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Abstract
We report on the first part of a study of electron-hydrogen scattering, using a method [Bhatia, Schneider, and Temkin, Phys. Rev. Lett. 70, 1936 (1993)], which allows for the ab initio calculation of total and elastic cross sections at higher energies. In its general form the method uses complex "radial" correlation functions, in a (Kohn) T-matrix formalism. The titled method, abbreviated CCKT, is reviewed, in the context of electron-hydrogen scattering, including the derivation of the equation for the (complex) scattering function, and the extraction of the scattering information from the latter. The calculation reported here is restricted to S-waves in the elastic region, where the correlation functions can be taken, without loss of generality, to be real. Phase shifts are calculated using Hylleraas-type correlation functions with up to 95 terms. Results are rigorous lower bounds; they are in general agreement with those of Schwartz [Phys. Rev. 124, 1468 (1961)], but they are more accurate and outside his error bounds at a couple of energies.
I. INTRODUCTION

In a previous paper [1], we developed a method for calculating elastic and total cross sections for electron-atom scattering. Called "complex correlation Kohn T", and abbreviated (CCKT) the method is intended for non-elastic scattering in low and medium energy range, by which we mean the energy domain where all the inelastic channels are open plus the continuum of ionization channels (up to a total available energy $E$, which is, say, less than five times the continuum threshold). Clearly no method can specify all such open channels individually, and usual approximations such as the Born and eikonal methods [2] are only reliable at high energies. Because our method is intended for the low continuum, it is a partial wave method where the partial wave expansion can still be expected to converge reasonably effectively.

The original formulation [1] was applied to an approximation of electron-hydrogen (e-H) scattering called the spherically symmetric model [3]. The S-wave part of that model reduces to the Temkin-Poet model [4,5], and it was only for the latter that the actual calculation was carried out. (The T-P model was later generalized in a different and more incisive way in what was called the "generalized exchange approximation" [6]).

In this paper we deal with the real (i.e. physical) e-H problem. The method is reviewed in Sec II. Specifically a more detailed derivation of the (complex) optical potential and scattering equations is given. As opposed to Ref. [1], the emphasis here is on the numerical solution of the scattering equation by decomposing it into its coupled real and imaginary parts and from the solution extracting the (non-unitary) $T$-matrix.

An actual calculation in its full generality, however, is still a very extensive undertaking. We shall therefore confine ourselves in this initial study, to elastic S-wave scattering. Here our aim is to check and increase the accuracy of the classic calculation of Schwartz [7]. Results are presented in Sec. III. Suffice it here to say that our results are rigorous lower bounds on the phase shifts. Thus where they exceed Schwartz' results, which they do at two energies, they are necessarily more accurate. In general our results are within the stated uncertainty of his remarkable calculation (considering when it was done). Here, given the convergence of our results, we would only claim greater accuracy.

II. THE COMPLEX CORRELATION KOHN T (CCKT) METHOD

A. General

Confined to the e-H partial wave (denoted by $L$) problem, the total spatial function is written:

$$
\Psi_L^{(\pm)}(r_1, r_2) = \frac{u_L^{(\pm)}(r_1)}{r_1} Y_{L0}(r_1 \phi_1) \phi_{10}(r_2) Y_{00}(r_2) \pm (1 \leftrightarrow 2) + \Phi_L^{(\pm)}(r_1, r_2)
$$

The superscript $(\pm)$ above refers to singlet (upper sign) or triplet (lower sign) scattering respectively. [That superscript will be dropped hereinafter, except where it is necessary.] Beyond the terms containing $u_L$ explicitly (those are the terms giving rise to the exchange approximation), the function $\Phi_L$ is the correlation function, which is our concern here. For arbitrary $L$ this function is most efficiently written in terms of symmetric Euler angles [8]:

$$
\Phi_L = \sum_{\epsilon} \sum_{\epsilon'} \left[ f_L^{\epsilon\epsilon'}(r_1, r_2, r_{12}) D_{L}^{\epsilon\epsilon'}(\theta, \phi, \psi) + f_L^{\epsilon'\epsilon}(r_2, r_1, r_{12}) D_{L}^{\epsilon'\epsilon}(\theta, \phi, \psi) \right]
$$

