

Lawrence Berkeley National Laboratory

Recent Work

Title

SULFUR 1s CORE-LEVEL PHOTOIONIZATION OF SF₆

Permalink

<https://escholarship.org/uc/item/9d6164vn>

Author

Ferrett, T.A.

Publication Date

1986-02-01



Lawrence Berkeley Laboratory

UNIVERSITY OF CALIFORNIA

Materials & Molecular Research Division

Submitted to Physical Review A

SULFUR 1s CORE-LEVEL PHOTOIONIZATION OF SF₆

T.A. Ferrett, D.W. Lindle, P.A. Heimann,
H.G. Kerkhoff, U.E. Becker, and D.A. Shirley

February 1986

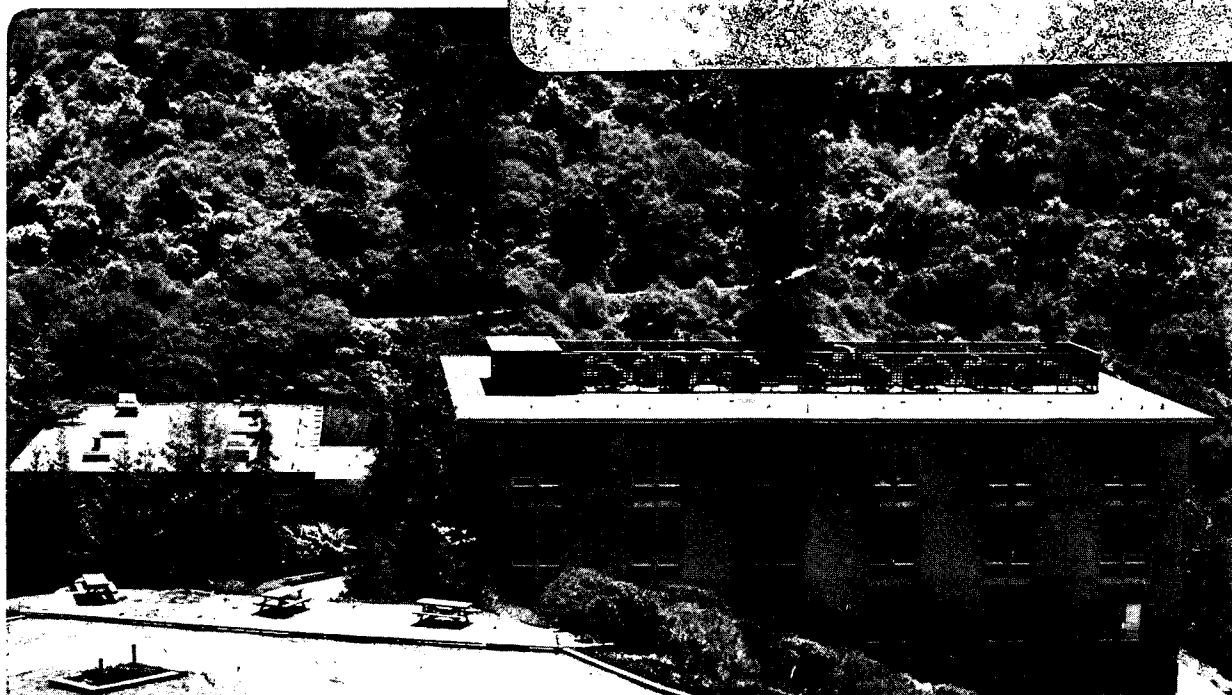
LAWRENCE
BERKELEY LABORATORY

FEB 27 1986

LIBRARY AND
DOCUMENTS SECTION

TWO-WEEK LOAN COPY

*This is a Library Circulating Copy
which may be borrowed for two weeks.*



LBL-18825
e

DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.

SULFUR 1s CORE-LEVEL PHOTOIONIZATION OF SF₆

T.A. Ferrett, D.W. Lindle, P.A. Heimann,
H.G. Kerkhoff,* U.E. Becker,* and D.A. Shirley

Materials and Molecular Research Division
Lawrence Berkeley Laboratory
and
Department of Chemistry
University of California
Berkeley, California 94720

Photoelectron spectra have been taken between 2460 and 2600 eV photon energy across the discrete and continuum resonances in the vicinity of the sulfur K edge in gaseous SF₆. Results at the below-threshold S 1s → 6t_{1u} resonance indicate that "highly excited" S 2p and S 2s satellites (with two core holes) are the primary autoionization final states of SF₆⁺. An observed asymmetric profile in the S(LVV) Auger angular distribution suggests interference effects in the alignment of these resonantly produced SF₆⁺ ions. Decay of the low-energy S 1s continuum resonances near 2507 eV photon energy into S 2p, S 2s, and/or valence photoemission channels indicates autoionizing character. These features are assigned as doubly excited states, leading to S 1s satellite thresholds observed here for the first time. At higher photon energies, between 2520 and 2570 eV, large oscillations in the S 1s cross section are reproduced well by MSM-X_α calculations, but are not

explained adequately by single-scattering plane-wave extended X-ray absorption fine structure (EXAFS) effects. We speculate that improvements in the description of both the electron-scattering process and the molecular potential are necessary to model the diffractive and nondiffractive (barrier interaction) effects in this energy region.

* Permanent address: Technische Universität Berlin, Institut für Strahlungs- und Kernphysik, Sekr. PN 3-2, Hardenbergstr. 36, D-1000 Berlin 12, West Germany.

I. Introduction

When the first gas-phase photoabsorption measurement near the sulfur K edge in SF₆ (reproduced in Fig. 1) was published in 1966, the appearance of the spectrum was described as being "unusually rugged" from the S 1s threshold (~2490 eV) up to about 2570 eV photon energy.¹ Several years later, a line-up of the S 1s, S 2p, and F 1s core-level photoabsorption spectra of SF₆ referenced to their respective ionization thresholds revealed an interesting correlation of the resonant features in kinetic energy to within a few eV.² This correlation led to the assignment of continuum features in these spectra as e_g and t_{2g} shape resonances,^{2,3} and in particular suggested that some of the "rugged" continuum features in the S 1s spectrum were related to potential-barrier effects.

Theoretically, there have been major advances in the basic understanding of potential-barrier effects in atoms⁴⁻⁷ and, initially with the application of the multiple-scattering method (MSM-X α), in molecules.⁸⁻¹³ In a simple one-electron model, a shape resonance is a pure final-state effect. It should thus occur at approximately the same kinetic energy for different core levels of a given molecule. The continuum electron can be pictured as being trapped temporarily by a centrifugal barrier, producing a resonance at a kinetic energy roughly comparable to the barrier height. The large observed photoabsorption intensities of discrete states also fit into this scheme; the unusual potential can enhance the discrete molecular

orbital transitions at the expense of Rydberg excitations and nonresonant continuum intensity. Because the core-level spectra of SF_6 exhibit strong resonances that have been associated with an unusually high potential barrier,^{2,3} SF_6 has become a prototypical example of potential-barrier effects and shape resonances.¹⁴

The present assignments for the S 1s absorption spectrum of SF_6 shown in Fig. 1 are as follows. We use the letter assignments in Fig. 1 throughout the paper to refer to features "a" through "g". The interpretation of the only intense below-threshold resonance (b) as a discrete excitation of a S 1s electron to the unoccupied $6t_{1u}$ level is straightforward.² The SF_6 energy-level diagram in Fig. 2 illustrates this transition. For the S 1s continuum, the shape-resonance model was used to assign two features at ~2493 (c) and 2506 eV (d) photon energy.² However, this interpretation is not so straightforward because these transitions [$\text{S } 1s(1a_{1g}) \rightarrow e_g, t_{2g}$] are symmetry-forbidden in the simple shape-resonance picture.² Furthermore, we note that assignment of resonance features based solely on photoabsorption data can be uncertain; effects due to satellite continua and doubly excited resonances may not be distinguishable from those caused by continuum shape resonances.^{15,16} Finally, the nature of the higher-energy features (f and g) remains unexplained.

To examine in more detail the core-level excitation of the "classic" potential-barrier molecule, we undertook a gas-phase photoemission study of the features near the sulfur K edge of SF_6 .

using synchrotron radiation and time-of-flight electron analysis. By investigating the behavior of the individual photoemission channels, we hoped to ascertain the nature of the "rugged" continuum structure and the autoionization decay characteristics of the $S\ 1s \rightarrow 6t_{1u}$ discrete resonance. No other photoemission measurements have been reported on the S and F core levels in SF_6 , but related photoemission experiments on the outer-valence orbitals in the photon-energy range 16-54 eV have been reported. These low-photon-energy results show that the assigned t_{2g} shape resonance exhibits unusual behavior by coupling to neighboring symmetry-forbidden valence channels.^{17,18} In addition, the lack of evidence for the e_g shape resonance in the valence subshells remains puzzling.

Related core-level photoemission results in other molecules include recent studies of autoionization below the carbon and nitrogen K edges of CO and N_2 . These studies have revealed interesting decay channels leading to singly-charged ions, identified by peaks termed "spectator" satellites.¹⁹⁻²² The dominance of these channels is rationalized as follows. A core-level electron is promoted to an unoccupied molecular orbital and remains as a spectator while the remaining electrons decay to fill the core hole.²³ The singly-charged ion configurations thus produced contain an excited electron and two valence holes; these final states are therefore valence satellites. This autoionization decay of the excited neutral in some cases may parallel the Auger decay of the corresponding ion

with the excited electron removed.

We observe similar "spectator" satellite decay channels at the $S\ 1s(1a_{1g}) \rightarrow 6t_{1u}$ excitation (b) below the $S\ 1s$ threshold of SF_6 . The satellites produced contain $S\ 2p$ and/or $S\ 2s$ vacancies. In addition, the subsequent relaxation of the resonantly produced SF_6^+ ions via Auger decay of the L holes permits the measurement of the Auger electron cross sections and, of more interest, the angular distribution asymmetry parameter β . We report the first measurement of the angular distribution of Auger electrons from an ion produced by an autoionization process. Autoionization can produce aligned $+1$ ions, and the subsequent Auger decay can retain some memory of this alignment, which may affect the Auger β .²⁴ No previous angular-distribution measurements have been performed at photon energies high enough such that the single ions produced by autoionization are energetically able to Auger decay.

Turning to the $S\ 1s$ continuum in SF_6 , we have examined the resonant features in the vicinity of 2507 eV (d and e) and at higher energies (f and g). The results near 2507 eV demonstrate that these resonant states decay into channels other than the $S\ 1s$ continuum (e.g. $S\ 2p$, $S\ 2s$, valence). This decay pattern, combined with the observation of $S\ 1s$ correlation-satellite thresholds slightly higher in energy, suggests assignment of these features as arising from doubly excited states.

In order to explain the higher-energy effects and the large oscillations in the cross section between 30 and 70 eV kinetic energy (f and g), the application of recent advances in extended x-ray

absorption fine structure (EXAFS) theory, including spherical-wave²⁵⁻²⁸ and multiple-scattering corrections,^{25,28-32} would probably be required. In addition, we propose that the details of the molecular potential also are important for intermediate-energy electrons which can interact significantly both with the diffuse electron cloud (large r) and with the localized atomic centers (small r). Our interpretation of the S 1s continuum generally indicates that the kinetic energy region 30-100 eV is especially complicated and probably displays neither simple shape-resonance nor EXAFS behavior.

The experimental details of this work are presented in Section II. In Section III, we lay the groundwork for interpretation of the results by describing our photoemission spectra. We discuss the results for the S 1s \rightarrow 6t_{1u} below-threshold resonance in Section IV, and for the S 1s continuum in Section V. Our conclusions appear in Section VI.

II. Experimental

The experiment was performed at the Stanford Synchrotron Radiation Laboratory (SSRL) using photons from "JUMBO", a double-crystal monochromator operating with Ge(III) crystals on Beam Line III-2. Synchrotron radiation photoionized an effusive beam of SF₆ molecules, and Auger and photoelectrons were detected at 0° and 54.7° relative to the photon polarization direction using the double-angle time-of-flight (DATOF) method.³³⁻³⁵ This method takes

advantage of the time structure of the SPEAR storage ring and allows determination of relative partial cross sections, branching ratios, and angular distributions for Auger and photoelectrons.

The angular distribution of photoelectrons from a randomly oriented sample using a linearly polarized photon source can be described in the dipole approximation by Yang's theorem as:³⁶

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

where θ is the angle between the photon polarization and the electron emission direction, $P_2(\cos\theta)$ is the second Legendre polynomial, and $\sigma(h\nu)$ and $\beta(h\nu)$ are the partial cross section and angular distribution asymmetry parameter, respectively. At the "magic angle" of $\theta=54.7^\circ$, peak intensities are proportional to the partial cross section $\sigma(h\nu)$. In order to measure $\beta(h\nu)$, it suffices to detect electrons at only one other angle, 0° in our case. The degree of linear polarization assumed in Eq. (1) is 100%. The polarization of the photons from "JUMBO" is unknown, but probably somewhat greater than 90%. This uncertainty leads to an absolute error in the β values reported here of less than ± 0.05 because of our calibration procedure.³⁴ Finally, though the dipole approximation is assumed valid for this experiment, deviations may occur with very high-energy photons in the keV range. However, such effects should be small for these particular measurements, where the dipole processes are strong.

Peak intensities obtained at 54.7° must be corrected for fluctuations in sample pressure and photon intensity in order to

determine relative partial cross sections. The photon intensity was monitored by measuring the total electron yield of graphite with a channeltron placed at the back of the sample chamber. A 1500 Å thick Al window separated the monochromator vacuum (10^{-10} torr) from the sample-chamber pressure (10^{-4} torr), which was recorded with a capacitance manometer. We estimate that systematic errors in our relative cross section (not represented by the statistical error bars in our plots) are 10% or less.

Time-of-flight spectra were taken from $h\nu=2460$ to 2600 eV with a monochromator bandpass of about 2.4 eV. Energy calibration of the monochromator was accomplished by scanning over the below-threshold resonance ($S\ 1s \rightarrow 6t_{1u}$) in SF_6 at 2486 eV.¹ The 54.7° analyzer transmission as a function of electron kinetic energy was determined using atomic argon, by comparing Ar 1s photoemission intensity to Ar LMM Auger intensity. Variations in this ratio were attributed to changes in the analyzer transmission at the kinetic energy of the Ar 1s peak. The asymmetry-parameter measurements were calibrated in the kinetic-energy range of about 7-100 eV by measuring the Ar 1s peak at varying kinetic energies, and assuming the $\beta(\text{Ar } 1s)=2.0$. At intermediate kinetic energies (110-200 eV), the S(LVV) Auger peak was used as a calibrant by assuming arbitrarily a nonresonant β value of zero. For very high kinetic energy electrons (>1800 eV), the F 1s photoemission peak in SF_6 was used for calibration with an assumed β of 2.0.

During the experiment there were significant fluctuations in the

position of the photon beam which changed the relative analyzer efficiency by as much as 20% and the 54.7° analyzer transmission by a factor of two. Within a set of spectra unaffected by beam movement, we observed that the intensity and asymmetry parameter for the F(KVV) Auger peak were relatively constant in the photon-energy range 2460-2600 eV. Thus, corrections were made in each spectrum for beam fluctuations using the F Auger peak as a standard. These experimental complications involving the movement of the photon beam also made calibration for the β measurements at low kinetic energies especially difficult, leading to possible systematic errors not represented by the statistical error bars shown in our asymmetry-parameter plots. We estimate that the uncertainty in the S 1s β between 2530 and 2550 eV is ± 0.15 . Likewise, at the lower kinetic energies (below 12 eV), the analyzer efficiency changes dramatically, also increasing the β uncertainty to ± 0.15 .

