Modeling electron dynamics coupled to continuum states in finite volumes with absorbing boundaries

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Abstract. Absorbing boundaries are frequently employed in real-time propagation of the Schrödinger equation to remove spurious reflections and efficiently emulate outgoing boundary conditions. These conditions are a fundamental ingredient for the calculation of observables involving infinitely extended continuum states in finite volumes. In the literature, several boundary absorbers have been proposed. They mostly fall into three main families: mask function absorbers, complex absorbing potentials, and exterior complex-scaled potentials. To date none of the proposed absorbers is perfect, and all present a certain degree of reflections. Characterization of such reflections is thus a critical task with strong implications for time-dependent simulations of atoms and molecules. We introduce a method to evaluate the reflection properties of a given absorber and present a comparison of selected samples for each family of absorbers. Further, we discuss the connections between members of each family and show how the same reflection curves can be obtained with very different absorption schemes.

1 Introduction

Continuum states play a central role in many physical processes involving atoms, molecules and extended low-dimensional systems. They are fundamental to the description of scattering problem with several applications ranging from ion collision and molecular damage [1] to electronic transport in molecular junctions [2]. Besides being important as final states there are situations where they play more articulated roles. This is the case, for instance, in strong laser physics where, in addition to direct ionization which involves the electronic continuum as final state, continuum states act as intermediate configurations before rescattering giving origin to peculiar features in photoemission [3,4] and high harmonic generation [5].

In many situations an explicit representation of the continuum wavefunctions is not necessary to calculate the relevant observables involving unbound states. Problems such as the description of ionization processes, the calculation of the absorption cross-section of atoms and molecules in the continuum, or electronic transport through molecular junctions, are just few prototypical examples. In these cases it is convenient to partition the space into two regions: an inner region containing the atomic or molecular system under study and an outer region defined as its complement. The problem is thus recast from finding an explicit representation of the continuum states to an implicit one where one solves the inner problem, imposing the appropriate conditions at the boundary between the two regions. Techniques used in these cases span from the R-matrix theory [9,10] to Green’s function methods [11] which have been applied both to the free [12–14] (e.g. for ionization processes) and bulk-continuum cases [15,16] (e.g. for electron transport).

In discrete and finite-size representations, either in real (e.g. finite differences) or Hilbert (e.g. basis set discretization) spaces, a detailed theoretical description of continuum states is a difficult task. The main difficulty arises from their extended nature spanning the whole space. In molecules the situation is further complicated by the presence of non-spherical multi-center potentials which quickly render any analytic approach impossible as the system grows in size. Several methods have been employed to address the problem; among the others, we mention the iterative Schwinger method [6], and B-spline expansions [7,8].
states is not needed. Their simplicity of implementation and limited computational cost make them, in many situations, preferable to the previously mentioned approaches. In this work we focus on three families of absorbers: mask function absorbers (MFAs), complex absorbing potentials (CAPs), and absorbers derived from smooth-exterior complex scaling (SES).

Mask function absorbers are the simplest type of absorber. At each time step one multiplies the wavefunction by a mask function, which has values between zero and one depending on the region in space. The function is generally chosen to be 1 within an inner region, then decay to zero near the boundary of the simulation box so as to damp any part of the wavefunction extending towards the boundary. MFAs have been employed in many different situations. We mention strong laser field studies on high harmonic generation [17], electron and proton emission [18], and above-threshold ionization [19]. They are the cornerstone of split-domain propagation schemes [20,21] where they act as matching layers between two different domains. In the literature they have been mainly associated with few-body problems (two or three particles), but in recent works they have also been used to study electron photoemission in multi-electron systems combined with time-dependent density functional theory [22–24].

The complex absorbing potential is, by far, the most popular type of absorber. The method consists in altering the Hamiltonian of a system by adding an artificial complex potential which is non-zero in a small portion of the simulation box close to the boundaries. Time propagation with this modified Hamiltonian results in wavepacket absorption or injection from the boundaries. Following their introduction as negative imaginary potentials in the work of Neuhauser and Baer [25], CAPs have been applied to a wealth of different situations ranging from relaxation of holes states in clusters [26] to charge injection for transport problems [27,28]. We refer the reader to an excellent review on the subject for further information [29].

The complex scaling method was originally developed to calculate the properties of metastable states or resonances. The method has a solid theoretical background with a long history of development, beginning with the original theorems by Aguilar and Combes [30] and Balslev and Combes [31]. The method continues to be the subject of much interest and has since been adapted to scalable computational approaches including configuration interaction [32] and DFT [33,34]. Complex scaling methods work by means of a scaling of the position variable of the Hamiltonian by a complex factor. This causes states that represent extended out-going waves or resonances to become normalizable and thus computationally accessible [35].

Exterior complex scaling (ECS), proposed by Simon [36], is a generalization where the scaling transformation is only applied outside a certain region rather than uniformly. ECS was originally introduced to avoid singularities in the Coulomb interaction at nuclear centers where the scaling method breaks down. In smooth-exterior scaling (SES) the scaling is governed by a smooth function to improve numerics.

Complex scaling methods can be applied to time-dependent problems as well as static ones. Under time-propagation, states extending into the complex-scaled (exterior) region will decay over time. The region thus acts as an absorber [37–39]. The method is also referred to as a reflection-free CAP (RF-CAP) even though it was shown to present reflections in combination with discretization in finite volumes [40]. It was later shown by Scrinzi [41] that such reflections can be reduced to machine precision in time propagations under strong infrared fields using a finite element plus a single infinite element approach.

As far as strictly finite or bounded volumes are employed, none of the ABs known in the literature is numerically reflection-free and, depending on the type of AB and the size of the buffer region, their absorption properties vary as a function of the impinging wavefunction’s kinetic energy. Characterization and control of the reflection properties of an AB is thus an important task with implications to many situations involving an implicit description of continuum states. By employing tailored Gaussian wavepackets we are able to map the reflection properties of a given absorber and display it as a function of different parameters. In the present work, we use these properties to illustrate differences and analogies between some of the most used absorbers.

In some cases different types of AB behave very similarly, indicating a degree of relation. In fact, it can be shown that all the three families are connected to one another as in Figure 1. While each of the single connections in the figure has been separately discussed in the literature, in this paper we aim at presenting the full picture in an homogeneous framework. Furthermore, besides illustrating the nature of each link, we show how it is possible to obtain the same absorption properties with very different types of ABs.

