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Elliptic and Circular Dichroism Effects in Two-Photon Double Ionization of Atoms

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An ab initio parametrization of the two-photon double ionization amplitude from an s2 subshell of an atom in a 1S state is presented and used to predict two light polarization effects on photoelectron angular distributions that do not exist in single-photon double ionization: (i) elliptic dichroism and (ii) circular dichroism at equal energy sharing. Estimates for He show large magnitudes for these effects, which provide a means for polarization control of double ionization by vacuum ultraviolet light.

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The physics of processes involving simultaneous ejection of several electrons from atoms and molecules has attracted interest since the 1960s owing to the insight into electron correlations that such studies provide. Single-photon double ionization (SPDI) of He (which can be measured in electron-electron or electron-recoil coincidence experiments) has been a research focus over the past decade. At present there exists a wealth of knowledge concerning electron correlations for this fundamental three-body problem [1]. Recent advances in generating vuv light of high intensity (than provided by synchrotrons) make possible studies of multielectron ejection due to multiphoton interactions of atomic electrons with vuv radiation. Two-photon double ionization (TPDI) of He has recently attracted much attention, as its study provides new insights into two-electron ejection dynamics (see, e.g., Refs. [2–5]). In particular, in contrast to SPDI, in which the P-wave channel of a two-electron continuum is populated, the TPDI process allows one to probe the S- and D-wave channels. The study of TPDI also provides a “bridge” between SPDI and multiphoton nonsequential double ionization (NSDI) in optical laser fields [6]. Experimentally, TPDI of He was investigated using the 27th harmonic of a Ti:sapphire laser with a photon energy of 41.8 eV and a peak intensity of 1014 W/cm2 [7]. So far all studies of TPDI have been limited to the case of linearly polarized light.

In this Letter, we study how the shape of the photoelectron angular distribution in TPDI [i.e., of the triply-differential cross section (TDCS)] depends upon the handedness of (in general elliptically polarized) vacuum ultraviolet (vuv) radiation. In SPDI, this dependence was established in 1992 [8], and is termed the circular dichroism (CD) effect; it is maximal for circularly polarized light and vanishes when two electrons are ejected with equal energies. In TPDI, we show that in addition to the CD effect, there exists also a qualitatively different, elliptic dichroism (ED) effect: in particular geometries that we specify, the shape of the TDCS depends on the helicity of the vuv light only for elliptic polarization (i.e., the effect disappears for pure circular polarization). We also show that in TPDI by circularly polarized light (i.e., when the ED effect vanishes), the CD effect persists even when both electrons are ejected with equal energies. In our numerical estimates of the TDCS for nonsequential TPDI of He, we consider photons of energy 45 eV (the 29th harmonic of a Ti:sapphire laser) and take into account their interaction with He perturbatively, which is adequate up to intensities of the order of 1014 W/cm2 [4].

The TDCS for TPDI of an atom by radiation having an electric field vector F(r, t) = F Re[e exp[i(k · r − ωt)]] (e · e′ = 1) and a photon flux density |j| = cF2/(8πℏω) (defined as dσ = dW/j2, where dW is the transition rate) is given in atomic units by

\[
\frac{d^3\sigma}{d\Omega_1 d\Omega_2 dE_2} = \sigma = |\mathcal{A}|^2,
\]

where \(\mathcal{A} = \frac{\pi}{\omega} p_1 p_2/|c(\omega^2)|\), \(p_1\) and \(p_2\) are the photoelectron momenta, and \(E_2 = p_2^2/2\). The amplitude \(A\) for a two-photon transition from an initial s2(1S) state \(\Phi_i\) to the two-electron continuum state \(\Psi^{(−)}_{p_1p_2}\) involves the two-electron Green’s function (TEGF) \(G(E_i + \omega)\),

\[
A = \langle \Psi^{(−)}_{p_1p_2} | (\mathbf{e} \cdot \mathbf{D}) G(E_i + \omega) (\mathbf{e} \cdot \mathbf{D}) | \Phi_i \rangle,
\]

where \((\mathbf{e} \cdot \mathbf{D}) = \mathbf{e} \cdot (\nabla_{\mathbf{r}_1} + \nabla_{\mathbf{r}_2})\) is the electric dipole operator (in velocity form) of the electron-photon interaction. Using the standard expansion of the final state wave function and the TEGF in Eq. (2) in bipolar harmonics [9], the evaluation of angular integrals in Eq. (2) by means of the Wigner-Eckart theorem yields

\[
A = \sum_{L=0,2} \sum_{L_1L_2} \frac{\delta^{L_2L_1}_0}{\sqrt{3(2L+1)}} \left[ (\mathbf{e} \cdot \mathbf{e})_{L_1} \cdot Y_{L_1}^{(L_2)}(\mathbf{\hat{p}}_1, \mathbf{\hat{p}}_2) \right],
\]

