Driven-dissipative dynamics of a strongly interacting Rydberg gas

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We study the non-equilibrium many-body dynamics of a cold gas of ground state alkali atoms weakly admixed by Rydberg states with laser light. On a timescale shorter than the lifetime of the dressed states, effective dipole-dipole or van der Waals interactions between atoms can lead to the formation of strongly correlated phases, such as atomic crystals. Using a semiclassical approach, we study the long-time dynamics where decoherence and dissipative processes due to spontaneous emission and blackbody radiation dominate, leading to heating and melting of atomic crystals as well as particle losses. These effects can be substantially mitigated by performing active laser cooling in the presence of atomic dressing.

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I. INTRODUCTION

Rydberg states are highly excited electronic states of atoms and molecules with large principle quantum numbers n. The remarkable properties of Rydberg states include their size r ∼ a0n2 with a0 the Bohr radius, implying huge polarizabilities α ∼ n7 and large electric dipole moments between Rydberg states, and thus strong coupling to external electric DC and microwave AC fields. Rydberg states can be excited from atomic or molecular ground states by laser light via absorption of one or more photons, with Rabi frequencies scaling as ∼ n−3/2. On the other hand, Rydberg states are long-lived, with lifetimes τ scaling as τ ∼ n5 (τ ∼ n3) for low (high) angular momentum states. For typical present experiments with n ∼ 50 these can be as large as tens of μs. Furthermore, atoms prepared in Rydberg states interact strongly, and these interactions can be controlled and enhanced by external fields. In particular, the van der Waals (vdW) interaction between ns states scales as V_{vdW} ∼ (ea0)4n11/r6, with e the electron charge, and can be made both attractive and repulsive. In the presence of external DC or AC electric fields the Rydberg-Rydberg interaction is a dipolar interaction, V_{dd} ∼ (ea0n2)2/r3. The latter is both long-range and anisotropic. Additional tunability can be achieved using, e.g., Förster resonances. These phenomena have been explored in recent experiments [6,20].

The remarkable properties of Rydberg states, and the tunability and strength of interactions between Rydberg atoms is reflected in novel many-particle physics of Rydberg gases [21,31], and provide the basis for applications in quantum information processing, e.g. in implementing (fast) quantum gates [32,37]. An underlying principle is the Rydberg-blockade mechanism, as first proposed in Ref. [32] and [33]. In the blockade regime the presence of a single atom excited to a Rydberg state shifts the energy levels of the surrounding atoms within a characteristic radius r_b (∼ μm), such that the laser excitation probability to the Rydberg state for any other atom within a volume ∼ r_b^3 is strongly suppressed. This excitation will be delocalized among all particles within r_b forming a “superatom”, with a collectively enhanced Rabi frequency √N_bΩ. Here, N_b is the number of atoms within r_b constituting a superatom and Ω is the single particle Rabi frequency. Various phenomena in the fields of many-body physics, quantum information applications and quantum optics related to the dipole-blockade mechanism have been discussed recently in Refs. [38–47].

Present experiments on Rydberg gases as a many-body system created by laser excitation from BECs [48,49], MOTs [6,18,20,50], atomic vapor cells [51] or optical lattices [19] have mainly explored the frozen gas regime. This corresponds to a short time dynamics, where the atomic motion can be ignored and (resonant) laser excitation of the Rydberg levels leads to large Rydberg-Rydberg interactions. In this limit atomic dissipation, e.g. spontaneous emission from Rydberg states, is negligible [52] and the dynamics maps to effective spin models, described by Hamiltonian dynamics [21,23,50,53].

An alternative regime is the Rydberg-admixed gas regime [25,28,54]: the idea is to admix the strong Rydberg-Rydberg interactions by off-resonant laser light weakly to the atomic ground states, thus providing an effectively much smaller, but still tunable vDW or dipolar ground state dynamics. That is, instead of kDebye dipole moments of Rydberg states one obtains effective ground state dipoles in the range of tens of Debyes [55]. Thus kinetic energies and effective interactions can become comparable, while at the same time spontaneous emission from the Rydberg state due to off-resonant laser tuning is strongly reduced. This effectively extends the lifetime of the gas, making it reminiscent of the dynamics of dipolar gases of polar molecules, as reviewed in [56,60].
by coupled Fokker-Planck equations for atoms effectively modeling the decay from the Rydberg state can be modeled in interaction between two Rydberg-dressed atoms. The dynamics of heating in such gases.

An interesting question emerging from the above discussion is, whether it is possible to form interesting condensed matter phases with these engineered \textit{atomic} dipolar gases; an example is provided by the formation of (stable) \textit{dipolar crystals of cold atomic gases} in analogy to dipolar crystals discussed for molecular particles [61]. In contrast to polar molecules [84] for Rydberg-admixed gases, dissipation and decoherence due to spontaneous emission is non-negligible for long times, and this raises the question of describing the long-time non-equilibrium dynamics of heating in such gases.

Understanding the long-term dynamics of Rydberg dressed atoms can also have important applications beyond many-body dynamics \textit{per se}. For example, recent proposals have investigated the possibility to efficiently couple cold atomic Rydberg gases to comparatively hot molecular ensembles via long-range dipolar or vdW interactions in order to achieve Doppler [62] and collisional Sisyphus-like [63] cooling schemes for molecules.

The paper is organized as follows: In Sec. II we give an overview of various atomic configurations studied, identify the main questions, and summarize the main results of the paper. Technical details of our calculations can be found in the remainder of the paper: In Sec. III we introduce the model and the notation which we will use throughout the paper. In Sec. IV A we discuss the Hamiltonian for a single Rydberg-dressed atom in the presence of external static and electromagnetic fields. The interaction Hamiltonian of two Rydberg-dressed atoms in the presence of external fields is considered in Sec. IV B where Born-Oppenheimer potentials are derived for both DC-electric and AC-microwave fields. In Sec. IV C we study the validity of the 2D treatment. In Sec. V we derive Fokker-Planck equations for laser cooled ground state atoms in the presence of Rydberg dressing. We find an additional two-body diffusion term due to the interaction between two Rydberg-dressed atoms. The dynamics of decay from the Rydberg state can be modeled by coupled Fokker-Planck equations for atoms effectively in the dressed and laser cooled ground state and atoms in one of the intermediate excited states. In Sec. VI A we study numerically spontaneous emission and blackbody radiation of a single Rydberg dressed atom. Population of intermediate excited states after decay from the Rydberg state will lead to a fluctuating dipole moment and heating of the motion due to photon recoil. The case of an interacting \textit{ensemble} of Rydberg dressed atoms is numerically studied in Sec. VI C. We find that the fluctuations of the dipole moment lead to strong mechanical effects and heating of the external motion, which strongly depends on the atomic density and the dressing scheme used. We study numerically the effect of laser cooling and melting in the presence of an optical lattice with the aim to extending the lifetime of the gas.

II. OVERVIEW OF RESULTS

Before presenting a detailed discussion of the dynamics of strongly interacting Rydberg dressed atoms, we find it worthwhile to summarize the main features of Rydberg dressing in a gas of (realistic) atoms, and the resulting many-particle dynamics. The processes we summarize here are derived in detail in Sec. III - VI.

Motivated by proposals for many-body physics with dipolar gases [61, 64], here we focus on the dynamics of a
Rydberg-dressed dipolar gas. We assume that the system is initially prepared in a crystalline state, e.g. by preparing a Mott insulator state of atoms in an optical lattice, and turning on adiabatically the Rydberg admixture while switching off the optical lattice, so that a dipolar crystal is formed. Similar to the setups of self-assembled crystals for polar molecules studied in Refs. [61, 64], we assume a two-dimensional (2D) configuration, i.e. that atoms are confined to the \((x,y)\)-plane by an external optical field, e.g., an optical lattice, in the z-direction (see Fig. 1), such that in-plane dipole-dipole interactions are purely repulsive. As explained below, this will minimize collisional losses linked to the attractive part of dipole interactions. This analysis is readily extended to, and actually simplified in the case of isotropic repulsive vdW interactions in three dimensions [65].

The main question we want to address is how heating and dissipation affect the many-body dynamics due to decay from the Rydberg state in the long-time limit. In fact, for times long enough, spontaneous emission and blackbody radiation will inevitably redistribute the population from the Rydberg state to various different excited states, which invalidates the simple two-level approximation for the internal dynamics of a single atom, mostly discussed so far [21, 31]. Given the fact that atoms in these other excited states can interact very differently from those prepared in the dressed ground state, e.g. they can have different dipole moments, decay from the Rydberg state can lead to strong mechanical effects and collisional losses in an ensemble of interacting Rydberg dressed atoms. In the following we will investigate this complex many-body dynamics. In particular, we will explore how to mitigate and control these effects using laser-cooling or in-plane optical lattices for a realistic scenario where each atom comprises a large number of internal states.

### A. Atomic configuration

The atomic configurations we have in mind are summarized in Fig. 2. (i) in panel (a) a DC-electric field with strength \(F\) polarizes each atom by splitting its energy levels into the Stark structure; the new Stark-split Rydberg state \(|r_e\rangle\) obtains an intrinsic dipole moment \(d_r\), which can be either parallel or antiparallel with respect to the external field. (ii) in panel (b) an AC-(microwave)-field of Rabi frequency \(\Omega_r\) is used to strongly couple two Rydberg states \(|r\rangle\) and \(|s\rangle\), which induces an oscillating dipole proportional to the transition dipole moment \(d_{rs}\).

In both configurations of Fig. 2 we weakly admix the ground state \(|g\rangle\) of each atom with the Rydberg state \(|r_e\rangle\) or \(|r\rangle\), respectively, using an off-resonant continuous wave laser with Rabi frequency \(\Omega_r\) and detuning \(\Delta_r \gg \Omega_r\). This immediately results in an effective dipole moment \(\langle \hat{d}_r \rangle \sim (\Omega_r/2\Delta_r)^2 d_0\) into the dressed ground state \(|\tilde{g}\rangle \sim |g\rangle + (\Omega_r/2\Delta_r)|r\rangle\), which can be tuned using the laser parameters, see Sec. IV B. Here, \(d_0 = d_r\) or \(d_0 = d_{rs}\) for DC-electric fields or AC-microwave fields, respectively, see Sec. IV B. This dressed ground state \(|\tilde{g}\rangle\) has now a finite, albeit comparatively long, lifetime \(\sim 1/\Gamma_{\tilde{g}}\), where \(\Gamma_{\tilde{g}} \sim (\Omega_r/2\Delta_r)^2 \Gamma_r\) with \(\Gamma_r\) the decay rate of the Rydberg state.

