

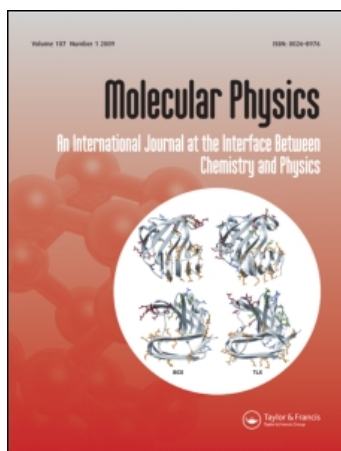
This article was downloaded by: [Australian National University]

On: 5 November 2010

Access details: Access Details: [subscription number 917491237]

Publisher Taylor & Francis

Informa Ltd Registered in England and Wales Registered Number: 1072954 Registered office: Mortimer House, 37-41 Mortimer Street, London W1T 3JH, UK



Molecular Physics

Publication details, including instructions for authors and subscription information:

<http://www.informaworld.com/smpp/title~content=t713395160>

Excited states of spherium

Pierre-François Loos^a; Peter M. W. Gill^a

^a Research School of Chemistry, Australian National University, Canberra, Australia

First published on: 02 September 2010

To cite this Article Loos, Pierre-François and Gill, Peter M. W.(2010) 'Excited states of spherium', Molecular Physics, 108: 19, 2527 – 2532, First published on: 02 September 2010 (iFirst)

To link to this Article: DOI: 10.1080/00268976.2010.508472

URL: <http://dx.doi.org/10.1080/00268976.2010.508472>

PLEASE SCROLL DOWN FOR ARTICLE

Full terms and conditions of use: <http://www.informaworld.com/terms-and-conditions-of-access.pdf>

This article may be used for research, teaching and private study purposes. Any substantial or systematic reproduction, re-distribution, re-selling, loan or sub-licensing, systematic supply or distribution in any form to anyone is expressly forbidden.

The publisher does not give any warranty express or implied or make any representation that the contents will be complete or accurate or up to date. The accuracy of any instructions, formulae and drug doses should be independently verified with primary sources. The publisher shall not be liable for any loss, actions, claims, proceedings, demand or costs or damages whatsoever or howsoever caused arising directly or indirectly in connection with or arising out of the use of this material.

INVITED ARTICLE

Excited states of spherium

Pierre-François Loos and Peter M.W. Gill*

Research School of Chemistry, Australian National University, Australian Capital Territory 0200, Canberra, Australia

(Received 21 April 2010; final version received 8 July 2010)

We report analytic solutions of a recently discovered quasi-exactly solvable model consisting of two electrons, interacting via a Coulomb potential, but restricted to remain on the surface of a \mathcal{D} -dimensional sphere. Polynomial solutions are found for the ground state, and for some higher ($L \leq 3$) states. Kato cusp conditions and interdimensional degeneracies are discussed.

Keywords: exact solution; excited states; spherium; cusp condition; interdimensional degeneracies

1. Introduction

A quasi-exactly solvable model is one for which it is possible to solve the Schrödinger equation exactly for a finite portion of the energy spectrum [1]. In quantum chemistry, a famous example of this is the Hooke's law atom [2–5], which consists of a pair of electrons, repelling Coulombically but trapped in a harmonic external potential. This model and others [6–12] have been used extensively to test various approximations [13–20] within density functional theory (DFT) [21–23] and explicitly correlated methods [24–28].

We have recently discovered [29] that a pair of electrons, repelling Coulombically but constrained to remain on the surface of a \mathcal{D} -sphere of radius R [17,30–37] is also quasi-exactly solvable and we have called this system \mathcal{D} -spherium. (We adopt the convention that a \mathcal{D} -sphere is the surface of a $(\mathcal{D}+1)$ -dimensional ball.) We have shown that the Schrödinger equation for the 1S and the 3P states of \mathcal{D} -spherium can be solved exactly for a countably infinite set of R values and that the resulting wave functions are polynomials in the interelectronic distance $0 \leq u \equiv |\mathbf{r}_1 - \mathbf{r}_2| \leq 2R$.