A representative S 1s time-of-flight spectrum is shown in Fig. 3. Count rates for the S 1s peak were 4-30 counts/sec with 1000 sec collection times. There are F 1s, S 2p, S 2s, and valence photoemission contributions to the high energy peak A, as well as S(KLL, KLV, and KVV) Auger intensity. The relative importance of these components will be discussed in more detail in Sec. III.

III. Peak Contributions in the S 1s Photoemission Spectra

Interpretation of experimental results in Sec. IV and V requires knowledge of the processes that contribute to the peaks in the S 1s

photoelectron spectra. These contributions must be considered together because some of them are not resolved in our spectra. The kinetic-energy resolution in the TOF spectra is limited by time dispersion due to the finite dimensions of the photon beam, and to a lesser extent by the time resolution inherent in the time-of-flight technique. For the measurements on the JUMBO monochromator, this corresponds to ~9% of the kinetic energy, or as much as 180 eV at 2000 eV kinetic energy.

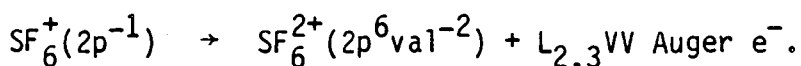
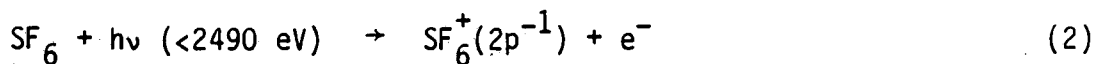
The TOF spectrum in Fig. 3, taken 91 eV above the S 1s threshold at 2490 eV, includes the S 1s main-line and satellite peaks, the S(LVV) Auger peak, the F(KVV) Auger peak, and peak A. Figure 4 illustrates a second spectrum taken below the S 1s threshold, showing the very weak S(L₁L_{2,3}V) Auger peak. Several of the peaks in Fig. 3 arise from only one process at all photon energies used in this study. Specifically, the S 1s main-line and satellite peaks and the peak comprised of the F(KVV) Auger transitions remain well-resolved from all other peaks.

There are, on the other hand, several dissimilar components in peak A and in the S(LVV) and S(L₁L_{2,3}V) Auger peaks. The processes that contribute to each of these observed peaks in three energy regions are listed in Table I. We define the energy ranges as follows: below the sulfur K edge, between the S 1s main-line and the first satellite threshold, and above the first S 1s satellite threshold.

Peak A has the most complicated structure because it includes all

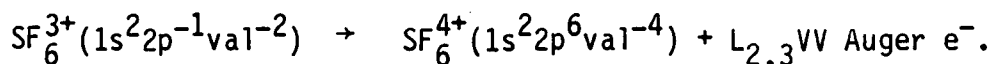
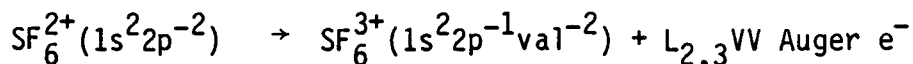
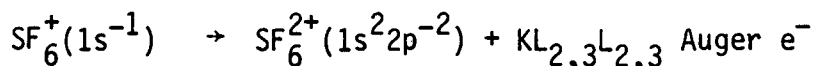
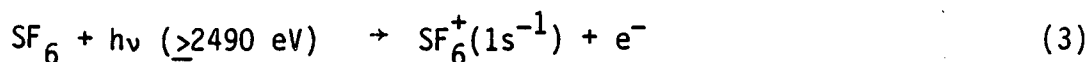
photoemission main lines and satellites with binding energies less than 700 eV. Below the sulfur K edge, the photoemission contributions to peak A are from F 1s, S 2s, S 2p, and valence main lines and satellites. Some "highly excited" S 2p and S 2s satellite configurations ($2p^{-2}v^*$, $2s^{-2}v^*$, and $2s^{-1}2p^{-1}v^*$, where the valence state v^* is probably either the $6t_{1u}$ or $6a_{1g}$ molecular orbital) also contribute to peak A in the vicinity of the S 1s $\rightarrow 6t_{1u}$ resonance at 2486 eV. There is a slight double peak structure to peak A, as shown in the spectrum in Fig. 4, because the F 1s peak is at a lower kinetic energy, while the remaining resonant satellite contributions are unresolved at higher energy. Finally, as the photon energy is increased to above the S 1s main-line and S 1s satellite thresholds, the primary Auger decay from S 1s hole states produces high-kinetic-energy S(KLL, KLV, and KVV) Auger electrons which also contribute to the peak A intensity.

Below the sulfur K edge, the $S(L_{1,2,3}VV)$ and $S(L_1L_{2,3}V)$ peaks result from Auger decay of resonantly and nonresonantly produced SF_6^+ ions ($2s^{-1}$, $2p^{-1}$, $2p^{-2}v^*$, etc...) with S 2p and/or S 2s holes, e.g.,



The especially complicated Auger decay cascades occurring at the discrete $S\ 1s \rightarrow 6t_{1u}$ resonance will be presented in the next section. In particular, the characteristics of the Auger decay of aligned SF_6^+ ions produced by autoionization will be discussed.

Moving above the $S\ 1s$ main-line and satellite thresholds, the $S(LVV)$ and $S(L_1L_{2,3}V)$ Auger peaks include the subsequent decay of $S\ 1s$ hole states, e.g.,



Note that in the particular example given above, it is also possible to produce a tertiary $S(L_{2,3}VV)$ Auger electron in the decay of SF_6^{3+} to SF_6^{4+} . Because tertiary and occasionally quaternary Auger decay processes can occur in filling the $S\ 2p$ and $S\ 2s$ core holes, the observed $S(L_1L_{2,3}V)$ and $S(LVV)$ Auger peaks will include all of these emitted Auger electrons. By writing these simple illustrative processes, we do not wish to imply that the complex Auger decay necessarily occurs stepwise. Dissociation has also been ignored in the above scheme, but will be considered later.

The reader is referred to Table I for specific peak contributions in each energy range appropriate to the discussion of results in Secs. IV and V.

IV. The Below-Threshold $S\ 1s \rightarrow 6t_{1u}$ Resonance

For the $S\ 1s \rightarrow 6t_{1u}$ resonance, the possible decay channels to SF_6^+ are described in Sec. A. We examine in Sec. B the energies and shapes of peak A and the $S(L_1L_{2,3}V)$ and $S(LVV)$ Auger peaks to deduce qualitatively the important decay channels leading to SF_6^+ . We show in Sec. C that an analysis of the sulfur Auger cross-section ratio LVV/LLV implies a dominance of a particular single-ion resonant configuration, the $2p^{-2}_v^*$ "spectator" satellite. In Sec. D, we present asymmetry-parameter results for peak A and the LVV Auger peak in the $S\ 1s \rightarrow 6t_{1u}$ resonance region. The observed asymmetric profile for the $S(LVV)$ Auger β is discussed with respect to ion alignment²⁴ and its implications for the Auger-electron angular distribution.

A. Resonant Decay Channels to SF_6^+

At this resonance, the excited neutral state $1s^{-1}6t_{1u}$ will decay to any continua that are energetically accessible: $F\ 1s$, $S\ 2p$, $S\ 2s$, and valence main-line and satellite final states of SF_6^+ .³⁷ The intensity in peak A below the $S\ 1s$ threshold

includes contributions from all of these channels. The below-threshold $S\ 1s(1a_{1g}) \rightarrow 6t_{1u}$ resonance appears in the cross section for both peak A and S(LVV) Auger (Fig. 5). The energy-level diagram in Fig. 2 illustrates these decay channels (solid lines).

The many SF_6^+ states to which the excited $1s^{-1}6t_{1u}$ state can autoionize are listed generically in Table II. Our photoemission spectra indicate no resonant enhancement in the F 1s channel [as mirrored in the F(KVV) Auger peak], and qualitatively there is very little enhancement of peaks with binding energies below 150 eV. Therefore, we have considered only the SF_6^+ decay channels which have S 2p and/or S 2s holes. Columns 1-4 of Table II describe these available continuum channels in terms of the configurations, decay types, and approximate binding energies of the states. We use Auger notation in column 3 only to denote the type of autoionization decay to each SF_6^+ photoemission channel.³⁸

Each available channel can be described as a S 2s or S 2p main line or satellite. The satellites can be split further into two groups. First, the S 2p and S 2s satellite configurations with binding energies about 10-40 eV above the respective main-line thresholds have a valence electron promoted to an unoccupied molecular orbital. Decay to the main lines and to these low-excitation photoemission satellites can be viewed as KLV Auger-type transitions. Furthermore, the transition can either leave the initially excited $6t_{1u}$ electron as a spectator (KLV) or involve it as a participant (KLV*) in the decay. In this context, decay to the main-line and

some satellite channels must involve the excited $6t_{1u}$ electron, while the satellite configurations with a $6t_{1u}$ electron are "spectator" satellites. The nonresonant intensity for low-excitation-energy satellites may vary from a few percent to as much as 25% relative to the main-line intensity.³⁹

The second class of satellites, which we shall call "highly excited" satellites, possess two core holes (S 2p and/or S 2s) and an excited electron, and have binding energies in the 350-500 eV range. The excited electron is likely to be in the $6t_{1u}$ orbital, although our limited resolution does not permit confirmation of this. These states are S 2p and S 2s satellites in the sense that they could be formed from SF_6 by core-hole ionization plus excitation from a core orbital to a $6t_{1u}$ or other valence molecular orbital. These "highly excited" satellites are produced from KLL-type decay and can also be termed "spectator" satellites if the excited electron is in the $6t_{1u}$ level. In general, we expect that "highly excited" satellites with excitation energies greater than 100 eV will have negligible nonresonant intensity.

B. Resonant Peak Shapes and Energy Shifts

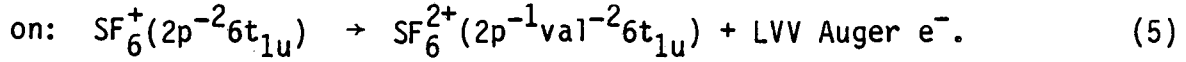
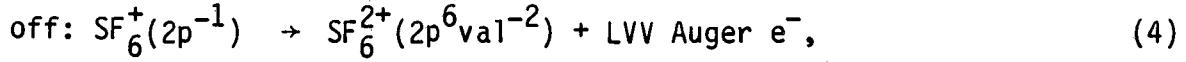
Conclusions about the important decay channels of SF_6^+ in the vicinity of the S 1s $\rightarrow 6t_{1u}$ resonance can be drawn by examining the structure of peak A with reference to Table II. Off resonance (2460 - 2470 eV), we find that peak A consists of a F 1s peak with a

binding energy of ~ 700 eV and a second much less intense peak with a binding energy of approximately 200 eV. This lower binding-energy peak is composed primarily of S 2p and S 2s main lines, with probable small contributions from low-excitation-energy satellites and valence peaks. At 2486 eV (on resonance), peak A appears as a single intense peak with a binding energy of 400(50) eV. The shape of the peak indicates very little intensity at binding energies less than 200 eV. The F 1s peak in Fig. 4 can be seen, partly resolved from the more intense contributions at binding energy ~ 400 eV. We conclude directly from the above qualitative observations that "highly excited" satellites produced by KLL-type transitions are the most important autoionization decay channels for the S 1s \rightarrow 6t_{1u} resonance.

Another indication that the photoemission channels of SF₆⁺ change at the S 1s \rightarrow 6t_{1u} resonance energy can be found in the kinetic-energy shift of the S(LVV) and S(LLV) Auger peaks. The kinetic energies of these broad peaks are higher on resonance than off resonance. Some additional structure on the high kinetic-energy side of the S(LVV) Auger peak also is observed on resonance. The size of the overall energy shift strongly suggests the accessibility of some new SF₆⁺ states other than the S 2p and S 2s main lines. The fact that the shift is to higher kinetic energy is qualitatively rationalized below to be consistent with production of "highly excited" ions.

As an example, we consider the S(L_{2,3}VV) Auger decay of SF₆⁺ to SF₆²⁺ for the single-ion configurations with 2p holes

postulated to be important off and on resonance, e.g.,



The kinetic energies (ϵ) of the S(LVV) Auger electrons in processes (4) and (5) are just the differences in energy between the ions:

$$(\epsilon_{LVV})_{\text{off}} = E(2p^{-1}) - E(2p^6va1^{-2}), \quad (6)$$

$$(\epsilon_{LVV})_{\text{on}} = E(2p^{-2}6t_{1u}) - E(2p^{-1}va1^{-2}6t_{1u}). \quad (7)$$

The observable kinetic-energy shift ($\delta\epsilon$) can be defined as:

$$\begin{aligned} \delta\epsilon &= (\epsilon_{LVV})_{\text{on}} - (\epsilon_{LVV})_{\text{off}} \\ &= [E(2p^{-2}6t_{1u}) - E(2p^{-1})] - \\ &\quad [E(2p^{-1}va1^{-2}6t_{1u}) - E(2p^6va1^{-2})] \\ &= [E_1] - [E_2]. \end{aligned} \quad (8)$$

The energies E_1 and E_2 both correspond superficially to the energy for promotion of a S 2p electron to a $6t_{1u}$ orbital, but E_1 is expected to be greater than E_2 because it requires more energy to remove a molecular core electron (i.e. 2p) from its orbital in the presence of another core hole than in the presence of one or two

diffuse holes in the valence shell, due to differences in screening. This clearly would suggest that E_1 is greater than E_2 . However, the difference in the energy of the $6t_{1u}$ level in the two configurations $2p^{-2}6t_{1u}$ and $2p^{-1}val^{-2}6t_{1u}$ also must be considered.⁴⁰ Because the $6t_{1u}$ orbital is least tightly bound, the dominant effect on its binding energy should be the Coulomb interaction, which is proportional to the ionic charge. Thus, the $6t_{1u}$ orbital in the $SF_6^{2+}(2p^{-1}val^{-2}6t_{1u})$ configuration will experience a larger contraction than in the $SF_6^+(2p^{-2}6t_{1u})$ configuration, with the result that E_2 would be smaller than E_1 . We conclude that $\delta\epsilon$ is positive for the initial Auger decay of 2p hole states [processes (4) and (5)].

This positive shift for $\delta\epsilon$ is in contrast to a shift to lower kinetic energy for Auger satellite transitions from 2-hole initial states to 3-hole final states.⁴¹ However, the initial state for many Auger satellites has the second hole "exterior" to the deepest hole (e.g. KL-LLL in Ne).⁴² The major effect in this case is an increased binding energy for the outer electrons which fill the core, resulting in a lower Auger kinetic energy relative to the "parent" KLL line. For Auger decay of $SF_6^+(2p^{-2}6t_{1u})$, the second 2p hole is in the same shell as the deepest hole, significantly affecting the screening in the core shell as well as in the valence shells.

The presence of Auger electrons from subsequent cascades to higher SF_6^{n+} ions ($n=3-5$) significantly complicates the above discussion. Autoionization to "highly excited" satellites results in

ions with two L-shell core holes, thus allowing for several Auger cascades. The secondary and higher Auger decay steps produce electrons with kinetic energies different from the primary Auger electron due to core and valence screening differences and the degree of involvement of the excited $6t_{1u}$ electron. The additional structure in the S(LVV) Auger peak may arise from such effects. Further explanation of these issues will be warranted when higher resolution spectra become available.

