

University of Nebraska - Lincoln

DigitalCommons@University of Nebraska - Lincoln

Anthony F. Starace Publications

Research Papers in Physics and Astronomy

May 1975

Effects of anisotropic electron-ion interactions in atomic photoelectron angular distributions

Dan Dill

Boston University, Boston, Massachusetts

Anthony F. Starace

University of Nebraska-Lincoln, astarace1@unl.edu

Steven T. Manson

Georgia State University, Atlanta, Georgia

Follow this and additional works at: <https://digitalcommons.unl.edu/physicsstarace>



Part of the [Physics Commons](#)

Dill, Dan ; Starace, Anthony F.; and Manson, Steven T., "Effects of anisotropic electron-ion interactions in atomic photoelectron angular distributions" (1975). *Anthony F. Starace Publications*. 5.

<https://digitalcommons.unl.edu/physicsstarace/5>

This Article is brought to you for free and open access by the Research Papers in Physics and Astronomy at DigitalCommons@University of Nebraska - Lincoln. It has been accepted for inclusion in Anthony F. Starace Publications by an authorized administrator of DigitalCommons@University of Nebraska - Lincoln.

Effects of anisotropic electron-ion interactions in atomic photoelectron angular distributions

Dan Dill

Department of Chemistry, Boston University, Boston, Massachusetts 02215

Anthony F. Starace*

Behlen Laboratory of Physics, The University of Nebraska, Lincoln, Nebraska 68508

Steven T. Manson†

Department of Physics, Georgia State University, Atlanta, Georgia 30303

(Received 3 October 1974)

The photoelectron asymmetry parameter β in LS coupling is obtained as an expansion into contributions from alternative angular-momentum transfers j_t . The physical significance of this expansion of β is shown to be that (i) the electric-dipole interaction transfers to the atom a characteristic single angular momentum $j_t = l_0$, where l_0 is the photoelectron's initial orbital momentum, whereas (ii) angular-momentum transfers $j_t \neq l_0$ indicate the presence of anisotropic (i.e., term-dependent) interaction of the outgoing photoelectron with the residual ion. For open-shell atoms the photoelectron-ion interaction is generally anisotropic; photoelectron phase shifts and electric-dipole matrix elements depend on both the multiplet term of the residual ion and the total orbital momentum of the ion-photoelectron final-state channel. Consequently, β depends on the term levels of the residual ion and contains contributions from all allowed values of j_t . These findings contradict the independent-particle-model theory for β , which ignores the final-state electron-ion interaction and to which our expressions reduce in the limiting cases for which only $j_t = l_0$ is allowed, namely (a) spherically symmetric atoms (e.g., closed-shell atoms) and (b) open-shell atoms for which the electron-ion interaction is isotropic (e.g., very light elements). Numerical calculations of the asymmetry parameters and partial cross sections for photoionization of atomic sulfur are presented to illustrate the theory and to demonstrate the information on electron-ion dynamics that can be obtained from the theoretical and experimental study of β for open-shell atoms.

I. INTRODUCTION

We obtain in this paper explicit expressions, in LS coupling, for the angular distribution of photoelectrons produced by electric dipole ionization of an arbitrary open- or closed-shell atom. Our treatment is based on the angular momentum transfer expansion for the differential photoionization cross section¹⁻³ and is intended to provide a theoretical framework that allows angular distribution calculations comparable in accuracy to the best calculations of total photoionization cross sections. The formulas we obtain show explicitly the influence of anisotropic electron-ion interactions on the electron angular distribution and at the same time explain the success of the Cooper-Zare⁴ independent-particle-model theory in predicting such distributions for closed-shell atoms.^{5,6} For other than the lightest open-shell atoms,⁷ however, we expect anisotropic electron-ion interactions to produce photoelectron angular distributions that deviate significantly from the predictions of the Cooper-Zare theory.

Our conclusions, described above, are contained

implicitly in the LS -coupling formulas for the angular distribution asymmetry parameter β obtained by Lipsky^{8,9} and by Jacobs and Burke,¹⁰ whose formulas are in principle equivalent to ours. The advantage of the angular momentum transfer expansion employed in this paper, however, is that such conclusions follow explicitly from our formulation. Hartree-Fock calculations of the angular distribution of electrons photoionized from atomic sulfur, a typical open-shell atom, are presented to illustrate our theoretical predictions.

In Sec. II we summarize the angular momentum transfer formulation of the differential photoionization cross section. We also exhibit how the angular momentum transfer probes anisotropic electron-ion interactions. The formulas in this section depend on the amplitude for photoionization with a particular value of the angular momentum transfer. The form of this amplitude in LS coupling, a main result of this paper, is obtained in Sec. III. In Sec. IV we illustrate the theory by calculating the photoelectron angular distribution of atomic sulfur. Lastly, we discuss our conclusions in Sec. V. A brief report of these results has been published elsewhere.¹¹

II. SUMMARY OF THE ANGULAR MOMENTUM TRANSFER FORMULATION

The ejection of an electron e^- from an *unpolarized* atomic target A by electric dipole interaction with an incident photon γ may be represented schematically as

$$A(J_0\pi_0) + \gamma(j_\gamma = 1, \pi_\gamma = -1) - A^+(J_c\pi_c) + e^-[lsj, \pi_e = (-1)^l]. \quad (1)$$

The differential cross section for this process can be separated into contributions characterized by alternative values of the angular momentum transfer,

$$\vec{j}_t = \vec{j}_\gamma - \vec{l} = \vec{J}_c + \vec{s} - \vec{J}_0, \quad (2)$$

provided no measurement is made of either the photoelectron spin or the orientation of the residual ion. The vector \vec{j}_t is the angular momentum transferred between the *unobserved* initial and final angular momenta in the reaction, i.e., between the total angular momentum \vec{J}_0 of the target A and the combined angular momenta of the residual ion A^+ and the photoelectron spin \vec{s} , which we denote $\vec{J}_{cs} \equiv \vec{J}_c + \vec{s}$. Allowed values of j_t are determined by conservation of angular momentum \vec{J} and parity π in reaction (1):

$$\vec{J} = \vec{J}_0 + \vec{j}_\gamma = \vec{J}_c + \vec{s} + \vec{l}, \quad (3)$$

$$\pi = -\pi_0 = \pi_c(-1)^l. \quad (4)$$

The general form of the differential cross section for reaction (1) is¹²

$$\frac{d\sigma}{d\Omega} = \frac{\sigma}{4\pi} [1 + \beta P_2(\cos\theta)]. \quad (5)$$

Here σ is the total cross section, θ is the angle between the axis of linear polarization of the incident light and the direction of the outgoing photoelectron, and β is the asymmetry parameter. The dynamical features of the angular distribution are thus contained in β , which may assume values in the range $-1 \leq \beta \leq 2$, corresponding to distributions varying from $\sin^2\theta$ to $\cos^2\theta$. [Though Eq. (5) assumes linearly polarized incident light, unpolarized,¹³ partially polarized,¹⁴ and elliptically polarized¹⁵ incident light produce angular distributions that may be expressed in terms of β .]

