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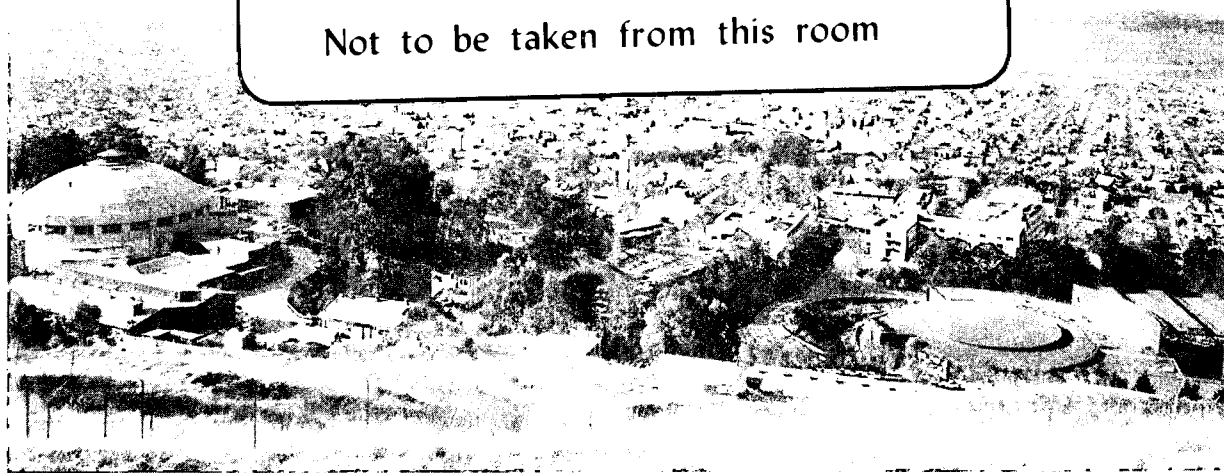
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**DYNAMICS OF ETHYLENE PHOTODISSOCIATION FROM
RO-VIBRATIONAL AND TRANSLATION ENERGY DISTRIBUTIONS
OF H₂ PRODUCTS**

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ABSTRACT

The dynamics of H₂ elimination from ethylene was studied via a pump and probe technique utilizing an ultrahigh resolution vacuum ultraviolet laser system. H₂ product internal and translational energy distributions were obtained for the photodissociation at 193nm. The distribution of energy in H₂ product from the dissociation of (1,1)-dideuteroethylene is also presented. Two separate H₂ elimination channels are inferred: a 1,1 elimination producing the vinylidene radical and a 1,2 elimination producing the acetylene molecule. Differences between the vibrational, rotational and translational energy distributions for these two channels are discussed and correlations between product internal and translational energy are presented. A comparison with ab initio calculations of the transition state configurations for these processes is made. We suggest that the H₂ elimination process may be non-statistical in nature. The D atom elimination from C₂D₄ was examined and kinetic energy distribution for this product measured.

I. INTRODUCTION

The study of the unimolecular decomposition of ethylene (C_2H_4) has been of great interest to the scientific community for more than 30 years. The importance of this molecule comes not only from its worth to industry as a precursor to many widely used materials but also from its value as a test of ab initio computation for a 6-atom system.

Early experiments determined the major unimolecular reaction pathways for C_2H_4 to be elimination of H (reaction 1) and elimination of H_2 (reaction 2); both forming acetylene.



These experiments were typically performed in a "bulb" using a variety of excitation sources including radiolysis,^{1,2} mercury sensitization,³ flash photolysis⁴ and discharge lamps.^{5,6,7,8,9}

It was found that the above two channels are roughly equiprobable for 193nm dissociation. However, a number of important questions remained regarding the structures of the intermediates and the electronic potential energy surfaces involved in the different reactive processes.

The next important steps came with the detection of two radical intermediates in the ethylene reaction scheme: the vinyl ($CH_2CH\cdot$) and vinylidene ($CH_2C:$) species. These species had both been predicted theoretically but it was not known whether their lifetimes would be long enough to permit detection. The presence

of $\text{CH}_2\text{CH}\cdot$ was confirmed initially by detection of products which could only result from the presence of $\text{CH}_2\text{CH}\cdot$ in the reactive process.^{4,10} The existence of the biradical, $\text{CH}_2\text{C}\cdot$, was not confirmed until much later when appropriate laser light sources had been developed in order to detect it by spectroscopic means.^{11,12,13} The vinylidene formed was found to be in an electronically excited triplet state ($^3\text{B}_2$), instead of its ground singlet state, $\text{A}(^1\text{B}_{1u})$. The temporal evolution of the triplet species suggested that it was not produced by the primary reactive process: this has been recently reinterpreted as being due to secondary photodissociation of the nascent vinyl radical¹⁴.

The mechanism of H-atom elimination from C_2H_4 has been well characterized.^{14,17,35} Following excitation with a vacuum ultraviolet (vuv) photon, C_2H_4^* can internally convert to the ground electronic state after which H-atom elimination occurs via simple bond rupture leading to the formation of energized $\text{CH}_2\text{CH}\cdot$. The excited vinyl radical can then either subsequently decay into acetylene plus another H-atom (provided the initial excitation energy is high enough) or react with other species that might be present. The mechanism for the elimination of molecular hydrogen from C_2H_4 , however, is not as well determined. The initially proposed reaction mechanism provided for two pathways: a 1,1 hydrogen elimination (1,1E) from one of the carbons leaving behind $\text{CH}_2\text{C}\cdot$ (reaction 3), or a 1,2 hydrogen elimination (1,2E) across the carbon-carbon double bond forming acetylene (reaction

4).



From studies of deuterated species the ratio for H_2 production between these two pathways, (3) to (4), was determined to be approximately 3:2.^{1,6,14} The total amount of energy available to the H_2 formed from reaction (3) is only 50-60 kcal/mole because of the high ΔH_f° of vinylidene (≈ 100 kcal/mole).¹⁵ The energy available to H_2 from reaction (4) is approximately twice as large at 106.3 kcal/mole. (The ΔH_f° 's for all pertinent species in the dissociation of C_2H_4 are given in Figure 1.)

The spectroscopy and excited states of ethylene have been reviewed in some detail.¹⁶ The planar ground state is labeled $N(^1\Sigma_g^+)$ while the first excited singlet state is designated $V(^1\Sigma_u^+)$. The broad, diffuse V-N band system extends from 215nm into the vuv with the intensity rapidly rising towards shorter wavelengths. It is considered to be the prototypical $\pi^* \leftarrow \pi$ transition. The first Rydberg transitions appear at 174.4nm. The V state of ethylene is twisted; it has a potential energy minimum for the methylene groups in a perpendicular configuration. It is capable, however, of free internal rotation for 193nm (148 kcal/mole) excitation. A third state predicted to be involved in the photochemistry of ethylene is the $Z(^1A_1)$ state

which has a π^* character. In this state the two methylene groups are pyramidalized¹⁷ and in that configuration the Z state becomes the lowest lying electronic state.

A considerable amount of theoretical work has been done relating to the problem of ethylene photochemistry.^{18,19,20,21} In Figure 2, we show a schematic correlation diagram²² for the relevant electronic states in ethylene. The excited V state of ethylene correlates directly, with small barrier, to the ground singlet state of vinylidene plus H₂. However, the V state correlates with neither the singlet ground state of acetylene nor the singlet ground state of the vinyl radical but does correlate with the singlet ground state of ethylidene (CHCH₃). Therefore, the 1,2E of H₂ and the H atom elimination should be viewed as occurring on the ground state surface. Ethylidene, initially postulated as an intermediate, has been shown to be unstable with respect to rearrangement. Recently, ab initio calculations, by M.Yoshimine, of some transition structures involved in the elimination of molecular hydrogen from C₂H₄ have been communicated to the authors.²² These results will be discussed later.

The experiments reported in this paper examine the rotational, vibrational, and translational energy distributions of the nascent H₂ product from the photodissociation of C₂H₄ at 193nm. The D-atom elimination channel from C₂D₄ was also examined for information about the translational energy of the D product. In order to distinguish between the two H₂ elimination channels

further experiments with deuterated ethylene were carried out. H_2 product from the dissociation of 1,1 dideuteroethylene (1,1- d_2) was analyzed in the same manner, allowing for a direct comparison between the products from the 1,1E and 1,2E channels in ethylene. The data presented below are consistent with two distinct channels for formation of H_2 each of which distributes differently the available energy amongst product degrees of freedom.

II. EXPERIMENTAL

The molecular beam apparatus and the vuv-xuv laser system used in these experiments have been described previously.^{23,24} A pump-and-probe technique was used to photolyze ethylene molecules in a molecular beam and then to probe the nasawnt H_2 products after a short time delay. The pump and probe beams were collinear and counterpropagating. The probe laser ionized the product H_2 molecules via (1+1) resonance enhanced multi-photon ionization (REMPI).^{25,26} The resultant ions were detected by time-of-flight mass spectrometry using a Daly type ion detector.

The probe laser system employed pulsed amplification of the output from a ring dye laser (Coherent 699-29). Computer controlled scanning of the ring laser provided the tuning ability of the probe laser. This high energy, near-transform limited pulse of visible light was then frequency doubled, forming ultraviolet (uv) light, and subsequently tripled or sum-frequency

mixed in a pulsed rare gas jet to produce vuv or extreme ultraviolet (xuv) radiation. A schematic view of the set-up of the laser and molecular beams in the main reaction chamber is shown in Figure 3.

A 193nm ArF excimer laser (Lambda-Physik EMG-103-MS) was used as the photolysis source. The temporal pulse width of the laser output was 20ns and the pulse to pulse jitter was ± 1 ns. An unstable resonator laser cavity was utilized in order to provide a uniform photolysis beam intensity in the interaction region and to maximize the brightness (watts/steradian) of the laser beam. An 8mm diameter section of the beam containing up to 10mJ of energy was selected and sent into the main chamber where it was then focused to a 2mm spot size with a 50cm CaF₂ lens. The excimer laser output was randomly polarized.

The beam steering optics were dichroic high reflecting, 45° mirrors coated for 193nm (Acton Research Inc.). The input window to the main chamber was a 25mm diameter, 3mm thick MgF₂ laser grade window which was rotated occasionally to avoid the attenuation of the laser due to the formation of color centers. All optics inside the main chamber were kept hot with a quartz heat lamp in order to minimize burn spot formation due to photolysis of adsorbed diffusion pump oil.

The 699-29 laser used in these experiments is extremely sensitive to external sources of vibration which can cause unstable operation. Since the excimer laser was on an optical table in contact with the 699-29 laser, it was necessary to

vibrationally isolate the excimer laser head (which contained a mechanically "noisy" gas processing unit) from the table. Electrical noise from the discharge in the laser chamber did not adversely affect the 699-29 laser.

It is essential, when determining product state distributions, to accurately normalize signals with respect to drift and fluctuation in output of both the pump and probe lasers. Therefore, careful measurement of the vuv-xuv and excimer laser pulse energies was required. A pyroelectric detector (Molectron J3-09) measured the 193nm pulse energy, as transmitted through one of the high reflectance mirrors. A sand-blasted fused silica window, used as a diffuser, was placed in front of the pyroelectric device as an attenuation source.

For xuv-vuv normalization, a platinum photodiode (p.d.) was used to monitor the probe laser intensity after the interaction region. The use of this type of system has been noted by Tonkyn and White.²⁷ A slightly different design was developed in this laboratory, a schematic drawing of which is shown in Figure 4. The vuv light impinges upon a disk of Pt metal ejecting electrons (by the photoelectric effect) which are collected by a biased metal ring. The charge lost from the Pt disk is then amplified via a charge sensitive pre-amplifier. The uv light is discriminated against by two methods. The high work function of the Pt metal (around 5.65eV) prohibits electrons from being ejected by single uv photons. However, in practice there was a fair amount of background electrons from the uv, perhaps due to

multi-photon effects. In order to further reduce this background, a negative (retarding) bias was applied to the collector ring. Since electrons generated by uv light will generally be of much lower kinetic energy than those generated by vuv photons, they will not be able to overcome as high a potential difference. Application of a -0.5 to -2.0 volts bias was enough to reduce the uv background signal. This bias was adjusted to maximize the ratio between vuv and uv signal. Under typical conditions, the ratio between the vuv and uv signals was 8:1.

