Resonant enhancements of high-order harmonic generation

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Solving the one-dimensional time-dependent Schrödinger equation for simple model potentials, we investigate resonance-enhanced high-order harmonic generation, with emphasis on the physical mechanism of the enhancement. By truncating a long-range potential, we investigate the significance of the long-range tail, the Rydberg series, and the existence of highly excited states for the enhancements in question. We conclude that the channel closings typical of a short-range or zero-range potential are capable of generating essentially the same effects.

I. INTRODUCTION

A very intriguing feature of above-threshold ionization or high-order harmonic generation concerns the dependence of these phenomena on the intensity of the driving field. The photo-electron or high-order harmonic peaks, as functions of the driving-field intensity, present resonance-like enhancements, such that a variation of a few percent in the external-field strength may drive up the spectral intensity by an order of magnitude. These enhancements have been observed experimentally by several groups for above-threshold ionization (ATI)\textsuperscript{[4,5]}, and, recently, for high-order harmonic generation (HHG)\textsuperscript{[3]}. A concomitant effect in ATI is a variation of the contrast of the spectrum\textsuperscript{[5]}

Early numerical observations that enhancements of ATI go hand in hand with enhancements of HHG were reported in Ref.\textsuperscript{[4]}. The existence of enhancements for both phenomena is not surprising, and is related to their common physical origin. Indeed, HHG and ATI present very similar spectral features, which are explained by similar physical pictures. These features are a wide energy range with approximately equally strong harmonics or photo-electron peaks, known as “the plateau”, followed by a sharp decrease in the harmonic or photo-electron signal, known as “the cutoff”. HHG is described by the so-called “three-step model”, in which an electron is ionized through tunneling or multiphoton ionization, is accelerated by the field and driven back to its parent ion, where it recombines to the ground state, emitting its energy as one harmonic photon\textsuperscript{[6]}. A similar process is responsible for the plateau in above-threshold ionization, with the main difference that, instead of recombining with the parent ion, the electron is elastically rescattered off it\textsuperscript{[5]}. In either case, the precise shape of the atomic potential is not very important, and it can be approximated, for instance, by a zero-range potential. This approximation describes very well the spectral features near the high-energy end of the plateau.

The resonance-like enhancements, however, primarily occur in the first half of the plateau, for which both the external driving field and the atomic potential are expected to influence the harmonic or photo-electron emission\textsuperscript{[7]}. In fact, very different arguments have been put forward to explain these enhancements. Several studies attribute these features to Rydberg states that, for an appropriate ponderomotive upshift, become multiphoton resonant with the ground state. Namely, a free electron in a laser field acquires a field-dependent energy shift by the ponderomotive potential $U_p = e^2 \langle A^2(t) \rangle / 2m$, where $A(t)$ is the vector potential of the laser field and $\langle \ldots \rangle$ denotes the average over a field cycle. Highly excited Rydberg states tend to undergo about the same shift as free electrons\textsuperscript{[8]}. The result of such a multiphoton resonance is either an increase in ionization, or the electronic wave packet is trapped near its parent ion for relatively long times, originating resonance-like structures in the spectra\textsuperscript{[9]}. The mechanism is quite similar to the Freeman resonances\textsuperscript{[10]}, which dominate the low-energy ATI spectrum where rescattering plays no role. According to this physical picture, the presence of high-lying Rydberg states is essential for the existence of the enhancements.

