The spectral decomposition of the helium atom two-electron configuration in terms of hydrogenic orbitals

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Abstract
The two-electron configuration in the helium atom is known to very high precision. Yet, we tend to refer to this configuration as a $1s^\uparrow 1s^\downarrow$ singlet, where the designations refer to hydrogen orbitals. The high precision calculations utilize basis sets that are suited for high accuracy and ease of calculation, but do not really aid in our understanding of the electron configuration in terms of product states of hydrogen orbitals. Since undergraduate students are generally taught to think of helium, and indeed the rest of the periodic table, in terms of hydrogenic orbitals, we present in this paper a detailed spectral decomposition of the two-electron ground state for helium in terms of these basis states. The $1s^\uparrow 1s^\downarrow$ singlet contributes less than 93\% to the ground state configuration, with other contributions coming from both bound and continuum hydrogenic states.

(Some figures may appear in colour only in the online journal)

1. Introduction
As early as 1928, Hylleraas \cite{Hylleraas1, Hylleraas2} recognized that using hydrogenic orbitals to describe the ground state electron configuration of the helium atom was not a good idea. In the introduction to his seminal paper he says, ‘It thereby appeared that the use of hydrogen eigenfunctions, as done by Dr Biemüller, leads to erroneous results, which obviously has to do with the fact that they do not form a complete functional system.’ \cite{Hylleraas1}. He then goes on to introduce a set of coordinates and variational wavefunctions that more accurately describe the two-electron ground state of the helium atom \cite{Hylleraas1, Hylleraas3}.

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To our knowledge, the results of Dr Biemüller were never published. Most students nowadays, if asked about the electronic ground state configuration of the helium atom, will respond to the fact that it is the singlet state\(^4\) of \(1s \uparrow 1s \downarrow\), as listed, for example, in many periodic tables. Even more senior colleagues, while realizing that this simple characterization omits interactions, become disturbed at the suggestion that other orbitals enter the ground state configuration, since this appears, at first glance, to be at odds with the ‘closed-shell’ character of He, and the fact that it is an ‘inert’ element. This misunderstanding tends to be reinforced in the way we teach undergraduate quantum mechanics: since the two-electron problem is already very difficult, helium is often used as an example case study for many approximate methods, including perturbation theory and the variational method. Usually only corrections to the energy (and not the wavefunction) are described, and even with the variational method, often a simple extension of a (singlet) \(1s \uparrow 1s \downarrow\) is used. This tends to reinforce the incorrect idea that the singlet \(1s \uparrow 1s \downarrow\) hydrogenic product wavefunction is the entire picture\(^4\).

The solution to this problem attained by Hylleraas was to adopt a set of basis functions that more accurately capture the electron correlations that exist in the helium two-electron ground state. We want to emphasize here that there is no question that this works extremely well\(^5\). Nonetheless, physicists and chemists still tend to think of atomic electronic configurations in terms of hydrogenic orbitals (the periodic table guides us in this direction), and this is how our intuition is formed. Thus, in spite of the lesson of Dr Biemüller’s (erroneous) calculation, we think it is important to answer the question ‘just how much of the helium two-electron ground state consists of the singlet \(1s \uparrow 1s \downarrow\) configuration?’ and, as an obvious follow-up, ‘what other hydrogenic states contribute to the ground state configuration?’\(^5\). Most (non-expert) colleagues are shocked when the answer requires continuum states\(^6\).

Aside from the pedagogical value, the requirement of more than one electronic configuration has also become a potentially important ingredient in models of metals and superconductors within the field of condensed matter\(^6–8\). The electronic ground state of helium has served as the ‘poster child’ for atoms in a solid state environment, where the electron occupation can fluctuate. An important ingredient in electron conduction is the fact that most lattice models in condensed matter physics focus on one set of orbitals from each atomic site; these orbitals overlap to form a conduction band, and the band that crosses the Fermi energy, i.e. the energy separating occupied from unoccupied bands, is the one band of interest for low energy excitations. But this reasoning is primarily based on single-electron ideas, and as a remedy, this model (tight-binding, or Hückel) is often generalized to include electron interactions with one another, especially when two electrons occupy the same orbital on the same site (the so-called Hubbard model\(^9\)). The Hubbard model and, in general, Hubbard-like models are the subject of intensive investigation in condensed matter\(^7\). However, almost all these models miss the important ingredient that the same single-electron orbitals that are pertinent to single-electron occupation are not adequate to describe a doubly occupied situation. That is, the situation, ‘when two electrons occupy the same orbital on the same

\(^4\) We use the phrase, ‘singlet state \(1s \uparrow 1s \downarrow\)’ to refer to the spin wavefunction \(\frac{1}{\sqrt{2}}[1s \uparrow 1s \downarrow - 1s \downarrow 1s \uparrow]\).

\(^5\) We should note that this work was started without knowledge of Hylleraas’ remark concerning Dr Biemüller quoted in the opening paragraph, and we initially expected that convergence would be attained with inclusion of a few shells beyond the singlet state \(1s \uparrow 1s \downarrow\).

\(^6\) Again, for those that have reflected on this problem, the need to include continuum (or scattering) states comes as no surprise, and they may even be inclined to dismiss the entire exercise as ‘bad results due to starting with a poor basis set.’ But we reiterate, the ‘poor basis set’ with which we start is the one with which we tend to think, and therefore these questions are worth answering. Furthermore, as outlined in the subsequent discussion, in some research-type problems we are forced to adopt a poor basis set, and then the present analysis may provide some useful guidance for these problems.

\(^7\) A search through APS journals alone reveals over 5000 articles that contain the keyword ‘Hubbard’ in their title or abstract.
atom" is simply not possible, without a modification of those orbitals. In reality, two electrons that find themselves on the same atom will spread out to avoid one another, and therefore inevitably occupy (at least partially) other single-particle orbitals (like the 2s and 3s orbitals in the case of helium). This study of helium, while avoiding more complicated ideas like Wannier bases and Hartree–Fock orbitals, will emphasize this very point. As such, the two-electron configuration in the helium atom serves as a stepping stone between a standard quantum mechanical "textbook" problem and a more research-oriented set of problems.

We first outline the general problem of solving for the eigenstates of two electrons bound to a central nucleus. This will first be done using 'natural' basis states consisting of product states of the bound state hydrogenic states. This is presumably the path followed by Dr Biemüller, but since, to our knowledge, this exercise has never been published, we provide an outline of the method, with most of the details relegated to an appendix. As already remarked, this procedure does not succeed in converging to the known answers for the helium atom, for the reason that the continuum states are required, and we arrive at the surprising (for some) conclusion that an accurate description of the helium ground state cannot be provided by using just the hydrogenic bound states.

We thus turn to a more direct approach, where, starting with the exact wavefunction, one can compute the coefficients of the various basis states of interest. For helium, while the exact ground state wavefunction is known numerically to many significant digits, we instead use the so-called Hylleraas wave-function, consisting of only three variational parameters, in the interest of simplicity and transparency. This will serve as our 'exact' wavefunction, and, consistent with what we stated above, the sum of the magnitudes of the coefficients of the basis states consisting of product hydrogenic orbitals will fall short of unity, with the remainder coming from the continuum states.

