

Molecular attosecond dynamics

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The generation and characterization of single attosecond pulses have been achieved through several efforts in the field of ultrafast intense laser sources over the last 20 years [1] and through theoretical developments on the interaction of intense light pulses with atomic and molecular systems [2]. The duration of attosecond pulses is rapidly approaching the atomic unit of time [3] that represents, in the classical description of the atomic model, the natural time scale of the electronic motion; also in quantum mechanics the attosecond regime is the relevant time domain for electrons as the inverse of the energy spacing among electronic levels, (that determines the time constant for non-stationary states), lies typically in this range. First applications of such pulses have been mainly focused on simple atoms or molecules to validate new experimental approaches and to gain first information on electron-electron correlation.

Experimental and theoretical results on the ultrafast dynamics initiated by single attosecond pulses in helium and H₂ and D₂ will be shown.

In hydrogen several states of the neutral molecule (autoionizing states) or of the molecular ion (1sσ_g and 2pσ_u) can be accessed due to the large bandwidth of the attosecond pulses. The electron dynamics can be probed and controlled using a synchronized infrared few-cycle pulse [4].

New directions for the investigation of attosecond dynamics in more complex molecular systems will be discussed. In particular the experimental challenges related to the complete characterization of complex dynamics involving electron and nuclear degree of freedom will be analyzed.

References

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