An – at least at first glance – completely different view relates these enhancements to channel closings that, by the same ponderomotive-upshift mechanism, may move into multiphoton resonance with the ground state. If, due to this shift, $N$ photons are no longer sufficient for the electron to reach the continuum, one refers to the $N$-photon channel as having closed. At an intensity corresponding to a channel closing, the electron is released in the continuum with a vanishing drift momentum. In consequence, in the course of its oscillatory motion in the laser field, it will return many times to its parent ion and upon each revisit have the opportunity to rescatter. Quantum mechanically, the corresponding probability amplitudes interfere, and a constructive interference

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manifests itself as an enhancement [3]. Such an effect
does not require the existence of excited states or Ryd-
berg states so that the atom can be modeled by a binding
potential of zero range, which neither supports excited
bound states nor resonances in the continuum [13,14].
The zero-range potential affords the concept of “quan-
tum orbits” which allows for an almost analytical ap-
proach to intense-laser–atom phenomena [13]. However,
one has to keep in mind that real atoms do have long-
range-potential tails, so that channel closings are diffuse
owing to the presence of the Rydberg series. In addition,
they support various bound states, whose influence on
the enhancements is not entirely clear.

In this paper, we perform a systematic study of the
influence of both the laser field and the atomic potential
on these enhancements, for high-order harmonic gener-
ation, by means of simplified atomic models. We address
the question of which one of the existing physical inter-
pretations is ultimately correct or whether both pictures
are complementary aspects of a more complete descrip-
tion. In particular, we investigate the importance of the
highly excited states in the process, and whether or not
they are crucial for the feature in question.

II. TRUNCATED SOFT-CORE POTENTIALS

We compute the harmonic spectra using the numerical
solution of the time-dependent Schrödinger equation
\[
\frac{d}{dt} |\psi(t)\rangle = \left[ \frac{\hbar^2}{2} + V(x) - p \cdot A(t) \right] |\psi(t)\rangle,
\]
for a one-dimensional model atom initially in the
ground state of a binding potential \( V(x) \) and subject to
a laser pulse with the field \( E(t) = -dA(t)/dt \). Atomic
units are used throughout. We consider a monochromatic
laser field
\[
E(t) = E_0 \sin \omega t,
\]
and the harmonic spectra are calculated from the dipole
acceleration \( \ddot{x} = \langle \psi(t) \mid -dV(x)/dx + E(t) \mid \psi(t) \rangle \).
We take the smoothly truncated soft-core potential
\[
V(x) = -\frac{\beta}{\sqrt{(\frac{x}{\sigma})^2 + 1}} f(x),
\]
with
\[
f(x) = \begin{cases} 
1 & (|x| < a_0), \\
\cos^7[\pi (|x|/a_0 - 1/2)] & (a_0 < |x| < L), \\
0 & (|x| > L), 
\end{cases}
\]
so that \( V(x) = 0 \) for \( |x| > L \). We choose \( a_0 \) of the order
of a few atomic units, and \( L \) of the order of the electron
excursion amplitude, so that \( L = r \alpha = r E_0 / \omega^2 \) with
the parameter \( r \) of order unity. Setting \( f(x) = 1 \) in [3] gives
the untruncated soft-core potential. By an appropriate
choice of the parameters \( L \) and \( \alpha_0 \), it is possible to alter
the highly excited bound states leaving the ground state
and the low excited states practically unaffected.

Let us assume that an atom, initially in the ground
state with energy \( \varepsilon_0 \), is ionized by \( N \) photons of frequency
\( \omega \), such that the electron reaches the continuum with the
lowest energy possible, that is, with a drift momentum
(outside the range of the binding potential) of zero. The
energy of the \( N \) photons must account for the binding
energy and the kinetic energy of the oscillatory motion
(the ponderomotive energy \( U_p \)), so that
\[
|\varepsilon_0| + U_p = N \omega.
\]
For intensities slightly larger than specified by the condition [3], at least \( N + 1 \) photons will be necessary for ion-
lization, such that Eq. [3] defines the \( N \)-photon channel-
closing intensity. The intensities that solve the channel-
closing condition [3] form a comb whose teeth as a func-
tion of \( \eta = U_p / \omega \) are separated by unity. If there is an
excited bound state with the (field-free) energy \( \varepsilon_n \) and
if this state undergoes the same ponderomotive upshift
as the continuum [3], then multiphoton resonance with
the ground state occurs for intensities such that Eq. [3]
is satisfied with \( \varepsilon_0 \) replaced by \( \varepsilon_0 - \varepsilon_n \),
\[
|\varepsilon_0 - \varepsilon_n| + U_p = N \omega.
\]
For a long-range potential [such as, in one dimension,
our untruncated potential [3], the true continuum is pre-
ceded by the Rydberg series so that one may question
the significance of the channel-closing condition [3] for
any physical phenomenon. For a finite-range potential
[such as the truncated potential (3)] the Rydberg series
is replaced by a finite sequence of bound states whose
number decreases with decreasing \( L \).

