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FRAGMENTATION BRANCHING RATIOS OF HIGHLY EXCITED HYDROCARBON MOLECULES C_nH AND THEIR CATIONS C_nH^+ ($n \leq 4$)

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Abstract

We have measured fragmentation branching ratios of neutral C_nH and C_nH^+ cations produced in high velocity (4.5 a.u) collisions between incident C_nH^+ cations and helium atoms. Electron capture gives rise to excited neutral species C_nH and electronic excitation to excited cations C_nH^+ . Thanks to a dedicated set-up, based on coincident detection of all fragments, the dissociation of the neutral and cationic parents were recorded separately and in a complete way. For the fragmentation of C_nH , the H-loss channel is found to be dominant, as already observed by other authors. By contrast, the H-loss and C-loss channels equally dominate the two-fragment break up of C_nH^+ species. For these cations, we provide the first fragmentation data ($n > 2$). Results are also discussed in the context of astrochemistry.

I-INTRODUCTION

Hydrocarbon radicals C_nH have been observed in a number of environments in the interstellar medium : diffuse clouds¹, cold dense molecular clouds and star forming regions², photodissociation regions^{3,4} and circumstellar envelopes⁵. Mostly neutral species have been detected, with the exception of recently discovered C_nH^- anions⁶. However, cations are included in astrochemical models⁷. Fragmentation of these species under UV photons³, cosmic rays⁸ and dissociative recombination⁹ will play a major role in the ongoing chemistry. For all of these processes, parent molecules are formed in electronically excited states before they decay into daughter fragments.

To our knowledge, all measurements concerning the fragmentation of electronically excited $C_nH^{(+)}$ species have been performed with impinging fast or slow electrons. In measurements based on crossed C_nH^+ ($n = 1,2$) ions with fast electron beams, some charged fragment production cross sections have been reported over a broad energy domain¹⁰⁻¹². These cross sections result generally, with some exceptions¹², from the contribution of both the dissociative excitation and ionization of the parent. Mass spectra of charged fragments following impact of 70 eV electrons on C_4H ¹³ and C_5H ¹⁴ have also been reported. They result from the sum of single and double ionization. Recent measurements of dissociative recombination branching ratios have been conducted at storage rings on C_2H^+ ¹⁵, C_3H^+ ¹⁶ and C_4H^+ ¹⁷. Contrary to previous measurements, the process of dissociative recombination is well isolated in these experiments.

From the theoretical side, studies concerning the structure of neutral^{18,19} and cationic²⁰ hydrocarbon molecules have been conducted. In recent works, electronic excited states above the dissociation threshold have been computed for C_nH ($n = 4-6$)^{21,22} and C_5H^+ ²³. In the case of C_4H , energies of all dissociation reactions were calculated using multireference MRSCF calculations with ZPE (zero point energy) corrections²¹. However, this type of information is relatively rare in the literature. Indeed, dissociation energies need consistent calculations and are very sensitive to the level of the quantum calculations. With that respect, information brought by branching ratio measurements, often connected to the values of dissociation energies, may be very useful.

In this work, the excitation and fragmentation of small C_nH^+ molecules ($n = 1-4$) colliding with helium atoms at high velocity ($v = 4.5$ atomic units) has been investigated experimentally. In this fast collision ($\sim 10^{-16}$ s), the excitation is exclusively of electronic origin. Cross sections of various electronic processes (excitation, ionisation, electron capture)

have been measured and the subsequent fragmentation totally recorded. The experimental technique, similar to the one used for the study of small carbon clusters²⁴, has been adapted to the detection of light (H) and hydrogenated (C_pH) fragments. With appropriate modifications, we could extract results at the same level of precision than previously done, *i.e* measurements of all individual branching ratios associated with each dissociative process. In this paper, only the fragmentation of neutral and singly charged molecules will be presented. Apart from its own interest, this experimental study is expected to bring valuable information to be used in an astrochemical context, as discussed later. The plan of the paper is the following. In section II, we describe the experimental method. Results of branching ratios are presented and discussed in section III. In section IV they are discussed in the context of astrochemistry.

II-EXPERIMENTS

The experiments have been performed at the Tandem accelerator in Orsay. The set-up is composed of three parts: the source of C_nH⁻ anions, the C_nH⁺ production system and the AGAT (AGrégat-ATome) set-up where the collision is performed and analysed.

II-1 The anion source

The anions were produced by a sputtering source: a 20 keV Cs⁺ beam impinged onto a rod made of 90% of coronene (C₂₄H₁₂) and 10% of Ag. Measured intensities of the anions at the exit of the source showed that mostly carbon clusters are formed. A small proportion of C_nH⁻ molecules (~10%) and a smaller proportion of C_nH₂⁻ species (~1%) were also produced.

The fact that we obtained a low hydrogenation of the molecules is due to the temperature of the source. The latter could be estimated, as in the case of sputtering from a graphite rod²⁵, to 3500 K. Beams of C_nH⁻ are contaminated by ¹³C¹²C_{n-1} clusters, of equal mass. Since intensities of C_n⁻ peaks were much larger than those of C_nH⁻, the contamination of these latter by ¹³C¹²C_{n-1} beams was important. Taking into account the 1% natural abundance of ¹³C with respect to ¹²C, we found that this contamination varied between 15% and 50% depending on n. After subtraction of this contamination, we get C_nH⁻ intensities around 5 nA, 6 nA, 200 pA and 900 pA for n = 1,2,3,4 respectively. The fact that even n species were found more abundant is in agreement with measurements from other authors¹⁹. It is explained by a higher electron affinity of the even n species.

II-2 Stripping and selection of C_nH^+ species

Mass selected C_nH^- molecules were injected at 0.2MeV in the Tandem accelerator. They were accelerated up to the high voltage terminal where they passed through a low pressure N_2 gas cell. The C_nH^+ cations, formed by the loss of two electrons in the gas cell, were accelerated again in the second part of the accelerator, magnetically selected and sent towards the experimental set-up AGAT. The high voltage was adjusted for each value of n in order to have beams with the same velocity of 4.5 atomic units. At this velocity, the flight time between the terminal and the AGAT set-up is $2\mu s$.

The C_nH^+ molecules possessed some internal energy, coming from the temperature of the source and the stripping process. The thermal internal energy of C_nH^- was calculated within the canonical ensemble for $T=3500\text{ K}$ ²⁶. In the stripping process, the energy deposit comes from the fact that ejection of electrons from various outer valence shells of C_nH^- can occur. As to ejection of inner valence electrons, it leads to fragmentation of the molecule. In the case of carbon clusters, we estimated the energy deposit due to outer valence electron ionization to $1.5\pm 0.5\text{ eV}$ whatever n , using photoemission spectra of C_n ²⁵. For C_nH species, due to the lack of such quantities, we assumed the same energy deposit. Altogether we found mean internal energies of C_nH^+ of the order of 2 eV for $n = 1$, 3 eV for $n = 2-3$ and 4 eV for $n = 4$ with standard deviations of 1eV for $n = 1$ and 2 eV for $n = 2-4$.

With such internal energies E_i , various isomers of C_nH^+ are populated. In a first approximation, we used the Boltzmann factor, $\exp(-\Delta E/E_i)$, for estimating the population of isomers placed at ΔE above the ground state of C_nH^+ . For C_2H^+ , we used MCSCF calculations of the five lowest excited states, all below 2 eV²⁷, and found that 48% of the ions were in a linear geometry and 52% in a quasi-linear (bent) geometry. For C_3H^+ , on the basis of CASSCF and CCSD(T) calculations performed on the two lowest isomers²⁸, we found that 57% of the ions should be in the linear ground state and 43% in the cyclic (C_3 -ring) isomer situated 0.8 eV above the ground state. For C_4H^+ , only information on the two lowest energy isomers is available¹⁷. According to these authors, the energy of the linear form is greater than that of the bi-cyclic isomer by 8 meV. An equal probability of the linear and bi-cyclic isomer is then deduced.

