R-matrix calculation of differential cross sections for low-energy electron collisions with ground-state and electronically excited-state O2 molecules

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Differential cross sections for electron collisions with the O2 molecule in its ground X 3Σg− state, as well as excited a 1Δg and b 1Σg+ states are calculated. As previously, the fixed-bond R-matrix method based on state-averaged complete active space self-consistent-field orbitals is employed. In addition to elastic scattering of electron with the O2 X 3Σg−, a 1Δg, and b 1Σg+ states, electron impact excitation from the X 3Σg− state to the a 1Δg and b 1Σg+ states as well as 6 eV states of c 1Σg−, A′ 1Δg, and A 3Σg+ states is studied. Differential cross sections for excitation to the 6 eV states have not been calculated previously. Electron impact excitation to the b 1Σg+ state from the metastable a 1Δg state is also studied. For electron impact excitation from the O2 X 3Σg− state to the b 1Σg+ state, our results agree better with the experimental measurements than previous theoretical calculations. Our cross sections show angular behavior similar to the experimental ones for transitions from the X 3Σg− state to the 6 eV states, although the calculated cross sections are up to a factor of 2 larger at large scattering angles. For the excitation from the a 1Δg state to the b 1Σg+ state, our results marginally agree with the experimental data except for the forward scattering direction.

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I. INTRODUCTION

A detailed knowledge of electron collisions with the oxygen molecule is important for the physics and chemistry of both laboratory and astrophysical plasmas. In particular, recent attempts to understand the electrical discharge oxygen-iodine laser have suggested that excited electronic states of O2 molecule play an important role. [1,2]. In a previous paper [3] (hereafter denoted paper I), we studied integral cross sections for electron collisions with the O2 molecule in its excited a 1Δg and b 1Σg+ states, in addition to the much studied electron scattering by the O2 X 3Σg− ground state. We used the fixed-bond R-matrix method with 13 target states represented by valence configuration interaction wave functions. State-averaged complete active space self-consistent-field (SA-CASSCF) orbitals with Gaussian-type basis functions were employed. The calculated cross sections for electron impact excitation from the a 1Δg state to the b 1Σg+ state at 4.5 eV agree well with the available experimental data of Hall and Trajmar [4]. Although elastic scattering of electrons by the a 1Δg and b 1Σg+ states was also studied, we could not find any experimental data for comparison.

In paper I, theoretical and experimental integral cross sections were compared. However, differential cross sections (DCSs) provide a more stringent test of theory and are often easier to measure reliably than integral cross sections. For electron impact electronic excitations, calculations which give good integral cross sections often give DCSs which differ significantly from those observed experimentally. In this paper, we present DCSs for the corresponding processes calculated using the same R-matrix model.

Previous experimental and theoretical studies in the field of electron O2 collisions are well summarized by Brung and Buckman [5]. Here we only review works relevant to this paper. The DCSs of electron collisions with O2 molecule have been measured by many experimental groups. In particular, electron impact excitation to the low-lying a 1Δg and b 1Σg+ states have been studied experimentally by Trajmar et al. [6], Shyn and Sweeney [7], Allan [8], Middleton et al. [9], and Linert et al. [10]. In contrast to these experimental works, only Middleton et al. [9] report calculations of DCSs for these excitation processes. Some of the more recent measurements have focused on electron impact excitations from the X 3Σg− ground state to the 6 eV states, i.e., the c 1Σg−, A′ 1Δg, and A 3Σg+ states which are also called the Herzberg pseudocontinuum [11–13]. Although these DCSs are not state resolved in most cases, Shyn and Sweeney [13] obtained cross sections for excitation to the individual electronic state within the 6 eV states. In this paper, we also calculate the DCSs of this process using the fixed-bond R-matrix method, since no previous theoretical calculation exists. Up to now, there is only one measurement of DCSs for electron collisions with electronically excited O2 molecule. Hall and Trajmar [4] obtained excitation cross sections from the O2 a 1Δg to the b 1Σg+ state at electron impact energy of 4.5 eV. Their integral cross section was compared with our R-matrix results in paper I.

In this paper, details of the calculations are presented in Sec. II, and we discuss the results in Sec. III comparing our results with previous theoretical and available experiments. Then the summary is given in Sec. IV.