The resolution of Eq. (5) into contributions corresponding to alternative values of j_t requires first that one determine the allowed values of j_t from Eqs. (2)–(4). Second, each value of j_t is characterized as being either parity favored or parity unfavored,² corresponding to whether the parity change of the target $\pi_0\pi_c$ is equal to $+(-1)^{j_t}$ or $-(-1)^{j_t}$, respectively. The total cross section σ and the asymmetry parameter β may then be ex-

pressed in terms of cross sections $\sigma(j_t)$ and asymmetry parameters $\beta(j_t)$ for a particular value of j_t as follows³:

$$\sigma = \sum_{j_t, J_{cs}} \sigma(j_t), \quad (6)$$

$$\sigma\beta = \sum_{J_{cs}} \left(\sum_{j_t}^{\text{fav}} \sigma(j_t)_{\text{fav}} \beta(j_t)_{\text{fav}} - \sum_{j_t}^{\text{unf}} \sigma(j_t)_{\text{unf}} \right). \quad (7)$$

In Eq. (7) we have summed the favored and unfavored values of j_t separately, but, as seen from Eq. (11) below, Eq. (7) represents β as a weighted average of $\beta(j_t)$. Note that while Eqs. (6) and (7) also have sums over J_{cs} (cf. Ref. 3, p. 1981) we do not indicate the dependence of $\sigma(j_t)$ and $\beta(j_t)$ on quantum numbers other than j_t until Sec. III of this paper. This dependence is hidden in the scattering matrix amplitudes $\bar{S}_l(j_t)$, in terms of which $\sigma(j_t)$ and $\beta(j_t)$ are given by³

$$\sigma(j_t)_{\text{fav}} = \pi\chi^2 \frac{2j_t+1}{2J_0+1} [|\bar{S}_+(j_t)|^2 + |\bar{S}_-(j_t)|^2], \quad (8)$$

$$\sigma(j_t)_{\text{unf}} = \pi\chi^2 \frac{2j_t+1}{2J_0+1} |\bar{S}_0(j_t)|^2, \quad (9)$$

$$\begin{aligned} \beta(j_t)_{\text{fav}} = & \{(j_t+2)|\bar{S}_+(j_t)|^2 + (j_t-1)|\bar{S}_-(j_t)|^2 \\ & - 3[j_t(j_t+1)]^{1/2} [\bar{S}_+(j_t)\bar{S}_-(j_t)^* + \text{c.c.}]\} / \\ & \{(2j_t+1)[|\bar{S}_+(j_t)|^2 + |\bar{S}_-(j_t)|^2]\}, \quad (10) \end{aligned}$$

$$\beta(j_t)_{\text{unf}} = -1. \quad (11)$$

In these equations, χ is the photon wavelength divided by 2π and c.c. denotes complex conjugate. The parity-favored cross sections and asymmetry parameters, Eqs. (8) and (10), depend on photoionization amplitudes $\bar{S}_\pm(j_t)$, the “ \pm ” denoting the value of the photoelectron’s orbital angular momentum, $l = j_t \pm 1$. The parity-unfavored partial cross sections in Eq. (9) depend on the amplitudes $\bar{S}_0(j_t)$, the “0” denoting $l = j_t$. That the asymmetry parameter for any parity-unfavored value of j_t is -1 independent of dynamics, as indicated in Eq. (11), is discussed in Ref. 2.

The scattering amplitudes $\bar{S}_l(j_t)$ may be expressed as a sum of reduced electric dipole matrix elements, each corresponding to a given total angular momentum \vec{J} ³:

$$\begin{aligned} \bar{S}_l(j_t) = & ((J_c s) J_{cs} l | \bar{S}(j_t) | \alpha_\sigma J_0 j_\gamma = 1) \\ = & n(\chi) \sum_J (-1)^{J_0 - J - 1} \hat{J} \begin{Bmatrix} J_{cs} & l & J \\ 1 & J_0 & j_t \end{Bmatrix} \\ & \times ((J_c s) J_{cs} l, J - \| P^{[-1]} \| \alpha_\sigma J_0). \quad (12) \end{aligned}$$

Here, $n(\chi) \equiv 4\pi\alpha\hbar\omega/3\chi^2$, $\hat{J} \equiv (2J+1)^{1/2}$, α_σ denotes the set of quantum numbers necessary to uniquely

specify the initial state, and the minus sign indicates that the final state is normalized according to incoming-wave boundary conditions. In Eq. (12) the “±” and “0” subscripts of the scattering-matrix amplitudes are replaced by the appropriate value of l . As it stands, Eq. (12) is completely general. The LS -coupling form of $\bar{S}_i(j_i)$ in terms of reduced electric dipole matrix elements is derived in Sec. III.

Before continuing with this formal development, however, we discuss in the rest of this section the physical significance of the angular momentum transfer and, in particular, its role as a probe of anisotropic electron-ion interactions.

The physical significance of angular momentum transfer as a direct probe of anisotropic electron-ion interactions is illustrated in Fig. 1. In this analysis it is convenient to think of the photoionization process as having two stages, namely, an initial stage A of photoabsorption proper and a subsequent stage B of escape of the photoelectron from the rest of the atom. The angular momentum transfer is always equal to the difference between the angular momentum input to the atom (namely, the angular momentum $j_\gamma = 1$ of the electric dipole interaction) and the angular momentum output from the atom (namely, the photoelectron's final-state orbital momentum \vec{l}). Thus the angular momentum transfer, $\vec{j}_t = \vec{j}_\gamma - \vec{l}$, is the net angular momentum transferred to (or deposited in) the target by the photoionization process. (Note that since we consider experiments in which the photoelectron's spin \vec{s} is not measured, \vec{s} is included as part of the angular momentum of the residual target.) The allowed values of j_t , however, are different in the two stages of the photoionization process.

In the initial stage A [illustrated in Fig. 1(a)] the photoabsorption imparts $j_\gamma = 1$ unit of orbital momentum to the photoelectron, which has initial orbital momentum \vec{l}_0 (in an independent-particle model), yielding a final orbital momentum $\vec{l}' = \vec{l}_0 + \vec{j}_\gamma$. Therefore in stage A the angular momentum transferred to the target is

$$\vec{j}_t = \vec{j}_\gamma - \vec{l}' = -\vec{l}_0, \quad (13)$$

where the magnitude j_t' has the single value $j_t' = l_0$. Furthermore, owing to parity conservation, $l' = l_0 \pm 1$, and hence $j_t' = l_0$ is a parity-favored angular momentum transfer.

During the subsequent escape of the photoelectron in stage B additional angular momentum transfers can arise, within the allowed range determined by Eq. (2), from anisotropic interactions of the photoelectron with the rest of the target. In this report we consider only spin-independent interactions in LS coupling. Therefore the interaction in stage B is that between the orbital motion

of the photoelectron and the net orbital motion of the electrons of the residual ion core, as illustrated in Fig. 1(b). This interaction produces a dynamical coupling of the respective orbital momenta \vec{l}' and \vec{L}_c . Owing to the resulting angular momentum exchanges \vec{k} between the photoelectron and the core, only the total angular momentum \vec{L} is conserved. (It is because of this dependence on L that we call these interactions anisotropic.) In particular, the photoelectron orbital momentum can change from \vec{l}' to \vec{l} during the departure of the photoelectron from the atom, in which case the angular momentum transfer is no longer $j_t' = -l_0$ but

$$\vec{j}_t \equiv \vec{j}_\gamma - \vec{l} = \vec{j}_t' - \vec{k}, \quad (14)$$

as illustrated in Fig. 1(c). Note that even if the magnitudes of \vec{l}' and \vec{L}_c remain unchanged, a precession (albeit quantized in units of \hbar) about the total orbital momentum \vec{L} is sufficient to produce a change in the magnitude of \vec{j}_t .

It is at this point that the connection between the present formulation and that of the Cooper-Zare independent-particle model⁴ emerges most clearly. The Cooper-Zare model treats the residual ion core as a spectator to the photoionization process. That is, stage B is ignored altogether, in which case only the single (parity-favored) angular momentum transfer $j_t = l_0$ arises. In addition, the amplitudes $\bar{S}_\pm(j_t = l_0)$ assume limiting forms [cf. Eq. (35) below] which, when substituted in Eq. (10), give the Cooper-Zare formula for the asymmetry parameter. These points are developed in detail in the following sections.