where \(\delta^{L_2L_1}_0 = \langle p_1p_2(l_1l_2L)|\mathbf{D}_{L_1}^{(L_2)}(E_i + \omega)\mathbf{D}|\Phi_i \rangle\) is the second-order two-electron reduced matrix element in which \(g_1^{L_2}(E)\) describes “intermediate” states of the electron pair of 1P\(^o\) symmetry with orbital angular momenta \(s_1\) and \(s_2\) (\(s_1 + s_2 = \text{odd}\) and \(l_1 + l_2 = \text{even}\) due to parity conservation). After rewriting the bipolar harmonic \(Y_{L_2}^{(L_1)}(\mathbf{\hat{p}}_1, \mathbf{\hat{p}}_2)\) in Eq. (3) in terms of irreducible tensors composed of vectors \(\mathbf{\hat{p}}_1\) and \(\mathbf{\hat{p}}_2\) [10] and evaluating the...
scalar product of these tensors with photon polarization tensors $(\mathbf{e} \otimes \mathbf{e})_b$ [9], we obtain the following invariant form for the TPDI amplitude:

$$A = f_0(\mathbf{e} \cdot \mathbf{e}) + f_1(\mathbf{e} \cdot \mathbf{p}_1)^2 + f_2(\mathbf{e} \cdot \mathbf{p}_2)^2 + f_3(\mathbf{e} \cdot \mathbf{p}_1)(\mathbf{e} \cdot \mathbf{p}_2),$$

(4)

where $f_0(p_1, p_2), f_1(p_1, p_2), f_2(p_1, p_2) \equiv f_1(p_2, p_1), and f_3(p_1, p_2)$ are four polarization-invariant amplitudes that depend only upon the photoelectron energies and $x \equiv \cos \theta = (\mathbf{p}_1 \cdot \mathbf{p}_2)$. They have the following general expressions in terms of Legendre polynomials, $P_l(x)$, and their derivatives, $P_l^{(k)}(x) = (d/dx)^k P_l(x)$:

$$f_1 = \sum_{l=0}^{\infty} \frac{2(l+1)(l+1)(l+1)(l+2)}{(2l)(2l+1)(2l+2)} M_{ll}^2 - \frac{1}{\sqrt{3}(l+1)(l+2)(2l+3)} M_{ll+2}^2 - \frac{1}{\sqrt{3}(l-1)(l-2)(2l+1)} M_{l-2}^2 P_l^{(2)}(x),$$

(5)

$$f_2 = \sum_{l=1}^{\infty} \frac{2}{\sqrt{l(l+1)(2l+1)}} \left[ \frac{2l+1}{\sqrt{2l(2l+1)(2l+3)}} M_{ll}^2 [2xP_l^{(2)}(x) - P_l^{(1)}(x)] + \frac{1}{\sqrt{3}} (M_{l-1}^2 + M_{l+1}^2) P_l^{(2)}(x) \right],$$

(6)

$$f_3 = -\frac{1}{3} \left[ \sum_{l=0}^{\infty} \sqrt{2l+1} M_{ll}^0 P_l(x) + f_1 + f_2 + xf_3 \right].$$

(7)

In Eqs. (5)–(7), $M_{ll}^0 = (4\pi)^{-1} \sum_{s=-l}^{s+l} \sum_{s_1=-l}^{s_1+l} d_{l_1 l_2}^{s_1 s_2} d_{s_1}^{s_2} d_{s_2}^{s_1}$ is dynamical factor relevant to the L-wave continuum channel. It depends only upon photoelectron energies and the dynamical model used to describe electron correlations. Note that the S-wave continuum channel factors, $M_{ll}^0$, enter only the amplitude $f_0$, which does not contribute in the case of circular polarization [since $(\mathbf{e} \cdot \mathbf{e}) = 0$ for this case]. The amplitudes $f_1$ and $f_2$ are symmetric in $p_1, p_2$: $f_0(p_1, p_2) = f_0(p_2, p_1), f_1(p_1, p_2) = f_1(p_2, p_1)$. By introducing $g_{s,a}$, we obtain an alternative set of four symmetrized amplitudes, where the antisymmetric amplitude, $g_{s,a}$, vanishes at equal energy sharing, $p_1 = p_2$. The parametrization of $A$ in Eq. (4) is similar to that for absorption of two photons in laser-assisted electron-atom scattering [11] (because the same set of vectors occurs in both problems). Note that an alternative parametrization of the TPDI amplitude [5] involves five parameters, indicating an implicit redundancy.

