Dynamics of ultracold dipolar particles in a confined geometry and tilted fields

Gouven Quéméner, Maxence Lepers, Olivier Dulieu
Laboratoire Aimé Cotton, CNRS, Université Paris-Sud, ENS Cachan, 91405 Orsay, France
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We develop a collisional formalism adapted for the dynamics of ultracold dipolar particles in a confined geometry and in fields tilted relative to the confinement axis. Using tesseral harmonics instead of the usual spherical harmonics to expand the scattering wavefunction, we recover a good quantum number \( \xi = \pm 1 \) which is conserved during the collision. We derive the general expression of the dipole-dipole interaction in this convenient basis set as a function of the polar and azimuthal angles of the fields. We apply the formalism to the collision of fermionic and bosonic polar KRb molecules in a tilted electric field and in a one-dimensional optical lattice. The presence of a tilted field drastically changes the magnitude of the reactive and inelastic rates as well as the inelastic threshold properties at vanishing collision energies. Setting an appropriate strength of the confinement for the fermionic system, we show that the ultracold particles can even further reduce their kinetic energy by inelastic excitation to higher states of the confinement trap.

I. INTRODUCTION

The field of ultracold gases composed of dipolar particles has generated tremendous interest during the past years \([1-6]\). One major goal is to shape at will the quantum properties of ultracold gases using the high degree of controllability available in experiments. Different kinds of dipolar particles are concerned by the quest for manifestation of dipole-induced effects. The first category of interest contains particles with electric dipoles such as ground state molecules of KRb \([7,8]\), RbCs \([9,10]\), NaK \([11]\) and many others experimentally under way... The second category includes particles with magnetic dipoles such as Cr \([12,14]\), Dy \([15,16]\), Er \([17,18]\) atoms and Er\(_2\) molecules \([19]\)... The third category consists of particles with both electric and magnetic dipoles such as molecules of OH \([20]\), SrF \([21]\), YO \([22,23]\), RbSr \([24]\)... All these particles can be manipulated by different configurations of electric and/or magnetic fields. They can also be loaded in optical lattices of different dimensions such as one dimensional (1D) lattices \([25,26]\), two dimensional (2D) lattices \([30,31]\) and three dimensional (3D) lattices \([32]\). Due to the wide and numerous domains of application of dipolar particles \([33]\), it is therefore important to understand how to control the interactions and the collisional properties of these particles under such configurations of fields and lattices. For example, it has been shown that chemical reactivity of molecules can be suppressed by an electric field in a confined 1D optical lattice \([26]\) or by selecting a particular electric field and appropriate quantum states of the molecules in a non-confined space \([34]\).

In this study, we investigate two-body collisions in a confining 1D optical lattice in electric and/or magnetic fields tilted with respect to the \(z\) and \(x\) axis, as illustrated in Fig. 1. We consider here “classical” dipoles aligned along the field, for which the angular internal structure of the particles (rotational angular momentum for the electric dipoles and electronic angular momentum for the magnetic dipoles) is not taken into account. We choose an effective value \(d\) or \(\mu\) of the electric or magnetic dipole moment, corresponding to their expectation value along the direction of the electric or magnetic field. For particles without angular internal structure, the total angular momentum \(J\) of the two-body colliding system reduces to the orbital angular momentum \(L\) associated with the quantum number \(l\), which is conserved in free space. When a field is applied parallel to the quantization axis \(z\), it is not conserved anymore but its projection \(L_z\) associated with the quantum number \(m_l\) still is. When the field is tilted and no more parallel with respect to the quantization axis, its projection is not conserved anymore. The scattering problem becomes challenging as \(l, m_l\) are all mixed. We show in this study that it is still possible to define a good quantum number, provided that the collision is described using the so-called “tesseral harmonics” \([35]\) instead of the standard spherical harmonics for the partial wave expansion of the scattering wavefunction. The problem is thus split in two sub-problems of smaller size. As an example of a dipolar system, we study...
fermionic and bosonic KRb + KRb collisions in a tilted electric field. The present theoretical formalism was also successfully used recently to understand the experimental observation of dipolar collisions of bosonic Feshbach Er$_2$ molecules in a 1D optical lattice in a tilted magnetic field [10].

The paper is structured as follows. In Section II, we develop the theoretical formalism for collision of particles in a tilted field and confined geometry where the appropriate basis set for partial wave expansion is introduced. In Section III, we show how the collisional rate coefficients and their threshold behaviours are strongly affected by tilted fields revealing the complexity of the mechanism. The role of the confinement trap is also explored and could be used to reduce the kinetic energy of the particles. Finally, we conclude in Section IV.

II. THEORETICAL FORMALISM

Ultracold dipolar collisions in a tilted field have already been studied in the past including microwave fields [13, 38] but without confinement, in crossed electric and magnetic fields [39, 40], or considering strong 1D confinement such that the particles are bound to collide in pure 2D [11]. By pure 2D we mean that the characteristic strength of the particles confinement is much stronger than the characteristic strength of their interaction (the dipole-dipole interaction here). However, this regime of pure 2D collisions is not yet reached in ongoing experiments as the required confinement strength is too strong. Instead, quasi-2D collisions occur. The particles start to collide at large distances in pure 2D but there is a point as they approach each other where the increasing magnitude of their interaction gets much bigger than the confinement strength. The particles do not feel anymore the presence of a 2D confinement and collide as if they were in a non-confined space. Therefore to reproduce the conditions of ongoing experiments, we describe the quasi-2D collisions of two ultracold particles of mass $m_1$, $m_2$ carrying tilted dipole moments and trapped in a 1D optical lattice of arbitrary confinement strength. We assume that the particles cannot hop from one potential well to another so that we approximate a well by a harmonic oscillator for particle 1 and 2, $V_{ho} = \frac{1}{2} n_1 \omega^2 z_1^2 + \frac{1}{2} m_2 \omega^2 z_2^2$. The angular frequency $\omega = 2\pi \nu$, $\nu$ governs the strength of the confinement. The particles are initially in a given state of the harmonic oscillator $n_1, n_2$ of energy $\varepsilon_{n_1} = h\nu(n_1 + 1/2), \varepsilon_{n_2} = h\nu(n_2 + 1/2)$.

The electric or magnetic fields make an angle $\theta_{E,B}$ and $\varphi_{E,B}$ with the $z$ and $x$ axis respectively as depicted in Fig. 2.a. The classical dipole approximation has been shown convenient for modelling electric dipoles [34] and magnetic dipoles [12] interactions in ultracold dipolar gases. It is an appropriate way to avoid the inclusion of the particles internal structure in the collisional formalism, thus sparing computational effort. No Stark nor Zeeman term appears in our formalism then.