In this article, we extend our earlier results [29] to higher angular momentum (up to $L=3$) states of \mathcal{D} -spherium ($\mathcal{D} \geq 2$) for both the singlet and triplet manifolds. The $\mathcal{D}=1$ case is anomalous and, for brevity, is not discussed here. We use atomic units throughout.

2. Wave function

The Hamiltonian of \mathcal{D} -spherium is

$$\hat{H} = -\frac{1}{2}(\nabla_1^2 + \nabla_2^2) + \frac{1}{u}, \quad (1)$$

where the two first terms represent the kinetic contribution of each electron, and u^{-1} is the Coulomb operator.

Following Breit [38], we write the total wave function as the product

$$\Phi(\{s_1, s_2\}, \{\mathbf{\Omega}_1, \mathbf{\Omega}_2\}, u) = \Xi(s_1, s_2)\chi(\mathbf{\Omega}_1, \mathbf{\Omega}_2)\Psi(u), \quad (2)$$

where Ξ , χ and Ψ are the spin, angular and interparticle wave functions, respectively, and s_i and $\mathbf{\Omega}_i$ are the spin and hyperspherical coordinates [39] of the i th electron. The singlet and triplet wave functions are given by the familiar [40] forms

$$^1\Xi(s_1, s_2) = \frac{1}{2^{1/2}}[\alpha(s_1)\beta(s_2) - \beta(s_1)\alpha(s_2)], \quad (3)$$

$$^3\Xi(s_1, s_2) = \begin{cases} \alpha(s_1)\alpha(s_2), \\ \frac{1}{2^{1/2}}[\alpha(s_1)\beta(s_2) + \beta(s_1)\alpha(s_2)], \\ \beta(s_1)\beta(s_2). \end{cases} \quad (4)$$

The angular part is associated with an energy

$$E_\chi = \frac{A}{R^2}, \quad (5)$$

$$A = \frac{\ell_1(\ell_1 + \mathcal{D} - 1)}{2} + \frac{\ell_2(\ell_2 + \mathcal{D} - 1)}{2}, \quad (6)$$

*Corresponding author. Email: peter.gill@anu.edu.au

Table 1. Ground state and excited states of \mathcal{D} -spherium.

State	Configuration	$\chi(\Omega_1, \Omega_2)$	δ	γ^{-1}	A	κ	Transformation
$^1S^e$	s^2	1	$2\mathcal{D}-1$	$\mathcal{D}-1$	0	0	$^3P^e$
$^3P^o$	sp	$\cos \theta_1 - \cos \theta_2$	$2\mathcal{D}+1$	$\mathcal{D}+1$	$\mathcal{D}/2$	1	$^1D^o$
$^1P^o$	sp	$\cos \theta_1 + \cos \theta_2$	$2\mathcal{D}+1$	$\mathcal{D}-1$	$\mathcal{D}/2$	0	$^3D^o$
$^3P^e$	p^2	$\sin \theta_1 \sin \theta_2 \sin(\phi_1 - \phi_2)$	$2\mathcal{D}+3$	$\mathcal{D}+1$	\mathcal{D}	1	
$^3D^e$	sd	$\chi^3_{p^o} \cdot \chi^1_{p^o}$	$2\mathcal{D}+3$	$\mathcal{D}+1$	$\mathcal{D}+1$	1	$^1F^e$
$^1D^o$	pd	$\chi^3_{p^o} \cdot \chi^3_{p^e}$	$2\mathcal{D}+5$	$\mathcal{D}+3$	$3\mathcal{D}/2+1$	2	
$^3D^o$	pd	$\chi^1_{p^o} \cdot \chi^3_{p^e}$	$2\mathcal{D}+5$	$\mathcal{D}+1$	$3\mathcal{D}/2+1$	1	
$^1F^e$	pf	$\chi^3_{p^e} \cdot \chi^3_{D^e}$	$2\mathcal{D}+7$	$\mathcal{D}+3$	$2\mathcal{D}+3$	2	

where ℓ_1 and ℓ_2 are the angular momentum quantum numbers of the corresponding one-electron configuration ($s=0, p=1, d=2, f=3, \dots$). The functions χ_i , which are dependent on the nature of the state considered [38,41], are gathered in Table 1, where $\theta_i \in [0, \pi]$ and $\phi_i \in [0, 2\pi]$ are the $(\mathcal{D}-1)$ th and \mathcal{D} th hyperspherical angles of the electron i . The corresponding one-electron configurations are also reported. In Table 1, the suffixes e (even) and o (odd) are related to the parity of the states, which is given by $(-1)^{\ell_1+\ell_2}$. Hence, we label the states with the notation $^{1,3}L^{e,o}$, where $L=S, P, D, F, \dots$