Below, we will seek to answer the following questions:
Is the very existence of an enhancement contingent on
the shape of the binding potential? If an enhancement
exists, does it occur at a channel-closing intensity [3]
for some \( N \), or is related to a multiphoton resonance [3]
with a certain excited state \( |n\rangle \), or is it unrelated to either?
For a truncated potential, do enhancements occur both
at the channel-closing intensities and at intensities where
an excited state becomes multiphoton resonant? To what
extent does the harmonic spectrum depend on the shape
of the potential?

For all examples presented in this paper, we choose
\( \beta = 2.1 \) and \( \sigma = 0.2 \) in the soft-core potential [3].
In this case, the field-free energies of the ground state
and the first four excited states are listed in Table [3] for
the untruncated potential as well as for various truncations.
The Table shows that the ground-state energy is virtu-
ally unaltered by the truncations we consider while the
excited states are more and more affected. For the un-
truncated potential, the first excited states are followed
by the Rydberg series. For all truncations considered,
the states listed are the only ones that survived.
large, for the parameter range used, the truncation eliminates the Rydberg series, changes the energy of the third excited state, and leaves the more deeply bound states unchanged. The shape of the effective potential barrier $V_{\text{eff}} = V(x) - xE(t)$ also remains very similar for all cases.

III. RESULTS

As a first step, we investigate a harmonic spectrum as a function of the laser intensity with regard to the existence of enhancements for particular intensities. The frequency of the laser field is taken as $\omega = 0.07600$ a.u., such that Eq. (5) predicts channel-closing intensities for near-integer values of $\eta = U_p/\omega$. The other intensities that correspond to resonances with excited states are also listed in Table I. For the frequency and the intensity range considered in this paper, the Keldysh parameter $\gamma = \sqrt{|\varepsilon_0|/2U_p}$ lies within the interval $1.66 < \gamma < 0.86$, which is mostly within the multiphoton regime.

Figure I displays two harmonic spectra, computed for the untruncated (a) and a truncated (b) soft-core potential. Both spectra are very similar, with pronounced enhancements near the 13th harmonic. For the untruncated and truncated cases, we observe maximal enhancements at $\eta = 4.3$ and $\eta = 4.1$, respectively. These values do not coincide with the channel-closing intensity $\eta = 4.04$ (modulo any integer) predicted by Eq. (5). This shows a clear influence of the binding potential on the absolute intensity for which these features occur. However, comparing parts (a) and (b) of Fig. I we notice a very remarkable fact: the spectra of the untruncated (a) and the truncated (b) potential are almost identical provided we compare a spectrum at the intensity $\eta$ for the former with a spectrum at the intensity $\eta - 0.2$ for the latter. This holds for all intensities considered. Table I shows that the truncated potential no longer supports the Rydberg series and its excited state $|3\rangle$ is very close to the continuum. We conclude that the Rydberg series has no visible impact on the shape of the harmonic spectrum. Checking again Table I we further conclude that the intensities where the enhancements are maximal are compatible with Eq. (6) with $n = 3$.