We distinguish three types of collisional processes: (i) Elastic processes for which the molecules remain in the same external state of the harmonic oscillator after the collision: $(n_1', n_2') = (n_1, n_2)$; (ii) Inelastic processes for which the molecules change their external state of the harmonic oscillator $(n_1', n_2') \neq (n_1, n_2)$ (note that a change of the internal states is not possible as the internal structure of the particles is not treated); (iii) Loss processes due for instance to chemical reactions occurring at ultralow energy, leading to products with high kinetic energy which are expelled from the trap.

A. Quasi-2D collisions in parallel field

We shall briefly recall the formalism for fields parallel to the quantization axis when $\theta_{E,B} = 0$. It is presented in more details in Ref. [28]. First, it is convenient to transform the motion of the individual particles 1 and 2 with position coordinates $\vec{r}_1, \vec{r}_2$ of masses $m_1, m_2$ in external states $n_1, n_2$, into the motion of two effective particles described by the center-of-mass and relative coordinates $\vec{R}, \vec{r}$ of masses $m_{tot}, m_{red}$ in external states $N, n$. For confinements modelled by harmonic oscillators, it can be shown then that the center-of-mass coordinate $\vec{R}$ is decoupled from the relative coordinate $\vec{r}$.

The potential energy term for the relative coordinate $\vec{r}$ is given by a van der Waals interaction

$$V_{vdW} = -C_6/r^6,$$

the confinement interaction given by a harmonic oscillator for the relative motion along $z$

$$V_{ho} = \frac{1}{2} m_{red} \omega^2 z^2,$$
and a dipole-dipole interaction composed of an electric and magnetic term

\[ V_{dd} = V_{dd}^e + V_{dd}^m \]

(3)

where

\[ V_{dd}^{e,m} = -X_{e,m}^2(1 - 3\cos^2\theta)/r^3 \]

(4)

with for electric dipoles

\[ X_e \equiv d/\sqrt{4\pi\varepsilon_0} \]

(5)

and for magnetic dipoles

\[ X_m \equiv \mu/\sqrt{(4\pi/\mu_0)}. \]

(6)

The coordinate \( \vec{r} \) is expressed in spherical coordinates \( \vec{r} = (r, \theta, \varphi) \) when \( V_{vdW} \) and \( V_{dd}^{e,m} \) are dominant (Fig. 2b), or in cylindrical coordinates \( \vec{r}' = (\rho, z, \varphi) \) when \( V_{ho} \) is dominant at very large distances.

We decompose the total wavefunction \( \psi(\vec{r}) \) of the relative motion into a basis set of spherical harmonics \( Y_l^{m_l}(\theta, \varphi) \equiv \langle \vec{r}|l, m_l \rangle \)

\begin{equation}
\psi(r, \theta, \varphi) = \frac{1}{r} \sum_{l,m_l} f_{l,m_l}(r) Y_l^{m_l}(\theta, \varphi)
\end{equation}

(7)

with \(-l \leq m_l \leq +l\). In this basis set, the expression of the potential energy terms become in bra-ket notation

\[ \langle l m_l|V_{vdW}|l'|m_l' \rangle = -\frac{C_6}{r^6} \delta_{m_l,m_l'} \delta_{l,l'} \]

(8)

\[ \langle l m_l|V_{ho}|l'|m_l' \rangle = \delta_{m_l,m_l'} \frac{1}{r^2} r^2 \}
\frac{1}{3} \left\{ (1)^{m_l} \sqrt{2l + 1} \sqrt{2l' + 1} \right\} \]
\times 2 \left( \begin{array}{ccc} l & 2 & l' \\ 0 & 0 & 0 \end{array} \right) \left( \begin{array}{ccc} l & 2 & l' \\ m_l & 0 & m_l' \end{array} \right) + \delta_{l,l'} \}
\end{eqnarray}

(9)

\[ \langle l m_l|V_{dd}^{e,m}|l'|m_l' \rangle = -\frac{\sqrt{30} X_{e,m}^2}{r^3} \]
\frac{1}{3} \left\{ (1)^{m_l} \sqrt{2l + 1} \sqrt{2l' + 1} \right\} \]
\times \left( \begin{array}{ccc} 1 & 1 & 2 \\ 0 & 0 & 0 \end{array} \right) \left( \begin{array}{ccc} l & 2 & l' \\ 0 & 0 & 0 \end{array} \right) \left( \begin{array}{ccc} l & 2 & l' \\ -m_l & 0 & m_l' \end{array} \right) \}
\end{eqnarray}

(10)

The first expression is diagonal in \( m_l \) and \( l \) while the last two expressions are diagonal in \( m_l \) but couple different values of \( l \). The expression for \( V_{ho} \) arises from the fact that \( s^2 = r^2 \cos^2 \theta = r^2 [4 \sqrt{\pi/5} Y_2^2 + 1]/3 \).

The time-independent Schrödinger equation is solved for a fixed total energy \( E_{tot} = E_{n_1} + E_{n_2} + E_{coll} \). Equations 8 to 10 lead to a set of coupled differential equations. To solve this system of equations, we use a diabatic-by-sector method \([28]\) which generates a set of adiabatic energy curves as a function of the inter-particle distance \( r \), and we propagate the log-derivative of the wavefunction \([42, 43]\). The boundary condition at short distance, where the propagation of the wavefunction is started, is set up by a tunable, diagonal log-derivative matrix given in Ref. \([34]\) for which we can control the amount of loss in this short-range region. For the boundary condition at large distance in the asymptotic region where the propagation of the wavefunction is ended, we use the asymptotic form of the cylindrical wavefunction which is a linear combination of regular and irregular Bessel functions of the cylindrical problem. Using a transformation matrix from cylindrical to spherical coordinates \([28]\), we obtain the asymptotic form of the wavefunction from which we deduce the \( K \), \( S \), and \( T \) matrices using the expression of the log-derivative matrix in spherical coordinates. The \( S \) matrix in the center-of-mass and relative coordinates is then expressed back into the individual coordinates \([28]\) yielding the elastic, inelastic and loss cross sections and rate coefficients for two particles starting in a given initial state \((n_1, n_2)\).