II. THEORETICAL METHODS

The R-matrix method itself has been described extensively in the literature [14–16] as well as in paper I. Thus we
do not repeat a general explanation of the method here. We used a modified version of the polyatomic programs in the UK molecular R-matrix codes [14] to extract T-matrix elements of the electron O₂ scatterings. These programs utilize Gaussian-type orbitals (GTO) to represent target molecule as well as a scattering electron. Although most of the past R-matrix works on electron O₂ collisions have employed Slater-type orbitals (STO), we select GTOs mainly because of the simplicity of the input and availability of basis functions. The SA-CASSCF orbitals are imported from the calculations with the MOLPRO suites of programs [17]. The use of SA-CASSCF orbitals improves the vertical excitation energies of the O₂ target states compared to the energies from HF orbitals as shown in paper I. These target orbitals are constructed from the [5s,3p] contracted basis of Dun-ning [18] augmented by a d function with exponent 1.8846, as in Sarpal et al. [19]. In the R-matrix calculations, we included 13 target states; \( \sum_{\sigma} c_a^* \Delta_{\sigma}, b \sum_{\sigma} c^* \sum_{\sigma} A^* \Delta_{\sigma}, A^* \sum_{\sigma}, B \sum_{\sigma}, \Delta_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma}, f^* \sum_{\sigma} \), where the last 4 \( \Delta_{\sigma} \) states were not included in the previous R-matrix studies performed by other groups. These target states were represented by valence configuration interaction wave functions constructed from the SA-CASSCF orbitals. In our fixed-bond R-matrix calculations, these target states are evaluated at the equilibrium bond length \( R=2.3a_0 \) of the \( \mathrm{O}_2 \) \( X^2 \Sigma_g^- \) ground electronic state.

The radius of the R-matrix sphere was chosen to be 10\( a_0 \) in our calculations. In order to represent the scattering electron, we included diffuse Gaussian functions up to \( l=5 \) with 9 functions for \( l=0 \), seven functions for \( l=1 \ldots 4 \) and six functions for \( l=4 \) and 5. The exponents of these diffuse Gaussian were fitted using the GTOBAS program [20] in the UK R-matrix codes. Details of the fitting procedure are the same as in Faure et al. [20]. In addition to these continuum orbitals, we included eight extra virtual orbitals, one for each symmetry. The construction of the 17 electrons CSFs for the total system is the same as in paper I. The R-matrix calculations were performed for all 8 irreducible representations of the \( D_{2h} \) symmetry, \( \mathrm{A}_u, \mathrm{B}_2 \), \( \mathrm{B}_3 \), \( \mathrm{B}_1u \), \( \mathrm{B}_3u \), \( \mathrm{B}_2g \), and \( \mathrm{A}_g \), for both doublet and quartet spin multiplicity of the electron plus target system.

The DCSs are evaluated from the T-matrix elements obtained by the R-matrix calculations. As in Gianturco and Jain [21] and Malegat [22], the DCS is expanded using the Legendre polynomials,

\[
\frac{d\sigma}{d\Omega}_{ij} = \sum_k A_k P_k(\cos \theta),
\]

where \( i \) and \( j \) denote the initial and final electronic states of the target, respectively. In exactly the same way as in Malegat [22], but for \( D_{2h} \) symmetry instead of \( D_{4h} \) symmetry in her paper, we can derive an expression of the expansion coefficients \( A_k \), which is

\[
A_k = \sum_{l|m|j} \sum_{l'|n|j'} \frac{(-1)^{\mu+\nu l'-l-\mu j'-j}(2k+1)}{8(2S_z+1)k^2} \delta_{l',\lambda',\mu';\lambda,\mu,j,l} \left( \begin{array}{lll} l & l' & k \\ -\lambda & \lambda' & \lambda - \lambda' \end{array} \right) \left( \begin{array}{lll} j & l' & k \\ -\mu & \mu' & \mu - \mu' \end{array} \right) \left( \begin{array}{lll} j & l & k \\ 0 & 0 & 0 \end{array} \right) \sum_{S} \left( \begin{array}{lll} S & S_z & \gamma_{m,n,m'} \gamma_{l,n,l'} \gamma_{\lambda,n,\lambda'} \gamma_{\mu,n,\mu'} \gamma_{\lambda',n,\lambda''} \gamma_{\mu',n,\mu''} \gamma_{\lambda'',n,\lambda'''} \gamma_{\mu'',n,\mu'''} \gamma_{\lambda''',n,\lambda''''} \gamma_{\mu''',n,\mu''''} \gamma_{\lambda'''',n,\lambda'''''} \gamma_{\mu'''',n,\mu'''''} \gamma_{\lambda''''',n,\lambda''''''} \gamma_{\mu''''',n,\mu''''''} \gamma_{\lambda'''''',n,\lambda'''''''} \gamma_{\mu'''''',n,\mu'''''''}} \end{array} \right).
\]

