

Investigation of the Higher Order Zeeman Effect

By

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DEDICATION

I would like to dedicate this thesis to ...

ACKNOWLEDGEMENTS

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CHAPTER 1

Introduction

- 1.1 Background and Motivation
- 1.2 Overview of Research Projects
- 1.3 Methodology Overview
- 1.4 Research Significance
- 1.5 Dissertation Structure

CHAPTER 2

Theoretical Methods

2.1 Overview

This chapter will provide an overview of all theoretical methods used throughout my dissertation. It will provide the theoretical building blocks upon which the dissertation is constructed, ranging from the set of units used in the problem, to perturbation equation solutions and methods for verifying calculations¹. As stated in the introduction, this thesis is based upon calculating matrix elements between two sets of wavefunctions connected by an operator. These matrix elements arise from perturbation theory and are necessary for computing energy shifts due to external fields such as those present in the higher-order Zeeman effect. The foundation of theory will be presented here, but the more specific calculations related to solving for the Zeeman effect will be presented in the relevant chapter below.

I will begin by first discussing atomic units in Sec. 2.2, then move on and introduce the Schrödinger equation for the one-electron problem as well as its solution in Sec. 2.3. In this section I will also discuss the significance and impact of the canonical momentum, and emphasize its importance in solving for the Zeeman effect. I will then introduce Perturbation Theory in Sec. 2.5, and describe how it can be used to approximate the new eigenvalues and eigenstates of the Zeeman-perturbed Hamiltonian. Sec. 2.4 will discuss various integration techniques that will be implemented both analytically and programmatically in order to simplify the calculation process, followed by an introduction to solving recursion relations in Sec. 2.6. This is the

¹Majority of the material discussed throughout this chapter are inspired and shaped by G. W. F. Drake. References will be included where needed.

core step in determining the perturbed wavefunctions used throughout the rest of the study. Finally, this chapter will conclude with the discussion of the Dalgarno Interchange Theorem and how it can serve as a critical check on the correctness of the obtained perturbed wavefunctions.

2.2 Atomic Units

First proposed by Hartree in November of 1928, atomic units have since become the standard across all of atomic physics. Atomic units were designed with the purpose of *”eliminating various universal constants from equations and also to avoid high powers of 10 in numerical work”* ???. Hartree proposed that we measure the mass in terms of the mass of the electron m_e , charge in terms of the electron charge, e , the vacuum permittivity in terms of $4\pi\epsilon_0$, and the angular momentum in terms of \hbar . Written mathematically we can say

$$\hbar = m_e = e = 4\pi\epsilon_0 = 1 \tag{1}$$

Scaling these units out of the problem greatly simplifies our calculation, and once a result is found, one can multiply the unscaled units back into the answer to retrieve the desired result in standard SI units. While this was proposed long before the field of computational physics emerged, it is of key importance that one implements this technique for any atomic physics simulation. Machine error is an unavoidable reality of programmed computation and working in units which avoids extreme values both large and small can help mitigate the uncertainty present in high precision calculations.

A consequence of utilizing atomic units is the need to redefine key fundamental physical quantities in terms of these scaled values. An example of key importance in the investigation of the higher-order Zeeman effect is the atomic unit of magnetic field strength, otherwise referred to as the atomic unit of magnetic flux density ??.

$$B_{\text{a.u.}} = \frac{\hbar}{ea_0^2} \quad (2)$$

This value is composed of the Bohr radius, which is defined as the most probable distance between an electron in the ground state of hydrogen and its nucleus. It can be described by the following equation ??

$$a_0 = \frac{4\pi\epsilon_0\hbar^2}{e^2m_e} = \frac{\hbar}{m_e c \alpha} \quad (3)$$

The Bohr radius itself is defined using a similar combination of fundamental constants which after applying the rules for atomic units, should equal 1. We can see now that all units in the atomic unit of field strength also become 1 as intended. At the end of our problem, we can simply multiply the determined result by the atomic unit of magnetic field strength when its constants are not scaled to 1, and retrieve our desired result in SI units. This is equivalent to multiplying the result in atomic units by ??.