III. PHOTOIONIZATION AMPLITUDES $\bar{S}_i(j_i)$ IN LS COUPLING

Our task in this section is to obtain the LS -coupling form of the reduced electric dipole matrix element in Eq. (12). Before specializing to LS coupling, however, let us consider the problem in general. The form of the reduced dipole matrix element in Eq. (12) is inconvenient for numerical

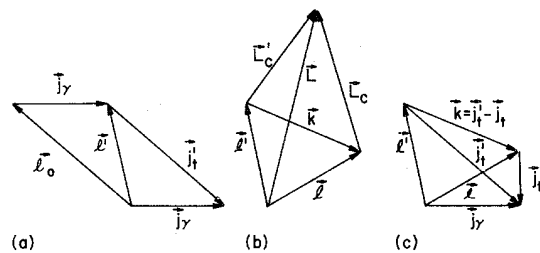


FIG. 1. Illustration of the origin of multiple angular momentum transfers in atomic photoionization reactions. See text for discussion.

calculation for two reasons. First, the final state $((J_c s)J_{cs}l, J-)$ is defined in terms of the dissociation-channel quantum numbers appropriate to the electron-ion system at infinite separation. In general it is more convenient to calculate the electric dipole matrix element for transition to one of the electron-ion eigenchannel states $(\alpha J|$, where α denotes the eigenchannel coupling scheme. Second, it is much more convenient to calculate *real* matrix elements, and for this reason a transformation to the standing-wave representation is desirable.

For these reasons, we expand the dipole matrix element in Eq. (12) as follows³:

$$\begin{aligned} & ((J_c s)J_{cs}l, J- \| P^{[1]} \| \alpha_0 J_0) \\ &= i^{-1} e^{i\sigma(J_c l)} \sum_{\alpha} ((J_c s)J_{cs}l | \alpha)^J e^{i\delta(\alpha)} (\alpha J \| P^{[1]} \| \alpha_0 J_0). \end{aligned} \quad (15)$$

Each term in the summation in Eq. (15) comprises three elements:

(i) The *phase factor* $i^{-1} e^{i[\sigma(J_c l) + \delta(\alpha)]}$, which effects the change from incoming-wave to standing-wave normalization. Here $\sigma(J_c l)$ is the Coulomb phase,

$$\sigma(J_c l) = \arg \Gamma(l+1-i/\sqrt{\epsilon}), \quad (16)$$

which depends on the binding energy $I(J_c)$ of the residual ion fine-structure level J_c through the photoelectron kinetic energy ϵ measured in Rydbergs:

$$\epsilon = \hbar\omega - I(J_c). \quad (17)$$

The phase $\delta(\alpha)$ is the photoelectron phase shift with respect to Coulomb waves in the eigenchannel α and represents the effect of short-range electron-ion interactions.

(ii) The *real transformation coefficients* $((J_c s)J_{cs}l | \alpha)^J$ which relate the eigenchannel coupling scheme to the dissociation channel coupling scheme.

(iii) The *real reduced dipole matrix elements* $(\alpha J \| P^{[1]} \| \alpha_0 J_0)$.

Thus far Eq. (15) and all preceding equations are exact for electric dipole transitions. Approximations must be made, however, in the representation of the eigenchannels $(\alpha J|$ and their phase shifts $\delta(\alpha)$ as well as in the representation of the initial state $|\alpha_0 J_0\rangle$. We proceed in the rest of this section to derive the *LS*-coupling form of Eq. (15) and then to reduce that further by assuming the use of radial one-electron wave functions appropriate for given term levels of the ion core and the electron-ion system.

In *LS* coupling α_0 and α are given by

$$\alpha_0 = L_0 S_0, \quad \alpha = (L_c l) L (S_c s) S. \quad (18)$$

Implicit in the definition of α_0 is that we have an atomic configuration having a single open shell l_0^N , where l_0 is the orbital angular momentum and N is the occupation number. Similarly α implicitly indicates the configuration of the final state after photoionization, which is of the form $l_0^{N-1} l$. The transformation coefficient in Eq. (15) may be found either algebraically¹⁶ or graphically¹⁷ to be

$$\begin{aligned} & ((L_c S_c) J_{cs} l | (L_c l) L (S_c s) S)^J \\ &= (-1)^{2J_{cs} + (L_c + l + L) + (S_c + s + S)} \hat{J}_c \hat{J}_{cs} \hat{L} \hat{S} \\ & \times \begin{Bmatrix} L_c & S_c & J_c \\ s & J_{cs} & S \end{Bmatrix} \begin{Bmatrix} L & S & J \\ J_{cs} & l & L_c \end{Bmatrix}. \end{aligned} \quad (19)$$

Finally, we must evaluate the *LS*-coupling form of the reduced dipole matrix element

$$(\alpha J \| P^{[1]} \| \alpha_0 J_0) = ((L_c l) L (S_c s) S, J \| P^{[1]} \| L_0 S_0, J_0). \quad (20)$$

This evaluation may be carried out graphically,¹⁸ but in what follows we shall proceed algebraically.

The first step in the evaluation of Eq. (20) is to make a fractional parentage expansion of the initial state,

$$\begin{aligned} |L_0 S_0, J_0\rangle &= \sum_{\bar{L}_0 \bar{S}_0} |(\bar{L}_0 l_0) L_0 (\bar{S}_0 s) S_0, J_0\rangle \\ & \times (l_0^{N-1} \bar{L}_0 \bar{S}_0, l_0 \| l_0^N L_0 S_0). \end{aligned} \quad (21)$$

Since the ionization process is spin independent, the second step is to split off the geometrical dependence of the matrix element in Eq. (20) on spin and total angular momentum quantum numbers¹⁹:

$$\begin{aligned} & ((L_c l) L (S_c s) S, J \| P^{[1]} \| (\bar{L}_0 l_0) L_0 (\bar{S}_0 s) S_0, J_0) \\ &= (-1)^{L+S+J_0+1} \hat{J} \hat{J}_0 \begin{Bmatrix} J & J_0 & 1 \\ L_c & L & S_0 \end{Bmatrix} \delta(S_c, \bar{S}_0) \delta(S, S_0) \\ & \times ((L_c l) L \| P^{[1]} \| (\bar{L}_0 l_0) L_0). \end{aligned} \quad (22)$$

The third step is to reduce the matrix element of the electric dipole operator to a one-electron matrix element by factoring out the geometrical dependence on core and total orbital momenta²⁰:

$$\begin{aligned} & ((L_c l) L \| P^{[1]} \| (\bar{L}_0 l_0) L_0) \\ &= N^{1/2} (-1)^{L_c + l_0 + L + 1} \hat{L} \hat{L}_0 \begin{Bmatrix} L & L_0 & 1 \\ l_0 & l & L_c \end{Bmatrix} \\ & \times \delta(L_c, \bar{L}_0) (l \| P^{[1]} \| l_0). \end{aligned} \quad (23)$$

In this equation the factor $N^{1/2}$ is a weight factor due to the presence of N equivalent electrons in the initial state.^{18, 21} The last step is to factor the reduced one-electron matrix element into its radial and angular parts²²:

$$(l \| P^{[1]} \| l_0) = (l \| C^{[1]} \| l_0) R_{\epsilon l}^{L_c S_c L}, \quad (24)$$

$$R_{\epsilon l}^{L_c S_c L} \equiv \int_0^\infty dr \psi_r((L_c S_c) \epsilon l, LS | r) \mathcal{W} \psi_t(n_0 l_0, L_0 S_0 | r). \quad (26)$$

where the angular part is

$$(l \| C^{[1]} \| l_0) = (-1)^l \tilde{u}_0 \begin{pmatrix} l & 1 & l_0 \\ 0 & 0 & 0 \end{pmatrix} \quad (25)$$

and the radial part is

Note particularly that this radial part is calculated using radial wave functions dependent *dynamically* on the angular momentum and spin quantum numbers of the initial state, of the final state, and of the residual ion core.

