

Energy spectrum of diatomic molecules in the Quantum Drude Oscillator model

Matthieu Sarkis,^{*} Loris di Cairano,[†] and Alexandre Tkatchenko[‡]

Department of Physics and Materials Science
University of Luxembourg, L-1511,
Luxembourg City, Luxembourg.

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We compute the energy spectrum of a system consisting of a pair of Quantum Drude Oscillators coupled through the Coulomb interaction up to Quadrupole-Quadrupole and Dipole-Octupole order in the multipolar expansion by re-expressing the Schrödinger equation in terms of a tri-confluent Heun equation.

INTRODUCTION

MBD, non-covalent interaction, linear response,

A Quantum Drude Oscillator (QDO) is a minimalist atomistic model in which the properties of an atom are encompassed in a small number of parameters. The model consists in assimilating the atom to a point particle of mass m and electric charge $-q$ attached to a fixed (infinite mass) center of charge $+q$ by a harmonic spring characterized by a frequency ω . Molecules are then defined as a collection of QDOs interacting through Coulomb interactions.

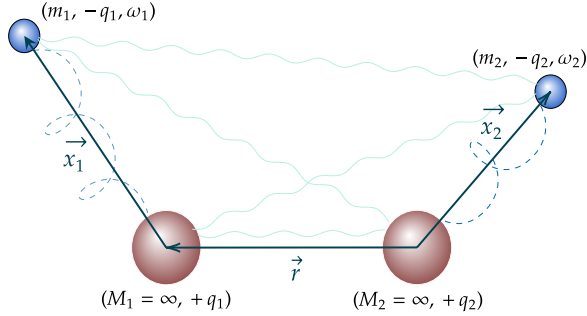


FIG. 1. Pair of interacting QDOs.

This simple model has been extensively used in various contexts, for instance in order to tackle the electronic structure problem for isolated molecules, in particular long range interactions, as well as to study the impact of an ambient bath or of an external electric field on molecular properties [1].

Even though the harmonic oscillator nature of the atomic constituents in this model drastically simplify the complexity of problem, the all-body Coulomb interaction remains challenging. One possible way to proceed is to treat the Coulomb interaction by performing a multipolar expansion.

By truncating the multipolar expansion up to Dipole-Dipole, one obtains a quadratic Hamiltonian. In the case of a closed linear chain, one can analytically solve for the spectrum of the Hamiltonian [2]. In the case of a general

geometry, one can solve for the London-van der Waals interaction energy can be calculated through a perturbative approach [3]. Though simple, this system was shown to capture long-range phenomena in large biomolecular systems [4].

In the present work, we focus instead on small (diatomic) molecules in 1d, and treat in an *analytical* way the Coulombic interactions up to Quadrupole-Quadrupole and Dipole-Octupole order, paving the way towards generalization for larger systems.

GitHub repository: The reader will find an open source code accompanying this paper following this link.

QUARTIC HAMILTONIAN FOR DIATOMIC MOLECULES

Multipolar expansion for the Coulomb potential

The Hamiltonian describing a system of N QDOs in 3d is given by:

$$H = \sum_{i=1}^N \left[\frac{1}{2} m_i \dot{\vec{x}}_i^2 + \frac{1}{2} m_i \omega_i^2 \vec{x}_i^2 \right] + \sum_{i < j} V_{\text{Coul}}(\vec{x}_i, \vec{x}_j), \quad (1)$$

with the Coulomb interaction given by

$$\frac{V_{\text{Coul}}(\vec{x}_1, \vec{x}_2)}{q_1 q_2 (4\pi\epsilon_0)^{-1}} = \frac{1}{r} - \frac{1}{|\vec{r} + \vec{x}_1|} - \frac{1}{|\vec{r} - \vec{x}_2|} + \frac{1}{|\vec{r} - \vec{x}_2 + \vec{x}_1|}$$

In the multipolar expansion, this can be expressed as:

$$V_{\text{Coul}}(\vec{x}_1, \vec{x}_2) = \sum_{n \geq 0} V_n(\vec{x}_1, \vec{x}_2), \quad (2)$$

with the following scaling behavior in terms of the distance between the centers:

$$V_n(\vec{x}_1, \vec{x}_2) \propto r^{-n+3}. \quad (3)$$

The first terms in that multipolar expansion have the following interpretation:

- $n = 0$: Dipole-Dipole interaction (DD),

- $n = 1$: Dipole-Quadrupole interaction (DQ),
- $n = 2$: Quadrupole-Quadrupole and Dipole-

Octupole interaction (QQ+DO).