\section*{B. Quasi-2D collisions in an arbitrary field}

\textit{In the spherical harmonics basis set}

As depicted in Fig. 2 when the fields are tilted, the expression of the dipole-dipole interactions become

\[ V_{dd}^{e,m} = -X_{e,m}^2(1 - 3\cos^2|\theta - \theta_{E,B}|)/r^3. \]

(11)

This implies a more complicated expression in the spherical harmonic basis set

\[ \langle l m_l|V_{dd}^{e,m}|l'|m_l' \rangle = -\frac{\sqrt{30} X_{e,m}^2}{r^3} \]
\times \left( \frac{1}{3} \left\{ (1)^{m_l} \sqrt{2l + 1} \sqrt{2l' + 1} \right\} \]
\times \left( \begin{array}{ccc} l & 2 & l' \\ 0 & 0 & 0 \end{array} \right) \left( \begin{array}{ccc} l & 2 & l' \\ -m_l & 0 & m_l' \end{array} \right) \}
\end{eqnarray}

(12)

For the special case \( \theta_{E,B} = 0 \), \( p_1 = p_2 = p = 0 \) and we recover the case \( m_l = m_l' = 0 \). For the special case \( \theta_{E,B} = \pi/2, p_1 = \pm 1, p_2 = \pm 1, p = 0, \pm 2 \) then we have \( m_l = m_l' \pm 0, \pm 2 \). For any other \( \theta_{E,B} \) we have \( m_l - m_l' = 0, \pm 1, \pm 2 \), increasing the number of coupled equations and leaving no good quantum numbers in Eq. 12.

\textit{In the tesseral harmonics basis set}

By properly symmetrizing the basis set of spherical harmonics, we can still recover a good quantum number.
We use the following symmetrized spherical harmonics in ket notation

\[
|l, \bar{m}_l, \xi\rangle = \frac{\delta_{l1} + l \delta_{l-1}}{\sqrt{2\Delta}} \times \left\{|l, -|m_l|\rangle + \xi (-1)^{|m_l|} |l, |m_l|\rangle\right\}
\]

(13)

where \(\bar{m}_l \equiv |m_l|\) and \(\Delta \equiv 1 + \delta_{m_l,0}\). The new quantum number \(\xi\) takes the values \(\xi = \pm 1\), while \(\bar{m}_l = 0, 1, 2, 3, ...\) when \(\xi = 1\) and \(\bar{m}_l = 1, 2, 3, ...\) when \(\xi = -1\). This new basis set are often called the tesserar harmonics 33. We note them \(Y_{l,\bar{m}_l,\xi}\) with

\[
\begin{align*}
Y_{l,\bar{m}_l,0,\xi=+1} & = \frac{1}{\sqrt{2}} \left\{ Y_{l,2\bar{m}_l} + (-1)^\bar{m}_l Y_{l,0\bar{m}_l} \right\} \\
& \propto P_{l\bar{m}_l}^0(\cos\theta) \cos \bar{m}_l \varphi \\
Y_{l,\bar{m}_l,0,\xi=+1} & = Y_{l,2\bar{m}_l} \propto P_{l\bar{m}_l}^0(\cos\theta) \\
Y_{l,\bar{m}_l,0,\xi=+1} & = \frac{1}{\sqrt{2}} \left\{ Y_{l,2\bar{m}_l} - (-1)^\bar{m}_l Y_{l,0\bar{m}_l} \right\} \\
& \propto P_{l\bar{m}_l}^0(\cos\theta) \sin \bar{m}_l \varphi .
\end{align*}
\]

(14)

In this new basis set, the potential energy matrix elements are given by

\[
\langle l, \bar{m}_l, \xi | V_{vdW} | l', \bar{m}_l', \xi'\rangle = -\frac{C_6}{r^6} \times \delta_{l,l'} \delta_{\bar{m}_l,\bar{m}_l'} \bar{c}_{l,l'} .
\]

(15)

\[
\langle l, \bar{m}_l, \xi | V_{ho} | l', \bar{m}_l', \xi'\rangle = \frac{1}{\sqrt{\Delta \bar{\Delta} \bar{\Delta}}} \delta_{\xi,\xi'} \delta_{\bar{m}_l,\bar{m}_l'}
\]

\[
\frac{1}{3} \bar{m}_{\text{red}} \omega^2 \frac{r^2}{2} \left\{ (-1)^{\bar{m}_l} \sqrt{2l+1} \sqrt{2l'+1} \right\}
\]

\[
\times \left[ 2 \left( \begin{array}{c} l \ 2 \ l' \\ 0 \ 0 \ 0 \end{array} \right) \left[ \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ 0 \ -\bar{m}_l' \end{array} \right] \delta_{\bar{m}_l,0} \delta_{\xi,1} \\
+ \bar{c}_{l,l'} + \delta_{l,l'} \xi' (-1)^{\bar{m}_l} \delta_{\bar{m}_l,0} \delta_{\xi,1} \right\},
\]

(16)

and after reducing Eq. 12

\[
\langle l, \bar{m}_l, \xi | V_{vdw} | l', \bar{m}_l', \xi'\rangle = -\frac{\sqrt{30} X_{vdw}}{r^6} \frac{1}{\sqrt{\Delta \bar{\Delta} \bar{\Delta}} \bar{\Delta}} (-1)^{\bar{m}_l} \sqrt{2l+1} \sqrt{2l'+1} \left( \begin{array}{c} l \ 2 \ l' \\ 0 \ 0 \ 0 \end{array} \right)
\]

\[
\left\{ (c_0)^2 \left( \begin{array}{c} 1 \ 1 \ 2 \\ 1 \ 1 \ -2 \end{array} \right) (\delta_{\xi,\xi'} + i \xi \delta_{\xi,\xi'-1}) \left( e^{-i2\bar{c}_{l,l'}} + \xi \xi' e^{i2\bar{c}_{l,l'}} \right) \right\}
\]

\[
\times \left[ \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ 0 \ -\bar{m}_l' \end{array} \right) + \xi' (-1)^{\bar{m}_l} \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ -2 \ -\bar{m}_l' \end{array} \right) \right]
\]

\[
+ 2 \bar{c}_{l,l'} \left( \begin{array}{c} 1 \ 1 \ 2 \\ 0 \ 1 \ -1 \end{array} \right) (\delta_{\xi,\xi'} + i \xi \delta_{\xi,\xi'-1}) \left( e^{-i2\bar{c}_{l,l'}} + \xi \xi' e^{i2\bar{c}_{l,l'}} \right)
\]

\[
\left[ \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ 1 \ -\bar{m}_l' \end{array} \right) - \xi' \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ 0 \ -\bar{m}_l' \end{array} \right) \right] + \xi' (-1)^{\bar{m}_l} \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ -2 \ -\bar{m}_l' \end{array} \right)
\]

\[
+ \delta_{\xi,\xi'} \delta_{\bar{m}_l,\bar{m}_l'} \left\{ (c_0)^2 \left( \begin{array}{c} 1 \ 1 \ 2 \\ 0 \ 0 \ 0 \end{array} \right) - 2 (c_1)^2 \left( \begin{array}{c} 1 \ 1 \ 2 \\ 1 \ 1 \ 0 \end{array} \right) \right\} \left( \begin{array}{c} l \ 2 \ l' \\ -\bar{m}_l \ 0 \ -\bar{m}_l' \end{array} \right) \delta_{\bar{m}_l,0} \delta_{\xi,1} \right\},
\]

