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MÜNCHEN

Bachelor's Thesis

# The influence of excited atomic states in multiphoton ionization

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Supervised by:

apl. Prof. Vladislav Yakovlev

Prof. Ulrich Schollwöck

July 16, 2025





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# Der Einfluss angeregter atomarer Zustände auf die Mehrphotonenionisation

Johannes Porsch



FAKULTÄT FÜR PHYSIK

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## **Abstract**

Multiphoton ionization of atoms in strong laser fields is a fundamental process in attosecond physics. In this work, we extend the strong-field approximation (SFA) by incorporating the influence of excited atomic states on ionization rates. Standard SFA formulations neglect these excited states, assuming that the laser field has no effect on the atom before ionization. However, in intense few-cycle laser pulses, the Stark shift and transient population of excited states can significantly modify ionization dynamics. We numerically solve the time-dependent Schrödinger equation (TDSE) using the tRecX code to extract time-dependent probability amplitudes for hydrogen's ground and excited states. These amplitudes are then integrated into the SFA formalism to evaluate their impact on ionization rates.

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# 1. Introduction

Ionization with an intense laser is one of the most fundamental processes in attosecond physics and quantum mechanics in general and has a broad range of applications such as medical physics or . Whether you want your system to ionize or not, it may be important to understand, how ionization works. However since it is a quantum mechanical process, one has to first ask what information are even experimentally accessible. Quantum mechanics at its very fundamental level tells us one can only measure probabilities and how likely the electron get ionized. Unfortunately, in an actual experiment that's the only option you got. Further one likes to know more about the "path" the electron takes after it gets struck by the laser pulse i.e the response of the electron to the laser pulse.

Everytime you hear "path" in quantum mechanics, the first name to think of is Heisenberg. His uncertainty principle sets the boundaries for these type of questions and does not let measure all the information you wish to had. Further to see the electrons path, one has to probe the system just as lightwaves probe the matter around you letting you see. However probing a quantum mechanical system effectively changes the quantum mechanical state of the electron so eventually one has to restrict oneself to something different.

The challenge of a theoretical model is now to predict ionization probabilities and further explain the underlying mechanisms and in the best case allowing some insights what happens between initial state of the atom and measured electron. A clever way to do this is by establishing the concept of an instantaneous ionization rate (IIR) via the property  $P_{\text{ion}} = \int_{\mathbb{R}} \Gamma(t) dt$  effectively telling you when and where an electron reacts to an incoming laser pulse. At first sight, "instantaneous" seems to conflict with the energy time uncertainty relation  $\Delta E \Delta t \geq \hbar/2$ .



## 2. Theory

The structure of this chapter largely follows [10], with several modifications and adaptations to the context of this thesis.

The aim of this chapter is to derive an expression for the instantaneous ionization rate (IIR) by analytically solving the time-dependent Schrödinger equation (TDSE) within the framework of the strong-field approximation (SFA).

### 2.1 Basic Formalism

The goal is to describe the time evolution of a quantum system under an external, time-dependent electromagnetic field in a form suitable for the strong-field approximation. To achieve this, a working formalism must be developed. The starting point is the time-dependent Schrödinger equation for a given Hamiltonian and the Maxwell equations, from which the light-matter interaction can be derived.

#### 2.1.1 Schrödinger Equation

The time evolution of a quantum system  $|\Psi(t)\rangle$  is determined by the time-dependent Schrödinger equation, a partial differential equation that describes the relationship between the change in time of  $|\Psi(t)\rangle$  with a general Hamiltonian  $\mathcal{H}(t)$ :

$$i\hbar \frac{\partial}{\partial t} |\Psi(t)\rangle \stackrel{\text{a.u.}}{=} i \frac{\partial}{\partial t} |\Psi(t)\rangle = \hat{\mathcal{H}}(t) |\Psi(t)\rangle. \quad (2.1)$$

$\hat{\mathcal{H}}$  contains all relevant information for the time evolution, meaning it can be understood as being generated by the Hamiltonian. The formal solution of (2.1) depends on the time dependence of the Hamiltonian and the physical system under consideration. In the general case where the Hamiltonian does not commute with itself at different times, the solution

to (2.1) can be written as a Dyson series:

$$|\Psi(t)\rangle = \hat{\mathcal{U}}(t, t_0) |\Psi(t_0)\rangle = \hat{1} + \sum_{n=1}^{\infty} (-i)^n \int_{t_0}^t dt_1 \int_{t_0}^{t_1} dt_2 \cdots \int_{t_0}^{t_{n-1}} dt_n \hat{\mathcal{H}}(t_n) \hat{\mathcal{H}}(t_{n-1}) \cdots \hat{\mathcal{H}}(t_1) |\Psi(t_0)\rangle. \quad (2.2)$$

**Physical setting** This thesis focuses only on the ionization dynamics of a hydrogen atom exposed to an ultrashort and intense laser pulse. The Coulomb potential in atomic units is given by

$$\hat{\mathcal{H}}_C \stackrel{\text{a.u.}}{=} \frac{1}{|\hat{\mathbf{r}} - \hat{\mathbf{r}}_0|}.$$

Since the TDSE is a *linear* PDE, the Hamiltonian can be partitioned into separate parts. This partitioning can be performed in various ways, all of which are mathematically equivalent. For instance, a division into the unperturbed part and the interaction part yields:

$$\hat{\mathcal{H}}(t) \stackrel{\text{a.u.}}{=} \underbrace{\frac{\hat{\mathbf{P}}^2}{2} + \frac{1}{|\hat{\mathbf{r}} - \hat{\mathbf{r}}_0|}}_{\hat{\mathcal{H}}_0} + \hat{\mathcal{H}}_I(t).$$

Solving the TDSE with  $\hat{\mathcal{H}}_0$  provides the bound states of the hydrogen atom. The calculation can be found, for instance, in [16] and will not be discussed here.

### 2.1.2 Light-Matter Interaction

A light wave consists of an electric field  $\mathbf{E}(\underline{x}, t)$  and a magnetic field  $\mathbf{B}(\underline{x}, t)$ , which propagate through vacuum at the speed of light. These fields are defined by Maxwell's equations in empty space:

$$\begin{aligned} \nabla \cdot \mathbf{E} &= 0 & \nabla \times \mathbf{E} &= -\frac{\partial \mathbf{B}}{\partial t} \\ \nabla \cdot \mathbf{B} &= 0 & \nabla \times \mathbf{B} &= \mu_0 \epsilon_0 \frac{\partial \mathbf{E}}{\partial t} \end{aligned}$$

These equations can be solved using the potentials

$$\begin{aligned} \mathbf{E} &= -\nabla \varphi - \frac{\partial \mathbf{A}}{\partial t}, \\ \mathbf{B} &= \nabla \times \mathbf{A}, \end{aligned} \quad (2.3)$$

where  $\mathbf{A}(\underline{x}, t)$  is the vector potential and  $\varphi(\underline{x}, t)$  is the scalar potential. These potentials are not uniquely determined, as different choices can lead to the same physical fields. Specifically, the transformations

$$\mathbf{A} \rightarrow \mathbf{A} + \nabla \chi, \quad \varphi \rightarrow \varphi - \frac{\partial \chi}{\partial t}, \quad (2.4)$$

with  $\chi(t)$  being an arbitrary smooth scalar function, also satisfy Maxwell's equations. This non-uniqueness is referred to as gauge freedom and arises directly from the structure of the equations. Selecting a particular gauge (i.e., fixing a certain  $\chi$ ) can be just a matter of convenience and simplify calculations, as demonstrated below.

### 2.1.3 Dipole Approximation

The dipole approximation is a widely used simplification in light-matter interaction. It applies when the wavelength of the optical field is significantly larger than both the size of the relevant bound electron states and the maximum displacement of a free electron. Additionally, it assumes that the magnetic field of the light has a negligible effect on the electron dynamics, requiring nonrelativistic particle velocities.

To derive this approximation, we first express Maxwell's equations in terms of the vector and scalar potentials, as defined in (2.3). This results in two coupled differential equations, which complicates the analysis. However, a simpler expression for the vector potential  $\mathbf{A}$  can be obtained by selecting a specific gauge—the Lorentz gauge:

$$\partial_\mu \mathbf{A}^\mu = 0 \quad \text{or} \quad \nabla \cdot \mathbf{A} + \frac{\partial \varphi}{\partial t} = 0.$$

This gauge choice is achieved by solving the inhomogeneous wave equation for  $\chi$ , which arises in the explicit calculation and is feasible when  $\mathbf{A}$  and  $\varphi$  are known. With this gauge, Maxwell's equations decouple into:

$$\begin{aligned} \nabla^2 \varphi - \frac{\partial^2 \varphi}{\partial t^2} &= 0, \\ \nabla^2 \mathbf{A} - \frac{\partial^2 \mathbf{A}}{\partial t^2} &= 0. \end{aligned}$$

The second equation is of particular interest, as it is a wave equation describing plane-wave solutions for  $\mathbf{A}$ :

$$\mathbf{A}(\underline{x}, t) = \mathbf{A}_0 e^{\pm i(\mathbf{k} \cdot \mathbf{r} - \omega t)}.$$

Mathematically, the dipole approximation corresponds to the leading-order term in the Taylor expansion of  $e^{i\mathbf{k} \cdot \mathbf{r}}$ . Consequently, the vector potential becomes spatially independent and can be approximated as [5]:

$$\mathbf{A}(\underline{x}, t) = \mathbf{A}_0 e^{\mp i\omega t} \exp \left\{ \pm 2\pi i \frac{|\mathbf{r}|}{\lambda} \mathbf{e}_{\mathbf{k}} \cdot \mathbf{e}_{\mathbf{r}} \right\} \approx \mathbf{A}_0 e^{\mp i\omega t} \left( 1 + \mathcal{O} \left( \frac{|\mathbf{r}|}{\lambda} \right) \right) = \mathbf{A}(t)$$

This approximation holds as long as the wavelength is sufficiently large. Under these

conditions, the magnetic field simplifies to:

$$\mathbf{B} = \nabla \times \mathbf{A} \approx 0.$$

Although a different gauge may be chosen later, the physical description of the system remains unchanged. The dipole approximation itself is gauge-independent, ensuring that  $\mathbf{B}$  remains approximately zero in any gauge. The Lorentz gauge is used here for convenience, as it provides an intuitive framework for the vector potential expansion.

