# Quantum dynamics with wave packet approaches: applications to the Infrared spectroscopy of protonated molecules

### Fabien Gatti<sup>\*</sup>

Orsay, France, Institut des Sciences Moléculaires d'Orsay (ISMO),

UMR 8214, Université Paris-Saclay

\*E-mail: <u>fabien.gatti@universite-paris-saclay.fr</u>

## David Lauvergnat

Orsay, France, Institut de Chimie Physique (ICP),

Université Paris-Saclay

### Markus Schröder, David Mendive Tapia, H.-D. Meyer and O. Vendrell

Theoretische Chemie, Heidelberg Universität, Germany

#### **ABSTRACT**

Many molecular processes, ranging from fundamental to applied problems, are known today to be impacted by strong nuclear quantum mechanical effects, including phenomena like tunneling, zero point energy effects, or non-adiabatic transitions. Recent success in helping to understand experimental observations in fields like heterogeneous catalysis, photochemistry, reactive scattering, optical spectroscopy, or femto- and attosecond chemistry and spectroscopy underlines that nuclear quantum mechanical effects affect many areas of chemical and physical research. The correct theory to describe the corresponding dynamics is Molecular Quantum Dynamics [1,2]. In contrast to standard quantum chemistry calculations, where the nuclei are treated classically, molecular quantum dynamics can cover quantum mechanical effects in their motion. New strategies have been developed to extend the studies to systems of increasing size. In particular, we present here several applications of the MultiConfiguration Time-Dependent Hartree method (MCTDH) [3,4,5] to the understanding and the control of molecular processes involving quantum effects. MCTDH can be seen as a timedependent MCSCF approach for the nuclei where wavepackets are propagated on one or several potential energy surfaces. Several examples will be presented highlighting the presence and the exploitation of quantum effects in molecular processes. Special emphasis will be placed on the spectroscopy in full dimensionality of protonated molecules: Eigen cation (33D), extended Zundel cation (51D), protonated acethylene (9D). We will sketch the results of current studies of water clusters such as water dimer and trimer (collaboration with Steve Ndengué, ICTP, Kigali, Rwanda). We will also illustrate the possibility to simulate the coherent behavior of molecules  $(CO_2)$  at the attosecond time scale (collaboration with Lou barreau, France and John Stanton, University of Florida).

#### References

[1] Molecular Quantum Dynamics, From Theory to Applications, Ed. F . Gatti, Springer, (2014) Heidelberg.

[2] Quantum Physics, Applications to Chemistry, F. Gatti, B. Lasorne, H.-D. Meyer and A. Nauts, Lectures Notes in Chemistry, Springer, 2017.

[3] H.-D. Meyer, U. Manthe, and L.S. Cederbaum, The multi-configurational time-dependent Hartree approach. Chem. Phys. Lett. 165 (1990), 73.

[4] M. H. Beck, A. Jäckle, G. A. Worth and H.-D. Meyer, The multiconfiguration time-dependent Hartree method:

A highly efficient algorithm for propagating wavepackets, Physics Reports 324 (2000),

[5] H.-D. Meyer, F. Gatti, and G. A. Worth, editors, Multidimensional Quantum Dynamics: MCTDH Theory and Applications, Wiley-VCH,(2009), Weinheim.