While much of this discussion is suited for undergraduates, some of the mathematics is more suited to undergraduate projects and/or graduate courses. In either case we feel that this approach to the problem provides a useful connection between course work and research-type problems. We have tried to keep much of the mathematics (which we have done in part analytically and in part with software packages like Mathematica or MAPLE) in appendices.

2. The helium two-electron problem

2.1. Preliminaries

The energetics of the two-electron helium atom are very well known. Here we concern ourselves with just the non-relativistic interactions, so the Hamiltonian governing this system is

\[
H = -\frac{\hbar^2}{2m} (\nabla_1^2 + \nabla_2^2) - \frac{Ze^2}{4\pi \epsilon_0} \left( \frac{1}{r_1} + \frac{1}{r_2} \right) + \frac{e^2}{4\pi \epsilon_0 \overline{r_{12}}},
\]

where \( r_{12} \equiv |\vec{r}_1 - \vec{r}_2| \) is the separation between the two electrons and \( r_1 \equiv |\vec{r}_1| \) (\( r_2 \equiv |\vec{r}_2| \)). Note that we have already used an important approximation in writing down equation (1)—we have adopted the Born–Oppenheimer approximation, which essentially assumes that the mass of the nucleus is infinite. Hence the only degrees of freedom are those of the two electrons, each with charge \(-e\) and mass \(m\). The factor of \( Z \) in the middle term is present because in general the charge of the nucleus is \( Ze \); obviously for helium \( Z = 2 \). It is the last term, representing the

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8 This is reminiscent of the situation in general relativity, where the presence of mass alters the spacetime structure in which the mass is situated.

9 Actually the method used to determine the 'exact' ground state energy and wavefunction (e.g. in [5]) is a straightforward generalization of the simplest Hylleraas wavefunction quoted here.
electron–electron repulsion, that causes the difficulty; this is the term which is often ignored (at least conceptually) in a student’s first exposure to the helium atom (and the periodic table, for that matter) in his/her undergraduate education.

The usual procedure is to first ignore the electron–electron repulsion; the problem is then readily solved since it now consists of essentially a hydrogen problem that has to accommodate two electrons. The ground state solution is then

$$\psi_{1}(\vec{r}_{1}, \vec{r}_{2}) = \phi_{100}(\vec{r}_{1})\phi_{100}(\vec{r}_{2}),$$

(2)

which is a product state of two single-electron solutions [10, 11],

$$\phi_{n\ell m}(\vec{r}) = R_{n\ell}(r) Y_{m}^{\ell}(\theta, \phi),$$

(3)

where \((n\ell m)\) are the usual quantum numbers and \(R_{n\ell}(r)\) is the radial [angular] part of the wavefunction, which can be written in terms of associated Laguerre polynomials and spherical harmonics, respectively. These functions are tabulated in most texts; for reference we write down the ground state result needed in equation (2):

$$\phi_{100}(\vec{r}) = 2 \left( \frac{Z}{a_{0}} \right)^{3/2} \frac{1}{\sqrt{4\pi}} e^{-Zr_{1}/a_{0}}.$$  

(4)

where \(a_{0}\) is the Bohr radius and it is understood that \(Z = 2\) for helium, but we have left it general in equation (4).

As it stands, equation (2) is deceptively simple; we have left out the spin degree of freedom, and therefore we have omitted any discussion of parahelium states (symmetric spatial and antisymmetric spin wavefunction components) and orthohelium (antisymmetric spatial and symmetric spin wavefunction components). In the absence of spin–orbit coupling, these two classes of states are not coupled by the Hamiltonian (1), and one can focus on one or the other. Since we are interested in the ground state configuration, we will focus on the parahelium states—these contain the lowest energy basis state and therefore the ground state. This means that all the two-particle states that we consider below should be understood to include the singlet spin state,

$$|\chi(s_{1}, s_{2})\rangle = \frac{1}{\sqrt{2}} [|\uparrow\rangle_{1} |\downarrow\rangle_{2} - |\downarrow\rangle_{1} |\uparrow\rangle_{2}],$$

(5)

where the subscripts 1 and 2 refer to particle 1 and 2. From this point on spin is no longer included in the discussion, but its presence has indeed dictated that we treat only symmetric orbital states, of which the state (2) is one. Other examples are listed in the appendix, and will be used below.

2.2. Matrix mechanics

Because this approach was so successful in other problems, our first line of attack is to utilize matrix mechanics; we decompose the ground state into a complete set of simple well-known basis states,

$$|\psi\rangle = \sum_{j=1}^{\infty} a_{j} |\psi_{j}\rangle,$$

(6)

where \(a_{j}\) are the unknown coefficients. Inserting this into the time-independent Schrödinger equation and taking inner products with the same basis states yields the eigenvalue equation

$$\sum_{j=1}^{\infty} H_{ij} a_{j} = E a_{i},$$

(7)

10 Ironically, we use particle labels to avoid particle labelling, i.e. to enforce indistinguishability. See footnote 4 for equivalent expressions of the spin singlet state.
where the matrix elements are given by

$$H_{ij} = \langle \psi_i | H | \psi_j \rangle. \quad (8)$$

As already remarked, a complete basis set of hydrogenic orbitals consists of an infinite number of bound states (of which equations (A.1a)–(A.1f) are the first few) and an infinite continuum of unbound states, for which a much more complicated expression is required (though it is essentially the analytical continuation of the bound states). But our course will be to forge ahead, and simply truncate the expansion to include only the low-lying bound states, thus excluding a (infinite) number of bound states and all the continuum states. Such a truncation scheme worked extremely well for the harmonic oscillator [12], where, in the square well basis used for that problem, only the first ten or so states were required to give fully converged results.

One calculational advantage of this choice of basis states is that the one-electron parts of the Hamiltonian (all but the last term in equation (1)) return eigenvalues of the hydrogen spectrum when operating on these states. That is, focusing on a product, $\phi_{n_1,\ell_1,m_1}(\vec{r}_1)\phi_{n_2,\ell_2,m_2}(\vec{r}_2)$, which covers the most general case encountered in equations (A.1a)–(A.1f), we have

$$H\phi_{n_1,\ell_1,m_1}(\vec{r}_1)\phi_{n_2,\ell_2,m_2}(\vec{r}_2) = \left[-Z^2E_0 \left(\frac{1}{n_1^2} + \frac{1}{n_2^2}\right) + \hat{H}_{\text{int}}\right]\phi_{n_1,\ell_1,m_1}(\vec{r}_1)\phi_{n_2,\ell_2,m_2}(\vec{r}_2), \quad (9)$$

where

$$\hat{H}_{\text{int}} = \frac{e^2}{4\pi \varepsilon_0} \frac{1}{|\vec{r}_1 - \vec{r}_2|} \quad (10)$$
is the ‘hard part’, i.e. the electron–electron interaction potential. In equation (9) and below we use $Z = 2$ and $E_0 \equiv \hbar^2/(2ma^2) \approx 13.606$ eV. To consider an actual matrix element between any general two-electron states, we require 12 quantum numbers; hence to avoid proliferation of indices we use the shorthand $i_1 = n_1, \ell_1, m_1$. Then

$$\langle \phi_{i_1} | H | \phi_{i_2} \rangle = -Z^2E_0\delta_{i_1,1}\delta_{i_2,1} \left(\frac{1}{n_1^2} + \frac{1}{n_2^2}\right) + \langle \phi_{i_1} | H_{\text{int}} | \phi_{i_2} \rangle, \quad (11)$$

where the last matrix element requires more careful analysis; details of the calculation of this matrix element, including some examples, are provided in appendix A. There a number of states are listed (see equations (A.1a)–(A.1f)), and a software package like MAPLE or Mathematica can be tasked with evaluating all the required matrix elements up to some cutoff, $i_{\text{max}}$; the resulting finite matrix can be easily diagonalized, and we can obtain the corresponding ground state eigenvalue and eigenvector.