3. Polynomial solutions

Substituting the ansatz (2) into the Hamiltonian (1) yields the Schrödinger equation

$$\left(\frac{u^2}{4R^2} - 1\right) \frac{d^2\Psi}{du^2} + \left(\frac{\delta u}{4R^2} - \frac{1}{\gamma u}\right) \frac{d\Psi}{du} + \frac{\Psi}{u} = E\Psi, \quad (7)$$

where the parameters δ and γ are tabulated for each state in Table 1.

By introducing the dimensionless variable $x = u/2R$, Equation (7) can be recast as a Heun's differential equation [42] with singular points at $x = -1, 0, +1$. Following the known solutions of this equation [43], we seek wave functions of the form

$$\Psi(u) = \sum_{k=0}^{\infty} a_k u^k, \quad (8)$$

and substitution into (7) yields the three-step recurrence relation

$$a_{k+2} = \frac{\gamma}{(k+2)[(k+1)\gamma+1]} \times \left\{ a_{k+1} + \left[\frac{k(k+\delta-1)}{4R^2} - E \right] a_k \right\}, \quad (9)$$

with the starting values $a_0 = 1$ and $a_1 = \gamma$.

If the series (8) terminates at a certain $k=n$, we obtain the exact wave function

$$\Psi_{n,m}(u) = \sum_{k=0}^n a_k u^k, \quad (10)$$

for a particular radius $R_{n,m}$ and energy $E_{n,m}$. This is an n th degree polynomial with m nodes between 0 and $2R$ ($0 \leq m \leq \lfloor (n+1)/2 \rfloor$) and requires that a_{n+1} and a_{n+2} vanish. If $a_{n+1} = 0$, the relation

$$R_{n,m}^2 E_{n,m} = \frac{n}{2} \left(\frac{n}{2} + \frac{\delta-1}{2} \right) \quad (11)$$

ensures that $a_{n+2} = 0$. For a given n , the energies are thus determined by finding the roots of the equation $a_{n+1} = 0$, which is a polynomial in E , of degree $\lfloor (n+1)/2 \rfloor$.

For the $^1D^e$ state, we have not been able to obtain polynomial solutions because the Hamiltonian (1) is not separable using the ansatz (2) and the wave function satisfies exchange coupled equations [44]. This applies also to some other states of higher angular momentum.

4. Results and discussion

Numerical values of the energies and radii for the $^1P^o$ and $^3P^e$ states are reported in Tables 2 and 3. Tables containing results for the $^1S^e$ and $^3P^o$ states can be found in [29]. Numerical values of the energies and radii for the higher angular momentum states can be determined using the interdimensional degeneracies (see Section 4.5).

For any given state, as n increases, the radius increases and the energy decreases. The opposite behaviour is observed with respect to m . Furthermore, as R (or, equivalently, n) increases, the electrons tend to localize on opposite sides of the sphere due to the dominance of the Coulomb interaction as the density decreases [34,35]. Such Wigner crystallization [45] has also been observed in other systems [5,10,46].

Table 2. Radii $R_{n,m}$ and energies $E_{n,m}$ for $^1P^o$ states of two electrons on a D -sphere ($D=2, 3, 4$).

