

# PHYS 3313 – Section 001

## Lecture #19

*Monday, Nov. 18, 2013*

*Dr. Jaehoon Yu*

- Orbital Angular Momentum Quantum Number
- Magnetic Quantum Number
- The Zeeman Effect
- Intrinsic Spin
- Electron Energy Levels
- Selection Rules
- Probability Distributions
- Statistics

Monday, Nov. 18, 2013



PHYS 3313-001, Fall 2013  
Dr. Jaehoon Yu

# Announcements

- Homework #7
  - CH7 end of chapter problems: 7, 8, 9, 12, 17 and 29
  - Due on Monday, Nov. 25, in class
- Quiz number 4
  - At the beginning of the class Monday, Nov. 25
  - Covers CH7 and what we finish this Wednesday
- Reading assignments
  - Entire CH8 (in particular CH8.1), CH9.4 and CH9.7
- Class is cancelled on Wednesday, Nov. 27



Group Number	Research Group Members	Research Topic	Presentation Date and Order
1	Z.Citty, S. Lagerson, K. McElvain, J. Vellarreal	Michelson-Morley Experiment	#6, Dec. 2
2	W. Brown, C. Hair, R. Reyes, H. Zapata	The Photo-Electric Effect	#2, Dec. 2
3	R. Clark, M. Kruse, C. Nguyen, B. Watson	The Unification of Electromagnetic and Weak Forces	#3, Dec. 2
4	J. Bolton, J. Day, B. Nuar,	Discovery of Electron	#6, Dec. 4
5	J. Bowerman, C. McNutt, M. Obiang, E. Perez	The property of molecules - the Brownian Motions	#4, Dec. 2
6	N. Boseman, V. Hopkins, S. Moorman, S. Moriaty	Black-body Radiation	#1, Dec. 4
7	E. Bainglass, J. Chavez, K. Izuagbe,	Rutherford Scattering	#3, Dec. 4
8	E. Blomberg, E. Duran, J. Grandinatti, R. Loew	The Discovery of Radioactivity	#1, Dec. 2
9	P. Conlin, J. Guevara, F. Islam, A. Nelson	Special Relativity	#5, Dec. 2
10	G. Brown, G. Collier, B. Ferguson, R. Subramaniam	Compton Effect	#2, Dec. 4
11	K. Brackney, C. Dunn, S. Schroeder, S. Sheladia	Super-Conductivity	#4, Dec. 4
12	A. Patten, C. Jay, C. Smith, J. Umphress	PHYS 3313-001, Fall 2013. The discovery of the Higgs particle Dr. Jaehoon Yu	<b>3</b> #5, Dec. 4

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# Research Project Report

1. Must contain the following at the minimum
  - Original theory or Original observation
  - Experimental proofs or Theoretical prediction + subsequent experimental proofs
  - Importance and the impact of the theory/experiment
  - Conclusions
2. Each member of the group writes a 10 (max) page report, including figures
  - 10% of the total grade
  - Can share the theme and facts but you must write your own!
  - Text of the report must NOT be a copy
  - **Due Mon., Dec. 2, 2013**



# Research Presentations

- Each of the 10 research groups makes a 10min presentation
  - 8min presentation + 2min Q&A
  - All presentations must be in power point
  - I must receive all final presentation files by 8pm, Sunday, Dec. 1
    - No changes are allowed afterward
  - The representative of the group makes the presentation followed by all group members' participation in the Q&A session
- Date and time:
  - In class Monday, Dec. 2 or in class Wednesday, Dec. 4
- Important metrics
  - Contents of the presentation: 60%
    - Inclusion of all important points as mentioned in the report
    - The quality of the research and making the right points
  - Quality of the presentation itself: 15%
  - Presentation manner: 10%
  - Q&A handling: 10%
  - Staying in the allotted presentation time: 5%
  - Judging participation and sincerity: 5%