II-3 The AGAT set-up

The AGAT set-up has already been described²⁵. This set-up is made of three parts: the collision chamber, the fragment's electrostatic deflector and the detection chamber. In the collision chamber, an effusive helium jet with a known profile is operating under single collision conditions and absolute cross sections are derived as explained previously²⁹. The electrostatic deflector is made of two parallel plates, 15 cm long and 3 cm apart. The electric field was set to 25kV/cm in order to separate fragments of various charge over mass ratios (q/m).

II-4 Detection of fragments and resolution of the fragmentation channels

In the detection chamber, seven silicon solid-state detectors operating in coincidence intercepted all $C_nH_p^{(q+)}$ fragments ($n = 0-4$, $p = 0-1$, $q = 0-3$), see Table I. Three detectors (numbered 1, 2, 3) were "home made" silicon surface barrier detectors mounted at the Institut de Physique Nucléaire in Orsay. They were made of N-type epitaxial silicon of large resistivity ($7 \text{ k}\Omega\cdot\text{cm}$) grown on N^+ type silicon substrate with a gold entrance layer (40nm) for the p^+N junction. Their areas were 36mm^2 for detector 1 and 72mm^2 for detectors 2 and 3. We used fast current preamplifiers designed also at the IPNO. Three detectors (numbered 4,5,6) were commercial ion-implanted silicon detectors of surface area 600mm^2 equipped with commercial charge preamplifiers. The detector for H^+ fragments was a commercial ion-implanted detector of large dimensions (50 mm x 50 mm) made of 4 independent parallel bands equipped with four commercial charge preamplifiers.

In the standard utilization of silicon solid state detectors, the charge delivered by the detector provides the kinetic energy of the fragment *i.e* its mass for fragments of constant velocity³⁰. With this property and the (q/m) analysis, the charge of each fragment was recorded. Specific electronic processes could then be recorded separately. Indeed, charge transfer gives rise to events with only neutral fragments, electronic excitation to events with only one singly charged fragment, single ionization to events with two singly charged fragments or one doubly charged fragment and so on. It has to be noted that non dissociative processes were also recorded, with the exception of the non-dissociative electronic excitation. In the latter case, the excited ion remains in the incident beam and cannot be separated.

Recently we showed that analysing the shape of transient currents allowed resolving, in number and in mass, pile up of many fragments in a detector ³¹. This technique, applied successfully to the fragmentation of carbon clusters ²⁹, does not work for H fragments. The reason is that the H fragment, of much lower energy than other fragments, does not contribute significantly to the signal. Therefore, a CH molecular fragment or two atomic C, H fragments give rise to current signals having almost the same shape within the present level of detector noise. Illustration of this is given in figure 1, which shows a two-dimensional representation of signals delivered by the neutral detector. The x axis represents the integral of the current signal (proportional to the neutral mass), the y axis its peak value ³¹. Each point refers to an event. In this figure, brackets indicate summed signals: {C/H} means two C/H fragments or one CH fragment; C/{C/H} means three C/C/H fragments or two C/CH fragments

In order to overcome this problem, we added to the shape analysis the so-called grid method. The grid method was introduced a long time ago in order to resolve the fragmentation of fast H₃ molecules ³². It is still used in dissociative recombination studies carried out at storage rings³³. The method consists in placing in front of the detector a grid of known transmission and to record mass spectra with and without the grid. The problem is solvable if the number of recorded intensities is equal to or exceeds the number of unknown branching ratios. We applied this technique and placed a grid (hole sizes and wire diameters 20μ) in front of the neutral detector where pile up is involved. The transmission of the grid, t, was measured by recording the attenuation of intensity from a radioactive alpha source (²⁴¹Am) and also by recording the fragmentation pattern of C₄ clusters with and without grid. Both methods agreed and we could extract a precise determination of t = 0.269 (± 0.004). As mentioned before, mass spectra and current shapes obtained with and without grid are used for extraction of branching ratios. As an illustration of the method, the dissociation of C₂H is treated in annex.

III-RESULTS AND DISCUSSION

III-1-Cross sections of electronic processes

Cross sections of electronic processes measured in the collision of C_nH⁺ projectiles with helium are reported in figure 2 as a function of n. The ionization process always dominates, followed by dissociative electronic excitation, and electron capture. The relative importance of these processes is as expected in this velocity range ^{34,35}.

The electronic excitation and ionization cross sections exhibit the same evolution with n . This trend is well reproduced by the Independent Atom and Electron (IAE) model^{30,36} in the case of ionization. In the IAE calculations, shown by a broken line, classical trajectory Monte Carlo (CTMC) impact parameter probabilities $P(b)$ were used for valence ionization in C and C^+ as in previous works^{25,37}. Calculations within the SCA (semiclassical approximation) model³⁸ were performed for $P(b)$ shapes due to inner shell ionization in C and C^+ and to ionization of H. Normalizations of $P(b)$ distributions were performed on measured ionization cross sections in C^+ ³⁵ and H³⁹.

Electron capture cross sections evolve in a smoother way with n . The slight increase as a function of n was also observed with incident carbon clusters²⁴. It was tentatively interpreted by an increase of the cluster electronic density of states with its size.

III-2-Relaxation of C_nH species

Tables II, III, IV and V present measured branching ratios for all dissociative and non dissociative relaxation channels of CH, C_2H , C_3H and C_4H respectively. Dissociation energies are also reported. In column 3, they were derived by using binding energies of H and C calculated by Pan et al.¹⁹. In this work, authors used DFT formalism and found ground states to have a quasi-linear geometry. Their dissociation energies were obtained without consideration of symmetry. With the use of symmetries reported in Table VI, we determined - and commented through footnotes in Tables II-V- cases where dissociation from the electronic ground state of the parent to electronic ground states of fragments is forbidden according to correlation rules⁴⁰. When dissociation was forbidden, we considered first electronic excited states of the parent and/or fragments.

a-Number of emitted fragments N_f and internal energy distribution

Figure 3 shows, for all species, probabilities of dissociation into a given number of fragments $P(N_f)$. These have been obtained by summing branching ratios of Table II, III, IV and V for each N_f . We observe, apart from CH, similar probability distributions peaked at two-fragment break up ($N_f=2$).

The number of emitted fragments reflects the internal energy of parent molecules. Indeed, as seen from dissociation energies reported in Tables III to V, a two, three, four, fragments break-up requires in average 6 eV, 12 eV, 19 eV respectively. These formation

energies are rather independent of the molecule size. In fact, as we have shown recently on carbon clusters, dissipation in fragments kinetic motion has to be taken into account to correctly connect internal energy and number of emitted fragments. We have:

$$P(N_f) = \frac{\int_{E_1}^{E_2} f(E) dE}{\int_0^{\infty} f(E) dE} \quad (1)$$

where $f(E)$ is the internal energy distribution and boundary E_1 and E_2 express as:

$$E_1(N_f) = E_{\text{diss_low}}(N_f) + (N_f - 2) * E_{\text{TVR}} \quad \text{for } N_f \geq 2 \quad (2)$$

$$E_2(N_f) = E_1(N_f + 1)$$

$$E_1(1) = 0$$

In (2), $E_{\text{diss_low}}(N_f)$ is the lowest dissociation energy for a given number of fragments (from Tables II to V), and E_{TVR} is the average energy dissipated in fragment's kinetic motion (translation, vibration, rotation). We assumed 1eV for E_{TVR} as in the case of C_n ^{41,42}.

In figure 4 is shown, for C_4H , the internal energy distribution extracted from data using a step function or the analytical form $E^{a_1} \exp(-a_2(E-a_3)^{a_4})$ for $f(E)$. This last form, depending on four parameters a_1 - a_4 , is known to reproduce the energy distribution due to electron capture in any velocity range²⁹. We note that the internal energy distribution is very broad, extending from low energies under the dissociation limit up to large energies, well above the ionization potential of C_4H . We obtained a similar result in the case of carbon clusters and explained this broad distribution by the electron capture process itself which, in high velocity ion-atom collisions, tends to populate an extended range of excited states^{24,34}. Mean internal energies and standard deviations of $f(E)$, reported in parentheses, were found respectively equal to 10 eV (4 eV), 9 eV (6 eV) and 10.5 eV (5 eV) for C_2H , C_3H and C_4H .

b-Branching Ratios for fixed N_f

The number of emitted fragments reflects the energy deposited in the system and may be very different from an experiment to another. Branching ratios for a given N_f value are less sensitive to the energy distribution and reflects more intrinsic properties of the parent molecule. In figure 5 are reported branching ratios normalised to 100% for each N_f value for C_2H , C_3H and C_4H .