	n/m	$D=2$				$D=3$				$D=4$			
		0	1	2	3	0	1	2	3	0	1	2	3
Radius	1	1.118				1.871				2.598			
	2	3.162				4.637				6.083			
	3	6.226	1.656			8.376	2.520			10.52	3.303		
	4	10.30	4.232			13.11	5.888			15.93	7.440		
	5	15.38	7.847	2.159		18.84	10.21	3.127		22.32	12.49	3.966	
	6	21.46	12.49	5.246		25.57	15.53	7.077		29.72	18.51	8.732	
	7	28.54	18.14	9.397	2.639	33.30	21.84	11.97	3.707	38.11	25.51	14.39	4.599
	8	36.63	24.80	14.59	6.222	42.03	29.15	17.86	8.220	47.49	33.49	21.01	9.976
Energy	1	1.000				0.5000				0.3333			
	2	0.3000				0.1860				0.1351			
	3	0.1355	1.914			0.09622	1.063			0.07460	0.7562		
	4	0.07541	0.4467			0.05820	0.2884			0.04731	0.2168		
	5	0.04757	0.1827	2.414		0.03874	0.1319	1.406		0.03260	0.1041	1.033	
	6	0.03257	0.09620	0.5450		0.02753	0.07468	0.3594		0.02378	0.06129	0.2754	
	7	0.02363	0.05851	0.2180	2.764	0.02051	0.04771	0.1587	1.656	0.01808	0.04035	0.1267	1.241
	8	0.01789	0.03903	0.1127	0.6200	0.01585	0.03295	0.08781	0.4144	0.01419	0.02853	0.07252	0.3215

Table 3. Radii $R_{n,m}$ and energies $E_{n,m}$ for $^3P^e$ states of two electrons on a D -sphere ($D=2, 3, 4$).

	n/m	$D=2$				$D=3$				$D=4$			
		0	1	2	3	0	1	2	3	0	1	2	3
Radius	1	2.291				3.000				3.708			
	2	5.477				6.892				8.307			
	3	9.616	3.006			11.72	3.748			13.84	4.478		
	4	14.73	6.851			17.52	8.334			20.32	9.797		
	5	20.84	11.62	3.676		24.30	13.82	4.453		27.78	16.01	5.208	
	6	27.94	17.35	8.156		32.07	20.26	9.708		36.22	23.15	11.22	
	7	36.04	24.05	13.54	4.315	40.83	27.66	15.84	5.128	45.65	31.25	18.10	5.909
	8	45.13	31.75	19.86	9.412	50.58	36.04	22.90	11.03	56.07	40.32	25.89	12.60
Energy	1	0.3333				0.2500				0.2000			
	2	0.1333				0.1053				0.08696			
	3	0.07300	0.7472			0.06002	0.5874			0.05093	0.4862		
	4	0.04607	0.2131			0.03908	0.1728			0.03390	0.1459		
	5	0.03166	0.1019	1.018		0.02752	0.08503	0.8196		0.02429	0.07315	0.6913	
	6	0.02306	0.05983	0.2706		0.02042	0.05117	0.2228		0.01829	0.04477	0.1906	
	7	0.01752	0.03932	0.1242	1.222	0.01575	0.03432	0.1046	0.9983	0.01428	0.03046	0.09079	0.8522
	8	0.01375	0.02778	0.07096	0.3161	0.01251	0.02464	0.06102	0.2629	0.01145	0.02215	0.05370	0.2269

The energies of the S , P and D states ($m=0$) for 3-spherium are plotted in Figure 1 (the quasi-exact solutions are indicated by markers), while density plots of 2-spherium ($n=1$ and $m=0$) are represented on Figure 2.

4.1. Natural/unnatural parity

In attempting to explain Hund's rules [47] and the 'alternating' rule [48,49] (see also [50,51]), Morgan and

Kutzelnigg [52–54] have proposed that the two-electron atomic states be classified thus: *a two-electron state, composed of one-electron spatial orbitals with individual parities $(-1)^{\ell_1}$ and $(-1)^{\ell_2}$ and hence with overall parities $(-1)^{\ell_1+\ell_2}$, is said to have natural parity if its parity is $(-1)^L$. [...] If the parity of the two-electron state is $-(-1)^L$, the state is said to be of unnatural parity.* [54].