The $D^{\epsilon\epsilon'}$ functions $(\epsilon = \pm 1, -1)$ are exchange adapted combinations of Wigner functions. (The $D$ functions, in general have been called rotational harmonics [9].) The $f$'s above are generalized
"radial" functions, which depend on the 3 residual coordinates that are required (beyond the Euler angles) to define the two vectors $r_1$ and $r_2$ (Further technical details of the symmetry and other properties of the symmetric Euler angle description are given in [8]). What distinguishes this CCKT approach is the fact that the "radial" functions are taken to be complex. That is different from ordinary variational methods, wherein the radial functions are taken to be real. In the latter case, when one forms the variational functional to be varied [cf. Eq.(5) below], one uses the function $\Psi_L^*$ on the left. That is so because the $f$ functions would be real, so that only the $D$ functions in (2) would be complex conjugated. When the radial functions are complex, however, then - in order to have a variational principle for the $T$ matrix - only the $D$ but not the radial functions are to be complex conjugated in the left-hand function. That is indicated by using the tilde rather than star for that function, which is used in the functional $I_L$ of the variational principle (below). Explicitly

$$\tilde{\Psi}_L = \Psi_L - \Phi_L + \tilde{\Phi}_L$$

where

$$\tilde{\Phi}_L = \sum \sum \sum \left[ f_L^{\epsilon^*}(r_1, r_2, r_{12}) (D_L^{\epsilon^*}(\theta, \phi, \psi))^* + f_L^{\epsilon^*}(r_2, r_1, r_{12}) (D_L^{\epsilon^*}(\theta, \phi, \psi))^* \right]$$

The Kohn variational principle [10] for $T_L$ reads:

$$\delta \left[ I_L - (-1)^L T_L \right] = 0$$

where $I_L$ is the (Kohn) functional

$$I_L = \int [\tilde{\Psi}_L (H - E) \Psi_L] dr_1 dr_2$$

Carrying out the variation leads to scattering equation for $u_L^{(\pm)}(r)$ (Rydberg units throughout):

$$\left[ -\frac{d^2}{dr^2} - \frac{L(L+1)}{r^2} + V_d \pm V_{ex} + V_{op}^{(\pm)} - k^2 \right] u_L^{(\pm)} = 0$$

$V_d$ and $V_{ex}$ are the well known direct and exchange potentials of the "exchange approximation" [11]. Those potentials (the latter being non-local) are real, however the (remaining) optical potential acting on $u_L(r)$ is complex:

$$V_{op} u_L = r \left( Y_{L0} PHQ \frac{1}{E - QHQ} QHP \Psi_L \right),$$

Thus, $u_L(r)$ is nontrivially complex, which leads to a nonunitary $T_L$ matrix, which is derivable from the asymptotic form of $u_L$

$$\lim_{r \to \infty} u_L(r) = \frac{i^L \sin(kr - \pi L/2)}{k} + T_L e^{ikr}$$

In defining the optical potential we have used the Feshbach approach [12], involving projection operators $P$ and $Q$, which for the hydrogenic (i.e. one-electron) target can be written explicitly [13]
\[ P = P_1 + P_2 - P_1 P_2 \]
\[ Q = 1 - P \]

where the spatial projectors are:

\[ P_i = e^{-r_i} Y_{00}(r_i) \langle < e^{-r_i} Y_{00}(r_i) \]

Note, \( P_1 \) and \( P_2 \) commute and are each idempotent, hence the complete \( P \) and \( Q \) operators are idempotent (\( P^2 = P \); \( Q^2 = Q \)) and orthogonal (\( PQ = 0 \)).

