The Hydrogen Atom

Thornton and Rex, Ch. 7

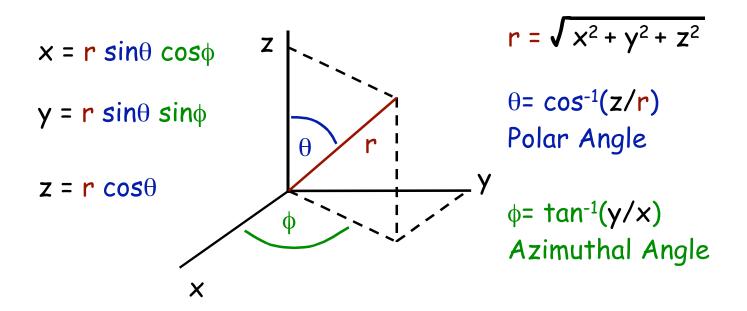
Applying Schrodinger's Eqn to the Hydrogen Atom

The potential:
$$V(r) = \frac{-1}{4\pi \epsilon_0} \frac{e^2}{r}$$

Use spherical polar coordinates (with $\psi(x,y,z) \Rightarrow \psi(r,\theta,\phi)$):

$$\frac{1}{r^2}\frac{\partial}{\partial r}\left(r^2\frac{\partial\psi}{\partial r}\right) + \frac{1}{r^2\sin\theta}\frac{\partial}{\partial\theta}\left(\sin\theta\frac{\partial\psi}{\partial\theta}\right)$$

$$+ \frac{1}{r^2 \sin^2 \theta} \frac{\partial^2 \psi}{\partial \phi^2} + \frac{2m}{\hbar^2} (E-V) \psi = 0$$



 $\psi(r,\theta,\phi)$ is separable:

$$\Rightarrow \psi(r,\theta,\phi) = R(r) f(\theta) g(\phi)$$

Substitute this into S Eqn and apply appropriate boundary conditions to R, f, g.

 \Rightarrow 3 separate equations and 3 quantum numbers. (For more, see section 7.2.)

$$\frac{d^2g}{d\phi^2} = -m_\ell^2g$$

Azimuthal Eqn.

$$\frac{1}{r^2}\frac{d}{dr}\left(r^2\frac{dR}{dr}\right) + \frac{2m}{\hbar^2}\left(E-V-\frac{\hbar}{2m}\frac{\ell(\ell+1)}{r^2}\right)R = 0$$

Radial Egn.

$$\frac{1}{\sin\theta} \frac{d}{d\theta} \left(\sin\theta \frac{df}{d\theta} \right) + \left(\ell \left(\ell + 1 \right) - \frac{m_{\ell}^2}{\sin^2\theta} \right) f = 0$$

Angular Eqn.

 m_{ℓ} and ℓ are quantum numbers.

The Radial Equation

The Radial Equation is the Associated Laguerre Equation.

We will find the ground-state solution. Require $m_{\ell} = 0$, $\ell = 0$.

$$\Rightarrow \frac{1}{r^2} \frac{d}{dr} \left(r^2 \frac{dR}{dr} \right) + \frac{2m}{\hbar^2} (E-V) R = 0$$

Substitute $V = -e^2/(4\pi\epsilon_0 r)$ and insert a trial solution:

$$R = A e^{-r/a_0}$$

This works if

$$a_0 = \frac{4\pi\epsilon_0 \, \hbar^2}{m \, e^2} \qquad \qquad \text{(Bohr Radius)}$$

and
$$E = -\frac{\hbar^2}{(2ma_0^2)} = -E_0 = -13.6 \text{ eV}$$

Higher order solutions can be found in terms of associated Laguerre functions.

They are labeled by a quantum number n (called the <u>principal quantum number</u>).

Energies are

$$E = - E_0 / n^2$$

(just like the Bohr prediction.)

Angular and Azimuthal Equations

The azimuthal equation is just a SHO equation with solution $g = A e^{im\phi}$.