### 2.3. Results

In figure 1, we show the ground state energy as a function of the inverse of $i_{\text{max}}$, with square symbols, along with a guide to the eye; it is clear that the energy has essentially saturated as $i_{\text{max}}$ increases, but to a value much higher than the known exact value, which is indicated by the large (red) square at the origin. The reason for this discrepancy has already been noted: including only the bound states is not sufficient—they do not form a complete basis set, and the contribution from the continuum states is significant. At first glance, it may seem odd that continuum states contribute to the ground state for the helium atom; however, one must keep in mind that (i) the right combination of plane waves, for example, can indeed describe a very localized state and (ii) with a poor choice of basis states (which, indeed, the hydrogenic states are), the bound states will simply not be sufficient to describe the helium atom. This latter point is important, as a truncation in the (infinite) basis for the problem in [12] did not deteriorate the solution there, as the set of basis states used in that reference were sufficiently ‘good’ to
describe the problem at hand. Later we shall demonstrate the role of continuum states in this problem.

Also shown in figure 1 is the probability for the singlet $(1s \uparrow 1s \downarrow)$ basis state (right-hand scale) as a function of $1/\text{max}$ (blue asterisks). The horizontal line indicates the saturated value, about 91%, indicating that a significant fraction (close to 10%) comes from other contributions. However, because of the problem mentioned above, this value is not reliable, and we will tackle this issue in the following section.

To summarize this section, we have attempted to describe the configuration of the two-electron ground state in helium, using only the bound state hydrogenic orbitals. The reason for doing this is that these orbitals are the ones with which we have the most intuition, and they are the ones we tend to use to gain a preliminary understanding of the periodic table. Our hope was that, while this set of basis states is infinite, a finite set used through some truncation scheme would capture the essence of the ground state configuration; in reality not only did this not work well, it is clear that even if one extrapolates to the use of all the hydrogenic bound states, this incompleteness will not work.

In fact, the continuum states are required for a proper description. The reason for this is that the continuum states are able to describe high resolution spatial correlations that the bound states cannot. The bound states become more extended as their quantum numbers increase, and so any finer scale spatial correlations will have to be properly described by the continuum states. This conclusion will be reinforced in the following section, where we examine projections of the various basis states on a very accurate but simple representation of the exact two-electron wavefunction.
3. Projections

3.1. Hylleraas wavefunction: bound states

One could try to include the continuum states in the preceding calculation. To avoid an infinite matrix would require a judicious selection of these states, presumably based on their energies. In our opinion this procedure is fraught with difficulties and various choices are possible, so we do not pursue it. What we really want is a good estimate of the contribution of the ‘naive’ \((1s \uparrow 1s \downarrow)\) basis state to the actual electronic ground state of helium. The contributions from the continuum states are of importance only insofar as they influence the value \(a_1^2 \approx 0.91\) obtained in the previous section. Thus we take a different approach and find that the conclusion of the previous section is reinforced quantitatively.

Using equation (6), we can formally rewrite this as

\[
\Psi_{\text{exact}} = \sum_{m=0}^{\infty} a_m |\psi_m\rangle + \int dp a_p |\psi_p^{\text{cont}}\rangle,
\]

where now it is clear that a complete set of states is being used; the \(|\psi_m\rangle\) are those enumerated in equations (A.1a)–(A.1f) and beyond, while the \(|\psi_p^{\text{cont}}\rangle\) refer to the (as yet unspecified) continuum basis states. These states, taken together, in fact form a complete orthonormal set.

Multiplying on the left by \(\langle \psi_n |\) and forming the inner product, therefore, projects out the contribution from the \(n\)th basis state:

\[
a_n = \int \psi_n \Psi_{\text{exact}}.
\]

For equation (13) to be useful we need to know \(\Psi_{\text{exact}}\); as mentioned in the introduction, this is known numerically to many digits precision [5], using a Hylleraas-type basis [3, 13]:

\[
\Psi_{\text{exact}} = \sum_{i,j,\ell} c_{i,j,\ell} e^{-ks^{2}/2} s^{j} t^{\ell} u^{i},
\]

where the Hylleraas coordinates are defined

\[
s = (r_1 + r_2)/a_0, \quad t = (r_2 - r_1)/a_0, \quad u = r_{12} = |r_1 - r_2|/a_0
\]

and the summation occurs over all non-negative integers for \(j\) and \(\ell\), and only even non-negative integers for \(i\). This variational basis set has been expanded in a variety of ways, as summarized in [5], but in what follows we take only three terms from equation (14), with \((i, j, \ell) = (0, 0, 0), (i, j, \ell) = (2, 0, 0)\) and \((i, j, \ell) = (0, 0, 1)\). Then we define the ‘Hylleraas wavefunction’ to be the three-parameter wavefunction,

\[
\Psi_{\text{Hy}}(r_1, r_2) = \frac{2}{\pi} \left(\frac{ks}{a_0}\right)^3 e^{-Zs^2}[1 + 2Zc_1 ku + c_2 (2Zkt)^2],
\]

where \(k, c_1\) and \(c_2\) are the three parameters to be determined by minimizing the energy, and \(Z\) will eventually be set to be equal to 2. Equation (16) will serve as our ‘exact’ wavefunction. In reality, minimization of the energy with \(\Psi_{\text{Hy}}\) yields \(E_{\text{Hy}} \approx -78.979\) eV, with variational parameters \(c_1 \approx 0.0803\) and \(c_2 \approx 0.0099\) and \(k \approx 0.908\). Evaluating the necessary integrals and obtaining these results is well documented [3, 13], and so these are simply summarized in appendix B. The attained energy is within 0.05% of the exact result (−79.014 eV); we consider this sufficiently close to justify our adoption for the present purposes of \(\frac{1}{\sqrt{N}} |\Psi_{\text{Hy}}\rangle\) as the ‘exact’ normalized helium wavefunction, with \(N\) given below. The calculation of the overlap integrals for the bound states can be done for the general case—some details are provided in appendix B, along with a simple example. Our results are summarized in table 1.
Table 1. Results for some overlaps, $a_i$.

| $i$ | Basis state | $a_i$ | $|a_i|^2$ | Total Probability |
|-----|-------------|------|---------|------------------|
| 1   | 100 100     | 0.9624 | 0.9263 | 0.9263           |
| 2   | 100 200     | -0.2148 | 0.0461 | 0.9725           |
| 3   | 100 300     | -0.0752 | 0.0057 | 0.9781           |
| 4   | 100 400     | -0.0427 | 0.0018 | 0.9799           |
| 5   | 100 500     | -0.0289 | 0.0008 | 0.9807           |
| 6   | 100 600     | -0.0213 | 0.0005 | 0.9812           |
| 7   | 100 700     | -0.0166 | 0.0003 | 0.9815           |
| 8   | 101 211     | 0.0260  | 0.0007 | 0.9822           |
| 9   | 210 210     | -0.0184 | 0.0003 | 0.9825           |
| 10  | 200 300     | -0.0146 | 0.0002 | 0.9827           |
| 11  | 100 320     | -0.0090 | 0.0001 | 0.9828           |
| 12  | 100 320     | 0       | 0       | 0.9828           |

It is clear that the largest contributions arise from states in which one of the electrons is in the $\phi_{100}(r)$ state, i.e. the single-electron ground state. In particular, more than 92% comes from the $1s \uparrow \downarrow$ singlet, but a sizeable contribution comes from states other than this one. Further (small) contributions arise from states not listed. While the total probability (fifth column in Table 1) is close to unity, inclusion of the remaining bound states (not shown) still gives a total probability that falls short of 0.99. The remaining probability arises from continuum states, to which we turn in the following subsection.