The dipole approximation is valid for most experimental configurations, including the calculations in this work. However, it has limitations—for example, it breaks down in cases where strong-field ionization generates very fast electrons ([5], [14]).

This was the essence of the dipole approximation. To further simplify the description of the laser field in the Hamiltonian, the choice of gauge must be considered more carefully.

#### 2.1.4 Length and velocity gauge

Calculating ionization rates in strong-field physics presents challenges due to the choice of gauge. While gauge transformations do not alter physical reality in classical electrodynamics, they can affect the practical computation of formulas derived in this chapter, requiring careful consideration.

First, we derive two basic expressions for the Hamiltonian in the velocity gauge and length gauge using the dipole approximation. The semiclassical Hamilton function for a free electron in an electric field<sup>1</sup> is:

$$\hat{\mathcal{H}}(\underline{x}, t) = \frac{1}{2m}(\hat{\mathbf{P}} - e\mathbf{A}(\underline{x}, t))^2 - e\varphi(\underline{x}, t) \quad (2.5)$$

Under the dipole approximation, this simplifies to:

$$\hat{\mathcal{H}}(\underline{x}, t) = \frac{\hat{\mathbf{P}}^2}{2m} - \frac{e}{m}\hat{\mathbf{P}} \cdot \mathbf{A}(t) + \frac{e^2}{2m}\mathbf{A}^2(t) - e\varphi(\underline{x}, t)$$

In most literature,  $\varphi$  is set to zero at this stage of the derivation because the sources of the electromagnetic wave lie outside the region of interest. While the dipole approximation does not strictly necessitate this assumption,  $\varphi$  will be left as is.

A common assumption in semiclassical Hamiltonians is that the vector potential alone drives electron state transitions, with no back-reaction from the electron [5]. This approximation holds for sufficiently high laser intensities, though its validity may depend on specific conditions.

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<sup>1</sup>Derivation can be found in [13]



Next, we perform a gauge transformation called length gauge via:

$$\chi = -\mathbf{A}(t) \cdot \underline{x} \quad (2.6)$$

This transformation sets  $\mathbf{A}$  to zero, modifying  $\varphi$  as follows:

$$\nabla\varphi \rightarrow \nabla \cdot (\varphi + \mathbf{x} \cdot \frac{\partial \mathbf{A}}{\partial t}) = \nabla\varphi + \frac{\partial \mathbf{A}}{\partial t} = -\mathbf{E} \quad (2.7)$$

Integrating this equation from the origin to  $\mathbf{x}$  yields the electric potential in the length gauge. With  $\mathbf{r}$  now quantized, the Hamiltonian becomes:

$$\hat{\mathcal{H}}(\underline{x}, t) = \frac{\hat{\mathbf{P}}^2}{2m} - e\hat{\mathbf{x}} \cdot \mathbf{E}$$

The time-dependent part  $\hat{\mathcal{H}}(t)$  of the Hamiltonian can be expressed as:

$$\hat{\mathcal{H}}_L(t) = -\hat{\mathbf{d}} \cdot \mathbf{E}(t) \quad (2.8)$$

where  $\hat{\mathbf{d}} = e\hat{\mathbf{x}}$  is the dipole operator and  $\mathbf{E}(t)$  is the electric field.

An alternative gauge transformation, the velocity gauge, is also relevant. Using:

$$\chi = -\frac{e^2}{2m} \int_{-\infty}^t \mathbf{A}^2(t') dt'$$

the dipole approximation leaves  $\mathbf{A}$  unchanged but modifies  $\varphi$  as:

$$\varphi \rightarrow \varphi - \frac{e^2}{2m} \mathbf{A}^2(t)$$

resulting in the Hamiltonian:

$$\hat{\mathcal{H}}(\underline{x}, t) = \frac{\hat{\mathbf{P}}^2}{2m} - \frac{e}{m} \hat{\mathbf{P}} \cdot \mathbf{A}(t)$$

and

$$\hat{\mathcal{H}}_V(t) = -\frac{e}{m} \hat{\mathbf{P}} \cdot \mathbf{A}(t) \quad (2.9)$$

where  $\varphi$  is set to zero.

These are standard formulations of the interaction Hamiltonian for light-matter interactions. As with any gauge transformation, certain scenarios favor one gauge over another (see Chapter 4).

## 2.2 Strong Field Formalism

The difficulty with ionization arises from the existence of in some sense two Hilbert spaces: one for the bound states in the Hydrogen atom that accounts for distortion of the wavefunction due to the laser field, and another for the continuum states that are primarily affected by the laser field but also influenced by the binding potential.

It will be shown that SFA provides a treatment for the second Hilbert space, simplifying its description. However, previous SFA formulations oversimplify the first Hilbert subspace to such an extent that there remains one Hilbert space for eigenstates unaffected by the laser field and another for continuum states unaffected by the binding potential.

SFA, in principle, concerns only the second part; neglecting the dynamics before ionization is not truly part of it. This section derives an expression that avoids excessive simplification of the first Hilbert space. Additionally, it will be demonstrated that there are two equivalent ways of solving the TDSE, which later diverge when approximations are introduced. Nevertheless, it becomes evident that these approaches nevertheless yield the same analytical result.

### 2.2.1 Projection into Subspaces

The full time-dependent Hamiltonian  $\hat{\mathcal{H}}(t)$  is projected onto subspaces using projection operators defined by:

$$\hat{X} = \sum_n |\Psi_n\rangle \langle \Psi_n| \quad \text{and} \quad \hat{Y} = \hat{1} - \hat{X}$$

The use of projection operators for this type of problem is inspired by [17]. Projection operators are idempotent, i.e.  $\hat{X}^2 = \hat{X}$ . This property can be considered as the defining characteristic of projection operators, so applying them twice produces nothing new.

Here,  $|\Psi_n\rangle$  represents the bound states of the atom. Using  $\hat{X} + \hat{Y} = \hat{1}$ , any partition of the Hamiltonian can be chosen. The most natural partitioning is first written as:

$$\hat{\mathcal{H}}(t) = \underbrace{\hat{X}\hat{\mathcal{H}}(t)\hat{X}}_{\hat{\mathcal{H}}_{\text{Bound}}(t)=\hat{\mathcal{H}}_0^{\text{Sub}}(t)} + \underbrace{\hat{Y}\hat{\mathcal{H}}(t)\hat{Y} + \hat{X}\hat{\mathcal{H}}(t)\hat{Y} + \hat{Y}\hat{\mathcal{H}}(t)\hat{X}}_{\hat{\mathcal{H}}_{\text{Cont}}(t)+\hat{\mathcal{H}}_{\text{Rec}}(t)+\hat{\mathcal{H}}_{\text{Ion}}(t)=\hat{\mathcal{H}}_1^{\text{Sub}}(t)} = \hat{\mathcal{H}}_0^{\text{Sub}}(t) + \hat{\mathcal{H}}_1^{\text{Sub}}(t)$$

Similar to  $\hat{\mathcal{H}}(t) = \hat{\mathcal{H}}_0(t) + \hat{\mathcal{H}}_1(t)$ , where  $\hat{\mathcal{H}}_1(t)$  represents the interacting perturbation (for instance, in the length gauge (2.8)), but now applied to the full Hamiltonian projected onto subspaces. For reasons that will be explained later, this is referred to as the Subspace partition. While the notation may appear arbitrary now, its significance will become clear later.

The Hamiltonian can alternatively be partitioned into a different form, the full partition:

$$\hat{\mathcal{H}}(t) = \underbrace{\hat{X}\hat{\mathcal{H}}(t)\hat{X} + \hat{Y}\hat{\mathcal{H}}(t)\hat{Y} + \hat{X}\hat{\mathcal{H}}(t)\hat{Y}}_{\hat{\mathcal{H}}_{\text{Bound}}(t) + \hat{\mathcal{H}}_{\text{Cont}}(t) + \hat{\mathcal{H}}_{\text{Rec}}(t) = \hat{\mathcal{H}}_0^{\text{Full}}(t)} + \underbrace{\hat{Y}\hat{\mathcal{H}}(t)\hat{X}}_{\hat{\mathcal{H}}_{\text{Ion}}(t) = \hat{\mathcal{H}}_1^{\text{Full}}(t)} = \hat{\mathcal{H}}_0^{\text{Full}}(t) + \hat{\mathcal{H}}_1^{\text{Full}}(t)$$

The physical meaning of these terms can be interpreted as follows: In  $\hat{\mathcal{H}}_{\text{Bound}}(t)$ , the electron remains within the subspace and can always be described by a superposition of bound states. Without ionization, this would be the only term governing the time evolution. In previous work, this part of the Hamiltonian was often simplified to the extent that effects like Stark shift or coupling to excited states were neglected.

$\hat{\mathcal{H}}_{\text{Cont}}(t)$  describes similar dynamics, but in the subspace of continuum states rather than bound states. While in reality these two spaces cannot be completely separated, this distinction provides conceptual clarity. This is the part where the SFA will later be applied.