As further support for the importance of the resonant SF_6^+ "highly excited" satellite channels and subsequent Auger cascades, we can compare the S(LVV) Auger peak observed at the S 1s $\rightarrow 6t_{1u}$ resonance with the S(LVV) peak observed above the S 1s threshold. We find that the S(LVV) peak shapes are very similar in these two cases. The primary contributions to the S(LVV) peak above the S 1s threshold (see Table I) must be from secondary and higher-order Auger decay following S 1s ionization, because sulfur K-shell ionization is stronger at these energies than the sulfur L-shell ionization processes. Furthermore, the initial Auger decay of the S 1s hole will be mostly KLL because the S 2p and S 2s orbitals (more than F 1s and valence) reside on the sulfur atom, providing good overlap with the S 1s orbital. This result for the initial decay step also can be deduced by comparison with Ar,⁴³ and shows that higher-order Auger decay from SF_6^{2+} starts primarily from configurations with two holes in the sulfur L shell ($2p^{-2}$, $2s^{-1}2p^{-1}$, and $2s^{-2}$). Still another way to reach the same conclusion is to note that the K-shell

natural linewidth greatly exceeds those in the L shell, and KLL decay is the principle de-excitation process in light elements. The similarity of the S(LVV) peaks above and below threshold indicates that configurations with two L-shell holes (e.g. $2p^{-2}6t_{1u}$) probably provide important decay channels for the S $1s \rightarrow 6t_{1u}$ resonance.

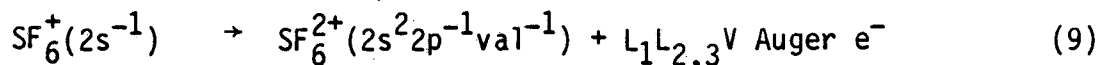
C. Sulfur LVV/LLV Auger Intensity Ratio

The qualitative assertion that "highly excited" satellites dominate the decay of the $1s^{-1}6t_{1u}$ state can be documented further using the observable sulfur Auger intensity ratio LVV/LLV.⁴⁴ Experimentally, the LVV/LLV intensity ratio changes from 3.0(3) off resonance to 25(5) at 2486 eV (Fig. 6). The enhancement in the Auger intensity ratio depends directly on the relative resonant cross sections for the "highly excited" satellite states. In fact, careful examination of the contributions to the observable Auger ratio leads to the identification of the most important "highly excited" satellite channel.

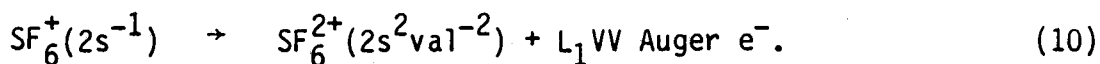
We shall now show that it is possible to express the LVV/LLV Auger ratio in terms of the dominant SF_6^+ cross sections provided that all subsequent Auger decay steps are accounted for. Columns 5-7 of Table II outline the primary through quaternary decay steps to form SF_6^{n+} ($n=2-5$) for each photoemission channel. For this estimate, we first assume that the ions Auger decay quickly enough to fill all

core holes (S 2p and S 2s) before dissociation can occur. A comparison of SF₆ ground-state vibrational lifetimes⁴⁵ with S 2p and S 2s core-hole lifetimes indicates that Auger decay of these holes is probably at least ten times faster than the dissociation rate. If infrequent dissociation does occur, leaving an ion fragment SF_n⁺ (n<6) with a sulfur L hole, the fragment itself will Auger decay to fill the core hole. Thus, the issue of dissociation does not significantly affect the following analysis of the S(LLV) and S(LVV) Auger peaks.

An additional factor necessary for deriving the LVV/LLV ratio is the fraction of S 2s holes which decays via the Coster-Kronig S(L₁L_{2,3}V) Auger pathway,



rather than the S(L₁VV) pathway,



In our analysis, we have assumed that this partitioning into the S(L₁L_{2,3}V) and S(L₁VV) Auger decay pathways is independent of the configuration containing the S 2s hole. For example, the fraction f of S 2s holes decaying via the Coster-Kronig S(L₁L_{2,3}V) channel is assumed to be the same for the configurations SF₆⁺(2s⁻¹) and SF₆⁺(2s⁻²6t_{1u}). The presence of these two possible

Auger-decay channels is evident in column 5 of Table II for each SF_6^+ state containing a S 2s hole.

The intensity ratio LVV/LLV in general (on and off resonance) can be derived in terms of f by adding up all the singly-charged ion cross sections which can subsequently Auger decay via LVV and LLV pathways as:

$$\left[\frac{LVV}{LLV} \right] = \frac{1}{f} \left\{ \frac{\sigma(2p) + \sigma(2p \text{ sat}) + \sigma(2s) + \sigma(2s \text{ sat}) + 2[\sigma(2p^{-2}v^*) + \sigma(2s^{-2}v^*) + \sigma(2p^{-1}2s^{-1}v^*)]}{\sigma(2s) + \sigma(2s \text{ sat}) + \sigma(2p^{-1}2s^{-1}v^*) + 2\sigma(2s^{-2}v^*)} \right\} \quad (11)$$

where $\sigma(x)$ represents the cross section for the SF_6^+ photoemission channel x.

The fraction f can be determined from the nonresonant LVV/LLV Auger ratio and the $\sigma(2p)/\sigma(2s)$ cross-section ratio. Away from the S 1s \rightarrow 6t_{1u} resonance ($h\nu < 2480$ eV), the expression for the LVV/LLV ratio can be simplified with the assumption that the S 2s and S 2p main lines are the only SF_6^+ channels which contribute to the S(LVV) and S(LLV) Auger peaks. With all satellite cross sections assumed negligible, the nonresonant ratio becomes:

$$\left[\frac{LVV}{LLV} \right]_{\text{off}} = \frac{1}{f} \left[\frac{\sigma(2s) + \sigma(2p)}{\sigma(2s)} \right] = \frac{1}{f} \left[1 + \frac{\sigma(2p)}{\sigma(2s)} \right]. \quad (12)$$

The $\sigma(2p)/\sigma(2s)$ cross-section ratio in Eq. (12) can best be estimated from measurements of this ratio in atoms excited with high-energy x-ray sources. From these trends in low-Z elements, we expect that for sulfur, the $\sigma(2s)/\sigma(2p)$ ratio should be between 1.0

and 3.0.⁴⁶ This assumption leads ultimately to a value of f greater than 0.4. This crude result can be compared to the Ar $L_1L_{2,3}M$ and L_1MM Auger intensities which show that the Coster-Kronig channel in Eq. (9) is the overwhelmingly favored decay pathway for an Ar 2s hole.⁴⁷ The fraction f for Ar has been determined from experiment and theory to be greater than 0.95.⁴⁷

Considering Eq. (11) for the resonant LVV/LLV ratio, we use the assertion from Sec. B that the "highly excited" satellite channels dominate the decay of the $S\ 1s \rightarrow 6t_{1u}$ resonance to simplify the expression for the LVV/LLV Auger ratio:

$$\begin{aligned} \left[\frac{LVV}{LLV} \right]_{on} &= \frac{2}{f} \left[\frac{\sigma(2p^{-2}v^*) + \sigma(2s^{-2}v^*) + \sigma(2p^{-1}2s^{-1}v^*)}{\sigma(2p^{-1}2s^{-1}v^*) + 2\sigma(2s^{-2}v^*)} \right] \\ &= \frac{2}{f} \left[\frac{\sigma_{Total}}{\sigma(2p^{-1}2s^{-1}v^*) + 2\sigma(2s^{-2}v^*)} \right]. \end{aligned} \quad (13)$$

It is clear upon inspection of Eq. (13) that in order for $[LVV/LLV]_{on}$ to be as large as the observed ratio on resonance [25(5)], the cross sections $\sigma(2s^{-2}v^*)$ and $\sigma(2p^{-1}2s^{-1}v^*)$ must be relatively small. In fact, the expression in brackets on the right-hand side of Eq. (13) must lie in the range 4-15 (using $0.4 \leq f \leq 1.0$). Even with this large uncertainty, these values indicate that at least 75% of the $S\ 1s \rightarrow 6t_{1u}$ resonant cross section appears in the $2p^{-2}v^*$ channel.

The dominance of the resonant $SF_6^+(2p^{-2}v^*)$ channel is not surprising if one examines the expected partitioning of the cross

sections within an Auger-like decay scheme. In other words, if the $6t_{1u}$ electron is left always as a "spectator", we can consider the core decay to be much like normal K-hole Auger decay. The KLL-type decay channels ("highly excited" satellites) should then dominate over the $KLV(V^*)$ decay pathways (main lines and low-excitation-energy satellites), which is observed qualitatively in our analysis of peak A on resonance. Furthermore, the partitioning into the three important KLL-type channels can be compared with three schemes: 1) statistical, 2) Ar KLL Auger decay,^{43,48} and 3) SF_6 S(KLL) Auger decay.⁴⁹ The partitioning in these scenarios is shown in Table III. It is clear that an Auger-like decay mechanism (similar to Ar and SF_6 KLL Auger rates) with dominant decay to the $2p^{-2}6t_{1u}$ state is consistent with our interpretation of the LVV/LLV ratio on resonance, within the assumptions made.

In summary, the $1s^{-1}6t_{1u}$ resonant state decays predominantly to the "highly excited" satellite configurations of SF_6^+ ($2p^{-2}v^*$, $2p^{-1}2s^{-1}v^*$, and $2s^{-2}v^*$). The decay of the excited state is such that the $6t_{1u}$ electron (v^*) probably remains as a "spectator", and the S 1s hole decays in a manner similar to Ar and SF_6 KLL Auger decay. Consistent with this, we find that at least 75% of the resonant cross section appears in the SF_6^+ ($2p^{-2}v^*$) channel.

D. Resonant Asymmetry Parameters

The asymmetry parameters for the S(LVV) peak and peak A provide

additional information on the photoemission and subsequent Auger-decay channels at the $S\ 1s \rightarrow 6t_{1u}$ resonance. Figure 7 shows β values for the $S(LVV)$ Auger peak and peak A, both demonstrating marked changes on resonance. We note that the behavior of the $\beta(\text{peak A})$ mostly is due to the photon-energy-dependent changes in the cross sections of the unresolved components of peak A. Below the $S\ 1s \rightarrow 6t_{1u}$ resonance, about 85% of the peak A intensity is $F\ 1s$ photoemission, yielding β near 2.0. On resonance, the $F\ 1s$ cross section remains unaltered while the other components of peak A exhibit a large increase in cross section, as seen in Fig. 4. The dominance of the $S\ 2p$ and $S\ 2s$ "highly excited" spectator-satellite cross sections causes the $\beta(\text{peak A})$ to mimic the β value for these resonant levels, which appears to be near zero or slightly negative on resonance. The presence of unresolved contributions (spectator satellites) in peak A prohibits the measurement of the individual-channel asymmetry parameters.

Unlike the oscillations in $\beta(\text{peak A})$, the asymmetric profile for the $\beta(LVV)$ at the $S\ 1s \rightarrow 6t_{1u}$ resonance cannot be dismissed as arising from variations in peak-component cross sections. Although the oscillation is small (≈ 0.12) and more data on the wings of the resonance are needed to confirm the exact shape, the $\beta(LVV)$ clearly is affected. In addition, the maximum in $\sigma(\text{peak A})$ in Fig. 5, $\sigma(LVV)$ and $\sigma(LLV)$ in Fig. 6, and the minimum in $\beta(\text{peak A})$ in Fig. 7 all lie at 2486.0(5) eV, the energy at which the $\beta(LVV)$ crosses its background value, further confirming the asymmetric shape. This result for an

Auger angular distribution is particularly intriguing because no previous studies have reported angle-resolved Auger decay emanating from ions resonantly produced by autoionization below a deep core-level threshold. Even though this unusual resonant profile for an Auger β should be considered tentative, some implications of an asymmetric shape are worthy of discussion.

An asymmetric profile in β ordinarily signals an interference effect. However, the interference in the autoionization process occurs one step previous to the observed Auger decay. That is, the direct-ionization and excited-state autoionization pathways leading to SF_6^+ interfere, causing oscillations in the σ and β values for the single-ion channels. Subsequent Auger decay cannot experience this interference phenomenon directly, but may retain "memory" of the process due to ion alignment.

Following up the idea of ion alignment, the Auger electron β is, in a sense, a "snapshot" of the molecular orientation of the SF_6^+ ion prior to Auger decay. According to Dehmer and Dill's formalism, β_{Auger} can be expressed as:²⁴

$$\beta_{Auger} = c \beta_m \quad (14)$$

where c is a constant characteristic of each individual Auger-decay channel, and β_m is the asymmetry of the molecular ion orientation following photionization. Because autoionization to $2p^{-2}v^*$ dominates the $S\ 1s \rightarrow 6t_{1u}$ resonance, the oscillation in $\beta(LVV)$

should reflect primarily the orientation of this particular SF_6^+ ion configuration in the resonance region. A varying energy dependence of single-ion alignment due to autoionization has been observed previously in atomic Cd for the double excitations above the 4d threshold $[4d^9(5s5p^3p)6s^1p]^{50}$ and over the $4d \rightarrow np$ ($n \geq 8$) and nf ($n \geq 5$) Rydberg series.⁵¹

A remaining point to consider is the relationship between the molecular-ion asymmetry, β_m , and the resonant photoelectron asymmetry parameter for the major channel $2p^{-2}v^*$. The asymmetry in the alignment of the $2p^{-2}v^*$ ion suggested by the S(LVV) β results also should appear in the β of the $2p^{-2}v^*$ photoelectron, which is convoluted in the total β (peak A). However, the individual profile cannot be determined from the β (peak A) because of the unresolved components and the rapidly changing cross sections.

To summarize, the asymmetric profile for the S(LVV) Auger β over the $S\ 1s \rightarrow 6t_{1u}$ resonance, though tentative, is highly unusual. Though the conceptual link between the Auger electron β and single-ion alignment is provided by Dehmer and Dill's theory,²⁴ there are no available calculations for the specific shape of the molecular orientation asymmetry, β_m , over a discrete resonance.

V. Above the S 1s Threshold

For the data taken above the sulfur K edge, both σ (peak A) and σ (LVV) cross sections (Fig. 5) show characteristic undulations in the

S 1s continuum. In particular, the peak near 2507 eV is evident in our data (the unresolved d and e features in Fig. 1), as are broader features at ~2525 and 2555 eV (f and g in Fig. 1). Oscillations also appear in $\sigma(S\ 1s)$ and $\beta(S\ 1s)$ shown in Fig. 8. For the sake of illustration, the $\sigma(S\ 1s)$ data for SF_6 have been normalized to the absorption curve.¹ The MSM-X α $\sigma(S\ 1s)$ and $\beta(S\ 1s)$ are also shown.¹¹

The S 1s continuum results will be discussed in two parts. In Sec. A we examine our data near the 2507 eV resonances and interpret sharp changes in the cross-section ratios (peak A)/1s and LVV/1s as indications of resonant enhancement in the S 2p, S 2s, and valence main-line and satellite channels. This behavior and the observation of S 1s satellites lead to an assignment of the two resonances visible in the photoabsorption cross section at 2506 and 2511 eV (d and e) as doubly excited states.