To complete this review of the CCKT method, the optical potential is expanded in terms of the eigenspectrum of the \( QHQ \) problem:

\[ \delta \left[ \frac{\langle \Phi_L QHQ \Phi_L \rangle}{\langle \Phi_L Q \Phi_L \rangle} \right] = 0 \]

This leads to complex radial eigenfunctions \( \Phi_L^{(s)} \) and complex eigenvalues \( \mathcal{E}_s \). Inserting a complete set of these functions (understood to be approximated by a discrete set arising from (13) using an ansatz with a finite number of variational parameters) allows the optical potential to be expanded:

\[ V_{op} u_L(r_1) = r_1 \sum_{s}^{N \omega} \frac{Y_{L0}(r_1) \phi_{10}(r_2) \frac{2}{r_{12}} Q \Phi_L^{(s)} > < Q \Phi_L^{(s)} \frac{2}{r_{12}} P \Psi_L >}{E - \mathcal{E}_s} \]

To repeat the main point of the CCKT method: because the radial functions \( f_L^{p,c} (r_1, r_2, r_{12}) \) in (2) are complex, the resulting \( T_L \) matrix (which in this method is actually a number) will be complex and nonunitary. This means that the elastic cross section, calculated from

\[ \sigma_L = 4\pi (2L + 1)|T_L|^2 \]

will be different from the total cross section

\[ \sigma_L = \frac{4\pi}{k} (2L + 1) \text{Im}(T_L) \]

The effect of a nonunitary \( T_L \), is of course well known from the literature (cf., for example Ref.[14]). What is new about the CCKT is that it provides an \textit{ab initio} method for calculating \( T_L \), rather than parametrizing it, as was done for example in the "clouded crystal ball" model of neutron-nucleus scattering and reactions [15].

B. Some Details of the CCKT Method

The Kohn variational principle is generally applied in such a way, as in Ref.[1], that the function \( u_L(r) \) of Eq.(7) is parametrized by an expansion in terms of known functions, and the functional in Eq.(6) is analytically evaluated; it is a bilinear expression in terms of the (linear) expansion coefficients. The (Kohn) principle then reduces to differentiation with respect to these parameters,
which in turn leads to a set of linear equations, the last being an inhomogeneous equation from the
differentiation with respect to $T_L$. The solution of the set of equations then provides values for all
the (linear) constants plus the value of $T_L$. The latter can be improved by subtracting the value
of the functional itself, which in general will not be zero when the calculation is performed in this
way. In that case an improved $T_L$ can be obtained by subtraction:

$$T_L^{\text{improved}} = T_L - I_L$$

When, however, the calculation is carried out (as it is here) by solving directly for $u_L(r)$, then this
is equivalent to making $I_L = 0$. Thus the $T_L$ that emerges is automatically the improved (i.e. the
second order) result. What we shall now do is to outline briefly how the calculation is performed
when the potentials and hence the solutions are complex. Specifically, the radial equation can be
written:

$$\left[ \mathcal{L}_L^{(\pm)} + V_{op}^{(\pm)} \right] u_L^{(\pm)}(r) = 0$$

where

$$\mathcal{L}_L^{(\pm)} = -\frac{d^2}{dr^2} + \frac{L(L+1)}{r^2} + V_d \pm V_{ex} - k^2$$

The $V_d$ is the (local) Hartree potential [$V_d(r) = -2\exp(-2r)(1 + 1/r)$] and $V_{ex}$ is the well-known
(non-local) static exchange potential [11]. The remaining (non-local) potential is what is generally
called the optical potential, Eq.(5).

We now give a few details of how one solves the radial equation (18) in the general case. Dropping
sub (super) scripts, we write the solution of (4)

$$u(r) = u_R(r) + iu_I(r)$$

where $u_R(r)$ and $u_I(r)$ are real functions which, substituted in (18), lead to a coupled set of (real)
differential equations.

$$(\mathcal{L} + V_{op}^{(R)})u_R = V_{op}^{(I)}u_I$$

$$(\mathcal{L} + V_{op}^{(R)})u_I = -V_{op}^{(I)}u_R$$

$V_{op}^{(R)}$ and $V_{op}^{(I)}$ above are the real and imaginary parts of the optical potential, respectively. (Both
are real, non-local potentials). The coupled Eqs.(21, 22) have two sets of solutions (i.e. both regular
at the origin), labeled $u_R^{(i)}(r)$, $u_I^{(i)}(r)$ [$i = 1,2$]. From these two solutions one determines two real
constants $C_1$ and $C_2$ such that the real and imaginary parts of the asymptotic form (9) is satisfied,
that the equation for $L=0$ reduces to