The remaining terms describe different physical processes. Two processes can occur: ionization of the electron or recombination with the atom. These processes are governed by the terms  $\hat{\mathcal{H}}_{\text{Rec}}(t)$  and  $\hat{\mathcal{H}}_{\text{Ion}}(t)$ . In the present case, only one of these terms is relevant, and it will be shown later that  $\hat{\mathcal{H}}_{\text{Ion}}(t)$  is responsible for ionization, effectively "kicking" the electron out of the atom [10].

While this approach may initially appear unnecessary, it establishes boundaries for the electron's behavior and provides useful intuition.

### 2.2.2 Dyson equation

For determining the time evolution, care must be taken since the full Hamiltonian does not commute with itself at different times, requiring an exact treatment. As the full Dyson series (2.2) can be cumbersome to deal with, an alternative approach is used. This approach is enabled by the fact that the Hamiltonian can be split into two parts by projecting it into subspaces. The ansatz chosen here is with the Dyson equation. It relies on the time evolution operator and is completely equivalent to the TDSE and therefore to the Dyson series (2.2). It can be written in two forms that are equivalent in their raw formulation but may yield different results when approximations such as SFA are introduced.

The more intuitive form of the Dyson equation for this physical setting is given by:

$$\hat{\mathcal{U}}(t, t_0) = \hat{\mathcal{U}}_0^{\text{Sub}}(t, t_0) - i \int_{t_0}^t \hat{\mathcal{U}}(t, t') \hat{\mathcal{H}}_1^{\text{Sub}}(t') \hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) dt' \quad (2.10)$$

Note that the "Sub" propagators are not identical to the "Full" propagators (2.11), but within the Dyson ansatz, the full propagator  $\hat{\mathcal{U}}$  remains unchanged. This will change later when SFA is introduced. Most formulas in this thesis can be best read from right to

left, such as this one. The time evolution starts with the Hamiltonian projected onto the bound states. The interaction Hamiltonian then effectively ionizes the electron, inducing a transition to a virtual state, followed by propagation with the full time evolution. For more insights, refer to [10].

The Dyson equation can also be expressed in a slightly different form:

$$\hat{\mathcal{U}}(t, t_0) = \hat{\mathcal{U}}_0^{\text{Full}}(t, t_0) - i \int_{t_0}^t \hat{\mathcal{U}}_0^{\text{Full}}(t, t') \hat{\mathcal{H}}_I^{\text{Full}}(t') \hat{\mathcal{U}}(t', t_0) dt' \quad (2.11)$$

It can be verified that both equations satisfy the TDSE by direct substitution of  $\hat{\mathcal{U}}(t, t_0)$ . The main difference between the “Full” and “Sub” formulations is that the “initial” propagator on the right side is now the full time propagator instead of a reduced one. That motivates the acronym for both variants.

To simplify the analysis, the physical setting is further specified. Starting from the ground state of the atom, the time evolution is given by  $\hat{\mathcal{U}}(t, t_0) |\Psi_0(t_0)\rangle$ . For additional simplification, a projection onto a continuum state  $|\Pi(t_c)\rangle$  at time  $t_c$  is considered.  $|\Pi(t_c)\rangle$  does not represent a plane wave but rather an abstract state characterized by its momentum. Naturally, there is no overlap between  $|\Pi(t_c)\rangle$  and  $\hat{\mathcal{U}}_0^{\text{Sub}}(t, t_0) |\Psi_0(t_0)\rangle$  or with  $\hat{\mathcal{U}}_0^{\text{Full}}(t, t_0) |\Psi_0(t_0)\rangle$ , as the electron has not yet been ionized and no initial population in the continuum is assumed [10]. Furthermore, when  $\hat{\mathcal{H}}_I^{\text{Sub}}(t')$  and  $\hat{\mathcal{H}}_I^{\text{Full}}(t')$  are expanded, and the orthogonality of bound states and continuum states is taken into account, most terms of  $\hat{\mathcal{H}}_I(t')$  vanish. The remaining expressions are:

$$\langle \Pi(t_c) | \hat{\mathcal{U}}(t, t_0) | \Psi_0(t_0) \rangle = -i \int_{t_0}^t \langle \Pi(t_c) | \hat{\mathcal{U}}(t, t') \hat{Y} \hat{\mathcal{H}}(t') \hat{X} \hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) | \Psi_0(t_0) \rangle dt' \quad (2.12)$$

and

$$\langle \Pi(t_c) | \hat{\mathcal{U}}(t, t_0) | \Psi_0(t_0) \rangle = -i \int_{t_0}^t \langle \Pi(t_c) | \hat{\mathcal{U}}_0^{\text{Full}}(t, t') \hat{Y} \hat{\mathcal{H}}(t') \hat{X} \hat{\mathcal{U}}(t', t_0) | \Psi_0(t_0) \rangle dt' \quad (2.13)$$

Note: No approximations have been made; (2.12) and (2.13) are exact.

### 2.2.3 Strong Field Approximation

In principle, the strong field approximation (SFA) involves neglecting the binding potential after ionization. This approximation is intuitively reasonable because the laser field becomes the dominant force acting on the electron, exerting significantly greater influence on the ionization dynamics than the Coulomb potential. It should be noted that SFA is not restricted to high-intensity laser pulses; even for arbitrary small ionization probabilities, SFA implies that once ionization occurs, the laser pulse dominates the subsequent dynamics.

First, the “Sub” partitioning must be addressed. The time evolution operator after ionization can be expressed as:

$$\hat{\mathcal{U}}(t, t') \approx \hat{\mathcal{U}}_{\text{SFA}}(t, t') = e^{-i \int_{t'}^t \hat{\mathcal{H}}_{\text{SFA}}(t'') dt''} \quad \text{and} \quad \mathcal{H}_{\text{SFA}}(t') = \hat{\mathcal{H}}(t) - \hat{\mathcal{H}}_{\text{C}}$$

Similarly, for the “Full” partitioning:

$$\hat{\mathcal{U}}_0^{\text{Full}}(t, t') \approx \hat{\mathcal{U}}_{\text{SFA}}(t, t') = e^{-i \int_{t'}^t \hat{\mathcal{H}}_{\text{SFA}}(t'') dt''} \quad \text{and} \quad \mathcal{H}_{\text{SFA}}(t') = \hat{\mathcal{H}}(t) - \hat{\mathcal{H}}_{\text{C}}$$

An exact expression for the SFA time evolution operator can be written, as it now commutes with itself at different times.

The SFA is particularly useful because the eigenstates of  $\hat{H}_{\text{SFA}}$  have an exact analytical solution: the Volkov states. To see this, consider the semiclassical Hamiltonian in (2.5).

Classically, the physics driven by the momentum operator  $\hat{\mathbf{P}}$  is known as the canonical momentum, given by:

$$\frac{\partial \mathcal{L}}{\partial \dot{\mathbf{x}}} = \mathbf{P} = m\dot{\mathbf{x}} + \frac{e}{c} \mathbf{A} \stackrel{\text{a.u.}}{=} \mathbf{p} + \mathbf{A} \quad (2.14)$$

where  $\mathcal{L}$  is the Lagrangian of the system.

In this case, the canonical momentum is conserved. To verify this, setting  $\varphi = 0$  (justified earlier) gives  $\mathbf{E} = -\frac{\partial \mathbf{A}}{\partial t}$ . The equation of motion for a charged particle in an electromagnetic field [13] is:

$$\frac{d\mathbf{p}}{dt} = \mathbf{E} + (\dot{\mathbf{x}} \times \mathbf{B}) \approx -\frac{\partial \mathbf{A}}{\partial t} = -\frac{d\mathbf{A}}{dt}$$

Thus,  $\frac{d}{dt} \mathbf{P} = 0$ .

And also the energy of the system is clear:

$$E(t) = \dot{\mathbf{x}} \frac{\partial \mathcal{L}}{\partial \dot{\mathbf{x}}} - \mathcal{L} = \frac{\mathbf{p}^2}{2} \quad (2.15)$$

Note that the energy is not conserved, as the previous argument holds only for the canonical momentum, not the kinetic momentum.

The key observation is that  $\langle \Pi(t_c) |$  (not  $\langle \mathbf{p}(t_c) |$ ) is an eigenstate of  $e^{-i \int_{t'}^t \hat{H}_{\text{SFA}}(t'') dt''}$ . Combining (2.14) and (2.15) yields:

$$\langle \Pi(t_c) | e^{-i \int_{t'}^t \hat{H}_{\text{SFA}}(t'') dt''} = \langle \Pi(t_c) | e^{-i \int_{t'}^t (\mathbf{p} + \mathbf{A}(t''))^2 dt''}$$

Here,  $\mathbf{P}$  is time-independent, while  $\mathbf{A}$  is not. Using the conservation of canonical momentum, the momentum at other times can be calculated. Of particular interest is the

asymptotic limit when the laser field vanishes:

$$\mathbf{P} = \mathbf{p}(t'') + \mathbf{A}(t'') = \mathbf{p}(t \rightarrow \infty) + \mathbf{A}(t \rightarrow \infty) = \mathbf{p}$$

Furthermore,  $|\Pi\rangle$  represents now a plane wave solution(how??),

$$|\Pi\rangle = |\mathbf{p} + \mathbf{A}\rangle$$

Combining these results leads to the following expressions for both “Full” and “Sub”:

$$\langle \Pi(t_c) | \hat{\mathcal{U}}(t, t_0) | \Psi_0(t_0) \rangle = -i \int_{t_0}^t e^{-i \int_{t'}^t (\mathbf{p} + \mathbf{A}(t''))^2 dt''} \langle \mathbf{p} + \mathbf{A} | \hat{\mathcal{H}}(t') \hat{X} \hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) | \Psi_0(t_0) \rangle dt' \quad (2.16)$$

and

$$\langle \Pi(t_c) | \hat{\mathcal{U}}(t, t_0) | \Psi_0(t_0) \rangle = -i \int_{t_0}^t e^{-i \int_{t'}^t (\mathbf{p} + \mathbf{A}(t''))^2 dt''} \langle \mathbf{p} + \mathbf{A} | \hat{\mathcal{H}}(t') \hat{X} \hat{\mathcal{U}}(t', t_0) | \Psi_0(t_0) \rangle dt' \quad (2.17)$$

Note that the SFA eliminates the difference in the left part of the integrand between “Sub” and “Full”, making the two equations distinct. The final step introduces a new approximation for  $\hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) | \Psi_0(t_0) \rangle$  and  $\hat{\mathcal{U}}(t', t_0) | \Psi_0(t_0) \rangle$ .