To analyze the TDCS, we parametrize the photon polarization vector as $\mathbf{e} = (\mathbf{e} + i\eta \hat{\mathbf{e}})/\sqrt{1 + \eta^2} (-1 \leq \eta \leq 1)$, where $\hat{\mathbf{e}} = [\mathbf{k} \times \mathbf{e}]$ and $\mathbf{e}$ and $\mathbf{k}$ indicate the directions of the major axis of the polarization ellipse and the photon wave vector, $\mathbf{k}$. The ellipticity $\eta$ is related to the degrees of linear and circular polarization, $l$ and $\xi$: $l = (1 - \eta^2)/(1 + \eta^2)$, $\xi = 2\eta/(1 + \eta^2) = i(\mathbf{k} \cdot [\mathbf{e}^* \times \mathbf{e}])$; $l^2 + \xi^2 = 1$. With these definitions and Eq. (4), the TDCS in Eq. (1) may be expressed as (cf. [11]):

$$\sigma = \sigma_0 + \xi\sigma_{CD} + \xi\sigma_{ED},$$

(8)

where $\sigma_0$ is invariant under the transformation $\mathbf{e} \rightarrow \mathbf{e}^*$ (i.e., under a change in sign of $\xi$), while the other two terms are dichroic (i.e., they change sign when $\xi \rightarrow -\xi$). The term proportional to $\xi$ is maximal at $\xi = \pm 1$ and describes the CD effect, while the term proportional to $\xi l$ vanishes for purely circular polarization ($\xi = \pm 1, l = 0$) and describes the ED effect; it is maximal for $l = [\xi] = 1/\sqrt{2}$. The general expressions for $\sigma_0$, $\sigma_{CD}$, and $\sigma_{ED}$ in Eq. (8) for unequal energy sharing will be published elsewhere; $\sigma_{CD}$ involves only the amplitudes $f_1$, $f_2$, while $\sigma_{ED}$ also involves $f_0$. Here we present explicit expressions for $\sigma_{CD}$ and $\sigma_{ED}$ for the case of equal energy sharing $[p_1 = p_2, \mathbf{p}_\pm = (\mathbf{p}_1 \pm \mathbf{p}_2)/2, (\mathbf{p}_+ \cdot \mathbf{p}_-) = 0]$:

$$\sigma_{CD}^{(eq)} = 2.4\text{Im}(f_{1b}^* g_{s,a})(\mathbf{k} \cdot [\mathbf{p}_+ \times \mathbf{p}_-])(\mathbf{k} \cdot \mathbf{p}_+)(\mathbf{k} \cdot \mathbf{p}_-),$$

(9)

$$\sigma_{ED}^{(eq)} = 4.4\text{Im}(g_{s,a}^* f_{1b})(\mathbf{k} \cdot [\mathbf{p}_+ \times \mathbf{p}_-])(\mathbf{e} \cdot \mathbf{p}_+)(\mathbf{e} \cdot \mathbf{p}_-) + 2.4\text{Im}[f_{1b}^* f_{1b}]$$

$$\times [(\mathbf{e} \cdot \mathbf{p}_+)(\mathbf{e} \cdot \mathbf{p}_-) + (\mathbf{e} \cdot \mathbf{p}_-)(\mathbf{e} \cdot \mathbf{p}_+)] - \sigma_{CD}^{(eq)}.$$  

(10)

One sees that dichroic effects in TPDI originate from interference of real and imaginary parts of polarization-invariant components of the generally non-Hermitian TPDI amplitude, in agreement with arguments in Ref. [12] on the origin of dichroic effects in photoprocesses involving unpolarized atoms. The parametrizations (4)–(10) constitute the main results of this Letter. In general, the dichroic terms (9) and (10) are not small compared to $\sigma_0$ in (8). Evaluating these terms is theoretically challenging. Existing calculations on TPDI treat only linearly polarized light, for which numerical solution of the Schrödinger equation represents a spatial five-dimensional problem; the various total cross section results for $\omega = 45$ eV differ by more than a factor of 2 (cf. Fig. 1 of [5]). A direct numerical solution for elliptically polarized light represents a spatial six-dimensional problem. Accurate experimental measurements thus may provide a valuable benchmark for theory. In order to estimate the magnitudes of the CD and ED effects in TPDI of He for such experiments, we use lowest-order perturbation theory (LOPT) in the interelectron interaction, $1/r_{12}$, to account for electron correlations, as done successfully to treat both NSDI by strong laser fields [6] and SPDI [13]. In SPDI, it was shown [13] that the knocknot mechanism is dominant for excess energies of tens of eV and asymmetric energy sharing. In the TPDI amplitude, there are two knocknot terms, in which both
photos are absorbed by the same electron; their diagrams are shown in Fig. 1. The diagram in Fig. 1(b) turns out to be negligible. Besides the knockout diagrams, the LOPT TPDI amplitude involves also diagrams corresponding to absorption of a single photon by each of the two electrons. Our estimates show that, for the photon energy considered in this Letter (~45 eV), these diagrams are suppressed compared to the knockout ones except in the narrow interval of angles θ near θ = 0. Moreover, to obtain correct results for this region of θ, the inclusion of high-order effects in 1/r_{12} is necessary owing to the divergence of these LOPT diagrams at equal energy sharing. Such calculations require the development of an appropriate regularization procedure and significant computational effort; thus we confine our estimates of dichroic effects in TPDI to those originating from knockout processes. These give the right order of magnitude for the TDCS in the case of linearly polarized photons, as shown in Fig. 2 where we make comparisons with the time-dependent close coupling (TDCC) [3] and convergent close coupling (CCC) [5] results. (Note that in Ref. [5] and here, the CCC results are scaled by a factor of 7). Our absolute LOPT estimates agree in order of magnitude with both the absolute TDCC and the scaled CCC results. (Our total TPDI cross section results for 40 eV ≤ ω ≤ 45 eV—not shown—agree to within ±50% with the absolute TDCC results.) Our predicted angular structure agrees qualitatively with both results for θ₁ = 0° and 30°, while it agrees only with the scaled CCC results for θ₁ = 90°. For θ₁ = 60° our LOPT