Section B will address the nature of the features at 2525 and 2555 eV photon energy. We compare the experimental data with a single-scattering plane-wave EXAFS calculation and with MSM-X α results. This is followed by a discussion in terms of approximations to the electron-scattering process and the description of the molecular potential in EXAFS and MSM-X α calculations which attempt to model the effects in this energy region.

A. The 2507 eV Resonances and S 1s Satellites

The $\sigma(S\ 1s)$ data in Fig. 8, in contrast to $\sigma(LVV)$ and $\sigma(\text{peak A})$

cross sections in Fig. 5, do not show a sharp feature at 2507 eV. To illustrate the differing behavior at this resonance, the cross-section ratios (peak A)/(S 1s) and (LVV)/(S 1s) are plotted in Fig. 9. In the ratio plots, the behavior at 2507 eV is much more pronounced. This peak appears as one resonance in Fig. 9 and is 4-5 eV full width at half maximum. Both the 2506 and 2511 eV resonances discernable in the photoabsorption measurement¹ (see Fig. 1, d and e) probably are present in our lower-resolution results. Because of the contributions included in the measured peaks (see Table I), we stress that any sharp changes in the ratios below the S 1s satellite thresholds are due to resonant contributions of the S 2p, S 2s, and valence main lines and satellites (low-excitation and "highly excited" satellites). Furthermore, the peak at 2506 eV and its shoulder at 2511 eV in the photoabsorption curve do not appear in the MSM-X α theory curve, indicating that they probably are caused by a multi-electron resonance or a symmetry-forbidden process.

The β (S 1s) in Fig. 8 also shows a dramatic effect in this low-energy region. The β falls from ~ 2 at high photon energy to a minimum at 2507 eV of 1.1. The MSM calculation shows a decrease in β_{1s} in this region, even though it does not predict the photoabsorption features at 2506 and 2511 eV. Thus, it is not clear from comparison with theory whether the β (S 1s) effect is associated with the 2507 eV feature. The departure of β (S 1s) from the atomic value of 2.0 for an s orbital is a direct indication of the anisotropy of the molecular potential.

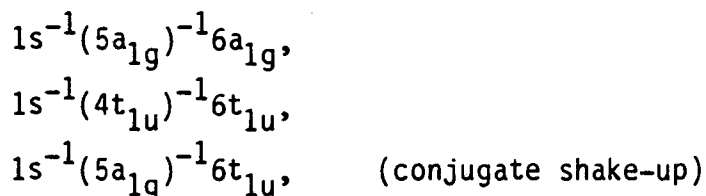
Pertinent to this resonance region, two S 1s correlation satellite peaks were observed directly at higher photon energies. Figure 3 shows the satellites at a representative photon energy of 2581 eV. We report an average branching ratio for the total satellite intensity relative to the S 1s main line of 15(3)% in the photon-energy range between 2570 and 2590 eV. The satellite thresholds were measured from a series of spectra to be 2510(1) and 2514(1) eV, 20 and 24 eV above the S 1s threshold, respectively. Note that the satellite binding energies are located 3-4 eV above the 2506 and 2511 eV photoabsorption features (d and e).

The observed satellite binding energies, combined with the sharp changes in the cross-section ratios discussed above, present a strong case for assignment of the resonant features as doubly excited autoionizing states preceding the satellite thresholds. The general assignment of features d and e as multiple excitations has been suggested previously.³ The possible decay channels of the postulated doubly excited resonances are depicted in Fig. 2 (dotted lines). The two resonances somewhat resolved in the absorption measurement may be leading to the two observed satellite thresholds. In analogy to the S 1s \rightarrow 6t_{1u} resonance, the configurations for the neutral doubly excited states could be $1s^{-1}(\text{val})^{-1}\text{v}^*6\text{t}_{1\text{u}}$, leading to the satellite ionic states $1s^{-1}(\text{val})^{-1}\text{v}^*$ of SF₆⁺. The 6t_{1u} electron is included in the excited-neutral configuration because the energy spacing of the resonance below the satellite thresholds (3-4 eV) is similar to the spacing of the

S 1s \rightarrow 6t_{1u} transition below the S 1s edge. However, multiplet splitting of the resonant states can be significant and will complicate the details of this simple assignment.

The intensity of these doubly excited resonances seems reasonable by comparison to the S 1s \rightarrow 6t_{1u} resonant intensity. The increase in the total cross section at one of the doubly-excited resonances is about 3(1)% of the increase at the S 1s \rightarrow 6t_{1u} resonance, while the total photoemission intensity of the S 1s satellites relative to the S 1s main-line photoemission is approximately 15%.

There have been no calculations to help identify the possible S 1s satellite configurations. Therefore, using the current valence ordering of Dehmer et al.¹⁸ and the 4t_{1u} \rightarrow 6a_{1g} transition energy of ~17 eV in the neutral molecule,¹⁸ we crudely approximated the possible shake-up and conjugate shake-up satellites in the energy region of interest.⁵² We have considered only promotion of an outer-valence electron to the 6a_{1g} and 6t_{1u} unoccupied orbitals. Based on energetics alone, this approach indicates that the observed S 1s satellites have the possible SF₆⁺ configurations:



though the conjugate shake-up state seems less likely. These configurations correspond to excitation from the deepest outer-valence

orbitals.

The general identification of these resonances as doubly excited states is reasonably clear, but in disagreement with the earlier assignment of one of them (2506 eV, feature d) as a symmetry-forbidden shape resonance.² Detailed theoretical calculations are needed to determine the energy positions of both the autoionizing states and satellite thresholds, and the intensity effects at the resonances.

B. High-energy Features

For the two higher-energy maxima at about 2525 and 2555 eV in $\sigma(S\ 1s)$, there is good agreement with the MSM calculation (Fig. 8). The S 1s kinetic energies at these maxima are about 35 and 65 eV, respectively. The magnitude of the increase in the experimental S 1s cross section near 2555 eV is quite large, about a factor of two over an energy range of 15 eV. The cross-section effect is probably accentuated by the suppression of the continuum intensity related to the huge enhancement of the S 1s $\rightarrow 6t_{1u}$ discrete resonance.

The significant scatter and uncertainty in the $\beta(S\ 1s)$ measurements above 2530 eV preclude any strong statements about an effect in the angular distribution associated with the rise in the cross section around 2545 eV. The MSM curve shows a minimum in $\beta(S\ 1s)$ which coincides with the S 1s cross-section minimum.

Because of the agreement between the MSM and experimental cross sections, we further consider the general MSM results for SF₆. The

nature of these high kinetic-energy oscillations was mentioned briefly by Wallace as originating from EXAFS behavior.¹¹ The MSM theory to some extent includes all single- and multiple-scattering events in the calculation of partial cross sections, so the particular physical effect(s) producing the oscillations is not clear when only cross section results are available. Wallace noted that there are no calculated symmetry-allowed shape-resonant states in this energy region; the presence of such quasi-bound states would be unusual considering the required barrier height ($>35, 65$ eV) needed for trapping the photoelectron. However, the eigenphase sum for $S\ 1s(1a_{1g}) \rightarrow \epsilon t_{1u}$ photoionization rises by $\sim\pi/3$ over an energy range of 15 eV centered at ~ 2549 eV photon energy,⁵³ which is the center of the rise in the $S\ 1s$ cross section. There is also a marked similarity among the calculated MSM cross sections for the various core levels of SF_6 in this kinetic-energy range. We will not elaborate on this similarity except to note that the intensity changes around 60 eV kinetic energy may be caused by a single phenomenon.

Because Wallace¹¹ interpreted the high-energy features as EXAFS, we have performed a single-scattering plane-wave EXAFS calculation for comparison with experiment to help determine the physical origin of these features by identifying the EXAFS portion of the continuum oscillations. The factors of short bond distance ($r = 1.58$ Å) and strong back-scattering amplitude which make single-scattering EXAFS pronounced are indeed present in SF_6 . The calculated EXAFS oscillatory amplitudes using a Debye-Waller factor of

$\sigma=0$ (best case for large amplitude) and two different central-atom phase shifts (sulfur and "adjusted" phase shifts) are shown in Fig. 10, plotted with the absorption curve (where the below-threshold nonresonant intensity has been subtracted) and the MSM- $X\alpha$ S 1s partial cross section. For the F backscattering atoms, the published Clementi-Roetti phase shifts and amplitudes were used.^{54,55} Various central-atom phase shifts other than those for sulfur were tried until the calculated high kinetic-energy oscillations coincided with the experimental energies of approximately 140, 215, and 295 eV (see Fig. 10, vertical lines). The resulting "adjusted" central-atom phase shift corresponds to a curve between that for Na and Mg (the Clementi-Roetti $l=1$ phase shifts with the $Z+1$ approximation); quantitatively, this equals the Si phase shift minus a value of 1.25 radians.⁵⁴ Though the variation of the central-atom phase shift did serve to line up the high kinetic-energy EXAFS features, it also reduced the effect in the cross section by a factor of two. For this reason, we also show in Fig. 10 the calculated EXAFS curve using the unadjusted sulfur phase shifts (dashed curve, bottom).

A comparison between the calculated EXAFS curve (with "adjusted" phase shifts) and experiment shows good agreement above ~ 120 eV kinetic energy in the amplitude of the EXAFS oscillations when an estimated background cross section is used (see top of Fig. 10, dotted line). The features below 100 eV, however, are not as well reproduced. It is difficult to assess the relative intensity effect for SF_6 in this region because of the uncertainty in the

nondiffractive experimental "background", though we can set a lower limit on the effect in the S 1s cross section at about 50%. The EXAFS calculation with the "adjusted" phase shift shows a 25% effect in this energy region, while the EXAFS curve using the sulfur-atom phase shift shows a 50% effect. We believe that the uncertainties in both the experimental and calculated amplitudes below 100 eV kinetic energy do not permit any conclusions based on intensity arguments.

Using the "adjusted" phase shifts to reproduce the energies of the high-kinetic-energy wiggles, the maximum of the large peak at 65 eV kinetic energy is off by ~20 eV in energy. This particularly poor energy agreement is reinforced when similar calculations on Br₂ and GeCl₄ are examined. The calculations for these molecules reproduce the corresponding experimental energies at both high energies and between 4 and 5 Å⁻¹ (60-100 eV).⁵⁵ It is known that atoms less electronegative than fluorine are not as effective in creating a barrier in the molecular potential which can modify the atomic effects.² These facts point to the possible importance of molecular effects in SF₆. Considering all of these factors, we conclude that the cross-section features at 35 and 65 eV kinetic energy in SF₆ (f and g in Fig. 1) do not arise exclusively from a simple single-scattering plane-wave EXAFS phenomenon.

Summarizing the results to this point, our conclusions are that the MSM-X_α calculation¹¹ reproduces the experimental S 1s cross section well below 100 eV kinetic energy (Fig. 10, top, dot-dashed curve), whereas our single-scattering plane-wave EXAFS calculation

does not. The next question is how to improve the general theoretical treatment of this problem. Two types of "fine-tuning" of the theory involve first, a better description of the electron-scattering process, and second, improvement in the treatment of the molecular potential. These two improvements are not necessarily separate developments.

EXAFS theorists recently have been interested in improving the description of the scattering process by introducing spherical waves into the single-scattering calculations.²⁵⁻²⁸ One can make an intuitive argument that the curvature of the wave front will be more important at low kinetic energies and for molecules with short bond distances.²⁸ From a more quantitative point of view, Lee and Pendry have done plane- and spherical-wave calculations on crystalline Cu.²⁵ The spherical-wave corrections to the Cu EXAFS curve shift the oscillations by an energy on the order of 20 eV and reduce the amplitude by a factor of two below 100 eV. Based on these crude estimates, we think that curved-wavefront corrections (which we have neglected in our calculation) should be important for SF₆ in the kinetic-energy range below 100 eV, affecting both the amplitude and phase of the EXAFS features.

The complication of multiple-scattering events should also be considered in this energy range (MSM theory of course includes these events). Multiple-scattering contributions to EXAFS have been calculated on a variety of systems with the following trends having been exhibited. First, the most important effect occurs with an

arrangement of collinear atoms where the intervening atom serves to "focus" amplitude back onto the central atom.³⁰⁻³² This focussing effect usually involves only one backscattering event and can result in a significantly enhanced amplitude and a phase shift at all energies.³⁰ A similar multiple-scattering path for SF₆ can be denoted S-F₁-S-F₂-S, where F₁ and F₂ are fluorine atoms collinear with the sulfur atom. Secondly, large-angle scattering can be significant, especially at low kinetic energy where electron scattering becomes more isotropic. Bunker and Stern³² estimated that below 30 eV kinetic energy for Mn 1s ionization of KMnO₄, the large-angle multiple-scattering amplitude relative to single scattering is about 25-50% (and about 10-20% at higher energy).

However, even with spherical-wave and multiple-scattering corrections included which improve the treatment of the electron-scattering process, there may still be significant interaction with the more diffuse molecular potential below 100 eV kinetic energy. A recent experimental study on the oxygen K-edge EXAFS spectra of O₂, CO, and CO₂ concludes that the observed disagreement between these experiments and single-scattering EXAFS theory is probably caused by inappropriate atomic parameters in the EXAFS calculation.⁵⁶ The atomic parameters and potential were thought to be inaccurate for modeling the electron scattering in the molecular orbitals.⁵⁶ The current degree of modification of EXAFS theory fails to include the complicated electronic structure and interaction due to the molecular nature of the problem. For example,

in the atomic phase-shift EXAFS calculations of Lee and Beni,⁵⁷ two atomic muffin-tin potentials are calculated separately and overlapped. Of course, the details of the potential between the atoms are not muffin-tin-like, as the authors note. The inaccuracies in this region should not affect the EXAFS calculations at high energy because any complexity will not be experienced by a high-energy electron. It is exactly in this region, however, where low-energy electrons may interact more with the details of the molecular potential.

MSM- $X\alpha$ calculations use as a starting point atomic potentials similar to those used in EXAFS theory.^{11,13} The molecular detail between the atoms is picked up, albeit indirectly, by a self-consistent treatment of the molecular potential. The possible nature of the interaction with the molecular field above approximately 30 eV kinetic energy has not been considered previously. The same interaction at lower kinetic energy can result in shape resonances, which are reasonably well understood. However, even in this respect SF_6 seems to be an especially complicated case.^{17,18,58}

In order to qualitatively imagine how a high-energy (30-60 eV kinetic energy) interaction might occur, it is useful to think of the simple square-barrier potential problem where:⁵⁹

$$\begin{aligned} V &= V_0 & 0 < r < r_0, \\ V &= 0 & r < 0 \text{ and } r > r_0. \end{aligned} \tag{15}$$

When the energy of a wave is greater than the barrier height V_0 , the wave is quantum-mechanically transmitted and reflected, giving oscillations in the continuum cross section, frequently called transmission resonances. The largest effect occurs with a large barrier width and/or height. Certainly, the realistic addition of a potential well and a repulsive wall may perturb even the qualitative aspects of this effect, but we use this simple example to illustrate the possible nature of an interaction between the photoelectron and a possibly large barrier induced by the electronegative fluorines in SF_6 .