Using the following identity for generic vectors \vec{r} and $\vec{\epsilon}$:

$$\frac{1}{|\vec{r} - \vec{\epsilon}|} = \frac{1}{r} \left[1 + \frac{\vec{\epsilon} \cdot \hat{r}}{r} - \frac{\epsilon^2 - 3(\vec{\epsilon} \cdot \hat{r})^2}{2r^2} + \frac{5(\vec{\epsilon} \cdot \hat{r})^3 - 3\epsilon^2(\vec{\epsilon} \cdot \hat{r})}{2r^3} + \frac{35(\vec{\epsilon} \cdot \hat{r})^4 - 30\epsilon^2(\vec{\epsilon} \cdot \hat{r})^2 + 3\epsilon^4}{8r^4} + \mathcal{O}\left(\frac{\epsilon^5}{r^5}\right) \right] \quad (4)$$

one can explicitly compute V_0 , V_1 and V_2 . Instead of writting out the expression of these potentials in full glory, let us now restrict ourselves to a system composed of $N = 2$ QDOs.

Reduced particle system

Let us move to the center of mass frame, in which the dynamics reduces to that of an effective reduced particle:

$$\vec{x} := \vec{x}_1 - \vec{x}_2 \quad (5a)$$

$$m_1 \vec{x}_1 + m_2 \vec{x}_2 = \vec{0}. \quad (5b)$$

In terms of the reduced variable, a straightforward computation gives us:

$$V_0(\vec{x}) = -\frac{m_1 m_2}{(m_1 + m_2)^2} \cdot \frac{x^2 - 3(\vec{x} \cdot \hat{r})^2}{4\pi\epsilon_0 r^3}, \quad (6a)$$

$$V_1(\vec{x}) = \frac{3m_1 m_2}{2(m_1 + m_2)^2} \cdot \frac{(\vec{x} \cdot \hat{r})(3x^2 - 5(\vec{x} \cdot \hat{r})^2)}{4\pi\epsilon_0 r^4}, \quad (6b)$$

$$V_2(\vec{x}) = \frac{m_1 m_2 (2m_1^2 + 3m_1 m_2 + 2m_2^2)}{4(m_1 + m_2)^4} \times \frac{3x^4 - 30(\vec{x} \cdot \hat{r})^2 x^2 + 35(\vec{x} \cdot \hat{r})^4}{4\pi\epsilon_0 r^5}. \quad (6c)$$

Let us now assume that the system is effectively one-dimensional, aligned along the direction on which the centers of the two QDOs are sitting. One obtain the following one-dimensional potentials for an effective particle of mass $m_1 m_2 / (m_1 + m_2)$:

$$V_0(x) = \frac{2m_1 m_2}{(m_1 + m_2)^2} \cdot \frac{2x^2}{4\pi\epsilon_0 r^3}, \quad (7a)$$

$$V_1(x) = -\frac{3m_1 m_2}{(m_1 + m_2)^2} \cdot \frac{x^3}{4\pi\epsilon_0 r^4}, \quad (7b)$$

$$V_2(x) = \frac{2m_1 m_2 (2m_1^2 + 3m_1 m_2 + 2m_2^2)}{(m_1 + m_2)^4} \cdot \frac{x^4}{4\pi\epsilon_0 r^5}. \quad (7c)$$

