Visualization of the hydrogen atom orbitals

Derek W. Harrison

February 1, 2021

## Introduction

The orbitals of the hydrogen atom are visualized using the wave equations obtained from analytical solution of the Schrödinger equation for the hydrogen atom.

## Model equations

The three-dimensional time independent Schrödinger equation for the hydrogen atom is:

$$E\psi = -\frac{\hbar^2}{2m}\nabla^2\psi - \frac{k}{r}\psi\tag{1}$$

Where  $\hbar$  is a constant, m the particle mass, k a constant depending on the charge of the electron, r the radial coordinate,  $\psi$  the wave equation and E the associated energy. Expanding the Laplacian in (1) in spherical coordinates gives:

$$E\psi = -\frac{\hbar^2}{2m} \left[ \frac{1}{r^2} \frac{\partial}{\partial r} \left( \frac{r^2 \partial \psi}{\partial r} \right) + \frac{1}{r^2 \sin(\phi)} \frac{\partial}{\partial \phi} \left( \sin(\phi) \frac{\partial \psi}{\partial \phi} \right) + \frac{1}{r^2 \sin^2(\phi)} \frac{\partial^2 \psi}{\partial \theta^2} \right] - \frac{k}{r} \psi$$
(2)

Where  $\phi$  is the polar angle and  $\theta$  the azimuthal angle. Note that often  $\theta$  is used instead of  $\phi$  to denote the polar angle and similarly  $\phi$  is used to denote the azimuthal angle, which is the opposite convention used here. The boundary conditions of the system are  $\psi = 0$  at r = 0 and as r grows large.

Analytical solution of equation (2) produces three quantum numbers n, l and m where n is the principal quantum number, l is the azimuthal quantum number and m is the magnetic quantum number. The resulting wave equations depend on these quantum numbers in addition to the spatial coordinates  $r, \theta$  and  $\phi$ . The wave equations are:

$$\psi_{nlm}(r,\theta,\phi) = Ce^{-\rho/2} \rho^{l} L_{n-l-1}^{2l+1}(\rho) e^{im\theta} P_{l}^{m}(\cos\phi)$$
 (3)

Where C is a normalization constant,  $\rho = \frac{2r}{na_0}$  with  $a_0$  the reduced Bohr radius,  $L_{n-l-1}^{2l+1}(\rho)$  is the generalized Laguerre polynomial, i the imaginary number and  $P_l^m(\cos\phi)$  the associated Legendre polynomial. The normalization constant is obtained by integrating the product of  $\psi$ , as given by (3), and its complex conjugate  $\psi^*$  over the domain V, equating the integral with 1 and solving for C:

$$1 = \int_{V} \psi \psi^* dV \tag{4}$$

The generalized Laguerre polynomials are given by the following recurrence relation:

$$L_k^{\alpha}(x) = \frac{2k - 1 + \alpha - x}{k} L_{k-1}^{\alpha}(x) - \frac{k - 1 + \alpha}{k} L_{k-2}^{\alpha}(x)$$
 (5)

Where  $\alpha$  and k are integers and x is the independent variable. The first two Laguerre polynomials, which form the base cases for the recurrence relation, are:

$$L_0^{\alpha}(x) = 1 \tag{6}$$

$$L_1^{\alpha}(x) = 1 + \alpha - x \tag{7}$$

The associated Legendre polynomials are computed using the following relation:

$$P_l^m(x) = (-1)^m (1 - x^2)^{m/2} \frac{d^m}{dx^m} P_l(x)$$
(8)

With  $P_l(x)$  the Legendre polynomial, which can be computed using the Rodrigues formula:

$$P_l(x) = \frac{1}{2^l l!} \frac{d^l}{dx^l} (x^2 - 1)^l \tag{9}$$

Equation (9) can be expanded to:

$$P_l(x) = \frac{1}{2^l} \sum_{k=0}^{\lfloor l/2 \rfloor} \frac{(-1)^k (2l-2k)!}{k! (l-k)! (l-2k)!} x^{l-2k}$$
(10)

Should the magnetic quantum number m be negative the associated Legendre polynomial is computed using:

$$P_l^{-|m|}(x) = (-1)^m \frac{(l-|m|)!}{(l+|m|)!} P_l^{|m|}(x)$$
(11)

## Visualization

The probability density is computed for various values of the quantum numbers n, l and m. Results are shown in figures 1 to 5. Note that some regions in the graphs are colored white when they should be colored dark blue, corresponding to  $\psi^2 = 0$ . The graphs represent cross-sections at  $\theta = 0$ .