We believe that the future understanding of this phenomenon will come primarily from detailed theoretical work which examines the origins of the dipole matrix-element changes in this energy range. The results of a step-by-step EXAFS calculation for SF_6 , where spherical-wave and multiple-scattering corrections are added sequentially, would certainly help to estimate the importance of interaction with the atomic cores (EXAFS) relative to that with the more diffuse aspects of the molecular potential. Improvements in the potential itself are necessary also to pick up the possible nondiffractive (barrier) interactions that occur in this intermediate-energy range. Combined with this, a careful examination of the results of the MSM- $X\alpha$ calculation, which quantitatively reproduces the cross section between 25 and 100 eV kinetic energy, could yield some insight into the physical origin(s) of these high-energy features.

IV. Conclusions

To summarize, the specific conclusions which can be drawn from our data near the sulfur K edge in SF₆ are as follows:

1. For the S 1s ($1a_{1g}$) \rightarrow $6t_{1u}$ resonance, "highly excited" satellites with two core holes (S 2p, S 2s) are the important SF₆⁺ decay channels. These configurations ($2p^{-2}v^*$, $2s^{-2}v^*$, and $2p^{-1}2s^{-1}v^*$), if regarded as S 2s and S 2p satellites, have exceptionally high excitation energies (>150 eV) and, according to the spectator model, probably contain a $6t_{1u}$ electron in the excited v^* orbital. In general, our data suggest that the decay of the neutral excited state proceeds much like Ar KLL and SF₆ S(KLL) Auger decay, with the $2p^{-2}v^*$ channel as the dominant one. Furthermore, an asymmetric resonant profile for the S(LVV) Auger β is observed, probably caused by ion alignment in the previous photoemission step.
2. The resonances around 2507 eV are probably doubly excited autoionizing states (leading to observed satellite thresholds), because decay into S 2p, S 2s, and/or valence photoemission channels is observed. The interpretation emphasizes the general requirement for results on individual photoemission channels in order to distinguish satellite

continua and autoionization effects from shape resonances in the assignment of absorption features.

3. The data further above the S 1s threshold (30-100 eV kinetic energy) are difficult to interpret. The experimental results show a factor of two increase in the S 1s cross section near 2550 eV, but no conclusive effect in the S 1s β . MSM- $X\alpha$ calculations successfully reproduce the effects in the S 1s cross section, indicating their one-electron nature.¹¹ We conjecture that the large effects are caused by a combination of spherical-wave and multiple-scattering effects in EXAFS and an interaction of the photoelectron with the details of the molecular potential.

There is an obvious need for further experimental work in several areas. High kinetic-energy resolution would help to assign the SF_6^+ resonant states (i.e., the location of the excited electron) and possibly the structure of the broad sulfur LLV and LVV Auger peaks on and off resonance. The decay of core-level discrete states in atoms and molecules has been examined in just a few systems to date. We predict that future resonant work below deep core-level thresholds will confirm the predominance of "highly excited" satellites and the importance of spectator decay. The study of core levels of other octahedral molecules with electronegative ligands and of substituted hexafluorides (like SF_5X) with respect to the continuum effects

above 30 eV kinetic energy may help to determine the origin of the cross-section effects in this energy region.

On the theory side, resonance calculations are needed for individual cross sections and angular distributions. More general work on the theory for Auger-electron angular distributions is called for to elucidate how autoionization produces an aligned molecular ion prior to Auger decay. Finally, advances in EXAFS theory as applied to molecules, especially concerning the treatment of the electron-scattering process and of the molecular potential, are needed to investigate the nature of interactions with low-energy (<100 eV) electrons.

Acknowledgements

We would like to acknowledge helpful discussions with J.L. Dehmer, J.A. Stephens, J.J. Barton, and M.N. Piancastelli. This work was supported by the Director, Office of Energy Research, Office of Basic Energy Sciences, Chemical Sciences Division of the U.S. Department of Energy under Contract No. DE-AC03-76SF00098. It was performed at the Stanford Synchrotron Radiation Laboratory, which is supported by the Department of Energy's Office of Basic Energy Sciences.

References

1. R.E. LaVilla and R.D. Deslattes, J. Chem. Phys. 44, 4399 (1966).
2. J.L. Dehmer, J. Chem. Phys. 56, 4496 (1972) and references therein.
3. F.A. Gianturco, C. Guidotti, and U. Lamanna, J. Chem. Phys. 57, 840 (1972).
4. S.T. Manson and J.W. Cooper, Phys. Rev. 165, 126 (1968).
5. J.P. Connerade, Contemp. Phys. 19, 415 (1978).
6. K.T. Cheng and C. Froese-Fischer, Phys. Rev. A 28, 2811 (1983).
7. K.T. Cheng and W.R. Johnson, Phys. Rev. A 28, 2820 (1983).
8. D. Dill and J.L. Dehmer, J. Chem Phys. 61, 692 (1974).
9. J.L. Dehmer and D. Dill, Phys. Rev. Lett. 35, 213 (1975).
10. J.L. Dehmer and D. Dill, J. Chem. Phys. 65, 5327 (1977).
11. R.S. Wallace, Ph.D. thesis, Boston University, 1980.
12. D. Loomba, S. Wallace, D. Dill, and J.L. Dehmer, J. Chem. Phys. 75, 4546 (1981).
13. J.L. Dehmer, D. Dill, and A.C. Parr in Photophysics and Photochemistry in the Vacuum Ultraviolet, edited by S.P. McGlynn, G. Findley, and R. Huebner (D. Reidel Publishing Co., Dordrecht, Holland, 1985), p. 341.
14. J. Berkowitz, Photoabsorption, Photoionization, and Photoelectron Spectroscopy (Academic Press, New York, 1979).
15. A.P. Hitchcock and C.E. Brion, Chem. Phys. 37, 319 (1979).
16. R.C.C. Perera, J. Barth, R.E. LaVilla, R.D. Deslattes, and A. Henins, Phys. Rev. A 32, 1489 (1985).

17. T. Gustafsson, Phys. Rev. A 18, 1481 (1978).
18. J.L. Dehmer, A.C. Parr, S. Wallace, and D. Dill, Phys. Rev. A 26, 3283 (1982).
19. L. Ungier and T.D. Thomas, Chem. Phys. Lett., 96, 247 (1983).
20. H.W. Haak, G.A. Sawatzky, L. Ungier, J.K. Gimzewski, and T.D. Thomas, Rev. Sci. Instrum. 55, 696 (1984).
21. L. Ungier and T.D. Thomas, J. Chem. Phys. 82, 3146 (1985).
22. U. Becker, R. Hölzel, H.G. Kerkhoff, B. Langer, D. Szostak, and R. Wehlitz (unpublished results).
23. C.M. Truesdale, S.H. Southworth, P.H. Kobrin, U. Becker, D.W. Lindle, H.G. Kerkhoff, and D.A. Shirley, Phys. Rev. Lett. 50, 1265 (1983).
24. D. Dill, J.R. Swanson, S. Wallace, and J.L. Dehmer, Phys. Rev. Lett. 45, 1393 (1980).
25. P.A. Lee and J.B. Pendry, Phys. Rev. B 11, 2795 (1975).
26. J.E. Muller and W.L. Schaich, Phys. Rev. B 27, 6489 (1983).
27. W.L. Schaich, Phys. Rev. B 29, 6513 (1984).
28. J.J. Barton, Ph.D. thesis, University of California, Berkeley, 1985.
29. C.A. Ashley and S. Doniach, Phys. Rev. B 11, 1279 (1975).
30. B.K. Teo, J. Am. Chem. Soc. 103, 3990 (1981).
31. J.J. Boland, S.E. Crane, and J.D. Baldeschwieler, J. Chem. Phys. 77, 142 (1982).
32. G. Bunker and E.A. Stern, Phys. Rev. Lett. 52, 1990 (1984).
33. M.G. White, R.A. Rosenberg, G. Gabor, E.D. Poliakov, G.

- Thornton, S. Southworth, and D.A. Shirley, Rev. Sci. Instrum. 50, 1288 (1979).
34. S. Southworth, C.M. Truesdale, P.H. Kobrin, D.W. Lindle, W.D. Brewer, and D.A. Shirley, J. Chem. Phys. 76, 143 (1982).
35. S. Southworth, U. Becker, C.M. Truesdale, P.H. Kobrin, D.W. Lindle, S. Owaki, and D.A. Shirley, Phys. Rev. A 28, 261 (1983).
36. C.N. Yang, Phys. Rev. 74, 764 (1948).
37. We note that direct decay to double-ionization (shake-off) channels also is possible, resulting in SF_6^{2+} configurations such as $2p^{-2}$, val^{-2} , and $2p^{-1}val^{-1}$. The energy distribution of the emitted electrons would be very broad and somewhat peaked near 0 eV and the threshold energy for the shake-off process, with the high-energy part of the distribution unresolved from the other components in peak A in our TOF spectra. Thus, we cannot assess what percentage of excited neutrals decay directly to shake-off channels at the $S\ 1s \rightarrow 6t_{1u}$ resonance, although we speculate that this fraction probably is less than 20%.
38. It is important to recall that the peaks enhanced at the $S\ 1s \rightarrow 6t_{1u}$ resonance are not KLL Auger peaks, but rather autoionization to photoemission channels. The autoionizing decay of the neutral $SF_6(1s^{-1}6t_{1u})$ state can, however, be described in terms (using Auger notation) similar to those used to describe the Auger decay of the single-ion $SF_6^+(1s^{-1})$.
39. Some representative examples for relative intensities of atomic

and molecular satellites are given in U. Gelius, J. Electron Spectrosc. 5, 985 (1974).

40. For the Auger decay of the resonantly produced state, we assume that the $6t_{1u}$ electron is not involved in the decay because other t_{1u} orbitals have better overlap with the S $2p(2t_{1u})$ hole. If the $6t_{1u}$ electron is involved, the S(LVV) Auger kinetic energy will shift to even higher energy.
41. D.A. Shirley, Phys. Rev. A 9, 1549 (1974).
42. H. Korber and W. Melhorn, Z. Phys. 191, 217 (1966); M.O. Krause, T.A. Carlson, and W.E. Moddeman, J. Phys. (Paris) Colloq. C4, 139 (1971).
43. L. Asplund, P. Kelfve, B. Blomster, H. Siegbahn, and K. Siegbahn, Phys. Scr. 16, 268 (1977).
44. In the following analysis, we neglect the previously mentioned direct shake-off decay on resonance,³⁷ which would leave L-hole ions which subsequently Auger decay with intensity appearing in the observable S(LVV) and S(LLV) Auger peaks. As long as the relative abundance of two-core-hole configurations within the shake-off manifold ($2p^{-2}:2s^{-2}:2p^{-1}2s^{-1}$) is similar to that for the single-ion manifold reached by autoionization ($2p^{-2}v^*:2s^{-2}v^*:2p^{-1}2s^{-1}v^*$), the qualitative conclusions about the SF_6^+ decay channels are unaffected. Furthermore, double-Auger processes such as $SF_6^+(2p^{-1}) \rightarrow SF_6^{3+}(val^{-3}) + 2e^-$ are not included in the observed S(LVV) and S(LLV) Auger peaks. We estimate that off resonance these Auger shake-off processes

account for only ~10% of the total L-hole Auger decay, based on the measured double-Auger processes for L holes in Ar.⁶⁰ On resonance, double-Auger decay may occur somewhat more frequently (though single-Auger decay probably will still dominate) because "highly excited" ions like $2p^{-2}6t_{1u}$ undergoing Auger decay may have a greater probability for shaking off the $6t_{1u}$ electron relative to shake-off of a valence electron in the Auger decay of a $2p^{-1}$ ion.

45. G. Herzberg, Electronic Spectra and Electronic Structure of Polyatomic Molecules (D. Van Nostrand Co., Princeton, New Jersey, 1966).
46. K. Siegbahn, C. Nordling, G. Johansson, J. Hedman, P.F. Hedén, K. Hamrin, U. Gelius, T. Bergmark, L.O. Werme, R. Manne, and Y. Baer, ESCA Applied to Free Molecules (North-Holland, Amsterdam, London, 1969).
47. W. Mehlhorn, Z. Phys. 208, 1 (1968).
48. J. Vayrynen, R.N. Sodhi, and R.G. Cavell, J. Chem. Phys. 79, 5329 (1983); M.O. Krause, Phys. Rev. Lett. 34, 633 (1975).
49. L. Asplund, P. Kelfve, B. Blomster, H. Siegbahn, K. Siegbahn, R.L. Lozes, and U.I. Wahlgren, Phys. Scr. 16, 273 (1977).
50. W. Kronast, R. Huster, and W. Mehlhorn, J. Phys. B 17, L51 (1984).
51. Z.M. Goodman, C.D. Caldwell, and M.G. White, Phys. Rev. Lett. 54, 1156 (1985); 54, 2381 (1985).
52. The latest valence ordering, though supported by many-body calculations,⁶¹ is not completely certain due to the presence

- of a possible shape resonance affecting several valence cross sections at about 23 eV photon energy.^{17,18}
53. J.A. Stephens and P. Dittman (private communication of calculations by R.S. Wallace).
 54. B.K. Teo and P.A. Lee, J. Am. Chem. Soc. 101, 2815 (1979).
 55. B.K. Teo, P.A. Lee, A.L. Simons, P. Eisenberger, and B.M. Kincaid, J. Am. Chem. Soc. 99, 3854 (1977); P.A. Lee, B.K. Teo, and A.L. Simons, 99, 3856 (1977).
 56. B.X. Yang, J. Kirz, and T.K. Sham, Phys. Lett. 110A, 301 (1985).
 57. P.A. Lee and G. Beni, Phys. Rev. B 15, 2862 (1977).
 58. T.A. Ferrett, D.W.-Lindle, P.A. Heimann, M.N. Piancastelli, P.H. Kobrin, H.G. Kerkhoff, U.E. Becker, W.D. Brewer, and D.A. Shirley (unpublished results).
 59. G. Baym, Lectures on Quantum Mechanics (Benjamin Cummings, Reading, Massachusetts, 1969); A. Messiah, Quantum Mechanics Vol. 1 (North-Holland, Amsterdam, and John Wiley and Sons, Inc., New York, 1961).
 60. T.A. Carlson and M.O. Krause, Bull. Am. Phys. Soc. 10, 455 (1965); T.A. Carlson and M.O. Krause, Phys. Rev. Lett. 17, 1079 (1966).
 61. W. von Niessen, L.S. Cederbaum, G.H.F. Diercksen, and G. Hohlneicher, Chem. Phys. 11, 399 (1975); P.J. Hay, J. Am. Chem. Soc. 99, 1013 (1977); W. von Niessen, W.P. Kraemer, and G.H.F. Diercksen, Chem. Phys. Lett. 63, 65 (1979).

TABLE I: Peak Contributions for the SF₆ S 1s Photoelectron Spectra.

Photon-Energy ^a Range	Peak		
	A	S(LVV) Auger	S(L ₁ L _{2,3} V) Auger
<2490 eV	F 1s, S 2p, S 2s, valence (main lines and satellites)	L _{1,2,3} VV Auger decay of S 2p, S 2s holes (main lines and satellites)	Auger decay of S 2s holes (main line and satellites)
2490 - 2510 eV	all of above <u>plus</u> KLL, KLV, KVV Auger decay of S 1s hole (main line)	all of above <u>plus</u> L _{1,2,3} VV secondary Auger decay of S 1s hole (main line)	all of above <u>plus</u> secondary Auger decay of S 1s hole (main line)
>2510 eV	all of above <u>plus</u> KLL, KLV, KVV Auger decay of S 1s hole (satellites)	all of above <u>plus</u> L _{1,2,3} VV secondary Auger decay of S 1s hole (satellites)	all of above <u>plus</u> secondary Auger decay of S 1s hole (satellites)

^a The energy ranges above designate the regions: below the S 1s threshold (<2490 eV), between the S 1s main-line and first satellite threshold (2490-2510 eV), above the first S 1s satellite threshold (>2510 eV).