The paper is organized as follows. We summarize the scattering theory from complex potentials in Section 2 and
introduce the quantities used to assess the reflection properties of ABs in finite volumes. In Section 3 we describe the details of the numerical calculations. We discuss the reflection properties of MFAs in Section 4. In Section 5 we present the reflection properties of polynomial and \sin^2 CAPs for different potential heights and illustrate by calculating the absorption cross-section of a one-dimensional hydrogen atom. The connection between MFAs and CAPs is discussed in Section 6. Finally in Section 7 we discuss SES in connection with CAPs.

2 Theory

Owing to the position they cover in Figure 1, CAPs play a central role in the present work. Below we briefly review the scattering theory from complex potentials following the review by Muga et al. [29]. Besides introducing the reader to the main theoretical formalism the purpose of the present section is that of formally derive \( \epsilon(k_0) \), a quantity assessing the error committed by an AB that is suitable for numerical evaluation in finite volumes. Atomic units will be used throughout \((m_e = e = \hbar = 1)\).

In one-dimension, the stationary Schrödinger equation for a free electron in the presence of a CAP may be written as

\[
H \psi(x) = H_0 \psi(x) + V_{\text{CAP}}(x) \psi(x) = E \psi(x) \quad (1)
\]

with \(H_0\) being the physical Hamiltonian and \(V_{\text{CAP}}(x)\) an absorbing potential with support in \([0, L]\) as in Figure 2a. For simplicity we consider here the case where \(H_0\) describes free electrons \(H_0 = -\nabla^2/2\).

The non-normalizable solutions of (1) corresponding to stationary scattering states are obtained by imposing one of the following asymptotic boundary conditions,

\[
\psi^+(k, x) = \begin{cases} 
  e^{ikx} + R^r(k)e^{-ikx} & x \to -\infty \\
  T^l(k)e^{ikx} & x \to \infty
\end{cases} \quad (2)
\]

\[
\psi^-(k, x) = \begin{cases} 
  T^r(k)e^{-ikx} + R^l(k)e^{ikx} & x \to -\infty \\
  e^{-ikx} & x \to \infty
\end{cases} \quad (3)
\]

For \(k > 0\), they enforce the condition corresponding respectively to a right-travelling wave \(\psi^+(k, x)\) and a left-travelling wave \(\psi^-(k, x)\). The functions \(R^l,r(k)\) and \(T^l,r(k)\) indicate the (left/right) reflection and transmission coefficients. These coefficients are, in general, complex functions of \(k\) and their square modulus represents the probability to reflect and transmit a wave with a given momentum. If the initial wavepacket is localized on the left side of the potential we can simplify the notation by imposing only the left asymptotic condition, discarding the right quantities.

By imposing the left asymptotic boundary conditions of (2) on the time-dependent Schrödinger equation

\[
i \frac{\partial \psi(t, x)}{\partial t} = H \psi(t, x), \quad (4)
\]
a unique solution is obtained for every single initial condition. For a right-moving wavepacket \(\psi_0(x)\) initially localized to the left region \([-\infty, 0]\), this solution is given by

\[
\psi(t, x) = \int_0^{\infty} dk \psi^+(k, x)e^{-ik^2t/2}\psi_0(k), \quad (5)
\]

where space is restricted to \(x \in [-\infty, 0] \cup [L, \infty]\), and \(\psi_0(k)\) represents the Fourier transform of \(\psi_0(x)\). Here, \(U(t) = e^{-ik^2t/2}\) is the time evolution operator associated with the free Hamiltonian \(H_0\). Note that, although reflection and absorption coefficients are strictly defined for \(k > 0\), they can be analytically continued to negative \(k\) and the integral (5) can be extended to the whole real axis.

The most appropriate quantity to assess the absorption properties of a CAP is the survival probability \(S(k)\). It is defined as the sum of transmission \(|T(k)|^2\) and reflection \(|R(k)|^2\) probabilities: \(S(k) := |T(k)|^2 + |R(k)|^2\). According to this definition, an ideal absorber, preventing reflections regardless of the incoming wavevector, would have \(S(k) = 0\) for all \(k > 0\). As already mentioned, none of the known CAPs is free from reflections, and the evaluation of \(S(k)\) is of great importance for practical applications.

While from (5) it is possible to evaluate the time evolution of an arbitrary wavepacket knowing the reflection and transmission coefficients (and thus \(S(k)\)), the calculation of these coefficients starting from the knowledge of \(\psi(t, x)\) is a more complicated task. A widely used approach consists in calculating \(R(k)\) and \(T(k)\) by numerically solving the static one-dimensional scattering problem [42–44].
This solution is however difficult to apply in the present work where we aim at addressing ABs expressed with quite different formulations. We instead use a different approach based on direct time propagation in a finite volume.

We focus on the situation where the solution of (1) is confined to the volume $[-X, L]$ like in Figure 2b. In this case it is customary to impose zero boundary conditions at the border of the region: i.e. $\psi(t, x = -X) = \psi(t, x = L) = 0$ for all $t$. The CAP therefore effectively includes an infinite barrier at $x = L$. CAPs so defined are also known as Type I potentials [29,45]. A wavefunction is thus either absorbed or reflected, the transmission coefficient $T(k)$ is therefore equal to zero and the survival probability is just equal to the square modulus of the reflection coefficient: $S(k) = |R(k)|^2$.

The situation is similar to what illustrated in Figure 2b: a left-moving wavepacket $\psi_0(x)$ initially localized in $[-X, 0]$ propagates as the sum of a free propagating and a reflected wave

$$\psi(t, x) = \psi_0(t, x) + \psi_R(t, x)$$

$$= \int_{-\infty}^{\infty} \frac{dk}{\sqrt{2\pi}} e^{ikx} e^{-ik^2t/2} \psi_0(k)$$

$$+ \int_{-\infty}^{\infty} \frac{dk}{\sqrt{2\pi}} e^{-ikx} R(k) e^{-ik^2t/2} \psi_0(k). \quad (6)$$

To obtain (6) we extended the integral in (5) over negative $k$ and imposed the normalization on incoming and outgoing waves.

All the informations regarding the reflection coefficient, and therefore $S(k)$, are contained in $\psi_R(t, x)$. The reflection error [46], defined as

$$\tilde{\epsilon}(t) = \int_{-X}^{0} dx [\psi(t, x) - \psi_0(t, x)]^2$$

$$= \int_{-X}^{0} dx [\psi_R(t, x)]^2, \quad (7)$$

is the closest quantity related to $\psi_R(t, x)$ that can be easily accessed from the direct time evolution of $\psi(t, x)$. In the long time limit $t \geq \tau$, when the free propagating wavefunction has left the simulation region, it represents a direct measure of reflections. Provided that the initial wavepacket $\psi_0(k)$ is well localized around $k_0$, it directly approximates $S(k_0)$ and, in the limit where $\psi_0(k)$ is a Dirac delta function centered at $k_0$ and $X \to \infty$, it coincides with the survival probability: $\tilde{\epsilon}(t \geq \tau) = S(k_0)$.

