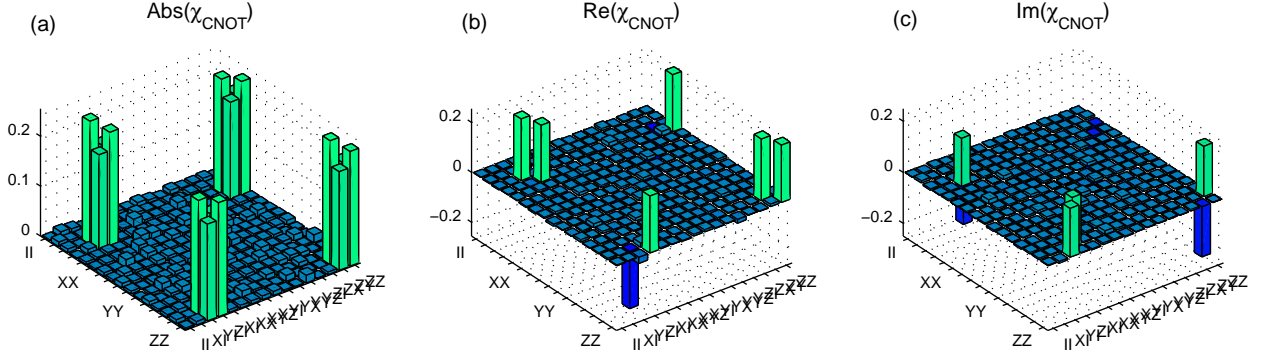


FIG. 5: Process tomography on the CNOT gate. (a), (b), (c) shows the absolute, real, and imaginary parts of the  $\chi$ -matrix respectively.

Error source	Magnitude	approx. contribution
off-resonant excitations	1%	10%
laser frequency fluctuations	300 Hz	5%
laser intensity fluctuations	1%	1%
Total		15%

TABLE IV: Error budget listing the major sources of errors on the process fidelity of the single CNOT gate, obtained by simulation. Each error source is assumed to be independent. The total error is calculated as the product of individual errors.

Square pulses on the blue sideband transition contain many higher harmonics, which causes residual excitation of the carrier transition. The carrier transition oscillations caused by this can be measured directly, averaged over many scans due to their small amplitude. Although the measured amount of off-resonant excitation is small ( $\sim 1\%$ ) for the laser intensity and secular frequencies used for our gates, both our simulations and previous work [36] have found that up to 10% improvement in gate fidelity can be gained by implementing amplitude pulse shaping.

## VII. CONCLUSION

In summary, we have developed a cryogenic, micro-fabricated ion trap system and demonstrated coherent

control of a single ion. The cryogenic environment suppresses anomalous heating of the motional state, as well as enables the use of a compact form of magnetic field stabilization using superconducting rings. We perform Stark shift correction in the pulse sequencer, removing the requirement for a separate laser path and acousto-optical modulator. A complete set of pulse sequences for performing quantum process tomography on a single ion's atomic and motional state is implemented. These components are sufficient to perform a CNOT gate on the atomic state and motional state of a single ion. It is expected that amplitude pulse shaping would further improve the gate fidelity. These techniques, realized in a relatively simple experimental system, make the single ion a possible tool for studying other interesting quantum mechanical systems, as well as offering a viable option for realizing a large-scale quantum processor.

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- [1] H. Häffner, C. F. Roos, and R. Blatt, *Physics Reports* **469** (2008).
  - [2] R. Blatt and D. Wineland, *Nature* **453**, 1008 (2008).
  - [3] J. Benhelm, G. Kirchmair, C. F. Roos, and R. Blatt, *Nature Physics* **4**, 463 (2008).
  - [4] D. Leibfried, E. Knill, S. Seidelin, J. Britton, R. Blakestad, J. Chiaverini, D. Hume, W. Itano, J. Jost, C. Langer, et al., *Nature* **438**, 639 (2005).
  - [5] H. Häffner, W. Hänsel, C. Roos, J. Benhelm, D. C. al kar, M. Chwalla, T. Körber, U. Rapol, M. Riebe, P. Schmidt, et al., *Nature* **438**, 643 (2005).
  - [6] S. Seidelin, J. Chiaverini, R. Reichle, J. J. Bollinger, D. Leibfried, J. Britton, J. H. Wesenberg, R. B. Blakestad, R. J. Epstein, D. B. Hume, et al., *Phys. Rev. Lett.* **96**, 253003 (2006).
  - [7] D. Stick, W. K. Hensinger, S. Olmschenk, M. J. Madsen, K. Schwab, and C. Monroe, *Nature Physics* **2**, 36 (2006).
  - [8] R. J. Epstein, S. Seidelin, D. Leibfried, J. H. Wesenberg,