For two-fragment break up ($N_f=2$), we see that in all cases the H emission is the most probable dissociation channel. This is not surprising since this channel is always predicted to

have the lowest dissociation energy (Tables III-V). The second most populated channel is the C-loss channel. This is in agreement with dissociation energies, except for C₄H. For this molecule though, the ordering of the channels is not well established. We also note that the CH-loss is always a small channel.

III-3 Fragmentation of C_nH⁺ species

In Tables VII, VIII, IX and X are reported measured branching ratios for all dissociative channels of CH⁺, C₂H⁺, C₃H⁺ and C₄H⁺ respectively. Dissociation energies are also reported. These were calculated using (3):

$$E_{\text{diss}}^+ = E_{\text{diss}} - \text{IP}_{\text{parent}} + \text{IP}_{\text{fragment}} \quad (3)$$

where E_{diss} is the dissociation energy of the corresponding neutral channel

$\text{IP}_{\text{parent}}$ is the adiabatic ionization potential of the neutral parent C_nH

$\text{IP}_{\text{fragment}}$ is the adiabatic ionization potential of the neutral fragment that is singly charged in the exit channel.

E_{diss} were taken from Tables II-V (third column). IP values were taken from the literature and are reported in Table VI. As in the case of neutrals, we indicated in Tables VII to X cases for which ground state to ground state dissociation is not permitted by correlation rules⁴⁰. Symmetries of the cationic states are given in Table VI. One notes that there are many more forbidden transitions with cations than with neutrals. This is often due to non conservation of the spin.

a-Number of emitted fragments N_f and internal energy distribution

Figure 6 presents, for each C_nH⁺ parent, probabilities of dissociation into a given number of fragments P(N_f). As only the dissociative part of excitation is measured in the experiment, measurements begin at N_f= 2. We used the method described in III-2 in order to extract internal energies of C_nH⁺ cations. We found 18 eV (9 eV), 18 eV (7 eV) and 18 eV (7 eV) for mean internal energies (and standard deviations of f(E)) for C₂H⁺, C₃H⁺ and C₄H⁺ respectively. These values are very close to those obtained with C_n⁺ clusters⁴³.

b-Branching Ratios for fixed N_f

In figure 7 are reported dissociation branching ratios, normalised to 100% for each N_f value, for CH^+ , C_2H^+ , C_3H^+ and C_4H^+ .

For two-fragment break up, we remark that the H-loss and C-loss channels are almost equally probable. This is not in accordance with dissociation energies, which are always predicted lower for the H-loss channel. For instance, the C_2^+/H and CH^+/C dissociation energies differ by as much as 2 eV although they have equal branching ratios. The relatively low branching ratio observed for C_2^+/H could come from the forbidden ground-state to ground state dissociation (see footnote in Table VIII). It could also be due to dynamical effects - i.e non statistical- since collisional excitation is more probable on the C-C bond than on the C-H bond according to the number of present electrons.

Another difficulty for interpreting these results is the poor confidence in dissociation energies. The IP used in (3) are not well known. For instance, recent determinations of IP in C_3 ⁴⁴ and C_4 ⁴⁵ are well below all previous values, in particular measurements of Ramanathan et al.⁴⁶.

IV- ASTROCHEMICAL CONTEXT

IV-1 Branching ratios in a statistical approach

In astrochemical models, reactions rates – i.e. total cross sections - can be, in most cases, safely estimated. Calculations exist and can rely on numerous experimental measurements. On the contrary, branching ratios of different exit channels are much more difficult to obtain. From the experimental point of view, it is because neutral fragments detection is necessary. From the theoretical point of view, it is because *ab initio* dynamical calculations are very difficult and time consuming for systems composed of more than a few atoms⁴⁷.

A “simple” theoretical way to predict branching ratios is to use a statistical approach⁴⁸. In such approach, the dynamics vanishes and the fragmentation is governed only by the amount of deposited energy. Statistical branching ratios can be derived from more or less approximated calculations such as Rice–Ramsperger–Kassel–Marcus (RRKM)⁴⁹, Weisskopf⁵⁰, Phase Space Theory (PST)⁵¹, Microcanonical Metropolis Monte Carlo (MMMC)²⁹. Moreover,

if the fragmentation is statistical, measurements of branching ratios with a given process can be extended to other processes.

For the fragmentation of small carbon clusters C_n ($n=5-9$) in high velocity collisions, the statistical approach has been shown to provide satisfactory results^{29,42}. The same conclusion was reached on smaller systems studied with photons⁵². In C_4 , fragmentation branching ratios measured in high velocity collisions³⁴ are close to those obtained in dissociative recombination⁵³. As to the fragmentation of cationic carbon clusters C_n^+ ($n=4-10$), similar results were found for measured branching ratios in photodissociation and in high velocity collisions⁵⁴. All these results tend to prove that, at least for these carbon species, the statistical approach is valid.

IV-2 Comparison between fragmentation branching ratios of C_nH species following electron capture in high velocity collisions and dissociative recombination

In Table XI are presented dissociative recombination branching ratios of C_2H^+ ¹⁵, C_3H^+ ¹⁶ and C_4H^+ ¹⁷ measured at storage rings. Only two-fragment channels are reported. Indeed, none of the three-fragment channels were observed in C_3H and C_4H fragmentation, due to detection limitation. For the same reason, some channels were not resolved in the fragmentation of C_3H and C_4H and are presented summed in the Table XI. Also reported in Table XI are two-fragment branching ratios that we obtained in this work.

One notices a general good agreement in the ordering of branching ratios obtained in dissociative recombination and high velocity collision experiments. The channel C_n/H is found slightly more populated in high velocity collisions than in dissociative recombination for C_2H and C_4H . This can be due to the sensitivity of branching ratios to the shape of the internal energy distribution. Indeed, in dissociative recombination, narrow energy distributions -peaked around the neutral parent's IP- vary from one molecule to another while a broad and constant energy distribution is involved in high velocity collisions. The good agreement between branching ratios obtained in dissociative recombination and high velocity collisions could be explained by a statistical fragmentation of C_nH species.

CONCLUSION

We have presented results concerning the fragmentation of neutral and cationic $C_nH^{(+)}$ hydrocarbons ($n \leq 4$) electronically excited in high velocity collisions with helium atoms. In

the case of neutrals, the H-loss channel was found to be the most probable, followed by the C-loss channel and the CH-loss channel. This ordering follows reasonably well dissociation energies and was already obtained by other authors. In contrast, the H-loss and C-loss channels both dominate two-fragment dissociation in case of C_nH^+ cations. This result is unexpected since dissociation energies are always predicted lower for H-loss as compared to C-loss. It could be due to dynamical (non statistical) effects. With that respect, it would be interesting to have results obtained in a different experimental situation, for instance, following excitation by another projectile or photon.

We discussed in the paper the astrochemical interest of these measurements. Branching Ratios for two-fragment break-up of C_nH molecules were compared to results of dissociative recombination obtained by other authors. We observed a general good agreement which could be explained by a statistical fragmentation behaviour in these species, as observed in carbon clusters.

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Detector	1	2	3	4	5	6	7
Fragments	H C CH C ₂ C ₂ H C ₃ C ₃ H C ₄ C ₄ H	C ₄ H ⁺ C ₄ ⁺ C ₃ H ⁺ C ₃ ⁺	C ₂ H ⁺ C ₂ ⁺ C ₄ H ⁺⁺ C ₄ ⁺⁺ C ₃ H ⁺⁺ C ₃ ⁺⁺	CH ⁺ C ⁺	CH ⁺⁺ C ⁺⁺	C ⁺⁺⁺	H ⁺

Table I:

Fragments collected by detectors in the experiment

Channel	BR(error) (%)	Ediss (eV) ¹	Ediss (eV) ²
CH	64(4)		
C/H	36(4)	3.65	3.45(0.01)

Table II:

Measured branching ratios (BR) of relaxation channels of CH molecules. In last columns, are reported channel dissociation energies (E_{diss}).