The time evolution described by (6) is defined only outside the CAP region, i.e. in $[-X, 0]$, hence the integration range in the definition of $\tilde{\epsilon}(t)$ in (7). For times bigger than $\tau$ such that $\psi_0(t = \tau, x)$ is mostly localized around this region and $\psi_R(t = \tau, x)$ is mostly localized in $[-X, 0]$, the integral can be extended to the whole range $[-X, L]$. We can thus extend the definition of (7) to the entire simulation box including the absorbing boundary region. For a given initial wavepacket $\psi_0(x)$ localized in $[-X, 0]$ with a peak around $k_0$, the reflection error can be redefined as

$$\epsilon(k_0) = \int_{-X}^{L} dx [\psi(t = \tau, x) - \psi_0(t = \tau, x)]^2, \quad (8)$$

where $\tau$ must be chosen big enough to satisfy the localization conditions on $\psi_0(t, x)$ and $\psi_R(t, x)$, but smaller than the time needed for $\psi_R(t, x)$ to bounce at $x = -X$ back into the absorption region.

This extended definition is preferable to that of (7) as it correctly handles pathological situations such as the one occurring when the CAP is just a perfect reflecting wall placed at $x = L$. In this case a normalized wavepacket would just bounce back at $x = L$ and reach $[-X, 0]$ after a time that depends on $k_0$ and on the width of the absorber region $L$. The unmodified reflection error $\epsilon(t)$ would then report either high or negligible values depending on such time while $\epsilon(k_0)$ would be a constant strictly equal to 1 hence indicating maximum reflection.

3 Results

In order to evaluate the reflection error for a range of energies we need to specify a family of initial wavefunctions $\psi_0(x)$. The Gaussian wavepackets

$$G(x, x_0, k_0, \sigma) = \pi^{-\frac{1}{4}} \sigma^{-\frac{1}{2}} e^{-\frac{(x-x_0)^2}{2\sigma^2}} + ik_0 x \quad (9)$$

are a natural choice, being localized in both real and Fourier space. Similar choices have been made in the literature [46–48]. Here (9) describes a normalized Gaussian wavepacket with initial velocity $k_0$, width $\sigma$, centered in $x_0$. We specifically choose the parameters

$$\sigma = 4\sqrt{2}/k_0, \quad (10)$$

$$x_0 = -3\sigma = -12\sqrt{2}/k_0. \quad (11)$$

Given an absorber width $L$, we then sample the reflection error $\epsilon(k_0, L)$ by performing time evolutions of Gaussian wavepackets with different $k_0$.

In each propagation $G(x, x_0, k_0, \sigma)$ is centered in $x_0 = -3\sigma$ and is evolved for a total time $\tau = 2(3\sigma + L)/k_0$ in a simulation box with $X = 6\sigma$. The total propagation time is chosen so that the free wavepacket travels a distance of $d = 6\sigma + 2L$. During the time evolution, $G(x, x_0, k_0, \sigma)$ spreads with an asymptotic velocity that depends on $\sigma$: $v_\sigma = 1/\sqrt{2\sigma}$. The wavepacket is thus characterized by the two velocities $k_0$ and $v_\sigma$. The choice $\sigma(k_0) = 4\sqrt{2}/k_0$ therefore ensures an expansion velocity slower than the translational one: $v_\sigma = k_0/8$. The chosen parameters also impose negligible left-moving ($k < 0$) components and ensure that the wavepackets have well-defined energies also when the energy is close to 0. With these choices the only free parameters left are $k_0, L$.

All the numerical calculations have been performed using finite differences in real space and real time with the Octopus [49–51] and QuantumPy 1 software packages. The spatial coordinates are discretized on a grid with spacing $\Delta x = 0.1$ a.u. and the time step $\Delta t = 0.01$ a.u. ($\Delta t = 0.05$ a.u. and time step $\Delta t = 0.001$ a.u. for kinetic energies greater than 2 KeV). These parameters secure an

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1. The code is released under GNU public license at https://github.com/gimunu/QuantumPy
accurate description of the time propagation for all the wavepackets considered. Additionally, they are directly comparable to the typical ones used in three-dimensional atomic and molecular calculations. To take a step further in the direction of application to real systems, in what follows, we express the reflection error $\epsilon(E, L)$ as a function of the wavepacket kinetic energy $E = 5k_0^2/4$ (in eV) and absorber width $L$ (in atomic units).

### 4 Mask function absorbers (MFAs)

MFAs [17] are commonly used in numerical propagation schemes where the infinitesimal time-evolution operator $U(t + \Delta t, t)$, connecting $t$ to $t + \Delta t$, is repeatedly applied to an initial wavefunction. One chooses a mask function $0 \leq M(x) \leq 1$ and apply it multiplicatively on each time step, i.e.

$$\psi(x, t + \Delta t) = M(x)U(t + \Delta t, t)\psi(x, t).$$

If the function $M(x)$ is chosen to smoothly decay from one in the buffer region $[0, L]$, the iterative application of (12) results in a damping of any part of the wave extending into this region. In order to absorb, the mask function has to be smaller than 1 but is free to assume any value from 0 to 1 at the border $x = L$. The resulting absorption properties depend on the specific functional form of $M(x)$.

In Figure 3 we show the reflection error $\epsilon(E, L)$ as function of kinetic energy $E$ and boundary width $L$ for a mask function defined as

$$M(x) = \begin{cases} 1 & \text{if } x < 0 \\ 1 - \sin^2 \left( \frac{\pi x}{2L} \right) & \text{if } 0 \leq x \leq L. \end{cases}$$

Reflections smoothly decay from almost unity at low energies to negligible values for high energies. The range of energies with minimal reflection is connected to the boundary width $L$: the greater $L$, the wider the region of minimal reflection. The high reflectivity for low energies can be diminished by increasing $L$, but slowly moving wavepackets, associated with large wavelengths, are difficult to remove with finite size absorbers.

Besides being easy to implement, the main attraction of MFAs derives from the central role they play in split-domain propagation schemes [20, 21]. At the foundation of these schemes lies the idea that it is always possible to perform a spatial partitioning of a wavefunction as a sum of two partially overlapping components

$$\psi(x, t) = M(x)\psi(x, t) + (1 - M(x))\psi(x, t).$$

In this partitioning, one piece of the wavefunction is localized in $[-\infty, L]$ and the other in the partially overlapping region $[0, \infty]$. The two parts can then be separately propagated with different Hamiltonians using different approximations. This decomposition is at the core of a recent method developed for the calculation of electron photoemission in molecular systems [22]. For further details we refer to reference [52].