For timescales which are comparable to, or even larger than \(1/\Gamma_{\tilde{g}}\) population in each of the Rydberg states \(|r_e\rangle\) or \(|r\rangle\) and \(|s\rangle\) will be redistributed due to spontaneous emission and blackbody radiation among several excited states \(|m\rangle\). In the long-time limit, after, e.g., a spontaneous emission event from the Rydberg states, the atomic state will in general not return to the ground state directly, but via a cascade process where several \(|m\rangle\)-states are populated. Since each \(|m\rangle\)-state has a finite lifetime, the cascade will not happen instantaneously. Population of these intermediate states can induce strong mechanical effects on the gas dynamics, leading to heating and losses, as explained below in Sec. IV B. It turns out that these effects depend crucially on how the dipole moment \(d_0\) in the Rydberg states is created, see Fig. 2 (a) and (b).

We analyze these effects in detail by performing semi-classical molecular dynamics simulations for an ensemble of interacting Rydberg-dressed atoms confined to a 2D geometry including a large number of internal states. In Figs. 3 (a) and (b), we provide representative examples for the cases (i) and (ii) above, respectively, for a system comprising \(N = 67\) atoms in a box with hard walls. The external dynamics is treated classically, while quantum jumps from the Rydberg state account for spontaneous emission and blackbody radiation, leading to a time-dependent dipole moment of the atoms. In the simulation we use model-atoms with several thousands of internal electronic states with the level structure and mass \(M\) of \(^{85}\)Rb. At time \(t = 0\) the atoms are prepared in a triangular crystal structure with a given density \(n_{2D}\) (see figure caption) at zero temperature \(T = 0\). In Fig. 3 the thin dashed lines are the initial melting temperatures \(T_M = 0.089 \frac{\Delta E}{k_B^3/n_{2D}^{3/2}}\) as determined in Ref. [66], with \(k_B\) Boltzmann’s constant. The solid blue and dotted green lines correspond to the mean atomic kinetic energy \(E_{\text{kin}}(t)\) and the total atom number \(N(t)\), respectively, plotted as a function of time \(t\). In each plot, the simulation time corresponds to \(1/\Gamma_{\tilde{g}}\), for the given choice of parameters.

### B. Rydberg dressing with a DC-electric field

Figure 3 (a) shows an example where we choose a very strong DC-electric field with strength \(F = 3\) kV/cm in order to polarize \(^{85}\)Rb atoms. Such a high electric field is motivated by studying mixtures of polar molecules and Rydberg atoms where both species are polarized by the same field [67]. In order to avoid, e.g., field-ionization with threshold scaling as \(F_{\text{ion}} \sim n^{-4}\), the choice of Rydberg states is thus limited to those with a low principal quantum number, e.g. \(n \sim 16\). Here, we use \(|\tilde{r}\rangle = |16d, m = 0\rangle\), with lifetime \(\tau_r \sim 5.5\) \(\mu s\) and field-induced dipole mo-
FIG. 3: (color online) Single trajectory of a semi-classical molecular dynamics simulation studying the non-equilibrium and melting dynamics of a 2D Rydberg-dressed crystal as a function of time $t$. Heating due to decay from the Rydberg states and population of intermediate states $|m_{(F)}\rangle$ leads to a rapid increase of the mean kinetic energy, $E_{\text{kin}}(t)$ (solid blue line, left axis) and a reduction of the particle number due to losses (dotted green line, right axis). In panel (a) the dipole moment in the Rydberg state is created using the DC-electric field dressing scheme of Sec. II B and the same atomic parameters as given there. Initially we choose a density $n_{2D} = 1 \mu m^{-2}$. Two insets show histograms of the momentum distribution at $t = 27 \mu s$ and $t = 40 \mu s$. In panel (b) the dipole moment in the Rydberg state is created using the AC-microwave dressing scheme of Sec. II C with the same atomic parameters as given in the text. The initial density is $n_{2D} = 0.2 \mu m^{-2}$. In both panels the simulation time corresponds to the effective lifetime of the dressed ground state atoms, $\tau_\vartheta = 125 \mu s$ or $\tau_\vartheta = 12.8 \text{ ms}$, respectively. The thin dashed line is the initial melting temperature, $T_M$, of the crystal.

C. Rydberg dressing with a AC-microwave field

In Fig. 3(b) we choose a strong and resonant AC-microwave field to couple the Rydberg states $|r\rangle = |50s\rangle$ and $|s\rangle = |49p\rangle$ (see level scheme of Fig. 2(b)), which have a transition dipole moment of $d_{rg} \sim 5 \text{ kDebye}$. A dressing laser with $\Omega_r/\Delta_r = 0.14$ yields dressed ground states with $d_\vartheta \sim 30 \text{ Debye}$ and $\tau_\vartheta \sim 12.8 \text{ ms}$. The comparatively long lifetime is due to the choice of a state with a larger principal quantum number $n \sim 50$. Since $F = 0$ intermediate Rydberg states $|m\rangle$ have essentially no dipole moment and we consider them as non-interacting.

Figure 3(b) shows that $E_{\text{kin}}(t)$ increases with $t$. This is again due to fluctuations of the atomic dipole moment between the value $d_\vartheta$ and zero, corresponding to the atom being in the dressed ground state $|\vartheta\rangle$ or in one of the intermediate states $|m\rangle$, respectively, together with photon recoil after a decay event. As discussed before, heating comes from the redistribution of the energy associated to local crystal distortions due to dipole fluctuations among all particles. For $t \lesssim 1.5 \text{ ms}$ the Rydberg dressed atoms are in a crystal-like phase, with $E_{\text{kin}}(t) < k_B T_M$. 

The heating and loss processes described above are interaction-dependent (since the interaction strength scales as $\sim n_{2D}^{3/2}$) and thus decrease considerably for a smaller atomic density. In Sec. VTC below we show how active laser cooling and an additional in-plane optical lattice can affect these processes.

| Number of particles | Time t [μs] | Kin. Energy [2 \pi kHz] |
|---------------------|-------------|------------------------|
|                     | 0           | 100                    |
|                     | 20          | 300                    |
|                     | 40          | 400                    |
|                     | 60          | 550                    |
|                     | 80          | 650                    |
|                     | 100         | 750                    |
|                     | 120         | 850                    |

| Number of particles | Time t [ms] | Kin. Energy [2 \pi kHz] |
|---------------------|-------------|------------------------|
|                     | 2           | 40                     |
|                     | 4           | 20                     |
|                     | 6           | 10                     |
|                     | 8           | 5                      |
|                     | 10          | 2                      |
|                     | 12          | 1                      |
In comparison with the DC-electric field case of Fig. 3a, the heating rate $E_{\text{kin}}(\tau_g)\gamma_g$ of the AC-microwave-dressing scheme is approximately one order of magnitude smaller while the particle loss rate $N(\tau_g)\gamma_g$ is approximately equal. This is because the initial density is smaller and the intermediate states have negligible dipole moments for AC-microwave scheme.

In the remainder of this work we derive a microscopic model for the long-time heating dynamics of Rydberg dressed atoms and discuss quantitatively the dependence of the heating and particle loss rates on the system parameters, e.g., the atomic density. We state under what conditions standard laser cooling can be performed in the presence of Rydberg dressing, and numerically investigate how it counters heating effects.

### III. THE MODEL

The purpose of this section is to introduce the physical model and the notation that we will use throughout the paper. The system is illustrated in Fig. 1: $N$ identical (alkali) atoms with momenta $\mathbf{p}_i$ at positions $\mathbf{r}_i$ ($1 \leq i \leq N$) are confined to a quasi two-dimensional (2D) geometry in the $(x,y)$-plane by a strong confinement along the $z$-axis.

The Hamiltonian dynamics of the system is governed by

$$H = \sum_{i=1}^{N} H^{(i)} + \sum_{i<j}^{N} H_{\text{int}}^{(ij)},$$  

which consists of a sum over single-particle terms $H^{(i)}$ (see Sec. IV) and two-particle interaction terms

$$H_{\text{int}}^{(ij)} = \frac{\mathbf{d}_i \cdot \mathbf{d}_j - 3(\mathbf{d}_i \cdot \mathbf{r})(\mathbf{d}_j \cdot \mathbf{r})}{4\pi\epsilon_0 r^3},$$

which account for the dipole-dipole interaction between two atoms. Here, $r = r_i - r_j = \mathbf{r}$ is the relative distance between the atoms, $\mathbf{d}_i$ the dipole operator of the $i$-th atom and $\epsilon_0$ is the vacuum permittivity.

#### A. Internal level structure and setup

We denote $|\alpha\rangle = |n_{\alpha}, \ell_{\alpha}, j_{\alpha}, m_{\alpha}\rangle$ and $\hbar\omega_{\alpha} = E_{n_{\alpha}, \ell_{\alpha}, j_{\alpha}}$ as the unperturbed eigenfunctions and eigenenergies, respectively, of the atomic Hamiltonian $H_{\text{at}}$. Here, $n_{\alpha}$ is the principal quantum number, $\ell_{\alpha}$ the orbital angular momentum, $j_{\alpha}$ the total angular momentum and $m_{\alpha}$ projection of the total angular momentum along a specified axis.

In this manifold of states we focus on three specific states, see Fig. 4: (i) the energetic ground state of the atom denoted by $|g\rangle$. All internal energies will be measured relative to the energy of this state, e.g. $\hbar\omega_g = 0$. (ii) a highly excited Rydberg states $|r\rangle$ with energy $\hbar\omega_r$ and (iii) a lower-lying excited state $|e\rangle$ with energy $\hbar\omega_e \ll \hbar\omega_r$, which will be utilized for laser cooling. All other states $|\alpha\rangle$ can be occupied via spontaneous emission or blackbody radiation from the high-lying Rydberg states. As we have already mentioned in Sec. II, the ground state of each atom is off-resonantly coupled to a high-lying Rydberg state $|r\rangle$ using a continuous wave laser with a Rabi frequency $\Omega_r$ and detuning $\Delta_r (\gg \Omega_r)$, see Fig. 4. Typically, in alkali atom experiments, the atoms are excited to the Rydberg state via two-photon transitions in which $\Delta_r$ and $\Omega_r$ are the effective detuning and Rabi frequency. In addition, we use a laser field with Rabi frequency $\Omega_s$ and detuning $\Delta_s$ to couple $|g\rangle$ to a nearby excited state $|e\rangle$ ($\hbar\omega_e \ll \hbar\omega_r$) forming a closed cycle for laser-cooling.