(17)
with

\[ c_{p,0,1} = \frac{\sqrt{4\pi}}{3} Y^{p}_{1} \left( \frac{\theta_E}{B}, \varphi_E, B = 0 \right), \]

namely \( c_{0} = \cos \theta_E, B \) and \( c_{1} = \sin \theta_E, B / \sqrt{2} \). Equation (15) is diagonal in \( \xi, \bar{m}_{l}, l \) and Eq. (16) is diagonal in \( \xi, \bar{m}_{l} \). Equation (17) is diagonal in \( \xi \) if \( \varphi_E = 0 \) and/or \( \varphi_B = 0 \), and it reduces in this case to

\[
\langle l, \bar{m}_{l}, \xi | V_{dd}^{e,m} | l', \bar{m}_{l}', \xi' \rangle = \frac{-\sqrt{30} X_{e,m}}{r^3} \frac{1}{\Delta \Delta_{l}} (-1)^{\bar{m}_{l}} \sqrt{2l+1} \sqrt{2l'+1} \left[ \begin{array}{c} 1 \bar{m}_{l} \bar{m}_{l}' \end{array} \right] \left[ \begin{array}{c} l \bar{m}_{l} \bar{m}_{l}' \end{array} \right] \delta_{\xi,\xi'}
\]

\[
\left\{ (c_{1})^{2} \left[ \begin{array}{c} 1 \ 1 \ 2 \\ 1 \ 1 \ -2 \end{array} \right] \left[ \begin{array}{c} \bar{m}_{l} \ 2 \ l' \\ -2 \ -\bar{m}_{l}' \end{array} \right] + \xi \xi' \left[ \begin{array}{c} 1 \ 2 \ l' \\ -2 \ -\bar{m}_{l}' \end{array} \right] + \xi' (-1)^{\bar{m}_{l}} \left[ \begin{array}{c} 1 \ 2 \ l' \\ -2 \ -\bar{m}_{l}' \end{array} \right] \right\} \times \left[ \begin{array}{c} (l \ 2 \ l' \\ -2 \ -\bar{m}_{l}' \end{array} \right] \left[ \begin{array}{c} 1 \ 2 \ l' \\ -2 \ -\bar{m}_{l}' \end{array} \right] \delta_{\bar{m}_{l},0} \delta_{\xi,1} \right\}. \quad (19)
\]

We can see that we recover a good quantum number \( \xi \) in this formalism which was not the case in the nonsymmetrized spherical harmonics basis set. When \( \varphi_E = \varphi_B \neq 0 \), or when a unique electric or magnetic field is used where \( \varphi_E \neq 0 \) or \( \varphi_B \neq 0 \), one can always recover the situation in Eq. (19) with good quantum numbers by a proper rotation of the \( x \) and \( y \) axis due to the cylindrical symmetry of the pancakes. Only in the more general case including both electric and magnetic fields where \( \varphi_E \neq \varphi_B \), one of the dipole-dipole expression is no more diagonal in \( \xi \) and Eq. (17) has to be used instead. To simplify the study in the following, we will consider the case where \( \varphi_E, B = 0 \) when the dipoles are only tilted in the \( xOz \) plane. Furthermore we will consider the case where we only apply a unique field (electric in this study).

In general, the quantum number \( \bar{m}_{l} = 0 \) is always automatically associated with the \( \xi = +1 \) one as it is not defined for the \( \xi = -1 \) one. For the \( \bar{m}_{l} > 0 \) values one has to use both \( \xi = \pm 1 \) values. For the special case of parallel field \( \theta_E, B = 0, \bar{m}_{l} \) is a good quantum number, \( c_{1} = 0 \) and Eq. (17) reduces to Eq. (10). Still for the parallel case for \( \bar{m}_{l} \neq 0 \), the \( \xi = -1 \) contribution is identical to the \( \xi = +1 \) one so that the total contribution is twice the \( \xi = +1 \) one. For \( \bar{m}_{l} = 0 \), only the \( \xi = +1 \) contribution is needed. Therefore, only the value of \( \xi = +1 \) is needed in the parallel field case for all \( \bar{m}_{l} \). The contribution of the different quantum numbers needed for different tilted configurations are summarized in Table 1.

The suitability of the tesseral harmonics can be qualitatively understood from their spatial shape. If \( \varphi = 0 \) or \( \pi \) which corresponds to the \( xOz \) plane (see Fig 2), \( Y_{l,m_{l},\xi = -1} \) vanishes there for any values of \( m_{l} \) since it is proportional to \( \sin m_{l} \varphi \) (see Eq. 13). We can therefore associate the \( \xi = -1 \) quantum number with an out-of-plane motion, excluding the collision in the \( xOz \) plane. If \( \varphi_E, B = 0, \) the electric or magnetic field is applied in the \( xOz \) plane only. As the dipoles are exclusively pointing in the \( xOz \) plane, the collisions of dipoles in the \( y \) direction start always side-by-side at long-range since there is no component of the dipole moment in the \( y \) direction. We thus expect that the manifold of the \( \xi = -1 \) curves is always more repulsive than its \( \xi = +1 \) counterpart due to the side-by-side, repulsive dipolar approach.

The tesseral harmonics representation of the partial waves is therefore a more appropriate basis set in the general case of collisions in arbitrary tilted electric and/or magnetic fields. Note that the present formalism is also adapted for collisions of particles in free space in crossed electric and magnetic fields, where one field (for example the magnetic field) is chosen as the quantization axis and the other (for example the electric field) is the tilted one. For example, the study done in Ref. 40 using spherical harmonics could be adapted using tesseral harmonics.