¹DFT calculations¹⁹

²experimental value from⁵⁵

Channel	BR(error) (%)	Ediss (eV) ¹	Ediss (eV) ²
C ₂ H	16.8(8)		
C ₂ /H	51.6(3)	5.01	5.10
C/CH	11.9(5)	7.86	7.88
2C/H	19.7(3)	11.51	

Table III:

Measured branching ratios (BR) of relaxation channels of C₂H molecules. In last columns, are reported channel dissociation energies (E_{diss}).

¹using DFT calculations¹⁹

²experimental values from⁵⁶ and⁵⁷ for C₂/H and C/CH respectively

channel	BR(error)(%)	Ediss (eV) ¹	Ediss (eV) ²
C ₃ H	25(3)		
C ₃ /H ^a	40(7)	3.46	3.08 (4.00)
C ₂ H/C	21(4)	6.21	5.60 (5.66)
C ₂ /CH	1(2)	7.57	6.96 (7.03)
C ₂ /C/H	6.5(3.5)	11.22	
2C/CH	4(2)	14.07	
3C/H	2.5(2.5)	17.72	

Table IV:

Measured branching ratios (BR) of relaxation channels of C₃H molecules. In last columns, are reported channel dissociation energies (E_{diss}).

¹using DFT calculations¹⁹

²calculations⁵⁸ for the linear l-C₃H isomer (and the cyclic c-C₃H one in parentheses)

a: dissociation towards ground state products possible for the linear isomer l-C₃H, when bent⁵⁸.

channel	BR(error)(%)	Ediss (eV) ¹	Ediss (eV) ²	Ediss (eV) ³
C ₄ H	11.8(3)			
C ₄ /H	29.9(3)	4.87	5.71	4.52
C ₃ H/C	13.3(5.5)	6.90	6.33	7.25
C ₂ /C ₂ H	7.9(2.5)	6.61	7.12	7.24
C ₃ /CH ^a	≤3	6.71	5.88	
C ₃ /C/H ^b	17.1(6.5)	10.36		
2C ₂ /H	11.1(3.3)	11.62		
C ₂ /C/CH	1.5(2)	14.47		
C ₂ H/2C	≤2	13.11		
C ₂ /2C/H	6.9(2.8)	18.12		
3C/CH	≤0.5	20.97		
4C/H	0.5(0.5)	24.62		

Table V:

Measured branching ratios (BR) of relaxation channels of C₄H molecules. In last columns, are reported channel dissociation energies (E_{diss}).

¹using DFT calculations¹⁹

²MCSCF multireference calculations for linear C₄H including ZPE (zero point energy) corrections²¹

³extracted from³³, by using (second and third values) the IP of C₄H (see Table VI).

a,b: ground state to ground state dissociation is not permitted for these channels but dissociation from the electronically excited state of C₄H, ²Π (+0.02eV⁵⁹) is possible.

Species (and g.s symmetry)	IP(error) (eV)		Cation (and g.s. symmetry)
H(2S_g)	13.6(<0.01)	60	H ⁺
C(3P_g)	11.26(<<0.01)	60	C ⁺ (2P_u)
CH($X^2\Pi, C_{\infty v}$)	10.64(0.01)*	55	CH ⁺ ($X^1\Sigma^+, C_{\infty v}$)
C ₂ ($X^1\Sigma_g^+, D_{\infty h}$)	11.41(0.3)**	61	C ₂ ⁺ ($X^4\Sigma_g^-, D_{\infty h}$)
C ₂ H($X^2\Sigma^+, C_{\infty v}$)	11.61(0.07)	62	C ₂ H ⁺ ($X^3\Pi, C_{\infty v}$)
C ₃ ($X^1\Sigma_g^+, D_{\infty h}$)	11.61(0.07)**	44	C ₃ ⁺ (X^2B_2, C_{2v})
c-C ₃ H(X^2B_2, C_{2v})	9.06	28	C ₃ H ⁺ ($X^1\Sigma^+, C_{\infty v}$)
l-C ₃ H($^2\Pi, C_{\infty v}$)(+0.04eV) ²⁸	8.36	28	
c-C ₄ (X^1A_g, D_{2h})	10.9(0.2)**	45	C ₄ ⁺ (X^2B_{1u}, D_{2h})
l-C ₄ ($^3\Sigma_g^-, D_{\infty h}$)(+0.04eV) ⁶³	11.0(0.2)**	45	
C ₄ H($X^2\Sigma^+, C_{\infty v}$)	12	17	C ₄ H ⁺ ($X^3\Sigma^-, C_{\infty v}$) ^a

Table VI:

Ground states and symmetries of neutral (first column) and cationic (third column) species relevant to this study. For C₃H and C₄, we reported also linear isomers which are very close in energy from cyclic ground states. In column 2 are reported ionization potentials used in equation (1); * : vertical IP; ** : adiabatic IP.

a: The C₄H⁺ molecular ion has been poorly studied and, to our knowledge, there is only one reference in which a $^1\Sigma^+$ state for its electronic ground state has been proposed⁶⁴. Ground state investigations have thus been performed using the Gaussian 2003 suite of programs⁶⁵. The calculations were performed at the B3LYP/cc-pTVZ and CCSD(T)/cc-pVTZ levels. Both were in agreement and predicted that the electronic ground state is in fact a $X^3\Sigma^-$ state.

Channel	BR(error) (%)	Ediss (eV)
C ⁺ /H	63.5(1)	4.27 ^a
C/H ^{+b}	36.5(1)	6.61

Table VII:

Measured branching ratios (BR) of dissociative channels of CH⁺ molecules. In column three are reported channel dissociation energies calculated with equation (1) using Table II and Table VI.

a: experimental value: 3.95 eV ($\pm 0.03\%$)⁶⁶

b: dissociation occurs from the electronic excited state $b^3\Sigma^-$ state of CH⁺ situated +4.5eV above the ground state¹¹.

Channel	BR(error) (%)	Ediss (eV)
C_2^+/H^a	16(2)	4.81
CH^+/C	14.5(2)	6.89
C_2/H^{+b}	2.6(1)	7.00
CH/C^+	2.4(2)	7.51
$C/H/C^+$	47.7(2)	11.16
$2C/H^+$	16.8(1)	13.50

Table VIII:

Measured branching ratios (BR) of dissociative channels of C_2H^+ molecules. In column three are reported channel dissociation energies calculated with equation (1) using Table III and Table VI.

a: ground state to ground state dissociation is not permitted for this channel; dissociation may occur from an electronic excited state of the parent ($^3\Sigma^-$ at +0.8eV²⁷) or lead to an excited state of C_2^+ ($^2\Pi_u$ at +0.8eV⁶⁷)

b: ground state to ground state dissociation is not permitted for this channel but dissociation may lead to an excited state of C_2 ($^3\Pi_u$ at +0.1eV⁶⁸)

Channel	BR(error) (%)	Ediss (eV)
C_2H^+/C	7.4(2)	8.76
C_3^+/H	10.5(1)	6.01
C_2H/C^+	4.7(1)	8.41
CH/C_2^{+a}	4.6(1)	9.91
C_2/CH^+	3.7(0.5)	9.14
C_3/H^+	0.8(0.1)	8.00
$C/CH/C^+$	13.6(2.5)	16.27
$C/H/C_2^+$	14.4(1)	13.56
$C_2/H/C^+$	9.2(1)	13.41
$C_2/C/H^{+b}$	4.3(1)	15.75
$2C/CH^+$	9.2(3)	15.65
$2C/H/C^+$	15.1(1.5)	21.28
$3C/H^{+c}$	2.5(1)	23.62

Table IX:

Measured branching ratios (BR) of dissociative channels of C_3H^+ molecules. In column three are reported channel dissociation energies calculated with equation (1) using Table IV and Table VI.