Single-valuedness requires $g(\phi) = g(\phi + 2\pi)$.

 \Rightarrow m_{ℓ} is an integer.

The angular equation is the <u>Associated</u> <u>Legendre Equation</u>.

It is customary to combine the θ and ϕ solutions together as Spherical Harmonics $Y(\theta,\phi)$

The quantum numbers satisfy

$$\ell = 0, 1, 2, 3, \dots$$

and

$$m_{\ell} = -\ell, -\ell+1, -\ell+2, \ldots, 0, \ldots, \ell-1, \ell$$

(They also satisfy $\ell < n$.)

$|Y(\theta,\phi)|^2$

$$m_{\ell}=0$$

$$m_{\ell}=\pm 1$$

$$m_{\ell}=\pm 2 \pm 1$$

$$m_{\ell}=\pm 3$$
 ± 2 ± 1 C

Atomic Quantum Numbers

- n Principal Quantum Number
- Orbital Angular Momentum

 Quantum Number
- m_e Magnetic Quantum Number

$$n = 1, 2, 3, ...$$

 $\ell = 0, 1, 2, 3, ...$
 $m_{\ell} = -\ell, -\ell + 1, -\ell + 2, ..., 0, ..., \ell - 1, \ell$

In summary:

$$n > 0$$
 $\ell < n$
 $|m_{\ell}| \le \ell$

Angular Momentum

Angular momentum of electron in the atom:

$$L = mvr = \sqrt{\ell(\ell+1)}$$
 ħ

(Note that this disagrees with Bohr's original guess of $L = n \pi$.)

For a given n, the energy is $E_n = -E_0/n^2$ independent of ℓ .

The different ℓ states are degenerate.

Historical Notation:

$$\ell = 0$$
 1 2 3 4 5
1 1 1 1 1
s p d f g h

States are usually labeled by the n number and the letter.

For example: n=3, $\ell=1$ \longrightarrow 3p state.

Magnetic Quantum Number

¿ determines the total angular momentum:

$$L = \sqrt{\ell(\ell+1)}$$
 ħ

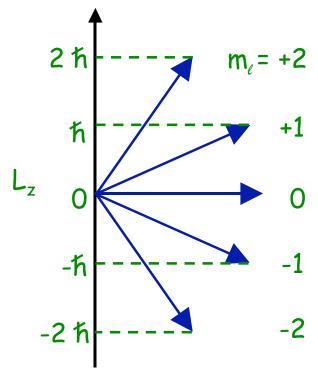
 m_{ℓ} gives the z component of L:

$$L_z = m_e \hbar$$

Direction of \vec{L} can never lie on z axis:

 $m_{\ell} < \sqrt{\ell(\ell+1)}$ always.

An example for $\ell = 2$ $\Rightarrow L = \sqrt{6} \, \hbar$

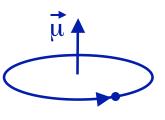


The choice of direction for the z axis is completely arbitrary.

 L_x and L_y are undetermined, except for $L_x^2 + L_y^2 = L^2 - L_z^2$

Magnetic Effects

An electron orbiting around a nucleus has magnetic moment $\vec{\mu}$:



$$\vec{\mu}$$
 = IA \hat{n} = $\frac{-e}{(2\pi r/v)} (\pi r^2) \hat{n}$ = $\frac{-erv}{2} \hat{n}$ = $\frac{-e}{2m} \vec{L}$

The component in the z direction is:

$$\mu_z = \frac{-e}{2m} L_z = \frac{-e}{2m} m_\ell \hbar = -m_\ell \mu_B$$

where
$$\mu_B = e \pi/2m$$

= 9.274×10⁻²⁴J/T
= 5.788×10⁻⁵ eV/T

is the Bohr magneton.