**Sub-Hilbertspace ansatz** Regardless of the final form, the expression can be expanded in the bound states  $|\Psi_n\rangle$ . This is clear by considering the defining equation of the time evolution operator and observing that any overlap of  $\hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) | \Psi_0(t_0) \rangle$  with a state outside the bound-state representation would cancel out due to  $\hat{X}$ . With this consideration, an ansatz using the interaction picture can be formulated as:

$$\hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) | \Psi_0(t_0) \rangle = \sum_n c_n(t') e^{-i E_n t'} |\Psi_n\rangle^2$$

The factors  $e^{-i E_n t'}$  are factored out to avoid potential numerical issues arising from their rapid oscillations.

In theory, this ansatz is exact but relies on the implicit assumption that the Hilbert spaces can be easily separated. Additionally, it assumes that the bound states remain unaffected by the laser field, which is not strictly true in reality. However, this approximation can be justified by the low laser intensity and the resulting ionization probability not exceeding 0.01.

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<sup>2</sup>Here,  $E_n$  are the eigenenergies and not the electric field. If it would be the electric field it would be denoted as a vector or the component of the vector.

**Full Hilbertspace ansatz** This case involves a more severe approximation. The primary advantage is that only the time evolution up to the ionization moment  $t'$  is required. The same assumptions regarding laser intensity and ionization probability apply here, justifying the approximation of the time evolution as a superposition of bound states:

$$\hat{\mathcal{U}}(t', t_0) |\Psi_0(t_0)\rangle = \sum_n c_n(t') e^{-iE_n t'} |\Psi_n\rangle$$

The key difference between the two ansatzes lies in the coefficients  $c_n(t')$ . In the subspace approach, the coefficients are obtained by solving the TDSE within the subspace spanned by the bound states. This approach effectively cuts off ionization, forcing the electron to the subspace regardless of laser intensity. Implementing this leads to a system of ordinary differential equations for the coefficients, hence the name. In contrast, the other approach is fundamentally different. Here, the TDSE must be solved in the full Hilbert space, with the coefficients extracted at times  $t'$ . In the atto/femtosecond regime, this demands substantial computational resources, as the electron exits the sub-Hilbert space during the process. For these numerical simulations, a solver called tRecX (discussed in Chapter 3) was employed.

Unlike other literature ([2], [10]), this ansatz does not neglect transitions between bound states prior to ionization. For example, the laser pulse may excite the electron without immediately ionizing it. The coefficients are expected to capture all relevant dynamics within the hydrogen atom before ionization, including effects such as AC Stark shifts and ground-state distortion.

The final expression can now be presented. By simplifying the dipole matrix element  $\mathbf{d}_n(\mathbf{p} + \mathbf{A}) = \langle \mathbf{p} + \mathbf{A} | \hat{\mathcal{H}}(t') | \Psi_n \rangle$  between the bound state and the continuum state, the dipole approximation (2.8) can be applied, yielding:

$$\langle \Pi(t_c) | \Psi_n(t) \rangle = -\frac{i}{2} \int_{t_0}^t e^{-i \int_{t'}^t (\mathbf{p} + A_z(t''))^2 dt''} E_z(t') \sum_n c_n(t) e^{-iE_n t'} \langle \mathbf{p} + A_z | \hat{d}_z | \Psi_n \rangle dt'$$

Here, the electric field is polarized along the z-axis. This equation corresponds to the strong-field S-matrix in the length gauge, as found in many papers [2], but with the distinction that excited states are not neglected.

## 2.3 Ionization rates

The concept of an instantaneous ionization rate (IIR) is a powerful and intuitive tool for understanding the physics of strong-field ionization. It is frequently used to provide a time-resolved picture of the ionization process, for instance, by assigning a probability for an electron to be released at a specific moment within the laser pulse [9]. This notion is

particularly useful in semiclassical models where the IIR helps to define the initial conditions for classical electron trajectories.

However, the concept of an ionization ‘event’ occurring at a precise instant in time is difficult to reconcile with conventional quantum mechanics. Strictly speaking, the only well-defined and directly measurable observable is the *total* ionization probability, which is the probability of finding the electron in a continuum state after the laser pulse has ended [9]. The IIR itself is not a direct quantum mechanical observable, and its definition is consequently not unique.

This ambiguity leads to some theoretical challenges, such as defining a working formalism for the tunneling delay time. Furthermore, an IIR defined by projecting the time-evolving wavefunction onto the subspace of bound states during the laser pulse interaction generally depends on whether the length gauge or velocity gauge is used. Since physical observables must be gauge-invariant, this poses a conceptual problem.

To address these challenges, alternative, unambiguous definitions have been proposed. One such approach defines the IIR as a functional derivative of the total ionization probability with respect to the waveform of the ionizing laser pulse [9]. This method is inherently gauge-invariant because it is based on physically measurable quantities. While this thesis will not follow this particular formalism, it is important to acknowledge the subtleties involved in defining an ionization rate. The subsequent derivation will proceed within the SFA framework, which provides a tractable, but gauge-dependent expression for the IIR.

### 2.3.1 Derivation of extended SFA Rate

The derivation mainly follows [2] with some modifications. First, the ionization probability is expressed as the amplitude of projection of the wavefunction onto the continuum subspace. The total ionization probability is obtained by integrating the ionization rate over the time domain. Therefore, the SFA rate (2.2.3) can be written as:

$$\begin{aligned}
 \lim_{t \rightarrow \infty} \langle \Psi(t) | \int d^3p |p\rangle \langle p| \Psi(t) \rangle &= \int_{-\infty}^{\infty} \Gamma_{\text{SFA}}(t) dt \\
 &= \int d^3p \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} dt_1 dt_2 e^{\frac{i}{2} \int_{t_1}^{t_2} (\mathbf{p} + A_z(t''))^2 dt''} E_z(t_1) E_z(t_2) \\
 &\quad \times \left( \sum_n e^{iE_n t_1} c_n^*(t_1) d_{z,n}^*(\mathbf{p} + A_z(t_1)) \right) \left( \sum_n e^{-iE_n t_2} c_n(t_2) d_{z,n}(\mathbf{p} + A_z(t_2)) \right)
 \end{aligned}$$



By changing variables to  $t = \frac{t_2+t_1}{2}$  and  $T = \frac{t_2-t_1}{2}$  and using the fact that the laser pulse is polarized along the z-axis, the expression becomes:

$$\begin{aligned}\Gamma_{\text{SFA}}(t) &= \int d^3p \int_{-\infty}^{\infty} dT e^{\frac{i}{2} \int_{t-T}^{t+T} (\mathbf{p} + A_z(t''))^2 dt''} E_z(t-T) E_z(t+T) \\ &\quad \times \left( \sum_n e^{i(t-T)E_n} c_n^*(t-T) d_{z,n}^*(\mathbf{p} + A_z(t-T)) \right) \left( \sum_n e^{-i(t+T)E_n} c_n(t+T) d_{z,n}(\mathbf{p} + A_z(t+T)) \right) \\ &= \sum_{n_1} \sum_{n_2} \int_0^{\infty} dp p^2 \int_0^{\pi} d\theta_p \sin \theta_p \int_0^{2\pi} d\phi_p \int_{-\infty}^{\infty} dT e^{\frac{i}{2} \int_{t-T}^{t+T} (\mathbf{p} + A_z(t''))^2 dt''} e^{i(t-T)E_{n_1} - i(t+T)E_{n_2}} \\ &\quad \times E_z(t-T) E_z(t+T) c_{n_1}^*(t-T) c_{n_2}(t+T) d_{z,n_1}^*(\mathbf{p} + A_z(t-T)) d_{z,n_2}(\mathbf{p} + A_z(t+T))\end{aligned}$$

The integration variable is changed from  $\mathbf{p}$  to  $\mathbf{p} + \mathbf{A}$ , and with  $\Delta(t') = \mathbf{A}(t+t') - \mathbf{A}(t)$ , the expression becomes:

$$\begin{aligned}\sum_{n_1} \sum_{n_2} \int_0^{\infty} dp p^2 \int_0^{\pi} d\theta_p \sin \theta_p \int_0^{2\pi} d\phi_p \int_{-\infty}^{\infty} dT e^{\frac{i}{2} \int_{-T}^T (\mathbf{p} + \Delta_z(t''))^2 dt''} e^{i(t-T)E_{n_1} - i(t+T)E_{n_2}} \\ \times E_z(t-T) E_z(t+T) c_{n_1}^*(t-T) c_{n_2}(t+T) d_{z,n_1}^*(\mathbf{p} + \Delta_z(-T)) d_{z,n_2}(\mathbf{p} + \Delta_z(T))\end{aligned}$$

Also the integrand in the exponential function can be further simplified:

$$\begin{aligned}\frac{i}{2} \int_{-T}^T (\mathbf{p} + \Delta_z(t''))^2 dt'' &= i\mathbf{p}^2 T + \frac{i}{2} \int_{-T}^T \Delta_z(t'')^2 + 2\mathbf{p} \cdot \Delta_z(t'') dt'' \\ &= i\mathbf{p}^2 T + ip \cos \theta_p \int_{-T}^T \Delta_z(t'') dt'' + \frac{i}{2} \int_{-T}^T \Delta_z(t'')^2 dt''\end{aligned}$$

Following the notation used in the code as well as in [2], where

$$\bar{\Delta}^k = \int_{-T}^T \Delta^k(t') dt' = \int_{-T}^T (\mathbf{A}(t+t') - \mathbf{A}(t))^k dt'$$

the final SFA rate to be implemented is given by:

$$\Gamma_{\text{SFA}}(t) = \sum_{n_1} \sum_{n_2} \int_0^{\infty} dp p^2 \int_0^{\pi} d\theta_p \sin \theta_p \int_0^{2\pi} d\phi_p \int_{-\infty}^{\infty} dT \quad (2.18)$$

$$\times \exp \left( i\mathbf{p}^2 T + ip \cos \theta_p \bar{\Delta}_z^k + \frac{i}{2} \bar{\Delta}_z^2 + i(t-T)E_{n_1} - i(t+T)E_{n_2} \right) \quad (2.19)$$

$$\times E_z(t-T) E_z(t+T) c_{n_1}^*(t-T) c_{n_2}(t+T) d_{z,n_1}^*(\mathbf{p} + \Delta_z(-T)) d_{z,n_2}(\mathbf{p} + \Delta_z(T)) \quad (2.20)$$

For numerical calculations, it can be convenient to split a complex integrand into its phase and absolute value. Although this is typically done with the given formula, the coefficients remain complex and may exhibit rapidly oscillating phase behavior.