Besides the general setup described above, we consider two different scenarios to create a strong dipole moment in the Rydberg state: (i) an homogeneous electric DC field of strength $F$ polarizes the atoms perpendicular to the 2D plane by splitting the energy levels into the Stark structure, see Fig. 4(a). The new Stark-split eigenstates $|\alpha_F\rangle$ obtain a large intrinsic dipole moment, e.g. $\mathbf{d}_0 = \langle r_F | \mathbf{d} | r_F \rangle$. (ii) a resonant microwave field with Rabi frequency $\Omega_s$ couples $|r\rangle$ to a nearby Rydberg state $|s\rangle$. Thereby the atom acquires a large oscillating dipole moment proportional to the transition dipole moment $\mathbf{d}_0 = \langle r | \mathbf{d} | s \rangle$. In both scenarios, the dipole moments can be as large as kilo-Debyes. The corresponding dipole-dipole interaction between two atoms in the Rydberg states is governed by $H_{\text{int}}$ of Eq. (2).

#### B. Master equation dynamics

The dynamics of the driven-dissipative many-body system made of interacting Rydberg-dressed atoms is described by the following master equation

$$\dot{\rho} = -i[H,\rho] + \mathcal{L}\rho,$$  

where the system density operator $\rho$ acts both on the Hilbert space of internal $|\alpha_1, \alpha_2, \ldots, \alpha_N\rangle$ and external $\{(r_1, p_1), \ldots, (r_N, p_N)\}$ degrees of freedom. This equation is readily written down as an extension of the familiar master equations of laser cooling for multi-level atoms including interaction between atoms [68, 69]. We denote $\mathcal{L}$ as the Lindblad operator, accounting for the decoherence due to spontaneous emission and blackbody radiation

$$\mathcal{L}\rho = \sum_{i=1}^{N} \sum_{\alpha, \beta} \Gamma_{\alpha\beta} D_{\alpha\beta}^{(i)} \rho,$$

which results from the coupling between the atomic system and the vacuum modes of the electromagnetic field, which have been adiabatically eliminated. The decay rate from state $|\alpha\rangle$ to $|\beta\rangle$,

$$\Gamma_{\alpha\beta} = \Gamma_{\alpha\beta}^{(\text{SE})} + \Gamma_{\alpha\beta}^{(\text{BBR})},$$
is obtained as the sum of respective contributions from spontaneous emission (SE) and blackbody radiation (BBR) \( \frac{70}{70} \), with

\[
\Gamma_{\alpha\beta}^{(BBR)} = \bar{n}_{\alpha\beta}(T)A_{\alpha\beta}, \quad (6a)
\]

\[
\Gamma_{\alpha\beta}^{(SE)} = \begin{cases} 
A_{\alpha\beta}, \quad \omega_{\alpha} > \omega_{\beta}; \\
0, \quad \omega_{\alpha} \leq \omega_{\beta}.
\end{cases} \quad (6b)
\]

Here, \( A_{\alpha\beta} \) is the Einstein A-coefficient and \( \bar{n}_{\alpha\beta} \) is the photon distribution, with

\[
\bar{n}_{\alpha\beta}(T) = \frac{1}{e^{\hbar(\omega_{\alpha} - \omega_{\beta})/k_{B}T} - 1}, \quad (7b)
\]

respectively. We use the notation

\[
\mathcal{D}_{\alpha\beta}^{(i)} = \int d^{2}k N_{\alpha\beta}(k)e^{-ik_{\alpha}a_{\beta}k_{\tau}}\rho_{\alpha\beta}^{(i)}e^{ik_{\alpha}a_{\beta}k_{\tau}}, \quad (8)
\]

to denote a general Lindblad term accounting for population redistribution from level \( |\alpha\rangle \) to \( |\beta\rangle \) of the \( i\)-th atom \((1 \leq i \leq N)\) with \( \sigma_{\beta\alpha}^{(i)} = |\alpha\rangle \langle \beta| \). With \( N_{\alpha\beta}(k) \) we denote the angular distribution of spontaneous emission from level \( \alpha \) to \( \beta \), which we assume to be a normalized and even function. The terms \( e^{ik_{\alpha}a_{\beta}k_{\tau}} \) describe the recoil of the atom due to a spontaneously emitted photon, where \( k_{\alpha\beta} = (\omega_{\alpha} - \omega_{\beta})/c \) is the wave number of the corresponding transition, with \( c \) the speed of light and \( k \) a unit-vector in the direction of spontaneous emission.

**IV. HAMILTONIAN DYNAMICS**

We now turn to analyze the single particle and twoparticle-interaction Hamiltonians of Eq. \((1)\). In particular, we derive the effective Born-Oppeheimer (BO) potential surfaces for two interacting Rydberg-dressed atoms in the presence of either a DC-electric-field (Sec. [IV B 1]) or an AC-microwave-field (Sec. [IV B 2]).

**A. Single particle Hamiltonian**

The single particle Hamiltonian \( H^{(i)} \) of Eq. \((1)\) consists of five terms

\[
H^{(i)}(t) = \hat{p}_{t}^{2}/2M + H_{at}^{(i)} + H_{dc}^{(i)} + H_{laser}^{(i)}(t) + H_{trap}. \quad (9)
\]

The first term in Eq. \((9)\) accounts for the kinetic energy of the \( i\)-th atom with mass \( M \). The second term

\[
H_{at}^{(i)} = \sum_{\alpha} \hbar\omega_{\alpha}\sigma_{\alpha\alpha}^{(i)}, \quad (10)
\]

accounts for the internal atomic energy levels, in the absence of external fields. The third term in Eq. \((9)\) reads

\[
H_{dc}^{(i)} = -\hat{d}^{(i)} \cdot \mathbf{E}_{dc} = -\hat{d}_{z}^{(i)} F, \quad (11)
\]

and describes the interaction of an atom with a static electric field, where \( \hat{d}^{(i)} \) is the atomic dipole operator of the \( i\)-th atom. The effect of a static electric field \( \mathbf{E}_{dc} = F \mathbf{e}_{z} \) is to polarize the atoms along the field direction \( \mathbf{e}_{z} \), by splitting the energies into the Stark structure \([71]\), with \( F \) the strength of the field. The new Stark-split eigenstates \( |\alpha F\rangle \) have an intrinsic dipole moment, which in the linear Stark regime is approximately given by \( d_{\alpha} = (3/2)e_{\alpha}a_{\alpha}(n_{1} - n_{2}) \) with \( a_{\alpha} \) Bohr’s radius and \( n_{1} \) and \( n_{2} \) the parabolic quantum numbers of state \( |\alpha F\rangle \). In this regime the energy levels are shifted proportional to the field strength, e.g. \( \Delta E_{\alpha} = \alpha_{F} F \). Coupling between adjacent \( n\)-manifolds can be neglected for field strength \( F < F_{\kappa} \), where \( F_{\kappa} \sim n^{-5} \) is the Inglis-Teller-limit \([1]\).

The term \( H_{laser}^{(i)} \) in Eq. \((9)\) describes the interaction of an atom with (e.g., microwave or optical) laser fields and consists of three terms

\[
H_{laser}^{(i)}(t) = H_{eg−laser}^{(i)}(t) + H_{rg−laser}^{(i)}(t) + H_{st−laser}^{(i)}(t), \quad (12)
\]

where each term has the form

\[
H_{eg−laser}^{(i)}(t) = \frac{\hbar\Omega_{\alpha}}{2}\sigma_{\alpha\beta}^{(i)} - i(k_{L\alpha}r_{\tau} - \omega_{L\alpha}t) + h.c., \quad (13)
\]

Here \( k_{L\alpha}/c \) is the wave-number of the laser with \( \omega_{L\alpha} \) the frequency of the laser, h.c. denotes the hermitian conjugate, and \( \Omega_{\alpha} \) is the Rabi frequency. The first term \( H_{eg−laser}^{(i)} \) in Eq. \((12)\) describes the coupling
of the atom to the cooling laser on the \((|g(F)| - |e(F)|)\)-transition, with Rabi frequency \(\Omega_r\) and frequency \(\omega_L\) detuned by \(\Delta_r\). The term \(H^{(i)}_{\text{rg-laser}}\) describes the coupling of the atom to the Rydberg-dressing laser on the \((|g(F)| - |r(F)|)\)-transition, with Rabi frequency \(\Omega_r\) and a frequency \(\omega_{L_r}\) detuned by \(\Delta_r\). In the following we are interested in the regime of large detuning \(\Delta_r \gg \Omega_r\) in order to weakly admix the Rydberg state to the ground state. Finally, \(H^{(i)}_{\text{sr-laser}}\) describes the coupling to a microwave field strongly mixing the Rydberg-states \(|r\rangle\) and \(|s\rangle\), with Rabi frequency \(\Omega_s\) and laser frequency \(\omega_{L_s}\) detuned by \(\Delta_s\).

In addition \(H^{(i)}_{\text{trap}}\) of Eq. (9) accounts for external trapping potentials.

B. Two-particle Hamiltonian: Born-Oppenheimer potentials

In this section we study the interaction between two Rydberg dressed atoms in the presence of an external static electric field (Sec. [IV B 1]) or a microwave field (Sec. [IV B 2]). For relative distances between the atoms larger than the size of the Rydberg atom \(r \sim a_0n_r^2\) and in the presence of external fields the atoms interact via dipole-dipole interaction governed by the Hamiltonian of Eq. (2). In particular, we derive the BO potential surfaces which, in the adiabatic approximation, play the role of effective interaction potentials \([61]\) (Eqs. (18) and (25)). They form the basis for the analysis of the time-dependent dynamics of Rydberg-dressed atoms which we discuss below in Sec. [V I] and [V].