### III. RESULTS

In the following, we study the collision of two indistinguishable, electric dipolar molecules of \( \text{KRb} \) in their absolute internal ground state for which the “classical dipoles” assumption is a good description. We use \( ^{40}\text{K}^{87}\text{Rb} \) for a fermionic system example and \( ^{41}\text{K}^{87}\text{Rb} \) for a bosonic one. We impose here \( \varphi_E = 0 \). They possess a permanent electric dipole moment of \( d_{p} \) for a bosonic field, \( \varphi_B = 0 \). The electric field can therefore induce a dipole moment \( d_{p} \) for a bosonic field, \( \varphi_B = 0 \). The results presented here will also be similar for collisions of magnetic dipolar particles with strong magnetic dipole moment, tuned by a tilted magnetic field \( B \), as performed in Ref. 19. The confinement in the \( z \) direction is described by a harmonic oscillator of
FIG. 3: (Color online) Adiabatic energy curves as a function of the intermolecular distance $r$ between two fermionic $^{40}\text{K}^{87}\text{Rb}$ molecules (left panels) and two bosonic $^{41}\text{K}^{87}\text{Rb}$ molecules (right panels) for a given induced dipole $d = 0.2$ D and tilt angle $\theta_E$. The curves of same color are coupled among each other. Only the curves corresponding to $\xi = +1$ are shown. For the $\xi = -1$ curves, see Appendix A. Up and down arrows indicate the variation trend of the curves when $d$ increases. Each adiabatic energy curve correlates to a state $n$ of the harmonic oscillator indicated at the right of each panel.

frequency $\nu = 20$ kHz which is a typical value employed in experiments. We assume that all the particles are in the ground state of the harmonic oscillator $n_1 = n_2 = 0$ before the collision, which is equivalent to the relative $n = 0$ quantum number [28]. We study those collisions by varying different parameters such as the collision energy, the tilted angle and the confinement frequency. The loss processes are described by a full loss condition at short range given in Ref. [34]. The full loss condition corresponds to either a chemical reaction with full probability
A. Interactions and adiabatic energy curves

The adiabatic energy curves are shown in Fig. 3 for the fermionic and the bosonic KRb molecules, for an induced dipole moment of $d = 0.2$ D. If there is a presence of a barrier relative to a given collision energy as $r$ decreases and if this barrier increases when $d$ increases (indicated by an arrow pointing upward in Fig. 3), we say that the corresponding curve is protective against possible short-range loss. In the absence of a barrier or if the barrier decreases (indicated by an arrow pointing downward), we say that the corresponding curve is non-protective. The adiabatic energy curves are presented only for the quantum number $\xi = +1$. For the $\xi = -1$ quantum number, the curves are equal to their $\xi = +1$ counterpart in the vanishing dipole moment limit, recovering the isotropic character of the long-range van der Waals interaction. For larger $d$, they are more repulsive due to the fact that the $\xi = -1$ manifold corresponds to side-by-side dipolar repulsive collisions as mentioned earlier. The $\xi = -1$ curves are shown in Appendix A.

The fermionic case for $\theta_E = 0$ is shown in the top left panel of Fig. 3. The thick black solid lines (resp. thin blue solid lines, thick red dashed lines, thin green dashed lines) represent the (unmixed) values of $\tilde{m}_l = 1$ (resp. $\tilde{m}_l = 3$, $\tilde{m}_l = 0$, $\tilde{m}_l = 2$). The lowest curve of the $n = 0$ harmonic oscillator ground state correlates to the protective, repulsive side-by-side $\tilde{m}_l = 1$ curve. For indistinguishable fermionic particles, we recall that the scattering takes place for odd $l$ partial waves. The lowest curve $l = 1$ features a p-wave barrier. There is an actual crossing between the lowest curves with $\tilde{m}_l = 1$ and $\tilde{m}_l = 0$ since the $\tilde{m}_l$ components do not couple to each other. The middle panel shows the case for $\theta_E = \pi/4$. The black solid lines represent the curves with mixed values $\tilde{m}_l = 0, 1, 2, 3$. Since all $\tilde{m}_l$ are coupled, the actual crossing at $\theta_E = 0$ becomes an avoided crossing. And the lowest $n = 0$ curve correlates to a non-protective curve. Finally, the bottom panel shows the case for $\theta_E = \pi/2$. The black solid lines represent the curves with mixed values $\tilde{m}_l = 1, 3$ and the red dashed lines the curves with mixed values $\tilde{m}_l = 0, 2$. Both series of curves do not couple to each other. The lowest $n = 0$ curve still correlates to the non-protective curve. So for particles in the lowest state, we can see how the adiabatic energy curve changes from a protective character coming from of a repulsive side-by-side interaction approach when $\theta_E = 0$ to a non-protective one coming from of an attractive head-to-tail interaction approach when the dipoles are tilted by $\theta_E = \pi/2$.

We get similar results in Fig. 3 for the bosonic symmetry. For indistinguishable bosonic particles, the scattering takes place in even $l$ partial waves. The lowest $l = 0$ curve is barrierless, in contrast with the fermionic particles. Now the lowest $n = 0$ curve correlates to the barrierless, non-protective curve at short range for all $\theta_E$ even $\theta_E = 0$. As $\theta_E$ increases from 0 to $\pi/2$, the $\tilde{m}_l = 2$ curve pushes slightly downwards the $\tilde{m}_l = 0$ curve so that the lowest $n = 0$ curve gets slightly more attractive.

The behaviour of these adiabatic energy curves have direct consequences on the behaviour of the collisional rate coefficients. This is presented below.

B. Collisions and rate coefficients

1. Rate coefficients versus the collision energy. Threshold laws

We show in Fig. 4 the rate coefficients for the fermionic and the bosonic case as a function of the collision energy for $d = 0.2$ D, starting with two particles in $n = 0$ relative motion state. The top, middle and bottom panels corresponds to $\theta_E = 0, \pi/4, \pi/2$ respectively. The total rates are reported as a thick solid line for each processes. One can see on Fig. 3 that when the collision energy is increased from the $n = 0$ threshold, different harmonic oscillator states become energetically open, resulting in sudden peaks in the inelastic rate coefficients (blue curves). In addition to the usual elastic rate coefficient (red curves), we also have reactive rate coefficients (black curves) corresponding to the full loss condition at short range.