Including the contribution of the harmonic potentials present in (1) and coupling the system to an external electric field, one finally obtain the following total one-dimensional potential:

$$V(x) = \alpha x + \beta x^2 + \gamma x^3 + \delta x^4, \quad (8)$$

with the following expression of the coupling constants:

$$\alpha = -\frac{m_1 q_2 + m_2 q_1}{m_1 + m_2} \cdot \mathcal{E} \quad (9a)$$

$$\beta = \frac{m_1 m_2}{(m_1 + m_2)^2} \cdot \frac{2q_1 q_2}{4\pi\epsilon_0 r^3} + \frac{1}{2} \cdot \frac{m_1 m_2}{m_1 + m_2} \cdot \frac{m_1 \omega_2^2 + m_2 \omega_1^2}{m_1 + m_2} \quad (9b)$$

$$\gamma = -\frac{m_1 m_2}{(m_1 + m_2)^2} \cdot \frac{3q_1 q_2}{4\pi\epsilon_0 r^4}, \quad (9c)$$

$$\delta = \frac{2m_1 m_2 (2m_1^2 + 3m_1 m_2 + 2m_2^2)}{(m_1 + m_2)^4} \cdot \frac{q_1 q_2}{4\pi\epsilon_0 r^5}. \quad (9d)$$

ANALYTIC SOLUTION OF THE SCHRÖDINGER EQUATION

In what follows, we define $k^2 := \frac{m_1 + m_2}{m_1 m_2} \frac{\hbar^2}{2}$. Let us consider the time-independent Schrödinger equation

$$\left[\frac{d^2}{dx^2} + \frac{\epsilon - V(x)}{k^2} \right] \psi(x) = 0, \quad (10)$$

where the potential V is provided by (8) and (9).

Following [5], we solve the problem separately on the positive and negative real axis, and impose that the two solutions (positive and negative) ψ_{\pm} and their first derivative match at the origin. For brevity reasons, we treat explicitly the case of the positive solution, the negative case being treated by performing the transformation $(x, \alpha, \beta, \gamma, \delta) \rightarrow (-x, -\alpha, \beta, -\gamma, \delta)$.

Performing the following change of variable:

$$\psi_+(x) = \exp \left[\frac{\gamma^2 - 4\beta\delta}{8k\delta^{3/2}} x - \frac{\gamma}{4k\delta^{1/2}} x^2 - \frac{\delta^{1/2}}{3k} x^3 \right] \phi_+(x) \quad (11)$$

we obtain:

$$\begin{aligned} \frac{d^2\phi_+}{dx^2}(x) + \left[\frac{\gamma^2 - 4\beta\delta}{4k\delta^{3/2}} - \frac{\gamma}{k\delta^{1/2}}x - \frac{2\delta^{1/2}}{k}x^2 \right] \frac{d\phi_+}{dx}(x) \\ + \left[\frac{16\delta^2(\beta^2 + 4\delta\epsilon) - 8\beta\gamma^2\delta + \gamma^4 - 32\gamma\delta^{5/2}k}{64\delta^3k^2} - \frac{8\alpha\delta^2 - 4\beta\gamma\delta + \gamma^3 + 16\delta^{5/2}k}{8\delta^2k^2}x \right] \phi_+(x) = 0, \end{aligned} \quad (12)$$

which is known as the so-called *tri-confluent Heun equation*. We further perform the following change of variable, the reason why to become clear soon:

$$\phi_+(x) = (1+x)^{\frac{4\beta\gamma\delta - \gamma^3 - 8\delta^2(\alpha + 2k\delta^{1/2})}{16k\delta^{5/2}}} f_+(x), \quad (13)$$

as well as map the positive real axis to the segment $[0, 1]$ through the change of argument

$$z = \frac{x}{x+1}, \quad (14)$$

and look for a solution of the obtained ODE in terms of a series expansion at zero of the form:

$$f_+(z) = \sum_{n \geq 0} c_n^+ z^n. \quad (15)$$

Plugging the ansatz into the differential equation, we obtain the following recurrence relation between the coefficients:

$$\begin{aligned} (n+1)(n+2)c_{n+2}^+ + (n+1)(A-4n)c_{n+1}^+ + [6(n-1)n + Bn + E]c_n^+ \\ + [C(n-1) + F - 4(n-2)(n-1)]c_{n-1}^+ + [(n-3)(n-2) + D(n-2) + G]c_{n-2}^+ = 0 \quad (n \geq 2) \end{aligned} \quad (16)$$