a: ground state to ground state dissociation is not permitted for this channel but dissociation leading to an excited state of C_2^+ ($^2\Pi_u$ at +0.8eV⁶⁷) is permitted

b: ground state to ground state dissociation is not permitted for this channel but dissociation leading to an excited state of C_2 ($^3\Pi_u$ at +0.1eV⁶⁸) is permitted

c: ground state to ground state dissociation is not permitted for this channel but dissociation leading to an excited state of C (for instance 1D_g at +1.2eV⁶⁹) is permitted

Channel	BR(error) (%)	Ediss (eV)
C/C ₃ H ⁺	9.2(1.3)	3.96
C ₄ ⁺ /H ^a	5.3(0.4)	3.77
C ₃ H/C ⁺	2.2(0.7)	6.16
C ₂ /C ₂ H ⁺ ^b	1.6(0.5)	6.21
C ₃ /CH ⁺ ^c	1.9(0.4)	5.35
C ₂ H/C ₂ ⁺	0.7(0.5)	6.01
C ₄ /H ⁺ ^d	0.3(0.1)	6.47
C ₃ ⁺ /CH	1.8(0.4)	6.32
C/H/C ₃ ⁺	15.1(1.2)	9.97
C ₃ /H/C ⁺ ^e	9.5(0.5)	9.62
C ₂ /H/C ₂ ⁺	7.0(1.1)	11.02
C ₂ H/C/C ⁺	2.9(0.8)	12.37
C ₂ /C/CH ⁺	3.6(0.5)	13.10
C ₃ /C/H ⁺	1.8(0.1)	11.96
2C/C ₂ H ⁺	0.7(0.7)	12.72
C/CH/C ₂ ⁺	0.7(0.7)	13.87
C ₂ /CH/C ⁺	2.3(0.5)	13.72
2C ₂ /H ⁺ ^f	0.9(0.1)	13.21
C ₂ /C/H/C ⁺	13.3(1.3)	17.37
2C/H/C ₂ ⁺	7.1(1.1)	17.52
2C/CH/C ⁺	2.1(0.7)	20.23
C ₂ /2C/H ⁺	1.7(0.2)	19.71
3C/CH ⁺	1.5(0.3)	19.61
3C/H/C ⁺	6.3(1)	25.24
4C/H ⁺	0.5(0.1)	27.58

Table X:

Measured branching ratios (BR) of dissociative channels of C₄H⁺ molecules. In column three are reported channel dissociation energies calculated with equation (1) using Table V and Table VI.

a: ground state to ground state dissociation is not permitted; dissociation towards excited C₄⁺ (⁴Σ_g⁻, +0.3eV⁷⁰) is possible.

b: ground state to ground state dissociation is not permitted; dissociation towards excited C₂ (³Π_u, +0.1eV⁶⁸) is possible.

c, e: ground state to ground state dissociation permitted towards cyclic isomer C₃ (³A₂[']) situated +0.8eV above the linear isomer⁵⁸.

d: ground state to ground state dissociation permitted towards l-C₄

f: ground state to ground state dissociation is not permitted; dissociation towards excited C₂ (³Σ_g⁻, +0.9eV⁶⁸) is possible.

Channel	DR	HV collision
C ₂ /H	52(4)	81(5)
C/CH	48(5)	19(8)
C ₃ /H	66(1.5)	65(11)
C ₂ H/C	} $\Sigma=34(1.7)$	33(6)
C ₂ /CH		≤ 2
C ₄ /H	44(1.2)	55(6)
C/C ₃ H	} $\Sigma=28(2)$	25(10)
C ₃ /CH		≤ 5
C ₂ /C ₂ H	28(2)	15(5)

Table XI:

Comparison, for two-fragment break-up, between measured branching ratios in dissociative recombination experiments (DR, adapted from ¹⁵, ¹⁶ and ¹⁷) and in high velocity collisions by electron capture (HV collision, this work); branching ratios and error bars (in parentheses) are given in %.