A 25mm diameter, 2mm thick LiF window was used to reflect a fraction of the probe beam and send it directly in to the Pt photodiode (see Figure 2). This photodiode provided excellent discrimination against the uv light, which was collinear with the vuv radiation and $>10^4$ times as intense. Care was taken to limit the amount of scattered 193nm radiation in the interaction chamber as this was the main source of background signal from the Pt p.d. In cases where sum-frequency mixing was employed to generate vuv radiation, an additional precaution was needed. Frequency tripling can occur simultaneously with sum-frequency mixing and therefore both vuv and xuv radiation may be present in the probe beam. Since the detected H_2 signal depends only on the vuv light but the photodiode signal responds to both, a LiF window was placed in front of the Pt p.d., acting as a filter which transmits only the vuv photons.

For the measurements of D atom elimination it was necessary to generate Lyman-alpha radiation. This radiation was generated by sum-frequency mixing in a pulsed jet of Kr. This process has been previously reported in a low pressure gas cell.²⁸ The D atoms were then ionized via (1+1) REMPI in a fashion analogous to the H₂ ionization. The detection sensitivity for D atoms was estimated to be 10⁴ atoms/cm³.

The normal ethylene was ultra-high purity grade (>99.995%) from Matheson. The 1,1-d₂ ethylene was obtained from MSD Isotopes and was 98% pure. The C₂D₄ was also obtained from MSD and was 99% pure. All gases were used neat and without further purification.

For the determination of the population distributions of H₂ products, 800 mbar of C₂H₄ was expanded through the 0.5mm nozzle of a pulsed valve.²⁹ The pulse duration was 150μsec. The photolysis laser was timed so as to intersect the molecular beam 80μsec after the start of the pulse. A 1.02mm diameter skimmer (Beam Dynamics) was located 1.5cm from the nozzle separating the source chamber from the differentially pumped main chamber. The distance from the skimmer to the interaction region was 5cm. The interaction region was located in the extractor of a Wiley-McLaren type TOF mass spectrometer. The source chamber pressure was 2x10⁻⁴ torr and the main chamber pressure was <2x10⁻⁶ torr when the valve was in operation. For the long photolysis/probe time delay studies the skimmer was changed to a low profile 2mm diameter model, located 2cm from the nozzle and 4.5cm from the

interaction region. When using the deuterated compounds, the backing pressures were typically around 200mbar and the molecular beam pulse duration was reduced to 100 μ sec (to reduce gas consumption). Although clusters could be detected via single-vuv photon ionization when C₂H₄ was seeded in a rare gas carrier, none could be seen in the expansion of neat ethylene when the valve backing pressure was below 1000mbar.

The molecular beam diameter was calculated to be 5mm in the interaction region and the probe beam spot size was estimated to be 100-200 μ m. The delay between the photolysis and probe beams was 20ns for the population scans. The H₂⁺ signals were normalized to both the vuv dependent signal from the Pt. p.d. and to the 193nm dependent signal from the pyroelectric detector. All acquired signals were processed by boxcar integrators (SRS 250) and averaged slightly (1/3 of the data collection time constant) before being ratioed by analog processors (SRS 225/235). This was done to prevent overloading of the dividing circuitry by the large fluctuations in the vuv intensity. The averaged output from the final analog processor was then recorded by the 699-29 computer (Apple II). All data was transferred from the 699-29 to a PC computer for analysis.

III. RESULTS AND ANALYSIS

A. POPULATIONS

The vibrational distribution of the H₂ product from the dissociation of C₂H₄ at 193nm is shown in Figure 5. The rotational distributions of the same product for the vibrational levels v''=0 through v''=3 are shown in Figure 6. Each H₂ transition was scanned twice and, whenever possible, P, Q, and R transitions were used. This provided up to six separate determinations of the population of a given rovibrational state.

The H₂ quantum state populations were derived in the following manner. The lineshape of each transition was fit by a non-linear least squares method to the function:³⁰

$$I \propto \frac{C}{v_D} * [1 + \beta * P_2(\cos\theta) * P_2(\chi)] \quad (5)$$

where v_D =maximum doppler shift, β =effective anisotropy parameter, P_2 =second order Legendre polynomial, θ =angle between the electric field vector of the photolysis laser and the direction of the probe laser (which was 90 in our experiment), χ =Doppler shift/ v_D , and C is proportional to the line intensity. In our study, this function was convoluted with a normalized Gaussian H₂ velocity distribution of a variable width. This is justified by the observation of Gaussian-like velocity distributions in the molecular beam photofragment translational spectroscopy experiment reported elsewhere¹⁴. For overlapping transitions partial fits were done to that part of the lineshape that exhibited no interference. Line intensity measurements from

these partial fits were used in the population distributions but the Doppler widths and anisotropy parameters were not considered reliable.

An in-depth study of the anisotropy parameters was not undertaken during this experiment. The reasons are twofold. The experimental detection geometry did not lend itself to measuring correlations between the electric field vector of the photolysis laser and the product velocity or angular momentum distributions. Furthermore, the observations of other correlations (such as a (\mathbf{v}, \mathbf{J}) correlation) would be complicated by the presence of two H_2 elimination channels. While non-zero anisotropy parameters were measured ($-0.5 < \beta < 0.5$), no systematic behavior of the parameters was found.

The relative H_2 populations were calculated from the measured line intensities in the manner described previously.²³ When comparing populations for a given H_2 quantum state that were measured from different transitions (e.g. P-branch versus R-branch), significant discrepancies, due to inaccuracies in the calculated linestrength factors, were noted. The large error bars seen for a few of the H_2 quantum state populations in Figure 6 are a result of these discrepancies. Such differences had been seen previously for different H_2 transitions when comparing the ratio of calculated H_2 emission probabilities to measured values.³¹ The difficulty in calculating accurate linestrengths arises from perturbation of the energy levels due to rotational coupling between the $\text{B}^1\Sigma_u^+$ and $\text{C}^1\Pi_u$ electronic states of H_2 .

Reproduceability for a given transition was very good, attesting to the effectiveness of the normalization techniques used.

There are a number of immediate observations to be made from the distributions shown in Figures 5 and 6. First, the H_2 product is vibrationally very hot and the distribution of energy among the vibrational quantum states appears statistical. This indicates that the H-H distance at the ethylene transition state(s) is elongated with respect to the equilibrium H-H distance of molecular hydrogen. There is also a significant amount of translational and rotational energy released in the dissociation, with H_2 observed up to the $J=13$ level (which corresponds to ≈ 30 kcal/mole). This amount of rotational excitation suggests an asymmetric transition state in which the C-H distances are significantly different. In addition, the H_2 rotational distributions have unusual shapes, especially for $v''=2$. The non-statistical distributions, which look somewhat bimodal, could be indicative of two competing H_2 elimination channels: namely 1,1E and 1,2E. In a later section, we speculate on the nature of the energy release for each of these channels.

The rotational state population distributions of H_2 product from 1,1- d_2 ethylene were measured in order to help understand the normal ethylene distributions. In previous studies of the photodissociation of deuterated ethylene compounds no isotopic scrambling was observed^{1,6,14} indicating that H_2 elimination occurs on a time scale that is fast compared to isotopic rearrangement. All H_2 product from the photodissociation of

CD_2CH_2 , therefore, is assumed to come from the 1,1E channel. These rotational distributions for the vibrational levels $v''=0$ and $v''=1$ are shown in Figure 7. The error bars for the $v''=0$ levels represent the standard deviations of the various population measurements. A statistical non-linear least squares fit to each of the vibrational levels²³ is also presented in Figure 7. The rotational temperatures calculated are $2500 \pm 200\text{K}$ for $v''=0$ and $2000 \pm 160\text{K}$ for $v''=1$. Unfortunately, it proved impossible to obtain a reliable H_2 vibrational distribution for this system due to high background contamination in the mass spectrometer from D^+ signal. For similar reasons, we were not able to measure rotational distributions for $v''=2$ and $v''=3$ for this isotopic species.

A statistical fit to the $v''=0$ and $v''=1$ H_2 rotational distributions for C_2H_4 shown in Fig. 6 yields "temperatures" of $2700 \pm 250\text{K}$ and $2850 \pm 250\text{K}$ respectively. While for the $v''=0$ level the measured rotational "temperature" for H_2 product from CD_2CH_2 is only slightly less than that from C_2H_4 , the $v''=1$ level shows a significant difference. The measured H_2 rotational distributions from the dissociation of 1,1- d_2 ethylene, which eliminates H_2 only by reaction (3), clearly show that the amount of energy partitioned into H_2 rotation for this channel is less than the amount of rotational energy measured in the H_2 product from the dissociation of C_2H_4 . It is important to note that this observation is indicative of the existence of two separate elimination channels. The amount of rotational energy in H_2 from

the 1,2E channel must be higher than that from the 1,1E channel in order to properly fit the total H₂ rotational distribution.

Energetically, one can understand the decrease in rotational energy for higher vibrational levels, especially for 1,1E, by considering that for $v''=3$ the H₂ vibrational energy is 33.6kcal/mole. Given the translational energy of ≈ 15 kcal/mole for this level, there is much less energy available for H₂ rotation than at the lower vibrational levels. Furthermore, the 1,2E channel has 106kcal/mole of energy available for products while the 1,1E channel has only 65kcal/mole available. This makes >50% more energy available to product from the 1,2E channel. This is one contributing factor to the higher H₂ rotational energy for 1,2 elimination.

The calculated transition state²² for the 1,1 elimination channel (see Fig. 1) is similar to that for H₂ elimination from formaldehyde. Rotational energy distributions for H₂ from the photodissociation of formaldehyde have been measured at a number of different excitation energies.³² While the formaldehyde experiments were done at energies much closer to the threshold for dissociation, the total amount of energy available to the H₂ product is comparable to that for the 1,1E channel (≈ 86 kcal/mole for H₂CO and 65kcal/mole for C₂H₄). The rotational distributions measured by Butenhoff et. al. for H₂ elimination from H₂CO were Boltzmann-like with temperatures of 1730K for the $v''=1$ level and 1240K for the $v''=3$ level. The amount and distribution of rotational energy imparted to H₂ in the two different

dissociative processes is comparable, as are the initial geometries.

Parallels can be drawn between the dynamics of the 1,1E channel of C_2H_4 and H_2 elimination from H_2CO . The formaldehyde dissociation has been successfully described by a modified impulsive model³³ with a large impact parameter. In this model, a single impulse between the departing fragments determines the rotational energy for a given impact parameter, direction of impulse, and available energy. The width of the rotational distributions is accounted for by vibrational motion of the parent molecule at the transition state. The H_2 rotational energy distributions appear statistical while the corresponding CO distributions are peaked at high J and are non-statistical in nature. For the case of H_2 elimination from ethylene via 193nm photolysis, the total energy (148kcal/mol) is much greater than the transition structure energies (< 100kcal/mol) and therefore we expect that an impulsive model applied to the calculated transition state geometries will not be meaningful. In other words, due to the excess internal energy, the transition structure is severely distorted with respect to the calculated saddle point minimum structure.

Experimental evidence for this point is revealed in the distribution of rotational energy observed for the H_2 product in the 1,2E channel. The transition structure in this case (see Fig. 1) was calculated to have equal C-H bond lengths - suggestive of low rotational excitation for the products.