FIG. 2 (color online). TDCS for TPDI of He by linearly polarized photons of energy ω = 45 eV. Full (red) curves, present LOPT knockout results; dotted (blue) curves, TDCC results [3]; dashed (green) curves, CCC results [5] scaled to the TDCC results as in [5].

FIG. 3 (color online). Geometries for measurement of the ED and equal energy sharing CD effects in TPDI. (a) Orthogonal geometry for measurement of ED at p₁ = p₂. (b) Coplanar geometry for measurement of ED, with one electron (p₁) detected in the xy plane and the other (p₂) detected in the plane spanned by k and p₁. (c) Geometry for measurement of CD at p₁ = p₂ with fixed ϕ₂.
results predict a more compressed angular structure than do the others.

In Figs. 4 and 5 we present our LOPT estimates for the ED and CD effects, respectively, for three geometries in which only one of the effects is present [cf. Fig. 3]. The left column of Fig. 4 shows polar plots of TDCSs for left-handed (χ > 0) and right-handed (χ < 0) photons, while the right column shows the relative ED parameter, δED ≡ [σ(±1) − σ(−1)]/[σ(±1) + σ(−1)], as a function of the corresponding polar angles. In Figs. 4(a)–4(d) we use the orthogonal geometry, while in Figs. 4(e) and 4(f) we use the coplanar geometry. Figure 4(d) has two geometric zeros determined by the condition that one of the vectors 𝑝_+ and 𝑝_- is orthogonal to either 𝑒 or 𝑝 (i.e., 𝜑_2 = 180°, 0°, or 𝜑_2 = 360°, 0°, so that 𝜑_2 = 135° or 315° for 𝜑_1 = 45°); there are also four dynamical zeros. These zeros are caused by cancellations of terms involving Im(0) in Eq. (10): their positions within the same geometry depend on the excess energy. Two zeros (symmetric with respect to θ = 180°) of δED in Fig. 4(f) also have a dynamical origin [since (𝐴 × 𝐵) = 0 for this geometry, they originate from a zero of the factor Im(0) in (9), whose position depends on the excess energy].

The important difference between the CD effect in SPDI and TPDI is that CD in SPDI vanishes for equal energy sharing, while in TPDI it remains nonzero (except in orthogonal and coplanar geometries). In Fig. 5, we present TDCSs and CD parameters, δCD ≡ [σ(±1) − σ(−1)]/[σ(±1) + σ(−1)], for two cases having the geometry in Fig. 3(c): 𝜑_2 = 45° and 𝜑_2 = 90°. All zeros of δCD have geometrical origins; that means that the term Im(0) in (9) does not change its sign in the interval 0° ≤ χ < 360° [for 𝜑_2 = 90°, this fact is obvious since the mutual angle θ is fixed (θ = 90°) for this case].

To conclude, we have presented model-independent representations for the transition amplitude and the TDCS for TPDI from the s² subshell of an atom in a 1S state. Based on these representations, we predict dichroic (ED and CD) effects that are nonzero for both symmetric and asymmetric excess energy sharing. Our analysis shows that the ED term in the TDCS involves a more feature-rich interference of polarization-invariant amplitudes than the CD term. Thus (in contrast to SPDI) measurements of TPDI with elliptically polarized light (with 0 < |ξ| < 1) are more informative. Our LOPT knockout estimates for He show that both CD and ED effects should be observable in TDCS measurements.

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