3.2. Overlap integrals: continuum states

For helium, including the full spectrum of continuum two-particle states into the problem of computing the ground state configuration is very complicated. Noting that the most important contributions from the bound states arise when one of the electrons is in the one-electron ground state (see Table 1), we will likewise compute only the contributions from the continuum states when one of the two electrons is in the one-electron (bound) ground state. These states, in the singlet configuration, are written as

$$\psi_p(r_1, r_2) = \frac{1}{\sqrt{2}} (\phi_p(r_1)\phi_{100}(r_2) + \phi_{100}(r_1)\phi_p(r_2)), \quad (17)$$

where only one label (the momentum $p \equiv |\vec{p}|$) is required because the other electron is in the $1s$ state so the continuum state also has $\ell = m = 0$, and we have written the right-hand side of equation (17) as depending only on the radial coordinates, $r_1$ and $r_2$. The single-particle state $\phi_p(r)$ has a radial part $R_p(r)$ given by

$$R_p(r) = \frac{Z}{a_0} \left\{ \frac{2\pi Z \rho_0}{i} e^{-2\pi \rho_0 Z} M \left( 1 + i \frac{Z \rho_0}{p a_0}, 2, 2i\rho r \right) \right\}, \quad (18)$$

where we have written this for general $Z$ (here we need only $Z = 2$), and

$$M(a, b, z) \equiv \sum_{m=0}^{\infty} \frac{(a)_m}{(b)_m} \frac{z^m}{m!} \quad (19)$$

is the so-called Kummer function [14] and $(a)_m \equiv a(a + 1)(a + 2) \cdots (a + m - 1)$ is the so-called Pochhammer symbol. Note that $(a)_0 \equiv 1$. Equation (18) is essentially the analytical...
continuation of the radial bound state wavefunctions. The standard [15] normalization condition for the continuum states,
\[ \int_0^{\infty} dr r^2 R_{\ell p}(r) R_{\ell p}(r) = \delta(p' - p), \]  
(20)
determines the coefficient in equation (18). Just as for the bound states we require
\[ a_p = \int d^3 r_1 \int d^3 r_2 \psi_p(r_1, r_2) \frac{1}{\sqrt{N}} \psi_{\ell p}(r_1, r_2). \]  
(21)
Because of symmetry the contributions from the two terms in equation (17) are identical; the integrals involved in equation (21) can be done, either with Mathematica or by hand. Some detail is provided in appendix C. With \( y \equiv p a_0 / Z \), the result is
\[ a_p = \sqrt{\frac{\pi}{N}} \frac{32 k^3}{(k + 1)^3} \frac{a_0}{Z} \frac{y}{1 - e^{-\pi y}} \left[ (2(k + 1)^2 I_2 \right. \\
+ c_1[16kI_1 + 4k(k + 1)^2 I_3 - 4k(k + 1)J_2 - 16kJ_1] \\
+ c_2[96k^2 I_2 + 8k^2(k + 1)^2 I_4 - 48k^2(k + 1)J_3] \right]. \]  
(22)
where the expressions for \( I_n \) and \( J_n \) are provided in appendix C. Then the total contribution for these continuum states,
\[ P_{\text{cont}} = \frac{2}{\pi} \int_0^{\infty} dp |a_p|^2, \]  
(23)
is evaluated numerically; the result is a further contribution of \( P_{\text{cont}} \approx 0.0117 \), which brings the total probability from the bound states in table 1 and the continuum states shown in equation (17) to \( \approx 0.995 \). The remaining probability required to reach unity comes from bound state contributions not included in table 1 and from continuum states beyond those not considered in equation (17).

4. Summary

The primary purpose of this paper was to demonstrate the degree to which the electron configuration in the helium atom is not simply the singlet \( 1s \uparrow 1s \downarrow \) state. In fact, we have shown, through two methods, diagonalization in a particular basis set, and through projection on this same basis, that approximately 8% of the wavefunction is not the \( 1s \uparrow 1s \downarrow \) singlet. As already noted, this has a very large effect on the energy. While we mentioned in the introduction that this fact may be of interest in the current research in strongly correlated electron systems, including superconductors, the emphasis here has been on pedagogical aspects. Further exploration of the consequences of this electron configuration in a solid can be the topic of specialized student research projects.

The diagonalization of the problem in a truncated basis was, in fact, not successful at reproducing the ground state energy; in being unsuccessful, this calculation has served to highlight important pedagogical points, and it is partly for this reason that we have included a detailed analysis of the problem here. The two-electron problem in helium served to highlight that the bound hydrogenic states do not form a complete basis set, and while one might have thought that the low-lying states would be sufficient to accurately describe the ground state, this work shows that this supposition is incorrect.

Setting up the electron configuration in helium as a matrix diagonalization problem also serves to provide a concrete example of matrix mechanics, which, for undergraduates at least, is often introduced only in a formal sense, with an abstract-only exposition of the Hilbert space and expansion in basis functions, etc. This problem also serves to provide the first exposure to
a realistic ‘many-body’ problem, and how one would construct many-particle wavefunctions. (Here, two electrons are the first stepping stone in this direction beyond the single-particle problem to which undergraduates are normally exposed.)

The second method we presented, projection of an accurate wavefunction onto an orthonormal basis set of ‘known’ and well-understood wavefunctions, requires a little more sophistication, because Hylleraas wavefunctions are usually only introduced in graduate studies. Nonetheless, the three-parameter Hylleraas wavefunction given in equation (16) is sufficiently simple to be suitable for undergraduates. The ensuing algebra for the bound states is also readily accessible, particularly if projections are computed on a case-by-case basis. For example, the evaluation of equation (B.11) requires knowledge only of elementary functions and standard integrations. We have included projections involving the continuum states as well, and these require more advanced mathematics, and can certainly be skipped at the undergraduate level.

The topic of even two-electron correlations has been traditionally confined to more advanced studies at the graduate level. Part of the reason for this is that there are better and more accurate methods available after a preliminary introduction to many-body methods has been assimilated by the student. However, most students see the electronic structure of atoms in the periodic table in general, and the helium atom in particular, in terms of hydrogen orbitals, and therefore it is desirable that a description be provided in terms of these same orbitals, as we have presented in this paper.