$$\lim_{r \to \infty} \left[ C_1 u_R^{(1)}(r) + C_2 u_R^{(2)}(r) \right] = \frac{\text{sink}r}{k} + T_R \cos kr - T_I \sin kr$$

$$\lim_{r \to \infty} \left[ C_1 u_I^{(1)}(r) + C_2 u_I^{(2)}(r) \right] = T_I \cos kr + T_R \sin kr$$
The two linearly independent solutions will necessarily have the asymptotic form

$$\lim_{r \to \infty} u_R^{(i)} = A_R^{(i)} \sin(kr + \eta_R^{(i)})$$

$$\lim_{r \to \infty} u_I^{(i)} = A_I^{(i)} \sin(kr + \eta_I^{(i)})$$  \hspace{1cm} (25)

in which one of the normalization constants, say $A_R^{(i)}$, is arbitrary. This is tantamount to letting $A_I^{(i)} = A_I^{(i)} / A_R^{(i)}$, which allows (23, 24) to be rewritten

$$C_1 \cos \eta_R^{(1)} + C_2 \cos \eta_R^{(2)} = \frac{1}{k} - T_I$$  \hspace{1cm} (27)

$$C_1 \sin \eta_R^{(1)} + C_2 \sin \eta_R^{(2)} = T_R$$  \hspace{1cm} (28)

$$C_1 A_I^{(1)} \cos \eta_I^{(1)} + C_2 A_I^{(2)} \cos \eta_I^{(2)} = T_R$$  \hspace{1cm} (29)

$$C_1 A_I^{(1)} \sin \eta_I^{(1)} + C_2 A_I^{(2)} \sin \eta_I^{(2)} = T_I$$  \hspace{1cm} (30)

The $\eta$'s and $A$'s are extracted from the solutions of (21, 22); thus (27-30) is a set four equations for four real constants: $C_1$, $C_2$ and in particular the real and imaginary parts of $T$: $T_R$ and $T_I$. From the latter one calculates the elastic and total cross sections, Eqs. (15), (16).

III. CALCULATION

As stated in the INTRODUCTION we calculate here only $S$-wave (i.e. $L=0$) elastic scattering, specifically phase shifts. This means, since $D_L = \text{constant}$, that the correlation function $\Phi_L$ is only a function of the "radial" coordinates. Here $\Phi_L$ is taken of Hylleraas form

$$\Phi_{L=0} = e^{-\gamma r_1 - \delta r_2} \sum_{lmn}^N C_{lmn} r_1^l r_2^m r_1^2 \pm (1 \leftrightarrow 2).$$  \hspace{1cm} (31)

where the sum includes all triples such that $l + m + n = \omega$ and $\omega = 0, 1, 2, \ldots, 8$. The total number of terms $N_\omega$ depends on spin and whether $\gamma = \delta$ or not. The values of $N_\omega$ as a function of $l, m, n$ is given in the following simple array:

<table>
<thead>
<tr>
<th>$\omega$</th>
<th>0</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>7</th>
<th>8</th>
</tr>
</thead>
<tbody>
<tr>
<td>$^1S$:</td>
<td>$N_\omega(\gamma = \delta)$</td>
<td>1</td>
<td>3</td>
<td>7</td>
<td>13</td>
<td>22</td>
<td>34</td>
<td>50</td>
<td>70</td>
</tr>
<tr>
<td>$^3S$:</td>
<td>$N_\omega(\gamma \neq \delta)$</td>
<td>1</td>
<td>4</td>
<td>10</td>
<td>20</td>
<td>35</td>
<td>56</td>
<td>84</td>
<td>120</td>
</tr>
</tbody>
</table>

To summarize the calculation: the $QHQ$ problem is solved (for a given $\gamma$ and $\delta$ and $N_\omega$). The result is a set of eigenvalues $\varepsilon_s = s=1, 2, \ldots, N_\omega$ and associated eigenfunctions $\Phi^{(s)}$. From them (the terms in) the optical potential, Eq.(14) is constructed, and the integro-differential Eq.(7) is solved noniteratively. Because all quantities are real, the solution is unique (up to an arbitrary normalization) with asymptotic form
From \( u^{(\pm)}(r) \) and its derivative the phase shift \( \eta \) is readily extracted.

