

Theoretical and Physical Chemistry Institute National Hellenic Research Foundation

Vass. Constantinou 48, Athens

LECTURE

"Transitions in the continuous spectrum of atoms induced by two ultrashort pulses with time delay"

Dr. Yannis Komninos

Theoretical and Physical Chemistry Institute, National Hellenic Research Foundation,

Athens, Greece

Thursday, May 10, 2018, 12:00 Seminar room, ground floor, NHRF

Transitions in the continuous spectrum of atoms induced by two ultrashort pulses

with time delay

Dr. Yannis Komninos

Theoretical and Physical Chemistry Institute, National Hellenic Research Foundation, Athens, Greece

There is a growing interest, both theoretically [1] and experimentally [2], in recent years for ultrafast processes in atoms induced by one or more coherent pulses. Motivated by this we have examined the scheme

discrete $|0\rangle \xrightarrow{\text{weak XUV}} \text{resonance } |1\rangle \xleftarrow{\text{moderately strong MIR}} \text{femtosec pulse } \omega_2 \rightarrow \text{resonance } |2\rangle$

where $|0\rangle$ is a discrete state, and $|1\rangle$ and $|2\rangle$ are resonance states.

The attosecond and femtosecond pulses with central frequencies ω_1 and ω_2 ($\omega_1 >> \omega_2$) act with an ultrashort time delay, t_D between them. In addition to the t_D , the external control parameters are the intensity and the full-width at half maximum (FWHM) of the femtosecond pulse.

As a specific application we examine the time-resolved dynamics associated with the processes $He_{1s^{2}} S \rightarrow (2s_{2}p)^{-1}P^{o} \leftrightarrow (2p^{2})^{-1}D$. For the transition $1s^{2} S \rightarrow (2s_{2}p)^{-1}P^{o}$, the FWHM of the XUV pulse is 160 attoseconds and its peak intensity at $10^{11} W / cm^{2}$ as it was used recently in the experiment of Kaldun et al [2]. The few-cycle mid-infrared (MIR) pulse couples the resonance states and its duration is varied in the range of a few decades of femtoseconds. Its intensity is in the range $5 \times 10^{10} - 5 \times 10^{11} W / cm^{2}$.

The results include the time-resolved formation of either the $(2s_2p)$ ¹ P^o or the $(2p^2)$ ¹D resonance, while they are coupled with each other. For the $(2s_2p)$ ¹ P^o state, comparison is made with the case where the time-dependent buildup of its asymmetric profile, excited from $1s^2$ ¹S by the same attosecond pulse, is calculated in the absence of coupling to another resonance.

References

[1] Th. Mercouris, Y. Komninos and C. A. Nicolaides, *Time-dependent formation of the profile of the* $He 2s2p \,^{1}P^{o}$ *state excited by a short laser pulse*, Phys. Rev. A **75**, 013407 (2007). Erratum, Phys. Rev. A **87**, 069905(E) (2013).

[2] A. Kaldun, A. Blättermann, V. Stooß, S. Donsa, H. Wei, R. Pazourek, S. Nagele, C. Ott, C. D. Lin, J. Burgdörfer, T. Pfeifer, *Observing the ultrafast buildup of a Fano resonance in the time domain*, Science **354**, 738 (2016).