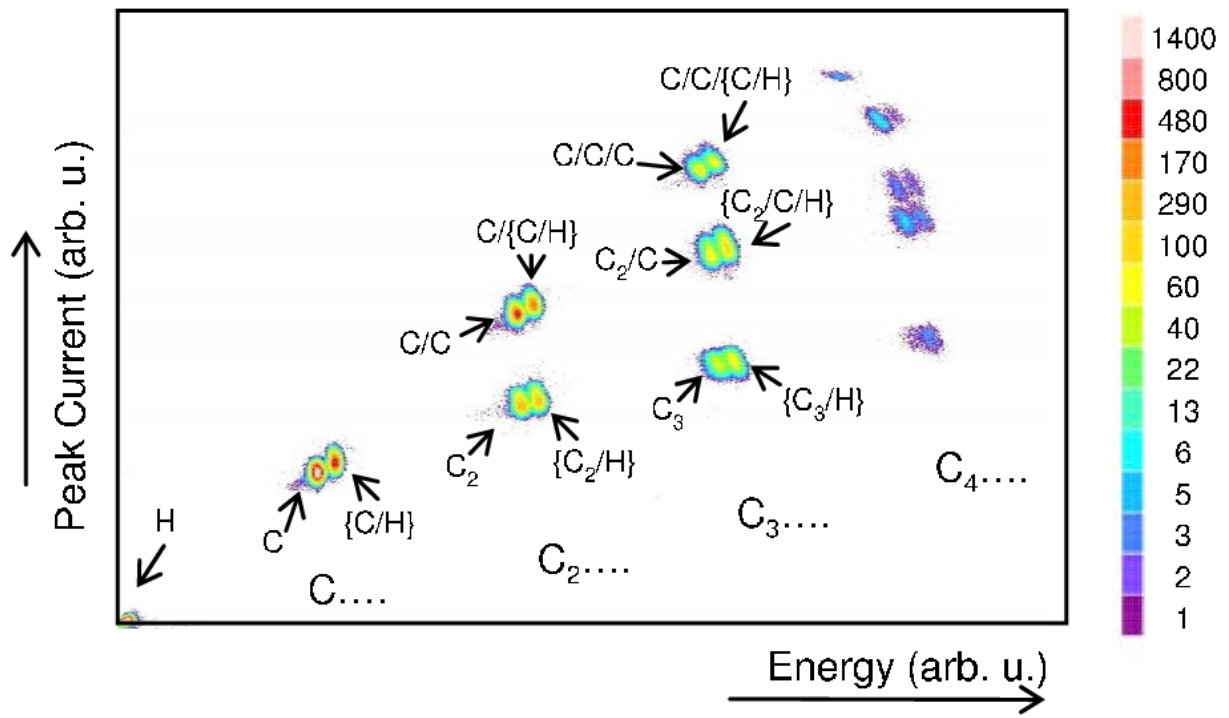


Figure 1

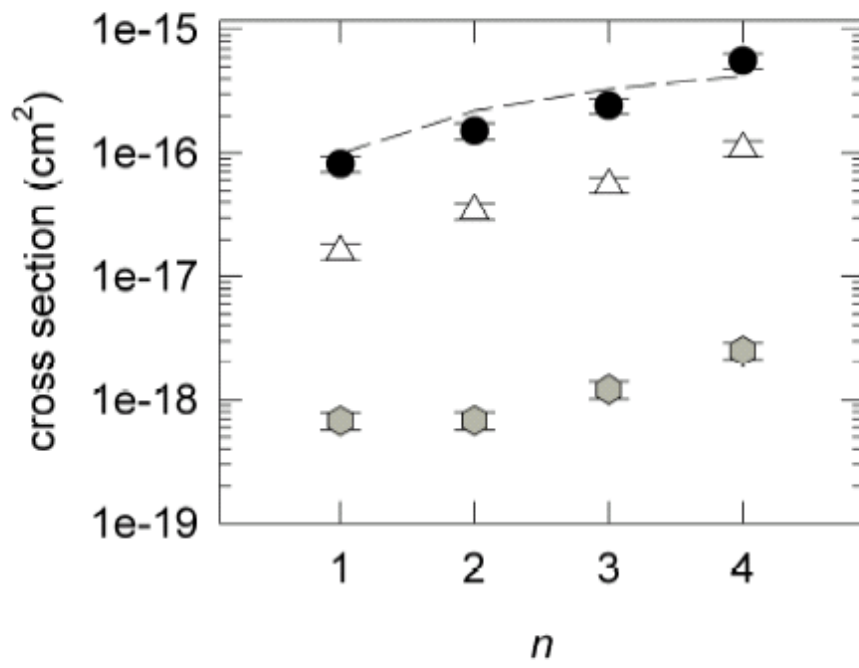


Figure 2

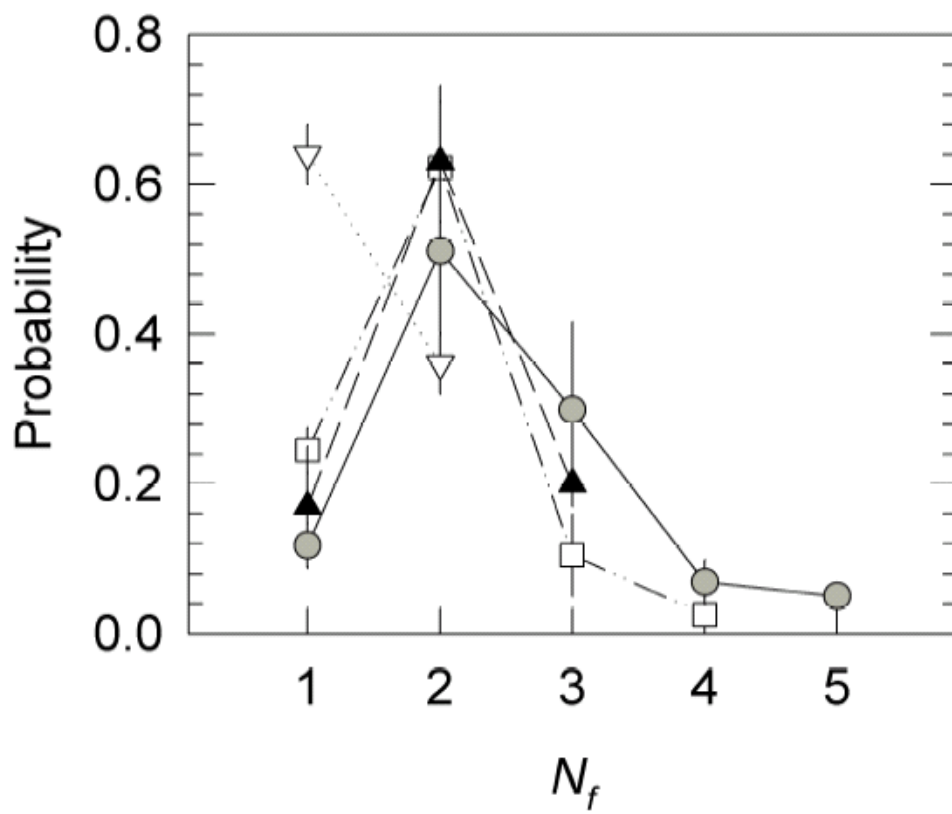


Figure 3

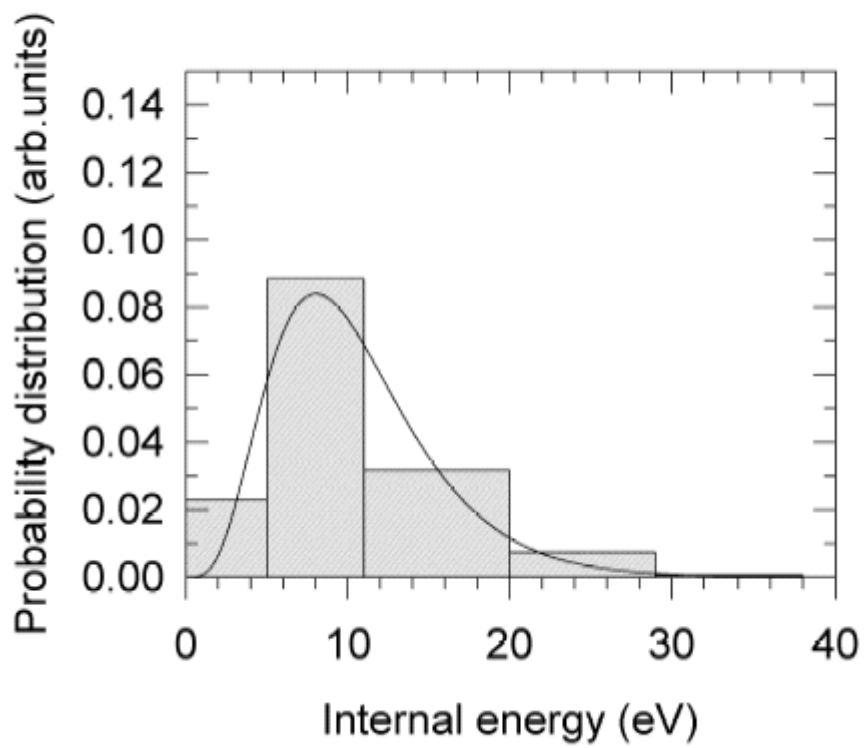


Figure 4

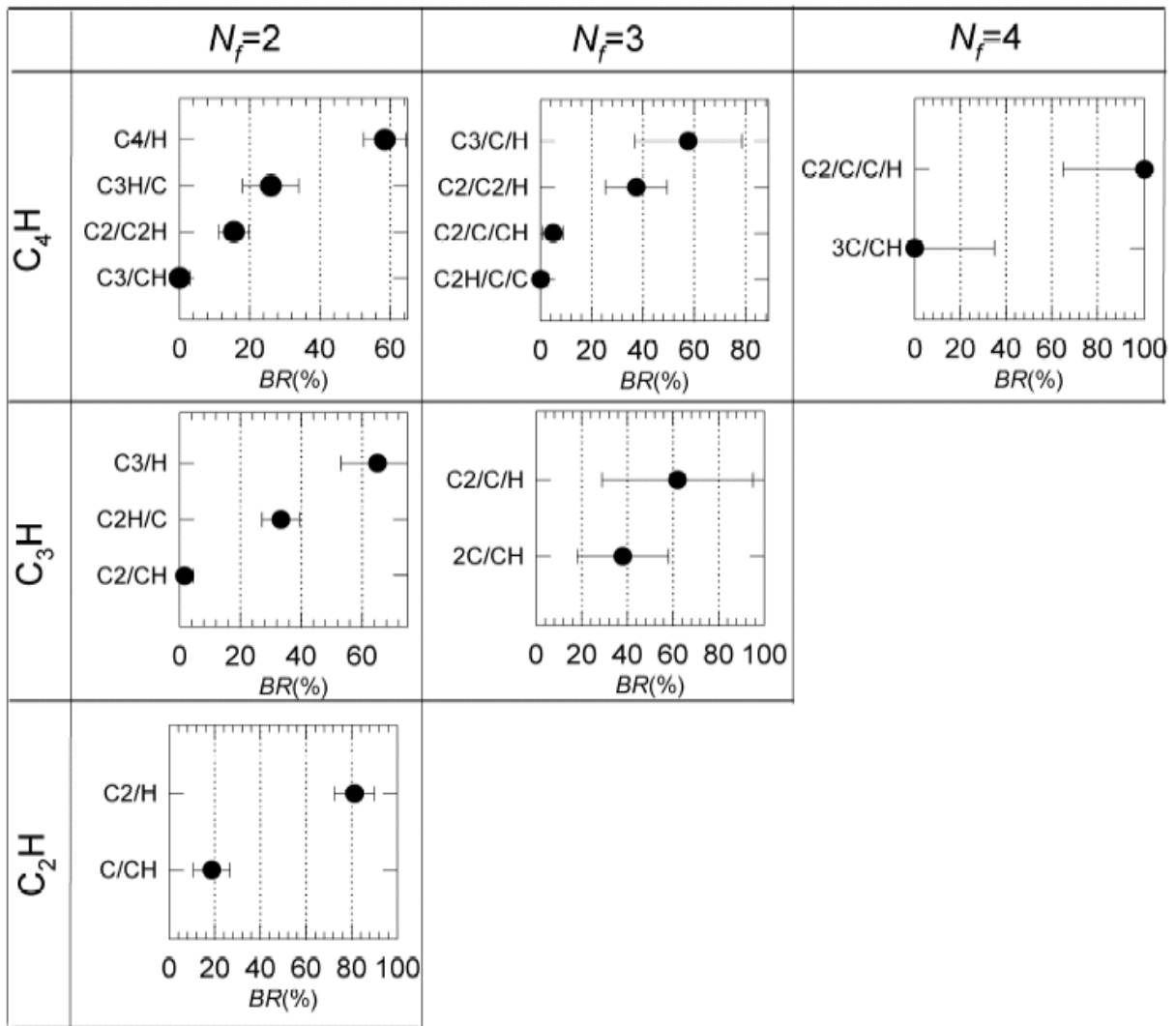


Figure 5

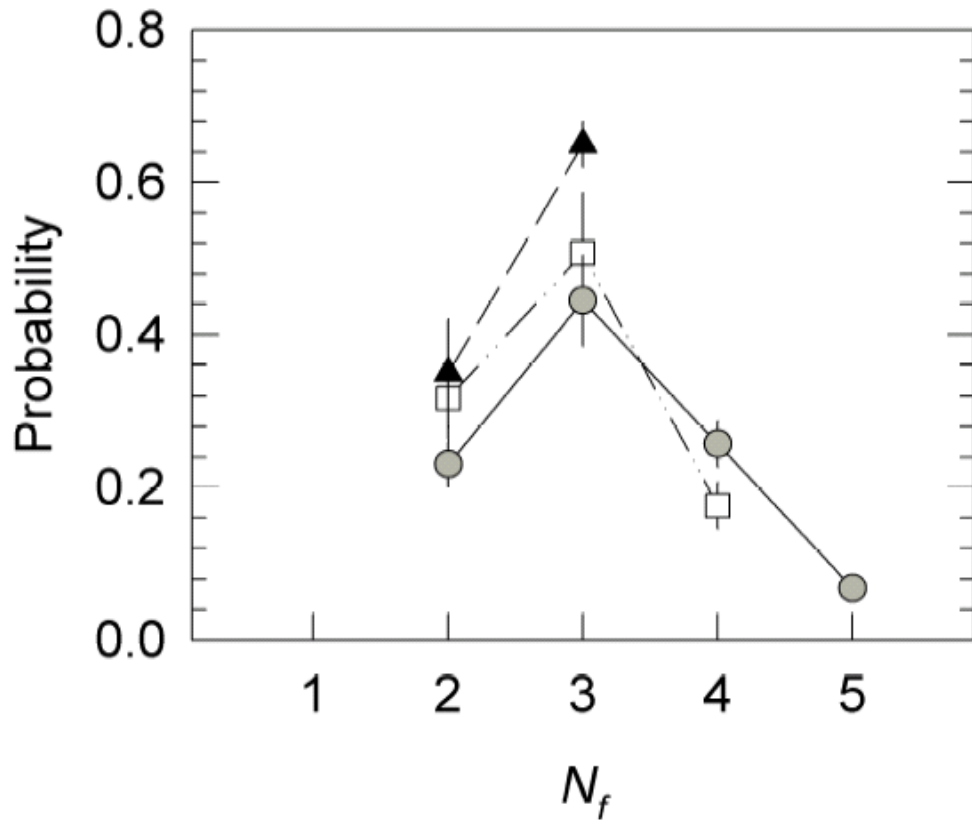


Figure 6

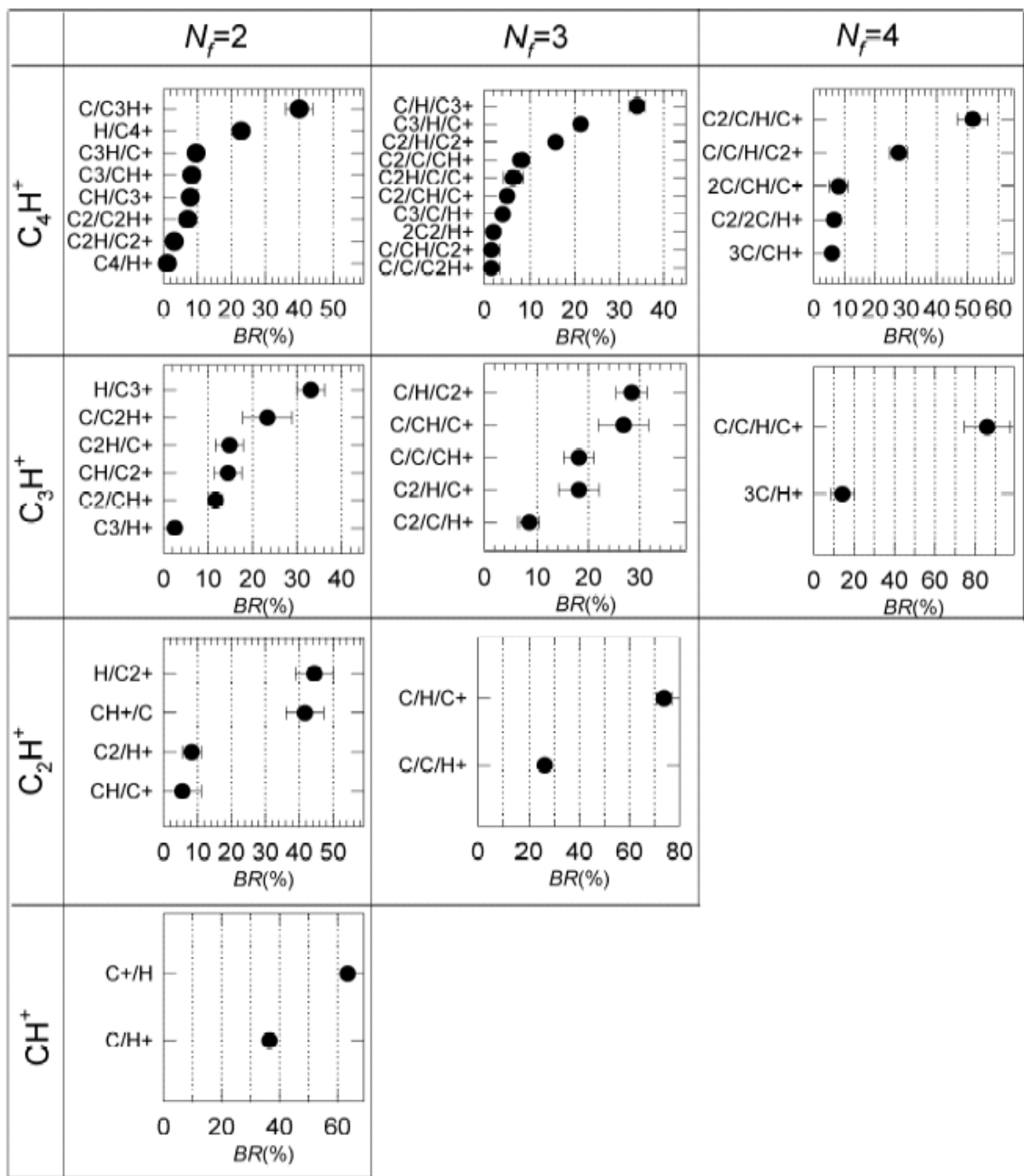


Figure 7

ANNEX :

Extraction of branching ratios by the grid method and current shape analysis

1- Expression of the transmission matrix; case of C₂H dissociation

A set of linear equations connect the intensities of the measured peaks to the intensities of the various dissociation channels (unknown quantities). We write TX=M where T is the transmission matrix, X is the vector containing the unknown quantities, M the vector containing the measured intensities (with grid and without grid). The probability that a fragment traverses the grid is t, the probability that it is stopped is s = (1-t) and since all fragments are independent, it is easy to write the form of the T matrix.