$$B_{\text{a.u.}} = 2.35051757077(73) \times 10^5 \text{ T} \quad (4)$$

The numbers in parenthesis here denote the uncertainty of the quantity. Another key benefit in atomic units is that it allows us to define almost all fundamental quantities in terms of two constants; the speed of light, c and the fine structure constant, α .²

The fine structure constant can be written as the following dimensionless quantity

²The fine structure constant was introduced in 1916 by Arnold Sommerfeld when he was investigating the gap in the spectral lines of the hydrogen atom. He compared his work with the Michelson Morley experiment from 1887 where the spectral lines were shown and it was deduced that the universe is not permeated by the aether ??

$$\alpha = \frac{e^2}{4\pi\epsilon_0\hbar c} = 7.2973525643(11) \times 10^{-3} \quad (5)$$

which in atomic units is just the following relationship with the speed of light

$$\alpha = \frac{1}{c} \quad (6)$$

The fine structure constant is often used as an essential expansion parameter, treating relativity as a perturbation in α . More on this will be discussed in Sec. 3.6.3. Applying the atomic unit scaling to the fine structure constant shown in equation (6) confirms that the second equivalence in equation (3) remains equal to 1.

Applying this technique to problems in atomic physics gives a powerful way of simplifying calculations while providing an easy conversion factor to SI units for comparison of theory and experiment. This dissertation will discuss all equations and quantities in atomic units unless further specified otherwise.

2.3 One-electron Schrödinger equation

Discovered by Schrödinger in 1926, the Schrödinger equation is a non-relativistic solution to the two body problem involving an orbiting electron and its nucleus³. The electron interacts with the nucleus via the Coulomb interaction described by the following potential

$$V(\vec{r}) = -\frac{Ze^2}{4\pi\epsilon_0\vec{r}} \quad (7)$$

In atomic units

³It is assumed here that the nucleus is a point particle with infinite mass.

$$V(\vec{r}) = -\frac{Z}{r} \quad (8)$$

We can write the energy of the system using the Hamiltonian, where our potential is the Coulomb potential. We then come up with

$$H = \frac{p^2}{2m} - \frac{Z}{r} \quad (9)$$

However, quantum mechanics dictates that all observable quantities must be determined by hermitian operators that act on the wavefunction ψ . The wavefunction ψ is described as a normalized vector in Hilbert space, \mathcal{H} containing all information pertaining to the system. This means that the momentum needs to be a hermitian operator that acts on the wavefunction, and that our Hamiltonian is also a hermitian operator that satisfies the following eigenvalue problem

$$\hat{H}|\psi\rangle = E_n|\psi\rangle \quad (10)$$

This is what is known as the Schrödinger equation. It can be applied to our two-body problem by substituting the momentum for its quantum operator analog. This results in the following expression in atomic units

$$\hat{H}\psi = \frac{1}{2}\nabla^2\psi - \frac{Z}{r}\psi = E_n\psi \quad (11)$$

We now have a second-order partial differential equation. Assuming a separable solution allows one to split the solution to any hydrogenic system into two parts; The radial part and the angular part. For our two body atomic system the solutions for

each can be written in terms of Laguerre polynomials and Spherical Harmonics⁴.

$$\psi_{nlm} = R_{nl}(r)\mathcal{Y}_m^l(\cos\theta) \quad (12)$$

The radial equation has the following solutions

The ability to separate the solution into two independent parts is a key feature which will be exploited when evaluating integrals involving these wavefunctions in Chapter 3

2.4 Integration Techniques

2.4.1 The total integral

2.4.2 The angular part

2.4.3 The radial part

2.5 Perturbation Theory

To quantify how a magnetic field affects atomic energy levels, we must determine the shifts induced by the Zeeman interaction. Since this interaction modifies the Hamiltonian slightly compared to the field-free case, we can utilize perturbation theory to provide a systematic way to approximate the new eigenvalues and eigenstates.

2.6 Recursion relations

2.7 The Dalgarno Interchange Theorem

⁴The derivation will be shown in the appendix for both the angular and radial parts of the wavefunction.