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Appendix A. Matrix mechanics for helium

A.1. General considerations

The first six two-particle states that need to be considered (dropping the spin part given in equation (5)) are, for example,

\[ \psi_1(r_1', r_2') = \phi_{100}(r_1')\phi_{100}(r_2'), \quad m_1 + m_2 = 0 \]  
(A.1a)

\[ \psi_2(r_1', r_2') = \frac{1}{\sqrt{2}} \left[ \phi_{100}(r_1')\phi_{200}(r_2') + \phi_{100}(r_2')\phi_{200}(r_1') \right], \quad m_1 + m_2 = 0 \]  
(A.1b)

\[ \psi_3(r_1', r_2') = \frac{1}{\sqrt{2}} \left[ \phi_{100}(r_1')\phi_{21-1}(r_2') + \phi_{100}(r_2')\phi_{21-1}(r_1') \right], \quad m_1 + m_2 = -1 \]  
(A.1c)

\[ \psi_4(r_1', r_2') = \frac{1}{\sqrt{2}} \left[ \phi_{100}(r_1')\phi_{210}(r_2') + \phi_{100}(r_2')\phi_{210}(r_1') \right], \quad m_1 + m_2 = 0 \]  
(A.1d)

\[ \psi_5(r_1', r_2') = \frac{1}{\sqrt{2}} \left[ \phi_{100}(r_1')\phi_{211}(r_2') + \phi_{100}(r_2')\phi_{211}(r_1') \right], \quad m_1 + m_2 = 1 \]  
(A.1e)

\[ \psi_6(r_1', r_2') = \phi_{200}(r_1')\phi_{200}(r_2'), \quad m_1 + m_2 = 0. \]  
(A.1f)
A.2. Simplifications and selection rules

Note that all these states are symmetric, and that we have indicated the quantum number \( m_1 + m_2 \) for each two-particle state. In any state in which this is not zero, that state will not contribute to the ground state. This selection rule will be derived below, along with some other rules that eliminate more of these states. These rules provide a major simplification, and to the degree that we anticipate that maybe the \((n\ell m) = (600)\) does not contribute very much to the ground state, the number of states that may be required might be anticipated to be small. Unfortunately, this is not the case, as the continuum states contribute as well; this fact ultimately thwarts this attempt to attain a spectral decomposition in terms of these basis states [3]. Nonetheless, we proceed along this line of investigation, first to demonstrate that this is the case and secondly to establish these helpful selection rules.

To evaluate the last element matrix in equation (11) we introduce the expansion [16, 17]

\[
\frac{1}{|r_1 - r_2|} = \sum_{k=0}^{\infty} \sum_{m_{n-k}} (k - |m_k|)! (k + |m_k|)! \int_{0}^{2\pi} \int_{0}^{r_k} r_k^{l_1 |m_1|} (\cos \theta_1) P^{l_1}_{l_1} (\cos \theta_1) e^{i\phi_1} \, dr_1 \, d\phi_1,
\]

(A.2)

where \( r_1 = r_1 \) and \( r_2 = r_2 \) if \( r_1 < r_2 \) and vice versa for \( r_1 > r_2 \). The \( P^{l_1}_{l_1} \) are the associated Legendre polynomials and the angles correspond to the spherical coordinates for each of the vectors, \( r_1 \) and \( r_2 \). Equation (A.2) should become familiar to students through problems in Electromagnetism as well as in Quantum Mechanics. We can expand each of the individual hydrogenic states (see equation (3)) and use

\[
Y^m_{\ell} (\theta, \phi) = \epsilon_m \sqrt{\frac{(2\ell + 1)(\ell - |m|)!}{4\pi (\ell + |m|)!}} e^{i\ell \phi} P^m_{\ell} (\cos \theta),
\]

(A.3)

where \( \epsilon_m = (-1)^m \) for \( m \geq 0 \) and \( \epsilon_m = 1 \) for \( m \leq 0 \). If we substitute into the last term of equation (11) we obtain

\[
\langle \phi_i, \phi_i | H_{\text{int}} | \phi_i, \phi_i \rangle = \sum_{k=0}^{\infty} \sum_{m_{n-k}} (k - |m_k|)! (k + |m_k|)! (-1)^{m_1 + m_2 + m_3 + m_4 + m_5 + m_6 + m_7 + m_8} / 2
\]

\[
\times \sqrt{\frac{(2l_1 + 1)(l_1 - |m_1|)!}{(l_1 + |m_1|)!}} \frac{(2l_2 + 1)(l_2 - |m_2|)!}{(l_2 + |m_2|)!} \frac{(2l_3 + 1)(l_3 - |m_3|)!}{(l_3 + |m_3|)!} \frac{(2l_4 + 1)(l_4 - |m_4|)!}{(l_4 + |m_4|)!}
\]

\[
\times \int_{0}^{2\pi} \int_{0}^{r_k} \int_{0}^{r_k} \frac{R_{n_1,l_1}(r_1)R_{n_1,l_2}(r_2)R_{n_1,l_3}(r_1)R_{n_1,l_4}(r_2)}{r_1^3 r_2^3} \times \int_{0}^{2\pi} \int_{0}^{\pi} P^{l_1}_{l_1}( \cos \theta_1) P^{l_2}_{l_2}( \cos \theta_2) \sin \theta_1 \, d\theta_1 \, d\phi_1
\]

\[
\times \frac{1}{2} \int_{0}^{\pi} P^{l_1}_{l_1}( \cos \theta_1) P^{l_2}_{l_2}( \cos \theta_2) \sin \theta_2 \, d\theta_2 \times \frac{1}{2\pi} \int_{0}^{2\pi} e^{i(m_1 + m_2 + m_4 + m_8) \phi} \, d\phi_1 \times \frac{1}{2\pi} \int_{0}^{2\pi} e^{i(m_2 + m_4 - m_6) \phi} \, d\phi_2.
\]

(A.4)

A.2. Simplifications and selection rules

The last two integrals in equation (A.4) require \( m_k \) to be fixed; moreover, compatibility between the two leads to a condition on quantum numbers of the states that lead to a non-zero expectation value of the interaction potential:

\[
m_1 + m_2 = m_3 + m_4.
\]

(A.5)
Inspection of the states $(A.1a)$–$(A.1f)$ and those beyond shows that many states (e.g. $(A.1c)$ and $(A.1e)$) do not contribute to the ground state, and hence can be discarded from further discussion.

