



OPEN ACCESS

EDITED BY
Yuichi Fujimura,
Tohoku University, Japan

REVIEWED BY
Xiaowei Sheng,
Anhui Normal University, China
Shu Ohmura,
Nagoya Institute of Technology, Japan

*CORRESPONDENCE
Takeshi Sato,
sato@atto.t.u-tokyo.ac.jp

SPECIALTY SECTION
This article was submitted to Physical
Chemistry and Chemical Physics,
a section of the journal
Frontiers in Chemistry

RECEIVED 30 June 2022
ACCEPTED 09 August 2022
PUBLISHED 13 September 2022

CITATION
Pathak H, Sato T and Ishikawa KL (2022),
Time-dependent optimized coupled-
cluster method with doubles and
perturbative triples for first principles
simulation of multielectron dynamics.
Front. Chem. 10:982120.
doi: 10.3389/fchem.2022.982120

COPYRIGHT
© 2022 Pathak, Sato and Ishikawa. This
is an open-access article distributed
under the terms of the [Creative
Commons Attribution License \(CC BY\)](#).
The use, distribution or reproduction in
other forums is permitted, provided the
original author(s) and the copyright
owner(s) are credited and that the
original publication in this journal is
cited, in accordance with accepted
academic practice. No use, distribution
or reproduction is permitted which does
not comply with these terms.

Time-dependent optimized coupled-cluster method with doubles and perturbative triples for first principles simulation of multielectron dynamics

Himadri Pathak¹, Takeshi Sato^{1,2,3*} and Kenichi L. Ishikawa^{1,2,3}

¹Department of Nuclear Engineering and Management, School of Engineering, The University of Tokyo, Tokyo, Japan, ²Photon Science Center, School of Engineering, The University of Tokyo, Tokyo, Japan, ³Research Institute for Photon Science and Laser Technology, The University of Tokyo, Tokyo, Japan

We report the formulation of a new, cost-effective approximation method in the time-dependent optimized coupled-cluster (TD-OCC) framework [T. Sato *et al.*, *J. Chem. Phys.* 148, 051101 (2018)] for first-principles simulations of multielectron dynamics in an intense laser field. The method, designated as TD-OCCD(T), is a time-dependent, orbital-optimized extension of the “gold-standard” CCSD(T) method in the ground-state electronic structure theory. The equations of motion for the orbital functions and the coupled-cluster amplitudes are derived based on the real-valued time-dependent variational principle using the fourth-order Lagrangian. The TD-OCCD(T) is size extensive and gauge invariant, and scales as $O(N^7)$ with respect to the number of active orbitals N . The pilot application of the TD-OCCD(T) method to the strong-field ionization and high-order harmonic generation from a Kr atom is reported in comparison with the results of the previously developed methods, such as the time-dependent complete-active-space self-consistent field (TD-CASSCF), TD-OCC with double and triple excitations (TD-OCCDT), TD-OCC with double excitations (TD-OCCD), and the time-dependent Hartree-Fock (TDHF) methods.

KEYWORDS

multielectron dynamics, time-dependent optimized coupled-cluster, high harmonic generation, strong laser field, strong field ionization

1 Introduction

Recent years witnessed unprecedented progress in laser technologies, which made it possible to observe the motions of electrons at the attosecond time scale (Itatani *et al.* (2004); Corkum and Krausz (2007); Krausz and Ivanov (2009); Baker *et al.* (2006)). On the other hand, various theoretical and numerical methods have been developed for interpreting, understanding, and predicting the experiments.

The multi-configuration time-dependent Hartree-Fock (MCTDHF) method (Caillat et al. (2005); Kato and Kono (2004); Nest et al. (2005); Haxton et al. (2011); Hochstuhl and Bonitz (2011)), and the time-dependent complete-active-space self-consistent-field (TD-CASSCF) method (Sato and Ishikawa (2013); Sato et al. (2016); Sato et al. (2018a)) are the most rigorous approaches to solve time-dependent Schrödinger equation (TDSE) of many-electron systems, where the wavefunction is given by the full configuration interaction (FCI) expansion,

$$\Psi(t) = \sum_I C_I(t) \Phi_I(t), \quad (1)$$

with both CI coefficients $\{C_I(t)\}$ and orbital functions $\{\psi_p(t)\}$ constituting Slater determinants $\{\Phi_I(t)\}$ are propagated in time according to the time-dependent variational principle (TDVP). The TD-CASSCF method broadens the applicability of the MCTDHF method by flexibly classifying the orbital subspace into frozen-core, dynamical-core, and active. Unfortunately, the factorial computational scaling impedes large-scale applications. There are reports of various affordable size-inextensive methods (Miyagi and Madsen (2013, 2014); Haxton and McCurdy (2015); Sato and Ishikawa (2015)) developed by limiting the CI expansion of the wavefunction. Alternatively, the size-extensive coupled-cluster method, which relies on an exponential wavefunction, is a superior choice to address these problems with a polynomial cost-scaling (Kümmel (2003); Shavitt and Bartlett (2009)). We have developed an explicitly time-dependent coupled-cluster method considering optimized orthonormal orbitals within the flexibly chosen active space, called the time-dependent optimized coupled-cluster (TD-OCC) method, (Sato et al. (2018b)) including double (TD-OCCD) and double and triple excitation amplitudes (TD-OCCDT). Our method is a time-dependent formulation of the stationary optimized coupled-cluster method (Scuseria and Schaefer (1987); Sherrill et al. (1998); Krylov et al. (1998)). Kvaal (Kvaal (2012)) also developed an orbital adaptive time-dependent coupled-cluster (OATDCC) method using biorthogonal orbitals. We take note of a few reports on the time-dependent coupled-cluster methods (Huber and Klamroth (2011); Pigg et al. (2012); Nascimento and DePrince (2016)), using time-independent orbitals, and their interpretation (Pedersen and Kvaal (2019); Pedersen et al. (2021)), including the very initial attempts (Schonhammer (1978); Hoodbhoy and Negele (1978, 1979)).