Details of the derivation are given in the Appendix. In Eq. (2),

\[
\left( \begin{array}{llll} l & l' & k \\ -\lambda & \lambda' & \lambda - \lambda' \end{array} \right),
\]

diffs from coefficients, \( k \), is the wave number of the incident electron, \( S_z \) is the spin quantum number of the initial target state, while \( S \) is the spin quantum number of the total system and \( M_S \) is the projection of the total spin. The indices \( \Gamma \) and \( \Gamma' \) run over the eight irreducible representations of the \( D_{2h} \) point group, since we employ the polyatomic version of the UK R-matrix code. The angular quantum numbers of the scattering electron, \( l_i \) and \( m_i \), etc., in the T-matrix element \( T^{\gamma SM_S}_{l_im_i,l'_m_i,j_m_i} \) specify the real spherical harmonics \( S^\gamma_l \) instead of the complex form \( Y^\gamma_l \), because the \( S^\gamma_l \) transform as irreducible representations under \( D_{2h} \) symmetry. This means, there

are transformation matrix elements \( C_{\lambda,m} \), etc., in the expression for \( A_k \) in order to convert the index of the scattering electron from the \( S^\gamma_l \) representation to the \( Y^\gamma_l \) representation. An expression of the matrix elements \( C_{\lambda,m} \) is given in the Appendix. Finally, we note that the summations over \( (\Gamma l_m) \) should satisfy the symmetry relation, \( \Gamma' = \text{IR} (\Gamma l_m) \times \text{IR}(S^\gamma_l) \), with \( \text{IR} \) and \( \text{IR}(S^\gamma_l) \) each being an irreducible representation of the \( D_{2h} \) group corresponding to the \( l \)th target state and the real spherical harmonic \( S^\gamma_l \), respectively. This relation also holds for the \( (\Gamma' l'_m) \), \( (\Gamma' \Gamma' l'_m m'_j) \), and \( (\Gamma' l'_m m') \).

In Eq. (2), the T-matrix elements belonging to different total symmetries are multiplied together. Since these matrix elements come from different calculations, overall phases of molecular orbitals and target CI vectors underlying these matrix elements may be inconsistent (see Ref. [23]), which may result in erroneous relative signs of these T-matrix elements. To avoid this inconsistency, we saved reference target CI
much lower than the distorted-wave approximation. Their results at 5 eV are
Schwinger variational iterative method combined with the orbitals was used in all these calculations.

values at 10°. This situation is the same in the previous calculations, according to this reference. The same set of molecular
other calculations, 

The same set of molecular

III. RESULTS AND DISCUSSION

A. Electron collisions with the O₂ X 3Σ⁻ ground state

Figure 1 shows DCSs for elastic electron scattering from the O₂ X 3Σ⁻ state compared with previous theoretical and experimental results. Our results are very similar to the previous R-matrix cross sections of Wöste et al. [24] and Machado et al. [25] were calculated using the Schwinger variational iterative method combined with the distorted-wave approximation. Their results at 5 eV are much lower than the R-matrix results at low scattering angle below 50 degrees. Our results agree reasonably well with the experimental cross sections at 10 eV, including the recent results of Linert et al. [26] for backward scattering. At 5 eV, our model significantly overestimates the cross sections for forward scattering compared to the experimental values. For example, our result is twice as large as the experimental values at 10°. This situation is the same in the previous R-matrix calculation of Wöste et al. [24]. As discussed by Wöste et al., this deviation can be attributed to a lack of long-range polarizability in the scattering model. For example, Gillan et al. [27] introduced polarized pseudostates to account for the long-range polarizability in electron-N₂ scattering and reduced the cross sections by 50% in the threshold energy region. The interaction potential of Machado et al. [25] includes the correlation-polarization term based on the free-electron-gas model. Probably the polarization introduced by this term is responsible for their better agreement with experiment at 5 eV. Since we are interested in electron collisions with the excited electronic states of O₂ in this work, we choose not to pursue precise accuracy further for the ground state elastic scattering. However, we must be mindful that similar long-range polarizability problems may exist in the other low-energy electron scattering processes, especially elastic electron scattering of the a ¹Δg and b ¹Σg° state O₂ molecules, which will be discussed below.