where the various coefficients are listed in full details in appendix. The change of variable (13) is chosen precisely such that the order of the above difference equation is as small as possible. The coefficients c_0^+ and c_1^+ are to be fixed through boundary conditions, and the coefficients c_2^+ and c_3^+ are in turn fixed by the following relations

$$\begin{aligned} c_2^+ &= -Ec_0^+ - Ac_1^+, \\ c_3^+ &= -\frac{Fc_0^+ + (B+E)c_1^+ + 2(A-4)c_2^+}{6}. \end{aligned} \quad (17)$$

Let us similarly denote by c_n^- the coefficients in the series expansion corresponding to the solution on the negative real axis. It has been shown in [5] and [6] that the smooth matching of the two solutions at the origin can be achieved by imposing the so-called *dominant solution* to the above defined difference equations vanishes. The conditions

$$\psi_+(0) = \psi_-(0) \quad \text{and} \quad \frac{d\psi_+}{dx}(0) = \frac{d\psi_-}{dx}(0) \quad (18)$$

lead to constraints on the initial conditions for the difference equations, namely that $c_0^+ = c_0^-$ and:

$$\begin{aligned} c_1^+ - c_0^+ \left[1 + \frac{8\delta^2(\alpha + \beta) - 2\gamma\delta(\gamma + 2\beta) + \gamma^3}{16\delta^{5/2}k} \right] \\ = -c_1^- + c_0^- \left[1 - \frac{8\delta^2(\alpha - \beta) + 2\gamma\delta(\gamma - 2\beta) + \gamma^3}{16\delta^{5/2}k} \right] \end{aligned} \quad (19)$$

The two sides of relation (19) are therefore equal to some quantity s , which parameterizes the solution. Now, the

vanishing of the dominant solution can be achieved by imposing that the series coefficients $c_n^+(\epsilon, s)$ and $c_n^-(\epsilon, s)$ are identically 0 asymptotically in n . This provides a pair of algebraic equations in the energy variable ϵ and the *shift* variable s .

Let us make a few remarks at this stage. Imposing the vanishing of c_n^\pm asymptotically in n means that practically one should fix a large value of n , denote it N , and impose that the polynomials c_N^\pm vanish. One observes that c_N^- is a linear function of s . One can therefore solve explicitly for the shift variable in terms of the energy ϵ . Plugging back into the other constraint, one obtains a rational function of the form:

$$c_N^+(\epsilon, s(\epsilon)) = \frac{P_N(\epsilon)}{Q_N(\epsilon)} \quad (20)$$

where P_N and Q_N are polynomial functions of ϵ , whose degree d_N is an non-decreasing function of N . The N zeros $\{\epsilon_i\}_{i=1}^N$ correspond then to first N energy levels of the system, provided they are not complex and that Q_N does not admit a zero of the same order.

RESULTS

First few energy levels

Ground state energy r vs. \mathcal{E}

TABLE I. Energy spectrum (in Ha) of the Ar-Ar system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	9.247342	32.416619	62.971511
1.0	2.981229	10.350087	20.248422
1.5	1.563464	5.442288	10.069702
2.0	1.023301	3.394336	6.337199
2.5	0.748161	2.413177	4.876881
3.0	0.590991	1.960271	4.289340

TABLE II. Energy spectrum (in Ha) of the Cs-Cs system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	4.076335	14.287104	27.755741
1.0	1.307214	4.543529	8.897871
1.5	0.673427	2.362932	4.396492
2.0	0.426306	1.441411	2.686432
2.5	0.296956	0.971662	1.957474
3.0	0.217699	0.719613	1.585340

TABLE III. Energy spectrum (in Ha) of the H-H system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	3.757546	13.234211	25.693826
1.0	1.213930	4.243743	8.293979
1.5	0.649664	2.239448	4.116225
2.0	0.436524	1.423565	2.744604
2.5	0.330635	1.081656	2.284054
3.0	0.279296	0.965099	2.204208