However, the large amount of rotational excitation suggests a highly distorted configuration at the transition state much different than the calculated minimum energy structure.

It is possible, with some speculation, to further analyze the H₂ distributions from C₂H₄. Specifically, we tried to estimate the contributions of 1,1E and 1,2E to the distributions for H₂(v''=0) and H₂(v''=1). This was done in the following manner. The ratio of the total (i.e. not state specified) H₂ product from the 1,1 elimination channel (reaction (3)) to the 1,2 elimination channel (reaction (4)) was taken (from previous measurements using deuterated compounds)^{1,6,14} as 3:2. However, we could not measure the relative contribution of each elimination channel to a given vibrational level, i.e. we did not have the vibrational population distributions for the two separate channels. In the ab initio studies of Yoshimine, discussed above, the H-H distance at the transition state for the 1,1E channel is calculated to be 0.878Å while the corresponding distance for the 1,2E channel is 0.941Å.²¹ We expect that a larger H-H distance at the transition state should correlate with greater vibrational excitation of the H₂ product. Using a harmonic approximation for the H-H extension at each transition state relative to a H₂(v''=0) product molecule, we estimate that the vibrational "temperature" for H₂ from the 1,2E channel should be approximately twice that for H₂ from the 1,1E channel. With the above assumptions, the percent contribution of the 1,1E channel to the H₂ product in the vibrational levels v''=0 to v''=3 was 67%, 49%, 32%, and 19%

respectively.

For the $v''=0$ and $v''=1$ levels, the rotational temperature of H_2 product from 1,1E (reaction (3)) is known from the deuterated ethylene study. The rotational distribution of H_2 product from the 1,2E channel for the $v''=0$ and $v''=1$ levels was estimated by making a non-linear least squared fit to the difference between the total H_2 distribution and the fit for 1,1E (as obtained from the dideuteroethylene study). The rotational "temperatures" obtained in this manner for the 1,2E channel are $3500 \pm 940K$ for $v''=0$ and $4000 \pm 800K$ for $v''=1$. It is noteworthy that these "temperatures" are significantly hotter than for the 1,1E case.

Due to the speculative nature of the above "temperatures", one must be cautious when making inferences from the data. The "temperatures" derived for the 1,2E channel have very large error bars. This might indicate that this channel produces a non-statistical distribution of rotational energy in the H_2 product. Further evidence for this comes from the $v''=2$ level - the level that showed the clearest evidence of bimodality. It is not possible to reasonably fit the $v''=2$ rotational distribution with two Boltzman distributions.

B. AVERAGE TRANSLATIONAL ENERGY

In the case of concerted decomposition there is a potential energy barrier in the exit valley and a coupling between relative translation of the products and their internal degrees of

freedom. This coupling is indicative of a "tight" transition state. In general, for a concerted decomposition, once the reaction proceeds past the critical configuration in the exit channel, the products formed near the top of the potential energy barrier are rapidly repelled from each other with enhanced translational energy. Additionally, translational energy may be gained by initial parent motions along the reaction coordinate and by the coupling of vibrations at the transition state into translation through exit channel interactions. The formation of H_2 in the photodissociation of ethylene can be described as proceeding through a "tight" transition state. For this case, we expect the potential energy release to be the dominant contribution. This has been observed previously for other concerted H_2 elimination reactions with large potential energy barriers.^{25,32}

The average H_2 velocity associated with each quantum state was calculated from the Doppler fits to the individual line shapes by the procedure described above. From these velocities the average H_2 translational energy ($E_{tr} = \frac{1}{2}mv^2$) was calculated. A plot of E_{tr} for H_2 versus the internal energy of the molecule is shown in Figure 8 along with a linear least squares fit to the data. Only those transitions which exhibited no interference from other nearby lines were used in this analysis. It is seen that the product translational energies are high (around 20kcal/mol) and that they decrease slightly with increasing internal energy.

It would be tempting to treat the elimination of H_2 from ethylene as the dissociation of a quasi-triatomic molecule (where the hydrocarbon fragment is the third particle) because the calculated transition states²² reveal that for both channels, the two H atoms recoil from a single carbon atom. In a real triatomic molecule there is a 1:1 trade-off between translational and internal energy since the total energy is fixed. In a plot such as that shown in Figure 8, all data points would lie on a straight line with slope -1. The reason why this is not observed in the current experiment is that the major part of the total energy remains within the internal degrees of freedom of the hydrocarbon fragment. In other words, the sum of internal plus translational energies for the H_2 fragment need not be constant.

The following simple model explains the observed correlation between the H_2 translational and vibrational energies. The model contains two features which are consistent with the observed correlations. At the transition structure, the repulsive energy is expected to depend on the H-H interatomic distance: as the H-H interatomic separation increases, the forming H_2 molecule looks less like a closed-shell molecule and, therefore, a weaker repulsion is anticipated. A negative correlation follows from the fact that transition structures with larger H-H interatomic distances are expected to correlate with vibrationally excited H_2 product; concomitantly, weaker repulsion is expected to result in lower translational energy.

A second contribution to the observed negative correlation arises from the fact that fairly close to the transition structure (on the exit valley side), the vibration of the H₂ product molecules is expected to become adiabatic. A quantum of H₂ vibration is very large compared to other frequencies in the problem. However, the H₂ vibrational frequency is at this point still significantly less than that of a free H₂ molecule. Therefore, as the H₂ molecule is repelled, some of the repulsive energy release appears as vibrational energy of the H₂ molecule and, consequently, less energy will be available for translation. Hence, a negative correlation is expected wherein the H₂ translational energy decreases as the vibrational energy increases.

It is also instructive to analyze the amount of H₂ translational energy as a function of rotational energy. The distribution of translational energies, shown in Fig. 8, is separated into the various vibrational levels for this purpose. For each vibrational level the translational energy decreases slightly with increasing rotational energy. The behavior for each v" level is somewhat linear. Linear least-squares fits to each vibrational level gives slopes of -0.15, -0.19, -0.10, and -0.10 for v"=0 to v"=3 respectively. There is a stronger correlation between vibrational and translational energy than between rotational and translational energy for H₂ elimination from ethylene. (There is undoubtedly an effect due to the presence of two H₂ elimination channels which complicates any

analysis).

Two transition state models which would lead to a correlation between rotational and translational energy are shown in Figure 9. These represent limiting cases for the approach to the transition state for H₂ elimination from a polyatomic molecule. In this picture, one H atom is much closer to the carbon center than the other. The C,H repulsion for the close H atom is much greater than for the other H atom and therefore makes the dominant contribution to the translational energy of the H₂ product. Due to the excess energy in the reaction, the transition state must be vibrationally excited. The vibrational motions at the transition state can affect the correlations between internal and translational energies of the products.

The first limiting case represents C-H bond rupture where the second H atom interacts only weakly with the carbon atom and the transition state is achieved through motions of the closer H atom. In this case, an increase in repulsive energy will lead to a simultaneous increase in translational energy and rotational energy (since a larger torque is exerted on the H₂ species). Therefore, a weak positive correlation would be expected between translational and rotational energy.

The second limiting case represents a situation analogous to a "fly-by pick-off" abstraction reaction. In this case, the transition state is achieved through motions of the farther H atom. As the farther H atom approaches the carbon atom, repulsion increases and enhanced translational energy is expected. However,

this same motion also decreases the H-H separation and therefore a smaller torque is exerted on the H₂ species, decreasing the rotational excitation. Consequently a weak negative correlation is expected.

In sum, if the range of motions at the transition state involves the closer of the two H atoms, we expect a weak, positive correlation and if the range of motions involves the farther H atom, then we expect a weak, negative, correlation between H₂ translational and rotational energy. Therefore, we suggest that the H₂ elimination from ethylene is more consistent with the limiting case shown in figure 9b. It should be pointed out that these models are consistent with the aforementioned discussion of vibrational correlations.

C. H₂ VELOCITY DISTRIBUTION

One of the interests of this experiment was to determine if a difference in translational energy of the H₂ product from the two elimination channels of C₂H₄ could be measured. Although the distribution of the H₂ internal energies suggests a bimodal distribution there is no clear evidence of this in the Doppler profiles themselves, even though the instrument resolution was <0.1cm⁻¹. The technique of velocity aligned Doppler Spectroscopy (VADS), first noted by Wittig³⁴ was used to improve the translational energy resolution of the experiment.

In VADS, a long time delay is introduced between the formation of the product and its measurement. In addition, the diameter of the probe and photolysis beams are kept as small as possible. Under these conditions any product with a velocity component directed perpendicular to the probe beam will have moved out of the region that is interrogated by the probe beam during the time between when the photolysis and probe beams are fired. This effectively discriminates against those molecules so the only product molecules that are detected are those directed at or away from the probe beam. The product velocity resolution of the experiment is then greatly enhanced and one can obtain a true speed distribution of the product. A diagram showing this effect is given in Figure 10.

A number of transitions for the $v''=1$ and $v''=2$ levels were measured under long time delay, or VADS, conditions. Those transitions which exhibited no interference from other lines and had large signals (the signal is significantly reduced under the VADS conditions) produced the best results. The delay between the photolysis and probe lasers needed to observe a VADS profile was dependent on the experimental parameters. The most critical of these was the size of the photolysis beam in the interaction region. The timing varied between 100ns for a small photolysis beam to 400ns for a larger beam. The delay was set at the point where no further depletion in signal at the line center could be observed.

For the $v''=1$ level the following H_2 transitions were observed under VADS conditions: B-X (1,1) R(1), B-X (2,1) P(3), and B-X (2,1) R(5). For the $v''=2$ VADS profiles were measured for the following transitions: B-X (2,2) R(0), B-X (2,2) R(1), and B-X (3,2) P(5). The measured Doppler profiles under normal and VADS conditions for the B-X (2,1) P(3) transition is shown in Figure 11. It is obvious, from Fig. 11, that the distribution of translational energy in the H_2 product is peaked away from zero. This is consistent with molecular beam time-of-flight measurements of this distribution.¹⁶

The VADS profiles provide information about the distribution of internal energy in the acetylene product. In order to calculate the amount of internal energy in the acetylene molecules that correspond to H_2 produced in the six quantum states for which the VADS profiles were obtained, the probe and photolysis lasers were approximated as overlapping lines and a $1/v^2$ weighting was applied to the H_2^+ intensities in the VADS profiles. While this is a fairly large approximation, it greatly simplifies the calculation and the average H_2 translational energies (measured from the peak of the weighted VADS profile) agree well with those measured from normal Doppler profiles. The calculated acetylene internal energy distributions, therefore, should be reasonable approximations to the true distributions. For each VADS profile the peak acetylene internal energy and the FWHM (full width at half maximum) of the distribution was calculated according to:

$$E_{C_2H_2}^{int} - E_{avail} - E_{H_2}^{int} - E_{H_2}^{tr} - E_{C_2H_2}^{tr}$$

These results are given in Table I. The FWHM's are roughly centered about the peak of the distributions.

TABLE I: C ₂ H ₂ INTERNAL ENERGY DISTRIBUTIONS AS MEASURED BY VADS			
H ₂ TRANSITION	V(0) (cm ⁻¹)	PEAK E _{int} (kcal/mole)	E _{int} FWHM (kcal/mole)
B-X(1,1)R(1)	87358.22	71.5	24.5
B-X(2,1)P(3)	88077.07	70.4	21.2
B-X(2,1)R(5)	87709.96	69.0	19.9
B-X(2,2)R(0)	84751.31	68.5	15.5
B-X(2,2)R(1)	84714.90	64.8	18.0
B-X(3,2)P(5)	84729.84	61.5	17.0

The v''=2 rotational distribution, shown in Fig.6, gives the clearest evidence of a bimodal product energy distribution. This lead us to undertake a very long time-delay VADS study of the B-X (2,2)R(1) transition with the hope that the improved velocity resolution (approx. 8%) would allow us to resolve the 1,1E and 1,2E contributions to the VADS lineshape. The result of this study is shown in Figure 12. The Doppler shift extends to almost 4cm⁻¹, corresponding to a H₂ translational energy of 46kcal/mol. Given the H₂ internal energy (v''=2, J''=1 ; 23kcal/mol) and the reaction exothermicities from Fig.1, this wing of the Doppler profile must correspond to the 1,2E channel. We might expect to see a slower component, due to 1,1E, superimposed on the 1,2E

lineshape. We do not see clear evidence of this despite the significant differences in exothermicity for the two channels. This is consistent with the majority of the available energy being partitioned to the hydrocarbon fragment, obscuring energetic differences as measured by the H_2 Doppler lineshape.