Further support for these conclusions comes from Fig. II where the yield of the 13th harmonic is plotted as a function of the scaled intensity $\eta = U_p/\omega$. In this figure, we also investigate the effect of the truncation in more detail, for a wider range of intensities and truncating parameters. The most striking feature is that the main effect of the various truncations is a rigid horizontal shift of the yield-versus-intensity curve. Remarkably, this statement includes the pronounced dips which are due to quantum interference between different quantum orbits [7,15]. While this may be plausible for the large values of the truncation parameter ($L = 2\alpha$) where the truncation mostly affects the potential outside of the classical electronic excursion, it is quite surprising for the small value $L = 0.3\alpha$. A similar shift pattern is observed for all low plateau harmonics (not shown).
FIG. 2. Intensity of the 13th harmonic as function of \( \eta = U_p/\omega \). Part (a): Comparison between the untruncated and various truncated soft-core potentials; the two values of the parameter \( L \), \( L = 31.78 \) a.u. and \( L = 4.77 \) a.u., correspond to \( 2a(\eta = 4.8) \) and \( 0.3a(\eta = 4.8) \), respectively. The binding energies for the various truncated potentials are given in Table I. Part (b): Enlargement of part (a), for \( 3.4 < \eta < 5.4 \). The values of this parameter, for which the enhancements are maximal, are marked by arrows.

Next, we investigate the quantitative amount of the afore-mentioned shift. Comparison with Table I shows that all of the enhancement peaks present in Fig. 2 are compatible with the intensities predicted by Eq. (6) for \( n = 3 \). As long as the state \(|3\rangle\) exists as a bound state, we conclude that the enhancement is due to a multiphoton resonance with this state, upshifted by the whole amount of the ponderomotive energy. However, the enhancement still exists (near integer \( \eta \)), though quantitatively smaller, in the case where the truncation (for \( L = 0.3a \)) has eliminated the third excited state. Here we have encountered a case of a pure channel-closing enhancement which is due to a multiphoton resonance with the ponderomotively upshifted continuum threshold.

Why does a multiphoton resonance with the third excited state (as long as it exists) lead to a pronounced resonance in the spectrum while a multiphoton resonance with other excited states does not? We can answer this question by inspecting the wave functions \(|x|n\rangle\) of the excited states. Please, notice that our calculations are still within the multiphoton regime of ionization. In this case, one expects that multiphoton resonance with an excited bound state is particularly relevant if the wave function of the latter is concentrated near the turning points of the wiggling motion of a classical electron with a drift momentum of zero, i.e., when \(|\langle x|n\rangle|^2\) has its maxima near \(|x| \approx E_0/\omega^2\). Figure 3 shows that indeed this is well satisfied by the state \(|3\rangle\).

FIG. 3. The probability distributions \(|\langle x|n\rangle|^2\) for the first four excited states of the untruncated potential (3). The range of excursion amplitudes \( \alpha = E_0/\omega^2 = 2\sqrt{\eta/\omega} \) that is covered in Fig. 2 (for \( 2.3 < \eta < 6.3 \)) is shaded. Truncation has a minor effect for \( n = 1 \) and 2. For \( n = 3 \) this effect is becoming noticeable, and for \( n = 4 \) it is strong. All higher excited states are eliminated by the truncations considered.

It is not surprising that ionization is affected by the resonances in a similar fashion as HHG. In Fig. 4 we plot the normalization of the time-dependent wave function at the end of the pulse, as a function of \( \eta \) for the truncation parameters used in Fig. 2. This normalization is smaller than unity, since part of the wave function is absorbed by a mask function, used in our computations to eliminate spurious reflection effects. It is a good measure of irreversible ionization, since the mask function is located at about ten times the electron excursion amplitude. In the figure, there are clear dips slightly preceding the intensities, for which the channel closings occur. The particular case, for which \(|3\rangle\) is absent, agrees with the results in [14] for a zero-range potential.
renders provided we employ an effective continuum threshold that resonant intensities are well described by Eq. (5), provided the role of the resonant bound state was taken over by the ponderomotively sufficient truncation of the potential, the effect on the harmonic spectrum is largely a horizontal shift of the yield-versus-intensity curve of the various harmonics.