TABLE IV. Energy spectrum (in Ha) of the He-He system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	4.838555	16.998352	32.990230
1.0	1.609674	5.563162	10.796479
1.5	0.928052	3.096340	5.617631
2.0	0.690066	2.190579	4.209331
2.5	0.597424	1.987363	4.173586
3.0	0.607965	2.159394	4.851867

TABLE V. Energy spectrum (in Ha) of the K-K system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	3.900846	13.629450	26.449767
1.0	1.254074	4.338725	8.463017
1.5	0.646062	2.260433	4.233126
2.0	0.409452	1.389547	2.556108
2.5	0.286775	0.935829	1.841376
3.0	0.211508	0.689674	1.476942

TABLE VI. Energy spectrum (in Ha) of the Kr-Kr system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	9.936548	34.839598	67.682787
1.0	3.196890	11.107797	21.743625
1.5	1.666467	5.817323	10.779216
2.0	1.078692	3.598874	6.723190
2.5	0.776461	2.514230	5.085759
3.0	0.599348	1.984794	4.360291

TABLE VII. Energy spectrum (in Ha) of the Li-Li system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	2.989579	10.393189	20.094511
1.0	0.965106	3.317199	6.411919
1.5	0.498061	1.728672	3.276409
2.0	0.316132	1.076633	1.951989
2.5	0.223280	0.727240	1.384589
3.0	0.166518	0.533844	1.098635

TABLE VIII. Energy spectrum (in Ha) of the Ne-Ne system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	8.067324	28.257747	54.866835
1.0	2.648128	9.141505	17.802546
1.5	1.463214	4.975572	9.086247
2.0	1.037985	3.321372	6.216936
2.5	0.839105	2.707087	5.494303
3.0	0.769833	2.636825	5.742950

TABLE IX. Energy spectrum (in Ha) of the Rb-Rb system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	4.031874	14.113400	27.409612
1.0	1.294297	4.489885	8.780383
1.5	0.666759	2.337266	4.359340
2.0	0.422361	1.430245	2.650383
2.5	0.294937	0.963709	1.922279
3.0	0.216807	0.712417	1.550805

TABLE X. Energy spectrum (in Ha) of the Xe-Xe system

spectrum r (a.u.)	ϵ_0	ϵ_1	ϵ_2
0.5	10.472412	36.762340	71.427248
1.0	3.360994	11.705550	22.941485
1.5	1.743863	6.103626	11.305873
2.0	1.117696	3.742926	7.034869
2.5	0.792188	2.580753	5.270133
3.0	0.598300	1.992258	4.438638

TABLE XI. Ground state energy (in Ha) of the Ar-Ar system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	9.247342	9.110717	8.952432
1.0	2.981229	2.633184	2.076356
1.5	1.563464	0.829677	-0.591543
2.0	1.023301	-0.325559	-2.992697
2.5	0.748161	-1.377726	-5.452778
3.0	0.590991	-2.381467	-7.976320

TABLE XII. Ground state energy (in Ha) of the Cs-Cs system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	4.076335	3.980857	3.863462
1.0	1.307214	1.035551	0.558141
1.5	0.673427	0.042066	-1.194431
2.0	0.426306	-0.786686	-3.026094
2.5	0.296956	-1.655817	-5.032131
3.0	0.217699	-2.584558	-7.136808

TABLE XIII. Ground state energy (in Ha) of the H-H system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	3.757546	3.680220	3.584748
1.0	1.213930	0.994879	0.606484
1.5	0.649664	0.155212	-0.841764
2.0	0.436524	-0.467646	-2.247426
2.5	0.330635	-1.037533	-3.672527
3.0	0.279296	-1.563088	-5.125057

TABLE XIV. Ground state energy (in Ha) of the He-He system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	4.838555	4.747673	4.637588
1.0	1.609674	1.370517	0.955271
1.5	0.928052	0.449201	-0.551122
2.0	0.690066	-0.067148	-1.750946
2.5	0.597424	-0.386765	-2.715690
3.0	0.607965	-0.544976	-3.452658