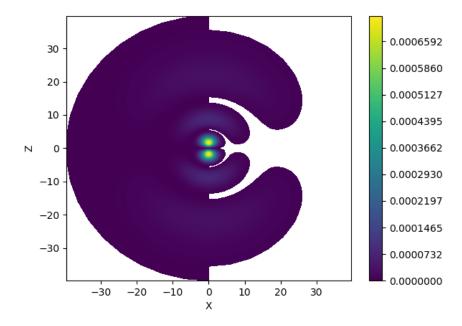


Figure 1: Probability density  $\psi_{410}^2$  of the 410 or  $4p_0$  orbital.

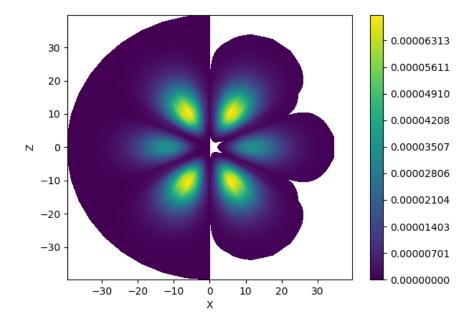


Figure 2: Probability density  $\psi_{431}^2$  of the 431 or  $4f_1$  orbital.

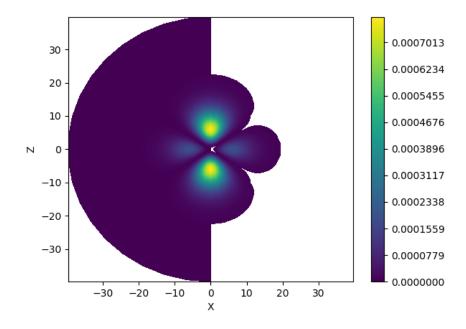


Figure 3: Probability density  $\psi_{320}^2$  of the 320 or  $3d_0$  orbital.

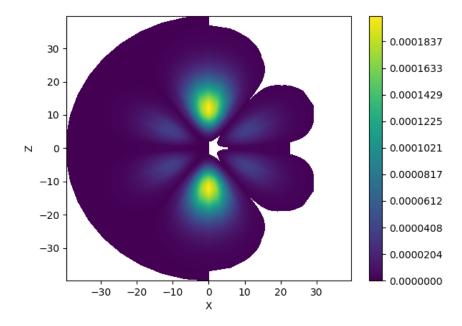


Figure 4: Probability density  $\psi_{430}^2$  of the 430 or  $4f_0$  orbital.

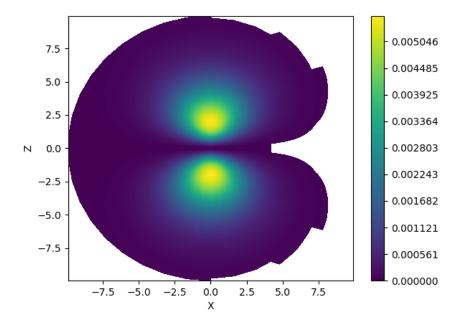


Figure 5: Probability density  $\psi_{210}^2$  of the 210 or  $2p_0$  orbital.

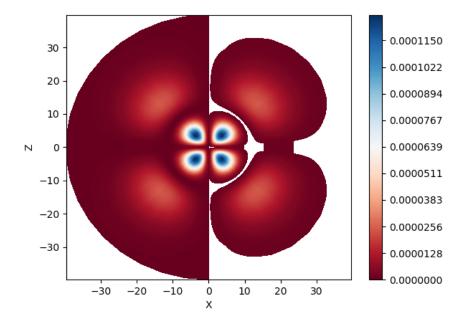


Figure 6: Probability density  $\psi_{421}^2$  of the 421 or  $4d_1$  orbital.

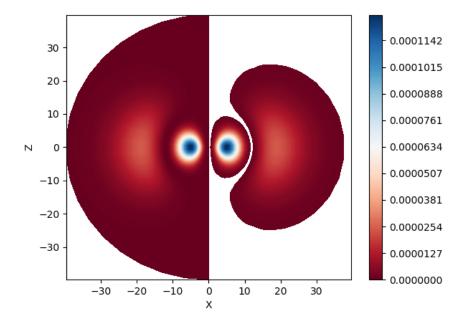


Figure 7: Probability density  $\psi_{422}^2$  of the 422 or  $4d_2$  orbital.