# Solution of the Angular and Azimuthal Equations

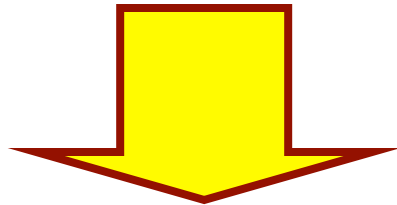
- The radial wave function  $R$  and the spherical harmonics  $Y$  determine the probability density for the various quantum states.
- Thus the total wave function  $\psi(r,\theta,\phi)$  depends on  $n$ ,  $\ell$ , and  $m_\ell$ . The wave function can be written as

$$\psi_{nlm_\ell}(r,\theta,\phi) = R_{nl}(r)Y_{lm_\ell}(\theta,\phi)$$



# Orbital Angular Momentum Quantum Number $\ell$

- It is associated with the  $R(r)$  and  $f(\theta)$  parts of the wave function.
- Classically, the orbital angular momentum  $\vec{L} = \vec{r} \times \vec{p}$  with  $L = mv_{\text{orbital}}r$ .
- $\ell$  is related to the magnitude of  $L$  by  $L = \sqrt{l(l+1)}\hbar$ .
- In an  $\ell = 0$  state,  $L = \sqrt{0(1)}\hbar = 0$ .



It disagrees with Bohr's semi-classical “planetary” model of electrons orbiting a nucleus  $L = n\hbar$ .

# Orbital Angular Momentum Quantum Number $\ell$

- Certain energy level is **degenerate** with respect to  $\ell$  when the energy is independent of  $\ell$ .
- Use letter names for the various  $\ell$  values
  - $\ell =$             0        1        2        3        4        5 ...
  - Letter =        s        p        d        f        g        h ...
- Atomic states are referred by their  $n$  and  $\ell$ 
  - **s**=sharp, **p**=principal, **d**=diffuse, **f**=fundamental, then alphabetical
- A state with  $n = 2$  and  $\ell = 1$  is called the  $2p$  state
  - Is  $2d$  state possible?
- The boundary conditions require  $n > \ell$



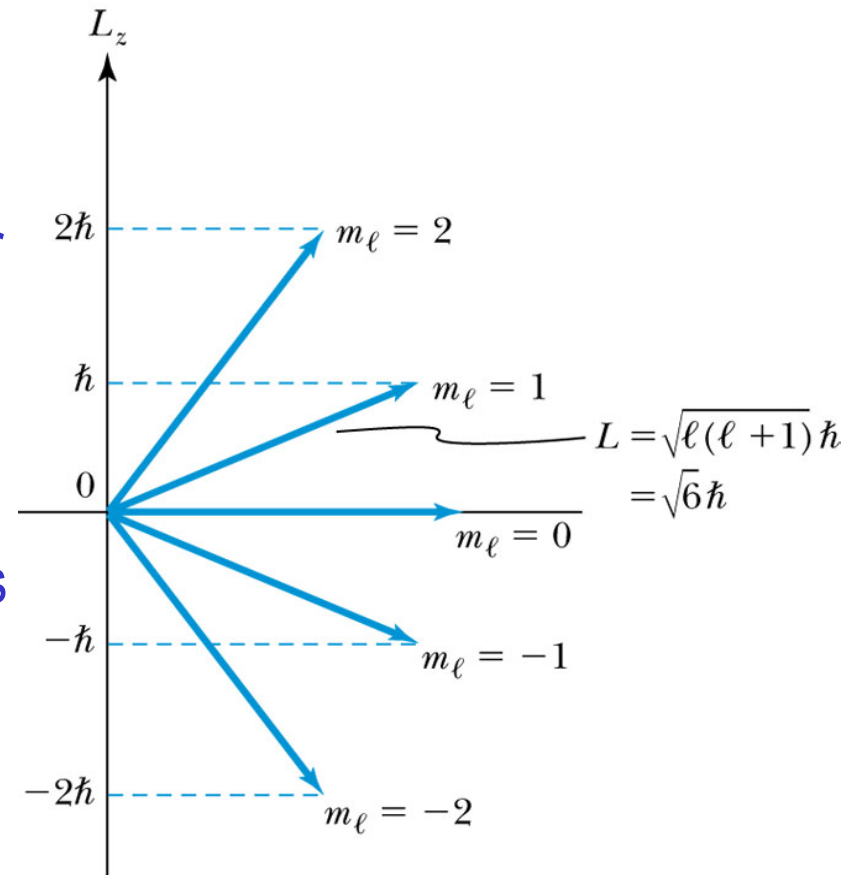


# Magnetic Quantum Number $m_\ell$

- The angle  $\phi$  is a measure of the rotation about the z axis.
- The solution for  $g(\phi)$  specifies that  $m_\ell$  is an integer and related to the z component of  $L$ .