This is what is done below (equation 1A) for the case of the dissociation of C₂H. For the M vector, the index g refers to measurements done with the grid. As to the bracket sign [], it indicates that this measurement is an unresolved sum of two terms (see below). We note that components of the X vector (unknown quantities) contain the four branching ratios of dissociation of C₂H (C₂H, C₂/H, C/CH and 2C/H) but also channels having an incomplete mass (C₂, 2C, CH, C/H, C and H). These incomplete channels come from dissociation of incident C₂H⁺ on the residual gas and, although small, have to be introduced in the equation especially for estimation of the errors.

$$\begin{array}{cccccccccc} t & t^2 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & & ([C_2H])_g \\ 0 & 0 & t^2 & t^3 & 0 & 0 & 0 & 0 & 0 & 0 & & (C/[CH])_g \\ 0 & ts & 0 & 0 & t & 0 & 0 & 0 & 0 & 0 & C_2H & (C_2)_g \\ 0 & 0 & 0 & t^2s & 0 & t^2 & 0 & 0 & 0 & 0 & C_2/H & (2C)_g \\ 0 & 0 & ts & 2t^2s & 0 & 0 & t & t^2 & 0 & 0 & C/CH & ([CH])_g \\ 0 & 0 & ts & 2t^2s & 0 & 2ts & 0 & ts & t & 0 & 2C/H & (C)_g \\ 0 & ts & 0 & ts^2 & 0 & 0 & 0 & ts & 0 & t * & C_2 & (H)_g \\ 1 & 1 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 2C & [C_2H] \\ 0 & 0 & 1 & 1 & 0 & 0 & 0 & 0 & 0 & 0 & CH & C/[CH] \\ 0 & 0 & 0 & 0 & 1 & 0 & 0 & 0 & 0 & 0 & C/H & C_2 \\ 0 & 0 & 0 & 0 & 0 & 1 & 0 & 0 & 0 & 0 & C & 2C \\ 0 & 0 & 0 & 0 & 0 & 0 & 1 & 1 & 0 & 0 & H & [CH] \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 1 & 0 & & C \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 0 & 1 & & H \end{array} \quad (1A)$$

As mentioned before, the shape analysis method allows to resolve the fragmentation of C_n species. Then, C_2 or $2C$ are separately measured (see components of vector M in equation 1A). On the contrary, the H fragmentation is not resolved. The notation $[C_2H]$ in the M vector means that intensities of C_2H and C_2/H are summed. Similarly $[CH]$ is the sum of CH and C/H intensities and $C/[CH]$ the sum of C/CH and $2C/H$ intensities.

2-Resolution of equations, branching ratios and error bars

As seen from (1A), 14 measurements (components of the M vector) are performed for 10 unknown quantities (components of the X vector) since t and s are known. Since the system is over-determined, the idea is to test and keep the equations which are the most sensitive to changes in the unknown branching ratios (or equivalently to remove the unsensitive equations which are only bringing uncertainties on the results). The error bars are calculated as follows: measured values are allowed to vary, within experimental error bars, by a Monte Carlo random procedure and unknown intensities of vector X extracted by resolving the equation (1A). The distribution of extracted values for each unknown intensity, obtained by varying the set of measured data, provides directly a mean value and error bar for this intensity. This treatment is performed also for the background (jet not overlapping with the beam), the final error bar taking into account this background subtraction.

3- Subtraction of the ^{13}C isotope contribution, final branching ratios and error bars

As mentioned in section II, a sizeable contribution of $^{13}C^{12}C_{n-1}^+$ exists in the C_nH^+ beams. This means that $^{13}C^{12}C_p^{q+}$ fragments may exist and will appear at the same mass than $C_{p+1}H^{q+}$. For the case of incident C_2H^+ beam and restricting to the neutralization (electron capture) process, it means that three new unknown quantities have to be considered in our equations: ^{13}C (same detector response as CH), $^{12}C^{13}C$ (as C_2H) and $^{12}C/^{13}C$ (as C/CH). We could have added these unknown quantities in the equation (1A) and resolve it but we choose to subtract the contribution due to ^{13}C at the final stage (from the already extracted intensities) since it involves quantities that are measured in the experiment. Indeed we get, for the final intensities:

$$I_{\text{final}}(C_2H) = I_{\text{extr}}(C_2H) - b * BR(^{13}C^{12}C) \quad (2A-a)$$

$$I_{\text{final}}(\text{C}_2/\text{H})=I_{\text{extr}}(\text{C}_2/\text{H}) \quad (2\text{A-b})$$

$$I_{\text{final}}(\text{C}/\text{CH})= I_{\text{extr}}(\text{C}/\text{CH})- b * \text{BR}({}^{12}\text{C}/{}^{13}\text{C}) \quad (2\text{A-c})$$

$$I_{\text{final}}(2\text{C}/\text{H})=I_{\text{extr}}(2\text{C}/\text{H}) \quad (2\text{A-d})$$

Where I_{extr} are obtained by the procedure described above, b is the percentage of ${}^{12}\text{C}{}^{13}\text{C}^+$ intensity in the C_2H^+ beam and BR are the branching ratios of de-excitation of excited ${}^{13}\text{C}{}^{12}\text{C}$ created by electron capture in the ${}^{13}\text{C}{}^{12}\text{C}^+$ -He collision. These quantities are measured in the experiment (recordings with C_n^+ beams of identical velocity were systematically performed) and it was assumed that these measurements with ${}^{12}\text{C}_n^+$ beams are applicable to ${}^{13}\text{C}{}^{12}\text{C}_{n-1}^+$ beams (*i.e.* there is no isotopic effect on cross section and dissociation patterns).

The b quantity was also measured in the experiment. Indeed, following conclusions of Ben-Itzhak et al ⁷¹, we assumed that no CH^{++} ions may survive after 70 ns. Then the peak at $M = 13$ near the fragment C^{++} (detector 5, see Table I) was assigned to ${}^{13}\text{C}^{++}$ and not to CH^{++} and b was extracted from equation (3A):

$$b = \frac{I_{\text{extr}}({}^{12}\text{C}^{++}{}^{13}\text{C}^{++})}{I_{\text{extr}}(M = 25, q = 4)} * I_{\text{extr}}(M = 25, q = 0) \quad (3\text{A})$$

Error bars for final branching ratios took into account the error bars for extracted values I_{extr} , the error bar on the b value and error bars on BR values, these last ones quite small since measurements with C_n^+ projectiles done without grid have large statistics.

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FIGURE CAPTION:

Figure 1 (Color):

Two-dimensional representation of signals from neutral fragments in the C_4H^+ -He collision; colors relate to the number of events, see color scale on the right side.

Figure 2 :

Measured cross sections in the C_nH^+ -He collisions as a function of n. Black circles: total ionisation cross sections; open triangles: electronic excitation (dissociative part); grey hexagons: electron capture. Broken line: IAE calculations for ionization (see text).

Figure 3:

Measured probabilities of dissociation, P, of C_nH as a function of the number of emitted fragments N_f . Open triangles: CH; full triangles: C_2H ; open squares: C_3H ; full circles: C_4H . Lines are to guide the eye.

Figure 4:

Internal energy distribution of C_4H after electron capture. Grey rectangles are obtained by assuming a step function for $f(E)$; the solid line is obtained by using the analytical form $f(E)=E^{a1}\exp(-a2(E-a3)^{a4})$ (see text).

Figure 5:

Measured branching ratios of dissociation of C_nH for given values of the number of emitted fragments N_f . From left to right: $N_f=2$ to $N_f=4$; from bottom to top: C_2H to C_4H .

Figure 6:

Measured probabilities of dissociation, P, of C_nH^+ species as a function of the number of emitted fragments N_f . Full triangles: C_2H^+ ; open squares: C_3H^+ ; full circles: C_4H^+ . Lines are to guide the eye.

Figure 7:

Measured branching ratios of C_nH^+ for given values of the number of emitted fragments N_f . From left to right: $N_f=2$ to $N_f=4$; from bottom to top: CH^+ to C_4H^+