After introducing spin, three classes emerge. In a three-dimensional space, the states with a cusp

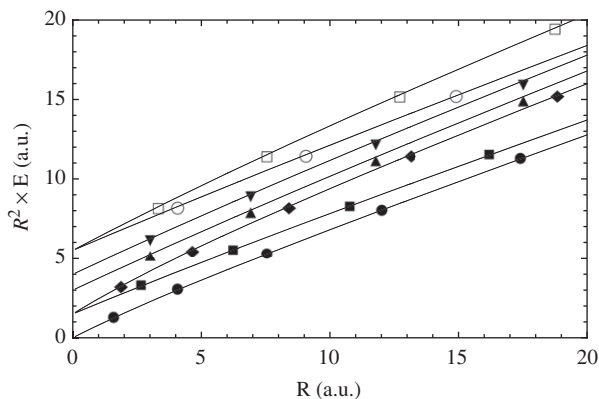


Figure 1. Energy of the S , P and D states of 3-spherium (${}^1S^e < {}^3P^o \leq {}^1P^o < {}^3P^e < {}^3D^e < {}^1D^o \leq {}^3D^o$). The quasi-exact solutions are shown by the markers.

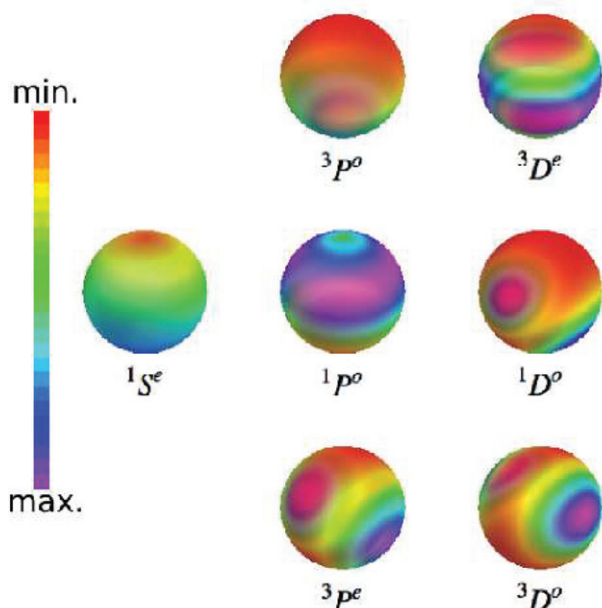


Figure 2. Density plots of the S , P and D states of 2-spherium. The squares of the wave functions when one electron is fixed at the north pole are represented. The radii are $3^{1/2}/2$, $15^{1/2}/2$, $5^{1/2}/2$, $21^{1/2}/2$, $21^{1/2}/2$, $3(5^{1/2})/2$ and $3(3^{1/2})/2$ for the ${}^1S^e$, ${}^3P^o$, ${}^1P^o$, ${}^3P^e$, ${}^3D^e$, ${}^1D^o$ and ${}^3D^o$ states, respectively.

value of $1/2$ [55,56] are known as the *natural parity singlet states*, those with a cusp value of $1/4$ [57] are the *natural and unnatural parity triplet states*, and those with a cusp value of $1/6$ [52], are the *unnatural parity singlet states*.

In previous work [29], we have observed that the ${}^1S^e$ ground state and the first excited ${}^3P^o$ state of 3-spherium possess the same singlet ($1/2$) and

triplet ($1/4$) cusp conditions as those for electrons moving in three-dimensional physical space and we have therefore argued that 3-spherium may be the most appropriate model for studying ‘real’ atomic or molecular systems. This is supported by the similarity of the correlation energy E_c of 3-spherium to that in other two-electron systems. Indeed, it can be shown [36] that, as $R \rightarrow 0$, E_c approaches -0.0476 , which is close to the corresponding values for the helium-like ions (-0.0467) [58], the Hooke’s law atom (-0.0497) [59], and two electrons in a ball (-0.0552) [46].

Most of the higher angular momentum states of 3-spherium, possess the ‘normal’ cusp values of $1/2$ and $1/4$. However, the unnatural ${}^1D^o$ and ${}^1F^e$ states have the cusp value of $1/6$.