For the fermionic particles for the $\theta_E = 0$ case, we see that reactive rates are suppressed compared to elastic one at ultralow energies. This has already been explained theoretically and experimentally [21, 28, 29]. As the lowest $n = 0$ state curve correlates to a repulsive protective barrier curve when $r$ decreases, the probability of the two particles to be close to each other is low, preventing short-range loss to take place. The inelastic rate coefficient emerges at a collision energy of $E_c \sim 2 \mu K$ corresponding to the $n = 2$ threshold opening. The
FIG. 4: (Color online) Elastic (e), inelastic (i) and reactive (r) rate coefficients (thick lines) for fermionic (left panels) and bosonic (right panels) K\textsubscript{Rb} + K\textsubscript{Rb} collisions as a function of the collision energy at $d=0.2$ D and $\nu=20$ kHz. Top panels: $\theta_E=0$, the thin solid lines are for $\tilde{m}_l = 1$ and $\xi = \pm 1$ ($\tilde{m}_l = 0$ and $\xi = +1$), the thin dashed lines are for $\tilde{m}_l = 3$ and $\xi = \pm 1$ ($\tilde{m}_l = 2$ and $\xi = \pm 1$) for fermions (bosons). Middle panels: $\theta_E = \pi/4$, the thin solid lines are for $\xi = +1$, $\tilde{m}_l = 0, 1, 2, 3$, the thin dashed lines are for $\xi = -1$, $\tilde{m}_l = 1, 2, 3$. Bottom panels: $\theta_E = \pi/2$, the thin solid lines are for $\xi = +1$, $\tilde{m}_l = 1, 3$ ($\tilde{m}_l = 0, 2$), the thin dashed lines are for $\xi = -1$, $\tilde{m}_l = 1, 3$ ($\tilde{m}_l = 2$) for fermions (bosons).

$n=1$ threshold is not open here at $E_c \sim 1 \mu$K since the $\tilde{m}_l = 0$ and 2 (red and green curves) do not mix with the $\tilde{m}_l = 1$ and 3 (black and blue curves). When open above $E_c \sim 2 \mu$K, the inelastic rates are bigger than the reactive ones and can easily amount or overcome the value of the elastic rates. For $\theta_E = \pi/4$, the reactive rates are bigger than the elastic ones at ultralow energies. This is explained again with the corresponding adiabatic
energy curve where the \( n = 0 \) state correlates to the attractive non-protective barrier due to the coupling of the \( \tilde{m}_i = 0 \) curve with the \( \tilde{m}_i = 1 \) one and the presence of the avoided crossing. The inelastic rate shows up now at \( E_c \sim 1 \mu K \) since the \( n = 1 \) threshold is now allowed and coupled with the \( n = 0 \) one. For \( \theta_E = \pi/2 \), we recover the head-to-tail collision with large reactive rate coefficient. Due to the mixing of \( \tilde{m}_i = 1,3 \) only, the \( n = 1 \) threshold is not open here too and only the \( n = 2 \) opens up at \( E_c \sim 2 \mu K \).

For the bosonic particles, as the collision takes place on a barrierless \( l = 0 \) curve, we see that the magnitude of the rates is bigger than the corresponding one for the fermions. For \( \theta_E = 0 \), elastic rates are comparable with the reactive ones (either smaller or bigger depending on the collision energy) but there is no protection against collisions here. As \( \theta_E \) increases for \( \theta_E = \pi/4 \) and \( \pi/2 \), the reactive rate increases as the lowest adiabatic energy curve decreases and becomes the dominant rate coefficient. For the same reasons than those presented above for the fermions, the inelastic rates show up as the \( n = 2 \) threshold opens up at \( E_c \sim 2 \mu K \) for \( \theta_E = 0 \) and \( \theta_E = \pi/2 \). For \( \theta_E = \pi/4 \), the \( n = 1 \) threshold also opens up at \( E_c \sim 1 \mu K \).

For all \( \theta_E \neq 0 \) plots, the contribution of the \( \xi = -1 \) curves (thin dashed lines) is marginal for an induced dipole moment of \( d = 0.2 \) D. This is due to the large repulsive character of the corresponding adiabatic energy curves which correspond to side-by-side repulsive dipolar collision in the \( y \) direction, shown in Appendix A.

For the inelastic collisions, there are striking differences at the threshold opening of the inelastic state depending on the field angle. This can be explained by the different behaviour of the quantum threshold laws studied in Ref. [46]. In this reference the threshold laws were derived for inelastic relaxation for \( (k_i \rightarrow 0) \ll k_f \) where \( k_i \) and \( k_f \) represent the initial and final wavevector. Here we consider that the threshold laws for excitation processes are the same than the relaxation ones by replacing \( k_i \) by \( k_f \) and the initial \( m_i \) by the final \( m_f \) [47]. Then the inelastic relaxation laws given by Eq. (11) of Ref. [46], where \( k_i \ll k_f \), for one or both of \( m_i \) and \( m_f \) equal to zero,

\[
\beta_{\text{in. relax.}}^{m_i \rightarrow m_f} \propto k_i^{2m_i} \propto \frac{E_c^{m_i}}{\ln^2 k_i} \quad (20)
\]

should translate explicitly for the excitation processes to

\[
\beta_{\text{in. exc.}}^{m_i \rightarrow m_f} \propto k_f^{2m_f} \propto \frac{(E_c - E_{th})^{m_f}}{\ln^2 \sqrt{2m_{red}(E_c - E_{th})/\hbar^2}} \quad (21)
\]

where \( k_f \ll k_i \), according to the behavior of the different elements of Eq. 10 of Ref. [46]. \( E_{th} \) corresponds to the excitation threshold energy. For both \( m_i \) and \( m_f \) different than zero, the inelastic rate from Ref. [46]

\[
\beta_{\text{in. relax.}}^{m_i \rightarrow m_f} \propto k_i^{2m_i} \propto E_c^{m_i} \quad (22)
\]

should translate to

\[
\beta_{\text{in. exc.}}^{m_i \rightarrow m_f} \propto k_f^{2m_f} \propto (E_c - E_{th})^{m_f}. \quad (23)
\]