$$L_z = m_\ell \hbar$$

- The relationship of  $L$ ,  $L_z$ ,  $\ell$ , and  $m_\ell$  for  $\ell = 2$ .
- $L = \sqrt{l(l+1)}\hbar = \sqrt{6}\hbar$  is fixed.
- Because  $L_z$  is quantized, only certain orientations of  $\vec{L}$  are possible and this is called **space quantization**.
- $m_\ell$  is called the magnetic moment since z axis is chosen customarily along the direction of magnetic field.



# Magnetic Quantum Number $m_\ell$

- Quantum mechanics allows  $\vec{L}$  to be quantized along only one direction in space and because of the relationship  $L^2 = L_x^2 + L_y^2 + L_z^2$ , once a second component is known, the third component will also be known.  $\rightarrow$  violation of uncertainty principle
  - One of the three components, such as  $L_z$ , can be known clearly but the other components will not be precisely known
- Now, since we know there is no preferred direction,

$$\langle L_x^2 \rangle = \langle L_y^2 \rangle = \langle L_z^2 \rangle$$

- We expect the average of the angular momentum components squared to be:  $\langle L^2 \rangle = 3\langle L_z^2 \rangle = \frac{3}{2l+1} \sum_{m_l=-l}^{+l} m_l^2 \hbar^2 = l(l+1)\hbar^2$

# Magnetic Effects on Atomic Spectra— The Normal Zeeman Effect

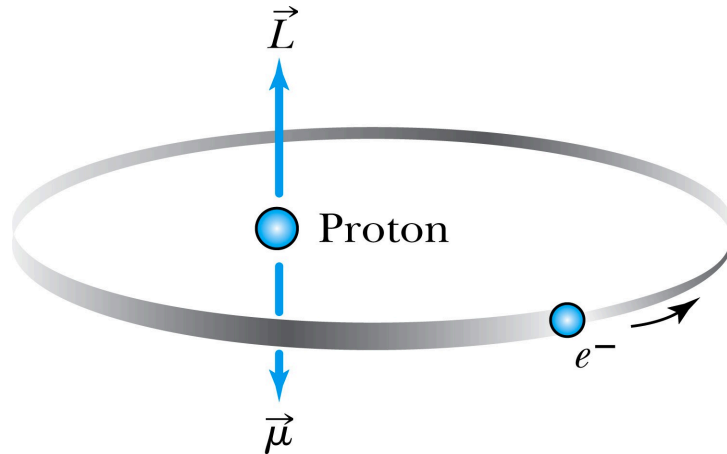
- A Dutch physicist Pieter Zeeman showed as early as 1896 that the spectral lines emitted by atoms in a magnetic field split into multiple energy levels. It is called the **Zeeman effect**.

## The **Normal** Zeeman effect:

- A spectral line of an atom is split into three lines.
- Consider the atom to behave like a small magnet.
- The current loop has a magnetic moment  $\mu = IA$  and the period  $T = 2\pi r / v$ . If an electron can be considered as orbiting a circular current loop of  $I = dq / dt$  around the nucleus, we obtain
$$\mu = IA = qA/T = \pi r^2 (-e)/(2\pi r/v) = -erv/2 = -\frac{e}{2m} mrv = -\frac{e}{2m} L$$
- $\vec{\mu} = -\frac{e}{2m} \vec{L}$  where  $L = mvr$  is the magnitude of the orbital angular momentum



# The Normal Zeeman Effect



- Since there is no magnetic field to align them,  $\vec{\mu}$  points in random directions.
- The dipole has a potential energy

$$V_B = -\vec{\mu} \cdot \vec{B}$$

- The angular momentum is aligned with the magnetic moment, and the torque between  $\vec{\mu}$  and  $\vec{B}$  causes a precession of  $\vec{\mu}$ .