CHAPTER 3

The Higher Order Zeeman Effect

3.1 Overview

3.2 History

3.3 Motivation

3.3.1 The $g - 2$ experiment

3.3.2 High-precision magnetometry

3.3.3 Connection to Atomic Physics

3.4 The Zeeman effect in Hydrogen

3.5 The Zeeman effect in Helium-3

3.6 The higher-order Zeeman effect

3.6.1 The quadratic Zeeman effect

The non-relativistic Hamiltonian for an n electron system in atomic units can be expressed as follows

$$\hat{H} = \sum_{i=1}^n \frac{\left(\vec{p}_i - e\vec{A}_i\right)^2}{2m} + V \quad (1)$$

using the canonical momentum instead of the classical momentum is essential to account for electromagnetic interactions. The vector potential \vec{A} can be described in terms of the magnetic field

$$\vec{A} = \frac{B}{2} (y\hat{x} - x\hat{y}) \quad (2)$$

which when expanded gives us the operator corresponding to the quadratic Zeeman effect

$$\hat{H}_Z^{(2)} = \frac{B^2 e^2}{8m} \sum_{i=1}^n (x_i^2 + y_i^2) \quad (3)$$

This operator can be expressed in spherical coordinates, written in terms of Legendre Polynomials

$$\hat{H}_Z^{(2)} = \frac{B^2 e^2}{12m} \sum_{i=1}^n r_i^2 (P_0(\cos \theta) - P_2(\cos \theta)) \quad (4)$$

${}^3\text{He}^+$ is a system which contains only one electron, so we can drop the summation which accounts for all electrons to get our final quadratic Zeeman operator

$$\hat{H}_Z^{(2)}({}^3\text{He}^+) = \frac{B^2 e^2}{12m} r^2 (P_0(\cos \theta) - P_2(\cos \theta)) \quad (5)$$

3.6.2 The magnetic dipole moment operator

The magnetic dipole moment operator represents the interaction of a magnetic dipole moment with an external magnetic field. It is described via the following relation

$$Q_{M1} = \mu_B \left(1 - \frac{2p^2}{3m^2 c^2} + \frac{Ze^2}{3mc^2 r} \right) \vec{\sigma} \cdot \vec{B} \quad (6)$$

Where μ_B is the Bohr magneton

$$\mu_B = \frac{e\hbar}{2mc} \quad (7)$$

The second term in the brackets of the magnetic dipole moment operator accounts

for the relativistic correction to the kinetic energy of the electron, and the third term is the potential energy due to the Coulomb interaction between the electron and the nucleus. The first term corresponds to the ordinary Zeeman Effect, which does not contribute to the sum over states due to orthogonality.

The ordinary Zeeman effect contributes to Q_{M1} in ${}^3\text{He}^+$ because it has non-zero spin due to the missing electron. For systems such as ${}^3\text{He}$, the ordinary Zeeman effect will not contribute.

3.6.3 The relativistic correction to ${}^3\text{He}^+$

Combining the magnetic dipole moment with the quadratic Zeeman operator, we can write down the relativistic corrections for ${}^3\text{He}^+$. Written in terms of pseudostates the relativistic correction is

$$C_{\text{rel}}^{(2)} = \sum_{\substack{n=-\infty \\ n \neq 0}}^{\infty} \frac{\langle \psi_0 | H_Z^{(2)} | \psi_n \rangle \langle \psi_n | Q_{M1} | \psi_0 \rangle}{E_0 - E_n} \quad (8)$$

CHAPTER 4

Conclusion and Future Work

4.1 Overview

4.2 Consolidation of Results

4.3 Synthesis of Conclusions

4.4 Future Work

4.5 Theoretical and Practical Impact

APPENDIX A

Methods of solving for higher-order perturbations programmatically

APPENDIX B

Derivation of the Hydrogen Atom

APPENDIX C

Derivation of the Recursion Relations

APPENDIX D

Special Functions

D.1 Overview

D.2 Laguerre Polynomials

D.3 Legendre Polynomials

D.4 Spherical Harmonics

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