One would expect that the Doppler shift observed for H_2 dissociating from ethylene through both 1,1 and 1,2 elimination would be larger than that for H_2 dissociating from 1,1- d_2 ethylene solely from 1,1 elimination. Although no VADS studies were performed for the deuterated species, a comparison of the H_2 normal Doppler profiles measured from C_2H_4 and $C_2H_2D_2$ reactions for the transitions C-X (1,0) Q(5), C-X (2,0) Q(7), and B-X (2,1) P(3) was made. The H_2 translational energies measured from these transitions are given in Table II. As one proceeds to higher energy H_2 quantum states, the excess energy for 1,1 elimination becomes considerably less. As a result of this, the H_2 translational energy decreases faster for product from the 1,1- d_2 ethylene than from C_2H_4 .

TABLE II: TRANSLATIONAL ENERGY OF H_2 DISSOCIATED FROM C_2H_4 AND CH_2CD_2 AT 193nm		
H_2 TRANSITION	$E_{tr}(H_2) / C_2H_4$ (kcal/mole)	$E_{tr}(H_2) / CH_2CD_2$ (kcal/mole)
C-X (1,0) Q(5)	21.8	22.1
C-X (2,0) Q(7)	21.3	19.3
B-X (2,1) P(3)	21.2	14

D. D-ATOM ELIMINATION

In addition to probing the H₂ elimination channels in the dissociation of C₂H₄, experiments were performed to gain information on the H atom elimination process in this molecule. Due to the presence of large background H⁺ signals it was necessary to measure D atoms eliminated from C₂D₄ photodissociated at 193nm.

Doppler scans of the D atom Lyman-alpha transition at both short and long (VADS condition) photolysis/probe time delays are shown in Figure 13. Both scans peak at zero Doppler shift and are fairly narrow in energy as compared with the H₂ translational energy. This is consistent with a simple bond rupture in which the translational energy distribution is peaked near zero and with previous measurements of the translational energy distribution in D (and H) atoms eliminated from C₂D₄ (and C₂H₄).^{14,35} The HWHM of the line profiles are 1.43cm⁻¹ and 1.57cm⁻¹ for the short and long time delay scans respectively, corresponding to translational energies of 6.5kcal/mole and 7.8kcal/mole respectively.

An attempt was made to measure the maximum D atom translational energy release in order to estimate the C-D bond strength in C₂D₄. D atoms were observed with up to ≈70kcal/mole of translational energy. This would correspond to a C-D bond strength of only ≈78kcal/mole, well below accepted values. Clearly, these fast D atoms cannot be the result of a single photon absorption. Photolysis power dependence studies done on

the fast D atom component showed a linear response over an order of magnitude of change in the photolysis energy density. However, this does not rule out a two photon absorption process, as has been previously assumed in other studies of this reaction.³⁶ If the cross section for absorption of a second photon by a product molecule (such as the vinyl radical) is much greater than that of the first photon this secondary absorption process can be saturated. If this is the case then power dependence studies will produce a linear result.

For the case of ethylene, where the absorption cross section of the product vinyl radical is much greater than that of ethylene,^{36,37} a careful photolysis laser power study as a function of D atom Doppler shift did reveal a non-linear behavior at low intensities, confirming the multiphoton nature of the process.³⁸ This has been observed previously for other concerted H₂ elimination reactions with large potential energy barriers.^{25,32} This suggests to us that the nascent vinyl radicals produced in the dissociation of ethylene can absorb a second 193nm photon to produce a second D (or H) atom.

IV. SUMMARY

The evidence presented in this paper clearly supports the existence of two distinct channels for the elimination of H₂ from C₂H₄. The existence of these channels, the 1,1 elimination shown in reaction (3) and the 1,2 elimination shown in reaction (4),

has been proposed previously and supported by numerous experiments. However, the information presented here provides an insight into differences in the dissociation dynamics of the two processes.

For the 1,1 elimination process, the amount of released energy is limited by the heat of formation of the vinylidene biradical. The average translational energy of H_2 from this channel is less than the total average H_2 translation energy observed. The amount of rotational energy imparted into the H_2 product from this dissociation pathway is also significantly lower than from the other channel. The critical configuration near the transition state for this reaction we suggest to be analogous to that for H_2 elimination from formaldehyde.

There is $\approx 50\%$ more energy available to H_2 product formed in the 1,2 elimination process. This larger amount of available energy is reflected in the rotational, vibrational, and translational energy of that H_2 product. The higher vibrational energy implies a longer H-H distance at the critical configuration of the 1,2 elimination reaction than of the 1,1 elimination reaction. This is consistent with recent ab initio calculations of the 1,1E and 1,2E transition states²². The higher rotational energy for the 1,2 elimination is due mostly to the larger kinetic energy release for this channel, despite the fact that the calculated transition state for 1,2E is more symmetric than for 1,1E. This implies that the critical configuration of the 1,2E transition state is significantly

distorted from the calculated minimum energy structure by the excess available energy. Furthermore, the poor fit of the 1,2E rotational distribution with a rotational temperature suggests that the actual distribution is non-Boltzmann in nature.

Recently, we learned that the barrier in the ground electronic state to 1,2 hydrogen exchange is calculated to be ≈ 75 kcal/mole, corresponding to the heat of formation of ethylidyne.²² This barrier is significantly lower than that for either 1,1E H₂ elimination (at 95 kcal/mol) or 1,2E H₂ elimination (at 110 kcal/mol). We would expect, therefore, that if the dynamics on the ground state were statistical in nature, then there should be complete isotopic scrambling on the timescale of reaction (i.e. no difference in HD yields between 1,1 dideuteroethylene and 1,2 dideuteroethylene). This is contrary to what is observed.¹⁴ We conclude, therefore, that the elimination reaction exhibits non-statistical dynamics. This is very interesting: as the molecule has 12 internal degrees of freedom and almost 150 kcal/mol of internal energy, one might expect intramolecular vibrational energy redistribution (IVR) to be extremely rapid and the dynamics, it follows, to be statistical. In the case of 1,1 elimination, excited state ethylene can move directly from the pyramidalized configuration at the crossing of the N and Z states (see Fig. 2) to the 1,1E transition state on the ground state by a simple twisting motion, thus avoiding hydrogen "scrambling". In the case of the 1,2E channel, calculations have not yet shown if excited state

ethylene can proceed directly to the 1,2E transition state on the ground state without going through an ethylidyne-like structure (where 1,2 hydrogen exchange may readily occur). However, it is also possible that the system does proceed through such a configuration but traverses quickly as it moves towards the 1,2E H₂ elimination transition state and the product channel.

Other information presented in this paper helps to further elucidate the dynamics of the dissociation processes. The VADS profiles confirm a concerted elimination mechanism and give an indication as to the distribution of internal energy in the hydrocarbon fragment. The D-atom Doppler profiles confirm that this atomic elimination process occurs on the ground state potential surface. Hopefully, these results will stimulate further experimental and theoretical work, providing a more complete picture of the dissociation dynamics of this important molecule.

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FIGURE CAPTIONS

Figure 1. Standard heat of formations of the pertinent species involved in the dissociation of C_2H_4 at 193nm. Energies are given in kcal/mole. Two transition structures, labeled 1,1E and 1,2E, are shown. Their geometries and energetics are based upon the work of Yoshimine (ref. 22).

Figure 2. State correlation diagram showing curve-crossing in ethylene, based upon the work of Yoshimine (ref. 22). As the methylene groups twist, the initially prepared V state intersects the Z state which, in turn, intersects the ground state (N) upon pyramidalization. The transition, V to Z to N, is an internal conversion process and is thought to be very fast. We expect that the photochemistry of ethylene should be described as occurring on the singlet ground state potential energy surface.

Figure 3. Schematic of the interaction area in the main chamber for the dissociation of C_2H_4 experiment. The substrates of the various optical components are indicated. Pt P.D.=platinum photodiode, QL = quartz heat lamp, and BD = Wood's Horn beam dump for 193nm beam reflection.

Figure 4. Schematic of the Platinum Photodiode used to measure vuv and xuv radiation intensity. The main body (shown roughly to scale) is constructed of glass and the Pt is held in place with

wire. -V = Low voltage power supply, CR = electron collector ring.

Figure 5. Distribution of vibrational energy in H_2 product from the dissociation of C_2H_4 (solid bars). A statistical fit to the distribution ($T = 4970 \pm 120K$) is also shown (hatched bars).

Figure 6. Distribution of rotational energy in H_2 product from the dissociation of C_2H_4 for the vibrational levels $v''=0$ to $v''=3$. The various rotational distributions as well as the vibrational distribution (Fig. 3) have the same relative scale. The error bars represent \pm one standard deviation of the measurements.

Figure 7. Distribution of rotational energy in H_2 product from the dissociation of 1,1- d_2 ethylene for the vibrational levels $v''=0$ and $v''=1$ (hatched bars). Statistical fits to the distributions ($T_r(v''=0) = 2500 \pm 200K$, $T_r(v''=1) = 2000 \pm 160K$) are also shown (solid bars). The error bars represent \pm one standard deviation of the measurements. The distributions are not scaled to each other.

Figure 8. Plot of translational energy versus internal energy for H_2 product from the dissociation of C_2H_4 . The various vibrational quantum states of the H_2 product are indicated.

Figure 9. Cartoon of a 3-centered transition state for H_2

elimination. (Top) Motion leading to a positive correlation between rotational and translational energy. (Bottom) Motion leading to a negative correlation between rotational and translational energy. The latter case is consistent with our observed correlations.

Figure 10. Pictorial representation of the time delay effect used in Velocity Aligned Doppler Spectroscopy (VADS). Shown are (Top) probe beam/ H_2 product overlap and expected line profile at zero time delay and (Bottom) probe beam/ H_2 product overlap and expected line profile at a long time delay (400ns). The cross hatched areas represent the spatial distribution of the H_2 dissociated from C_2H_4 .

Figure 11. H_2 Doppler profiles measured under VADS conditions for the $v''=1$ level transition H_2 B \leftarrow X (2,1) P(3). The VADS profile is shown with a heavy line while the normal Doppler profile is shown with a light line. The intensity scales are not related.

Figure 12. Red side of VADS profile for B-X (2,2) R(1) transition at 700ns (top trace) time delay. The trace is an average of 10 scans.

Figure 13. Scans of the Lyman-alpha transition ($\nu_0=82282.65 \text{ cm}^{-1}$) for D atoms eliminated from C_2D_4 dissociated at 193nm under

normal delay (top trace) and VADS conditions (bottom trace). The arrow indicates the maximum kinetic energy expected for a single photon event.