### 5 Complex absorbing potentials (CAPs)

ABs based on CAPs are constructed by adding an artificial complex potential $V_{\text{CAP}}(x)$ to the physical Hamiltonian $H_0$ as described by (1). By virtue of this change, the new Hamiltonian $H$ is no longer Hermitian, and the associated time propagation operator,

$$U_{\text{CAP}}(t + \Delta t, t) = e^{-iH\Delta t} = e^{-i[H_0+V_{\text{CAP}}(x)]\Delta t},$$

becomes non-unitary. When $V_{\text{CAP}}(x)$ is chosen to be non-zero only in the buffer region $[0, L]$, this non-unitarity becomes localized and, during time propagation, the wavefunctions have their norm altered (increased or decreased) when overlapping with that region. In order to absorb, the imaginary part of $V_{\text{CAP}}(x)$ must take negative values since the presence of a negative imaginary potential at the exponent of the propagation operator of (15) induces an exponential damping of wavepackets.

Among the many possible choices, the monomial CAP represents one of the most discussed in the literature [29, 46, 48, 53]. It consists of a purely imaginary potential of the form

$$V_{\text{CAP}}(x) = V_m(x) = \begin{cases} 0 & \text{if } x < 0 \\ i\alpha x^b & \text{if } 0 \leq x \leq L. \end{cases}$$

To act as an absorber, a monomial CAP must have $\alpha < 0$, while $b$ can be any number greater than one. The strength of induced damping depends on the kinetic energy of the colliding wavepacket and on the width of the absorption region as shown in Figure 4. In this case we select $\alpha = -0.003$ a.u. and $b = 3$.

Except for $L < 10$ a.u., the reflection error qualitatively follows the one previously discussed for the mask functions of Figure 3. For $L = 5$ a.u. the CAP is simply not large enough and at best capable of delivering a
Fig. 4. Reflection error $\epsilon(E, L)$ as a function of energy $E$ and boundary width $L$ for a monomial complex potential with $b = 3$ and $\alpha = -0.003$ a.u. The functional form of $V_{\text{CAP}}$ on the imaginary axis is shown in the inset.

minimum 10% reflection for $E \approx 5$ eV. Compared to the MFA, the monomial CAP is less sensitive to increase of $L$, and the reflection error $\epsilon(E, L)$ converges quickly to an asymptotic shape as $L \to \infty$. This is because the shape of the absorbing potential $\alpha x^3$ does not change with $L$. Once $L$ is large enough that a given wave packet does not reach $x = L$ due to the height of the potential, further increase of $L$ cannot reduce the reflection. The relationship between CAPs and mask functions will be studied in the next section. Generally if one lets the CAP diverge at the boundary, it will behave roughly like a mask function approaching zero.

In order to disentangle the effect of the boundary width $L$ from that of the maximum value of $V_{\text{CAP}}$ in $[0, L]$ we study the case of a $\sin^2$ potential

$$V_{\text{CAP}}(x) = V_s(x) = \begin{cases} 0 & \text{if } x < 0 \\ i \eta \sin^2 \left( \frac{x \pi}{2L} \right) & \text{if } 0 \leq x \leq L. \end{cases}$$

(17)

Compared with the CAP in (16) this potential smoothly increases from zero to its maximum value at the edge of the box $V_s(L) = \eta$, and, like the monomial one, it absorbs only for negative values of $\eta$.

The reflection error for a $\sin^2$ CAP with $\eta = -0.2$ is shown in Figure 5. Compared with the monomial CAP in Figure 4 the survival probability displays a quite different behavior presenting a flat minimum centered around $E = 10$ eV increasingly extending with $L$. The low energy absorption is globally better than the monomial CAP but quickly deteriorates for high kinetic energies, and $\epsilon(E, L)$ is more sensitive to the changes of $\eta$. The failure at high energies occurs when the wavepackets are fast enough to reflect at $x = L$ and exit the absorbing region before the CAP can absorb them.

The reflection error is shown in Figure 6 for selected values of the potential height $\eta$. For small $\eta$ the reflection error has a peaked minimum that widens up with $L$.

As $\eta$ increases, the absorption window widens and the minimum moves to higher energies quickly overrunning the plot range.

We conclude the current section by presenting an example where CAPs can be used in time-dependent calculations. The calculation of the continuous part of the absorption spectrum of a hydrogen atom is an illustrative application of CAPs to real-time propagation of the Schrödinger equation. We consider the simple case of a one-dimensional hydrogen atom with a softened Coulomb interaction described by the Hamiltonian

$$H_0 = -\frac{1}{2} \frac{\partial^2}{\partial x^2} - \frac{1}{\sqrt{x^2 + 2}}.$$  

(18)
Fig. 7. One-dimensional hydrogen atom absorption cross-section (bilogarithmic scale) in the continuum. The plot has been shifted so that the zero of the energy scale corresponds to hydrogen ionization threshold 13.6 eV. Dashed lines represent the spectrum for hard-wall boundary conditions (cyan) and a converged exact solution calculated with a box of size 6000 a.u. (yellow). Solid lines are the spectra calculated with a $\sin^2$ CAP defined by (17), with $L = 10$ a.u. and having different heights: $\eta = -0.01$ a.u. (red), $\eta = -0.2$ a.u. (green), $\eta = -1$ a.u. (blue), and $\eta = -10$ a.u. (purple).

In the spectrum, the absorption lines appear in correspondence with allowed transitions from ground to excited states. They are positioned at photon energies precisely equal to the energy difference between such states, and their strength is proportional to the dipole matrix element connecting the states. At photon energies above the first ionization threshold, the spectrum thus involves matrix elements connecting to continuum states. A good description of the spectrum in this region is therefore ultimately linked to a good representation of continuum states in the same energy region.

Instead of directly evaluating the dipole matrix elements we here follow the time evolution of the density. The optical absorption properties can be easily calculated, in the linear response, by analyzing the time dependence of the dipole moment subject to a small initial kick perturbation (see for instance Chap. 7 of Ref. [54]). To this end we discretize spatial and temporal coordinates with $\Delta x = 0.1$ a.u., $\Delta t = 0.01$ a.u., and perform time propagations with different boundary absorbers for a maximum time of 1000 a.u.

First we analyze the situation where no CAP is present. To this end we perform the time evolution in a 60 a.u. box with hard wall boundary conditions. The resulting spectrum centered in the continuum region (i.e. for photon energies above the ionization threshold) is shown in Figure 7 (cyan dashed line over pale green area). It is highly structured and presents a series of peaks where a smooth behavior should be expected. These peaks can be associated with transitions to unphysical box-states, i.e. the eigenstates associated with the infinite square well embedding the system. The continuum, constituted by such discrete states, is thus strongly dependent on the box size and far from being a good approximation for the real one.