The TD-OCCDT scales as $O(N^8)$ (N = the number of active orbitals), not ideally suited for applications to larger chemical systems. Therefore, we have developed a few lower cost methods in the TD-OCC framework (Pathak et al. (2020b,c,a, 2021)). We find triple excitations are necessary, including perfect optimization of the orbitals. Therefore, we are interested in developing affordable TD-OCC methods retaining a part of the triples. The most popular coupled-cluster method that treats the triple excitation amplitudes approximately is called CCSD(T) (Raghavachari et al. (1989);

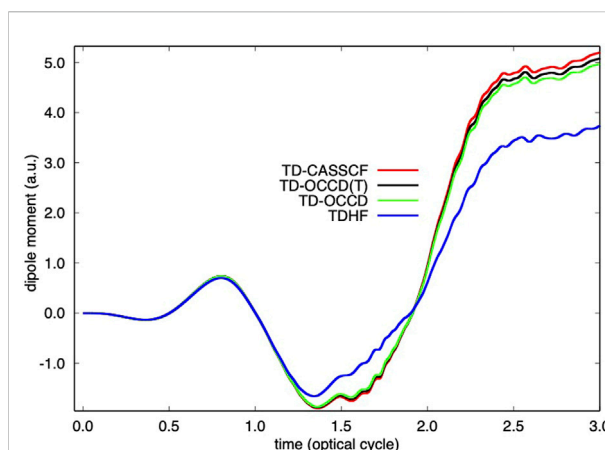


FIGURE 1

Time evolution of dipole moment of Kr irradiated by a laser pulse with a wavelength of 800 nm and a peak intensity of 2×10^{14} W/cm² calculated with TDHF, TD-OCCD, TD-OCCD(T), and TD-CASSCF methods.

Watts et al. (1993)). Bozkaya et al, (Bozkaya and Schaefer (2012)) included various symmetric and asymmetric triple excitation corrections to their optimized double (OD) method.

In this communication, we report the formulation and implementation of the CCSD(T) method in the time-dependent optimized coupled-cluster framework, TD-OCCD(T). Following our previous works (Sato et al. (2018b); Pathak et al. (2020b,c, 2021)), we exclude single excitation amplitudes but optimize the orbitals according to time-dependent variational principle (TDVP). As the first application of this method, we study electron dynamics in Kr using intense near-infrared laser fields.

2 Methods

The second quantization representation of the Hamiltonian, including the laser field, is as follows,

$$\hat{H} = h_v^\mu(t) \hat{c}_\mu^\dagger \hat{c}_\nu + \frac{1}{2} u_{\nu\lambda}^{\mu\gamma} \hat{c}_\mu^\dagger \hat{c}_\nu^\dagger \hat{c}_\lambda \hat{c}_\gamma \quad (2)$$

where \hat{c}_μ^\dagger (\hat{c}_μ) represents a creation (annihilation) operator in a complete, orthonormal set of $2n_{\text{bas}}$ time-dependent spin-orbitals $\{\psi_\mu(t)\}$. n_{bas} is the number of basis functions used for expanding the spatial part of ψ_μ , which, in the present real-space implementation, corresponds to the number of grid points, and

$$h_v^\mu(t) = \int dx_1 \psi_\mu^*(x_1) [h_0 + V_{\text{ext}}] \psi_\nu(x_1), \quad (3)$$

$$u_{\nu\lambda}^{\mu\gamma} = \iint dx_1 dx_2 \frac{\psi_\mu^*(x_1) \psi_\gamma^*(x_2) \psi_\nu(x_1) \psi_\lambda(x_2)}{|\mathbf{r}_1 - \mathbf{r}_2|}, \quad (4)$$

where $x_i = (\mathbf{r}_i, \sigma_i)$ represents a composite spatial-spin coordinate. h_0 is the field free one-electronic Hamiltonian and $V_{\text{ext}} = A(t)p_z$

in the velocity gauge, $A(t) = -\int^t E(t') dt'$ is the vector potential, with $E(t)$ being the laser electric field linearly polarized along the z axis.