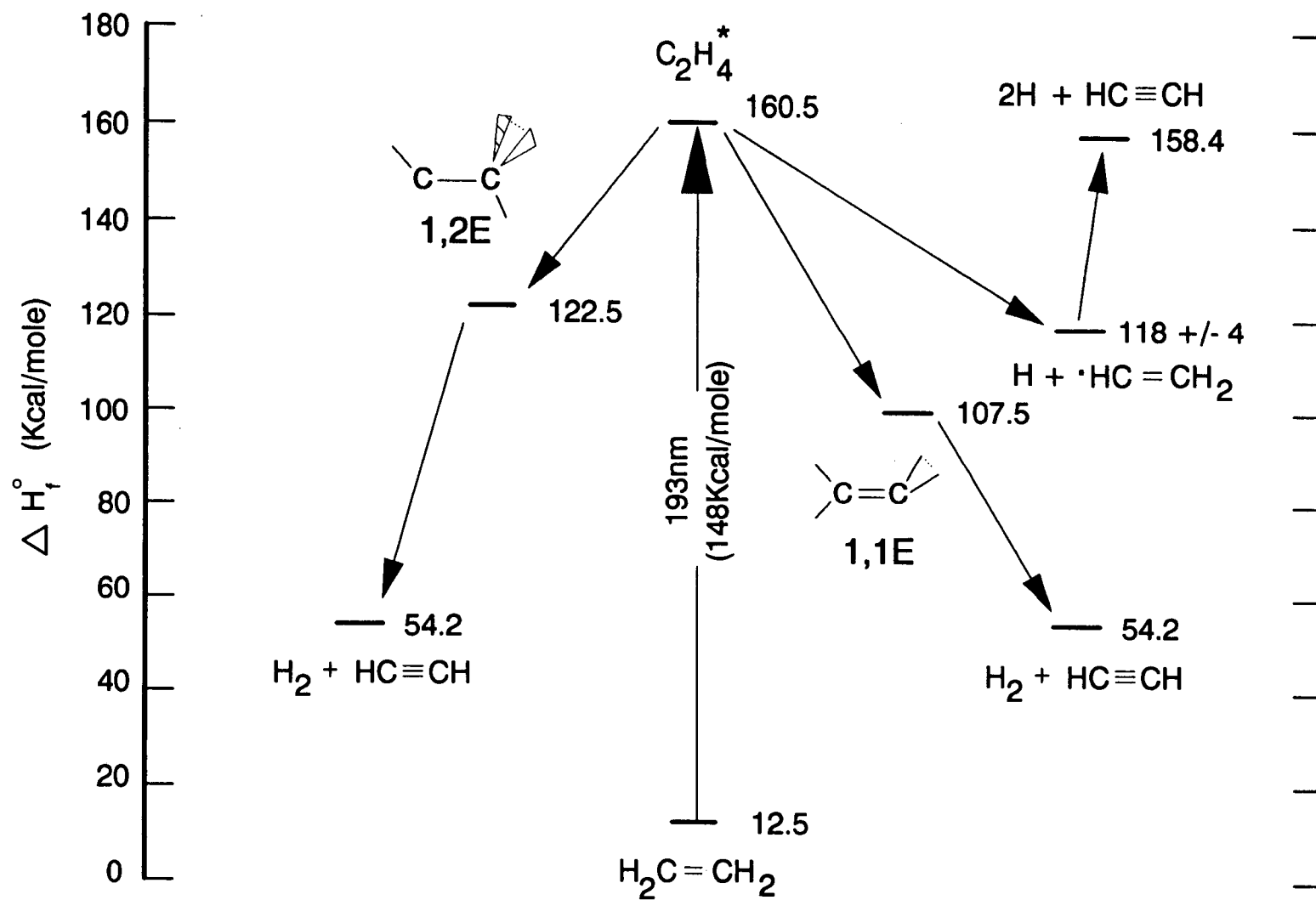


Fig. 1

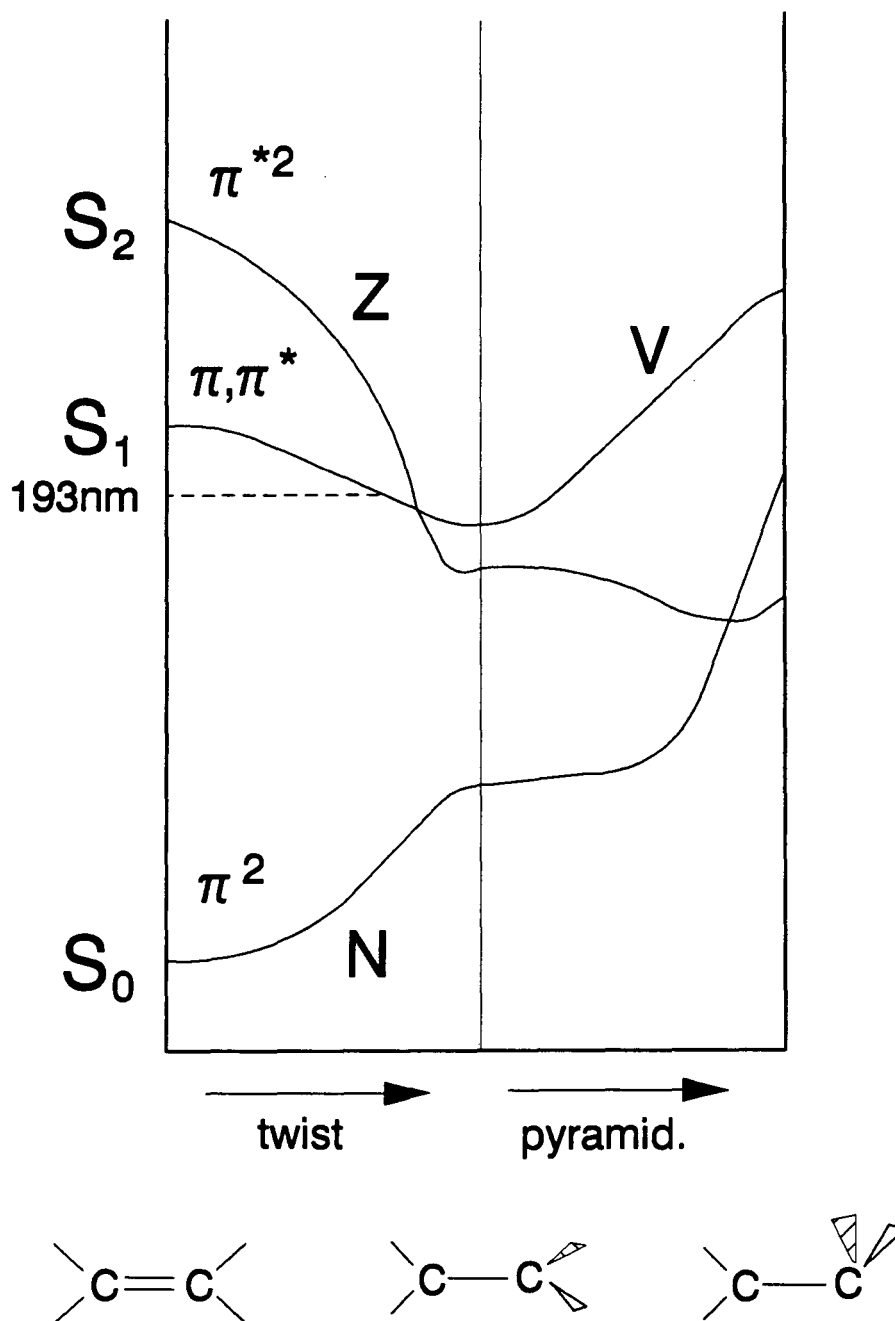


Fig. 2

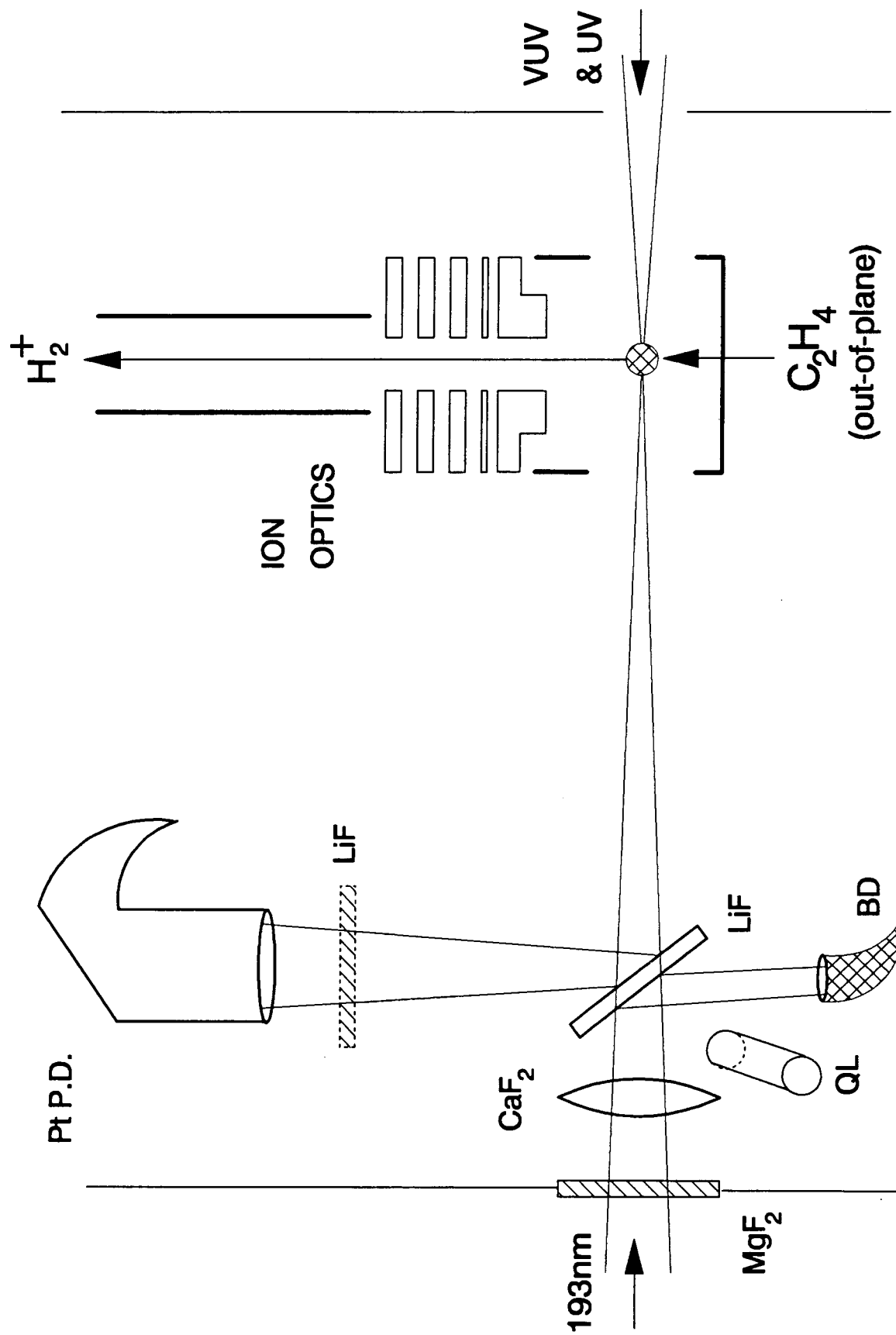


Fig. 3