4.2. First-order cusp condition

The wave function, radius and energy of the lowest states are given by

$$\Psi_{1,0}(u) = 1 + \gamma u, \quad R_{1,0}^2 = \frac{\delta}{4\gamma}, \quad E_{1,0} = \gamma, \quad (12)$$

which are closely related to the Kato cusp condition [56]

$$\frac{\Psi'(0)}{\Psi(0)} = \gamma. \quad (13)$$

We now generalize the Morgan–Kutzelnigg classification [53] to a D -dimensional space. Writing the interparticle wave function as

$$\Psi(u) = 1 + \frac{u}{2\kappa + D - 1} + O(u^2), \quad (14)$$

we have

$$\begin{aligned} \kappa = 0 & \text{ for natural parity singlet states,} \\ \kappa = 1 & \text{ for triplet states,} \\ \kappa = 2 & \text{ for unnatural parity singlet states.} \end{aligned} \quad (15)$$

The labels of the D -spherium states are given in Table 1.

4.3. Second-order cusp condition

The second solution is associated with

$$\Psi_{2,0}(u) = 1 + \gamma u + \frac{\gamma^2(\delta + 2)}{2\gamma(\delta + 2) + 4\delta + 6} u^2, \quad (16)$$

$$R_{2,0}^2 = \frac{(\gamma + 2)(\delta + 2) - 1}{2\gamma}, \quad (17)$$

$$E_{2,0} = \frac{\gamma(\delta + 1)}{(\gamma + 2)(\delta + 2) - 1}. \quad (18)$$

For \mathcal{D} -spherium, the second-order cusp condition is

$$\frac{\Psi''(0)}{\Psi(0)} = \frac{1}{2\mathcal{D}} \left(\frac{1}{\mathcal{D}-1} - E \right). \quad (19)$$

Following (19), the classification (15) can be extended to the second-order coalescence condition, where the wave function (correct up to second-order in u) is

$$\Psi(u) = 1 + \frac{u}{2\kappa + \mathcal{D} - 1} + \frac{u^2}{2(2\kappa + \mathcal{D})} \left(\frac{1}{2\kappa + \mathcal{D} - 1} - E \right) + O(u^3). \quad (20)$$

Thus, we have, for $\mathcal{D}=3$,

$$\frac{\Psi''(0)}{\Psi(0)} = \begin{cases} \frac{1}{6} \left(\frac{1}{2} - E \right), & \text{for } \kappa = 0, \\ \frac{1}{10} \left(\frac{1}{4} - E \right), & \text{for } \kappa = 1, \\ \frac{1}{14} \left(\frac{1}{6} - E \right), & \text{for } \kappa = 2. \end{cases} \quad (21)$$

For the natural parity singlet states ($\kappa=0$), the second-order cusp condition of 3-spherium is precisely the second-order coalescence condition derived by Tew [60], confirming that 3-spherium is an appropriate model for normal physical systems.

4.4. Third-order cusp condition

The third-order cusp condition of 3-spherium is

$$\frac{\Psi'''(0)}{\Psi(0)} = \frac{1}{18} \left(\frac{1}{8} - E + \frac{15}{16R^2} \right), \quad (22)$$

which is similar, but not strictly equivalent, to the one derived by Tew [60], due to the R -dependence of (22). The generalization to \mathcal{D} dimensions is straightforward.

4.5. Interdimensional degeneracies

As shown in Table 1, many states of \mathcal{D} -spherium have the same Hamiltonian (7) as lower angular momentum states of $(\mathcal{D}+2)$ -spherium.

Using the transformation $(\mathcal{D}, L) \rightarrow (\mathcal{D}+2, L-1)$ (see Table 1), one can see that the Hamiltonian of the ${}^3P^e$, ${}^1D^o$, ${}^3D^o$ and ${}^1F^e$ states for a given value of \mathcal{D} are respectively identical to those for ${}^1S^e$, ${}^3P^o$, ${}^1P^o$, and ${}^3D^e$ states at $\mathcal{D}+2$. The transformation $(\mathcal{D}, L) \rightarrow (\mathcal{D}+2, L-1)$, preserves the parity of the states, but ‘flips’ the spin configuration, thereby increasing by one unit the value of κ . In \mathcal{D} -spherium, we note that the Hamiltonians of the ${}^3P^e$ and ${}^3D^e$ states are identical.

Similar interdimensional degeneracies, first noticed by van Vleck [61], have been observed for various systems [44,62–65].