In order to avoid reflection, or equivalently to increase the number of box levels per unit of energy, a much larger simulation box has to be employed. In the current case, a box of 6000 a.u. is enough to avoid reflections and completely contain the wavefunction at the end of the time propagation. In contrast to the previous case, here the spectrum (yellow dashed line over pale yellow area) presents a smooth behavior characteristic of the continuum region. The goal of a CAP would be to reproduce the same spectrum with a much smaller simulation box. To this end we fix the box size to 60 a.u. and recalculate the spectrum by placing two $L = 10$ a.u. wide $\sin^2$ CAPs at the edges and vary $\eta$. For $\eta = -0.01$ a.u. the spectrum presents a marked oscillating structure with peaks located at box-state transition energies. The effect of this CAP is here limited to a smearing between peaks plus a small increase of the absorption. For $\eta = -0.2$ a.u. the spectrum becomes smoother for low energies. In this region it lies on top of the exact solution until it departs from it as the energy becomes larger than 10 eV. A further decrease of $\eta$ moves the smooth region forward to higher energies leaving oscillations in the lower energy end.

None of these CAPs is good enough to reproduce the exact results for the whole range of energies but only in a limited region that changes with $\eta$. This behavior corresponds well to the one we observed for the reflection error $\epsilon(E, L)$ in Figures 5 and 6. We therefore conclude that $\epsilon(E, L)$, although being an approximation to the survival probability $S(E, L)$, can provide useful information to assess and control reflections in practical calculations.

6 Connecting MFAs and CAPs

This section is devoted to the discussion of the mathematical connections $a$ and $b$ in the scheme of Figure 1. The existence of such a connection has been known in the literature since long time ago [20,55] but, to the best of the authors’ knowledge, has never been presented in a unified framework.

We begin with the problem of linking an MFA to a CAP and therefore with the problem of casting the first into the formalism of the second. The solution can be simply obtained by comparing the time propagation operators in the two formalisms. Starting from (12) we can define an infinitesimal mask function time propagation operator as

$$U_M(t + \Delta t, t) = M(x)U(t + \Delta t, t) = M(x)e^{-iH_0\Delta t}$$

$$= e^{\tilde{M}(x)}e^{-iH_0\Delta t},$$

where $H_0$ is the physical Hamiltonian and

$$\tilde{M}(x) = \ln |M(x)|$$

is a logarithmic mask function. By requiring the two propagators in (19) and (15) to be equal it is possible to derive an explicit form for the CAP $V_{\text{CAP}}(x)$ associated with the
mask function $M(x)$. The resulting equation can be expanded using the Baker-Campbell-Hausdorff formula for exponential operators (in Appendix) and, to first order in $\Delta t$, this CAP can be written as,

$$
V_{\text{CAP}}^{(1)}(x) = \frac{i}{\Delta t} \tilde{M} + \frac{1}{2!} \{\tilde{M}(x), H_0\} + \frac{1}{12} \{\tilde{M}(x), \{\tilde{M}(x), H_0\}\} - \frac{1}{720} \{\{\{H_0, \tilde{M}(x)\}, \tilde{M}(x)\}, \tilde{M}(x)\} + \ldots
$$

This expression can be simplified by evaluating the commutator $[H_0, \tilde{M}(x)]$ and the higher order ones. In one dimension, higher order commutators vanish above the second order in $\tilde{M}(x)$. The result is a closed formula for the mask function-CAP approximation to first order in $\Delta t$:

$$
V_{\text{CAP}}^{(1)}(x) = \frac{i}{\Delta t} \tilde{M}(x) - \frac{d\tilde{M}(x)}{dx} \frac{\partial}{\partial x} - \frac{1}{6} \left( \frac{d\tilde{M}(x)}{dx} \right)^2 - \frac{1}{2} \frac{d^2\tilde{M}(x)}{dx^2}.
$$

The mask function derived CAP therefore contains the differential operator $\partial/\partial x$ and is not purely imaginary, but has components also on the real axis. CAPs that are not strictly imaginary have been studied in the literature, and it has been pointed out that the inclusion of a real component in the potential may lead to enhanced absorption [29].

For small values of $\Delta t$ the CAP in (22) is dominated by the first term $i\tilde{M}(x)/\Delta t$ which is purely imaginary. For every mask function $0 \leq M(x) \leq 1$ this term is on the negative imaginary axis, and so $V_{\text{CAP}}^{(1)}(x)$ acts as an absorber.

In Figure 8a we present a comparison of the reflection error ($\epsilon(E, L)$ calculated with the MFAs defined in (13) (points) and the one obtained using the CAP of (22) (solid lines). The agreement between the curves for different $L$ underlines the good quality of the first-order approximation for the time step used. In this case $M(L) = 0$, and thus $\tilde{M}(x) \to -\infty$ as $x \to L$.

We now turn to the derivation of the inverse relation, i.e. the problem of expressing a CAP as a mask function. Once again, the correspondence is derived by requiring the evolution operators to be equal. Using Zassenhaus’ expansion [56] for $U_{\text{CAP}}(t + \Delta t, t)$ we have

$$
M_{\text{CAP}}(x) = e^{-iV_{\text{CAP}}(x)\Delta t} \times e^{\left[2\frac{dV_{\text{CAP}}(x)}{dx} \frac{\partial}{\partial x} + \frac{dV_{\text{CAP}}(x)}{dx} \frac{\partial^2}{\partial x^2}\right] \Delta t^2} \times \ldots
$$

Similar to the previous case, the CAP mask function is here complex-valued. This does, however, not affect the splitting property of (14) which also includes this situation. When $V_{\text{CAP}}(x)$ is a purely imaginary potential, the first-order term of (23),

$$
M_{\text{CAP}}^{(1)}(x) = e^{-i\Delta t V_{\text{CAP}}(x)},
$$

becomes a real function.

Fig. 8. Comparison of reflection errors $\epsilon(E, L)$ for the same absorber expressed as CAP or mask function. (a) The mask-function absorber of (13) (points) compared to its representation as CAP (22) (solid lines). The inset depicts $\tilde{M}(x)$; (b) sin$^2$ CAP as of (17) with $\eta = 0.2$ a.u. (points) expressed as a mask function through (24) (solid lines). In the inset the first-order mask function $M_{\text{CAP}}^{(1)}(x)$ is pictured.