Putting Eqs. (21)–(26) together, we find for the reduced dipole matrix element in Eq. (20)

$$\begin{aligned} & ((L_c l) L(S_c s) S, J \| P^{[1]} \| L_0 S_0, J_0) \\ &= N^{1/2} (l_0^{N-1} L_c S_c, l_0 \| l_0^N L_0 S_0) (-1)^{L_c + s_0 + J_0 + 1} \hat{J}_0 \hat{L}_0 \hat{L}_0 \tilde{u}_0 \begin{Bmatrix} J & J_0 & 1 \\ L_0 & L & S \end{Bmatrix} \begin{Bmatrix} L & L_0 & 1 \\ l_0 & l & L_c \end{Bmatrix} \begin{pmatrix} l & 1 & l_0 \\ 0 & 0 & 0 \end{pmatrix} R_{\epsilon l}^{L_c S_c L}. \end{aligned} \quad (27)$$

Finally, substituting Eqs. (15), (19), (20), and (27) into Eq. (14) and noting that $(-1)^{J_0 - J} = (-1)^{J - J_0}$, we find for the scattering amplitude the following result:

$$\begin{aligned} \bar{S}_l(j_t) &\equiv n(\lambda) N^{1/2} (l_0^{N-1} L_c S_c, l_0 \| l_0^N L_0 S_0) i^{-l} e^{i\sigma(J_c l)} (-1)^{S_c + s_0 + J_0} \tilde{u}_0 \begin{pmatrix} l & 1 & l_0 \\ 0 & 0 & 0 \end{pmatrix} \hat{J}_0 \hat{J}_c \hat{J}_{cs} \hat{L}_0 \hat{S}_0 \begin{Bmatrix} L_c & S_c & J_c \\ s & J_{cs} & S_0 \end{Bmatrix} \\ &\times \sum_L e^{i\delta_{\epsilon l}^{L_c S_c L} R_{\epsilon l}^{L_c S_c L}} \hat{L}^2 \begin{Bmatrix} L_0 & L_c & l_0 \\ l & 1 & L \end{Bmatrix} (-1)^{2J_{cs} + s_0 + l + L} \sum_J (-1)^J \hat{J}^2 \begin{Bmatrix} J_{cs} & J & l \\ 1 & j_t & J_0 \end{Bmatrix} \begin{Bmatrix} J_0 & J & 1 \\ L & L_0 & S_0 \end{Bmatrix} \begin{Bmatrix} S_0 & J & 1 \\ l & L_c & J_{cs} \end{Bmatrix}. \end{aligned} \quad (28)$$

Since we have assumed no dependence of the phase shifts and radial dipole matrix elements on total angular momentum J , the sum over J in Eq. (28) may be performed analytically using the Biedenharn identity²³ to yield the desired expression for the scattering amplitude in LS coupling:

$$\begin{aligned} \bar{S}_l(j_t) &= n(\lambda) N^{1/2} (l_0^{N-1} L_c S_c, l_0 \| l_0^N L_0 S_0) i^{-l} e^{i\delta_{\epsilon l}^{L_c S_c L}} Q(j_t, J_c, J_{cs}) (-1)^{(L_0 + L_c + 1) + (S_c + s_0) + (J_{cs} - J_0 - j_t)} \hat{J}_0 \tilde{u}_0 \begin{pmatrix} l & 1 & l_0 \\ 0 & 0 & 0 \end{pmatrix} \\ &\times \sum_L e^{i\delta_{\epsilon l}^{L_c S_c L} R_{\epsilon l}^{L_c S_c L}} \hat{L}^2 \begin{Bmatrix} L_0 & L_c & j_t \\ l & 1 & L \end{Bmatrix} \begin{Bmatrix} L_0 & L_c & l_0 \\ l & 1 & L \end{Bmatrix}, \end{aligned} \quad (29)$$

where

$$Q(j_t, J_c, J_{cs}) \equiv \hat{L}_0 \hat{S}_0 \hat{J}_c \hat{J}_{cs} \begin{Bmatrix} J_c & L_c & S_c \\ S_0 & s & J_{cs} \end{Bmatrix} \begin{Bmatrix} j_t & L_c & L_0 \\ S_0 & J_0 & J_{cs} \end{Bmatrix}. \quad (30)$$

Though Eq. (29) gives the LS -coupling form of the scattering amplitude, we see that there is a geometrical dependence on the quantum numbers J_{cs} and J_c relating to the fine-structure levels of the ionic core. (We have neglected any dependence of the phase shifts $\delta_{\epsilon l}^{L_c S_c L}$ and radial dipole matrix elements $R_{\epsilon l}^{L_c S_c L}$ on the fine-structure levels.) We consider this dependence on J_{cs} and J_c in turn.

All of the dependence on J_{cs} in Eq. (29) is contained in the geometrical factor $(-1)^{J_{cs} - J_0} \times Q(j_t, J_c, J_{cs})$, which depends additionally on quantum numbers that are either fixed for a given ionization process (e.g., $s, L_0, S_0, L_c, S_c, J_c$) or enter

incoherently in the differential and total cross sections (e.g., j_t in addition to J_{cs}). The square of this factor, with phase +1, enters into the definitions of the cross section [Eq. (6)] and the asymmetry parameter [Eq. (7)], each of which involves a summation over J_{cs} . Accordingly, it is convenient to define a new quantity

$$\bar{Q}(j_t, J_c) \equiv \sum_{J_{cs}} Q(j_t, J_c, J_{cs})^2, \quad (31)$$

which gives the statistical weight with which ionization probability for a given j_t is distributed

among the possible fine-structure levels J_c , since

$$\sum_{J_c} \bar{Q}(j_t, J_c)^2 = 1. \quad (32)$$

Note, however, that in Eq. (29) there is an additional dynamical dependence on J_c arising from the Coulomb phase $\sigma(J_c l)$. Often, though, the fine-structure separations of the residual ion are not resolved. Then $\sigma(J_c l)$ can be taken as independent of J_c , and the dependence of the cross sections and asymmetry parameter on J_c can be removed altogether by application of Eq. (32).

Having obtained the form of the scattering amplitude in LS coupling in Eq. (29), it is instructive to return to our discussion in the last section concerning the role of j_t as a probe of anisotropic interactions, as illustrated in Fig. 1. The allowed values of j_t are those consistent with the triangular relations $\{L_0 L_c j_t\}$ and $\{l j_t\}$, implied by the first $6j$ symbol in Eq. (29). The coupling of the electron to the residual ion [cf. Fig. 1(b)] is reflected in the dependence of the phase shifts $\delta_{el}^{L_c S_c L}$ and dipole matrix elements $R_{el}^{L_c S_c L}$ on the total angular momentum of the electron-ion complex. Only when these phases and matrix elements do not depend on L (i.e., when the electron-ion interaction is isotropic) is j_t restricted to the single value $j_t = l_0$. For in this case the dynamical weight factors in Eq. (29) may be extracted from the summation, since

$$e^{i\delta_{el}^{L_c S_c L}} R_{el}^{L_c S_c L} \xrightarrow{\text{isotropic interaction}} e^{i\delta_{el} R_{el}}, \quad (33)$$

and the summation over L may be performed analytically:

$$\sum_L \hat{L}^2 \begin{Bmatrix} L_0 & L_c & j_t \\ l & 1 & L \end{Bmatrix} \begin{Bmatrix} L_0 & L_c & l_0 \\ l & 1 & L \end{Bmatrix} = \hat{l}_0^{-2} \delta(j_t, l_0). \quad (34)$$