1. DC-electric field

We consider two atoms in the presence of a static DC-field of strength \(F\), driven by the Rydberg-dressing laser, in the configuration of Fig. 4b. The microwave and the cooling laser are absent, i.e. \(\Omega_m = \Omega_c = 0\). The new Stark-split eigenstates \(|g_F\rangle \sim |g\rangle\) and \(|r_F\rangle\) are obtained by diagonalizing \(H_{\text{at}} + H_{\text{dc}}\) and the detuning of the dressing laser \(\Delta_r\) is defined relative to the shifted energy levels. To obtain the BO potentials we first neglect dissipation and treat the position operators \(\hat{r}_i\) as parameters \(r_i\). Within this limit each atom can be described by a two-state model consisting of \(|g\rangle\) and \(|r_F\rangle\), coupled by a dressing laser. This is valid for distances larger than \(r_n \sim (D/\Delta E_n)^{1/3}\), where diabatic crossings between BO-surfaces of neighboring \(n\)-manifolds can be neglected. Here, \(D = \frac{d_0^2}{4\pi\epsilon_0}\) is the dipolar coupling strength and \(\Delta E_n \gg \hbar\Delta_r\) is the energy separation between neighboring states. In a rotating frame the single particle Hamiltonian describing this model system reduces to

\[
H^{(i)} = -\hbar\Delta, \sigma^{(i)}_{rr} + \hbar\Omega_r (\sigma^{(i)}_{rr} + \sigma^{(i)}_{rg}),
\]

where the operator \(\sigma^{(i)}_{rr} = |\alpha_F\rangle \langle \beta_F|\) acts on the new Stark-split eigenstates, \(i \in \{1, 2\}\), and \(\Delta_r = \omega_L - \omega_r\) is the detuning from the \((|r_F\rangle - |g\rangle)\)-resonance. Position dependent phases of Eq. (13) will be included in Sec. [V].

Since the DC-electric field aligns the dipoles of the atoms along the direction of the field and the dominant dipole moment is \(d_0 = \langle r_F | \hat{d} | r_F \rangle\), the interaction term in Eq. (2) is

\[
H^{(ij)}_{\text{int}} = \frac{D(1 - 3 \cos^2 \vartheta)}{|\mathbf{r}_i - \mathbf{r}_j|^3} \left[ \sigma^{(i)}_{rr} \otimes \sigma^{(j)}_{rr} \right],
\]

where \(\vartheta\) is the angle between the dipole axis and the radial vector between two atoms. When the atoms are confined in a 2D-plane, e.g. by a strong optical potential, the angle is fixed to \(\vartheta = \pi/2\) resulting in a purely repulsive interaction. Within this model, the total Hamiltonian \([\text{Eq. (1)}]\) for two atoms in the basis \(|g, g\rangle, |r_F, g\rangle, |g, r_F\rangle, |r_F, r_F\rangle\) reads as

\[
H = \hbar \left( \begin{array}{cccc}
0 & \frac{1}{2} \Omega_r & \frac{1}{2} \Omega_r & 0 \\
\frac{1}{2} \Omega_r & 0 & -\Delta_r & \frac{1}{2} \Omega_r \\
\frac{1}{2} \Omega_r & -\Delta_r & 0 & \frac{1}{2} \Omega_r \\
0 & \frac{1}{2} \Omega_r & \frac{1}{2} \Omega_r & V(r) - 2\Delta_r
\end{array} \right),
\]

with

\[
\hbar V(r) = \langle r_F, r_F | H^{(12)}_{\text{int}} | r_F, r_F \rangle = \frac{D}{r^3}.
\]

The BO potentials for the scattering of two Rydberg-dressed atoms are obtained by diagonalizing the Hamiltonian of Eq. (16) for fixed relative position and zero kinetic energy, see Fig. 5. We assume that the linewidths of the corresponding states are smaller than the energy separation between them, and that the relevant energies are small enough, such that Landau-Zener transitions between different BO surfaces can be neglected (secular approximation). In this case the resulting position dependent eigenvalues act as potentials in each state manifold \([61][72]\). Asymptotically (e.g., at large distances where the dipole interaction is negligible), the new dressed eigenstates are \(|\tilde{g}\rangle = |g\rangle + (\Omega_r/2\Delta_r) |r_F\rangle\) and \(|\tilde{r}_F\rangle = |r_F\rangle - (\Omega_r/2\Delta_r) |g\rangle\). We note that the dynamics governed by the Hamiltonian of Eq. (16) for atoms initially in the ground state is restricted to the symmetric subspace made of the three states which asymptotically connect to the states \(|\tilde{g}, \tilde{g}\rangle, (|\tilde{r}_F, \tilde{g}\rangle + |\tilde{g}, \tilde{r}_F\rangle)/\sqrt{2},\) \(|\tilde{r}_F, \tilde{r}_F\rangle\), with energies slightly perturbed by the dressing laser field; the antisymmetric state \((|\tilde{r}_F, \tilde{g}\rangle - |\tilde{g}, \tilde{r}_F\rangle)/\sqrt{2}\) is decoupled, and does not contribute to the dynamics.

In the following, we focus on blue detuning, e.g. \(\Delta_r > 0\). Fig. 5 shows the BO-potentials as a function of the interparticle distance \(r\) for a specific choice of parameters. Because of the choice of blue-detuning, the
Condon point at

\[ r_c = \left( \frac{d_0^2}{8\pi\epsilon_0\hbar\Delta_r} \right)^{1/3} \]  

between the ground state BO-potential and other energy surfaces. As a consequence, there is a sudden change in the slope of the energy surface for \( r \approx r_c \), where the ground state BO-potential inherits the character of the one that asymptotically connects to the energy of the state \( |\vec{r}_F, \tilde{r}_F\rangle \), and becomes strongly repulsive. This effect has been discussed in Refs. [61, 65, 72] in the context of so-called blue-shielding techniques, where the strong repulsion for \( r < r_c \) does not allow for particles to come close to each other in a scattering event, thus preventing collisional losses due to, e.g., collision-induced ionization at short distance. In the reminder of this work, we will be mostly interested in confining the dynamics to distances \( r > r_c \).

2. AC-microwave-field

In this section we consider the Hamiltonian dynamics of two atoms in the presence of a linearly polarized, near-resonant microwave field with Rabi frequency \( \Omega_s \) coupling the Rydberg states \( |r\rangle \) and \( |s\rangle \), see Fig. 3(b). The ground state is again weakly dressed with the state \( |r\rangle \) using an off-resonant laser with Rabi frequency \( \Omega_r \) and a large detuning \( \Delta_r \gg \Omega_r \). The DC-electric field and the cooling laser are absent, i.e. \( \Omega_c = 0 \) and \( F = 0 \). The explicit choice of a near-resonant microwave field is to obtain large dipoles for atoms in the dressed ground state, which scale as \( \sim (\Omega_r/\Delta_r)^2 \), similar to the DC-electric field case of Sec. [IV B 1]. Again neglecting dissipation and the external degrees of freedom for a moment we are left with a three level system consisting of the ground state \( |g\rangle \) and the two Rydberg states \( |r\rangle \) and \( |s\rangle \). Since the electric DC-field is absent, these states are the bare eigenstates of \( H_{st} \). Below we show that, by a judicious choice of systems parameters, it is possible to obtain an interaction strength and Condon radius of similar magnitude as in the previous scheme of Sec. [IV B 1] above.

In a frame rotating with the laser frequencies the single particle Hamiltonian of Eq. (1) for this model system reduces to

\[
H^{(i)} = -\hbar\Delta_r \sigma_{sr}^{(i)} + \hbar(\Delta_s + \Delta_s) \sigma_{ss}^{(i)} + \frac{\hbar\Omega_r}{2} (\sigma_{rg}^{(i)} + \sigma_{gr}^{(i)}) + \frac{\hbar\Omega_c}{2} (\sigma_{sr}^{(i)} + \sigma_{rs}^{(i)}),
\]

where the operator \( \sigma_{\alpha\beta}^{(i)} = |\alpha\rangle \langle \beta| \) acts on the bare eigenstates, \( i \in \{1, 2\} \), and \( \Delta_s = \omega_{Ls} - \omega_s \) is the detuning of the microwave laser from the \( |r\rangle - |s\rangle \)-resonance. The corresponding dipole-dipole interaction Hamiltonian of Eq. (2) reduces to

\[
H_{int}^{(ij)} = \frac{d_0^2}{4\pi\epsilon_0} \frac{1 - 3\cos^2 \vartheta}{|r_i - r_j|^3} \left[ \sigma_{sr}^{(i)} \sigma_{sr}^{(j)} + \sigma_{rs}^{(i)} \sigma_{rs}^{(j)} \right],
\]

Figure 3 shows that the energy of the dressed two-particle ground state state \( E_{\tilde{g}\tilde{g}} \) (solid green line) is strongly affected by the dipole-dipole interactions, and for \( V(r) \sim 2\Delta \) avoided crossings occur among the BO-potentials of all symmetric states. Note that the energy of the antisymmetric state \( (|\vec{r}_F\tilde{g}\tilde{g}⟩ - |\tilde{g}\vec{r}_F\tilde{g}⟩)/\sqrt{2} \) is uncoupled and not shown. In particular, there is a resonant state interaction potential

\[
E_{\tilde{g}\tilde{g}}(r) = \left( \frac{\Omega_r}{2\Delta_r} \right)^4 \frac{d_0^2}{r^3} \quad \text{as}
\]

where the second line is valid in the limit \( V(r) \ll \Delta_r \), i.e. \( r \gg r_c \) (see below). While the first two terms on the r.h.s. of Eq. (18) are simple light shifts, the equation shows that at large distances the effective ground state BO-potential has an effective spatial dependence

\[
V_{\tilde{g}\tilde{g}}(r) = \left( \frac{\Omega_r}{\Delta_r} \right)^4 \frac{d_0^2}{r^3}.
\]

As explained in Refs. [25, 28, 54], this means that by dressing the particles with the laser field we have achieved a dipole-dipole interaction for atoms prepared in their (dressed) ground state, and the strength of the interaction is tunable by varying the ratio \( \Omega_r/\Delta_r \).
with \( \vartheta \) defined as before.

In the case of a near-resonant microwave field, where \( \Delta_s \ll \Omega_s \) it is convenient to perform a unitary transformation \( \mathcal{U}_s \) of Eq. (22) which diagonalizes the Hamiltonian in the subspace \( \{ |r\rangle_i, |s\rangle \} \). The corresponding new eigenstates are

\[
a_{\pm} = \frac{1}{\sqrt{2}} \left( 1 \pm \frac{\Delta_s}{\sqrt{\Delta_s^2 + \Omega_s^2}} \right)^{1/2},
\]

with the corresponding eigenenergies

\[
E_{\pm} = -\Delta_r - \frac{1}{2} \left( \Delta_s \mp \sqrt{\Delta_s^2 + \Omega_s^2} \right).
\]  

After the transformation, the single particle Hamiltonian of Eq. (20) in the basis \( \{ |g\rangle, |+\rangle, |−\rangle \} \) is

\[
\mathcal{U}_s^{(i)} H^{(i)} \mathcal{U}_s^{(i)} = \hbar \begin{pmatrix}
0 & \frac{1}{2} \Omega_{+} & -\frac{1}{2} \Omega_{−} \\
\frac{1}{2} \Omega_{−} & E_{+} & 0 \\
−\frac{1}{2} \Omega_{+} & 0 & E_{−}
\end{pmatrix},
\]

with the effective Rabi frequencies \( \Omega_{\pm} = a_{\pm} \Omega_r \). In this transformed picture the new eigenstates \( |\pm\rangle \) are coupled to the ground state by lasers with Rabi-frequencies \( \Omega_{\pm} \) and have a dipole moment \( \mathbf{d}_{\pm} = \langle \pm | \mathbf{d} | \pm \rangle = \pm \langle r | \mathbf{d} | s \rangle = \pm \mathbf{d}_0 \), see Fig. 6.