The DCSs for excitation to the a ¹Δg state at electron impact energy 5 and 10 eV are compared in Fig. 2 with the previous theoretical calculation and the experimental measurements of Middleton et al. [9], Shyn and Sweeney [7], Allan [8], and Linert et al. [10]. The cross sections at 5 eV agrees well with the previous calculation and experimental data below 120°. However, our results are much lower than the previous calculation of Middleton et al. at scattering angle above 130°. At an electron scattering energy of 10 eV, our cross section deviates further from the previous calculation of Middleton et al. [9] especially at scattering angle below 60° and above 140°. In contrast to the backward enhanced cross sections of Middleton et al., our DCSs have a slightly forward enhanced character. Our results agree better with the experimental data at low scattering angles than the
previous calculation of Middleton et al. [9]. At large scattering angle, our results deviate from the experimental data of Shyn and Sweeney [7], but agrees rather well with the recent measurement of Linert et al. [10]. The R-matrix model of Middleton et al. [9] included the lowest nine O\textsubscript{2} target states and \( l = 0\)–5 scattering electron orbitals with \( \sigma, \pi, \) and \( \delta \) symmetry. In this work, we included 13 target states and all components of \( l = 0\)–5 scattering electron orbitals. In addition to these differences, Middleton et al. used HF STO orbitals where we employed CASSCF GTO orbitals. We carried out a test calculation with \( l = 0\)–3 scattering electron orbitals and got almost the same cross sections as in the \( l = 0\)–5 case, which suggests that difference in the number of target states may be important for the shape of these excitation cross sections.

Figure 3 compares DCSs for excitation to the \( b^1\Sigma^+_g \) state at electron impact energy of 5 and 10 eV with the previous R-matrix calculation and the experimental measurements of Middleton et al. [9], Shyn and Sweeney [7], and Allan [8]. Transitions between \( \Sigma^+ \) and \( \Sigma^- \) target states are forbidden at scattering angles of 0° and 180°, because the scattered electron wave function vanishes in the plane defined by incident electron beam and the molecular axis for any orientation of the molecule [28,29]. As a consequence, the DCSs decrease to be zero toward 0° and 180°. As is apparent from Fig. 3, our cross sections become zero at 0° and 180°, which is consistent with this selection rule. Compared to the previous R-matrix calculations of Middleton et al. [9], our cross sections have similar profile, but with slightly smaller magnitude at all scattering angles. Agreement with experiment is good at 5 eV below 120°, although our results underestimate the experimental cross sections at larger scattering angles. At 10 eV, the magnitude of the experimental cross sections of Middleton et al. [9] and Shyn and Sweeney [7] do not agree with each other; however, our cross sections are closer to the results of Shyn and Sweeney at low scattering angles below 50°. Between 60° and 90°, our results are closer to the results of Middleton et al.

Figure 4 shows DCSs for excitations to the 6 eV states for electron impact energies of 10 and 15 eV. Here the 6 eV states means the group of the \( c^1\Sigma^-_u, A^1\Delta_u, \) and \( \tilde{A}^3\Sigma^+_u \) states. The cross sections shown in Fig. 4 are a sum of individual excitation cross sections of these three electronic states, in line with most experimental measurements. The figure includes the recent experimental cross sections of Campbell et al. [11], Green et al. [12], and Shyn and Sweeney [13]. The individual cross sections are shown in Figs. 5 and 6 for impact energies of 10 and 15 eV, together with the state-resolved experimental cross sections of Shyn and Sweeney [13]. Our summed cross sections given in Fig. 4 are backward-enhanced for both the 10 and 15 eV cases, in accordance with the experimental cross sections Campbell et al. [11] and Shyn and Sweeney [13]. However, the forward enhancement of the DCSs at 10 eV observed by Green et al. [12] is not reproduced by our calculation. The individual cross sections in Figs. 5 and 6 show similar angular behavior compared to the experimental results of Shyn and Sweeney. However, our DCSs for excitation to the \( A^1\Delta_u \) state is more steeply backward direction. Also the peak in the \( A^3\Sigma^+_u \) state cross sections is more pronounced in our
calculation. Note that our results for the \( A^3\Sigma_u^- \) state become zero at 0° and 180° as dictated by the \( \Sigma^-\Sigma^+ \) selection rule.