The scattering amplitude then depends only on the final orbital angular momentum l of the photoelectron,

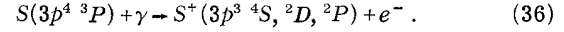
$$\bar{S}_l(j_t = l_0) \propto i^{-l} e^{i(\sigma_{el} + \delta_{el})} \tilde{U}_0 \begin{pmatrix} l & 1 & l_0 \\ 0 & 0 & 0 \end{pmatrix} R_{el}, \quad (35)$$

where the proportionality constant indicates that we have not written down all the other factors from Eq. (29) which depend on quantum numbers that are fixed for a given photoionization process. These other factors do not contribute to the asymmetry parameter in Eq. (10) since they occur in both numerator and denominator and thus cancel out. Setting $j_t = l_0$ in Eq. (10) and substituting the scattering amplitude from Eq. (35) leads to the asym-

metry parameter $\beta(l_0)$ of the Cooper-Zare independent-particle model.⁴

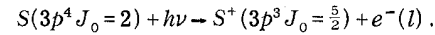
IV. APPLICATION TO SULFUR PHOTOIONIZATION

To illustrate the theory developed in the last two sections we calculate the angular distribution of photoelectrons ionized from atomic sulfur according to the reaction



For each of the residual ion terms $L_c S_c$ we present in Table I the allowed values of photoelectron angular momentum l , angular momentum transfer j_t , reaction parity (where parity change = +1 is favored and parity change = -1 is unfavored), and the allowed values of total angular and spin momenta for the electron-ion system. We see that the 4S -ion term has only the single angular momentum transfer $j_t = l_0 = 1$, but that the 2D and 2P ion terms both have other values of j_t , including parity-unfavored values. Note that for the 2D - and 2P -ion terms the $l=2$ states have two or more allowed values of LS , implying that there will be interference between phase shifts belonging to different final-state channels.

It should be noted that the values of the quantum numbers appearing in Table I are all those obtained from Eqs. (2)–(4) that are consistent with our approximations, given in Sec. III. In particular, angular momentum and parity conservation, represented by Eqs. (2)–(4), allow the photoelectron orbital angular momentum l to have the values 0, 2, 4, and 6 for the reaction



However, in our LS -coupling single-configuration approximation for the ground state the electric dipole operator restricts the photoelectron to $l=0, 2$ [cf. Eq. (25)]. One may obtain $l=4$ photoelectrons in LS coupling by considering the interaction of the ground $3p^4$ configuration with other highly excited configurations, as found by Conneely, Smith, and Lipsky.²⁴ To obtain $l=6$ photoelectrons one must either abandon LS coupling and consider relativistic interactions for which only the total angular momentum is conserved, or else, keep LS coupling but consider the photoionization of highly excited ground states to excited ion states that do not interact with the configuration $3p^3$. This promotion of the photoelectron to $l=4$ or $l=6$ is a specific example of anisotropic interaction as represented schematically in Fig. 1(b).

For conciseness we shall concentrate in what follows on the photoionization reaction leading to the 2D -ion term since it shows the strongest anisotropic electron-ion interactions. For this ion

term the scattering amplitudes $\bar{S}_t(j_t)$ in Eq. (29) for the allowed values of l and j_t listed in Table I are

$$\bar{S}_s(1) = C \frac{1}{3} e^{i(\sigma_s + \delta_s(^3D))} R_{\epsilon_s}(^3D), \quad (37a)$$

$$\bar{S}_d(1) = C(2^{1/2}/5) e^{i\sigma_d} \left[\frac{1}{3} e^{i\delta_d(^3S)} R_{\epsilon_d}(^3S) + \frac{3}{4} e^{i\delta_d(^3P)} R_{\epsilon_d}(^3P) + \frac{7}{12} e^{i\delta_d(^3D)} R_{\epsilon_d}(^3D) \right], \quad (37b)$$

$$\bar{S}_d(2) = C(2^{1/2}/5) e^{i\sigma_d} \left[-\frac{1}{3} e^{i\delta_d(^3S)} R_{\epsilon_d}(^3S) - \frac{1}{4} e^{i\delta_d(^3P)} R_{\epsilon_d}(^3P) + \frac{7}{12} e^{i\delta_d(^3D)} R_{\epsilon_d}(^3D) \right], \quad (37c)$$

$$\bar{S}_d(3) = C(2^{1/2}/5) e^{i\sigma_d} \left[\frac{1}{3} e^{i\delta_d(^3S)} R_{\epsilon_d}(^3S) - \frac{1}{2} e^{i\delta_d(^3P)} R_{\epsilon_d}(^3P) + \frac{1}{6} e^{i\delta_d(^3D)} R_{\epsilon_d}(^3D) \right]. \quad (37d)$$

In Eq. (37) C denotes those constant factors in Eq. (29) that are common to all channels, s and d denote $l=0$ and $l=2$, and $R_{\epsilon_d}(^3S)$, for example, denotes the radial dipole matrix element $R_{\epsilon_d}^{L_c S_c L}$ for $L_c S_c = ^2D$, $L=0$, and $l=2$. Note that we have ig-

TABLE I. Allowed values for the ion-core term level $L_c S_c$, photoelectron orbital angular momentum l , angular momentum transfer j_t , reaction parity $\pi_0 \pi_c (-1)^{j_t}$, and total orbital and spin angular momenta LS for the reaction $S(3p^4 \ ^3P) + h\nu \rightarrow S^+(3p^3 \ ^4S, \ ^2D, \ ^3P) + e^-$.

$L_c S_c$	l	j_t	Parity	LS
4S	0	1	+1 (fav)	3S
4S	2	1	+1 (fav)	3D
2D	0	1	+1 (fav)	3D
2D	2	1	+1 (fav)	$^3D, \ ^3P, \ ^3S$
2D	2	2	-1 (unf)	$^3D, \ ^3P, \ ^3S$
2D	2	3	+1 (fav)	$^3D, \ ^3P, \ ^3S$
2P	0	1	+1 (fav)	3P
2P	2	1	+1 (fav)	$^3D, \ ^3P$
2P	2	2	-1 (unf)	$^3D, \ ^3P$

nored the dependence of the Coulomb phase shifts σ_s and σ_d on ion-core fine-structure levels J_c , as discussed at the end of the previous section.

The asymmetry parameter for photoionization to the 2D -ion term is given by Eqs. (7)–(11):

$$\beta = \frac{3|\bar{S}_d(1)|^2 - 3 \times 2^{1/2} [\bar{S}_d(1)\bar{S}_s(1)^\dagger + \text{c.c.}] - 5|\bar{S}_d(2)|^2 + 2|\bar{S}_d(3)|^2}{3|\bar{S}_s(1)|^2 + 3|\bar{S}_d(1)|^2 + 5|\bar{S}_d(2)|^2 + 7|\bar{S}_d(3)|^2}. \quad (38)$$

In Eq. (38) the common factor C in Eq. (37) cancels in numerator and denominator. As pointed out in the last section, Eq. (38) reduces to the Cooper-Zare result for β^4 when the phase shifts $\delta_{\epsilon_d}^{L_c S_c L}$ and radial dipole matrix elements $R_{\epsilon_d}^{L_c S_c L}$ become independent of $L_c S_c L$. It is of interest to see how this occurs for this particular example. Note first that $\bar{S}_d(2) \rightarrow 0$ and $\bar{S}_d(3) \rightarrow 0$ in Eqs. (37c) and (37d) when the phase shifts and dipole matrix elements become identical. We also see that the squared modulus of each of these scattering amplitudes, having $j_t \neq l_0 = 1$, is nonzero partly because the resulting factors $\cos[\delta_d(^3S) - \delta_d(^3P)]$, etc., in the cross terms are not unity. These same factors also arise in cross terms of $|\bar{S}_d(1)|^2$ and are partly responsible for changing the value of this modulus from what its (nonzero) value would be in a Cooper-Zare model calculation. For these reasons we regard the magnitude of phase-shift differences $\delta_{\epsilon_d}^{L_c S_c L} - \delta_{\epsilon_d}^{L_c S_c L'}$ to be an indication of the strength of anisotropic electron-ion interactions and hence of the validity of the Cooper-Zare model for β .⁴