## 2.4 Strong Field Ionization

This section mostly follows [1]. While not directly necessary for this thesis, these terms are frequently encountered in the literature and are briefly mentioned here for completeness. Ionization can occur in different ways, depending on the laser parameters and the atom or material used. To distinguish these processes, several parameters must be defined. According to [12], strong field ionization primarily depends on three parameters: the photon energy, the binding potential, and the ponderomotive energy.

**Ponderomotive Energy** The ponderomotive energy is defined as the cycle-averaged quiver energy of a free electron in an electromagnetic field [cite wikipedia]. Since harmonic motion is experienced by the particle, the time-averaged kinetic energy is given by

$$\frac{1}{2}m\omega^2 \langle x^2 \rangle = \frac{e^2 E^2}{4m_e \omega^2} = \frac{E^2}{4\omega^2}$$

**Keldysh Parameter** The Keldysh parameter  $\gamma$  compares the ponderomotive energy with the ionization potential. It is defined as

$$\gamma = \sqrt{\frac{I_p}{2U_p}}$$

The ionization potential in the hydrogen atom is  $I_p = 13.6\text{eV} = 0.5\text{au}$ . Different regimes of ionization, each with distinct physical interpretations, can be defined based on the value of the Keldysh parameter.

### 2.4.1 Tunneling Ionization

This regime is typically characterized by  $\gamma \ll 1$ . Here, the laser pulse is sufficiently strong to distort the potential barrier. Alternatively, the tunneling regime can be defined as the regime where the quasi-static approximation holds, meaning the instantaneous ionization rate is entirely determined by the absolute value of the electric field at that time [1]. This allows the ionization rate for DC and AC fields in this regime to be calculated analytically [4].

$$\Gamma_{\text{SFA}}(t) \propto \exp\left(-\frac{2}{3} \frac{(2I_p)^{3/2}}{F}\right)$$

Here,  $F$  represents the peak of the electric field for an AC field.

### 2.4.2 Multiphoton Ionization

The regime where  $\gamma \gg 1$  is commonly referred to as multiphoton ionization. This occurs when the field strength is significantly weaker than in the tunneling regime. In other words, the multiphoton regime can be defined as the regime where

$$\Gamma_{\text{SFA}}(t) \propto \mathbf{E}^{2N} \quad \text{with} \quad N = \frac{I_p}{E_e}$$

holds [10]. This process can be visualized as the electron gaining energy while remaining under the barrier, effectively being heated up. Unlike the tunneling regime, this process is non-adiabatic. For further details, refer to [10] and [12].

### 2.4.3 Intermediate Regime

This regime is particularly relevant for this thesis. The previous regimes describe limiting cases where the complementary process is negligible. However, in reality, a coexistence of both processes often occurs. In the intermediate regime, the tunneling process is no longer adiabatic. The potential barrier undergoes significant oscillations during the electron's tunneling time, allowing energy to be absorbed from the laser field while tunneling takes place.

## 2.5 Stark shift

This section mainly follows [7] and [8].

**Phenomenology** The Stark shift describes the shifting and splitting of energy levels due to the presence of an external electric field. This phenomenon arises from the interaction between the electric field and the atom's electric dipole moment. Two types of Stark shifts exist: the DC Stark shift and the AC Stark shift. DC stands for direct current, meaning the electric field is constant in time, while AC stands for alternating current, corresponding to a (rapidly) oscillating electric field. Under certain conditions (e.g., a laser pulse far off resonance), the AC Stark shift can be approximated as the quadratic DC Stark shift [8].

When the laser pulse is not far off resonance, significant changes to the energy levels of the bound states and to the electron dynamics can occur.

**AC Stark Shift** The electron wave function can be expressed as a superposition of bound states with coefficients  $c_n(t)$ :

$$|\Psi(t)\rangle = \sum_n c_n(t) e^{-iE_n t} |\Psi_n\rangle$$

where  $E_n$  are the eigenenergies and  $c_n(t)$  are the complex coefficients. When considering electron dynamics before ionization, the coefficients can be interpreted as carrying two types of information. First, the amplitude of the coefficients represents the probability of finding the electron in a given state  $n$ . Second, the phase of the coefficients reflects changes in the energy level of state  $n$ . The second part can be interpreted as the subcycle AC Stark shift [7]. This can be expressed mathematically as:

$$|\Psi(t)\rangle = \sum_n |c_n(t)| e^{-i \int_{-\infty}^t E_n + \delta E_n(t') dt'} |\Psi_n\rangle \quad (2.21)$$

The dynamic stark energy shifts  $\delta E_n(t')$  are of primary interest.

In the context of this thesis, the influence of the AC Stark shift on the ionization process is of particular interest. In previous derivations of the SFA rate, both the transition to excited states and the effect of the laser field on the ground state were neglected. However, with the extended version of the SFA rate (2.20), it becomes possible to account not only for the influence of the laser field on excited states but also to distinguish between the effects of the Stark shift and the depletion of the ground state (i.e., changes in the probability amplitude). While including many physical effects is theoretically appealing, it is not always practical unless the impact of each approximation on the results is understood. The findings related to this question are presented in Chapter 4.

## 3. Methods

This chapter introduces the methods used throughout the thesis including numerical methods for solving or implementing certain types of equations, as well as physical methods used for result comparison or data interpretation.

To compare and verify the ionization rate from Chapter 2, the TDSE must be solved numerically without approximations. A reliable approach for comparison involves the use of the sampling method TIPTOE [15], where the ionization yield from both are methods being compared. When expanding an existing model, it is particularly important to perform such comparisons to assess whether modifications align with expected outcomes.

### 3.1 Numerical Methods

To implement formula 2.2.3, the TDSE must be solved numerically in two different ways to obtain the coefficients  $c_n(t)$ . Two methods were used (numerical solver and ODE), each with its own advantages and disadvantages.

For numerically solving the TDSE without approximations, a solver called tRecX was used.

#### 3.1.1 tRecX

tRecX is a C++ code designed for solving generalized inhomogeneous time-dependent Schrödinger-type equations in arbitrary dimensions and in a variety of coordinate systems [19]. Ionization is typically challenging for numerical solvers because the electron leaves the subspace of bound states, making calculations computationally expensive. Additionally, the Coulomb potential decays only as  $1/r$ , raising the question of how to discretize the space and set boundary conditions.

tRecX employs various techniques that make it particularly suitable for light-atom interactions. For example, the method called “infinite-range exterior complex scaling” applies a unitary transformation to complex-scale space beyond a certain distance from the nucleus. This can be likened to the analytic continuation of a function into the complex plane. Fur-

thermore, the wavefunction is ‘damped’ by the complex-scaled region, preventing reflections at the boundary that could lead to numerical issues [18].

Moreover, tRecX allows the gauge ((2.8), (2.9)) in which the TDSE is solved to be specified, increasing its flexibility. This feature will be important later.

Throughout this thesis, tRecX was employed for two purposes.

First, it was used to solve the TDSE in the entire Hilbert space to obtain the coefficients  $c_n(t)$ , which are required for the improved SFA rate. For this purpose, the source code was modified to calculate the coefficients and store them in an appropriate format. Further details regarding the implementation can be found in Chapter 4.

Second, tRecX served as a reference within the TIPTOE experiment to evaluate the performance of the SFA rate with excited states (2.20).

### 3.1.2 System of ODEs

The coefficients of the wavefunction within the bound-state subspace were obtained using the interaction picture, and the resulting coupled system of differential equations was solved numerically. The calculation was implemented in Python, utilizing the ODE solver `solve_ivp` from SciPy’s `integrate` module. The corresponding code is provided in the electronic appendix within the `HydrogenSolver` class. Further details on the implementation can be found in Chapter 4.

## 3.2 TIPTOE

This section mostly follows [15] and [1].

TIPTOE (Tunneling Ionization with a Perturbation for the Time-domain Observation of an Electric field) is a method for direct sampling of an electric pulse in the femtosecond to attosecond regime using quasistatic subcycle tunneling ionization in a gaseous medium or air.