5. Conclusion

In this article, we have reported exact solutions of a Coulomb correlation problem, consisting of two electrons on a \mathcal{D} -dimensional sphere. The Coulomb problem can be solved exactly for an infinite set of values of the radius R for both the ground and excited states, on both the singlet and triplet manifolds. The corresponding exact solutions are polynomials in the interelectronic distance u .

The cusp conditions (up to third-order in the interelectronic distance), which are related to the behaviour of the wave function at the electron–electron coalescence point, have been analysed and classified according to the natural or unnatural parity of the state considered.

Finally, we have shown seen that, as in other one-, two- or three-electron systems, there exist interdimensional degeneracies between some of the states of \mathcal{D} -spherium.

Acknowledgements

PMWG thanks the NCI National Facility for a generous grant of supercomputer time and the Australian Research Council (Grant DP0984806) for funding.

References

- [1] A.G. Ushveridze, *Quasi-Exactly Solvable Models in Quantum Mechanics* (Institute of Physics Publishing, Bristol, 1994).
- [2] N.R. Kestner and O. Sinanoglu, *Phys. Rev.* **128**, 2687 (1962).
- [3] R.J. White and W. Byers Brown, *J. Chem. Phys.* **53**, 3869 (1970).
- [4] S. Kais, D.R. Herschbach and R.D. Levine, *J. Chem. Phys.* **91**, 7791 (1989).
- [5] M. Taut, *Phys. Rev. A* **48**, 3561 (1993).
- [6] A. Alavi, *J. Chem. Phys.* **113**, 7735 (2000).
- [7] J. Jung and J.E. Alvarillos, *J. Chem. Phys.* **118**, 10825 (2003).
- [8] J. Jung, P. Garcia-Gonzalez, J.E. Alvarillos and R.W. Godby, *Phys. Rev. A* **69**, 052501 (2004).
- [9] D.C. Thompson and A. Alavi, *Phys. Rev. B* **66**, 235118 (2002).
- [10] D.C. Thompson and A. Alavi, *Phys. Rev. B* **69**, 201302 (2004).
- [11] D.C. Thompson and A. Alavi, *J. Phys.: Condens. Matter* **16**, 7979 (2004).