The scattering amplitudes in Eqs. (37a) and (37b), having $j_t = l_0 = 1$, contribute to the cross section and the asymmetry parameter β whether the phase shifts and matrix elements are identical or not. The scattering amplitudes in Eqs. (37c) and (37d), however, having $j_t \neq l_0 = 1$, contribute only when the phases and matrix elements are dif-

ferent from one another. An index of the strength of angular momentum transfers $j_t \neq l_0$ is thus the fraction $[\sigma - \sigma(j_t = l_0)]/\sigma$, where σ is the photoionization cross section and $\sigma(j_t = l_0)$ is the partial cross section corresponding to $j_t = l_0$. For photoionization to the 2D ion term of sulfur this ratio is expressed as

$$\begin{aligned} [\sigma - \sigma(1)]/\sigma = & [5|\bar{S}_d(2)|^2 + 7|\bar{S}_d(3)|^2 / \\ & [3|\bar{S}_s(1)|^2 + 3|\bar{S}_d(1)|^2 \\ & + 5|\bar{S}_d(2)|^2 + 7|\bar{S}_d(3)|^2]. \end{aligned} \quad (39)$$

To evaluate Eqs. (37)–(39) we used continuum Hartree-Fock (HF) wave functions obtained by solving the equations given by Dalgarno, Henry, and Stewart²⁵ using methods discussed fully by Kennedy and Manson.⁶ These continuum wave functions depend on both the ionic term level and the total orbital angular momentum. Discrete HF single-particle orbitals for the neutral atom and for the ion were obtained from the tabulation of Clementi.²⁶

For comparison, we have also carried out a Cooper-Zare type of calculation employing Herman-Skillman²⁷ (HS) wave functions. The continuum HS wave functions are calculated in the average sulfur potential appropriate to the ground configuration as tabulated by Herman and Skillman.²⁷ These wave functions depend neither on the ion

core level $L_c S_c$ nor on the total angular momentum L , and thus the phase shifts and radial dipole matrix elements depend only on ϵl , and Eq. (37) reduces to

$$\bar{S}_s(1) = C \frac{1}{3} e^{i(\sigma_s + \delta_s)} R_{\epsilon s}, \quad (40a)$$

$$\bar{S}_d(1) = C(2^{1/2}/3) e^{i(\sigma_d + \delta_d)} R_{\epsilon d}, \quad (40b)$$

$$\bar{S}_d(2) = \bar{S}_d(3) = 0. \quad (40c)$$

Note that since the HS continuum wave functions do not depend on the ionic term level the asymmetry parameters for each ion term, when plotted versus photoelectron kinetic energy ϵ , are identical. Discrete wave functions for both the ion and the atom were taken to be the tabulated HS neutral-atom discrete wave functions.

In Fig. 2 we have plotted HF phase shifts $\delta_{\epsilon d}^{L_c S_c L}$ for the ionic term level 2D as a function of photoelectron kinetic energy ϵ . The three allowed values of L are listed in Table I. These phase shifts differ by as much as 0.7 rad, indicating that anisotropic electron-ion interactions are significantly large.

In Fig. 3 we have plotted the three asymmetry parameters corresponding to the three alternative ionic term levels resulting from photoionization of the sulfur atom. Contrary to the Cooper-Zare model, these asymmetry parameters are significantly different from one another when plotted as a function of photoelectron kinetic energy. In this plot the length formula for electric dipole transitions has been used since this is the correct one for HF calculations.²⁸ In Table II, however, we list calculated HF asymmetry parameters using both length and velocity formulas for the dipole matrix elements in order to show that for most energies listed the differences between the asymmetry parameters for different ion terms are larger than the length and velocity difference for a given ion term. We also list for comparison the

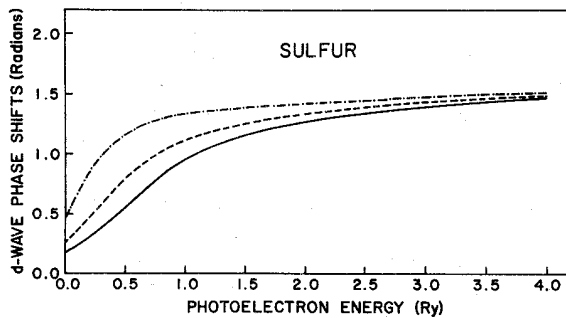


FIG. 2. Hartree-Fock d -wave phase shifts $\delta_{\epsilon d}^{L_c S_c L}$ for the 2D sulfur-ion term versus photoelectron kinetic energy ϵ for alternative allowed values of L . Solid line, $L=0$ [i.e., the state $3p^3(^2D)\epsilon d^3S$]; dashed line, $L=1(^3P)$; dot-dashed line, $L=2(^3D)$.

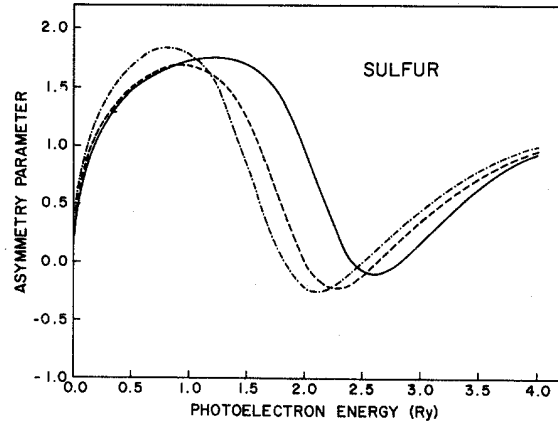


FIG. 3. Asymmetry parameters $\beta(^3P \rightarrow L_c S_c)$ for the photoionization reactions $S(3p^4^3P) \rightarrow S^+(3p^3 L_c S_c) + e^-$ as a function of photoelectron kinetic energy. Solid line, 4S ionic term; dashed line, 2D ; dot-dashed, 2P .

β parameter calculated using HS wave functions and the Cooper-Zare formula for β [i.e., Eq. (40)]. The HS wave functions are quite different from the HF wave functions, and thus the HS asymmetry parameter does not seem to be an "average" of the HF asymmetry parameters at low energies.

In Table III we have plotted HF and HS cross sections for photoionization of sulfur. Note that the HS cross section is a total cross section and would correspond to the sum of the three HF partial cross sections at a given photon energy. However, we have plotted the HF partial cross sections as functions of photoelectron kinetic energy for comparison with Table II. Comparing Tables II and III, we see that the largest differences in the asymmetry parameters occur for energies $1.5 \leq \epsilon \leq 2.1$ Ry. This is just before the Cooper minima²⁹ in the cross sections, which occur in the region $2.1 \leq \epsilon \leq 2.8$ Ry. The cross sections in the region $1.5 \leq \epsilon \leq 2.1$ Ry are of the order of 10^{-18} cm², and thus measurement of β for the different thresholds should be experimentally possible, if not for sulfur then for some other element. Simply put, we wish to emphasize that the differences we have found between the asymmetry parameters for the different ionic term levels are not dependent on being at a cross-section minimum. Indeed, as seen in Fig. 3 and Tables II and III there are measurable differences between $\beta(^3P \rightarrow ^4S)$ and $\beta(^3P \rightarrow ^2P)$ in the energy range $0.1 \leq \epsilon \leq 0.8$ Ry, where the cross sections are of the order of 10^{-17} cm².