TABLE II: Complete Description of Autoionization of the SF₆ (1s⁻¹6t_{1u}) State to SF₆⁺ Photoemission Channels, and Subsequent S(LVV and LLV) Auger Cascades to Form SF₆ⁿ⁺ (n=2-5)

Autoionization to SF ₆ ⁺ via 1s ⁻¹ 6t _{1u}				Auger decay of SF ₆ ⁺ to SF ₆ ⁿ⁺ (n=2-5)		
SF ₆ ⁺ photoemission channel	configuration ^a	autoionization decay "type" ^b	Binding energy(eV)	Auger transitions to fill all S 2p _c and S 2s holes	Total number of Auger electrons ^d	
					LLV	LVV
main lines:						
2p	2p ⁻¹	KL _{2,3} V [*]	181.0 ^e	L _{2,3} VV	0	1
2s	2s ⁻¹	KL ₁ V [*]	244.7 ^f	1. L ₁ VV 2. L ₁ L _{2,3} V + L _{2,3} VV	0 1	1 1
satellites:						
2p	2p ⁻¹ val ⁻¹ 6t _{1u} 2p ⁻¹ val ⁻¹ v [*] 1u	KL _{2,3} V and KL _{2,3} V [*]	~190-220 ^g	L _{2,3} VV	0	1
2s	2s ⁻¹ val ⁻¹ 6t _{1u} 2s ⁻¹ val ⁻¹ v [*] 1u	KL ₁ V and KL ₁ V [*]	~255-285 ^g	1. L ₁ VV 2. L ₁ L _{2,3} V + L _{2,3} VV	0 1	1 1

TABLE II: (cont.)

Autoionization to SF ₆ ⁺ via 1s ⁻¹ 6t _{1u}				Auger decay of SF ₆ ⁺ to SF ₆ ⁿ⁺ (n=2-5)		
SF ₆ ⁺ photoemission channel	configuration ^a	autoionization decay "type" ^b	Binding energy(eV)	Auger transitions to fill all S 2p _c and S 2s holes	Total number of Auger electrons. ^d	
					LLV	LVV
"highly excited" satellites:						
2p	2p ⁻² 6t _{1u} 2p ⁻² v [*] 1u	KL _{2,3} L _{2,3}	~360 ^h	2 L _{2,3} VV	0	2
2s	2s ⁻² 6t _{1u} 2s ⁻² v [*] 1u	KL ₁ L ₁	~490 ^h	1. L ₁ VV + L ₁ VV 2. L ₁ VV + L ₁ L _{2,3} V + L _{2,3} VV 3. L ₁ L _{2,3} V + L ₁ L _{2,3} V + 2 L _{2,3} VV	0 1 2	2 2 2
"mixed"	2p ⁻¹ 2s ⁻¹ 6t _{1u} 2p ⁻¹ 2s ⁻¹ v [*] 1u	KL ₁ L _{2,3}	~430 ^h	1. L ₁ VV + L _{2,3} VV 2. L ₁ L _{2,3} V + 2 L _{2,3} VV	0 1	2 2

TABLE II: (cont.)

- a Inner- and outer-valence orbitals (Binding energies 16 to 44 eV) are denoted by "val". Excitation to an unoccupied molecular orbital other than $6t_{1u}$ (probably $6a_{1g}$) is denoted by "v*".
- b The Auger notation is used here to describe the decay type to the SF_6^+ photoemission final state. V designates that the $6t_{1u}$ electron has remained as a "spectator"; V* denotes that the excited $6t_{1u}$ electron has participated in the decay to SF_6^+ .
- c All 2s holes can decay via $S(L_1L_{2,3}V)$ or $S(L_1VV)$ Auger channels with some partitioning assumed independent of the configuration containing the 2s hole state.
- d Auger electrons are divided into sulfur LLV and LVV categories corresponding to the two observable Auger peaks in our photoemission spectra. The totals here include all primary through quaternary decay to give Auger electrons within the kinetic-energy region of the two observable peaks.
- e The S 2p binding energy is an average of the $2p_{3/2}$ and $2p_{1/2}$ spin-orbit binding energies which are 180.4 and 181.7 eV, respectively.⁴⁶
- f Ref. 46.
- g This energy range is based on observed S 2p satellite excitation energies.⁵⁸
- h Binding energy is estimated crudely by summing the binding energies (in the neutral) of the two core holes. Relaxation has not been included.

TABLE III: Relative Intensities for the "Highly Excited" Satellite Channels for the $S\ 1s \rightarrow 6t_{1u}$ Resonance in SF_6 Compared to KLL Auger Decay.

Scheme	Relative Intensity		
	$2p^{-2}$	$2s^{-1}2p^{-1}$	$2s^{-2}$
1. statistical decay:	0.54	0.43	0.04
2. Ar KLL-like decay: ^{43,48}	0.74(5)	0.22(5)	0.04(1)
3. SF_6 S(KLL)- like decay: ⁴⁹	0.72	0.19	0.09
4. observed decay:	> 0.75	-	-

Figure Captions

Fig. 1 Photoabsorption spectrum of SF₆ near the S 1s threshold from Ref. 1. The features discussed in the text are: the S 1s → 6t_{1u} resonance (b), the resonances near 2507 eV (d and e), and the higher-energy features (f and g). Features a and c are not discussed in this work, but have been interpreted as the symmetry-forbidden transitions S 1s(1a_{1g}) → 6a_{1g} (a) and εt_{2g} (c).²

Fig. 2 Sulfur 1s resonance energy-level diagram for SF₆. The 1s⁻¹6t_{1u} neutral excited state and its available photoemission decay channels to SF₆⁺ are shown with solid lines. Note the presence of "highly excited" S 2p and S 2s satellites produced by decay of the 1s⁻¹6t_{1u} state. Two postulated doubly excited states (see text for details) of the neutral are shown also, with some of the corresponding decay channels illustrated by dotted lines. Some excitation and decay pathways have been omitted for clarity. The designations "val" and "v*" represent outer-valence orbitals, and either 6a_{1g} or 6t_{1u} orbitals, respectively.

Fig. 3 TOF spectrum of SF₆ taken at θ=0° and 2581 eV photon energy above the S 1s threshold at 2490 eV. The components for peak A are listed in Table I.

Fig. 4 A TOF spectrum of SF₆ taken at θ=54.7° and 2484 eV near the S 1s(1a_{1g}) → 6t_{1u} resonance.

Fig. 5 Peak A (top) and S(LVV) Auger (bottom) relative cross sections and the photoabsorption measurement (Ref. 1, solid curve). The below-threshold nonresonant intensity has been subtracted from the data and the absorption curve. The data have been scaled to the absorption curve at 2497 eV for comparison. The cross-section scale (in Mb) thus refers strictly only to the photoabsorption data, and not to the photoemission and Auger cross sections.

Fig. 6 Relative cross sections for the $S(L_{1,2,3}VV)$ and $S(L_1L_{2,3}V)$ Auger electrons over the $S\ 1s \rightarrow 6t_{1u}$ resonance. The scale for the two lowest-energy points has been expanded times 100.

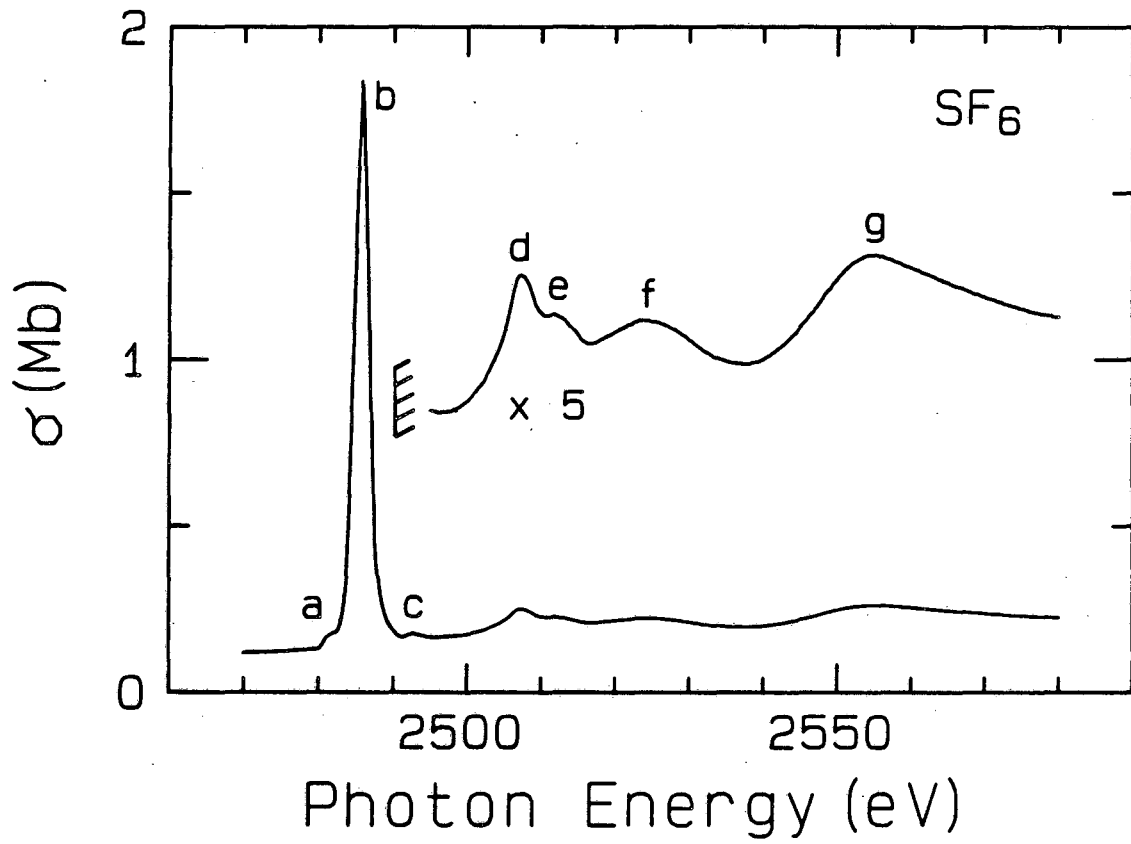
Fig. 7 Angular-distribution parameters for peak A (top) and S(LVV) Auger (bottom) over the $S\ 1s \rightarrow 6t_{1u}$ resonance. The oscillations in β (peak A) above the $S\ 1s$ threshold mainly are caused by added contributions from S(KLL, KLV, and KVV) Auger electrons.

Fig. 8 $S\ 1s$ partial cross section (top) and asymmetry parameter (bottom) plotted with MSM results (dashed curve).¹¹ The solid curve is the photoabsorption measurement¹ with the below-threshold nonresonant contributions subtracted. The cross-section data have been scaled to this "adjusted" absorption curve at 2535 eV.

Fig. 9 Cross-section ratios (peak A)/(S 1s) (top) and (LVV)/(S 1s) (bottom) in the vicinity of the 2507 eV resonances. For the

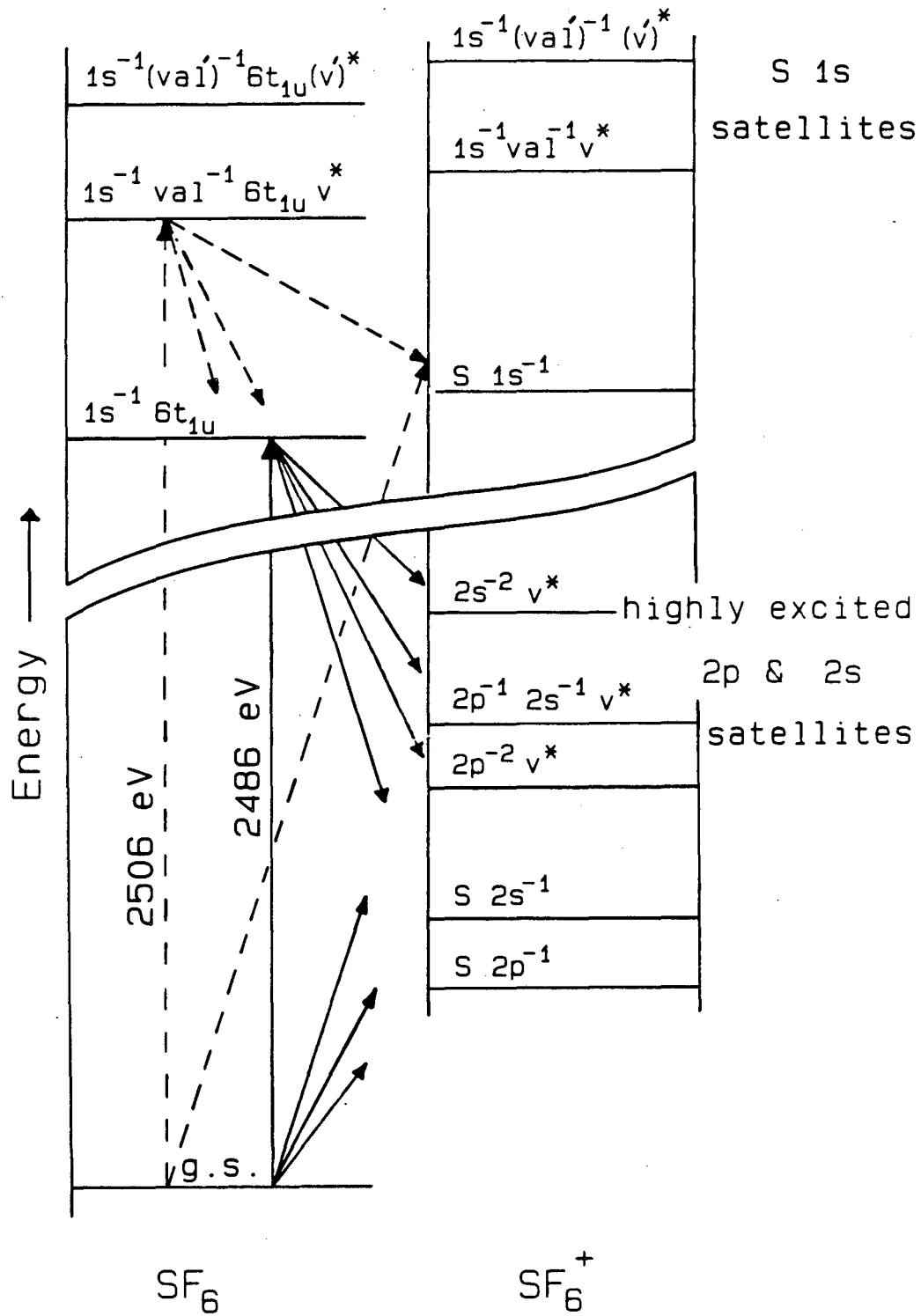
ratio (peak A)/(S 1s), the F 1s cross section has been subtracted from the peak A intensity using the F(KVV) Auger intensity in each spectrum. Observed S 1s satellite and main-line thresholds also are shown.