The complete set of $2n_{\text{bas}}$ spin-orbitals (labeled with $\mu, \nu, \gamma, \lambda$) is divided into n_{occ} occupied (o, p, q, r, s) and $2n_{\text{bas}} - n_{\text{occ}}$ virtual spin-orbitals. The coupled-cluster (or CI) wavefunction is constructed only with occupied spin-orbitals, which are time-dependent in general, and virtual spin-orbitals form the orthogonal complement of the occupied spin-orbital space. The occupied spin-orbitals are classified into n_{core} core spin-orbitals, which are occupied in the reference Φ and kept uncorrelated, and $N = n_{\text{occ}} - n_{\text{core}}$ active spin-orbitals (t, u, v, w) among which the active electrons are correlated. The active spin-orbitals are further split into those in the hole space (i, j, k, l) and the particle space (a, b, c, d), which are defined as those occupied and unoccupied, respectively, in the reference Φ . The core spin-orbitals can further be split into frozen-core space (i'', j''), fixed in time and the dynamical-core space (i', j'), propagated in time (Sato and Ishikawa (2013)) (See Figure 1 in Sato et al. (2018b) for a pictorial illustration).

The real action formulation of the TDVP with orthonormal orbitals is our guiding principle, (Sato et al. (2018b))

$$S = \text{Re} \int_{t_0}^{t_1} L dt = \frac{1}{2} \int_{t_0}^{t_1} (L + L^*) dt, \quad (5)$$

$$L = \langle \Phi | (1 + \hat{\Lambda}) e^{-\hat{T}} \left(\hat{H} - i \frac{\partial}{\partial t} \right) e^{\hat{T}} | \Phi \rangle, \quad (6)$$

$$\hat{T} = \hat{T}_2 + \hat{T}_3 \dots = \tau_{ij}^{ab} \hat{E}_{ij}^{ab} + \tau_{ijk}^{abc} \hat{E}_{ijk}^{abc} \dots, \quad (7)$$

$$\hat{\Lambda} = \hat{\Lambda}_2 + \hat{\Lambda}_3 \dots = \lambda_{ab}^{ij} \hat{E}_{ab}^{ij} + \lambda_{abc}^{ijk} \hat{E}_{abc}^{ijk} \dots, \quad (8)$$

where $\tau_{ij\dots}^{ab\dots}$ ($\lambda_{ab\dots}^{ij\dots}$) are (de-)excitation amplitudes, and $\hat{E}_{ij\dots}^{ab\dots} = \hat{c}_a^\dagger \hat{c}_b \dots \hat{c}_j \hat{c}_i$. The stationary conditions, $\delta S = 0$, with respect to the variation of the parameters of the wavefunction ($\delta \tau_{ij\dots}^{ab\dots}$, $\delta \lambda_{ab\dots}^{ij\dots}$, and $\delta \psi_\mu$) gives us the corresponding equations of motions (EOMs), $\delta \psi_\mu$ is orthonormality-conserving orbital variation.

For deriving the TD-OCCD(T) method, we first construct a fourth-order Lagrangian defined in Pathak et al. (2021). We make a further approximation to the Lagrangian and write separating it into two parts,

$$L_{\text{CCD(T)}}^{(4)} = L_0 + \langle \Phi | (1 + \hat{\Lambda}_2) [(\bar{f} + \hat{v}) e^{\bar{T}_2}]_c | \Phi \rangle - i \lambda_{ab}^{ij} \tau_{ij}^{ab} \quad (9a)$$

$$+ \langle \Phi | \hat{\Lambda}_2 [(\bar{f} + \hat{v}) \hat{T}_3]_c | \Phi \rangle + \langle \Phi | \hat{\Lambda}_3 (\bar{f} \hat{T}_3)_c | \Phi \rangle$$

$$+ \langle \Phi | \hat{\Lambda}_3 (\hat{v} \hat{T}_2)_c | \Phi \rangle - i \lambda_{abc}^{ijk} \tau_{ijk}^{abc}, \quad (9b)$$

where $\bar{f} = \hat{f} - i \hat{X}$, $\hat{f} = (h_q^p + v_{qj}^{pj}) \{\hat{E}_q^p\}$, $\hat{v} = v_{qs}^{pr} \{\hat{E}_{qs}^{pr}\}/4$, and $v_{qs}^{pr} = u_{qs}^{pr} - u_{sq}^{pr}$, $\hat{X} = X_\nu^\mu \hat{E}_\nu^\mu$, and $X_\nu^\mu = \langle \psi_\mu | \psi_\nu \rangle$ is anti-Hermitian. The double amplitudes are obtained by making $L_{\text{CCD(T)}}^{(4)}$ of Eq. 9a stationary with respect to $\delta S / \delta \lambda_{ab}^{ij}(t) = 0$, $\delta S / \delta \tau_{ij}^{ab}(t) = 0$, the triples by making Eq. 9b stationary with respect to $\delta S / \delta \lambda_{abc}^{ijk}(t) = 0$, and $\delta S / \delta \tau_{ijk}^{abc}(t) = 0$,

$$i \dot{\tau}_{ij}^{ab} = v_{ij}^{ab} - p(ij) \bar{f}_j^k \tau_{ik}^{ab} + p(ab) \bar{f}_c^a \tau_{ij}^{cb} + \frac{1}{2} v_{cd}^{ab} \tau_{ij}^{cd} + \frac{1}{2} v_{ij}^{kl} \tau_{kl}^{ab} + p(ij) p(ab) v_{ic}^{ak} \tau_{kj}^{cb} - \frac{1}{2} p(ij) \tau_{ik}^{ab} \tau_{jl}^{cd} \tau_{cd}^{kl} + \frac{1}{2} p(ab) \tau_{ij}^{bc} \tau_{kl}^{ad} \tau_{cd}^{kl} + \frac{1}{4} \tau_{kl}^{ab} \tau_{ij}^{cd} \tau_{cd}^{kl} + \frac{1}{2} p(ij) p(ab) \tau_{il}^{bc} \tau_{jk}^{ad} \tau_{cd}^{kl} \quad (10)$$