TABLE XV. Ground state energy (in Ha) of the K-K system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	3.900846	3.798718	3.672463
1.0	1.254074	0.961731	0.444593
1.5	0.646062	-0.035078	-1.362536
2.0	0.409452	-0.892356	-3.275553
2.5	0.286775	-1.798818	-5.377141
3.0	0.211508	-2.769875	-7.827196

TABLE XVI. Ground state energy (in Ha) of the Kr-Kr system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	9.936548	9.795814	9.633507
1.0	3.196890	2.840242	2.274883
1.5	1.666467	0.914564	-0.532113
2.0	1.078692	-0.316154	-3.056110
2.5	0.776461	-1.455630	-5.674813
3.0	0.599348	-2.572968	-8.404424

TABLE XVII. Ground state energy (in Ha) of the Li-Li system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	2.989579	2.886101	2.754734
1.0	0.965106	0.657280	0.096843
1.5	0.498061	-0.231815	-1.636857
2.0	0.316132	-1.061981	-3.526222
2.5	0.223280	-1.953835	-5.616848
3.0	0.166518	-2.914737	-11.972520

TABLE XVIII. Ground state energy (in Ha) of the Ne-Ne system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	8.067324	7.938880	7.788706
1.0	2.648128	2.324114	1.795116
1.5	1.463214	0.814542	-0.481886
2.0	1.037985	-0.043649	-2.342096
2.5	0.839105	-0.668123	-3.988911
3.0	0.769833	-1.095276	-5.406152

TABLE XIX. Ground state energy (in Ha) of the Rb-Rb system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	4.031874	3.933375	3.812070
1.0	1.294297	1.013631	0.519442
1.5	0.666759	0.014231	-1.261255
2.0	0.422361	-0.828604	-3.131122
2.5	0.294937	-1.715198	-5.181182
3.0	0.216807	-2.663303	-7.510481

Atomic data

We collect here the QDO parameters obtained in [7].

Details on the derivation

TABLE XX. Ground state energy (in Ha) of the Xe-Xe system

r (a.u.) \ \mathcal{E} (a.u.)	0.0	1.0	2.0
0.5	10.472412	10.333027	10.172863
1.0	3.360994	3.008892	2.455036
1.5	1.743863	1.001161	-0.421036
2.0	1.117696	-0.272030	-2.991068
2.5	0.792188	-1.461408	-5.680509
3.0	0.598300	-2.651579	-8.519189

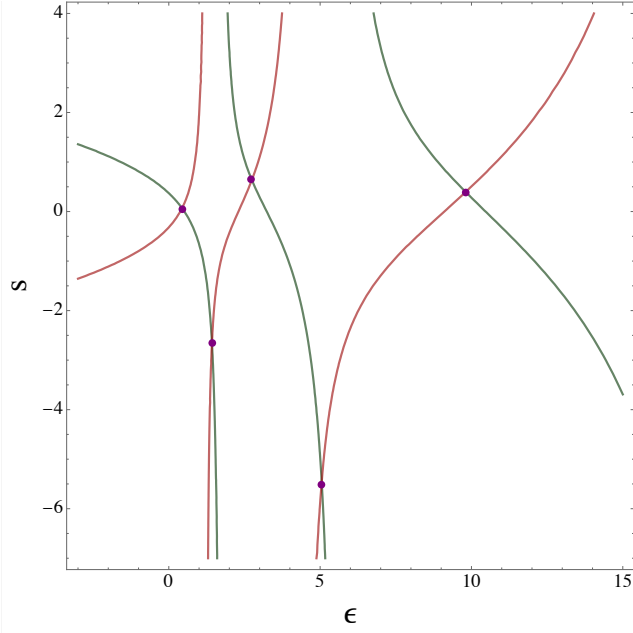


FIG. 2. Intersection loci of the three surfaces $z = c_{20}^+(\epsilon, s)$ and $z = c_{20}^-(\epsilon, s)$ and $z = 0$. The purple dots correspond to the first 5 excited states of the H-H system for $r = 2$ and $\mathcal{E} = 0$.