Again, the dynamics of the symmetric and antisymmetric states are decoupled. For two particles the dynamics of the symmetric subspace is governed by the Hamiltonian \( H = H^{(1)} + H^{(2)} + H^{(12)} \) which, represented in the basis \( \{ |g,g\rangle, \frac{1}{\sqrt{2}} (|g,+\rangle + |+,g\rangle), \frac{1}{\sqrt{2}} (|g,−\rangle + |−,g\rangle), \frac{1}{\sqrt{2}} (|+,−\rangle + |−,+\rangle), \frac{1}{\sqrt{2}} (|+,−\rangle + |−,+\rangle), \frac{1}{\sqrt{2}} (|−,−\rangle + |−,+\rangle) \} \), reads

\[
H_{\text{sym}} = \hbar \begin{pmatrix}
0 & \Omega_{+} / 2 & −\Omega_{−} / 2 & 0 & 0 & 0 \\
−\Omega_{−} / 2 & −\Delta_r + \Omega_{+} / 2 & 0 & −\Omega_{−} / 2 & 0 & 0 \\
0 & 0 & −\Delta_r − \Omega_{+} / 2 & −\Omega_{−} / 2 & 0 & 0 \\
−\Omega_{−} / 2 & 0 & −\Delta_r / 2 & −\Delta_r / 2 & −\Omega_{+} / 2 & −\Omega_{−} / 2 \\
0 & 0 & −\Omega_{−} / 2 & −\Omega_{+} / 2 & 0 & −\Omega_{s} \\
−\Omega_{−} / 2 & 0 & −\Omega_{−} / 2 & −\Omega_{+} / 2 & −\Omega_{s} & V − 2\Delta_r
\end{pmatrix},
\]

where we assumed exact resonance of the microwave field, i.e. \( \Delta_s = 0 \).

The BO-potentials are obtained by diagonalizing the Hamiltonian \( H_{\text{sym}} \), leading to the new dressed eigenstates which parametrically depend on \( r \). An analytic expression for the BO-potential of two atoms in the dressed ground state can be obtained perturbatively in the limit

\[
E_{gg}^{\text{MW}} = \frac{2\Delta_r \Delta_s^2}{4\Delta_s^2 + \Omega_s^2} \left[ 1 - \frac{\Omega_s^2 (4\Delta_r^2 + 3\Omega_s^2)}{(6\Delta_s^2 + \Omega_s^2)(4\Delta_s^2 + \Omega_s^2)} \right] - \frac{\Omega_s^2 (4\Delta_r^2 + 3\Omega_s^2)(4\Delta_s^2 - 4\Delta_r^2)}{(4\Delta_s^2 + \Omega_s^2)^2 (2V\Delta_r − 4\Delta_s^2 + \Omega_s^2)}.
\]

Note, that there are two resonances at \( \Delta_r = \Omega_s /2 \) and \( 2V(r_c^{\text{MW}})\Delta_r = 4\Delta_s^2 + \Omega_s^2 = 0 \). The first one corresponds to the level crossing between the states \( |g−\rangle \) and \( |−−\rangle \).
The latter inequality comes from the fact that we assume $\Omega_\alpha$ is the smallest frequency scale for the non-degenerate perturbation theory to be valid. Therefore, these two states must not be exactly degenerate. With this assumption we find for the ground state energy and the Condon radius

$$E_{gg}^M = E_{\text{const.}} + f(\alpha) \left( \frac{\Omega}{2\Delta_{\alpha}} \right)^4 V(r),$$

$$r_c^{MW} = g(\alpha) \sqrt{\frac{d_0^2}{8\pi \epsilon_0 \hbar \Delta_{\alpha}}},$$

with $E_{\text{const.}} = \Omega_\alpha^2/[2\Delta_{\alpha}(1-\alpha^2)] - \Omega_\alpha^4(1+\alpha^2)/[8\Delta_{\alpha}^3(1-\alpha^2)^3]$, $f(\alpha) = 2\alpha^2/(1-\alpha^2)^4$ and $g(\alpha) = 1/(1-\alpha^2)^{1/3}$. In the limit $\alpha \to 0$ the state $|s\rangle$ is not coupled to $|r\rangle$ and we obtain the same light shifts as in Eq. (18). Note that in this limit the interaction strength vanishes, $f(\alpha) \to 0$. This is due to the fact that in the absence of a DC-electric field the bare state $|r\rangle$ has no intrinsic dipole moment.

Figure 7(a) shows the BO-potentials for the symmetric states as a function of the interparticle distance $r$ for a specific set of parameters near resonance, as discussed above. The energy of the dressed two-particle states $|--\rangle$ and $|++\rangle$ is strongly shifted by the dipole-dipole interaction, and avoided crossings occur among the BO-potentials of all symmetric states. This leads to a sudden change in the slope of the energy surface of, e.g. $E_{gg}$, at the Condon radius $r_c^{MW}$.

The functions $f(\alpha)$ and $g(\alpha)$ versus $\alpha$ are shown in Fig. 7(b). Increasing $\alpha$ towards 1 will on one hand increase the effective ground state interaction potential according to $f(\alpha)$. On the other hand it will increase the Condon radius $r_c^{MW}$ according to $g(\alpha)$. Fig. 7(b) shows that there is a region, $\Delta\alpha$, for which $f(\alpha) > 1$ but $1 < g(\alpha) < 1.5$ between 0.45 < $\alpha$ < 0.84. Operating in this region leads to formal similar interaction strength and Condon radii as for the DC-electric field dressing scheme, e.g. Eq. (18) and Eq. (19).

C. Validity of the 2D treatment

In this section we examine in detail under what criteria we can treat the system of interacting Rydberg dressed atoms as purely 2D in nature. As mentioned before, the atoms are trapped in the $(x,y)$-plane by a strong harmonic confinement along the $z$-direction. The resulting three dimensional (3D) potential in relative coordinates reads

$$V_{3D}(r) = \frac{1}{4}M\omega_z^2z^2 + \hbar E_{gg}(r).$$

where the first term is the harmonic confinement with trapping-frequency $\omega_z$. The second term, $E_{gg}(r)$, is the BO potential of two interacting dressed ground state atoms, obtained by numerically diagonalizing the Hamiltonian of (10), taking into account the full 3D characteristic of the dipole-dipole interactions.
For relative distances larger than the Condon radius, $r_c$ [Eq. (19)], the interaction can be approximated by dipole-dipole interaction. In this case, the 3D potential can be rewritten as

$$V_{3D}^{\text{mol}}(r) = V_0 \left[ \frac{\kappa \tilde{z}^2}{\rho^2 + \tilde{z}^2} \right],$$

which is reminiscent of polar molecules [61, 64]. Here, $\tilde{z} = z/r_c$ and $\tilde{\rho} = (x^2 + y^2)^{1/2}/r_c$. The corresponding contour plot is shown in Fig. 8(b). The dimensionless parameter $\kappa$ characterizes the strength of the confinement relative to the interaction

$$\kappa = \left( \frac{2 \Delta_r}{\Omega_r} \right)^4 \frac{M \omega_0^2 r_c^2}{8 \hbar \Delta_r} = 2 \left( \frac{\Delta_r}{\Omega_r} \right)^3 \frac{\omega_\perp}{\Omega_r} \left( \frac{r_c}{a_{ho}} \right)^2,$$

where $a_{ho} = \sqrt{\hbar/M \omega_\perp}$ is the harmonic oscillator length in transversal direction. For $\kappa \gg 1$ the confinement in $z$-direction dominates over the dipole-dipole interaction.

Analytic analysis of Eq. (30) yields the saddle points

$$\rho_* = \pm 2 z_* \quad \text{and} \quad z_* = \frac{3^{1/5}}{\sqrt{5} \kappa^{1/5}}.$$  

The height of the potential at the saddle point, corresponding to the height of the energy barrier separating the repulsive region from the attractive region, is

$$V_{3D}^{\text{mol}}(z_*, \rho_*) = V_0 \left( \frac{\kappa}{3} \right)^{3/5},$$

which is independent of $\Delta_r$ for a fixed ratio of $\Omega_r/\Delta_r$. For the same parameters used above we obtain $\kappa = 30$ which yields a potential barrier at the saddle point corresponding to $T_s = 150 \mu K$. Comparing Fig. 8(a) and (b) one can see the rapid change of the interaction strength at the Condon radius for Eq. (29) (panel (a)) leads to a repulsive shield which leads to a slightly higher potential barrier corresponding to $T_s = 220 \mu K$ compared to the case of a pure inverse-cubic interaction potential (panel (b)) with $T_s = 150 \mu K$. Additionally we find that the saddle-point in panel (a) is shifted to larger $\rho$ values than in panel (b).

In a 3D scenario two particles undergo collapse when they have large enough collisional energy to overcome the energy barrier at the saddle points. In the numerical simulations of Sec. IV B we will treat the system as purely 2D with a $1/r^3$-potential (see Eq. (12)), assuming particles are being lost if they collapse in 3D. For a specific set of parameters we use the latter analysis to calculate the height of the potential at the saddle point, $V_{3D}(\rho_*, z_*)$, in a 3D scenario. The height of the saddle point will then be translated into a critical 2D distance, $\rho_{\text{loss}}$, at which $E_{gg}^{\text{loss}}(\rho_{\text{loss}}) = V_{3D}(\rho_*, z_*)$, with $E_{gg}$ defined in Eq. (18(b)).