Fig. 10 S 1s single-scattering plane-wave EXAFS oscillatory amplitudes χ (%) for SF₆ as a function of kinetic energy (bottom). The solid and dashed curves represent the χ (%) calculated with the "adjusted" and sulfur central-atom phase shifts, respectively. The photoabsorption curve¹ is plotted on an absolute scale for comparison (solid curve, top), along with the MSM-X α S 1s partial cross section¹¹ (dot-dashed curve, top). The "non-EXAFS" background cross section has not been subtracted from the experimental absorption data. Above 100 eV kinetic energy, an estimation of the sloping background can be made (dashed curve, top) for comparison of the amplitude effects for that energy range. The vertical lines are drawn to emphasize the energy agreement between the "adjusted" phase shift EXAFS calculation and the experimental absorption data.



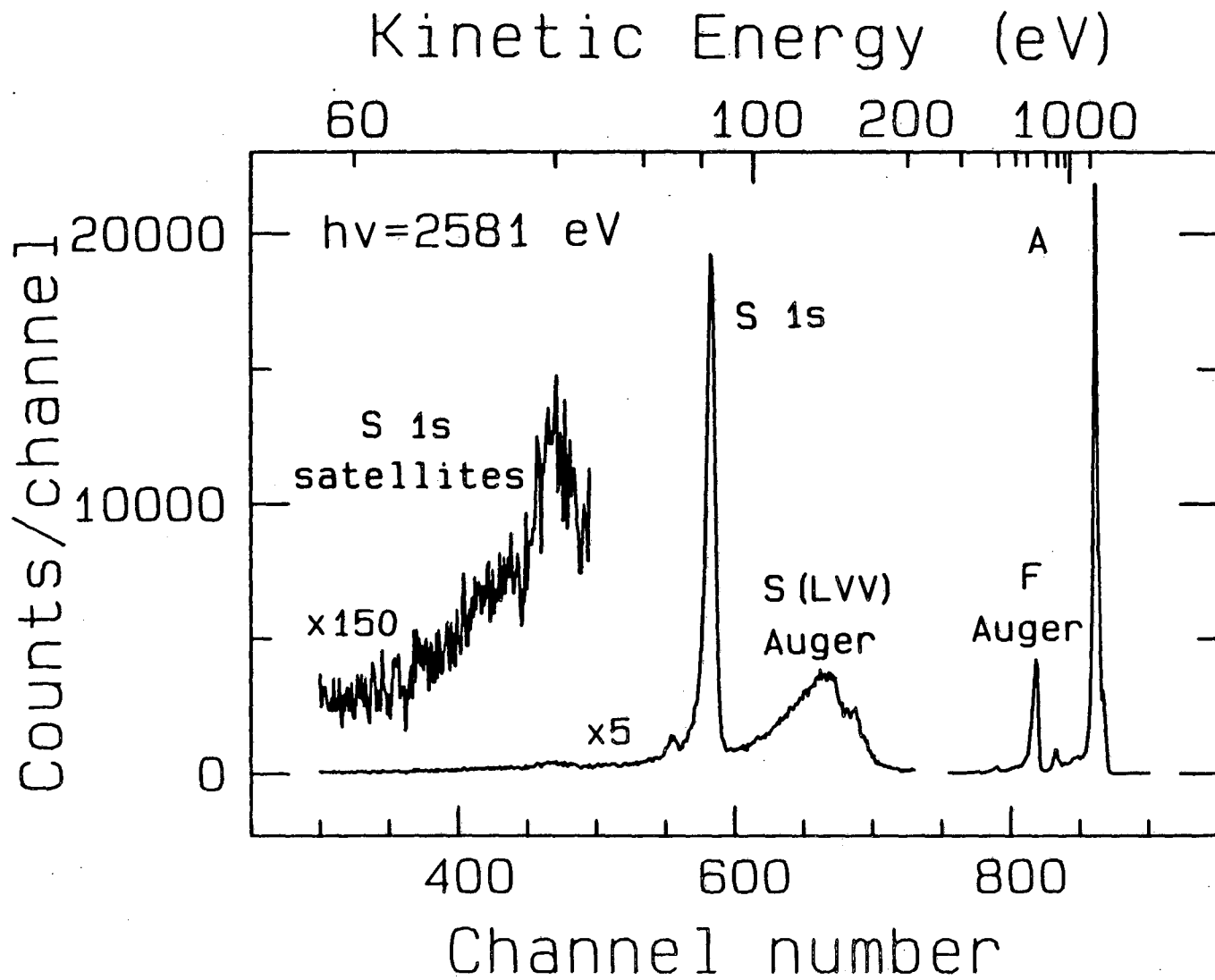
XBL 862-360

Figure 1



XBL 862-365

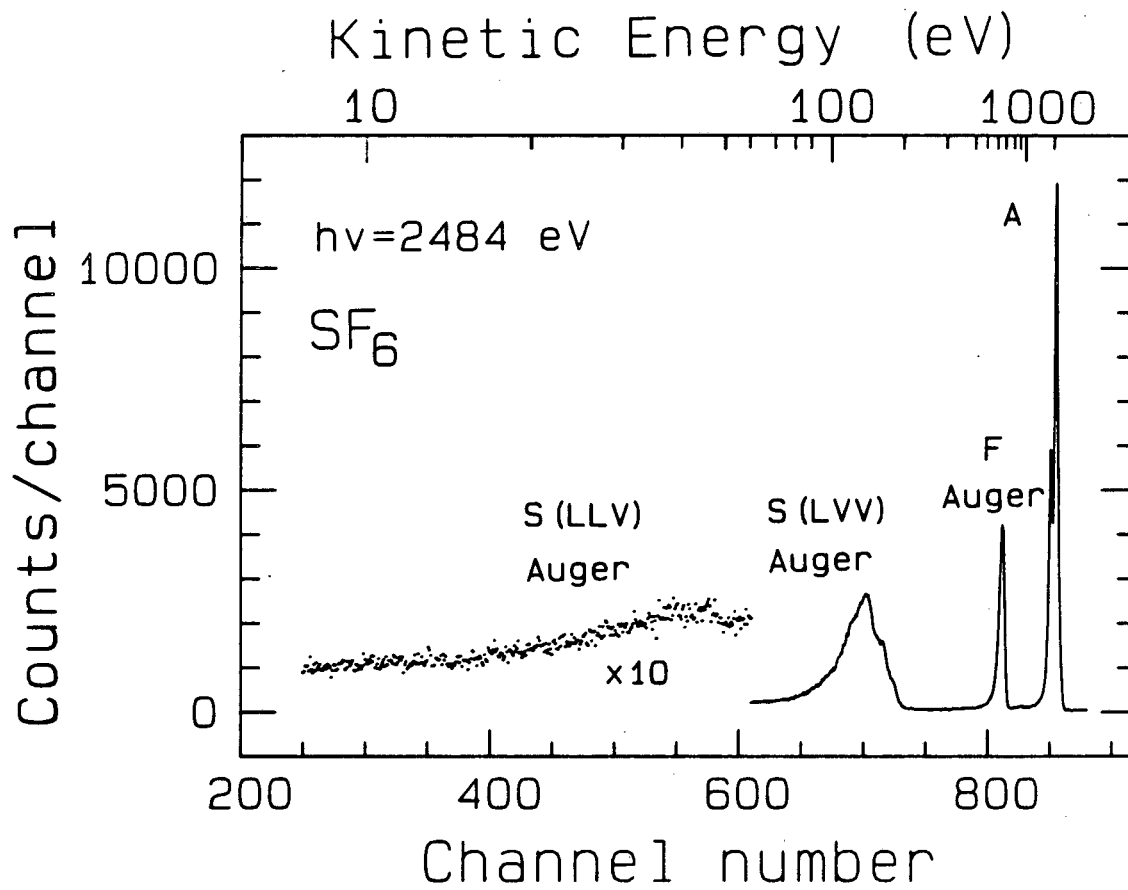
Figure 2



60

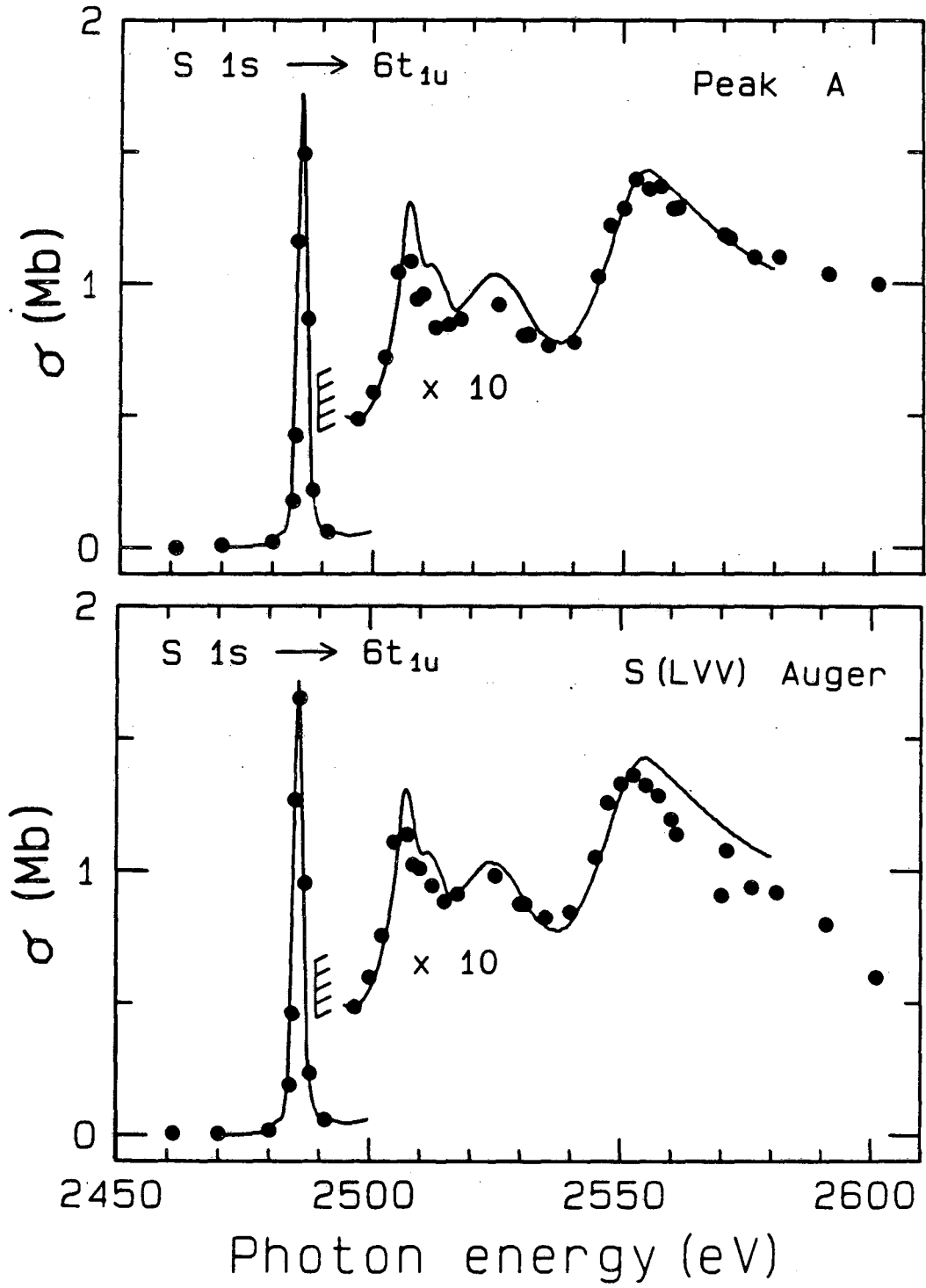