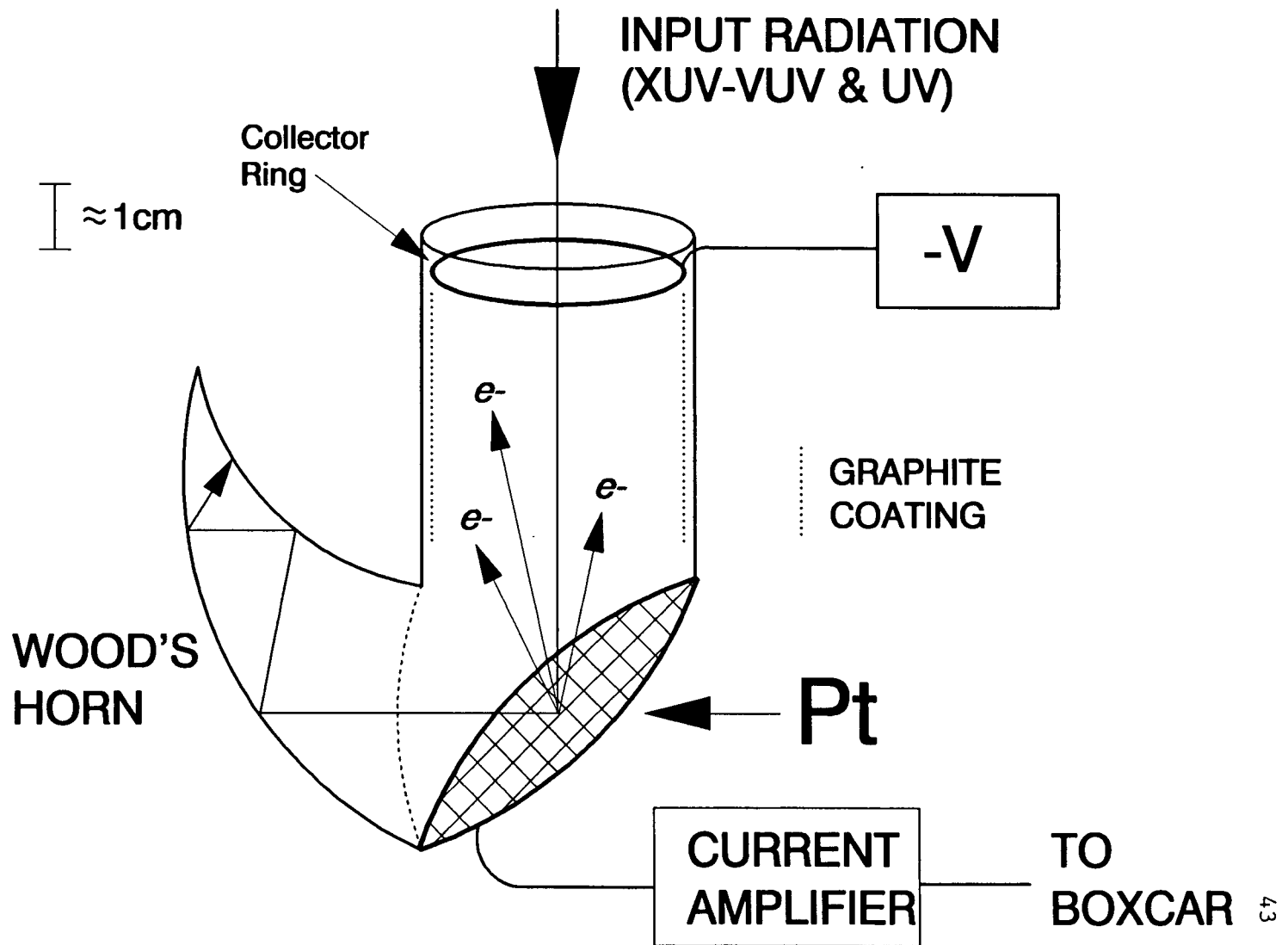


Fig. 4

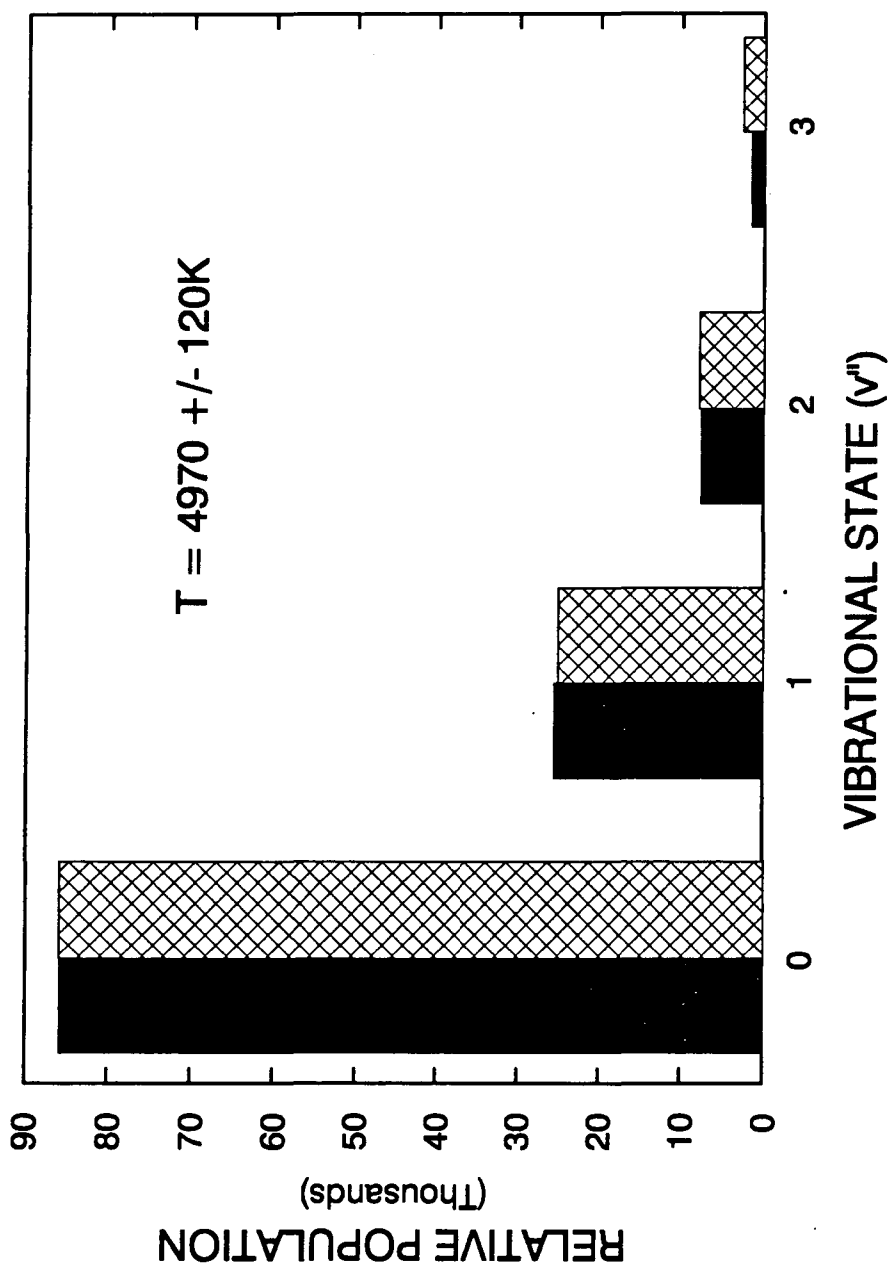


Fig. 5

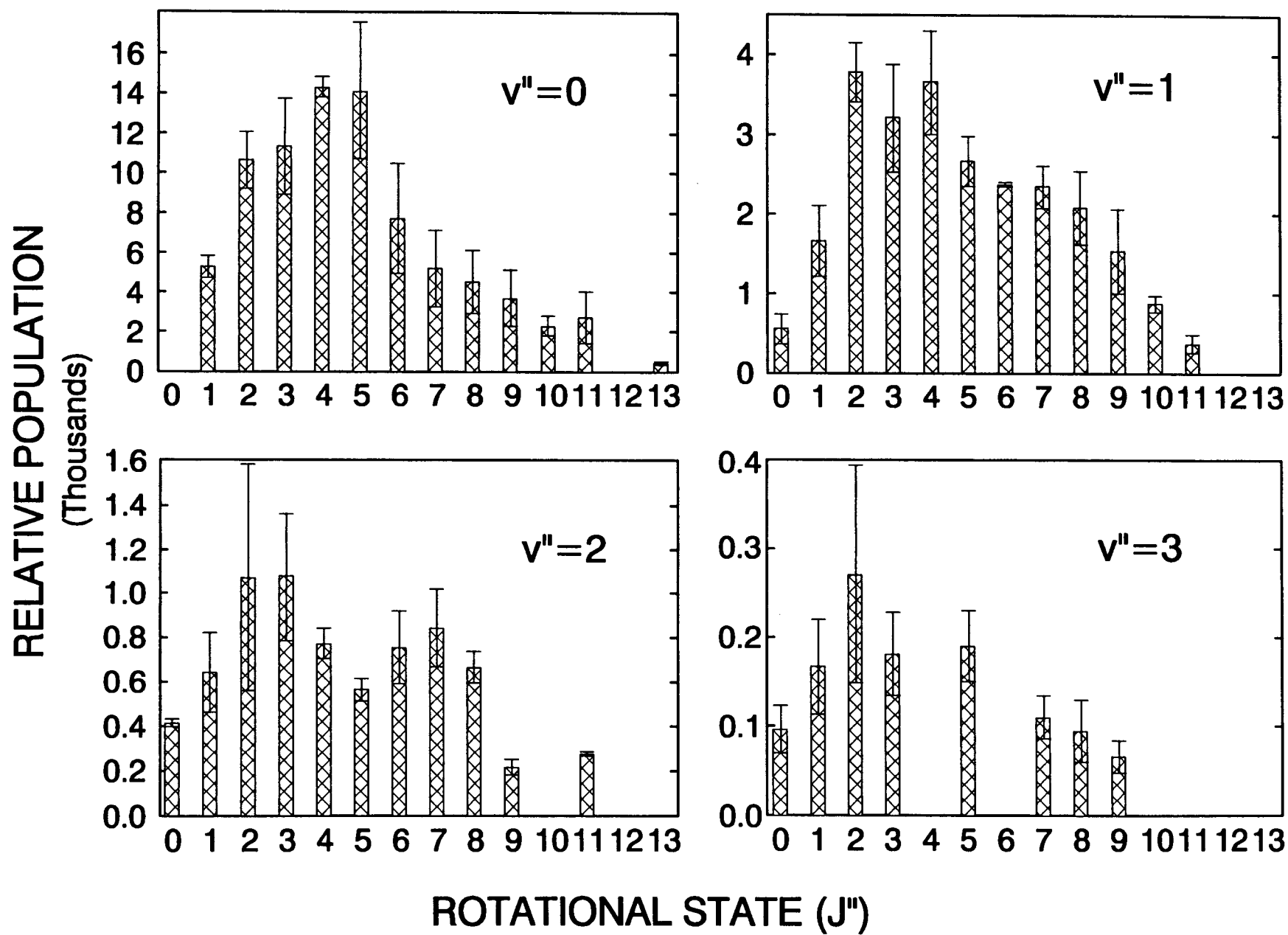


Fig. 6

RELATIVE POPULATION
(Thousands)