Finally, in Fig. 4 we examine the influence of angular momentum transfers $j_i \neq l_0$ on the asymmetry parameter and partial cross section for photoionization to the 2D ionic term level. The solid line represents the asymmetry parameter given by Eq. (38) and plotted also in Fig. 3. The

TABLE II. HF asymmetry parameters for the reactions $S(3p^4^3P) + h\nu \rightarrow S^+(3p^3^4S, ^2D, ^2P) + e^-$ as a function of photoelectron kinetic energy ϵ using dipole-length (velocity) formula and comparison with HS asymmetry parameter.

ϵ (Ry)	β ($^3P \rightarrow ^4S$)	β ($^3P \rightarrow ^2D$)	β ($^3P \rightarrow ^2P$)	HS β
0.00	0.144(0.054)	0.176(0.248)	0.274(0.360)	0.35
0.05	0.551(0.495)	0.584(0.614)	0.705(0.739)	0.74
0.10	0.769(0.734)	0.805(0.814)	0.942(0.950)	0.96
0.20	1.052(1.048)	1.090(1.082)	1.248(1.229)	1.25
0.40	1.365(1.407)	1.402(1.402)	1.594(1.564)	1.61
0.60	1.543(1.619)	1.574(1.604)	1.779(1.761)	1.78
0.80	1.659(1.761)	1.670(1.735)	1.851(1.856)	1.82
1.00	1.734(1.849)	1.693(1.782)	1.793(1.820)	1.68
1.25	1.772(1.860)	1.581(1.640)	1.472(1.488)	...
1.50	1.714(1.665)	1.234(1.131)	0.847(0.775)	0.10
1.80	1.408(0.944)	0.497(0.185)	0.044(-0.078)	-0.24
2.10	0.724(0.016)	-0.127(-0.270)	-0.250(-0.268)	-0.02
2.30	0.211(-0.210)	-0.223(-0.207)	-0.173(-0.136)	0.18
2.60	-0.108(-0.029)	-0.029(0.076)	0.094(0.155)	0.44
2.80	-0.013(0.189)	0.170(0.273)	0.281(0.339)	0.60
3.00	0.177(0.393)	0.360(0.447)	0.450(0.500)	0.74
4.00	0.933(1.003)	0.972(0.989)	0.996(1.009)	1.10
8.00	1.576(1.526)	1.513(1.523)	1.522(1.526)	1.54
15.00	1.697(1.629)	1.614(1.636)	1.622(1.636)	1.62
30.00	1.582(1.561)	1.534(1.564)	1.537(1.564)	1.54

dashed line represents β calculated according to Eq. (38), but setting $\bar{S}_a(2) = \bar{S}_a(3) = 0$. Note that the result is *not* the Cooper-Zare expression for β since we still use Eq. (37b) for $\bar{S}_a(1)$, and thus the dependence of phase shifts and radial dipole ma-

TABLE III. HF cross sections for the reactions $S(3p^4^3P) + h\nu \rightarrow S^+(3p^3^4S, ^2D, ^2P) + e^-$ as a function of photoelectron kinetic energy ϵ using dipole length (velocity) formula and comparison with HS cross section. All cross sections are in units of 10^{-18} cm^2 .

ϵ (Ry)	σ ($^3P \rightarrow ^4S$)	σ ($^3P \rightarrow ^2D$)	σ ($^3P \rightarrow ^2P$)	σ (HS)
0.00	13.82(8.92)	27.43(19.23)	20.87(15.40)	58.00
0.05	14.61(9.11)	28.00(19.18)	20.78(15.04)	58.78
0.10	15.03(9.10)	27.46(18.42)	19.73(14.04)	57.24
0.20	15.04(8.65)	24.21(15.65)	16.03(11.09)	48.33
0.40	12.85(6.85)	15.42(9.38)	8.43(5.62)	24.60
0.60	9.47(4.75)	8.70(5.05)	4.01(2.62)	11.84
0.80	6.27(2.97)	4.67(2.61)	1.92(1.23)	4.73
1.00	3.84(1.73)	2.47(1.34)	0.95(0.61)	2.00
1.25	1.96(0.82)	1.14(0.59)	0.44(0.27)	...
1.50	0.97(0.38)	0.56(0.29)	0.23(0.15)	0.58
1.80	0.42(0.17)	0.29(0.17)	0.15(0.10)	0.52
2.10	0.21(0.11)	0.20(0.15)	0.13(0.10)	0.57
2.30	0.16(0.10)	0.19(0.16)	0.13(0.10)	0.61
2.60	0.14(0.12)	0.20(0.18)	0.14(0.12)	0.68
2.80	0.14(0.13)	0.21(0.19)	0.15(0.12)	0.71
3.00	0.15(0.14)	0.22(0.20)	0.15(0.13)	0.75
4.00	0.22(0.17)	0.27(0.23)	0.17(0.14)	0.80
8.00	0.17(0.12)	0.18(0.15)	0.11(0.09)	0.42
15.00	0.05(0.04)	0.06(0.05)	0.04(0.03)	0.15
30.00	0.01(0.01)	0.01(0.01)	0.01(0.01)	0.03

trix elements on $L_c S_c L$ is still important. We see from Fig. 4 that values of $j_t \neq l_0 = 1$ lower β as much as 0.2 units in the neighborhood of $\epsilon \approx 1.0$ Ry. The dot-dashed curve is a plot of the ratio $[\sigma - \sigma(1)]/\sigma$ given in Eq. (39). Values of $j_t \neq l_0 = 1$ contribute as much as 8% to the partial cross section in the neighborhood of $\epsilon \approx 1.5$ Ry.

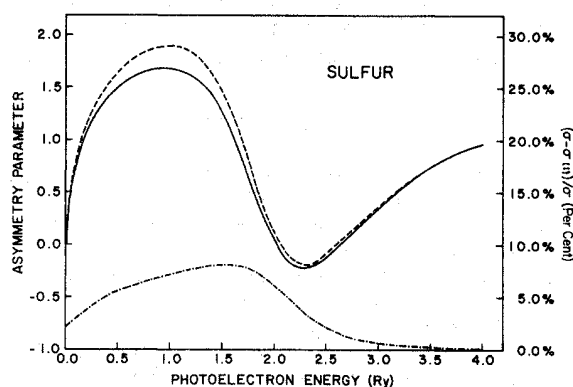


FIG. 4. Dependence of the asymmetry parameter β ($^3P \rightarrow ^2D$) and cross section σ ($^3P \rightarrow ^2D$) for the 2D ion term of sulfur on angular momentum transfers $j_t \neq l_0$ as a function of photoelectron kinetic energy. Left-hand scale refers to (i) the solid line denoting β and (ii) the dashed line denoting $\beta(j_t = l_0 = 1)$, both for the $^3P \rightarrow ^2D$ transition. Right-hand scale refers to the dot-dashed line which denotes the ratio $[\sigma - \sigma(j_t = 1)]/\sigma$ for the $^3P \rightarrow ^2D$ transition.