**B. Electron collisions with the O\(_2\) \( a^1\Delta_g \) and \( b^1\Sigma_g^+ \) excited states**

The DCSs for elastic electron scattering with the excited O\(_2\) \( a^1\Delta_g \) and \( b^1\Sigma_g^+ \) states are shown in Fig. 7. We cannot compare them with previous theoretical or experimental work, since there is no available data. These DCSs show strong similarity with those of the elastic electron scattering with the \( X^3\Sigma_g^- \) ground state in Fig. 1. The magnitude of these cross sections is almost the same for the 10 eV case. All of them have a large forward peak at 0°, a small rise in the cross sections at 180°. The location of the minimum moves inward from 140° to 90° as the electron scattering energy increases. This similarity is also reflected in the integral cross sections for elastic electron collisions with the \( X^3\Sigma_g^- \), \( a^1\Delta_g \), and \( b^1\Sigma_g^+ \) states. The profiles and magnitudes of the integral cross sections are basically the same for all these three electronic states as shown in paper I. The main configuration of these three electronic states has the form \( \pi^2\sigma^4 \), and this may be responsible for this similarity. Our R-matrix calculations tend to overestimate the elastic scattering cross sections of the \( X^3\Sigma_g^- \) state at low scattering angles, below 50°, compared to the experimental data. Considering the strong similarity of the cross section profiles for elastic scattering from excited states and the ground state, our calculations may also overestimate the cross section at low scattering angle at low electron impact energy.

In Table I, we show momentum transfer cross sections for electron elastic scattering by the \( X^3\Sigma_g^- \), \( a^1\Delta_g \), and \( b^1\Sigma_g^+ \) states. As a consequence of the similarity in DCSs, the momentum transfer cross sections have a similar magnitude. Compared to the experimental data of Shyn and Sharp [30].

**FIG. 5.** Differential cross sections for excitation from the O\(_2\) \( X^3\Sigma_g^- \) state to the individual state of the 6 eV states. (a) Shows excitation cross sections for the \( c^1\Sigma_u^- \) state, (b) is for the \( A^3\Sigma_u^- \) state, and (c) is for the \( A^3\Sigma_u^+ \) state. The electron impact energy is 10 eV. The experimental data was taken from Shyn and Sweeney [13].

**FIG. 6.** The same as in Fig. 5, but for an electron impact energy of 15 eV.

**FIG. 7.** Differential cross sections for elastic scattering of the O\(_2\) excited states. (a) The \( a^1\Delta_g \) state, (b) the \( b^1\Sigma_g^+ \) state. Each line corresponds to a different electron impact energy as shown in the legend.
TABLE I. Elastic momentum transfer cross sections in unit of $10^{-16}$ cm$^2$.

<table>
<thead>
<tr>
<th>Electron impact energy (eV)</th>
<th>$X^3\Sigma_g^-$</th>
<th>$a^1\Delta_g$</th>
<th>$b^1\Sigma_g^+$</th>
</tr>
</thead>
<tbody>
<tr>
<td>This work</td>
<td>8.18</td>
<td>8.16</td>
<td>7.95</td>
</tr>
<tr>
<td>$X^3\Sigma_g^-$</td>
<td>6.80</td>
<td>6.69</td>
<td>6.57</td>
</tr>
<tr>
<td>$a^1\Delta_g$</td>
<td>6.73</td>
<td>6.57</td>
<td>6.29</td>
</tr>
<tr>
<td>$b^1\Sigma_g^+$</td>
<td>6.04</td>
<td>5.90</td>
<td>5.83</td>
</tr>
<tr>
<td>$X^3\Sigma_g^-$</td>
<td>6.04</td>
<td>5.86</td>
<td>5.85</td>
</tr>
<tr>
<td>$a^1\Delta_g$</td>
<td>6.96</td>
<td>6.79</td>
<td>6.39</td>
</tr>
<tr>
<td>$b^1\Sigma_g^+$</td>
<td>6.73</td>
<td>6.57</td>
<td>6.29</td>
</tr>
</tbody>
</table>