Furthermore, Gaunt was able to evaluate the $\theta_1$ and $\theta_2$ integrals analytically [18]. We have actually found it simpler to evaluate the integrals as they are (either numerically or analytically with the aid of MAPLE or Mathematica); nonetheless, Gaunt’s formula leads to [17]\(^{11}\) the so-called triangular condition for the angular momenta. In the first ($\theta_1$) integration in equation $(A.4)$, for example, this requires

$$\ell_1 + \ell_3 \geq k \geq |\ell_1 - \ell_3|.$$  

Thus, the $k$-sum in equation $(A.4)$ is truncated at $k_{\text{max}}$, where

$$k_{\text{max}} = \min(\ell_1 + \ell_3, \ell_2 + \ell_4).$$

It is conventional [17, 19] to introduce coefficients defined as follows:

$$c^k(\ell m; \ell' m') = (-1)^{(m+m'+|m'|+m-|m'|)/2} \sqrt{\frac{(k-|m-m'|)!}{k + |m-m'|!}} \frac{(2\ell + 1)(\ell - |m|)!}{(\ell + |m|)!} \frac{(2\ell' + 1)(\ell' - |m'|)!}{(\ell' + |m'|)!} \times \frac{1}{2} \int_{-1}^{1+1} P_{\ell m}(\mu) P_{\ell' m'}^*(\mu) d\mu.$$ (A.8)

In terms of these coefficients the interaction matrix element can be written as

$$\langle \phi_{\ell_1} \phi_{\ell_2} | H_{\text{int}} | \phi_{\ell_3} \phi_{\ell_4} \rangle = \frac{e^2}{4\pi \epsilon_0} \delta_{m_1+m_2,m_3+m_4}$$

\[ \times \sum_{k=0}^{k_{\text{max}}} c^k(\ell_1 m_1; \ell_3 m_3)c^k(\ell_2 m_2; \ell_4 m_4) R_1^k(n_1 \ell_1, n_2 \ell_2; n_3 \ell_3, n_4 \ell_4), \]  

where

$$R_1^k(n_1 \ell_1, n_2 \ell_2; n_3 \ell_3, n_4 \ell_4) = \int_0^\infty \int_0^\infty R_{n_1 \ell_1}(r_1) R_{n_2 \ell_2}(r_2) R_{n_3 \ell_3}(r_1) R_{n_4 \ell_4}(r_2) \frac{r_1^{n_1} r_2^{n_2}}{\Gamma_{n_1+1} \Gamma_{n_2+1}} d^2r_1 d^2r_2.$$ (A.10)

This last double integration can be readily done by hand (though it is tedious) or can be done (symbolically) with Mathematica or MAPLE, since the radial wavefunctions can be readily expressed in terms of associated Laguerre polynomials [11]. Also note the reversed order of the arguments in the second $c^k$ in equation $(A.9)$; this is important since

$$c^k(\ell m; \ell' m') = (-1)^{(m'-|m|)} c^k(m'; \ell m).$$ (A.11)

Inspection of the states $(A.1a)$–$(A.1f)$ and those beyond not already omitted by condition $(A.5)$ indicates that three distinct possibilities remain (e.g. when $(n_1 \ell_1 m_1) = (n_2 \ell_2 m_2)$ and $(n_3 \ell_3 m_3) = (n_4 \ell_4 m_4)$, etc), but all of these can be handled through equation $(A.9)$.

One other selection rule is present in these results, though not readily apparent. In equation $(A.9)$, the same value of the index $k$ must work for both $c^k$ coefficients. These coefficients have been tabulated, for example, in the text by Slater [17], where it is clear that two even or two odd angular momenta (referring to $\ell$ and $\ell'$) couple to one another only through even values of $k$, while an even and an odd angular momentum couple to one another only through an odd value of $k$. For example $\ell = 0$ and $\ell' = 0$ (ss) results in a non-zero $c^k$ coefficient only if $k = 0$, while $\ell = 0$ and $\ell' = 2$ (sd) yields a non-zero $c^k$ coefficient only if $k = 2$; the case $\ell = 1$ \(^{11}\) The required formula is found on page 308.
and \( t' = 1 \) (pp) has a non-zero \( c^1 \) coefficient if \( k = 0 \) or \( k = 2 \). In contrast the sp, sf and pd coefficients are non-zero for \( k = 1 \), \( k = 3 \) and \( k = 1, 3 \), respectively. This means that \( \ell_1 + \ell_3 \) has to have the same parity as \( \ell_2 + \ell_4 \). Since the parity of the \( 1s \uparrow 1s \downarrow \) state is even, then, for example, we can further discard state (A.1d) from the list, leaving only three of the original six states listed.

### A.3. An example

By way of example we quote some steps for the first state (not listed) that does not utilize single-particle s states,

\[
\psi_{14}(\vec{r}_1, \vec{r}_2) = \frac{1}{\sqrt{2}}[\phi_{211}(\vec{r}_1)\phi_{211}(\vec{r}_2) + \phi_{21-1}(\vec{r}_2)\phi_{211}(\vec{r}_1)]. \tag{A.12}
\]

To compute the matrix element that couples the first (and primary) basis state with this one, we need to evaluate the overlap integral,

\[
H_{1,14} = \langle \psi_1 | H_{\text{int}} | \psi_{14} \rangle. \tag{A.13}
\]

This consists of two overlaps (equation (A.12) has two terms), but, through a change of variables \( \vec{r}_1 \leftrightarrow \vec{r}_2 \), these are readily seen to be equal to one another. We are thus left with

\[
H_{1,14} = \sqrt{2}(\phi_{100}\phi_{100}|H_{\text{int}}|\phi_{21-1}\phi_{211}). \tag{A.14}
\]

Equation (A.6) tells us that \( 1 \geq k \geq 1 \), i.e. only \( k = 1 \) needs to be considered. Then a straightforward evaluation of \( c^1(00; 1 - 1) \) and \( c^1(11; 00) \) (as required in equation (A.9)) gives us \(-1/\sqrt{3}\) for the first and \(+1/\sqrt{3}\) for the second (using equation (A.11)). Next only one double radial integral (equation (A.10) for \( k = 1 \)) is required, and a straightforward evaluation, readily done by hand, gives

\[
R^1(10; 10; 21; 21) = \frac{112}{2187} \frac{Z}{a_0}. \tag{A.15}
\]

Combining this with the \( c^1 \)'s and the \( \sqrt{2} \) from equation (A.14) gives

\[
H_{1,14} = -\sqrt{2} \frac{448}{6561} E_0, \tag{A.16}
\]

where \( E_0 \equiv \frac{h^2}{2ma_0^2} \approx 13.606 \text{ eV} \) is adopted as our unit of energy, and we have used \( a_0 = \frac{4\pi\varepsilon_0}{\epsilon} \frac{h^2}{m} \).

In practice, we have written a program in MAPLE to perform these tasks, for the bound basis states, up to some cutoff. That is, all the matrix elements, \( H_{ij} \), up to some cutoff, \( i_{\text{max}} \), are evaluated, and then this matrix is diagonalized. The cutoff \( i_{\text{max}} \) is defined as the ‘\( n \)’ quantum number up to which all states are included. For example, if \( i_{\text{max}} = 2 \), then only five basis states are considered. The rest of the states either do not contribute to the ground state, or, if one of the hydrogenic single-particle states has \( n = 3 \), only contributes to the next shell (\( i_{\text{max}} = 3 \)) and beyond. Matrix elements for the case \( i_{\text{max}} = 2 \) are tabulated in the following section.