Examples demonstrating the convergence of \( \eta \) for \( k=0.8 \) as a function of \( N \) are given in Table I. By virtue of the fact that \( \eta \)'s are rigorous lower bounds on the phase shift [13], the convergence then becomes a powerful indication of the accuracy of the result. Phase shifts as a function of \( k \) are given in Tables II and III. To repeat: they are rigorous lower bounds in all cases.

They are compared to the results of Schwartz [7]. Our \( ^1S \) phase shifts are seen to exceed Schwartz' (including his estimated error) at \( k=0.3, 0.4 \). Otherwise our results for both \( ^1S \) and \( ^3S \) are within his estimated uncertainty, which we find to be quite impressive considering that his calculation was carried out over 40 years ago! Nevertheless we believe that our present calculation is more accurate. The difficulty in making a secure extrapolation is due to the difficulty of estimating quantitatively the effects of polarization and other long-range potentials at non-zero \( k \). (In principle, one can optimally, rigorously include polarization by use of a polarization pseudostate [16]; however in practice those corrections are small, yet require a much more extensive calculation than is worthwhile here for our present purpose.) The convergence of the results suggests that they are accurate to one or two units in the fourth place after the decimal point. (We have given five digits because to that accuracy they are rigorous lower bounds.)

Scattering lengths are not included in our calculation, because at \( k=0 \) the Kohn variational principle, which applies to Schwartz' calculation, is well known to provide rigorous (in this case upper) bounds [17]. It is worth recalling that one of us [18] had earlier deduced that long range polarization would have decisive quantitative effect on the scattering length and had derived a formula

\[
a = a(R) - \alpha \left( \frac{1}{R} - \frac{a}{R^2} + O \left( \frac{1}{R^3} \right) \right)
\]

in which \( a(R) \) represents the value of scattering length including only the region of configuration space \( r \leq R \), and the remaining terms in (33) give the effect of the long range polarization (\( \alpha=\text{polarizability of the hydrogen atom}=4.5 \)). Using that formula led to the first correct estimate of the true scattering length [18]. Thus when Schwartz applied his calculation at \( k=0 \), he found it necessary to include the equivalent of long range polarization terms, as predicted [18], in addition to the Hylleraas terms in order to get adequate convergence. His results are: \( a(^1S)=5.965 \pm 0.003 \), \( a(^3S)=1.7686 \pm 0.0002 \). For these reasons we believe his scattering lengths are sufficiently secure and accurate as to not require further calculation.

Numerical results were obtained with Cray Y-MP computer of the NASA Center for Computation Science.
REFERENCES


TABLE I. Convergence$^a$ of S-wave phase shift as a function of $N(\omega)$ for $k=0.8$