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

Where  $\mu_B = e\hbar / 2m$  is called the **Bohr magneton**.

- $\vec{\mu}$  cannot align exactly in the z direction and has only certain allowed quantized orientations.

$$\vec{\mu} = -\frac{\mu_B \vec{L}}{\hbar}$$

# The Normal Zeeman Effect

- The potential energy is quantized due to the magnetic quantum number  $m_\ell$ .

$$V_B = -\mu_z B = +\mu_B m_l B$$

- When a magnetic field is applied, the  $2p$  level of atomic hydrogen is split into three different energy states with the electron energy difference of  $\Delta E = \mu_B B \Delta m_\ell$ .

$m_\ell$	Energy
1	$E_0 + \mu_B B$
0	$E_0$
-1	$E_0 - \mu_B B$

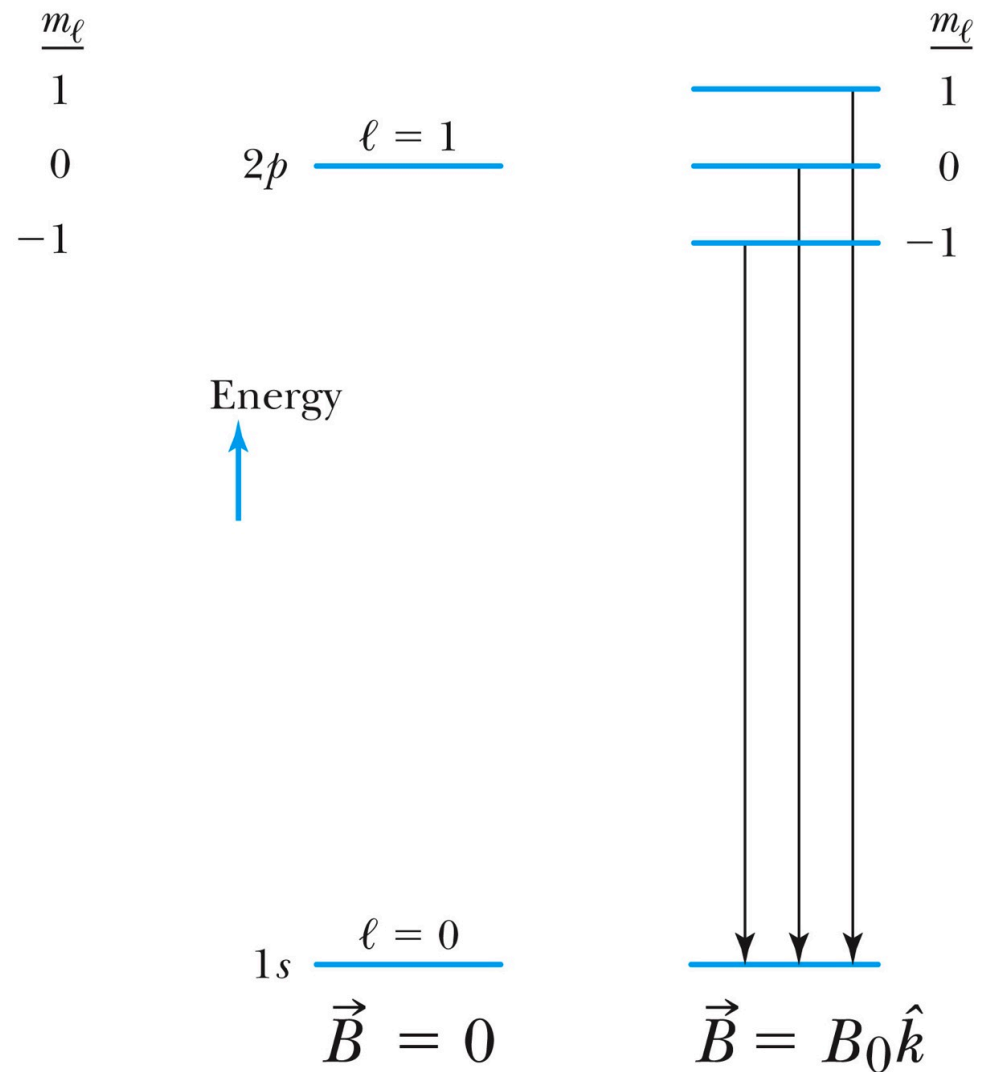
$$n = 2 \quad \underline{\ell = 1}$$

$$\vec{B} = 0$$

- So split is into a total of  $2\ell+1$  energy states

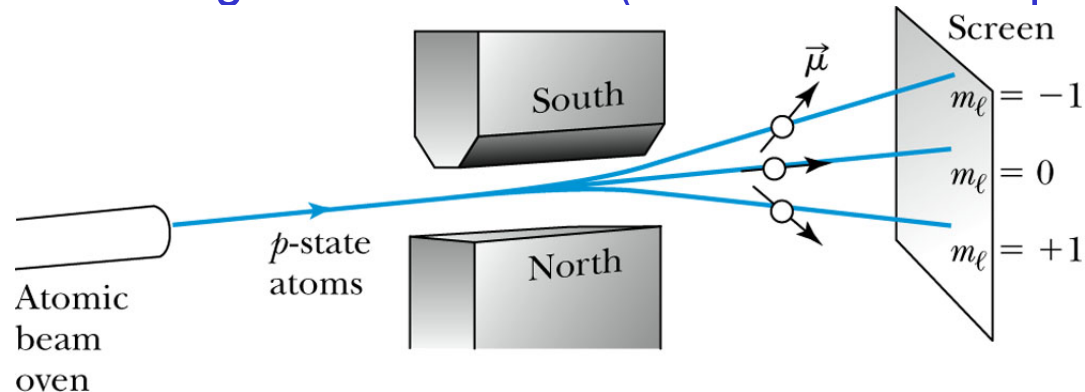
# The Normal Zeeman Effect

- A transition from 1s to 2p
- A transition from 2p to 1s



# The Normal Zeeman Effect

- An atomic beam of particles in the  $\ell = 1$  state pass through a magnetic field along the z direction. (Stern-Gerlach experiment)



- $V_B = -\mu_z B$
- $F_z = -(dV_B/dz) = \mu_z (dB/dz)$
- The  $m_\ell = +1$  state will be deflected down, the  $m_\ell = -1$  state up, and the  $m_\ell = 0$  state will be undeflected. → saw only 2 with silver atom