### A.4. Matrix equation for \( i_{\text{max}} = 2 \)

If we restrict basis states in section 2 to those with \( n = 2 \) or less, only five two-particle states need to be considered; referring to equations (A.1a)–(A.1f); these are \( \psi_1, \psi_2, \psi_6, \psi_{14} \) and \( \psi_{16} \), where \( \psi_{14} \) is given in equation (A.12), and

\[
\psi_{16}(\vec{r}_1, \vec{r}_2) = \phi_{210}(\vec{r}_1)\phi_{210}(\vec{r}_2). \tag{A.17}
\]
Writing the wavefunction in terms of these wavefunctions alone gives rise to a $5 \times 5$ matrix diagonalization problem (see equation (7)). The resulting equation is

\[
\begin{bmatrix}
\frac{1}{2} & -\frac{12768}{64827} & -\frac{64}{729} & -\frac{448}{6561} \sqrt{2} & -\frac{448}{6561} \\
-\frac{32768}{64827} & \frac{2969}{729} & -\frac{4096}{6561} & \frac{2048}{28125} \sqrt{2} & \frac{2048}{28125} \\
-\frac{64}{729} & -\frac{4096}{6561} & \frac{179}{128} & -\frac{15}{128} \sqrt{2} & -\frac{15}{128} \\
-\frac{448}{6561} \sqrt{2} & \frac{2048}{28125} \sqrt{2} & -\frac{15}{128} \sqrt{2} & \frac{47}{30} & -\frac{27}{6561} \sqrt{2} \\
-\frac{448}{6561} & \frac{2048}{28125} & -\frac{15}{128} & -\frac{27}{6561} \sqrt{2} & \frac{779}{6561} \\
\end{bmatrix}
\begin{bmatrix}
a_1 \\
a_2 \\
a_7 \\
a_{14} \\
a_{16}
\end{bmatrix}
= E
\begin{bmatrix}
a_1 \\
a_2 \\
a_7 \\
a_{14} \\
a_{16}
\end{bmatrix},
\]  
(A.18)

where $E_0 \approx 13.606$ and the subscripts on the coefficients correspond to the labels in the wavefunctions. The resulting ground state energy is $E_1 = -77.13$ eV, and the eigenvector has components

\[
\begin{align*}
a_{100,100} &= 0.9520 \\
a_{100,200} &= -0.3040 \\
a_{200,200} &= -0.0146 \\
a_{21-1,211} &= 0.0267 \\
a_{210,210} &= -0.0188 .
\end{align*}
\]  
(A.19)

Clearly, the $\psi_{1(100,100)}$ basis state dominates, and small adjustments occur as the number of basis states increases. Nonetheless, almost 10% of the wavefunction is comprised of components beyond the $1s \uparrow 1s \downarrow$ state. The matrix construction and diagonalization indicated here is repeated, with the aid of the software package MAPLE, for increasing values of $I_{\text{max}}$, and the results from these calculations are reported in the text.

### Appendix B. Hylleraas wavefunction

#### B.1. The expectation value of the energy

For reference, the expectation value of the energy, using the normalized Hylleraas wavefunction $\frac{1}{\sqrt{N}} |\psi_{H} \rangle$ with $|\psi_{H} \rangle$ given by equation (16), is

\[
E_{\text{Hy}} = -\frac{\hbar^2}{2m a_0^2} \langle kZ \rangle^2,
\]  
(B.1)

where $k$ can be determined analytically in terms of $c_1$ and $c_2$ (see equation (B.4), below),

\[
kZ = \frac{2L - L'}{2M},
\]  
(B.2)

and the unit of energy is the Rydberg,

\[
\frac{\hbar^2}{2m a_0^2} \equiv 1 \text{ Ryd} \approx 13.606 \text{ eV}.
\]  
(B.3)

The parameters $L, L', M$ and $N$, corresponding to different parts of the Hamiltonian, are given by

\[
\begin{align*}
L &= 4 + 30c_1 + 48c_2 + 72c_1^2 + 280c_1c_2 + 576c_2^2 \\
L' &= 5/4 + 8c_1 + 9c_2 + (35/2)c_1^2 + 48c_1c_2 + 78c_2^2 \\
M &= 2 + (25/2)c_1 + 24c_2 + 32c_1^2 + 146c_1c_2 + 480c_2^2 \\
N &= 4 + 35c_1 + 48c_2 + 96c_1^2 + 308c_1c_2 + 576c_2^2.
\end{align*}
\]  
(B.4)

Aside from units (most treatments use the so-called chemistry units), these all agree with results in the literature [3, 13]. Use of the optimally determined parameters, $c_1 \approx 0.0803$ and $c_2 \approx 0.0099$, and $k \approx 0.908$ determines the energy via equation (B.1).
B.2. Overlap integrals: bound states

With a very accurate wavefunction in hand, we can simply utilize equation (13) to determine the probability of each basis state in the ground state wavefunction. The general hydrogenic bound state can be written as

\[ \phi_{n,l,m}(r, \theta, \phi) = G_{n,l} e^{-Zr/na} r^l I_{n-\ell-1} \left( \frac{2Zr}{na} \right) Y^m_{\ell}(\theta, \phi), \]

where

\[ G_{n,l} = \left( \frac{2Z}{na} \right)^3 \frac{(n-\ell-1)!}{2n! (n+\ell)!} \left( \frac{2Z}{na} \ell \right), \]

and the required overlap integrals consist only of terms of the form

\[ I = \int \phi^*_{n_1, l_1, m_1} \phi^*_{n_2, l_2, m_2} \frac{1}{\sqrt{N}} \Psi_{Hy} \]

In the \( u, s, t \) coordinates defined above, and using the volume element,

\[ d\tau = \frac{\alpha^6}{8} (s^2 - t^2) u \, ds \, dr \, d\sin \theta_1 \, d\theta_1 \, d\phi_1 \, d\phi_2, \]

many of these integrals are straightforward to calculate analytically—an example will be shown below. However, it is of interest to compute these overlaps for general quantum numbers, in particular to check on selection rules, and to examine the convergent behaviour for large \( n_1 \) and \( n_2 \). To this end we use the same expansion, equation (A.2) used earlier to evaluate matrix elements of the Hamiltonian, and then evaluate equation (B.7). The details of the derivation for a general matrix element are a little too cumbersome to include here; all of the overlaps involving bound state wavefunctions can be expressed in terms of hypergeometric functions, which are readily evaluated using Mathematica, and with these we can readily sum the contributions from the bound states.

For students, however, it is more instructive to compute this overlap ‘by hand’ for a few of the most relevant states. We outline the procedure for the most important state, \( \psi_1(\vec{r}_1, \vec{r}_2) = \phi_{100}(\vec{r}_1)\phi_{100}(\vec{r}_2) \). Then

\[ a_1 = \int d^3r_1 \int d^3r_2 \phi_{100}(\vec{r}_1)\phi_{100}(\vec{r}_2) \frac{1}{\sqrt{N}} \Psi_{Hy}(\vec{r}_1, \vec{r}_2), \]

and the only angular dependence occurs in the \(|\vec{r}_1 - \vec{r}_2|\) term in the Hylleraas wavefunction. This requires the knowledge of the angle between the two vectors, \( \vec{r}_1 \) and \( \vec{r}_2 \), which we can call \( \theta_{12} \); this is given as [3]

\[ \cos \theta_{12} = \cos \theta_1 \cos \theta_2 + \sin \theta_1 \sin \theta_2 \cos (\phi_2 - \phi_1), \]

where \( \vec{r}_i = (r_i, \theta_i, \phi_i) \), for \( i = 1, 2 \) are the spherical coordinates for each vector. A simpler trick is to line up the \( z_2 \) axis with \( \vec{r}_1 \); then \( \theta_{12} \equiv \theta_2 \), and all six integrations are then straightforward. The result is

\[ a_1 = \frac{32 \alpha^6}{\sqrt{N} (k + 1)^6} \left[ 4 + 35c_1 \frac{k}{k + 1} + 96c_2 \left( \frac{k}{k + 1} \right)^2 \right], \]

\[ \approx 0.9624, \]

and therefore \( a_1^2 \approx 0.9262 \). The numerical values follow upon substitution of the optimal values of \( k, c_1 \) and \( c_2 \). Similar calculations can be performed for the other overlap coefficients; some results are tabulated in table 1, where the constituents of the basis states, suitably symmetrized, are listed. Note that the label provided under the ‘basis state’ column identifies the two single-particle wavefunctions involved, and the basis state is a singlet state, and therefore the spatial part is symmetrized, as enumerated in equations (A.1a)–(A.1f).
Appendix C. Evaluation of overlap integrals