A typical TIPTOE simulation consists of two linearly polarized laser pulses: a “fundamental” and a “signal” pulse, similar to common pump-probe experiments. The drive pulse is the pulse to be sampled, with the ionization yield of a certain medium providing the measurement. In first order, the ionization rate can be approximated as

$$\Gamma(E_F + E_S) \approx \Gamma(E_F) + E_S \left. \frac{d\Gamma(E_S)}{dE} \right|_{E=E_F} \quad (3.1)$$

In this approximation, depletion of the ground state is neglected. The total ionization yield

$N_{\text{total}}$  obtained by the two pulses is given by

$$N_{\text{total}} = N_{\text{F}} + \delta N = \int dt \Gamma(E_{\text{F}}(t)) + \int dt E_{\text{S}}(t) \left. \frac{d\Gamma(E_{\text{S}}(t))}{dE} \right|_{E=E_{\text{F}}(t)}$$

By varying the delay  $\tau$  between the two pulses, the ionization yield takes on different values. From this, it follows that:

$$\delta N(\tau) \propto E_{\text{S}}(\tau) \quad (3.2)$$

Thus, the field amplitude of the signal pulse can be sampled by measuring the ionization yield for different delays. The TIPTOE method can be applied across a broad spectral range of the signal pulse, as long as the fundamental pulse is shorter than 1.5 optical cycles.

This method provides a way to compare ionization dynamics predicted by a numerical solver and by the SFA rate (2.20). TIPTOE is particularly useful because numerical simulations can provide good predictions about ionization probabilities, while analytical models describe ionization rates. A TIPTOE simulation can help reconstruct the ionization dynamics, which is especially relevant in the context of this thesis. Later, the ionization rate  $\Gamma(E_{\text{F}} + E_{\text{S}})$  will be integrated over the full time domain, and the ionization yield for different delays will be compared with the results from the numerical solution of the TDSE. The results are shown in chapter 5.

For better visualization of the underlying physics in TIPTOE, the background ionization  $N_{\text{F}}$  from  $E_{\text{F}}(t)$  is subtracted, and the ionization yield is normalized, so the formula in plot 5.1 reads

$$\frac{N_{\text{total}} - N_{\text{F}}}{N_{\text{max}}} = \frac{\delta N(\tau)}{N_{\text{max}}} \quad (3.3)$$

However, interesting physics can also be observed by comparing only the total ionization yield, as discussed later in Chapter 5.

Typically, TIPTOE is not used for this kind of analysis but rather for its sampling capabilities. Instantaneous ionization rates are highly useful because TIPTOE enables sampling of the electric field of a laser pulse in the femtosecond to attosecond regime, which has broad applications in fields such as laser spectroscopy and medical physics.

### 3.3 GASFIR

GASFIR stands for General Approximator for Strong Field Ionization Rates. It is an analytical retrieval tool designed to reconstruct data obtained from numerical solutions of the TDSE. The model was validated for hydrogen and shows good agreement with existing theories in the quasi-static limit of tunneling ionization, not only for hydrogen but also for helium and neon [3].

The working principle of GASFIR is based on the use of ionization probabilities to retrieve ionization rates. The approach employs the idea from SFA that the rates can be expressed as  $\int dT K(t, T)$ , where  $K(t, T)$  is a kernel function. Later in the code, the kernel function is also visible, with the modifications applied there.

GASFIR is not part of this thesis; however, this thesis was partially motivated by the idea that an improvement of the SFA formalism could benefit the GASFIR approach.



## 4. Implementation

This chapter describes the implementation of the formula (2.20) and the associated challenges. The majority of the code was originally developed by the authors of [2]. Modifications made to both the tRecX source code and the SFA rate are documented in the electronic appendix [github zenodo]. The two main new components in this implementation are the coefficients  $c_n(t)$  and the dipole matrix elements  $d_{z,n}(\mathbf{p})$ , while the remaining parts were straightforward.

### 4.1 Coefficients

As mentioned in chapter 2, the Dyson equation can be written in two ways, resulting in two different expressions for the S-matrix. The difference between the two approaches lies only in the coefficients.

#### 4.1.1 Subspace Ansatz

To understand how the coefficients are determined, the defining equation can be examined:

$$\hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) |\Psi_0(t_0)\rangle = \sum_n c_n(t') e^{-iE_n t'} |\Psi_n\rangle$$

with

$$i\partial_{t'} \hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) = \hat{X} \hat{\mathcal{H}}(t') \hat{X} \hat{\mathcal{U}}_0^{\text{Sub}}(t', t_0) \quad (4.1)$$

This can be interpreted as the wavefunction being restricted to the subspace. The derivation for the coefficients is outlined below.

First,  $\hat{\mathcal{H}}(t)$  is split into two parts:  $\hat{\mathcal{H}}_0$  and  $\hat{\mathcal{H}}_I(t)$ , where the eigenstates of  $\hat{\mathcal{H}}_0$  are  $|\Psi_n\rangle$  and the eigenenergies are  $E_n$ . By substituting the ansatz above into (4.1) and multiplying with  $\langle\Psi_m|$ , the coefficients  $c_n(t)$  are obtained:

$$i\dot{c}_m(t) = \sum_n c_n(t) e^{-i\omega_{nm}t} \langle\Psi_m|\hat{\mathcal{H}}_I(t)|\Psi_n\rangle$$

with  $\omega_{nm} = E_n - E_m$ .

Several challenges arise when implementing this approach. First, while  $\sum_n$  theoretically extends from 0 to  $\infty$ , numerical simulations require a finite number of states. The justification for the chosen number of states must be considered, along with ensuring that truncating the sum does not introduce numerical instabilities.

Additionally, increasing the number of coefficients affects previously calculated values. For example, solving only for the ground state  $c_0(t)$  results in  $|c_0(t)|^2 = 1$ . However, the ground-state occupation decreases as more states are included in the sum, illustrating the *coupled* nature of the system of ODEs.

As the number of states increases, the results (e.g., the rates) should converge. However, since the electron is constrained to the subspace, including highly excited states may introduce numerical instabilities. For intensities around  $10^{14} \frac{\text{W}}{\text{cm}^2}$  and longer wavelengths (800nm – 1200nm), allowing occupation of higher states led to unphysical oscillations in the rates. Under typical conditions, the electron would ionize, but since ionization is suppressed in this approach, excitation to high-energy states can cause the model to fail. Thus, a balance must be struck between accuracy and numerical stability. In the simulations presented later, a maximum of  $n = 3$  was used for the bound-state calculations. This choice is justified by analyzing the population dynamics of the hydrogen atom over time, where most pre-ionization dynamics are governed by the first few bound states—particularly  $1s$ ,  $2s$ ,  $2p$ , and  $3p$ —meaning that including additional states has minimal impact on the lower-state coefficients.

Note: This discussion pertains not to the number of bound states included in the final simulations, but rather to the number considered when solving the ODE. While only the ground state could have been used in the SFA rate calculations, the influence of higher bound states on the ground state must still be assessed. This justification is provided here.

Further, equation (4.1.1) is gauge dependent because of  $\langle m | \hat{\mathcal{H}}_I(t) | n \rangle$ . Either the length gauge (2.8) or the velocity gauge (2.9) can be chosen. Which gauge should be used to obtain the most meaningful results?

To address this question, the scenario can be reconsidered. An electron initially resides in the ground state, and before ionization, its behavior is primarily governed by the Coulomb potential. After ionization, according to the SFA, it is described as a plane wave oscillating in the laser field.

From this perspective, both gauges can be useful. During the first part of the process—when the electron remains bound to the hydrogen atom—the length gauge is more appropriate, as the electron’s behavior is better described in terms of its position rather than its momentum. This is why, in most elementary introductions to light-matter interaction (such as discussions of Rabi oscillations), the length gauge is preferred for systems

where ionization does not occur. Conversely, after ionization, the velocity gauge becomes more suitable, as the electron is no longer influenced by the potential (due to the SFA) and is fully characterized by its momentum.

Thus, the length gauge is a fitting choice for the coefficients used in the ODE ansatz.

### 4.1.2 Full Hilbertspace

The coefficients from the subspace are not the only ones used. A numerical solver (in this case tRecX) was employed to solve the entire TDSE, and the coefficients were extracted from the wavefunction.

As mentioned earlier, there are two ways to consider  $\hat{U}(t', t_0) |\Psi\rangle$ . First, the TDSE can be solved in the subspace of the bound states (as done with the ODE), or the TDSE can be solved in the full Hilbert space and then projected onto bound states. tRecX implements the second approach, which is significantly more complex than the ODE method. Since far more physical effects influence the time evolution of the coefficients—and consequently the ionization rate—it becomes more challenging to isolate individual effects and their consequences. Nevertheless, having two independent methods that, in some sense, compute the same quantity is useful, allowing for interpretation based on their respective advantages and disadvantages.

For extracting the coefficients from tRecX—i.e., solving the full TDSE and projecting onto bound states—the source code was modified as follows: The occupation probability of specified bound states was already implemented, meaning the code outputs  $\langle \Psi(t) | \hat{P}_{\text{Occ}\{H0:n\}} | \Psi(t) \rangle = |\langle \Psi(t) | n \rangle|^2$ . The main modification involved implementing a new function that replaces the left bra  $\langle \Psi(t) |$  with the eigenstate used in  $\hat{P}_{\text{Occ}\{H0:n\}}$ , while ensuring the eigenstates were normalized. This results in:

$$\langle \Psi(t) | \hat{P}_{\text{Occ}\{H0:n\}} | \Psi(t) \rangle \rightarrow \langle n | \hat{P}_{\text{Occ}\{H0:n\}} | \Psi(t) \rangle = \langle n | n \rangle \times \langle n | \Psi(t) \rangle = c_n(t)$$

To achieve this, the eigenvalue problem had to be solved again, and the eigenstates were passed to the function calculating the expectation value. A more elegant and efficient implementation could likely be devised, especially since the eigenstates are already computed elsewhere, but the current approach suffices for now. A detailed description of the changes made to tRecX can be found in the appendix [GitHub/Zenodo].