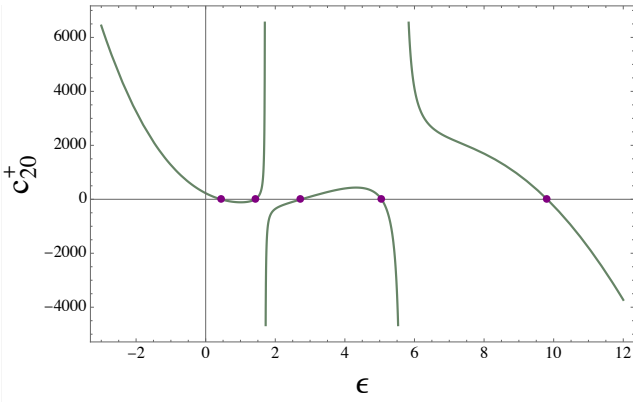


FIG. 3. Plot of $z = c_{20}^+(\epsilon, s(\epsilon))$ for the H-H system at $r = 2$ and $\mathcal{E} = 0$. The zeros are denoted by purple dots and correspond to the first excited states of the system.

TABLE XXI. Parameters for QDOs (in a.u.)

	ω	m	q
H	0.4273	0.6099	0.7080
Li	0.0687	1.2545	0.9848
K	0.0630	0.8101	0.9670
Rb	0.0603	0.7343	0.9274
Cs	0.0531	0.6939	0.8950
He	1.0187	0.5083	0.8532
Ne	1.2965	0.3491	1.2494
Ar	0.7272	0.3020	1.3314
Kr	0.6359	0.2796	1.3741
Xe	0.5152	0.2541	1.3570

$$\begin{aligned}
A &= -\frac{8\delta^2(\alpha + \beta) - 2\gamma\delta(2\beta + \gamma) + \gamma^3 + 32\delta^{5/2}k}{8\delta^{5/2}k} \\
B &= \frac{8\delta^2(3\alpha + 2\beta - \gamma) - 4\gamma\delta(3\beta + \gamma) + 3\gamma^3 + 96\delta^{5/2}k}{8\delta^{5/2}k} \\
C &= -\frac{8\delta^2(3\alpha + \beta - \gamma) - 2\gamma\delta(6\beta + \gamma) + 3\gamma^3 + 16\delta^3 + 96\delta^{5/2}k}{8\delta^{5/2}k} \\
D &= \frac{-4\beta\gamma\delta + \gamma^3 + 8\delta^2(\alpha + 4\sqrt{\delta}k)}{8\delta^{5/2}k} \\
E &= \frac{4\gamma^2\delta^2(\gamma(4\alpha + \gamma) + 4\beta^2 + 8\beta\gamma) - 32\gamma\delta^3(\alpha + \beta)(2\beta + \gamma) + 64\delta^4(\alpha + \beta)^2 - 4\gamma^4\delta(2\beta + \gamma)}{256\delta^5k^2} \\
&\quad + \frac{\gamma^6 + 256\delta^5(2k^2 + \epsilon) + 128\delta^{9/2}k(3\alpha + 2\beta - \gamma) - 64\gamma\delta^{7/2}k(3\beta + \gamma) + 48\gamma^3\delta^{5/2}k}{256\delta^5k^2} \\
F &= -\frac{\left(-4\beta\gamma\delta + \gamma^3 + 8\delta^2(\alpha + 2\sqrt{\delta}k)\right)\left(-4\gamma\delta(\beta + 2\delta) + \gamma^3 - 2\gamma^2\delta + 8\delta^2(\alpha + \beta + 2\delta + 4\sqrt{\delta}k)\right)}{128\delta^5k^2} \\
G &= \frac{\left(-4\beta\gamma\delta + \gamma^3 + 8\delta^2(\alpha + 2\sqrt{\delta}k)\right)\left(-4\beta\gamma\delta + \gamma^3 + 8\delta^2(\alpha + 4\sqrt{\delta}k)\right)}{256\delta^5k^2}
\end{aligned}$$

* matthieu.sarkis@uni.lu

† loris.dicairano@uni.lu

‡ alexandre.tkatchenko@uni.lu

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