We provide some of the details for the evaluation of equation (21). Because of the symmetry in the singlet state, and using the definition

$$a_p = \sqrt{\frac{2}{2\pi}} \left( \frac{kZ}{a_0} \right)^3 A_p$$  \hspace{0.5cm} (C.1)

we require the integral

$$A_p = \int d^3 r_1 \int d^3 r_2 \phi_p(r_1) \phi_{100}(r_2) e^{-Zk a_0 (r_1 + r_2)}$$

$$\times \left[ 1 + 2Zc_1 \frac{k}{a_0} |\vec{r}_1 - \vec{r}_2| + c_2 \left[ 2Z \frac{k}{a_0} (r_1 - r_2) \right]^2 \right]^3,$$  \hspace{0.5cm} (C.2)

where

$$\phi_p(r_1) = \frac{1}{\sqrt{4\pi}} \frac{Z}{a_0} \sqrt{\frac{2\pi}{pa_0}} e^{-iyp} M \left( 1 + \frac{Z}{pa_0}, 2, 2ipr \right),$$  \hspace{0.5cm} (C.3)

and $M(a, b, z)$ is the Kummer function [14]. Note that

$$\int_0^\pi d\theta_1 \sin \theta_1 \int_0^\pi d\theta_2 \sin \theta_2 \int_0^{2\pi} d\phi_1 \int_0^{2\pi} d\phi_2 [1] = (4\pi)^2,$$  \hspace{0.5cm} (C.4)

while

$$\int_0^\pi d\theta_1 \sin \theta_2 \int_0^\pi d\theta_2 \sin \theta_2 \int_0^{2\pi} d\phi_1 \int_0^{2\pi} d\phi_2 |\vec{r}_1 - \vec{r}_2| = (4\pi)^2 \left( r_+ + \frac{1}{3} r_\times \right),$$  \hspace{0.5cm} (C.5)

where $r_+$ ($r_\times$) refers to the smaller (larger) of $r_1$ or $r_2$. The first of these is trivial, while the second follows most readily by using the trick mentioned following equation (B.10).

Two integrations remain, and it is efficient to switch to dimensionless variables, $x_1 \equiv Zr_1/a_0$, $x_2 \equiv Zr_2/a_0$ and $y \equiv pa_0/Z$. Then, since the Kummer function depends only on $x_1$, the $x_2$ integration is elementary, and leaves behind polynomials in $x_1$. One arrives at the expression given by equation (22), with the definitions

$$I_n(y) \equiv B_n(y, k + iy) \quad \text{and} \quad J_n(y) \equiv B_n(y, 2k + 1 + iy),$$  \hspace{0.5cm} (C.6)

where

$$B_n(y, z) \equiv \int_0^\infty dx x^n e^{-\frac{z}{x}} M \left( 1 + \frac{i}{y}, 2, 2iyx \right).$$  \hspace{0.5cm} (C.7)

Now an expansion of the Kummer function (following equation (19)) allows us to do the integral and obtain an infinite summation which can be recognized as a hypergeometric function [14],

$$B_n(y, z) = \frac{n!}{z^{n+1}} F \left[ 1 + \frac{i}{y}, n + 1; \frac{2iy}{z} \right],$$  \hspace{0.5cm} (C.8)

with $n = 1, 2, 3, 4$ needed. The standard definition of the hypergeometric function uses the Gauss hypergeometric series [14], with the circle of convergence in the unit circle $|z| = 1$:

$$F(a, b; c; z) = \sum_{m=0}^\infty \frac{(a)_m(b)_m z^m}{(c)_m m!},$$  \hspace{0.5cm} (C.9)
where the \((a)_n\) are the Pochhammer symbols introduced earlier. Writing equation (C.8) would not normally be too useful (because hypergeometric functions are hard to evaluate numerically), except that one can use the identity [14]

\[
F[a, b; c; z] \equiv (1 - z)^{c-a-b}F[c - a, c - b; c; z].
\]

(C.10)

This identity is extremely helpful because the second variable in the hypergeometric function on the right-hand side is a non-positive integer for \(n = 1, 2, 3, 4\) and so, because of the definition of the Pochhammer symbol, the infinite sum in the definition of the hypergeometric function becomes truncated to \(n\) terms. We thus obtain, for \(n = 1, 2, 3, 4\),

\[
I_n(y) = \frac{1}{k^2 + y^2} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{k} \right) D_n
\]

and

\[
J_n(y) = \frac{1}{(2k + 1)^2 + y^2} \exp \left( -\frac{2}{y} \tan^{-1} \left( \frac{y}{2k + 1} \right) \right) E_n.
\]

(C.11, C.12)

where

\[
D_n = \frac{n!}{(k - iy)^{n-1}} F \left[ 1 - \frac{i}{y}, 1 - n; 2, \frac{2iy}{k + iy} \right].
\]

and

\[
E_n = \frac{n!}{(2k + 1 + iy)^{n-1}} F \left[ 1 - \frac{i}{y}, 1 - n; 2, \frac{2iy}{2k + 1 + iy} \right].
\]

(C.13, C.14)

With these definitions, straightforward evaluation gives

\[
D_1 = 1
\]

\[
D_2 = \frac{2(k - 1)}{k^2 + y^2}
\]

\[
D_3 = \frac{4(k - 1)(2k - 1)}{(k^2 + y^2)^2} - \frac{2}{k^2 + y^2}
\]

\[
D_4 = \frac{8(k - 1)(2k - 1)(3k - 1)}{(k^2 + y^2)^3} - \frac{8(3k - 2)}{(k^2 + y^2)^2}
\]

(C.15)

and

\[
E_1 = 1
\]

\[
E_2 = \frac{4k}{(2k + 1)^2 + y^2}.
\]

(C.16)

Summarizing, we have

\[
I_1 = \frac{1}{k^2 + y^2} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{k} \right)
\]

\[
I_2 = \frac{2(k - 1)}{(k^2 + y^2)^2} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{k} \right)
\]

\[
I_3 = \frac{4(k - 1)(2k - 1)}{(k^2 + y^2)^3} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{k} \right)
\]

\[
I_4 = \frac{8(k - 1)(2k - 1)(3k - 1)}{(k^2 + y^2)^4} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{k} \right)
\]

\[
J_1 = \frac{1}{(2k + 1)^2 + y^2} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{2k + 1} \right)
\]

\[
J_2 = \frac{4k}{(2k + 1)^2 + y^2} \exp \left( -\frac{2}{y} \tan^{-1} \frac{y}{2k + 1} \right)
\]

(C.17)

and these are to be substituted into equation (22).
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