$$-i \dot{\lambda}_{ab}^{ij} = \frac{v_{ab}^{ij}}{ab} - p(ij) \bar{f}_k^i \lambda_{ab}^{kj} + p(ab) \bar{f}_c^a \lambda_{cb}^{ij} + \frac{1}{2} v_{cd}^{ab} \lambda_{cd}^{ij} + \frac{1}{2} v_{ij}^{kl} \lambda_{kl}^{ab} + p(ij) p(ab) v_{kb}^{cj} \lambda_{ac}^{ik} - \frac{1}{2} p(ij) \lambda_{cd}^{ik} \tau_{kl}^{cd} \tau_{ab}^{jl} + \frac{1}{2} p(ab) \lambda_{bc}^{kl} \tau_{kl}^{cd} \tau_{ad}^{ij} + \frac{1}{4} \lambda_{kl}^{ij} \tau_{kl}^{cd} \tau_{cd}^{ij} + \frac{1}{2} p(ij) p(ab) \lambda_{ac}^{jk} \tau_{kl}^{cd} \tau_{bd}^{il} - \frac{1}{2} p(ij) \lambda_{ab}^{ik} \tau_{kl}^{cd} \tau_{cd}^{jl} + \frac{1}{2} p(ab) \lambda_{bc}^{ij} \tau_{kl}^{cd} \tau_{ad}^{kl} + \frac{1}{4} \lambda_{ij}^{ab} \tau_{cd}^{kl} \tau_{cd}^{kl} \tau_{ab}^{ij} \quad (11)$$

$$i \dot{\tau}_{ijk}^{abc} = p(k/ij) p(a/bc) v_{ik}^{ad} \tau_{ij}^{ab} - p(i/jk) p(c/ab) v_{jk}^{cd} \tau_{il}^{ab} - p(k/ij) \bar{f}_k^l \tau_{ijl}^{abc} + p(c/ab) \bar{f}_c^d \tau_{ijk}^{abd}, \quad (12)$$

$$-i \dot{\lambda}_{abc}^{ijk} = p(k/ij) p(a/bc) v_{bc}^{ad} \lambda_{ad}^{ij} - p(c/ab) p(i/jk) v_{ic}^{jk} \lambda_{ab}^{ij} + p(c/ab) \bar{f}_c^d \lambda_{abd}^{ijk} - p(k/ij) \bar{f}_i^c \lambda_{abc}^{ijl} + p(i/jk) p(a/bc) \bar{f}_a^l \lambda_{bc}^{ijk}, \quad (13)$$

where $p(\mu\nu)$ and $p(\mu|\nu\gamma)$ are the permutation operators; $p(\mu\nu) A_{\mu\nu} = A_{\mu\nu} - A_{\nu\mu}$, and $p(\mu|\nu\gamma) = 1 - p(\mu\nu) - p(\mu\gamma)$.

The EOM for the orbitals can be written down in the following form Sato et al. (2016),

$$i |\dot{\psi}_p\rangle = (\hat{I} - \hat{P}) \hat{F} |\psi_p\rangle + i |\psi_q\rangle X_p^q, \quad (14)$$

where $\hat{I} = \sum_\mu |\psi_\mu\rangle \langle \psi_\mu|$ is the identity operator within the space spanned by the given basis, $\hat{P} = \sum_q |\psi_q\rangle \langle \psi_q|$ is the projector onto the occupied spin-orbital space, and

$$\hat{F} |\psi_p\rangle = \hat{h} |\psi_p\rangle + \hat{W}_s^r |\psi_q\rangle P_{or}^{qs} (D^{-1})_p^o, \quad (15)$$

where D and P are Hermitalized one- (1RDM) and two- (2RDM) particle reduced density matrices defined in Sato et al. (2018b), and W_s^r is the mean-field operator (Sato and Ishikawa (2013)). The matrix element X_p^q includes orbital rotations among various subspaces. Non-redundant orbital rotations are determined by $i(\delta_b^a D_i^j - D_b^a \delta_i^j) X_j^b = F_p^a D_i^p - D_p^a F_i^{p*} - \frac{i}{8} \tau_{ijk}^{abc} \lambda_{bc}^{jk} - \frac{i}{8} \tau_{ijk}^{abc} \lambda_{bc}^{jk}$. Redundant orbital rotations $\{X_j^i\}$, $\{X_j^j\}$, and $\{X_b^a\}$ can be arbitrary antiHermitian matrix elements. The general expressions for the RDMs are the same as in the TD-OCCD(T) method (Pathak et al. (2021)).

3 Numerical results and discussion

Our numerical implementation has an interface with the Gaussian09 program (Frisch et al. (2009)) for checking ground state energy with the standard Gaussian basis results. We study BH molecule with double- ζ plus polarization (DZP). We have

TABLE 1 Comparison of the ground state energy of BH ($r_e=2.4$ bohr) molecule in DZP basis^a.

