

CHIRALEX: theory

Emil J. Zak^{1,*}

¹Center for Free-Electron Laser Science, Deutsches Elektronen-Synchrotron DESY, Notkestraße 85, 22607 Hamburg, Germany
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I. INTRODUCTION

II. BASIS FUNCTIONS AND GRIDS

A. Radial quadratures

B. Radial grid

C. Radial basis functions

1. Interpolated radial functions

D. Angular basis functions

III. INDEX MAPPING

IV. ROTATED ELECTROSTATIC POTENTIAL

Each molecular-frame orientation gives different electrostatic potential as seen from the laboratory frame, in which the basis set is defined. This means that for each orientation one must generate generally a different Lebe-

dev quadrature grid. Even if a global scheme is used, the values of the electrostatic potential will be different at these points for each orientation. This is because the lebedev grid is defined in the laboratory frame. Therefore at this stage, a separate Psi4 calculation must be initialized for each orientation. I do not see a way around it at the moment, other than performing multipole expansion and calculating the angular part (spherical tensor form) of the potential analytically. But finding this expansion is costly, even more than running Psi4 for every orientation.

V. KINETIC ENERGY OPERATOR

VI. HAMILTONIAN

We decided to keep the bound state Hamiltonian H_0 and the initial propagation Hamiltonian H_{init} as separate entities, calculated independently. The computational overhead related to calculating the bound-state part twice is marginal, as we assume that the full propagation basis is much bigger than the basis for the bound Hamiltonian $N_{bas} \gg N_{bas0}$

* emil.zak@cfel.de