XBL 862-361

Figure 3



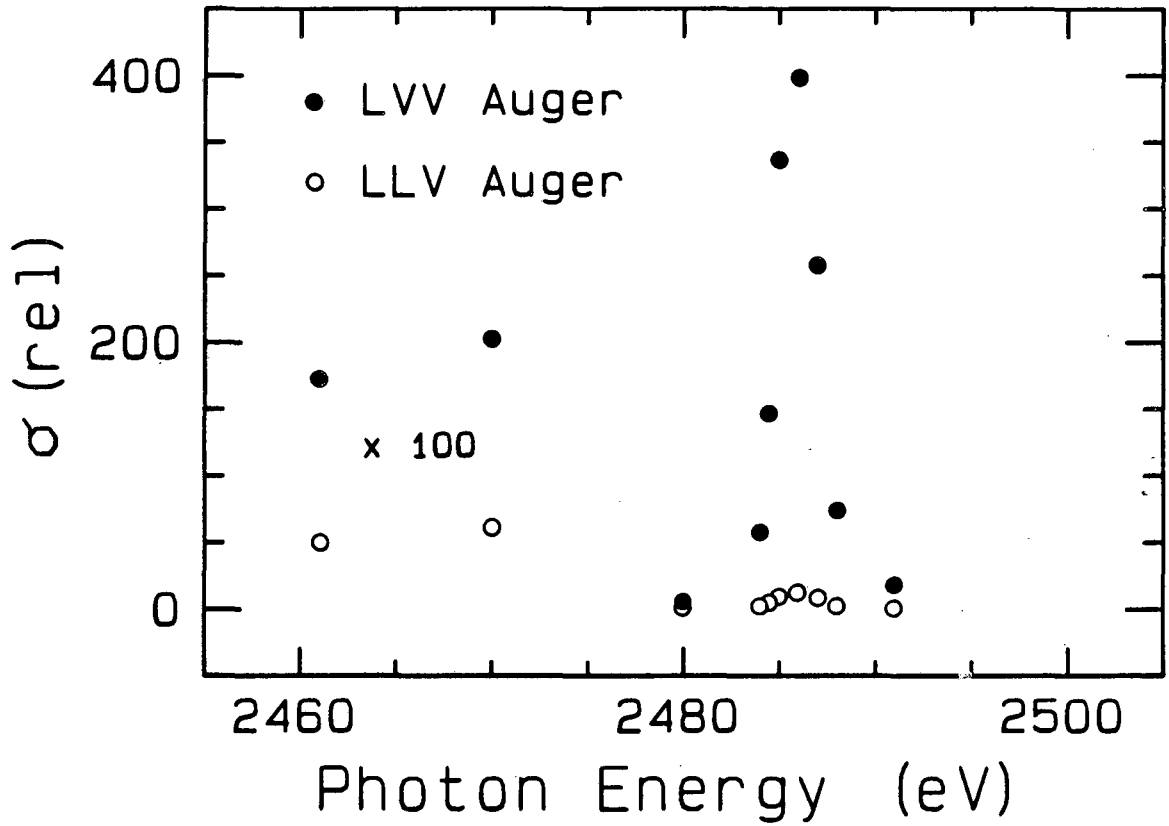
XBL 862-369

Figure 4



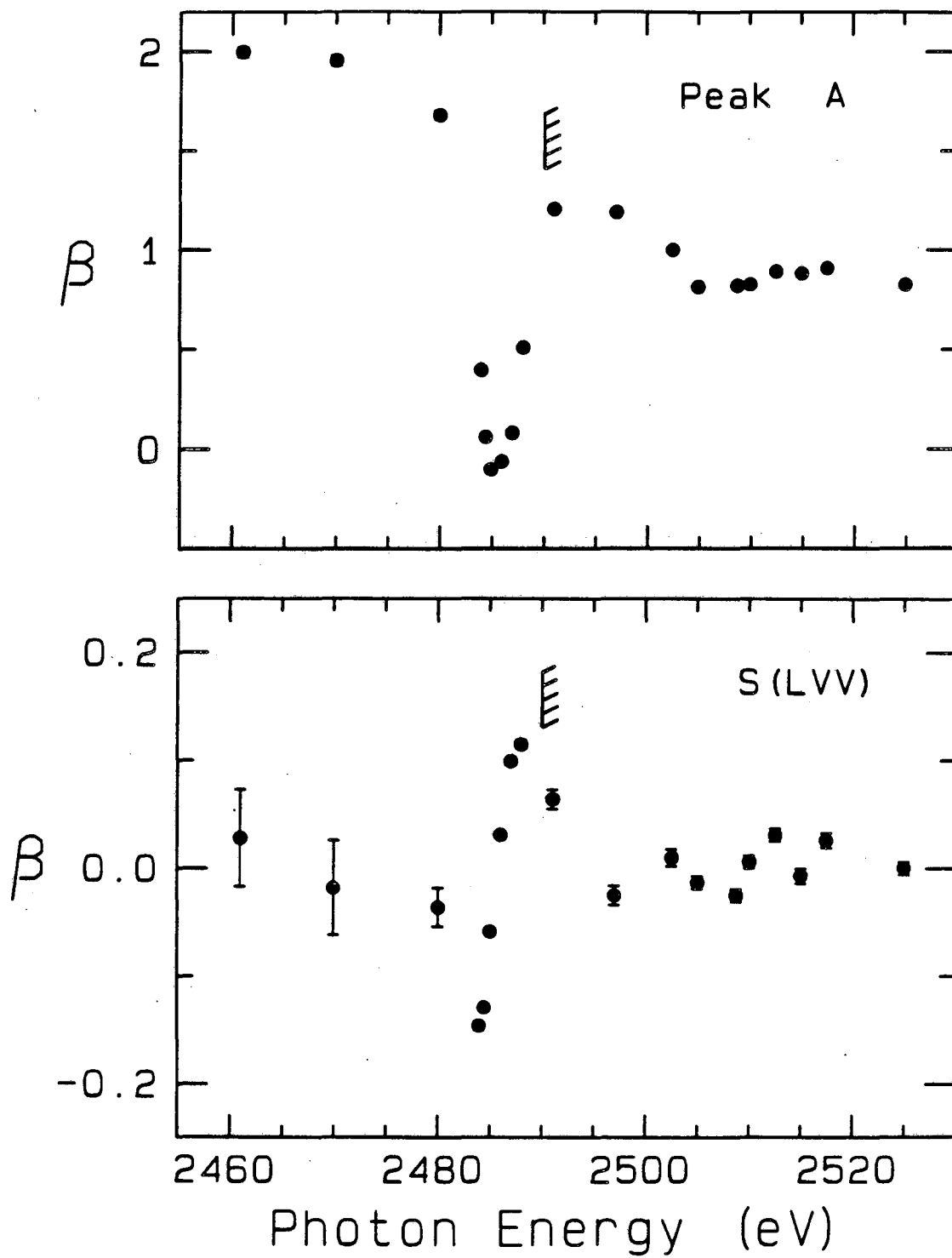
XBL 862-362

Figure 5



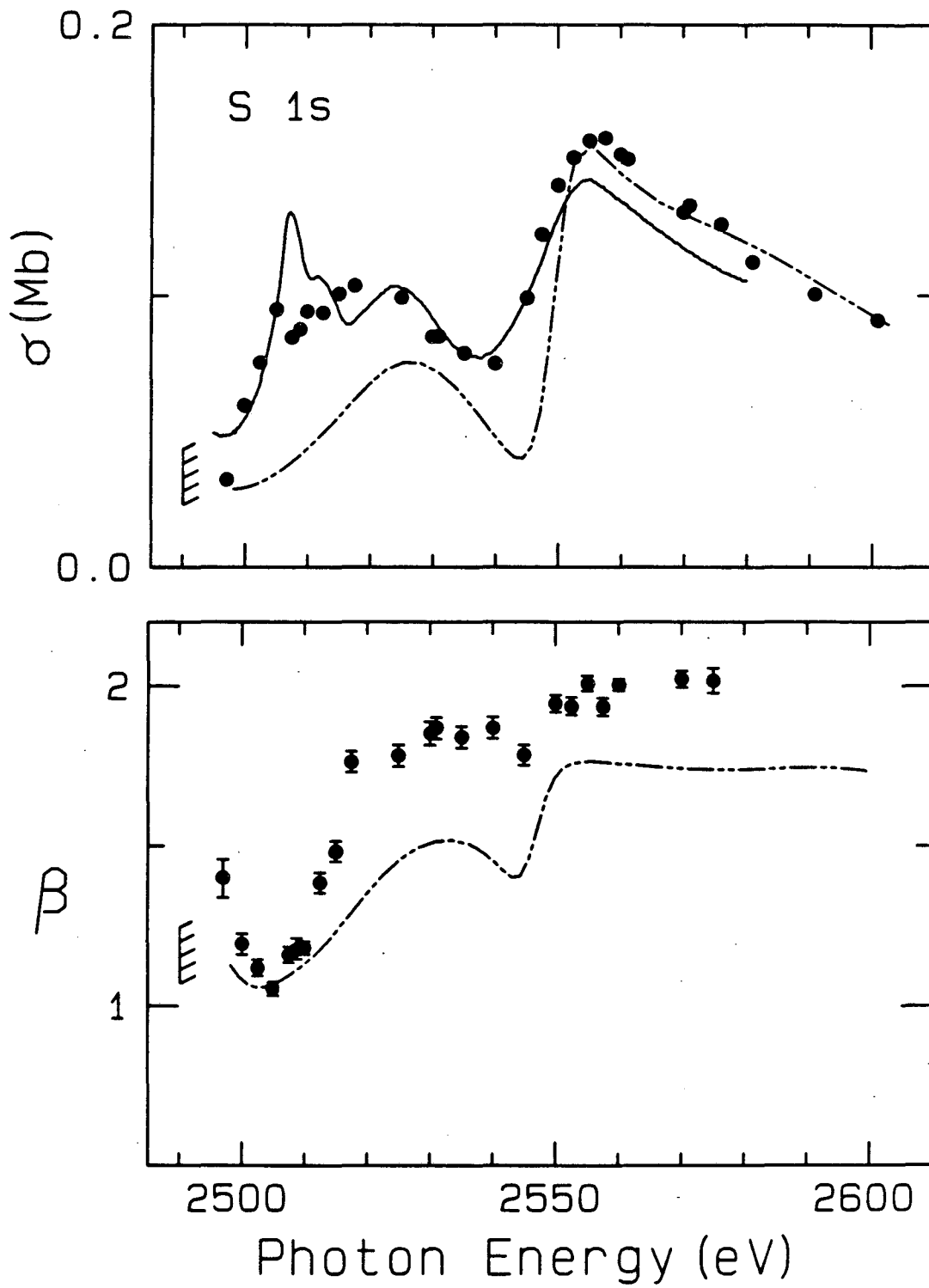
XBL 862-368

Figure 6



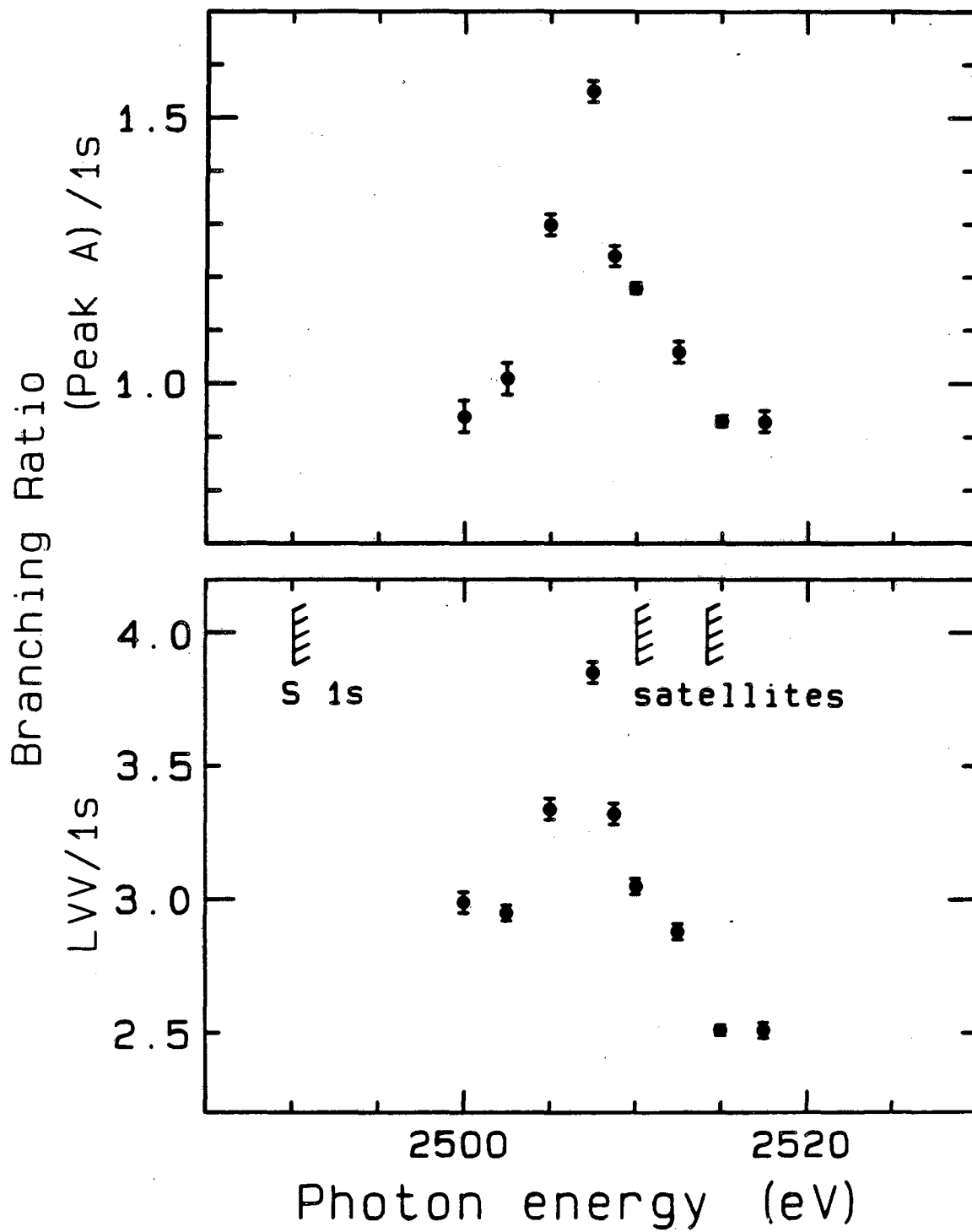
XBL 862-367

Figure 7



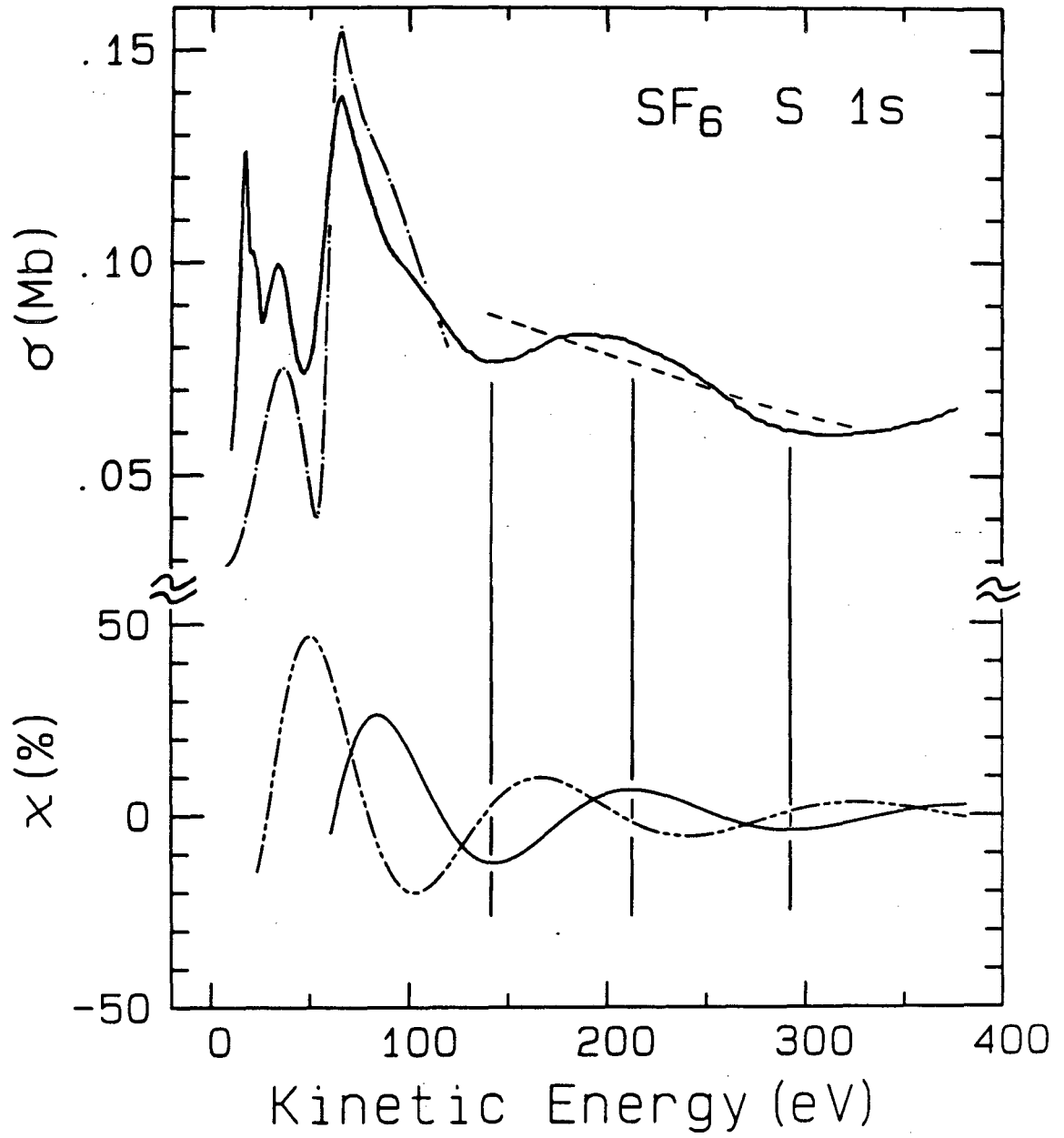
XBL 862-363

Figure 8



XBL 862-366

Figure 9



XBL 862-364

Figure 10

This report was done with support from the Department of Energy. Any conclusions or opinions expressed in this report represent solely those of the author(s) and not necessarily those of The Regents of the University of California, the Lawrence Berkeley Laboratory or the Department of Energy.

Reference to a company or product name does not imply approval or recommendation of the product by the University of California or the U.S. Department of Energy to the exclusion of others that may be suitable.

*LAWRENCE BERKELEY LABORATORY
TECHNICAL INFORMATION DEPARTMENT
UNIVERSITY OF CALIFORNIA
BERKELEY, CALIFORNIA 94720*