In principle, this is all that is required. However, simply using the coefficients without further scrutiny would be insufficient. Several issues can arise, particularly when extracting values from code not originally designed for this purpose. This is especially relevant when computing quantities that are not experimentally accessible or are gauge-dependent, as such results would typically be of limited utility. Thus, caution is necessary.

**Gauge dependence** The coefficients are gauge-dependent, similar to the subspace approach using the ODE. However, in tRecX, a "hybrid gauge" is employed. This means that within a certain radius  $R_g$  from the nucleus, length gauge is used, while outside this radius, velocity gauge is applied. Length gauge is better suited for pre-ionization dynamics, as the electron's behavior is more accurately described by its position rather than its momentum. The gauge radius must be chosen carefully to ensure that pre-ionization dynamics are not computed in velocity gauge. To verify the gauge consistency of the coefficients, they were compared with the ODE approach, which operates strictly in length gauge.

**States used in the SFA rate** A similar issue arises with the tRecX coefficients as with the ODE coefficients: determining how many should be included in the calculation. Unlike the subspace approach, ionization is permitted here, so including higher bound states is not expected to cause the same numerical difficulties as before. However, the gauge radius should not be exceeded, as beyond this point, the coefficients correspond to the velocity gauge, which may produce incorrect or unphysical results. To minimize numerical issues, only the first few bound states are typically used for the SFA rate. This choice can be justified by examining the populations of the hydrogen atom, where most dynamics occur within the  $1s$ ,  $2s$ ,  $2p$ , and  $3p$  states.

## 4.2 Dipole matrix Elements

The dipole transition matrix elements  $\mathbf{d}_{nlm}(\mathbf{p})$  from a certain bound state to the Volkov states are somewhat less ambiguous than the coefficients. Calculating them in general is difficult, but in the case of hydrogen, an analytical solution is possible. A detailed derivation can be found in Appendix A.

As mentioned in the theory section, this simplicity in calculation is only possible due to the strong approximation made with the SFA regarding the dipole matrix elements. The final state after ionization is not truly a plane wave but is approximated as such under the SFA.

As discussed in the coefficients section, not all states contribute equally to the ionization dynamics. Besides the ground state  $1s$ , this thesis restricts calculations to the states  $2p$  and  $3p$ . Most of the pre-ionization dynamics is determined by these states; including additional states would primarily refine the ionization rate rather than introduce significant new physics.

Simplifying the matrix elements is also crucial to avoid numerical instabilities. Particularly when integrating over the azimuthal angle  $\phi$  in the final rate, where exact cancellations are expected, limited numerical precision can lead to incorrect results.

The dipole matrix elements can be simplified by noting that the ground state is  $1s$  and the light wave is linearly polarized, ensuring that all other states also have  $m = 0$ . Since the derivation is lengthy, it is provided in Appendix A as well.

### 4.3 TIPTOE Simulations

As discussed in Chapter 3, TIPTOE is used in this thesis to compare the ionization yield obtained from the SFA rate and the numerical solution of the TDSE with tRecX.

In implementation, both approaches can be treated as a type of "machine," where an arbitrary laser field is input and an ionization probability is output. The laser field in this case consists of two pulses in a `cos8` envelope: a strong "fundamental" pulse and a weaker "signal" pulse shifted in time by  $\tau$ . In tRecX, two laser pulses can be implemented directly. In the SFA rate implemented in Python, the laser field is constructed as an object of the class `LaserField` from the file `field_functions.py`.

Particularly in the SFA rate, numerical challenges arise because the signal pulse acts only as a small perturbation to the fundamental pulse. Care must be taken to ensure that numerical methods accurately capture this perturbation; otherwise, TIPTOE will fail. This challenge is addressed in Chapter 5.



## 5. Results and Discussion

Before discussing of the improvement of the SFA rate worked out, first we have to formulate what we want to learn from this kind of generalisation. In principle we want to investigate how the excited states influence the ionisation process or in general.

We want to investigate what influences the ionization dynamics more, the stark shift or the distortion of the ground state. This question is being answered here.

However, the much bigger question is that there is quite a big difference between the overall ionization yield from previous simulations within SFA compared with the numerical solution of the TDSE. An extended version of the SFA model could tell, if the discrepancy is due to the simplification of the dynamics before or after ionization. Unfortunately, this thesis cannot answer this question to a satisfactory level, but it provides the theoretical framework and formulas needed to achieve this goal in future.

### 5.1 Comparison with TDSE using TIPTOE

First compare the ionization dynamics predicted from the standart SFA approach [2] with the results from tRecX (solving TDSE numerically). As can be seen in 5.1(a), the standart SFA does a good job reconstructing the ionization dynamics, but some parts it does not capture at all<sup>1</sup>. Especially in offcycle ionization dyanmics seems to be some kind of phase shift that does not match at all.

This is the starting point of this thesis since that is what was previously known. Now the question is how to improve this SFA model regarding the ionization dynamics.

What I implemented is the extended SFA model described in chapter 2 and chapter 4. As mentioned previously, there are two ways of determining the coefficients, either by solving the TDSE in the subspace of the hilbertspace using the system of ODE's, or numerically in the full hilbertspace using a numerical solver (here tRecX). In the plot 5.1(b) both the TIPTOE results from the Subspace as well as the Full hilbertspace approach are shown. Very important is that here, only the first coefficient and first dipole matrix element is used. In other words, in formula (2.20) only the first term is used, i.e.  $n_1 = n_2 = 1$ . In

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<sup>1</sup>Note that (3.2) is fulfilled and  $\delta N(\tau)$  is actually proportional to the signal pulse.

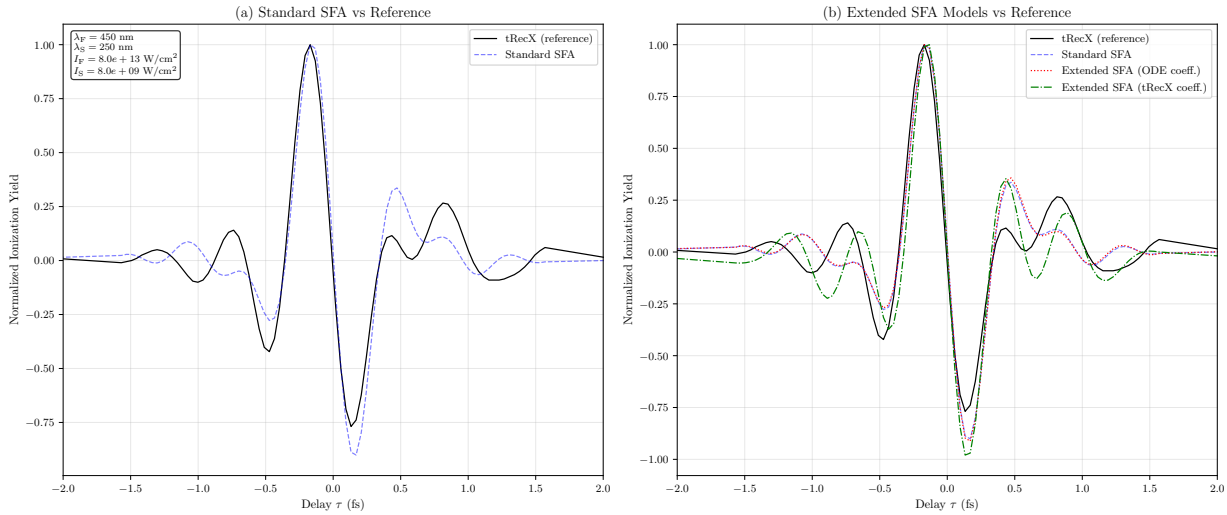


Figure 5.1: Comparison of ionization yields from TIPTOE simulations between different SFA models and reference data from tRecX. (a) Standart SFA overall does a good job reconstructing the ionization dynamics, but some parts it does not capture at all (b) Extended SFA to excited states indeed shows some improvements in the reconstructing

principle, almost the same formula was already implemented for simulating the standart SFA model, since there is also just the ground state involved. The only difference is that in the extended SFA model for one state, the coefficient  $c_0(t)$  is jused so all the difference between standart SFA and extended SFA in 5.1(b) is coming from the coefficients  $c_0(t)$ . That makes it possible to split .....

**Difference in ionization yield between SFA and TDSE** As mentioned in chapter 3, the Ionization yield in 5.1 is normalized, to see the actual difference better. Further, because we are mostly interested in the dynamics and not the plain ionization yield, its normalized according to (3.3). However, in the not normalized plot, one could see that the numerical solution of the TDSE is predicting a ionization probability orders of magnitude higher than the SFA model. In addition to that, small changes in the in the delay of the signal pulse leads to bigger changes in the TDSE results than the SFA results, so the ‘reference’ TIPTOE result is much more sensitive to changes in the laser field. First observation can be interpreted by the fact that in SFA we are assuming, the electron sits in a volkov state after ionization so the elecotrnr does not ‘feel’ the coulomb potential anymore. This is of course a very strong assumption, any much ‘harder’ to achcieve for the electron. Imagine the laser has to really catapult the electron out of the atom to make this assumption approximately valid. Therefore the overall ionization yield from the SFA model is much lower than the one from the TDSE. Thats is also one essential reason for restrictign the laser to small ionization probabilities. Second, the numerical solution of the



TDSE captures much more dynamics and physical effects of the ionization process. Many different processes can happen during that time, and that could justify why the ionization yield of the TDSE is much more sensitive to changes in the electric field.