Method	This work	References
OCCD ^b	- 25.225 591 67	- 25.225 592 Bozkaya and Schaefer (2012)
OCCD(T) ^b	- 25.226 913 29	- 25.226 913 Bozkaya and Schaefer (2012)
OCCD ^c	- 25.178 285 70	- 25.178 286 Krylov et al. (1998)
OCCD(T) ^c	- 25.178 301 00	

^aGaussian09 program ([Frisch et al. \(2009\)](#)) is used to generate the required one-electron, two-electron, and overlap integrals, required for the imaginary time propagation of EOMs in the orthonormalized Gaussian basis. A convergence cut-off of 10^{-15} Hartree of energy difference is chosen in subsequent time steps.

^bSix electrons correlated within the full basis set.

^cSix electrons correlated within the six optimized active orbitals.

reported ground state energy computed by propagating in the imaginary time for OCCD and OCCD(T) methods in [Table 1](#) and compared those with the optimized double and asymmetric triple excitation corrections for the orbital-optimized doubles method of [Bozkaya et al.](#), [Bozkaya and Schaefer \(2012\)](#). We also compare our OCCD ground state energy result with [Krylov et al.](#), [Krylov et al. \(1998\)](#) within the chosen active space of six electrons correlated among the six optimized active orbitals. We obtained a perfect agreement for all available values.

We have used a spherical-finite-element-discrete-variable representation (FEDVR) basis for representing orbital functions, [Sato et al. \(2016\)](#); [Orimo et al. \(2018\)](#) $\chi_{klm}(r, \theta, \psi) = \frac{1}{r} f_k(r) Y_{lm}(\theta, \phi)$ where Y_{lm} and $f_k(r)$ are spherical harmonics and the normalized radial-FEDVR basis function, respectively. The expansion of the spherical harmonics continued up to the maximum angular momentum L_{\max} , and the radial FEDVR basis supports the range of radial coordinate $0 \leq r \leq R_{\max}$ with $\cos^{1/4}$ mask function used as an absorbing boundary for avoiding unphysical reflection from the wall of the simulation box. We have used $l_{\max} = 72$, and the FEDVR basis supporting the radial coordinate $0 < r < 300$ using 78 finite elements each containing 25 DVR functions. The absorbing boundary is switched on at $r = 180$ in all our simulations. The Fourth-order exponential Runge-Kutta method ([Hochbruck and Ostermann \(2010\)](#)) is used to propagate the EOMs with 20000 time steps for each optical cycle. We run the simulations for a further 6,000 time steps after the end of the pulse. In all correlation calculations, eight electrons of $4s4p$ orbitals are considered as active and correlated among thirteen active orbitals. We report simulation results computed using a three-cycle laser pulse with a central wavelength of 800 nm having intensity 2×10^{14} W/cm² and a period of $T = 2\pi/\omega_0 \sim 2.67$ fs.

We report the time evolution of dipole moment of Kr in [Figure 1](#) and in [Figure 2](#) single electron ionization probability. Time-dependent dipole moment is evaluated as a trace $\langle \psi_p | \hat{z} | \psi_q \rangle D_p^q$ using 1RDMs. For the single electron ionization probability, we computed the probability of finding an electron

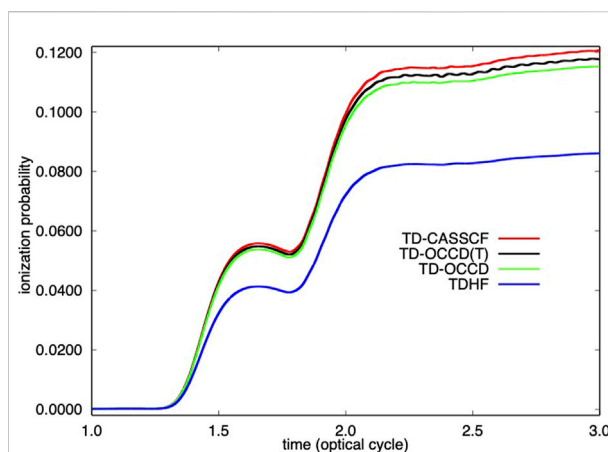


FIGURE 2

Time evolution of single ionization probability of Kr irradiated by a laser pulse with a wavelength of 800 nm and a peak intensity of 2×10^{14} W/cm² calculated with TDHF, TD-OCCD, TD-OCCD(T), and TD-CASSCF methods.

outside a sphere of a radius of 20 a.u. using RDMs defined in Refs. 19; 20; 37. We compare the results of TD-CASSCF, TD-OCCD(T), TD-OCCD, and TDHF methods.

We observe a substantial underestimation (both in [Figure 1](#), and [Figure 2](#)) by the TDHF method due to the lack of correlation treatment. All correlation methods perform according to their ability to treat electron correlation. We also computed results using the TD-OCCDT method but not reported here since those results are not identifiable from the TD-CASSCF results within the graphical resolution.