- [12] D.C. Thompson and A. Alavi, *J. Chem. Phys.* **122**, 124107 (2005).
- [13] C. Filippi, C. J. Umrigar and M. Taut, *J. Chem. Phys.* **100**, 1290 (1994).
- [14] M. Taut, A. Ernst and H. Eschrig, *J. Phys. B* **31**, 2689 (1998).
- [15] K. Ivanov, K. Burke and M. Levy, *J. Chem. Phys.* **110**, 10262 (1999).
- [16] M. Seidl, J.P. Perdew and S. Kurth, *Phys. Rev. Lett.* **84**, 5070 (2000).
- [17] M. Seidl, *Phys. Rev. A* **75**, 062506 (2007).
- [18] J. Sun, *J. Chem. Theor. Comput.* **5**, 708 (2009).
- [19] P. Gori-Giorgi and A. Savin, *Int. J. Quantum Chem.* **109**, 2410 (2009).
- [20] M. Seidl and P. Gori-Giorgi, *Phys. Rev. A* **81**, 012508 (2010).
- [21] P. Hohenberg and W. Kohn, *Phys. Rev.* **136**, B864 (1964).
- [22] W. Kohn and L. Sham, *Phys. Rev.* **140**, A1133 (1965).
- [23] R.G. Parr and W. Yang, *Density Functional Theory for Atoms and Molecules* (Oxford University Press, Oxford, 1989).
- [24] W. Kutzelnigg, *Theor. Chim. Acta* **68**, 445 (1985).
- [25] W. Kutzelnigg and W. Klopper, *J. Chem. Phys.* **94**, 1985 (1991).
- [26] T.M. Henderson, K. Runge and R.J. Bartlett, *Chem. Phys. Lett.* **337**, 138 (2001).
- [27] T.M. Henderson and R.J. Bartlett, *Phys. Rev. A* **70**, 022512 (2004).
- [28] D. Bokhan, S. Ten-no and J. Noga, *Phys. Chem. Chem. Phys.* **10**, 3320 (2008).
- [29] P.-F. Loos and P.M.W. Gill, *Phys. Rev. Lett.* **103**, 123008 (2009).
- [30] G.S. Ezra and R.S. Berry, *Phys. Rev. A* **25**, 1513 (1982).
- [31] G.S. Ezra and R.S. Berry, *Phys. Rev. A* **28**, 1989 (1983).
- [32] P.C. Ojha and R.S. Berry, *Phys. Rev. A* **36**, 1575 (1987).
- [33] R.J. Hinde and R.S. Berry, *Phys. Rev. A* **42**, 2259 (1990).
- [34] P.-F. Loos and P.M.W. Gill, *Phys. Rev. A* **79**, 062517 (2009).
- [35] P.-F. Loos and P.M.W. Gill, *Phys. Rev. A* **81**, 052510 (2010).
- [36] P.-F. Loos and P.M.W. Gill, *J. Chem. Phys.* **131**, 241101 (2009).
- [37] P.-F. Loos, *Phys. Rev. A* **81**, 032510 (2010).
- [38] G. Breit, *Phys. Rev.* **35**, 569 (1930).
- [39] J.D. Louck, *J. Mol. Spectrosc.* **4**, 298 (1960).
- [40] H.A. Bethe and E.E. Salpeter, *Quantum Mechanics of One- and Two-Electron Atoms* (Dover, Mineola, New-York, 1977).
- [41] H.F. King, *J. Chem. Phys.* **46**, 705 (1967).
- [42] A. Ronveaux, *Heun's Differential Equations* (Oxford University Press, Oxford, 1995).
- [43] A.D. Polyanin and V.F. Zaitsev, *Handbook of Exact solutions for Differential Equations* (Chapman & Hall/CRC, London, 2003).
- [44] D.R. Herrick, *J. Math. Phys.* **16**, 281 (1975).
- [45] E. Wigner, *Phys. Rev.* **46**, 1002 (1934).
- [46] P.-F. Loos and P.M.W. Gill, *J. Chem. Phys.* **132**, 234111 (2010).
- [47] F. Hund, *Z. Phys.* **33**, 345 (1925).
- [48] H.N. Russell and W.F. Meggers, *Sci. Pap. Bur. Stand.* **22**, 364 (1927).
- [49] E.U. Condon and G.H. Shortley, *The Theory of Atomic Spectra* (Cambridge University Press, Cambridge, 1935).
- [50] R.J. Boyd, *Nature* **310**, 480 (1984).
- [51] J.W. Warner and R.S. Berry, *Nature* **313**, 160 (1985).
- [52] W. Kutzelnigg and J.D. Morgan III, *J. Chem. Phys.* **96**, 4484 (1992).
- [53] J.D. Morgan III and W. Kutzelnigg, *J. Phys. Chem.* **97**, 2425 (1993).
- [54] W. Kutzelnigg and J.D. Morgan III, *Z. Phys. D* **36**, 197 (1996).
- [55] T. Kato, *Trans. Am. Math. Soc.* **70**, 195 (1951).
- [56] T. Kato, *Commun. Pure Appl. Math.* **10**, 151 (1957).
- [57] R.T. Pack and W. Byers-Brown, *J. Chem. Phys.* **45**, 556 (1966).
- [58] J.D. Baker, D.E. Freund, R.N. Hill and J.D. Morgan III, *Phys. Rev. A* **41**, 1241 (1990).
- [59] P.M.W. Gill and D.P. O'Neill, *J. Chem. Phys.* **122**, 094110 (2005).
- [60] D.P. Tew, *J. Chem. Phys.* **129**, 014104 (2008).
- [61] J.H. van Vleck, *Central Fields in Two Vis-à-Vis Three Dimensions: An Historical Divertissement* (Butterworths, London, 1973), p. 26.
- [62] D.R. Herrick and F.H. Stillinger, *Phys. Rev. A* **11**, 42 (1975).
- [63] D.J. Doren and D.R. Herschbach, *J. Chem. Phys.* **85**, 4557 (1986).
- [64] D.Z. Goodson, D.K. Watson, J.G. Loeser and D.R. Herschbach, *Phys. Rev. A* **44**, 97 (1991).
- [65] M. Dunn and D.K. Watson, *Phys. Rev. A* **59**, 1109 (1999).