Both these observations are interesting, but can be expected from how the SFA model is constructed. To make sure that the differences really do come from the ‘fundamentals’ of the SFA model (i.e. the assumption that the final state is a Volkov state), one has to make sure that is the only possible option for the differences. This is of course very difficult since there are many different processes that can have an influence on the ionization dynamics that were not considered in the SFA model. However by making



## 6. Conclusion and Outlook

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# A. Dipole transition matrix elements

Here, the general transition dipole matrix elements into the continuum for a hydrogen atom are derived. The general matrix element in this case is given by:

$$\underline{d}_{nlm}(\underline{p}) = \langle \Pi | \hat{\mathbf{d}} | \Psi_{nlm} \rangle \stackrel{\text{a.u.}}{=} \langle p | \hat{\mathbf{r}} | \Psi_{nlm} \rangle$$

where  $|p\rangle$  is a plane wave. The wave function for the hydrogen atom is well known:

$$\Psi_{nlm}(\underline{x}) = \langle \underline{x} | \Psi_{nlm} \rangle = R_{nl}(r) Y_{lm}(\theta, \phi) \quad (\text{A.1})$$

with  $R_{nl}(r)$  being the radial part of the wavefunction and  $Y_{lm}(\theta, \phi)$  being the spherical harmonics.

By partitioning the  $\hat{\mathbf{1}}$  operator and using the fact that  $\hat{\mathbf{r}} \rightarrow i\nabla_{\underline{p}}$  in momentum representation, a general formula for the transition is obtained:

$$\underline{d}_{nlm}(\underline{p}) = i\nabla_{\underline{p}} \int d^3\underline{x} \psi_{nlm}(\underline{x}) e^{-i\underline{p}\cdot\underline{x}} = i\nabla_{\underline{p}} \phi_{nlm}(\underline{p})$$

In principle, this integral (i.e the Fourier transform) is all that is required. Due to the structure of  $\psi_{nlm}$ , a result similar to (A.1) can be expected. A posteriori, it can be shown that:

$$\mathcal{F}\{\psi_{nlm}(\underline{x})\} = \phi_{nlm}(\underline{p}) = F_{nl}(p) Y_{lm}(\theta_p, \phi_p)$$

where  $F_{nl}(p)$  is the Fourier transform of the radial part of the wavefunction and  $Y_{lm}(\theta_p, \phi_p)$  are the spherical harmonics in momentum space, analogous to the hydrogen atom in position space.

## Momentum space

The so-called plane wave expansion [11] of the exponential part of the integral is given by:

$$e^{i\underline{p}\cdot\underline{x}} = \sum_{l'=0}^{\infty} (2l' + 1) i^{l'} j_{l'}(pr) P_{l'}(\underline{p} \cdot \underline{x}) = 4\pi \sum_{l'=0}^{\infty} \sum_{m'=-l'}^{l'} i^{l'} j_{l'}(pr) Y_{l'm'}(\theta_p, \phi_p) Y_{l'm'}^*(\theta_x, \phi_x)$$

Here,  $j_l(pr)$  represents the spherical Bessel functions, and the integration is performed over spherical coordinates. While the expression initially appears complex, the orthogonality of the spherical harmonics can be used to simplify the integral to:

$$\phi_{nlm}(\underline{p}) = 4\pi \sum_{m=-l}^l Y_{lm}(\theta_p, \phi_p) i^l \underbrace{\int_0^\infty dr r^2 j_l(pr) R_{nl}(r)}_{\tilde{R}_{nl}(p)}$$

This structure is the desired form. The focus now shifts to the radial part  $\tilde{R}_{nl}(p)$  of the integral. The term  $R_{nl}(r)$  corresponds to the radial function of the hydrogen atom in position space and is independent of the magnetic quantum number  $m$ . It consists of an exponential term dependent on  $r$ , a polynomial term dependent on  $r$ , the generalized Laguerre polynomials, and the normalization constant. A closed expression for the generalized Laguerre polynomials would be convenient. They are represented as:

$$L_n^l(r) = \sum_{\iota=0}^n \frac{(-1)^\iota}{\iota!} \binom{n+l}{n-\iota} r^\iota$$

Thus, the Laguerre polynomials depend only on an exponential term and finitely many polynomial terms.  $\tilde{R}_{nl}(p)$  can be expressed (without prefactors and summation over  $\iota$ ) as:

$$\int_0^\infty dr r^{2+l+\iota} e^{-\frac{Zr}{n}} j_l(pr)$$

Before solving the integral using computational methods, the spherical Bessel function must be transformed into ordinary Bessel functions:

$$j_l(pr) = \sqrt{\frac{\pi}{2pr}} J_{l+\frac{1}{2}}(pr)$$

At this stage, it is useful to combine all prefactors and summations into a single expression and examine the integral as a whole:

$$\begin{aligned} \phi_{nlm}(\underline{p}) &= \frac{\pi^{3/2}}{\sqrt{2p}} \sqrt{\left(\frac{2}{n}\right)^3 \frac{(n-l-1)!}{n(n+1)!}} \\ &\times \sum_{m=-l}^l \sum_{\iota=0}^{n-l-1} i^l \frac{(-1)^\iota}{\iota!} \left(\frac{2}{n}\right)^{l+\iota} \binom{n+l}{n-l-1} \underbrace{\int_0^\infty dr r^{l+\iota+\frac{3}{2}} e^{-\frac{Zr}{n}} J_{l+\frac{1}{2}}(pr) Y_{lm}(\theta_p, \phi_p)}_{(*)} \end{aligned} \quad (\text{A.2})$$

The remaining integral was computed using Mathematica, so a detailed derivation is not provided. Interestingly, an analytical solution exists. The result for  $(*)$  is:

$$(*) = {}_2\tilde{F}_1\left(2 + l + \frac{\iota}{2}, \frac{1}{2}(5 + 2l + \iota); \frac{3}{2} + l; -\frac{n^2 p^2}{Z^2}\right)$$

Here,  ${}_2\tilde{F}_1$  denotes the regularized hypergeometric function, defined as:

$${}_2\tilde{F}_1(a, b; c; z) = \frac{{}_2F_1(a, b; c; z)}{\Gamma(c)} = \frac{1}{\Gamma(a)\Gamma(b)} \sum_{n=0}^{\infty} \frac{\Gamma(a+n)\Gamma(b+n)}{\Gamma(c+n)} \frac{z^n}{n!}$$

The final expression for  $\phi_{nlm}(\underline{p})$ , which can also be found in [6] in a slightly different form, is given by:

$$\begin{aligned} \phi_{nlm}(\underline{p}) = & \sum_{\iota=0}^{2l+1} \frac{(-1)^\iota 2^{\iota+\frac{1}{2}} n (in)^l (p^2)^{l/2} Z^{-l-3} \Gamma(2l + \iota + 3)}{\iota!} \\ & \times \begin{pmatrix} l + n \\ -l + n - \iota - 1 \end{pmatrix} \sqrt{\frac{Z^3 \Gamma(n-l)}{\Gamma(l+n+1)}} \\ & \times Y_l^m(\theta_p, \phi_p) {}_2\tilde{F}_1\left(l + \frac{\iota}{2} + 2, \frac{1}{2}(2l + \iota + 3); l + \frac{3}{2}; -\frac{n^2 p^2}{Z^2}\right) \end{aligned} \quad (\text{A.3})$$

## Improved rate

Usually all that remains is to differentiate (A.3) with respect to  $\underline{p}$ . However, before that, the SFA rate can be improved directly by performing the integration over  $\phi_p$  analytically. The rate according to chapter 2 is given by:

$$\begin{aligned} \Gamma_{\text{SFA}}(t) = & \sum_{n_1} \sum_{n_2} \int_0^\infty dp p^2 \int_0^\pi d\theta_p \sin \theta_p \int_0^{2\pi} d\phi_p \int_{-\infty}^\infty dT \\ & \times \exp\left(i\mathbf{p}^2 T + ip \cos \theta_p \bar{\Delta}_z^k + \frac{i}{2} \bar{\Delta}_z^2 + i(t-T)E_{n_1} - i(t+T)E_{n_2}\right) \\ & \times E_z(t-T)E_z(t+T)c_{n_1}^*(t-T)c_{n_2}(t+T)d_{z,n_1}^*(\underline{p} + \Delta_z(-T))d_{z,n_2}(\underline{p} + \Delta_z(T)) \end{aligned}$$

As mentioned before, the dipole transition matrix element can be written in the following convenient form:

$$d_{z,nlm}(\underline{p}) = \left[ i \nabla_{\underline{p}} \phi_{nlm}(\underline{p}) \right]_z = i Y_{lm}(\theta_p, \phi_p) \left( \cos \theta_p \frac{\partial F_{nl}}{\partial p} - \frac{\sin \theta_p}{p} \frac{\partial F_{nl}}{\partial p} \right)$$

Note that here the  $z$  component of a gradient in spherical coordinates was taken. The exact formula can be easily computed using the transformation between Cartesian and spherical coordinates.

The integration over  $\theta_p$  is difficult and might not be possible to perform analytically. However, for the integral over  $\phi_p$ , only the following needs to be solved:

$$\begin{aligned} \int_0^{2\pi} d\phi_p Y_{l'm'}^*(\theta_p, \phi_p) Y_{lm}(\theta_p, \phi_p) &= \frac{1}{4\pi} \sqrt{(2l+1)(2l'+1) \frac{(l-m)!(l'-m')}{(l+m)!(l'+m')!}} \int_0^{2\pi} d\phi_p e^{i(m-m')\phi_p} \\ &= \frac{1}{2} \sqrt{(2l+1)(2l'+1) \frac{(l-m)!(l'-m)}{(l+m)!(l'+m)!}} \end{aligned}$$

This means that the magnetic quantum number is conserved during the transition between two arbitrary states. Since the initial state is the ground state  $1s$ ,  $m$  is zero here and therefore everywhere. This significantly simplifies the expression.



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# Declaration of Authorship

*Hiermit erkläre ich, die vorliegende Arbeit selbständig verfasst zu haben und keine anderen als die in der Arbeit angegebenen Quellen und Hilfsmittel benutzt zu haben.*

München, den 20.6.2025

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Unterschrift