Next, we report high-harmonic generation in [Figure 3](#). It is calculated by squaring the modulus $I(\omega) = |a(\omega)|^2$ of the Fourier transform of the expectation value of the dipole acceleration with a modified Ehrenfest expression ([Sato et al. \(2016\)](#)). In panel (c) of [Figure 3](#), we plot the absolute relative deviation ($\delta(\omega)$, of the spectral amplitude $a(\omega)$ from the TD-CASSCF value for each method. All methods qualitatively predict similar HHG spectra with TDHF underestimates the spectral intensity. The relative deviation of results from TD-CASSCF ones follows the general trend TDHF > TD-OCCD > TD-OCCD(T) > TD-OCCDT, the same as what we observe for the time-dependent dipole moment and single ionization probability. We also simulated results with lower and higher intensity. However, the trend remains the same.

Finally, we make a tally of computational costs for all the methods considered in this article. All simulations performed using an Intel(R) Xeon(R) Gold 6,230 central processing unit (CPU) with 40 processors with a clock speed of 2.10 GHz, and report total simulations time in [Table 2](#). Further, we report a reduction in the computational cost for various TD-OCC methods relative to the TD-CASSCF. We see a massive 63% cost reduction for the TD-OCCD(T) method, which is larger

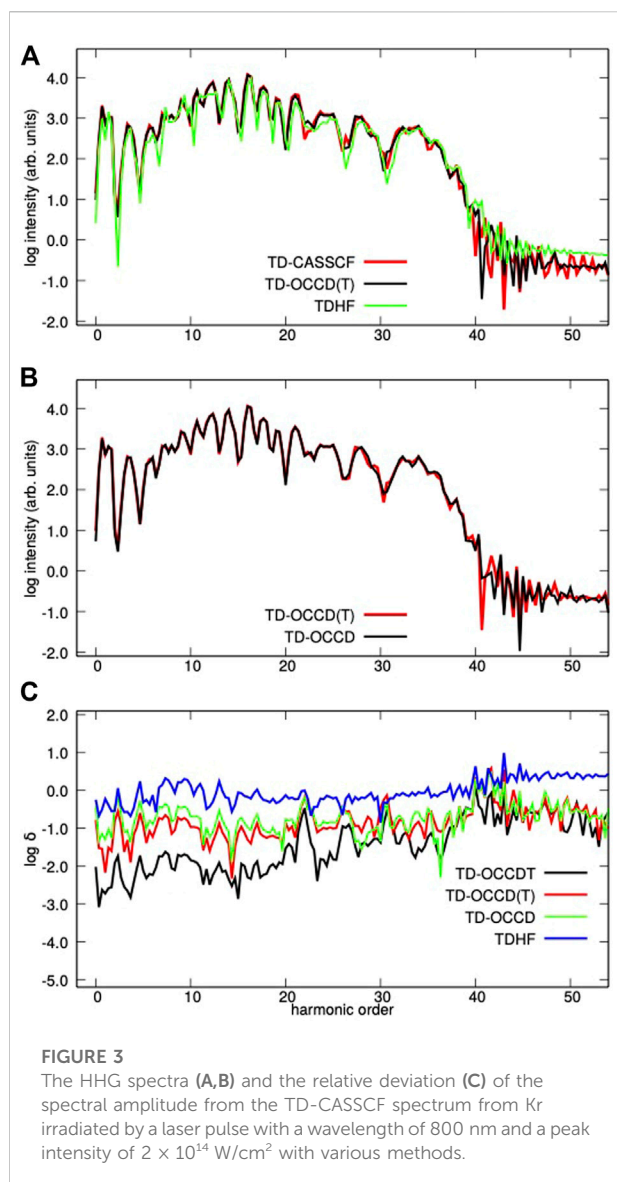


TABLE 2 Comparison of the total simulation time^a (in min) spent for TD-CASSCF, TD-OCCDT, TDCCD(T), and TD-OCCD methods.

Method	Time (min)	Cost reduction (%)
TD-CASSCF	47303	...
TD-OCCDT	19697	58
TD-OCCD(T)	17504	63
TD-OCCD	17494	63

^aTime spent for the simulation of Kr atom for 66000 time steps ($0 \leq t \leq 3.3T$) of a real-time simulation ($I_0 = 2 \times 10^{14}$ W/cm² and $\lambda = 800$ nm), using an Intel(R) Xeon(R) Gold 6230 CPU with 40 processors having a clock speed of 2.10 GHz.

than for the TD-OCCDT method (58%), and a minimal increase from the TD-OCCD method.

4 Concluding remarks

We have reported the formulation and implementation of the TD-OCCD(T) method. As the first application, we employed this method to study laser-driven dynamics in Kr exposed to an intense near-infrared laser pulse. We observe a 63% cost reduction in comparison to the TD-CASSCF method without losing much accuracy. Therefore, we conclude that TD-OCCD(T) method will certainly be beneficial in exploring highly accurate *ab initio* simulations of electron dynamics in larger chemical systems.

Data availability statement

The raw data supporting the conclusion of this article will be made available by the authors, without undue reservation.

Author contributions

HP and TS formulated the method. HP numerically implemented the method and performed simulations. All the authors analyzed the results and contributed to the submitted version of the manuscript.