Shyn and Sharp [30] $X^3\Sigma_g^-$ 6.7 8.4
Sullivan et al. [31] $X^3\Sigma_g^+$ 6.5 6.0 6.2 6.4

and Sullivan et al. [31], our calculation overestimates the $X^3\Sigma_g^-$ state momentum transfer cross section at 2 eV by 20%, but underestimates the cross section at 10 eV by 8% of Sullivan et al. value or 29% of the Shyn and Sharp value. Our momentum transfer cross sections for the $a^1\Delta_g$ and $b^1\Sigma_g^+$ states may similarly be overestimates or underestimates depending on the electron impact energy.

Figure 8 shows DCSs for electron impact excitation from the $a^1\Delta_g$ state to the $b^1\Sigma_g^+$ state. The figure also includes the experimental data of Hall and Trajmar [4] at impact energy of 4.5 eV. Our cross section profiles have characteristic features of minima around 10° and 90° and maxima around 50° and 150°. They agree with the experimental cross sections of Hall and Trajmar [4] within their error bars except at 20° and 30°. The cross sections of Hall and Trajmar appear to increase from 50° to 0° whereas our cross sections decrease from 60° toward 10°. In the 50°–140° angular region, Hall and Trajmar’s cross sections vary less than ours. However, a precise comparison is difficult because of large error bars and lack of other experimental data.

IV. SUMMARY

We have calculated differential cross sections for electron collisions with the O$_2$ molecule in its ground $X^3\Sigma_g^-$ state, as well as excited $a^1\Delta_g$ and $b^1\Sigma_g^+$ states. As in our previous work, we employed the fixed-bond $R$-matrix method based on state-averaged complete active space SCF orbitals. In addition to elastic scattering of electron with the O$_2$ $X^3\Sigma_g^-$, $a^1\Delta_g$ and $b^1\Sigma_g^+$ states, we studied electron impact excitations from the $X^3\Sigma_g^-$ state to the $a^1\Delta_g$ and $b^1\Sigma_g^+$ states as well as 6 eV states of $c^1\Sigma_u^-$, $A^2\Sigma_u^+$, and $A^2\Sigma_g^+$ states. DCSs for the excitations to the 6 eV states were not calculated previously. We also studied electron impact excitation to the $b^1\Sigma_g^+$ state from the metastable $a^1\Delta_g$ state. For electron impact excitation from the O$_2$ $X^3\Sigma_g^-$ state to the $b^1\Sigma_g^+$ state, our results agree better with the experimental measurements than the previous theoretical cross sections. Our cross sections show similar angular behavior to the experimental ones for transitions from the $X^3\Sigma_g^-$ state to the 6 eV states. For the excitation from the $a^1\Delta_g$ state to the $b^1\Sigma_g^+$ state, our results marginally agree with experimental data except for the forward scattering direction.

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APPENDIX: DERIVATION OF EQUATION (2)

Derivation of the DCS formula in Eq. (2) is similar to that of Malegat [22] except for the use of real spherical harmonics $S_\nu^\mu$ employed in the polyatomic version of UK molecular $R$-matrix codes instead of complex form $Y_\nu^\mu$. For convenience of the reader, brief derivation of the formula is given in this Appendix. In the expressions below, we follow the notation of Malegat [22].

The scattering wave function describing collision of an electron plane wave with a molecule is expressed as

$$
\Psi_f(x_1',...,x_N',\sigma',r) = \Psi_f(x_1',...,x_N') \chi_{(1/2)m_1}(\sigma') e^{i k z}
+ \sum \Psi_f(x_1',...,x_N') \chi_{(1/2)m_1}(\sigma')
\times F_{1/2}(\mathbf{r}) e^{i k / r}.
$$

(A1)

Here $x$ denotes the space and spin coordinates of the molecular electrons. The primed coordinates refer to the molecular frame with the $z'$ axis along the molecular symmetry axis, and the unprimed coordinates to the laboratory frame with the $z$ axis along the incident electron beam. The incident electron has wave number $k$, with spin projection $m_1$. The index $i$ represents quantum numbers of the electronic state of
the target molecule, $\Gamma$, $S$, and $M_S$, whereas the index $I$ refers to $(i, m_i)$ collectively.