A numerical comparison for a sin$^2$ CAP ($\eta = 0.2$ a.u.) expressed in the two formalisms is shown in Figure 8b. It shows that (24) yields two numerically identical absorbing properties and that, for practical applications, the two methods are equivalent. Further, the results indicate that the value of a mask function at the edge of the box is connected with the position of the minimum in $\epsilon(E, L)$. This can be easily understood by comparison with $V_{\text{CAP}}(L)$.

Finally, we observe that compared with the CAPs whose values may range from 0 to $-\infty$ in $[0, L]$, the mask function spans a much more limited range $[0, 1]$. As we will discuss in the next section, this fact has positive numerical implications possibly leading to stabler propagation schemes.

7 Smooth-exterior scaling (SES)

We conclude our inventory of ABs by discussing the effect of a smooth-exterior scaling of the Hamiltonian as a reflection absorber. This approach is based on a generalization of the original, uniform complex scaling method [30,31]. In SES the spatial coordinates of the Hamiltonian are rotated into the complex plane by an angle $\theta$ outside a certain region by means of the transformation [36,57,58]: $x \to F(x)$. This transformation has to be such that it asymptotically recovers the linear scaling transformation

$$
F(x) \to xe^{i\theta} \text{ for } x \to \infty.
$$

As can be easily shown, given a function $g(x)$ smoothly increasing from 0 to 1 close to the onset of the boundary
region \( x = 0 \), the family of transformations

\[
f(x) = \frac{dF(x)}{dx} = 1 + (e^{i\theta} - 1)g(x)
\]  

(26)

satisfies the limit condition of (25). For every choice of \( g(x) \), it is then possible to recover \( F(x) \) by direct integration of (26).

Following a SES transformation, the new Hamiltonian turns non-Hermitian and, like for CAPs, the time-evolution operator becomes non-unitary. It is further possible to push the connection with CAPs by defining a \( V_{\text{CAP}}^S(x) \) that added to the Hamiltonian \( H_0 \) implements the SES transformation [39]. The SES CAP is

\[
V_{\text{CAP}}^S(x) = V_0(x) + V_1(x) \frac{\partial}{\partial x} + V_2(x) \frac{\partial^2}{\partial x^2}
\]  

(27)

where

\[
V_0(x) = \frac{1}{4f^3(x)} \frac{d^2f(x)}{dx^2} - \frac{5}{8f^8(x)} \left( \frac{df(x)}{dx} \right)^2
\]  

(28)

\[
V_1(x) = \frac{1}{f^3(x)} \frac{df(x)}{dx}
\]  

(29)

\[
V_2(x) = \frac{1}{2} \left( 1 - \frac{1}{f^2(x)} \right)
\]  

(30)

derive from an evaluation of the kinetic operator on \( F(x) \) together with a scaling of the volume element \( dz = f(x)dx \).

The CAP defined in (27) contains the differential operators \( \partial/\partial x \) and \( \partial^2/\partial x^2 \) that make it actively change its shape as a wavepacket moves in. It constitute one of the simplest way to implement SES in real-space codes and explicitly illustrates the existence of the connection \( c \) in Figure 1. The existence of the inverse connection \( \partial/\partial c \) has been discussed in reference [59] indicating the possibility to map a given CAP with an SES transformation [60]. It is worth to note that such connection is limited to the absorption of outgoing wavepackets only, and that ECS provides more than just a simple absorber. In fact, a time propagation correctly accounting for the evolution of left and right eigenstates of the exterior complex scaled Hamiltonian would offer, in principle, the possibility to recover the electron dynamics in the outer region and allow incoming wavepackets to reenter the inner region [37,41].

A popular choice for the function \( g(x) \) in (26) is provided by

\[
g(x) = \frac{1}{2}(1 + \tanh[\lambda x]).
\]  

(31)

Integrating over \( x \) in (26) then yields

\[
F(x) = x + (e^{i\theta} - 1) \left( \frac{x}{2} + \frac{1}{2\lambda} \log[\cosh[\lambda x]] \right).
\]  

(32)

Similar SES has been employed in static calculations of resonance lifetimes [61] as well as boundary absorber in real-time propagations [40,46,62].

In Figure 9 (solid lines) we show the absorption properties of \( V_{\text{CAP}}^S(x) \) associated with a \( g(x) \) defined by (31) having \( \lambda = 1 \) and \( \theta = 0.2 \). The reflection error at any given \( L \) presents similar traits quickly decaying for large energies. As the energy is lowered, \( \epsilon(E, L) \) steeply increases and exceeds 1 indicating the presence of an instability region where the SES amplifies the norm instead of reducing it. This region is pushed towards lower energies as \( L \) increases; for \( L > 30 \) a.u. we observe numerical instability only for energies \( E < 0.063 \) eV.

The scaling transformation (32) implements a smooth transition of the wavefunction from \( \psi(x) \) for \( x \ll 0 \) to \( e^{i\theta/2}\psi(F(x)) \) for \( x \gg 0 \) (the factor \( e^{i\theta/2} \) is necessary to ensure that the scaling transformation appropriately conserves scalar products [41]) on a region with spatial extension proportional to \( 1/\lambda \). Spurious reflections are introduced when the grid is not fine enough to capture the phase jump of \( e^{i\theta/2} \) in the transition region. This can be identified with the appearance of a bump structure in the reflection curves.

The connections between ABs built so far, Figure 1, allows us to proceed a step further in the chain and recast SES as a MFA through the composition of \( b \) and \( c \) links. The points in Figure 9 represent the reflection error for an MFA associated to \( V_{\text{CAP}}^S(x) \) through (24). The behavior here is slightly different from what we observed in the previous section; for \( E > 0.15 \) eV, \( \epsilon(E, L) \) closely reproduces the results obtained with \( V_{\text{CAP}}^S(x) \), while for lower energies it absorbs better. This indicates a better numerical stability of the mask function formalism compared to the SES CAP one, at least in the present implementation.

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Since the SES transformation (32) extends beyond the boundary region \([0, L]\) into \([-X, 0]\) a direct comparison of the reflection curves in Figure 9 with the ones presented in the previous sections is unfair. In order to enable the
of the absorption properties of any absorber based on absorbing boundaries. In this paper we studied the problem of modeling the dynamics of extended states in finite volumes with the use of absorbing boundaries.

8 Conclusions

In this paper we studied the problem of modeling the dynamics of extended states in finite volumes with the use of absorbing boundaries. We derived a method that permits the evaluation of the absorption properties of any absorber based on the numerical time propagations of localized wavepackets. This method served to systematically illustrate the main features of the three most important boundary families known in the literature: mask function absorbers, complex absorbing potentials, and smooth exterior complex scaling potentials. All these absorbers can be easily implemented in existing codes performing time propagations of the Schrödinger equation with little computational costs and leaving unaffected the overall scaling, for instance, with the system size.