Funding

This research was supported in part by a Grant-in-Aid for Scientific Research (Grants No. JP18H03891 and No. JP19H00869) from the Ministry of Education, Culture, Sports, Science and Technology (MEXT) of Japan. This research was also partially supported by JST COI (Grant No. JPMJCE1313), JST CREST (Grant No. JPMJCR15N1), and by MEXT Quantum Leap Flagship Program (MEXT Q-LEAP) Grant Number JPMXS0118067246.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated

organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

References

- Baker, S., Robinson, J. S., Haworth, C., Teng, H., Smith, R., Chirilă, C., et al. (2006). Probing proton dynamics in molecules on an attosecond time scale. *Science* 312, 424–427. doi:10.1126/science.1123904
- Bozkaya, U., and Schaefer, H. F., III (2012). Symmetric and asymmetric triple excitation corrections for the orbital-optimized coupled-cluster doubles method: Improving upon ccSD (t) and ccSD (t) λ. Preliminary application. *J. Chem. Phys.* 136, 204114. doi:10.1063/1.4720382
- Caillat, J., Zanghellini, J., Kitzler, M., Koch, O., Kreuzer, W., and Scrinzi, A. (2005). Correlated multielectron systems in strong laser fields: A multiconfiguration time-dependent Hartree-Fock approach. *Phys. Rev. A . Coll. Park.* 71, 012712. doi:10.1103/physreva.71.012712
- Corkum, P. á., and Krausz, F. (2007). Attosecond science. *Nat. Phys.* 3, 381–387. doi:10.1038/nphys620
- [Dataset] Frisch, M., Trucks, G., Schlegel, H., Scuseria, G., Robb, M., Cheeseman, J., et al. (2009). *Gaussian 09, revision d. 01*.
- Haxton, D. J., Lawler, K. V., and McCurdy, C. W. (2011). Multiconfiguration time-dependent Hartree-Fock treatment of electronic and nuclear dynamics in diatomic molecules. *Phys. Rev. A . Coll. Park.* 83, 063416. doi:10.1103/physreva.83.063416
- Haxton, D. J., and McCurdy, C. W. (2015). Two methods for restricted configuration spaces within the multiconfiguration time-dependent Hartree-Fock method. *Phys. Rev. A . Coll. Park.* 91, 012509. doi:10.1103/physreva.91.012509
- Hochbruck, M., and Ostermann, A. (2010). Exponential integrators. *Acta Numer.* 19, 209–286. doi:10.1017/s0962492910000048
- Hochstuhl, D., and Bonitz, M. (2011). Two-photon ionization of helium studied with the multiconfigurational time-dependent Hartree-Fock method. *J. Chem. Phys.* 134, 084106. doi:10.1063/1.3553176
- Hoodbhoy, P., and Negele, J. W. (1978). Time-dependent coupled-cluster approximation to nuclear dynamics. i. application to a solvable model. *Phys. Rev. C* 18, 2380–2394. doi:10.1103/physrevc.18.2380
- Hoodbhoy, P., and Negele, J. W. (1979). Time-dependent coupled-cluster approximation to nuclear dynamics. ii. general formulation. *Phys. Rev. C* 19, 1971–1982. doi:10.1103/physrevc.19.1971
- Huber, C., and Klamroth, T. (2011). Explicitly time-dependent coupled cluster singles doubles calculations of laser-driven many-electron dynamics. *J. Chem. Phys.* 134, 054113. doi:10.1063/1.3530807
- Itatani, J., Levesque, J., Zeidler, D., Niikura, H., Pépin, H., Kieffer, J. C., et al. (2004). Tomographic imaging of molecular orbitals. *Nature* 432, 867–871. doi:10.1038/nature03183
- Kato, T., and Kono, H. (2004). Time-dependent multiconfiguration theory for electronic dynamics of molecules in an intense laser field. *Chem. Phys. Lett.* 392, 533–540. doi:10.1016/j.cplett.2004.05.106
- Krausz, F., and Ivanov, M. (2009). Attosecond physics. *Rev. Mod. Phys.* 81, 163–234. doi:10.1103/revmodphys.81.163
- Krylov, A. I., Sherrill, C. D., Byrd, E. F., and Head-Gordon, M. (1998). Size-consistent wave functions for nondynamical correlation energy: The valence active space optimized orbital coupled-cluster doubles model. *J. Chem. Phys.* 109, 10669–10678. doi:10.1063/1.477764
- Kümmel, H. G. (2003). A biography of the coupled cluster method. *Int. J. Mod. Phys. B* 17, 5311–5325. doi:10.1142/s0217979203020442
- Kvaal, S. (2012). *Ab initio* quantum dynamics using coupled-cluster. *J. Chem. Phys.* 136, 194109. doi:10.1063/1.4718427
- Miyagi, H., and Madsen, L. B. (2013). Time-dependent restricted-active-space self-consistent-field theory for laser-driven many-electron dynamics. *Phys. Rev. A . Coll. Park.* 87, 062511. doi:10.1103/physreva.87.062511
- Miyagi, H., and Madsen, L. B. (2014). Time-dependent restricted-active-space self-consistent-field theory for laser-driven many-electron dynamics. ii. extended formulation and numerical analysis. *Phys. Rev. A . Coll. Park.* 89, 063416. doi:10.1103/physreva.89.063416
- Nascimento, D. R., and DePrince, A. E., III (2016). Linear absorption spectra from explicitly time-dependent equation-of-motion coupled-cluster theory. *J. Chem. Theory Comput.* 12, 5834–5840. doi:10.1021/acs.jctc.6b00796
- Nest, M., Klamroth, T., and Saalfrank, P. (2005). The multiconfiguration time-dependent Hartree-Fock method for quantum chemical calculations. *J. Chem. Phys.* 122, 124102. doi:10.1063/1.1862243
- Orimo, Y., Sato, T., Scrinzi, A., and Ishikawa, K. L. (2018). Implementation of the infinite-range exterior complex scaling to the time-dependent complete-active-space self-consistent-field method. *Phys. Rev. A . Coll. Park.* 97, 023423. doi:10.1103/physreva.97.023423
- Pathak, H., Sato, T., and Ishikawa, K. L. (2020a). Study of laser-driven multielectron dynamics of ne atom using time-dependent optimised second-order many-body perturbation theory. *Mol. Phys.* 118, e1813910. doi:10.1080/00268976.2020.1813910
- Pathak, H., Sato, T., and Ishikawa, K. L. (2021). Time-dependent optimized coupled-cluster method for multielectron dynamics iv: Approximate consideration of the triple excitation amplitudes. *J. Chem. Phys.* 154, 234104. doi:10.1063/5.0054743
- Pathak, H., Sato, T., and Ishikawa, K. L. (2020b). Time-dependent optimized coupled-cluster method for multielectron dynamics. ii. a coupled electron-pair approximation. *J. Chem. Phys.* 152, 124115. doi:10.1063/1.5143747
- Pathak, H., Sato, T., and Ishikawa, K. L. (2020c). Time-dependent optimized coupled-cluster method for multielectron dynamics. iii. a second-order many-body perturbation approximation. *J. Chem. Phys.* 153, 034110. doi:10.1063/5.0008789
- Pedersen, T. B., Kristiansen, H. E., Bodenstien, T., Kvaal, S., and Schøyen, Ø. S. (2021). Interpretation of coupled-cluster many-electron dynamics in terms of stationary states. *J. Chem. Theory Comput.* 17, 388–404. doi:10.1021/acs.jctc.0c00977
- Pedersen, T. B., and Kvaal, S. (2019). Symplectic integration and physical interpretation of time-dependent coupled-cluster theory. *J. Chem. Phys.* 150, 144106. doi:10.1063/1.5085390
- Pigg, D. A., Hagen, G., Nam, H., and Papenbrock, T. (2012). Time-dependent coupled-cluster method for atomic nuclei. *Phys. Rev. C* 86, 014308. doi:10.1103/physrevc.86.014308
- Raghavachari, K., Trucks, G. W., Pople, J. A., and Head-Gordon, M. (1989). A fifth-order perturbation comparison of electron correlation theories. *Chem. Phys. Lett.* 157, 479–483. doi:10.1016/s0009-2614(89)87395-6
- Sato, T., Ishikawa, K. L., Březinová, I., Lackner, F., Nagele, S., and Burgdörfer, J. (2016). Time-dependent complete-active-space self-consistent-field method for atoms: Application to high-order harmonic generation. *Phys. Rev. A . Coll. Park.* 94, 023405. doi:10.1103/physreva.94.023405
- Sato, T., and Ishikawa, K. L. (2013). Time-dependent complete-active-space self-consistent-field method for multielectron dynamics in intense laser fields. *Phys. Rev. A . Coll. Park.* 88, 023402. doi:10.1103/physreva.88.023402
- Sato, T., and Ishikawa, K. L. (2015). Time-dependent multiconfiguration self-consistent-field method based on the occupation-restricted multiple-active-space model for multielectron dynamics in intense laser fields. *Phys. Rev. A . Coll. Park.* 91, 023417. doi:10.1103/physreva.91.023417
- Sato, T., Orimo, Y., Teramura, T., Tugs, O., and Ishikawa, K. L. (2018a). “Time-dependent complete-active-space self-consistent-field method for ultrafast intense laser science,” in *Progress in ultrafast intense laser science XIV* (Berlin, Germany: Springer), 143–171.