In an external magnetic field, B, the magnetic dipole feels a torque:

$$\vec{\tau} = \vec{\mu} \times \vec{B}$$

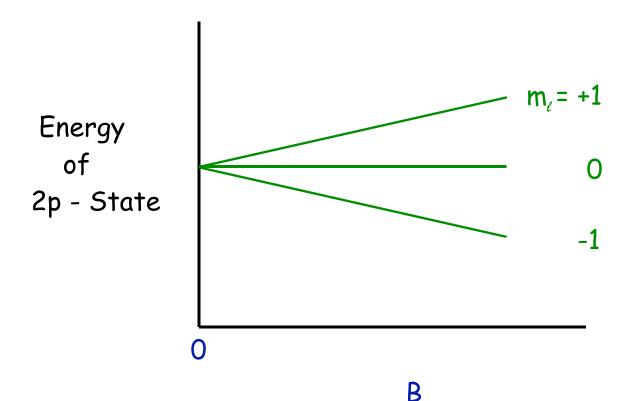
and has a potential energy: $V_B = -\vec{\mu} \cdot \vec{B}$

If \vec{B} is in the z direction then

$$V_B = -\mu_z B = +m_\ell \mu_B B$$

The energies for different m_{ℓ} , which were degenerate for B=0, are now separated into $2\ell+1$ different levels.

This is called the Normal Zeeman Effect.



The Stern-Gerlach Experiment

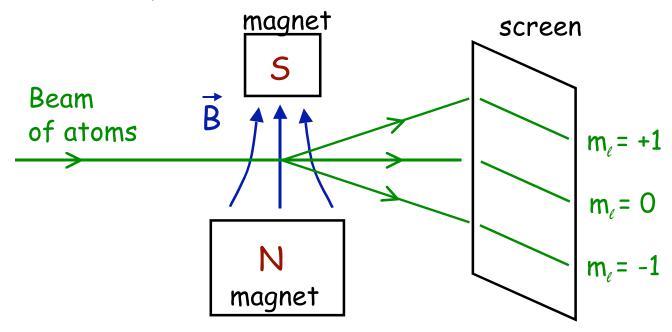
In the presence of an inhomogeneous magnetic field, there will be a net force on the atoms, depending on m_{ℓ} :

$$F_z = -\frac{dV_B}{dz} = m_\ell \mu_B \frac{dB}{dz}$$

i.e. In the +z direction for
$$+m_{\ell}$$

No force for $m_{\ell}=0$
In the -z direction for $-m_{\ell}$

Thus, one could split the atoms according to the quantum number m_{ℓ} :



Electron Spin

1922 - Stern and Gerlach did their experiment. The atoms split into two beams.

But the number of m_{ℓ} values is always odd: $(2\ell + 1)!$

1925 - Goudsmit and Uhlenbeck proposed that the electron had an intrinsic spin and an intrinsic magnetic moment.

In analogy with orbital angular momentum they proposed a magnetic spin quantum number:

$$m_s = \pm 1/2$$



The electron's spin can either be oriented "up" or "down":

$$+\frac{1}{2}\hbar$$
 $-\frac{1}{2}\hbar$

The total spin quantum number is s = 1/2:

$$|\vec{S}| = \sqrt{s(s+1)} \, \text{ h} = \sqrt{3/4} \, \text{ h}$$

$$\vec{\mu}_S = \frac{-e}{m} \vec{S} = -[2] \frac{\mu_B}{\hbar} \vec{S}$$

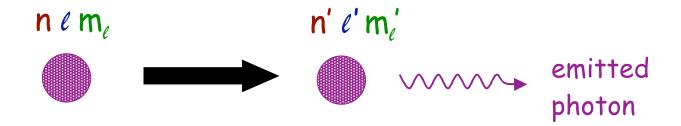
Compare with:

$$\vec{\mu}_{L} = \frac{-e}{2m} \vec{L} = -[1] \frac{\mu_{B}}{\hbar} \vec{L}$$

[*] are called gyromagnetic ratios:

$$g_s = 2$$
 $g_\ell = 1$

Selection Rules



Allowed transitions:

• lifetimes $\tau \sim 10^{-9}$ sec

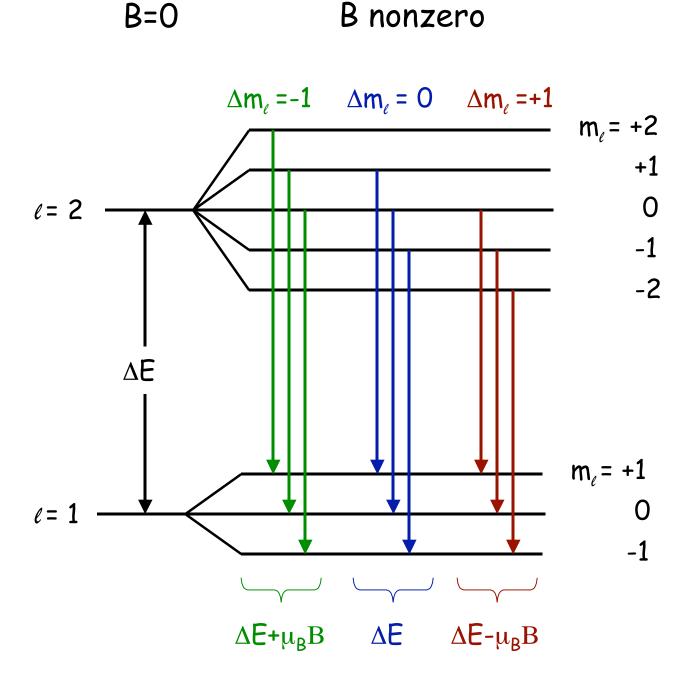
$$\Delta$$
 n = anything, $\Delta \ell = \pm 1$, $\Delta m_{\ell} = 0$, ± 1

Forbidden transitions:

lifetimes much longer

Ex.
$$2s \rightarrow 1s$$
, $\tau \sim 1/7$ sec

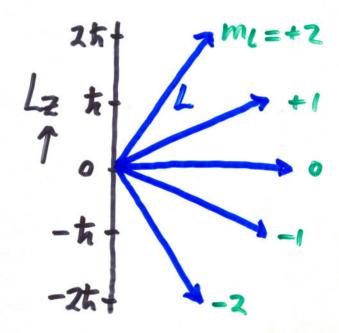
Selection Rules and Normal Zeeman Effect



B field always splits spectral lines into 3 for normal Zeeman effect.

7.14 FOR A 3d STATE DRAW ALL THE POSSIBLE ORIENTATIONS OF THE ANGULAR MOMENTUM VECTOR L.
WHAT IS $L_x^2 + L_y^2$ FOR THE $M_t = -1$ COMPONENT?

Possible values of $m_l = -2$ -1 0 +1 +2 with $L_2 = m_l t$



FOR
$$m_1 = -1$$
 $\Rightarrow L_2 = -\pi$

$$L^2 = L_x^2 + L_y^2 + L_z^2$$

$$\therefore 6\pi^2 = L_x^2 + L_y^2 + \pi^2$$

$$\therefore Lx^2 + Ly^2 = 5h^2$$

7.25 A HYDROGEN ATOM IN A 55 STATE IS IN A MAGNETIC FIELD OF 3T. WHAT IS THE ENERGY IN THE ABSENCE OF THE MAG. FIELD ? HOW MANY STATES ARE THERE . AND WHAT ARE THEIR ENERGIES IN THE MAG. FIELD? n=5 l=3 $m_{l}=0,\pm 1,\pm 2,\pm 3$ $E_5 = -\frac{13.6}{35} = -0.544 \text{ eV}$ VB = m, BMB = m, 3.5.79E-5 ex => = 0 FOR M1 = 0 = ±1.74E-4 eV FOR m1 = ±1 = ±3.47E-4 eV FOR ML = ±2 = ± 5.21 E-4 eV FOR m1 = ± 3