- If the space quantization were due to the magnetic quantum number  $m_\ell$ , the number of  $m_\ell$  states is always odd at  $(2\ell + 1)$  and should have produced an odd number of lines.

# Intrinsic Spin

- In 1920, to explain spectral line splitting of Stern-Gerlach experiment, Wolfgang Pauli proposed the forth quantum number assigned to electrons
- In 1925, Samuel Goudsmit and George Uhlenbeck in Holland proposed that *the electron must have an intrinsic angular momentum* and therefore a magnetic moment.
- Paul Ehrenfest showed that the surface of the spinning electron should be moving faster than the speed of light to obtain the needed angular momentum!!
- In order to explain experimental data, Goudsmit and Uhlenbeck proposed that the electron must have an **intrinsic spin quantum number  $s = \frac{1}{2}$** .





# Intrinsic Spin

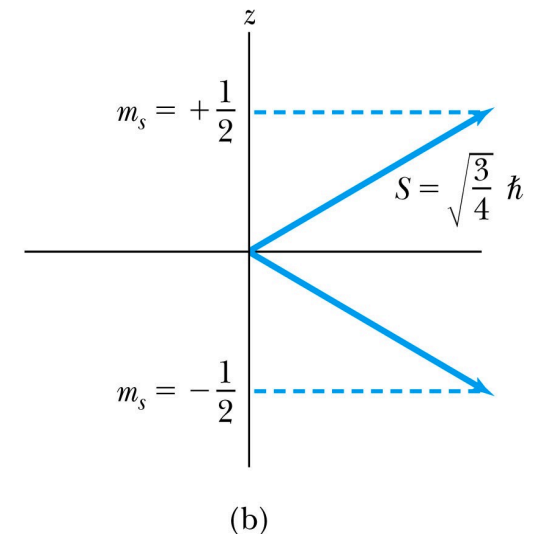
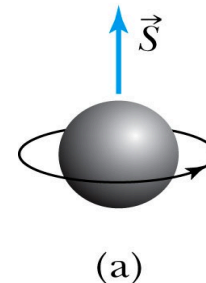
- The spinning electron reacts similarly to the orbiting electron in a magnetic field. (Dirac showed that this is necessary due to special relativity..)
- We should try to find  $L$ ,  $L_z$ ,  $\ell$ , and  $m_\ell$ .
- The **magnetic spin quantum number**  $m_s$  has only two values,  $m_s = \pm 1/2$ .

