

## $H_n^+$ Ejection from Aligned $CH_3Cl$ Molecules in Intense $fs$ Laser Fields

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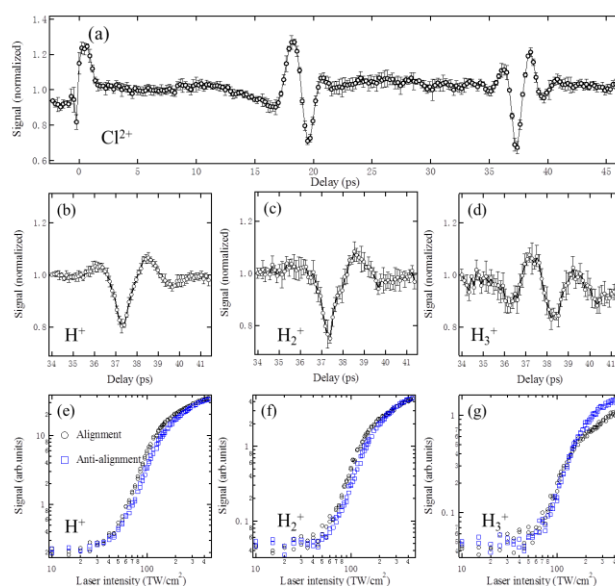
**Synopsis** We report experimental results on strong-field laser interact with aligned  $CH_3Cl$  molecules, the ejection of fragments  $H_n^+$  dependent on alignment of  $CH_3Cl$  molecules has been studied. The mechanism of fragmentation builds on the alignment sensitive ionization and dissociation from inner and outer valence orbitals or the excitation process of molecular have been discussed.

The study on the ejection of  $H_n^+$  from Hydrocarbon molecules is very important which is related to many photophysical processes such as proton migration, deformation, and other ultra-fast processes [1]. This phenomenon has been studied for different molecules in intense laser fields, such as  $CH_3OH$ ,  $C_mH_n$ , and  $CH_3Cl$  [2]. For further understanding of the underlying mechanism, a deep investigation with elaborated measurement is required.

We have studied the different fragment ejections, using  $CH_3Cl$  as an example, and focus on the effect of orbitals or the excitation process on the fragment ejection of polyatomic molecules in intense laser initiate dynamics. We performed the experiment for field-free aligned  $CH_3Cl$  molecules from  $fs$  laser impulsive interaction. We align the rotation state selected  $CH_3Cl$  from a hexapole Stark selector by using a linearly polarized laser, with a stretched pulse of 300  $fs$ , at the intensity of  $3 \times 10^{12} \text{ W/cm}^2$ . The aligned molecules were ionized or dissociated by a 50  $fs$  laser as a probe in the intensity range from  $1 \times 10^{13}$  -  $4 \times 10^{14} \text{ W/cm}^2$ , and the alignment dependence of strong field ionization and dissociation yields were measured, as shown in Figure 1.

The alignment dependence of fragment ions  $Cl^{2+}$ ,  $H_n^+$  ( $n=1,2,3$ ) are given in Fig. 1 (a)-(d), respectively. It is interesting to notice that  $H^+$  and  $H_2^+$  show the same alignment dependence as the case of  $Cl^{2+}$ , giving a larger yield in the direction of the probe laser polarization parallel to the molecular axis. But  $H_3^+$  gives a prefer yield in the direction of the probe laser polarization perpendicular to the molecular axis, just like the parent ion. The ion yields  $H_n^+$  of alignment and anti-alignment as a function of the laser intensity were also measured and the results are given in Figure 1(e)-(g). For  $H^+$  and  $H_2^+$ , the ion yield for the aligned molecules is

larger than that for anti-aligned molecules almost for the whole laser intensity region. The signal of  $H_3^+$  is almost the same when laser intensity below  $1.6 \times 10^{14} \text{ TW}$ , and as increasing the laser intensity the signal is stronger when probe laser is perpendicular than parallel to the molecular axis. Furthermore,  $H_n^+$  ejection was controlled by manipulating the molecular alignment. The same dependence of the ejected  $H_3^+$  and  $CH_3Cl^+$  on the alignment observed indicates that  $H_3^+$  is ejected from  $CH_3Cl^{2+}$ , produced by sequence double ionization.



**Figure 1.** The delay dependent of the yields of signal produced from different fragmentation channels  $Cl^{2+}$ ,  $H_n^+$  ( $n=1,2,3$ ) (a)-(d). (e)-(g) is ion intensities of  $H_n^+$  ( $n=1,2,3$ ) dissociated from aligned and anti-aligned molecules as a function of laser intensity.

## References

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- [2] E. Wells *et al* 2013 *Nat. Comms* 4 2895.

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