In our study, for the inelastic excitation rate \( n = 0 \rightarrow n = 2 \) at the \( E_c \sim 2 \mu K \) threshold for the specific cases \( \theta_E = 0 \) and \( \theta_E = \pi/2 \), we found a threshold law of

\[
\beta_{\text{in. exc.}}^{0 \rightarrow 2} \propto (E_c - E_{th}) \ln^2 \sqrt{2m_{red}(E_c - E_{th})/\hbar^2}
\]

for the bosons, in agreement with Eq. (23) and Eq. (21), where \( E_{th} = \epsilon_n = 2 \). At \( E_c \sim 1 \mu K \), the inelastic excitation rate \( n = 0 \rightarrow n = 1 \) for \( \theta_E = \pi/4 \) for the fermionic (resp. bosonic) case, gets the behaviour of the one for the bosonic (fermionic) case at \( \theta_E = 0 \) or \( \pi/2 \), with a sharper (rounder) shape. This is due to a change of parity in \( \tilde{m}_i \) in the inelastic transition. For the fermionic (bosonic) case, the inelastic transition \( n = 0 \rightarrow n = 1 \) corresponds to \( \tilde{m}_i = 1 \rightarrow \tilde{m}_i = 0 \) (\( \tilde{m}_i = 0 \rightarrow \tilde{m}_i = 1 \)). For the bosonic case the \( n = 0 \rightarrow n = 1 \) inelastic transition, we found a threshold law of

\[
\beta_{\text{in. exc.}}^{0 \rightarrow 1} \propto (E_c - E_{th}) \ln^2 \sqrt{2m_{red}(E_c - E_{th})/\hbar^2}
\]

(24) for the fermionic case, with a sharper (rounder) shape. This is in agreement with Eq. (21). Finally, for the elastic and reactive collisions, the quantum threshold laws are found to be \( \beta_{\text{cl}} \propto E_c^2 \) and \( \beta_{\text{cl}} \propto E_c \) for the fermions in agreement with Eq. (12) and (14) determined in Ref. [10] for dipolar collisions in quasi-2D. For the bosons we find \( \beta_3 \propto \beta_{\text{cl}} \propto \ln^2 \sqrt{2m_{red}E_c/\hbar^2} \) in agreement with Eq. (9) and (13) of the same reference.

2. Sensitivity of the rate coefficients versus the tilted field angle

The effect of the tilted field on the collision is directly seen in Fig. 5 for fermions (top panel) and bosons (bottom panel). This is plotted at a fixed collision energy of \( E_c = 1 \mu K \) and induced dipole of \( d = 0.2 \) D.

There is no fermionic and bosonic inelastic rate at \( \theta_E = 0 \) and \( \theta_E = \pi/2 \) at \( E_c = 1 \mu K \). Couplings between \( \tilde{m}_i = 1 \), \( \tilde{m}_i = 3 \) and \( \tilde{m}_i = 0 \); \( m_i = 2 \) curves are not allowed at these two angles so that the transition \( n = 0 \rightarrow n = 1 \) is forbidden there. Then the inelastic rates rise from \( \theta_E = 0 \) to \( \theta_E = \pi/4 \) due to the turning on of the couplings. At \( \theta_E = \pi/4 \) there is a maximal coupling between the \( \tilde{m}_i \) components: (i) a maximal coupling for \( \Delta \tilde{m}_i = 2 \) due to the presence of the \( \sin^2 \theta_E = 1/2 \) term in Eq. (19); (ii) a maximal coupling for \( \Delta \tilde{m}_i = 1 \) due to the presence of the \( \cos \theta_E \sin \theta_E = 1/2 \) term. Thus the inelastic rates reach their maximal value at \( \theta_E = \pi/4 \). Conversely, the couplings turn off as \( \theta_E \) passes from \( \pi/4 \) to \( \pi/2 \), and then the inelastic rates shut off.

The fermionic reactive rate increases as we pass from a side-by-side approach \( \theta_E = 0 \) to a head-to-tail one \( \theta_E = \pi/2 \). The continuous transition seen in the rate as \( \theta_E \) increases can be understood qualitatively by the continuous increase of the \( \Delta \tilde{m}_i = 1 \) coupling. At \( \theta_E = 0 \) the
lowest curve of symmetry $\tilde{m}_l = 1$ connects to the protective barrier and leads to a rate $\beta_0$, while at $\theta_E = \pi/2$ this curve connects to the non-protective barrier that leads to a rate $\beta_{\pi/2} \gg \beta_0$. Now, at $\theta_E = 0$ the lowest curve of symmetry $\tilde{m}_l = 0$ connects to the non-protective barrier, the same one that leads to the rate $\beta_{\pi/2}$. Therefore as the coupling between the two symmetries increases from $\theta_E = 0$ to $\pi/4$, the reactive rate is a combination of $\beta_0$ and $\beta_{\pi/2}$ with coefficients that decrease the contribution of $\beta_0$ and increase the one of $\beta_{\pi/2}$, hence increasing the total reactive rate. From $\theta_E = \pi/2$ to $\pi/4$ the reverse argument holds since now at $\theta_E = \pi/2$ the lowest curve of symmetry $\tilde{m}_l = 1$ ($\tilde{m}_l = 0$) connects to the non-protective (protective) barrier. The reactive rate is again a combination of $\beta_{\pi/2}$ and $\beta_0$. But as the coupling increases from $\theta_E = \pi/2$ to $\pi/4$, the coefficient of $\beta_{\pi/2}$ decreases while the one of $\beta_0$ increases, hence decreasing the total reactive rates and connecting to the trend between $\theta_E = 0$ and $\pi/4$.

The sensitivity of the fermionic reactive rate with the field angle for the KRb system is quite strong, since a small change of $\theta_E = \pi/10$ ($= 18^\circ$) gives rise to an order of magnitude increase in the rate. Therefore in experiments of fermionic KRb molecules, it is important for the electric field to be quite parallel to the 1D optical lattice confinement axis to avoid additional losses due to slight tilted angles. Note however that the range of this strong sensitivity of the rates may vary from one system to another.

In contrast for bosons, there is a very slight dependence of the reactive rate with $\theta_E$. This is due to the slight downward pushing of the $\tilde{m}_l = 2$ curve to the $\tilde{m}_l = 0$ ones when $\theta_E$ increases as can be barely seen on Fig. 4. The reactive rate value is anyway larger than the elastic and inelastic one.

Additionally, it is interesting to analyze the effect of the $\xi$ component. For fermions, the $\xi = -1$ rate coefficient turns out to be the same than the $\xi = +1$ one at $\theta_E = 0$ because Eq. (19) reduces to Eq. (10) which is an expression independent of the $\xi$ number. When departing from the angle $\theta_E = 0$, the $\xi = +1$ rate coefficients are dominant compared to the $\xi = -1$ ones. As a good approximation, one can neglect the contribution of the latter for large values of $\theta_E$ (note that as $d$ increases, this is less true though, see comment in Appendix A). For bosons at $\theta_E = 0$, the $\xi = +1$ term contains the $\tilde{m}_l = 0$ component responsible for a barrierless collision while intrinsically the $\xi = -1$ one does not. Therefore the latter case does not yield a higher rate than the $\xi = +1$ case, then it is a good approximation to neglect the $\xi = -1$ component for bosons for all angles $\theta_E$.