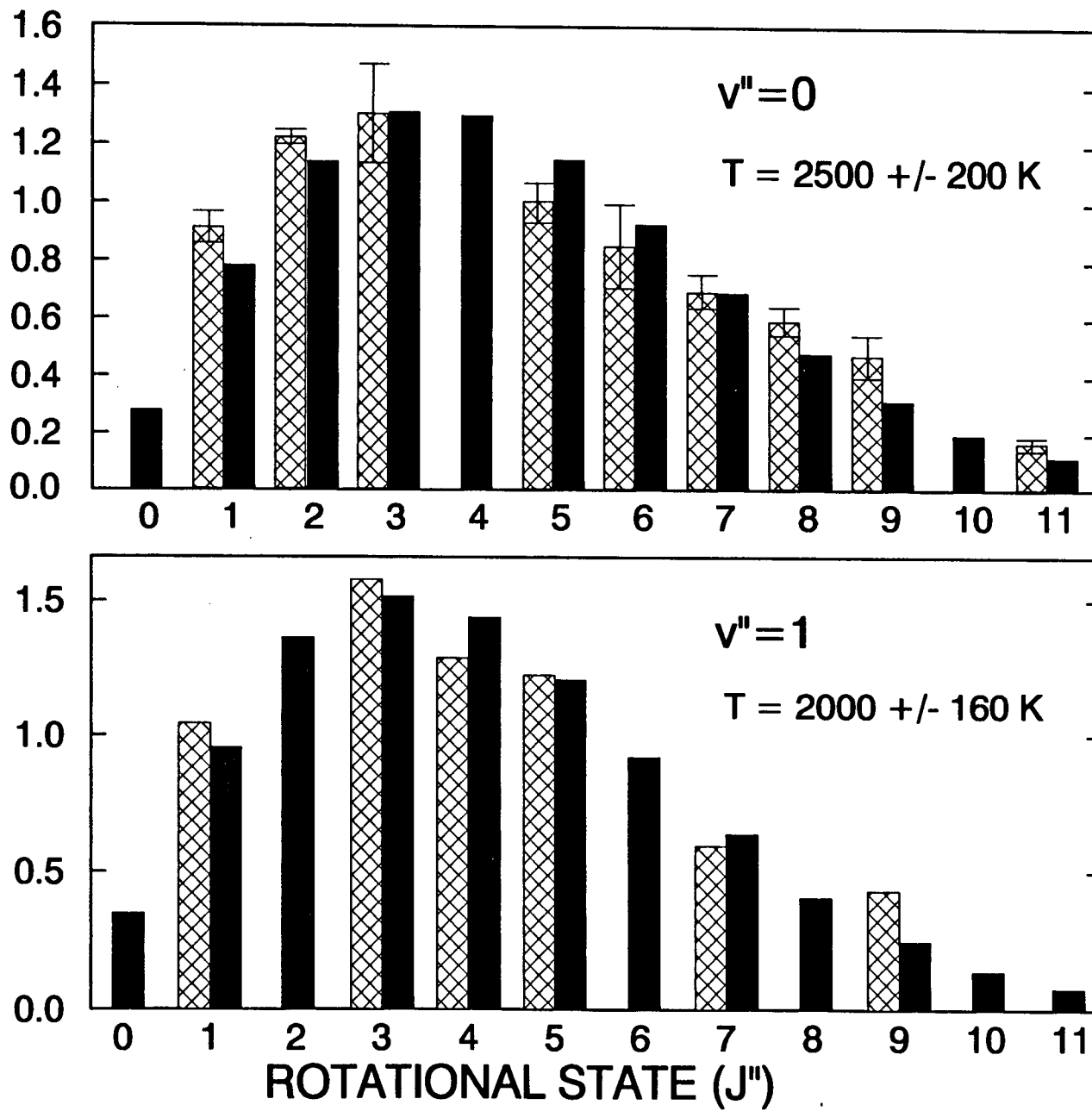


Fig. 7

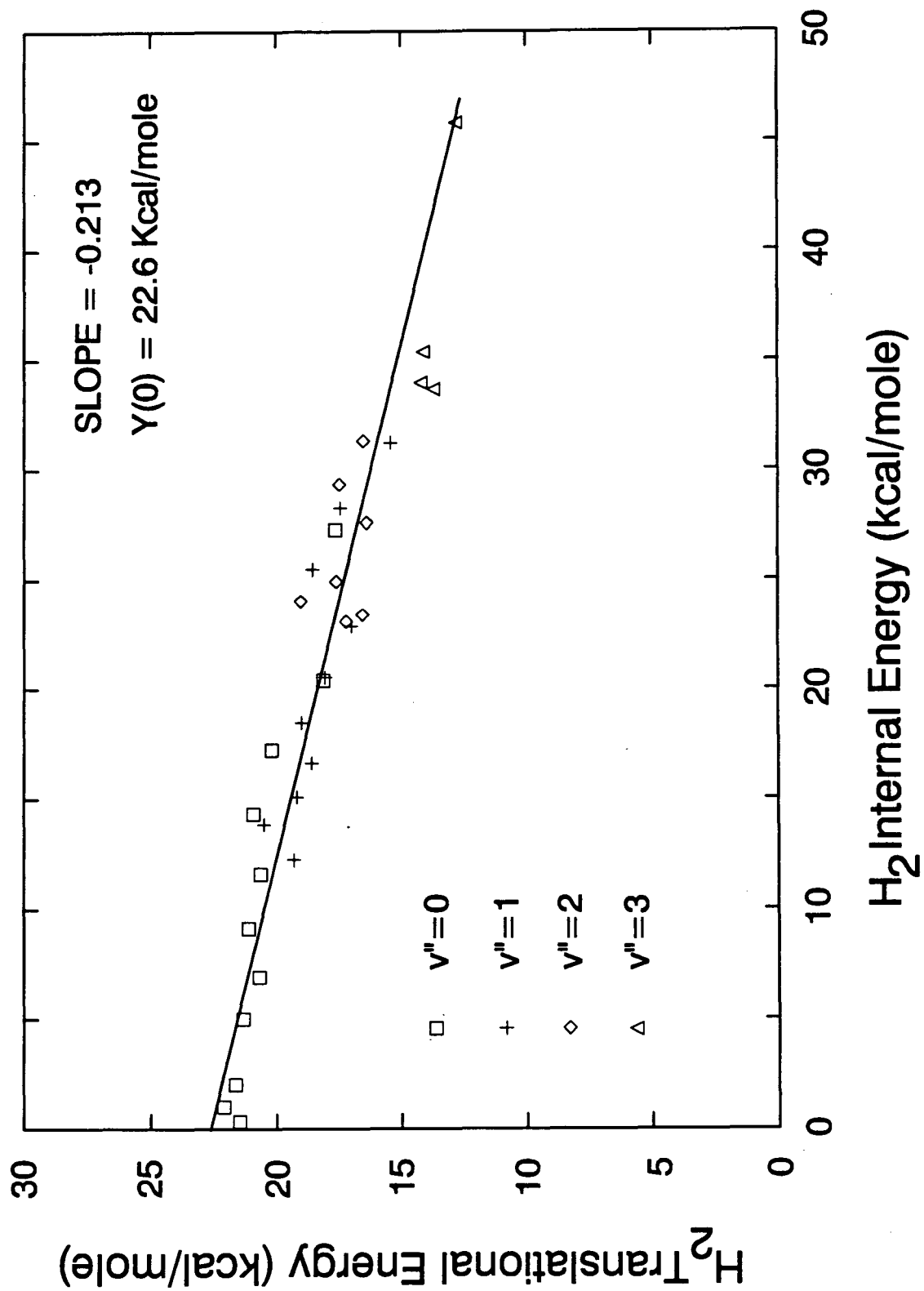
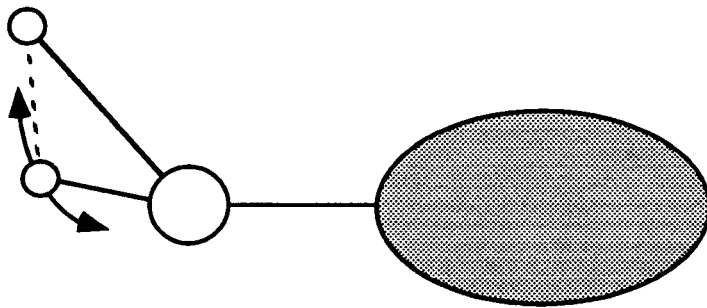