**V. LASER COOLING OF DRESSED RYDBERG ATOMS**

In this section we examine in detail the dissipative processes in Eq. (3) and study laser cooling of interacting
atoms in the presence of Rydberg dressing. As an example we consider Doppler cooling but the model can be extended to sub-Doppler cooling schemes. For simplicity of the analytic treatment in Sec. V A we will first consider model atoms where we neglect decay from the Rydberg state in order to derive semiclassical Fokker-Planck equations which describe the cooling dynamics in the presence of Rydberg-dressing and interactions. In Sec. VI we will include decay from the Rydberg state which couples the dressed ground state via rate equations to one of the intermediate states. Again for simplicity of the analytic treatment in this section, we will first consider model atoms with a single intermediate state \( \ket{m} \). In Sec. VI we will numerically investigate the interplay of laser-cooling and heating due to population of intermediate states of an ensemble of dressed Rydberg atoms taking into a large number of internal states. We show that laser cooling can alleviate the heating dynamics for interacting Rydberg atoms described above, extending the lifetime of strongly interacting phases in these systems, as for self-assembled crystals.

\[
W^{(N)}(r_1 \ldots r_N; p_1 \ldots p_N; t) = \int \frac{du_1}{h^3} \ldots \frac{du_N}{h^3} \langle r_1 + \frac{u_1}{2} \ldots r_N + \frac{u_N}{2} | \rho | r_1 - \frac{u_1}{2} \ldots r_N - \frac{u_N}{2} \rangle e^{-i u_1 p_1 / \hbar} \ldots e^{-i u_N p_N / \hbar}. \tag{34}
\]

(ii) We derive the equation of motion for \( W \) using the master equation Eq. (3) and expand the resulting equation of motion up to second order in terms of the photon momentum in order to obtain a positive probability function in the semiclassical limit. (iii) We adiabatically eliminate the Rydberg state \( \ket{r} \) in the limit of a large detuning \( \Delta_e \). The resulting two-level systems - consisting of a dressed ground state \( \ket{\tilde{g}} = \ket{g} + (\Omega_e / 2\Delta_e) \ket{r} \) with dipole moment \( d_3 \) and an excited state \( \ket{e} \) - interact only when the atoms are in the dressed ground state. (iv) Finally, we adiabatically eliminate the fast internal degrees of freedom of this effective two-level system in favor of the (much slower) external dynamics. Due to the Rabi oscillations between the dressed ground state \( \ket{\tilde{g}} \) (with dipole moment \( d_3 \)) and the lower-lying excited state \( \ket{e} \) (with negligible dipole moment) the dipole of the atom is fluctuating in time. This leads to an additional diffusion term in the equation of motion in addition to the standard Doppler-cooling diffusion terms. After a lengthy calculation one obtains for two atoms in their dressed ground state, \( W_{gg}^{(2)} = \langle gg | W | gg \rangle \), the following equation of motion

\[
\left( \frac{\partial}{\partial t} + \sum_{i=1}^2 \frac{p_i}{M} \frac{\partial}{\partial r_i} \right) W_{gg}^{(2)} = \sum_{i=1}^2 \left[ \frac{\partial}{\partial p_i} f_c + D_1^{(i)} \right] W_{gg}^{(2)} - f_{12} \cdot \left( \frac{\partial}{\partial p_1} - \frac{\partial}{\partial p_2} \right) W_{gg}^{(2)} + D_{12} W_{gg}^{(2)}, \tag{35}
\]

which is valid when \( \Delta_e \gg \Omega_e, \Gamma_e \gg \hbar k_{c_{eg}}^2 / m \), where \( k_{c_{eg}} \) is the wave number of the \((e)\)-(g)-transition. Additional to the free motion term on the left hand side, the first term on the right hand side describes single-particle Doppler cooling with the cooling force \( f_c = \beta_k L_{ke} \cdot \mathbf{p} \) and the standard diffusion operator \( D_1 = \sum_k D_k \hat{p}_k^2 \) accounting for spontaneous emission and diffusion due to the cooling laser in various spatial directions \( k \in \{ x, y, z \} \). Here we assumed two counter-propagating laser beams and expanded the resulting radiation pressure forces up to second order in \( k_{L_{ke}} \). The third term of Eq. (35) proportional to

\[
f_{12} = \frac{\Omega_r^4}{4\Delta_e^2 (V - 2\Delta_e)^2} \frac{\partial V}{\partial r_1} = - \frac{\partial E_{gg}}{\partial r_1}, \tag{36}
\]

accounts for the interaction between the two atoms. We note that this term can be rewritten, using the potential of the dressed ground state, Eq. (15), obtained in section A. Fokker-Planck equation

In the following we will derive equations of motion for the external dynamics (position and momentum) of model atoms consisting of a ground state \( \ket{g} \) which is coupled with a far detuned laser to a Rydberg state \( \ket{r} \) with Rabi frequency \( \Omega_r \) and detuning \( \Delta_r \gg \Omega_r \). Additionally, the ground state \( \ket{g} \) is coupled to a lower-lying excited state \( \ket{e} \) using a counter-propagating laser with Rabi frequency \( \Omega_e \) and detuning \( \Delta_e \). The state \( \ket{e} \) decays directly to the ground state \( \ket{g} \) with a fast decay rate \( \Gamma_e \), realizing a closed cycle transition. Atoms which are both in the Rydberg state interact via dipole-dipole interaction described by the Hamiltonian Eq. (15).

The derivation of equations of motion for the external degrees of freedom is done in four steps [69, 74]: (i) first we use the Wigner function formalism to map the density operator \( \rho \) onto a quasi-probability distribution in phase space...
Besides those standard laser cooling terms, the term
\[
\mathcal{D}_{12} = \frac{4\Omega_{\Gamma}^2}{(\Gamma_e^2 + 4\Delta_e^2)^2} \left[ \mathbf{r}_{12} \cdot \left( \frac{\partial}{\partial p_1} - \frac{\partial}{\partial p_2} \right) \right]^2
\]
is a two-body diffusion term which accounts for the fluctuations of the force (which is present only in the dressed ground state), due to Rabi-oscillations between \(|\alpha\rangle\) and \(|\beta\rangle\). For near-resonant laser light it is proportional to the population and lifetime of the excited state. In the limit \(\mathbf{r}_{12} \ll h\Gamma_e \mathbf{k}_e\), this diffusion term is small compared to the single-particle diffusion terms and one can approximate the system with two interacting atom, which are independently laser cooled.

**B. Quantum jumps and rate equations**

A quantum jump occurs when the atom in the Rydberg state \(|r\rangle\) decays to an intermediate state \(|m\rangle\) due to spontaneous emission or blackbody radiation. In the following, we allow for the possibility that the intermediate state \(|m\rangle\) is long-lived, with a lifetime \(\sim 1/\Gamma_{mg}\) comparable with the external dynamics and thus cannot be adiabatically eliminated. This will lead to two coupled equations of motion: (i) an equation for laser-cooled atoms in the dressed ground state, with dipole moment \(d_g = (\Omega_r/2\Delta_r)^2 d_r\), and (ii) an equation of motion for atoms in the \(|m\rangle\)-state (and possibly with a dipole moment \(d_m\)), which do not experience a cooling force.

To simplify the notation for the analytic treatment, we consider only a single atom in the presence of the cooling and dressing lasers. The generalization to two atoms is straightforward. After performing steps (i)-(iv) of the latter section we find the following coupled FPEs for the dynamics of the dressed ground state \(|\tilde{g}\rangle\) and the intermediate state \(|m\rangle\):

\[
\begin{align*}
\left( \frac{\partial}{\partial t} + \frac{P}{M} \cdot \frac{\partial}{\partial \mathbf{r}} \right) W_{\tilde{g}}^{(1)} &= -\Gamma_{\tilde{g}} W_{\tilde{g}}^{(1)} + (\Gamma_m + \mathcal{D}_{mg}) W_{m}^{(1)} + \left[ \frac{\partial}{\partial \mathbf{p}} \cdot \mathbf{f} + \mathcal{D} \right] W_{\tilde{g}}^{(1)}, \\
\left( \frac{\partial}{\partial t} + \frac{P}{M} \cdot \frac{\partial}{\partial \mathbf{r}} \right) W_{m}^{(1)} &= -\Gamma_m W_{m}^{(1)} + \left[ \Gamma_{\tilde{g}} + \left( \frac{\Omega_r}{2\Delta_r} \right)^2 \mathcal{D}_{rm} \right] W_{\tilde{g}}^{(1)},
\end{align*}
\]

where \(W_{\alpha}^{(1)} = \langle \alpha | W^{(1)} | \alpha \rangle\) is the single particle Wigner function for the atom in state \(|\alpha\rangle\) \((\alpha \in \{\tilde{g}, m\})\). The equations are coupled by the effective ground state decay-rate

\[
\Gamma_{\tilde{g}} = \left( \frac{\Omega_r}{2\Delta_r} \right)^2 \Gamma_{rm}
\]
due to optical pumping via the Rydberg state \(|r\rangle\). The terms \(\mathcal{D}_{\alpha\beta}\) are standard diffusion operators accounting for spontaneous emission from state \(|\alpha\rangle\) to \(|\beta\rangle\).

**VI. DISSIPATIVE DYNAMICS OF DRESSED RYDBERG ATOMS**

We will now analyze numerically the dissipative processes of Eq. [3], which are associated with the finite lifetime of excited Rydberg states. In Sec. [VI A] below, we consider the case of a single atom, in which we investigate the population of intermediate states due to spontaneous emission or blackbody radiation. As a consequence, the atom acquires a time-dependent dipole moment. In addition, spontaneous emission and blackbody radiation act as small heating sources due to the photon recoil. In Sec. [VI B] we will summarize our previous discussion of the driven-dissipative dynamics of laser-cooled and interacting Rydberg-dressed atoms as a prescription for a molecular dynamics simulation. The effect of population in intermediate states will be more substantial in the case of a large AC-microwave field \((F > 0)\) the states \(|\alpha_F\rangle\) can have a large parallel or antiparallel dipoles, \(\mathbf{d}_\alpha = (\alpha_F \hat{d} |\alpha_F\rangle)\), on the order of kilo-Debye, causing strong dipole-dipole interactions with other Rydberg-dressed atoms. Whereas, in the case of a AC-microwave field (and no DC-electric field) the intermediate states do not have any dipole moment.