Sato, T., Pathak, H., Orimo, Y., and Ishikawa, K. L. (2018b). Communication: Time-dependent optimized coupled-cluster method for multielectron dynamics. *J. Chem. Phys.* 148, 051101. doi:10.1063/1.5020633

Schonhammer, K., and Gunnarsson, O. (1978). Time-dependent approach to the calculation of spectral functions. *Phys. Rev. B* 18, 6606. doi:10.1103/physrevb.18.6606

Scuseria, G. E., and Schaefer, H. F., III (1987). The optimization of molecular orbitals for coupled cluster wavefunctions. *Chem. Phys. Lett.* 142, 354–358. doi:10.1016/0009-2614(87)85122-9

Shavitt, I., and Bartlett, R. J. (2009). *Many-body methods in chemistry and physics: MBPT and coupled-cluster theory*. Cambridge: Cambridge University Press.

Sherrill, C. D., Krylov, A. I., Byrd, E. F., and Head-Gordon, M. (1998). Energies and analytic gradients for a coupled-cluster doubles model using variational brueckner orbitals: Application to symmetry breaking in o 4+. *J. Chem. Phys.* 109, 4171–4181. doi:10.1063/1.477023

Watts, J. D., Gauss, J., and Bartlett, R. J. (1993). Coupled-cluster methods with noniterative triple excitations for restricted open-shell Hartree–Fock and other general single determinant reference functions. energies and analytical gradients. *J. Chem. Phys.* 98, 8718–8733. doi:10.1063/1.464480