We showed how the calculated numerical reflection properties can be directly employed to assess the quality of the absorption spectrum of the one-dimensional hydrogen atom in the continuum. This indicates the possibility to employ the reflection curves presented in a large number of situations involving real atoms and molecules. Although the presented results have been derived in one dimension the obtained curves could be directly applied in higher dimensional problems provided spherical symmetric boundaries are employed and the impinging wavepackets are perpendicular to the boundary surface. In the most general case, a study of the reflection properties as a function of the angle of incidence is, in principle, needed.

Finally we discussed the intimate connection between the three families of ABs. Although being piecewise discussed in the literature, in this work we illustrated this network of connections in a unified picture. This was done by pairing analytical derivations with numerical examples. Furthermore, we showed how in certain situations, mask function boundaries can provide a stabler formalism for numerical applications.

In the recent years lot of attention as been posed on the description of dynamical processes involving continuum states in different sectors of physics ranging from ultrafast, ultra-intense laser physics to molecular transport. From the theoretical standpoint this poses tremendous challenges that need to be overcome. The consolidation of a common ground for the illustration and classification of ABs presented in this work is an important step towards the development of numerical tools capable to accurately capture these complex phenomena.

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A.U.G. designed the study and wrote the first version of the paper. A.H.L. contributed to the writing and discussion, A.R. supervised and contributed to the discussion.

Appendix: MFA and CAP

The connection between CAPs and mask functions is established using the Baker-Campbell-Hausdorff formula which expresses the product of the exponentials of two operators \( X \) and \( Y \) as a single exponential

\[
e^X e^Y = e^{X+Y}
\]

Localized in the boundary region. The new transformation is thus turning into the complex plane only for \( x > 0 \) reaching the asymptotic condition \( \approx xe^{i\theta} \) at the border of the boundary region \([0,L]\):

\[
F(x) = \begin{cases} 
  x & \text{if } x < 0 \\
  x + (e^{i\theta} - 1) \left( \frac{x}{2} - \frac{L}{2\pi} \sin \left( \frac{\pi x}{L} \right) \right) & \text{if } 0 \leq x \leq L \\
  (x - \frac{L}{2})e^{i\theta} + \frac{L}{2} & \text{if } x > L.
\end{cases}
\]

\[
g(x) = \begin{cases} 
  0 & \text{if } x < 0 \\
  \sin^2 \left( \frac{\pi x}{L} \right) & \text{if } 0 \leq x \leq L, \\
  1 & \text{if } x > L.
\end{cases}
\]

\[
x=0 \quad L
\]

\[
\epsilon(E,L) \quad \text{solid lines} 
\]

\[
\epsilon(x,E) \quad \text{inset}
\]

\[
x \leq \frac{L}{2}
\]

\[
x \geq \frac{L}{2}
\]
with
\[
Z = X + Y + \frac{1}{2}[X,Y] + \frac{1}{12}\{X,[X,Y]\} + \frac{1}{12}[Y,[X,Y]] + \frac{1}{720}\left\{[[H_0,\hat{M}],\hat{M}],\hat{M}\} + \ldots\right\} + O(\Delta t^3)
\] (A.2)

Here \([X,Y] = XY - YX\) is a commutator. In (A.19) the operators are \(X = \hat{M}\) and \(Y = -i\hbar_0\Delta t\). Substituting and ordering by powers of \(\Delta t\),
\[
Z = -i\Delta t \left( \frac{i}{\Delta t} \hat{M} + H_0 \frac{1}{2} \left[ \hat{M}, H_0 \right] + \frac{1}{12} \left[ \hat{M}, \left[ \hat{M}, H_0 \right] \right] + \frac{1}{720} \left[ \left[ \left[ H_0, \hat{M} \right], \hat{M} \right], \hat{M} \right] + \ldots \right)
\]
\[
- \Delta t^2 \left( \frac{1}{12} \left[ H_0, \left[ \hat{M}, H_0 \right] \right] + \ldots \right) + O(\Delta t^3)
\] (A.3)

we obtain an explicit form for the exponent of the finite-difference time propagator. The formula in (21) can be easily obtained by comparison with (19) to first order in \(\Delta t\).

The results can be further simplified by direct evaluation of the commutators. Even in the presence of an external potential in \(H_0\), only the kinetic operator yields a contribution \([-\nabla^2/2, \hat{M}(x)\}] to the commutator. In one dimension, nested commutators containing a single instance of \(H_0\) and more than two \(\hat{M}(x)\) are zero, and (21) can be written in a closed form.

The reciprocal relation connecting CAP to an MFA can be obtained using Zassenhaus’ formula that expresses the decomposition of the exponential of an operator represented as a sum of two operators \(X\) and \(Y\) into the product of exponential operators
\[
e^{X+Y} = e^{X}e^{-\frac{i}{2}Y[X,Y]}e^{\frac{i}{2}[Y,[X,Y]] + [X,[X,Y]]} \times e^{-\Delta t\frac{i}{12}[[Y,[X,Y]],Y] + \frac{1}{12}[[X,Y],Y]Y] + \ldots}
\] (A.4)

This time we have \(X = -i\hbar_0\Delta t\) and \(Y = -i\nu_{\text{CAP}}\Delta t\) and, after substituting and ordering by powers of \(\Delta t\), we obtain
\[
e^{-i\left(H_0 + \nu_{\text{CAP}}\Delta t\right)} = e^{-i\hbar_0\Delta t} \times e^{-i\nu_{\text{CAP}}\Delta t} e^{\frac{i\Delta t^2}{12\hbar_0\nu_{\text{CAP}}}} \times e^{O(\Delta t^3)}
\] (A.5)

Equation (23) is easily obtained comparing the above expression with (19) and evaluating \([H_0, \nu_{\text{CAP}}]\) in one dimension.