**A. Decoherence of a single atom**

The dressed ground state \(|\tilde{g}\rangle\) has a finite lifetime \(\tau_{\tilde{g}} = \Gamma_{\tilde{g}}^{-1}\), where \(\Gamma_{\tilde{g}} = (\Omega_r/2\Delta_r)^2 \Gamma_r\), and \(1/\Gamma_r\) is the
The first method we calculated the decay matrix in- 
...ry [75–77] and a model potential method [78]. With 
...method we obtained the decay matrix up to n = 110 including s, p, d, f and g- 
...angular momentum states. With Γα = ∑βΓαβ we denote the total decay rate of the state |α⟩ and Γ0 ≡ Γg = (Ωr/2Δr)2Γr = (Ωr/2Δr)2∑βΓrβ is the effective decay rate of the dressed ground state. Fig. 9 shows the branching ratio of decay in the absence of external fields from the |16d⟩ state to |np⟩ and |nf⟩ states, where we summed over j and mj levels. For T = 0 K (upper panel) the only contribution to the decay rate comes from spontaneous emission, which favors decay to low-lying states. About 60 % of the population in the |16d⟩ state decays to the |5p⟩ state. The lower panel of Fig. 9 shows the branching ratio for T = 300 K. In this case the decay rate is determined not only by spontaneous emission but also by blackbody radiation, which in addition leads to substantial decay to neighboring states. Hence, only 51 % of the population in the |16d⟩ states decays directly to the |5p⟩ state. From this it is clear that in current cold atom experiments performed in room temperature environments, the role of high-lying intermediate states cannot be neglected in the long-time limit.

The internal dynamics corresponding to a cascade process of the electron towards the ground state following a decay event via the Rydberg states is given by

\[
\frac{d}{dt}\begin{pmatrix} p_0 \\ p_1 \\ p_2 \\ \vdots \\ \vdots \\ \vdots \\ \vdots \end{pmatrix} = \begin{pmatrix} -\Gamma_0 & \Gamma_{10} & \Gamma_{20} & \cdots & \vdots & \vdots & \vdots \\ -\Gamma_{01} & -\Gamma_1 & -\Gamma_{21} & \cdots & \vdots & \vdots & \vdots \\ -\Gamma_{02} & -\Gamma_{12} & -\Gamma_2 & \cdots & \vdots & \vdots & \vdots \\ \vdots & \vdots & \vdots & \ddots & \vdots & \vdots & \vdots \\ \vdots & \vdots & \vdots & \vdots & \ddots & \vdots & \vdots \\ \vdots & \vdots & \vdots & \vdots & \vdots & \ddots & \vdots \\ \vdots & \vdots & \vdots & \vdots & \vdots & \vdots & \ddots \end{pmatrix} \begin{pmatrix} p_0 \\ p_1 \\ p_2 \\ \vdots \\ \vdots \\ \vdots \end{pmatrix}, \quad \text{(40)}
\]

where \( p_\alpha \) is the probability of being in the state |\alpha(F)⟩ in the absence (presence) of a static electric field, \( F = 0 \) (\( F > 0 \)). Note that in the case of a DC-electric field...
(F > 0) the decay rates $\Gamma_{\alpha\beta\gamma}$ are a sum of several decay rates between bare states, $\Gamma_{\alpha\beta}$, according to the contributions of various different angular momentum states to the Stark split states $|\alpha\rangle$ and $|\beta\rangle$.

**DC-electric field:** The Stark structure of atomic states (including only $m = 0$ states) is shown in Fig. 10 for an electric field along $e_z$. States with an angular momentum $\ell > 3$ are almost degenerate and exhibit a linear Stark effect, while lower angular momentum states, which have a larger quantum defect which breaks the degeneracy, exhibit a quadratic Stark shift. Fig. 10 shows that the $|16d\rangle$ state (thick green line) is well separated from the neighboring states by an energy gap of $\sim 200$ GHz up to a field strength of $F \sim 3$ kV/cm and experiences a strong energy shift, corresponding to a large dipole moment. For an electric field $E_{\text{dc}} = F e_z$ only $m = 0$ states are coupled and the resulting dipole moments are polarized along the direction of the external field. Fig. 11 shows the $z$-component of the dipole moment, $d_z$, for different states $|\alpha\rangle$ and a field strength of $F = 3$ kV/cm. The figure shows that states $|\alpha\rangle$ gain a large dipole moment of hundreds of Debye which can be positive and negative. The dipole moment of the state which for $F \to 0$ connects to $|16d, m = 0\rangle$ is about 680 Debye.

The time evolution of Eq. (10) is readily simulated. We prepare the atom in the dressed ground state $|g\rangle$, that is $p_\beta = p_\alpha = 1$. After a time step $\Delta t$ much smaller than the timescales associated with the decay rates a random number determines if and to which internal state $|\alpha\rangle$ the atom makes a transition (quantum jump) according to the rates of Eq. (10). During each time step the dipole moment $d(t)$ of the atom is equal to the dipole moment of the current state, $d_{\alpha(t)}$. This new configuration then again propagated another time step $\Delta t$.

Figure 12(a) shows an example result for the time evolution of the atomic dipole moment $d(t)$ of a $^{85}$Rb atom polarized by a DC-electric field with strength $F = 3$ kV/cm. The ground state is coupled to the Rydberg state $|F\rangle$ which in the absence of a DC-electric field ($F = 0$) connects to the $|r\rangle = |16d, m = 0\rangle$-state using a Rydberg-laser with $(\Omega_r/2\Delta_r) = 0.21$. We initially prepare the atom in the dressed ground state $|g\rangle$, and compute the time evolution using Eq. (10). The red trajectory in Fig. 12(a) shows a typical result for the deviation of the atomic dipole from the dipole moment of the dressed ground state $d_\gamma = 30$ Debye as a function of time $t$, in units of the ground state lifetime $\tau_\gamma = 125$ µs. The figure shows large positive and negative spikes for the values of $d$, followed by long times where the system is in the dressed ground state, with dipole moment $d_\gamma$. The large “spikes” correspond to the population of intermediate states $|\alpha\rangle$, during the cascade process towards the ground state, following a spontaneous emission event. These large positive and negative fluctuations of $d(t)$ will cause strong dipole-dipole interactions and hence large mechanical effects when a gas of interacting Rydberg atoms is considered.

**Microwave dressing and F=0:** In this case we couple the ground state $|s\rangle$ of Rubidium to the state $|t\rangle$ using a laser with $(\Omega_r/2\Delta_r)^2 = 0.05$. An additional microwave field couples the states $|t\rangle$ and $|p\rangle$ such that the dressed ground state obtains a dipole moment of $d_\gamma = 30$ Debye. The blue trajectory in Fig. 12(b) shows a typical trajectory for the time evolution of the dipole moment as a function of time in units of $\tau_\gamma = 90$ µs. For this dressing scheme the intermediate states $|m\rangle$ do not possess a dipole, and thus the dipole fluctuates between long periods when it has the value $d_\gamma$, corresponding to the atom in the dressed ground state, to periods where the atom has no dipole, corresponding to the cascade processes following spontaneous emission.
like a “blinking dipole”.

In comparison, in the DC-electric field case (blue trajectory in panel (a)) the dipole fluctuations are much larger and can take both positive and negative values. Whereas in the microwave case (red trajectory in panel (b)) the dipoles fluctuate between zero and $d_g$.

In the following we examine the external dynamics of a single atom. It is dominated by small momentum fluctuations associated with the photon recoil following a series of cascade decay events. As an example, we simulate heating due to decay from the $|g⟩ = |16d⟩$ state using the same parameters as before. The result of a molecular dynamics simulation is shown in Fig. 13 where we plot single kinetic energy trajectories as a function of time (thin black lines). We propagate the internal dynamics according to Eq. (40). Each decay event is associated with a momentum kick in a random direction corresponding to the transition, see next Section. It is shown in the figure that as time progresses the kinetic energy increases due to photon recoils. We perform a statistical average over 50 runs of simulations (thick green line) which yields an average heating rate of $2π \times 106.3$ kHz/ms. It will be shown below that for high enough densities this single particle heating rate is much smaller than the heating rate due to interactions associated with fluctuations of the dipole moment in the many-body case, as discussed in the next section.

B. Molecular dynamics simulation

In our semiclassical description the state of each atom at time $t$ is specified by $\{r_i(t), p_i(t), α_i\}$, where $r_i(t)$ and $p_i(t)$ are the center-of-mass coordinate and momentum of the $i$-th atom, respectively, and $α \in \{g, m, m', \ldots\}$ denotes the internal electronic state with $i = 1, \ldots, N$. The internal state is either the ground state $|g⟩$, dressed by the admixture of the Rydberg state, and the excited state $|e⟩$ from the laser cooling (see Fig. H), or one of the many intermediate states $|m⟩$ populated during a decay cascade back to the ground state. The dynamics of the gas is described by a set of coupled equations, as written down for the case of one or two atoms in Eqs. (38) and (35), respectively. These results can be generalized immediately to $N$ atoms. Mathematically, they are a set of coupled Fokker-Planck equations for the external motion, and rate equations for the internal electronic states, describing a combined diffusive and jump Markov process for the probability density $W^{(N)}(r_1, p_1, α_1; \ldots; r_N, p_N, α_N; t)$. This stochastic process is readily simulated.

Consider atom $i$, which we assume to be in the dressed ground state $|g⟩$. Its center of mass motion obeys the Langevin equations

$$\dot{r}_i = \frac{p_i}{M},$$
$$\dot{p}_i = \sum_{j \neq i} f_{\text{int}}^{(ji)} - \beta p_i + f_{\text{ex}}^{(i)} + F^{(i)} \tag{41b}$$

where on the right hand side of Eq. (41b) we sum over the forces from all the other ground state atoms, as well as the forces from atoms in one of the intermediate states $|m⟩$, with

$$f_{\text{int}}^{(ji)}(t) = -\frac{d_{α_i}d_{α_j}}{4πε_0} \frac{1}{|r_i - r_j|^3}. \tag{42}$$

Here, $d_{α_i} = d_g$ is the effective dipole moment of the dressed ground state (see Eqs. (18) and (25) for the DC- and AC-field cases, respectively) and $d_{α_j}$ can either be equal to $d_g$ if the $j$-th atom is in the dressed ground state $|g⟩$, or equal to $d_m$ if the $j$-th atom is in the state $|m⟩$. In the DC-electric field case $d_m = ⟨m_p⟩|d(m_p)|$, while in the AC-microwave field case $d_m = 0$, see Sec. VIA. In addition, we have added in Eq. (41b) the familiar terms describing possible laser cooling [74], and an external trapping force $f_{\text{ex}}^{(i)}$. The last term in Eq. (41b) is a stochastic force [79] from quantum fluctuations due to the recoil of spontaneous emission events from both the Rydberg state repopulating the ground state, but also from laser cooling, obeying

$$⟨F_k^{(i)}(t)F_l^{(i)}(t')⟩ = D_k δ_{kl} δ_{ij} δ(t - t’), \tag{43}$$

with $k, l \in \{x, y, z\}$ and $i, j \in 1, \ldots, N$ and $D_k$ the diffusion coefficients of Eq. (35). We make the simplifying assumption that the cross-noise term of Eq. (37) arising from the fluctuating ground state dipole due to laser cooling discussed in Sec. V is negligible.