<table>
<thead>
<tr>
<th>$N(\omega)$</th>
<th>$\gamma = \delta$</th>
<th>$\eta(+)^{(+)}$</th>
<th>$N(\omega)$</th>
<th>$\gamma$</th>
<th>$\delta$</th>
<th>$\eta(-)^{-}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$E.A.$$^a$</td>
<td>0.65127</td>
<td>0.87528</td>
<td>10(2)</td>
<td>0.65</td>
<td>0.55</td>
<td>1.63845</td>
</tr>
<tr>
<td>3(1)</td>
<td>1.23</td>
<td>0.79028</td>
<td>20(3)</td>
<td>0.84</td>
<td>0.70</td>
<td>1.64205</td>
</tr>
<tr>
<td>7(2)</td>
<td>0.68</td>
<td>0.87684</td>
<td>35(4)</td>
<td>0.85</td>
<td>0.50</td>
<td>1.64294</td>
</tr>
<tr>
<td>13(3)</td>
<td>0.68</td>
<td>0.87684</td>
<td>56(5)</td>
<td>0.85</td>
<td>0.50</td>
<td>1.64344</td>
</tr>
<tr>
<td>22(4)</td>
<td>0.69</td>
<td>0.88362</td>
<td>84(6)</td>
<td>0.85</td>
<td>0.50</td>
<td>1.64379</td>
</tr>
<tr>
<td>34(5)</td>
<td>0.61</td>
<td>0.88584</td>
<td>84(6)</td>
<td>0.85</td>
<td>0.50</td>
<td>1.64379</td>
</tr>
<tr>
<td>50(6)</td>
<td>0.54</td>
<td>0.88616</td>
<td>56(5)</td>
<td>0.85</td>
<td>0.50</td>
<td>1.64344</td>
</tr>
<tr>
<td>70(7)</td>
<td>0.66</td>
<td>0.88687</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>95(8)</td>
<td>0.72</td>
<td>0.88718</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$^a$ The phase shifts (in radians) have been optimized with respect to $\gamma$ and $\delta$ for each $N(\omega)$.
$^b$ E. A. are the well-known exchange approximation phase shifts [11]; it corresponds to no correlation terms [$N(\omega)=0 \rightarrow V_{\mu}=0$ in Eq.(7)].
TABLE II. Phase shifts of $^1S$ for various $k$ for $N=95$

<table>
<thead>
<tr>
<th>$k$</th>
<th>$\gamma = \delta$</th>
<th>$\eta$</th>
<th>$\eta_{Schwartz}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.1</td>
<td>0.53</td>
<td>2.55358</td>
<td>2.553(1)</td>
</tr>
<tr>
<td>0.2</td>
<td>0.60</td>
<td>2.06673</td>
<td>2.0673(9)</td>
</tr>
<tr>
<td>0.3</td>
<td>0.72</td>
<td>1.69816</td>
<td>1.6964(5)</td>
</tr>
<tr>
<td>0.4</td>
<td>0.70</td>
<td>1.41540</td>
<td>1.4146(4)</td>
</tr>
<tr>
<td>0.5</td>
<td>0.70</td>
<td>1.20094</td>
<td>1.202(1)</td>
</tr>
<tr>
<td>0.6</td>
<td>0.70</td>
<td>1.04083</td>
<td>1.041(1)</td>
</tr>
<tr>
<td>0.7</td>
<td>0.71</td>
<td>0.93111</td>
<td>0.930(1)</td>
</tr>
<tr>
<td>0.8</td>
<td>0.72</td>
<td>0.88718</td>
<td>0.886(1)</td>
</tr>
</tbody>
</table>
TABLE III. Phase shifts of $^3S$ for various $k$ for $N=84$

<table>
<thead>
<tr>
<th>$k$</th>
<th>$\gamma$</th>
<th>$\delta$</th>
<th>$\eta$</th>
<th>$\eta_{Schwartz}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.1</td>
<td>0.85</td>
<td>0.12</td>
<td>2.93853</td>
<td>2.9388(4)</td>
</tr>
<tr>
<td>0.2</td>
<td>0.84</td>
<td>0.39</td>
<td>2.71741</td>
<td>2.7171(5)</td>
</tr>
<tr>
<td>0.3</td>
<td>0.90</td>
<td>0.37</td>
<td>2.49975</td>
<td>2.4996(8)</td>
</tr>
<tr>
<td>0.4</td>
<td>0.91</td>
<td>0.38</td>
<td>2.29408</td>
<td>2.2938(4)</td>
</tr>
<tr>
<td>0.5</td>
<td>0.88</td>
<td>0.44</td>
<td>2.10454</td>
<td>2.1046(4)</td>
</tr>
<tr>
<td>0.6</td>
<td>0.77</td>
<td>0.52</td>
<td>1.93272</td>
<td>1.9329(8)</td>
</tr>
<tr>
<td>0.7</td>
<td>0.70</td>
<td>0.55</td>
<td>1.77950</td>
<td>1.7797(6)</td>
</tr>
<tr>
<td>0.8</td>
<td>0.86</td>
<td>0.50</td>
<td>1.64379</td>
<td>1.643(3)</td>
</tr>
</tbody>
</table>