This picture allows one, for the purposes of HHG, to model the atom by a short-range or zero-range potential using a binding energy that is adapted to the energy difference between the ground state and the relevant excited state of the real atom that is supposed to be modeled. In this case, excited states need not be considered and the enhancement can be attributed to a multiphoton resonance with the continuum threshold and the generation – in the three-step model – of an electron with a vanishing drift momentum. The picture of an effective continuum threshold is also intuitively appealing: in the presence of a strong laser pulse, the highly excited states acquire finite widths owing to ionization and the finite pulse duration, so that the electron moves in a quasi-continuum.

In the case that we investigated (\(a_0 = 3, \ L = 0.3\alpha\)), cf. Fig. 6, the enhancement as well as the general harmonic yield were markedly reduced compared with the untruncated potential. This is not so for a comparison of a realistic single-active-electron binding potential to a (regularized) zero-range potential in three spatial dimensions, neither for HHG nor for ATI [18].

In conclusion, we have confirmed the significance of resonant enhancements for high-order harmonic generation. The resonances occur, when an appropriate highly excited state or, in the absence of such a state, the continuum threshold are ponderomotively upshifted so that they become multiphoton resonant with the ground state. The mechanism is similar to high-order above-threshold ionization. In the latter case, remarkably, the multiphoton resonance continues to be significant while the ionization process is already deeply in the tunneling regime [19]. For high-order harmonic generation, this remains to be investigated.

**IV. DISCUSSION AND CONCLUSIONS**

The picture that emerges from these calculations is this: high-order harmonic generation can be strongly enhanced by a multiphoton resonance with a ponderomotively upshifted excited bound state close to the continuum threshold. In the cases that we investigated the relevant (field-free) bound state \([n]\) was the state where \(x_{n,\text{max}}\) \(\approx \alpha \equiv E_0/\omega^2\) with \(x_{n,\text{max}}\) the value of \(x\) that renders \((x|n)\)’s maximal. Bound states higher than this do not lead to noticeable enhancements, nor do the existence or nonexistence of a Rydberg series have an effect, neither on the enhancements nor on the entire harmonic spectrum. In all cases considered, the relevant bound state was \([n = 3]\). When this state was eliminated by sufficient truncation of the potential, the role of the resonant bound state was taken over by the ponderomotively upshifted continuum threshold. Hence, in all cases, the resonant intensities are well described by Eq. (3), provided we employ an effective continuum threshold that is given by \(\tilde{\varepsilon} \equiv \varepsilon_0 - \varepsilon_n\) where \(\varepsilon_n\) is the energy of the crucial bound state if such a state exists and zero otherwise. Remarkably, if the value of \(\varepsilon_n\) changes owing to a change of the potential, the effect on the harmonic spectrum is largely a horizontal shift of the yield-versus-intensity curve of the various harmonics.

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| n  | untruncated ε<sub>n</sub> | Δη<sub>n</sub> | a<sub>0</sub> = 8 L = 31.78 ε<sub>n</sub> | Δη<sub>n</sub> | a<sub>0</sub> = 3 L = 31.78 ε<sub>n</sub> | Δη<sub>n</sub> | a<sub>0</sub> = 3 L = 4.77 ε<sub>n</sub> | Δη<sub>n</sub> |
|----|----------------|---------|-------------------------------|---------|-------------------------------|---------|-------------------------------|---------|
| 0  | 0.7566         | 0.04    | 0.7566                        | 0.04    | 0.7565                        | 0.05    | 0.7565                        | 0.05    |
| 1  | 0.0846         | 0.16    | 0.0845                        | 0.16    | 0.0825                        | 0.13    | 0.0524                        | 0.74    |
| 2  | 0.0465         | 0.67    | 0.0453                        | 0.64    | 0.0389                        | 0.55    | -                             | -       |
| 3  | 0.0216         | 0.33    | 0.0124                        | 0.21    | 0.0036                        | 0.09    | -                             | -       |
| 4  | 0.0156         | 0.25    | 0.0014                        | 0.06    | -                             | -       | -                             | -       |