- J. J. Bollinger, J. M. Amini, R. B. Blakestad, J. Britton, J. P. Home, W. M. Itano, et al., *Phys. Rev. A* **76**, 033411 (2007).
- [9] A. Steane, *Quantum Inf. Comput.* **7**, 171 (2004).
- [10] J. Labaziewicz, Y. Ge, P. Antohi, D. Leibbrandt, K. R. Brown, and I. L. Chuang, *Phys. Rev. Lett.* **100**, 013001 (2008).
- [11] G. Gabrielse, J. Tan, P. Clateman, L. A. Orozco, S. L. Rolston, C. H. Tseng, and R. L. Tjoelker, *Journal of Magnetic Resonance* **91**, 564 (1991).
- [12] H. Häffner, S. Gulde, M. Riebe, G. Lancaster, C. Becher, J. Eschner, F. Schmidt-Kaler, and R. Blatt, *Phys. Rev. Lett.* **90**, 143602 (2003).
- [13] L. M. K. Vandersypen and I. L. Chuang, *Rev. Mod. Phys.* **76**, 1037 (2005).
- [14] L. Tian, P. Rabl, R. Blatt, and P. Zoller, *Phys. Rev. Lett.* **92**, 247902 (2004).
- [15] W. K. Hensinger, D. W. Utami, H.-S. Goan, K. Schwab, C. Monroe, and G. J. Milburn, *Phys. Rev. A* **72**, 041405 (2005).
- [16] J. Kim and C. Kim, *Quantum Inf. Comput.* **9**, 0181 (2008).
- [17] N. Daniilidis, T. Lee, R. Clark, S. Narayanan, and H. Häffner, *J. Phys. B* **42**, 154012 (2009).
- [18] Y. Ge, S. X. Wang, J. Labaziewicz, and I. L. Chuang (2009), in preparation.
- [19] P. B. Antohi, D. Schuster, G. M. Akselrod, J. Labaziewicz, Y. Ge, Z. Lin, W. S. Bakr, and I. L. Chuang, *Rev. Sci. Instr.* **80**, 013103 (2009).
- [20] J. Labaziewicz, P. Richerme, K. R. Brown, I. L. Chuang, and K. Hayasaka, *Opt. Lett.* **32**, 572 (2007).
- [21] A. D. Ludlow, X. Huang, M. Notcutt, T. Zanon-Willette, S. M. Foreman, M. M. Boyd, S. Blatt, and J. Ye, *Opt. Lett.* **32**, 641 (2007).
- [22] P. Pham, Master's thesis, Massachusetts Institute of Technology (2005).
- [23] F. Schmidt-Kaler, S. Gulde, M. Riebe, T. Deuschle, A. Kreuter, G. Lancaster, C. Becher, J. Eschner, H. Häffner, and R. Blatt, *Journal of Physics B: Atomic, Molecular and Optical Physics* **36**, 623 (2003).
- [24] A. A. B. Brojeny and J. R. Clem, *Phys. Rev. B* **68**, 174514 (2003).
- [25] Q. A. Turchette, D. Kielpinski, B. E. King, D. Liebfried, D. M. Meekhof, C. J. Myatt, M. A. Rowe, C. A. Sackett, C. S. Wood, W. M. Itano, et al., *Phys. Rev. A* **61**, 063418 (2000).
- [26] S. Haze, R. Yamazaki, K. Toyoda, and S. Urabe, *Phys. Rev. A* **80**, 053408 (2009).
- [27] D. F. V. James, *Appl. Phys. B* **66**, 181 (1998).
- [28] J. I. Cirac and P. Zoller, *Phys. Rev. Lett.* **74**, 4091 (1995).
- [29] A. M. Childs and I. L. Chuang, *Phys. Rev. A* **63**, 012306 (2000).
- [30] C. Monroe, D. M. Meekhof, B. E. King, W. M. Itano, and D. J. Wineland, *Phys. Rev. Lett.* **75**, 4714 (1995).
- [31] D. F. V. James, P. G. Kwiat, W. J. Munro, and A. G. White, *Phys. Rev. A* **64**, 052312 (2001).
- [32] M. Ježek, J. Fiuráek, and Z. c. v. Hradil, *Phys. Rev. A* **68**, 012305 (2003).
- [33] F. Schmidt-Kaler, H. Häffner, M. Riebe, S. Gulde, G. P. T. Lancaster, T. Deuschle, C. Becher, C. Roos, J. Eschner, and R. Blatt, *Nature* **422**, 408 (2003).
- [34] J. L. O'Brien, G. J. Pryde, A. Gilchrist, D. F. V. James, N. K. Langford, T. C. Ralph, and A. G. White, *Phys. Rev. Lett.* **93**, 080502 (2004).
- [35] C. F. Roos, G. P. T. Lancaster, M. Riebe, H. Häffner, W. Hänsel, S. Gulde, C. Becher, J. Eschner, F. Schmidt-Kaler, and R. Blatt, *Phys. Rev. Lett.* **92**, 220402 (2004).
- [36] M. Riebe, K. Kim, P. Schindler, T. Monz, P. O. Schmidt, T. K. Körber, W. Hänsel, H. Häffner, C. F. Roos, and R. Blatt, *Phys. Rev. Lett.* **97**, 220407 (2006).