References
1. B. Boudaiffa, P. Cloutier, D. Hunting, M.A. Huels, L. Sanche, Science 287, 1658 (2000)
2. Introduction to Molecular Electronics, Lecture Notes in Physics, edited by G. Cuniberti, K. Richter, G. Fagas (Springer, Berlin, Heidelberg, 2006), Vol. 680
3. M. Meckel et al., Science 320, 1478 (2008)
4. Y. Huismans et al., Science 331, 61 (2011)
5. R.M. Lock, S. Ramakrishna, X. Zhou, H.C. Kapteyn, M.M. Murnane, T. Seideman, Phys. Rev. Lett. 108, 133901 (2012)
6. R.R. Lucchese, G. Raseev, V. McCoy, Phys. Rev. A 25, 2572 (1982)
7. I. Sánchez, F. Martín, J. Phys. B 30, 679 (1997)
8. H. Bachau, E. Cormier, P. Decleva, J.E. Hansen, F. Martin, Rep. Prog. Phys. 64, 1815 (2001)
9. A. Dora, J. Tennyson, L. Bryjko, T. van Mourik, J. Chem. Phys. 130, 164307 (2009)
10. P. Descouvemont, D. Baye, Rep. Prog. Phys. 73, 036301 (2010)
11. A. Ermolaev, I. Puzyrin, A. Selin, S. Vinitsky, Phys. Rev. A 60, 4831 (1999)
12. J. Inglesfield, J. Phys.: Condens. Matter 20, 095215 (2008)
13. J.E. Inglesfield, J. Phys.: Condens. Matter 23, 305004 (2011)
14. T. Nakatsukasa, K. Yabana, J. Chem. Phys. 114, 2550 (2001)
15. S. Kurth, G. Stefanucci, C.O. Almblad, A. Rubio, E.K.U. Gross, Phys. Rev. B 72, 035308 (2005)
16. G. Stefanucci, S. Kurth, A. Rubio, E.K.U. Gross, Phys. Rev. B 77, 075339 (2008)
17. J. Krause, K. Schafer, K. Kuklicer, Phys. Rev. A 45, 4998 (1992)
18. K. Kuklicer, F. Mies, K. Schafer, Phys. Rev. A 53, 2562 (1996)
19. M. Lein, J. Marangoz, P. Knight, Phys. Rev. A 66 (2002)
20. S. Chelkowski, C. Foisy, A.D. Bandrauk, Phys. Rev. A 57, 1176 (1998)
21. R. Grobe, S. Haan, J. Eberly, Comput. Phys. Commun. 117, 200 (1999)
22. U. De Giovannini, D. Varsano, M.A.L. Marques, H. Appel, E.K.U. Gross, A. Rubio, Phys. Rev. A 85, 062515 (2012)
23. U. De Giovannini, G. Brunetto, A. Castro, J. Walkenhorst, A. Rubio, ChemPhysChem 14, 1363 (2013)
24. A. Crawford-Uraga, U. De Giovannini, D.J. Mowbray, S. Kurth, A. Rubio, J. Phys. B 47, 124018 (2014)
25. D. Neuhauser, M. Baer, J. Chem. Phys. 91, 4651 (1989)
26. R. Sauntra, L.S. Cederbaum, Chem. Phys. 365, 1 (2002)
27. K. Varga, S. Pantelides, Phys. Rev. Lett. 88, 076804 (2007)
28. B.D. Wibking, K. Varga, Phys. Lett. A 376, 365 (2012)
29. J. Muga, J.P. Palao, B. Navarro, I.L. Egusquiza, Phys. Rep. 395, 357 (2004)
30. J. Aguilar, J. Combes, Commun. Math. Phys. 22, 269 (1971)
31. E. Balslev, J.M. Combes, Commun. Math. Phys. 22, 280 (1971)
32. J.L. Sanz-Vicario, E. Lindroth, N. Brandefelt, Phys. Rev. A 66, 052713 (2002)
33. D.L. Whitenack, A. Wasserman, Phys. Rev. Lett. 107, 163002 (2011)
34. A.H. Larsen, U. De Giovannini, D.L. Whitenack, A. Wasserman, A. Rubio, J. Phys. Chem. Lett. 4, 2734 (2013)
35. B. Simon, Ann. Math. 97, 247 (1973)
36. B. Simon, Phys. Lett. A 71, 211 (1979)
37. C. McCurdy, C. Stroud, M. Wisinski, Phys. Rev. A 43, 5980 (1991)
38. U.V. Riss, H.D. Meyer, J. Phys. B 28, 1475 (1995)
39. N. Moiseyev, J. Phys. B 31, 1431 (1999)
40. O. Shemer, D. Brisker, N. Moiseyev, Phys. Rev. A 71, 032716 (2005)
41. A. Scrinzi, Phys. Rev. A 81, 053845 (2010)
42. D. Macias, S. Brouard, J.G. Muga, Chem. Phys. Lett. 228, 672 (1994)
43. Á. Vibók, G.J. Halász, Phys. Chem. Chem. Phys. 3, 3048 (2001)
44. D.E. Manolopoulos, J. Chem. Phys. 117, 9552 (2002)
45. J.P. Palao, J. Muga, J. Phys. Chem. A 102, 9464 (1998)
46. R. Zavin, I. Vorobeichik, N. Moiseyev, Chem. Phys. Lett. 288, 413 (1998)
47. A. Vibók, G.G. Balint-Kurti, J. Phys. Chem. 96, 8712 (1992)
48. U.V. Riss, H.D. Meyer, J. Chem. Phys. 105, 1409 (1996)
49. M.A.L. Marques, A. Castro, G. Bertsch, A. Rubio, Comput. Phys. Commun. 151, 60 (2003)
50. X. Andrade, A. Castro, H. Appel, M. Oliveira, C.A. Rozzi, F. Lorenzen, M.A.L. Marques, E.K.U. Gross, A. Rubio, Phys. Stat. Sol. B 243, 2465 (2006)
51. X. Andrade et al., J. Phys.: Condens. Matter 24, 233202 (2012)
52. A.H. Larsen, U. De Giovannini, A. Rubio, Density-functional methods for excited states (Springer Berlin, Heidelberg, 2015), Topics in Current Chemistry
53. C. Jhala, M. Lein, I. Dreissigacker, Phys. Rev. A 82, 063415 (2010)
54. M.A.L. Marques, N.T. Maitra, F. Nogueira, E.K.U. Gross, A. Rubio, Fundamentals of Time-Dependent Density Functional Theory (Springer-Verlag, Berlin, 2011)
55. R. Kosloff, D. Kosloff, J. Comput. Phys. 63, 363 (1986)
56. W. Magnus, Commun. Pure Appl. Math. 7, 649 (1954)
57. N. Moiseyev, J.O. Hirschfelder, J. Chem. Phys. 88, 1063 (1988)
58. N. Moiseyev, Non-Hermitian Quantum Mechanics (Cambridge University Press, 2011)
59. U.V. Riss, H. Meyer, J. Phys. B 26, 4503 (1993)
60. R. Santra, Phys. Rev. A 74, 034701 (2006)
61. Y. Sajeev, M. Sindelka, N. Moiseyev, Chem. Phys. 329, 307 (2006)
62. D.J. Kalita, A.K. Gupta, J. Chem. Phys. 134, 094301 (2011)