3. Rate coefficients versus the confinement

The effect of the confinement strength is shown in Fig. 4 for a fixed collision energy of $E_c = 500$ nK, induced dipole of $d = 0.2$ D and field angle $\theta_E = \pi/10$ ($18^\circ$). At $\nu = 20$ kHz, the first threshold $n_1 = 0, n_2 = 1$ is located at an energy of $\sim 1\mu K$ above the energy of the initial state ($n_1 = 0, n_2 = 0$). Therefore a collision energy of 500 nK is not sufficient to open up inelastic collisions and there is no inelastic rate. When the confinement decreases, there is a given $\nu$ for which the first inelastic transition becomes open, when the first threshold ($n_1 = 0, n_2 = 1$) energy amounts the value of the collision energy. This is the case here at $\nu = 10.4$ kHz. These different values of $\nu$ are indicated with an arrow along with the different threshold openings ($n_1, n_2$). At $\nu = 10.4$ kHz, the first inelastic threshold opening (first arrow from the right) for the fermions is quite strong, recalling the sharp one seen in Fig. 4 for $\theta_E = \pi/4$. The first one for the bosons is rather weak and smooth as also seen in Fig. 4. The second opening (second arrow from the right) is now smooth for the fermions and sharp for the bosons. And so forth, the successive openings alternate between sharp and smooth patterns. As
The inset in the top panel sketches the excitation inelastic and reactive rate coefficients for fermionic (top panel) and bosonic (bottom panel) KRb + KRb collisions as a function of the tilt angle $\theta_E$, at $E_\nu=500 \, \text{nK}$, $d=0.2 \, \text{D}$ and $\nu = 20 \, \text{kHz}$. The numbers in the brackets represent the harmonic states ($n_1, n_2$) of particles 1 and 2 and the arrows the corresponding energy thresholds. The inset in the top panel sketches the excitation inelastic process $(0,0) \rightarrow (0,1)$ (see text for details).

An interesting feature is seen for the fermionic system. At $\nu = 9 \, \text{kHz}$ for example, the inelastic rate $(n_1 = 0, n_2 = 0) \rightarrow (n_1 = 0, n_2 = 1)$ reaches the value of the elastic rate ($\sim 2 \times 10^{-6} \, \text{cm}^2 \, \text{s}^{-1}$) while the reactive rate is about 3.3 times smaller. The inelastic process here is an excitation inelastic process (see the inset of Fig. 6). The total energy is set before the collision by $E_{\text{tot}} = \varepsilon_0 + \varepsilon_0 + E_c = 0.932 \, \text{µK}$. We also have $\varepsilon_0 = 0.216 \, \text{µK}$ and $\varepsilon_1 = 0.648 \, \text{µK}$. After the collision, the total energy is given by $E_{\text{tot}} = \varepsilon_0 + \varepsilon_1 + E_c$, where $E_c$ is the final kinetic energy. The conservation of the total energy implies $E_c = E_c - (\varepsilon_1 - \varepsilon_0) = E_c - h\nu = 500 \, \text{nK} - 432 \, \text{nK} = 68 \, \text{nK}$. This means that for an excitation inelastic process, the particles after the collision have a smaller kinetic energy than before the collision and therefore slowed down by this mechanism. Loss of molecules can happen but according to the rates, for each loss of one pair, three pairs get slowed down from 500 nK to 68 nK. Note that the final kinetic energy depends on the choice of the frequency trap (see inset): its value is even smaller when the frequency is closer to the frequency that shuts off the inelastic transition, here $\nu = 10.4 \, \text{kHz}$. Therefore, particles can be slowed down to arbitrary small kinetic values, as far as the reactive rates remain smaller than the inelastic ones. Of course the reverse relaxation inelastic process $(n_1 = 0, n_2 = 1) \rightarrow (n_1 = 0, n_2 = 0)$ can also happen after the particles have been excited, restoring back the 500 nK kinetic energy to the particles. But this can be prevented by tilting the electric field back to the parallel case $\theta_E = 0$. In such cases, the inelastic transition is forbidden as discussed previously, leaving the particles in the harmonic states 0 and 1 with small kinetic energy. Another possibility is to find a way to remove directly the particles in the harmonic state 1 leaving only the one in the ground state 0 with small kinetic energy.

IV. CONCLUSION

By implementing tesseral harmonics in place of spherical harmonics in the collisional formalism of two ultracold tilted dipolar particles in confined space, we showed that we can recover a good quantum number $\xi$. This separates the overall problem into two sub-problems of smaller size even when a field is tilted. Inelastic and reactive rates show dramatical changes in a tilted field. This is due to additional couplings between $m_l$ components when a tilt is applied. We also showed that fermionic dipolar particles can lose kinetic energy and slow down due to favourable trap excitation inelastic collision under an appropriate confinement strength of the 1D lattice in a tilted field. Future works will investigate whether this mechanism can be efficient to even further cool down the particles in such configuration by taking into account the initial kinetic energy distribution of the particles for a given temperature and their rethermalization due to the elastic collisions during this process.

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Appendix A: $\xi = \pm 1$ adiabatic energy curves

Fig. 7 presents the adiabatic energy curves for $d = 0.2$ D and $\theta_E = \pi/4$ for both the fermionic and bosonic system and both quantum numbers $\xi = \pm 1$. One can see that the $\xi = -1$ curves always corresponds to more repulsive curves than $\xi = +1$ ones. This is due to the fact that $\bar{m}_l \geq 1$ for $\xi = -1$ and it correlates to a $l \geq 1$ curve with a large centrifugal barrier. At weak $d$ the $\xi = -1$ manifold does not play an important role in the collision. Note that at large $d$, the $l \geq 1$ curve can turn attractive again and then the $\xi = -1$ curve can start to play a role in the collision [28].

![Diagram of adiabatic energy curves](image)

**FIG. 7:** (Color online) Adiabatic energy curves as a function of the intermolecular distance $r$ for $d = 0.2$ D, $\theta_E = \pi/4$ for the $\xi = +1$ manifold (black solid lines) and the $\xi = -1$ manifold (red dashed lines). Fermions: top panel, bosons: bottom panel. Each adiabatic energy curve correlates to a state $n$ of the harmonic oscillator.

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