V. DISCUSSION AND CONCLUSIONS

We have shown that anisotropic electron-ion interactions in atomic sulfur lead to measurable differences between photoelectron angular distribution asymmetry parameters corresponding to alternative ionic term levels. Similar effects are expected for most open-shell atoms. A measure of the strength of anisotropic electron-ion interactions is the difference between phase shifts for alternative final-state channels. In atomic sulfur these phase-shift differences are as large as 0.7 rad. A separate study of atomic oxygen⁷ found phase-shift differences of only 0.2 rad and asymmetry parameters that were nearly identical for each ionic term level. However, atomic oxygen and the other second-row elements are regarded as exceptions, since they are too light to have strong interactions, and atomic sulfur is regarded as more typical of open-shell atoms in general. Our choice of atomic sulfur for study was purely a matter of convenience. We know of no experimental data on photoelectron angular distributions for an open-shell atom. We emphasize, however, that we expect the magnitude of the difference between asymmetry parameters and the magnitude of the cross sections to be experimentally measurable for many open-shell atoms.

For the particular case of atomic sulfur we have found that angular momentum transfers $j_i \neq l_0$, which do not arise in the Cooper-Zare model,⁴ contribute only a small but nevertheless significant amount to the asymmetry parameters and to the cross sections. We simply do not know whether this will hold true for other open-shell atoms. The contributions to the asymmetry parameter from angular momentum transfers $j_i = l_0$ are, how-

ever, quite different from those in the Cooper-Zare model, which has *only* $j_i = l_0$ contributions, since the phase-shift differences are so large in the different final-state channels [cf. Eqs. (37b) and (40b)].

For closed-shell atoms our formulas reduce rigorously to those of the Cooper-Zare model. Unfortunately, nearly all experimental measurements of photoelectron angular distributions known to us are for closed-shell atoms. Considering the importance of photoelectron angular distributions to such diverse areas as radiation dosimetry (e.g., δ -ray spectrum)³⁰ and the physics of the upper atmosphere (e.g., conjugate point phenomena)³¹ as well as to theoretical physics, as emphasized in this paper, we feel that experimental data on photoelectron angular distributions for open-shell atoms would be most valuable.

Last, we point out that our formulas for photoelectron angular distributions have been derived for any electron-ion coupling scheme, but worked out in detail only for *LS* coupling. In general the electron-ion interaction is best described in an intermediate coupling scheme, particularly in semiempirical calculations.³ Nearly all *ab initio* atomic calculations, however, use the *LS*-coupling scheme, and it is for these calculations that our formulas have been worked out most fully. While we have calculated phase shifts and dipole matrix elements in HF approximation, other more accurate procedures (e.g., many-body perturbation theory, random-phase approximation, etc.) may be used to compute these quantities for use in our formulas for the asymmetry parameter. Similarly, while we have ignored fine-structure splittings of the ionic core, these may easily be included in angular-distribution calculations using our formulas as discussed at the end of Sec. III.

*Supported in part under National Aeronautics and Space Administration Grant No. NGR 28-004-021.

†Supported in part under National Science Foundation Grant No. GP-38905.

¹U. Fano and D. Dill, Phys. Rev. A **6**, 185 (1972).

²D. Dill and U. Fano, Phys. Rev. Lett. **29**, 1203 (1972).

³D. Dill, Phys. Rev. A **7**, 1976 (1973).

⁴J. Cooper and R. N. Zare, *Lectures in Theoretical Physics*, edited by S. Geltman, K. T. Mahanthappa, and W. E. Britten (Gordon and Breach, New York, 1969), Vol. XI-C, pp. 317-37.

⁵For representative experimental measurements see, e.g., T. A. Carlson, G. E. McGuire, A. E. Jonas, K. L. Cheng, C. P. Anderson, C. C. Lu, and B. P. Pullen, in *Proceedings of an International Conference on Electron Spectroscopy, Asilomar, California, 1971*, edited by D. A. Shirley (North-Holland, Amsterdam, 1972), pp. 207-231; P. Mitchell and K. Codling, Phys.

Lett. **38A**, 31 (1972); M. J. Lynch, A. B. Gardner, and K. Codling, Phys. Lett. **40A**, 349 (1972); A. Niehaus and M. W. Ruf, Z. Physik **252**, 84 (1972); M. J. Lynch, K. Codling, and A. B. Gardner, Phys. Lett. **43A**, 213 (1973); M. J. Van der Wiel and C. E. Brion, J. Electron Spectrosc. Related Phenomena **1**, 439 (1973); G. R. Branton and C. E. Brion, J. Electron Spectrosc. Related Phenomena **3**, 123 (1974).

⁶For representative theoretical calculations see, e.g., J. W. Cooper and S. T. Manson, Phys. Rev. **177**, 157 (1969); S. T. Manson and J. W. Cooper, Phys. Rev. A **2**, 2170 (1970); S. T. Manson, Phys. Rev. Lett. **26**, 219 (1971); D. J. Kennedy and S. T. Manson, Phys. Rev. A **5**, 227 (1972); S. T. Manson, J. Electron Spectrosc. Related Phenomena **1**, 413 (1973); S. T. Manson, Chem. Phys. Lett. **19**, 76 (1973).

⁷A. F. Starace, S. T. Manson, and D. J. Kennedy, Phys. Rev. A **9**, 2453 (1974).

- ⁸L. Lipsky, in *Fifth International Conference on the Physics of Electronic and Atomic Collisions: Abstracts of Papers* (Nauka, Leningrad, 1967), pp. 617-18.
- ⁹L. Lipsky (unpublished).
- ¹⁰V. L. Jacobs and P. G. Burke, *J. Phys. B* 5, L67 (1972).
- ¹¹D. Dill, S. T. Manson, and A. F. Starace, *Phys. Rev. Lett.* 32, 971 (1974).
- ¹²C. N. Yang, *Phys. Rev.* 74, 764 (1948).
- ¹³See Ref. 6, first paper.
- ¹⁴J. A. R. Samson, *J. Opt. Soc. Am.* 59, 356 (1969); *Philos. Trans. R. Soc. Lond. A* 268, 141 (1970); J. A. R. Samson and J. L. Gardner, *J. Opt. Soc. Am.* 62, 856 (1972).
- ¹⁵V. Schmidt, *Phys. Lett.* 45A, 63 (1973).
- ¹⁶U. Fano and G. Racah, *Irreducible Tensorial Sets* (Academic, New York, 1959), Chap. 12.
- ¹⁷A. P. Yutsis, I. B. Levinson, and V. V. Vanagas, *Theory of Angular Momentum* (Israel Program for Scientific Translations, Jerusalem, 1962), Chapter V.
- ¹⁸J. S. Briggs, *Rev. Mod. Phys.* 43, 189 (1971).
- ¹⁹Reference 16, Eq. (15.7).
- ²⁰Reference 16, Eq. (15.7').
- ²¹U. Fano, *Phys. Rev.* 140, A67 (1965).
- ²²B. W. Shore and D. H. Menzel, *Principles of Atomic Spectra* (Wiley, New York, 1968), Eq. (10.4).
- ²³See Ref. (16), Eq. (1.2) or Ref. (17), Eq. (A6.53).
- ²⁴M. J. Conneely, K. Smith, and L. Lipsky, *J. Phys. B* 3, 493 (1970).
- ²⁵A. Dalgarno, R. J. W. Henry, and A. L. Stewart, *Planet. Space Sci.* 12, 235 (1964).
- ²⁶E. Clementi and C. Roetti, *At. Data Nucl. Data Tables* 14, 177 (1974).
- ²⁷F. Herman and S. Skillman, *Atomic Structure Calculations* (Prentice-Hall, Englewood Cliffs, N. J., 1963).
- ²⁸A. F. Starace, *Phys. Rev. A* 3, 1242 (1971); 8, 1141 (1973).
- ²⁹J. W. Cooper, *Phys. Rev.* 128, 681 (1962).
- ³⁰H. Bichsel (private communication).
- ³¹S. T. Manson, D. J. Kennedy, A. F. Starace, and D. Dill, *Planet. Space Sci.* 22, 1535 (1974).