In order to expand the wave function in Eq. (A1), a symmetry adapted $(N+1)$-electron wave function is prepared as

$$
\Psi_{i \mu m_j}^{GSM_S}(x_1', \ldots, x_N', \alpha', r) = \sum_{\mu m_j} \Psi_{j \mu m_j}^{GSM_S}(x_1, \ldots, x_N, \alpha') S_{j \mu m_j}^m(r')
\times f_{i \mu m_j}^{GSM_S}(r)/r,
$$

(A2)

where $\Gamma$, $S$, and $M$ stand for symmetry of the $(N+1)$-electron system, i.e., an irreducible representation of the $D_{2h}$ group in this work, spin quantum number and its projection to the symmetry axis. The orbital angular momentum of the scattering electron and its projection are represented by $l$ and $m_l$. In case of $m > 0$, the real spherical harmonics $Y^m_{i-1}$ is related to the complex form of spherical harmonics $Y^m_i$ as [33]

$$
\begin{pmatrix}
Y^m_i \\
Y^{-m}_i
\end{pmatrix} = \frac{1}{\sqrt{2}} \begin{pmatrix}
(-1)^m & -1 \\
1 & i
\end{pmatrix} \begin{pmatrix}
S^m_i \\
S^{-m}_i
\end{pmatrix}.
$$

(A3)

In the $m=0$ case, we only have $Y^0_i$ and $S^0_i$ and thus the matrix element is 1. Note that $S^m_i$ behaves as an irreducible representation under $D_{2h}$ symmetry operations, whereas $Y^m_i$ does not. The spin coupled function in Eq. (A2) is obtained by integrating over the asymptotic region by

$$
\lim_{r \to \infty} f_{i \mu m_j}^{GSM_S}(r) = \frac{1}{\sqrt{k_j}} \left[ e^{-i(k_j r - (1/2)\pi)} \delta_{i \mu m_j} + e^{i(k_j r - (1/2)\pi)} i^{i \mu m_j} \right].
$$

(A5)

Expanding Eq. (A1) in the symmetry adapted functions of Eq. (A2) gives

$$
\Psi_I(x_1', \ldots, x_N', \alpha', r) = \sum_{i \mu m_j} \sum_{\Gamma} a^i_{\mu m_j} \Gamma_{\alpha \beta \gamma} \psi_{GSM_S}(x_1', \ldots, x_N', \alpha', r).
$$

(A6)

By comparing the incoming parts on the right- and left-hand sides, we obtain the expansion coefficient

$$
\begin{align*}
\Gamma_{\alpha \beta \gamma} &= -i/\sqrt{4\pi(2l+1)} \sum_{\lambda} D^{\lambda \gamma}_{\alpha \beta} C_{\lambda, m_i} \\
&\times \left( S_{MS} S_{i \mu m_j}^m \right),
\end{align*}
$$

(A7)

where $D^{\lambda \gamma}_{\alpha \beta} C_{\lambda, m_i}$ is the rotation matrix with the Euler angles $(\alpha, \beta, \gamma)$ representing rotation of the laboratory frame to the molecular frame. The matrix element $C_{\lambda, m_i}$ defined in Eq. (A3), relates the spherical harmonics $Y^m_{i \gamma}$ and $S^m_{i \gamma}$. The collision amplitude can then be obtained by equating the outgoing parts,

$$
F_I(r) = \sum_{i \mu m_j} \sum_{\Gamma} \sqrt{\pi(2l+1)} \left( S_{MS} S_{i \mu m_j}^m \right) \\
\times \left( S_{MS} S_{i \mu m_j}^m \right)
\times D^{\lambda \gamma}_{\alpha \beta} D^{\lambda \gamma}_{\alpha \beta} Y^m_{i \gamma}(r) C_{\lambda, m_i} C_{\lambda, m_i}^* T_{i \mu m_j}^{GSM_S}.
$$

(A8)

Here we use the $T$-matrix elements $T_{i \mu m_j}^{GSM_S}$ instead of the $S$-matrix.

By summing over the final states and averaging over the initial states and the molecular orientation $(\alpha, \beta, \gamma)$, the differential cross section is expressed by the Legendre polynomials expansion (1) with expansion coefficients given by Eq. (2).