Similarly, atom $i$ in one of the intermediate states $|m⟩$ obeys the equation of motion

$$\dot{r}_i = \frac{p_i}{M}, \quad \dot{p}_i = \sum_{j \neq i} f_{\text{int}}^{(ji)}, \tag{44}$$

where again $f_{\text{int}}^{(ji)}$ are the forces of Eq. (42) and $d_{α_i} = d_m$. Note that atoms in one of the (intermediate) Rydberg
states \(|m\rangle\) are assumed to be neither trapped, \(f^{(i)}_{\alpha\beta} = 0\), nor laser cooled, \(\beta = 0\).

Optical pumping from \(|\tilde{g}\rangle\) to one of the intermediate states \(|m\rangle\) and the following cascaded decay, \(|\tilde{g}\rangle \rightarrow |m\rangle \rightarrow \ldots \rightarrow |m'\rangle \rightarrow |\tilde{g}\rangle\), back to the dressed ground state will redistribute the atomic populations according to the rate equations \(40\). We can simulate this many-body dynamics by starting with a given atomic configuration, and propagating Eqs. \(41\) and \(44\) for a time step \(\Delta t\), much smaller than the time scales corresponding to the above mentioned rates. We then determine the probabilities for atom \(i\) in state \(\alpha\), to make a transition (jump) to another internal state according to the rate equations \(40\) for optical pumping and decay, and pick a new configuration according to these probabilities. Each cascade of quantum jump is associated with momentum kicks \(\hbar k_m\), \(\ldots\), \(\hbar k_{m'\alpha}\) in random spatial directions according to the distribution \(N_{\alpha\beta}\) of Eq. \(8\) and with \(k_{\alpha\beta}\) the wavevector of the corresponding transition. This new configuration is then again propagated another time step according to equations Eqs. \(41\) and \(44\).

Due to population of intermediate states in the long-time limit and the resulting time-dependent dipole moments we expect a rapid heating caused by fluctuations of the interaction force described above. In addition we also expect atomic losses. For example, when a ground state atom collides with an atom in an intermediate \(|\alpha\rangle\)-state with a large dipole moment the kinetic energy released in the collision can (i) be of the order of the height of the saddle point (see Sec. \(\text{IV C}\)). In this case the atoms will overcome the potential barrier and experience an attractive interaction. Or, (ii) the energy released can be of the order of the depth of external confinement in the transversal direction. A third loss mechanism is be Fenning ionization at distances smaller than \(4\pi a_0\). Each of these processes leads to two-body or one-body losses. In the simulations, we utilize a model including two-particle losses, where two particles are lost whenever their relative distance \(r\) becomes smaller than a certain critical radius \(r_{\text{loss}}\) (see Sec. \(\text{IV C}\)).

C. Effects of decoherence on the dynamics of many interacting Rydberg atoms

In this section we numerically investigate the effects of spontaneous emission and blackbody radiation on the long-time dynamics of an ensemble of interacting Rydberg atoms by performing semi-classical molecular dynamics simulations. As discussed above population of intermediate states \(|m\rangle\) with different interaction properties, e.g. different dipole moments, will lead to strong fluctuations of the interparticle forces in the many-body case.

Using semi-classical molecular dynamics simulations we study the resulting non-equilibrium dynamics of an ensemble constitution of \(N = 67\) Rydberg-dressed \(^85\text{Rb}\) atoms. Initially they are prepared in a perfect triangular crystal in a box with hard walls at \(T = 0\). For both dressing schemes we choose system parameters such that the ground state dipole moment is \(d_g = 30\) Debye, corresponding to a melting temperature of \(T_M = 0.6\ \mu\text{K}\) (\(2\pi \times 12\ \text{kHz}\)) for \(n_{2D} = 1\ \mu\text{m}^{-2}\). Additionally, the effect of laser-cooling and a triangular in-plane optical lattice is analyzed in order to stabilize the atomic crystal. Note that, we assume only dressed ground state atoms experience both the cooling laser and the lattice with a depth of \(50\ E_R\), with \(E_R = 2\pi \times 3.8\ \text{kHz}\), while atoms in one of the intermediate states \(|m\rangle\) do not, see Eqs. \(41\) and \(44\).

1. DC-electric field dressing

Fig.\(14\) analyzes heating in a gas of Rydberg dressed atoms using the DC-electric field scheme of Fig.\(4\) (b) with the same parameters as in Sec. \(\text{II B}\). For \(\Delta r = 2\pi \times 250\ \text{MHz}\) the Condon point is at \(r_c \sim 520\ \text{nm}\). For a transversal trapping frequency of \(\omega_{\perp} = 2\pi \times 200\ \text{kHz}\) the height of the potential barrier at the saddle point is \(\sim 150\ \mu\text{K}\). Each marker in Fig.\(14\) is an average over 50
runs of the simulation, where we simulated the dynamics for a time $\tau_3$. 

Fig. 15(a) shows the heating rate, $\gamma_3 E_{\text{kin}}(t = \tau_3)$ as a function of the atomic density for different cooling and lattice parameter, with $E_{\text{kin}}(t)$ the mean kinetic energy of all $N$ atoms. In the case of no laser cooling, $\beta = 0$, there is a strong dependence of the heating rate on the density, while for a cooling rate of $\beta = 2\pi \times 30$ kHz the heating rate is almost zero and a steady state is realized on a timescale $\beta^{-1} \ll \tau_3$. This steady state is due to the interplay of laser cooling and heating together with loss of high energy particles from the 2D confinement. This is shown in Fig. 14(b), where the number of remaining particles after a time $\tau_3$ is shown as a function of the density. Again, there is a strong dependence on the density and also on the laser cooling rate $\beta$. For densities $n_{2D} \lesssim 0.04 \, \mu\text{m}^{-2}$ the effect of the fluctuating dipoles can be neglected on surrounding atoms, while for densities $n_{2D} \sim 1 \, \mu\text{m}^{-2}$ the particle number has been decreased by 3% (with cooling) or 15% (no cooling) after a time $\tau_3 = 120 \, \mu\text{s}$. Similar to Fig. 15(a) we observe an accelerated loss of particles, resulting in a dramatic decrease of the particle number for times $t > \tau_3$. The effect of an in-plane optical lattice is small and hardly changes the heating rate or particle number shown in Fig. 14.

2. AC-microwave field dressing

Fig. 15(a) shows heating dynamics in a gas of Rydberg dressed atoms using the microwave dressing scheme of Fig. 14(c) with the same parameters as in Sec. IV C. For $\Delta_r = 2\pi \times 1.0$ GHz and $\alpha = 0.5$ we find $r_c \sim 1.38 \, \mu\text{m}$. With $\omega_\perp = 2\pi \times 100$ kHz the height of the barrier at the saddle points is $\sim 70 \, \mu\text{K}$.

In our treatment, the basic heating process of a crystal comes from its spatial rearrangement, once one of the atoms decays to an intermediate state $|m\rangle$ with zero dipole moment. We explain this as follows: Due to the lack of dipole moment, particles in the intermediate state $|m\rangle$ can travel a distance $v_R/\Gamma_m$ without interacting with neighboring atoms, where $v_R$ is the recoil velocity and $\Gamma_m$ the mean lifetime of an intermediate state. (i) if $v_R/\Gamma_m$ is of the order of the mean particle distance in the crystal $n_{2D}^{-1/2}$ two particles will most probably come closer than the critical radius $r_{\text{loss}}$ and get lost. We find that this latter process can result in an unusual collisional evaporative cooling effect. (ii) if $v_R/\Gamma_m$ is smaller than the mean particle distance in the crystal the atom may return to the ground state with a finite dipole moment, resulting in large time-dependent fluctuations of the dipole-dipole interaction between neighboring atoms. The net heating rate of the crystal is a competition between processes (i) and (ii).

We note that accidental vdW interactions may occur between atoms in one of the intermediate states $|m\rangle$, which may lead to additional heating. However, these processes are not considered here.

Fig. 15(a) shows the mean heating rate, $\gamma_3 E_{\text{kin}}(t = \tau_3)$ as a function of the atomic density for different cooling and lattice parameter, with $E_{\text{kin}}(t)$ the mean kinetic energy of all $N$ atoms. Again, each marker in Fig. 15(a) is an average over 50 runs of the simulation, where we simulated the dynamics for a time $\tau_3 = 12.8 \, \mu\text{s}$. Similar as in the DC-electric field case of Fig. 14 there is a strong dependence of the heating rate on the density and on the laser cooling rate $\beta$, while the effect of an in-plane optical lattice is weak. Additional laser cooling of the atoms with a rate $\beta = 30 \, \text{kHz}$ again leads to a steady state of the mean kinetic energy (see panel (a)) and diminishes the particle loss rate, which is shown in Fig. 14(b). The number of remaining particles after a time $\tau_3$ is shown as a function of the atomic density. Remarkably, more than 70% (50%) of the particles are left with (without) laser cooling and densities of $n_{2D} = 0.2/\mu\text{m}^2$ after $t = \tau_3$. 

![Graph showing heating rate and particle number as functions of density](image-url)
VII. CONCLUSIONS AND OUTLOOK

In this work we have investigated the long-time nonequilibrium dynamics of an ensemble of cold ground state atoms, weakly admixed with a Rydberg state using laser light. For times comparable to or larger than the effective lifetime of the ground state, the population of intermediate Rydberg states following spontaneous emission significantly affects the atomic motion in a strongly interacting gas, by providing a dominant heating and loss mechanism. Numerical simulations indicate that, due to the absence of dipole moments in the intermediate Rydberg states, ensembles of atoms polarized by AC-microwave fields exhibit a significantly different long-time dynamics, compared to atoms dressed by DC-electric fields.

Understanding this long-time dynamics is relevant for the experimental demonstration of laser-dressing techniques [32] as well as the creation of long-lived strongly correlated atomic phases such as self-assembled atomic crystals. Related to recent work on mixtures of polar molecules and Rydberg atoms [62, 63], we speculate that long-lived cold crystals may be used as cold reservoirs for achieving, e.g., sympathetic cooling of polar molecules [67].

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