Fig. 8

POSITIVE CORRELATION



NEGATIVE CORRELATION

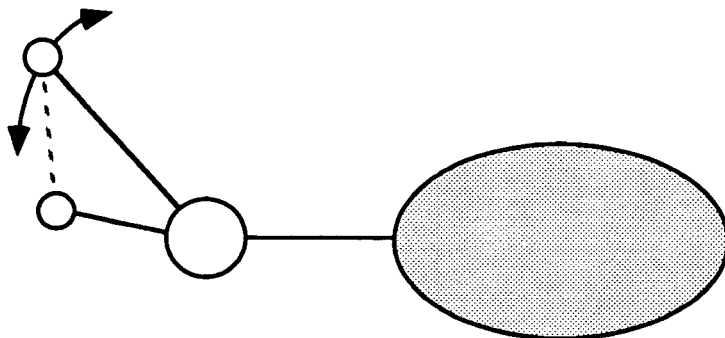


Fig. 9

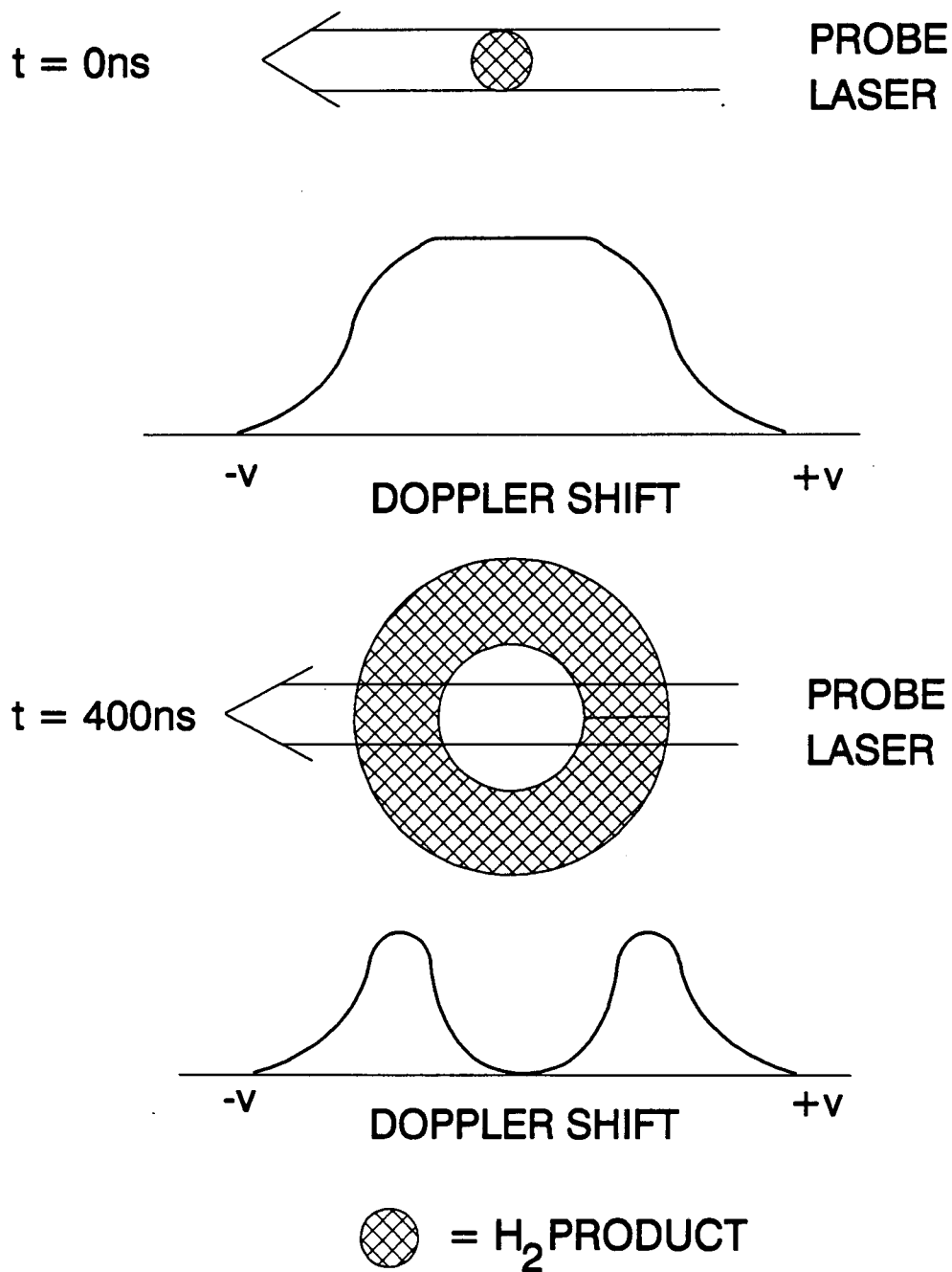


Fig. 10

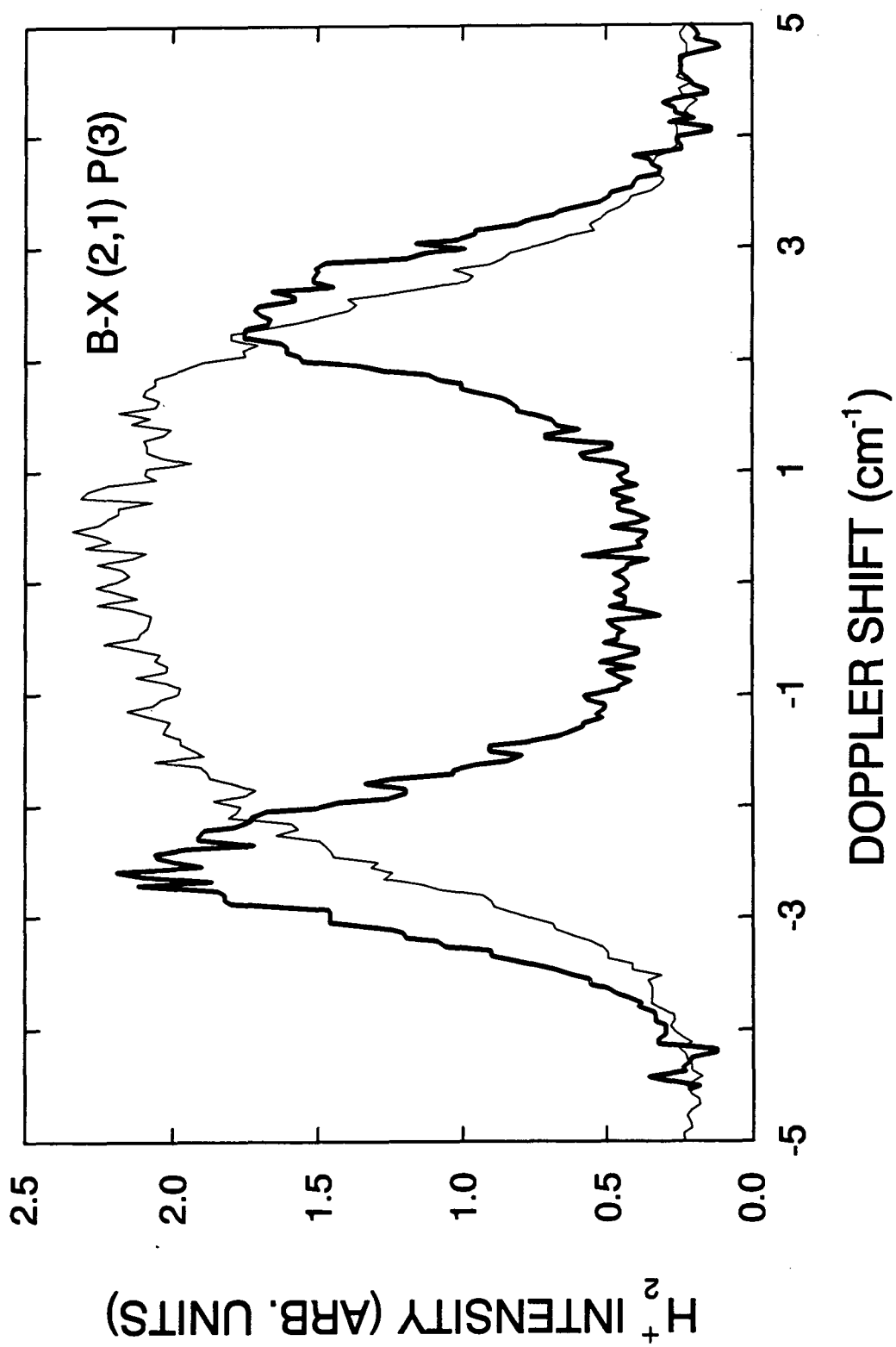


Fig. 11

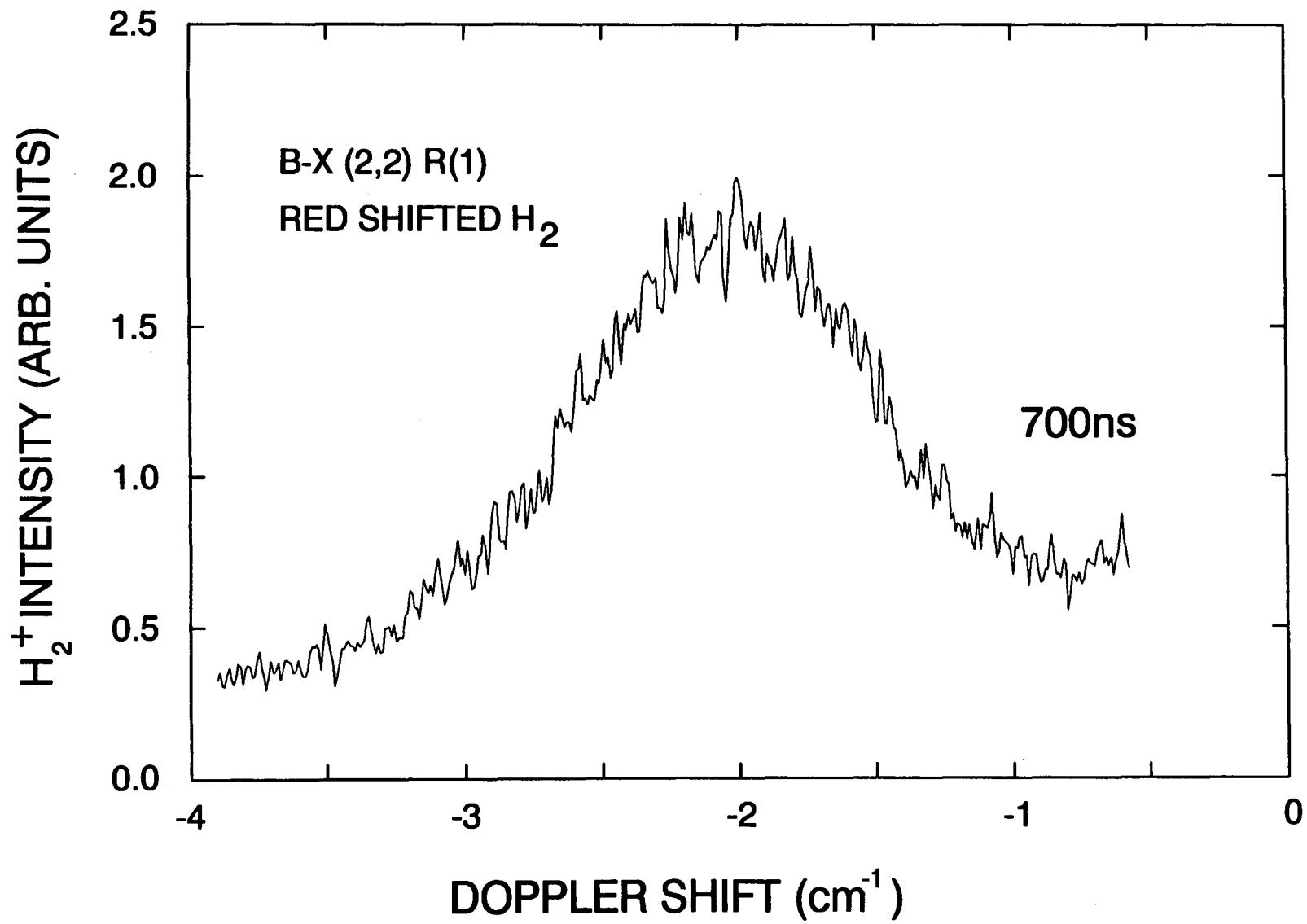


Fig. 12

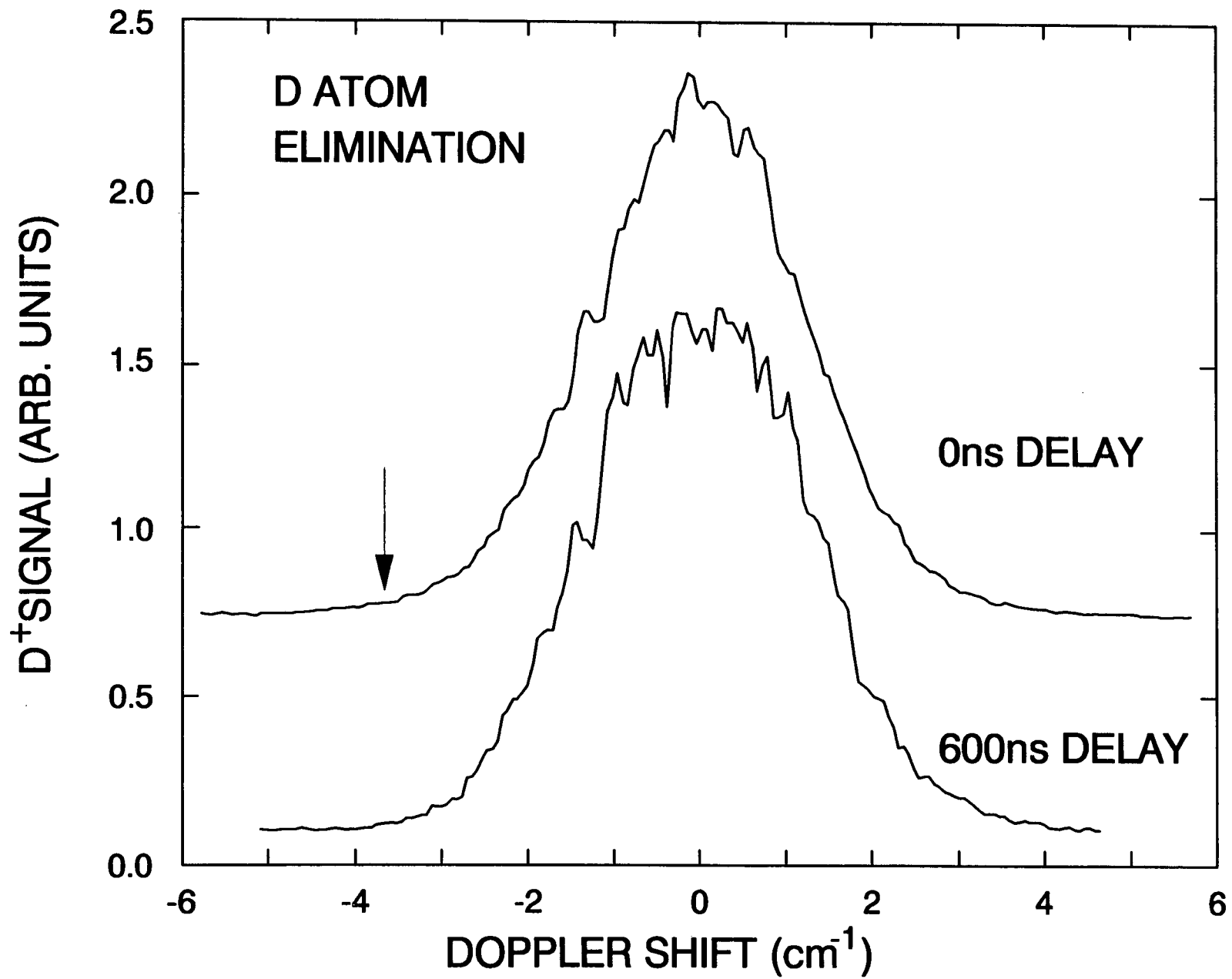


Fig. 13

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