The electron's spin will be either “up” or “down” and can never be spinning with its magnetic moment  $\mu_s$  exactly along the z axis.

For each state of the other quantum numbers, there are two spins values

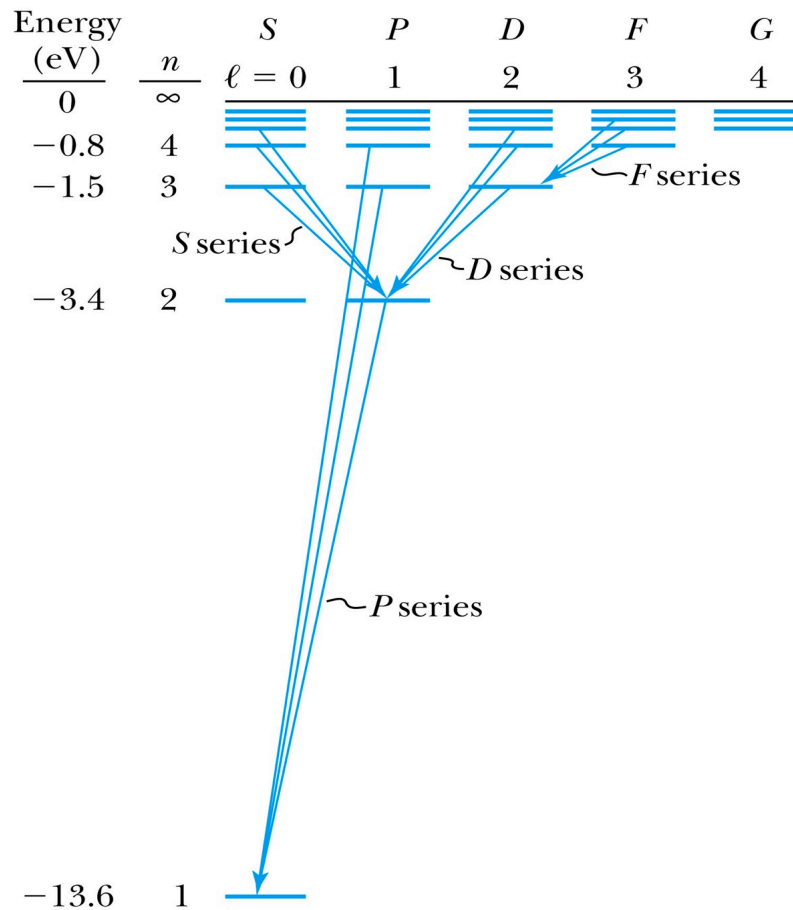
The **intrinsic spin angular momentum**

vector  $|\vec{S}| = \sqrt{s(s+1)}\hbar = \sqrt{3/4}\hbar$



# Energy Levels and Electron Probabilities

- For hydrogen, the energy level depends on the principle quantum number  $n$ .



- In ground state an atom cannot emit radiation. It can absorb electromagnetic radiation, or gain energy through inelastic bombardment by particles.

# Selection Rules

- We can use the wave functions to calculate transition probabilities for the electron to change from one state to another.

**Allowed transitions:** Electrons absorbing or emitting photons can change states when  $\Delta\ell = \pm 1$ . (Evidence for the photon carrying one unit of angular momentum!)

$$\Delta n = \text{anything}$$

$$\Delta\ell = \pm 1$$

$$\Delta m_\ell = 0, \pm 1$$

**Forbidden transitions:** Other transitions possible but occur with much smaller probabilities when  $\Delta\ell \neq \pm 1$ .



# Probability Distribution Functions

- We must use wave functions to calculate the probability distributions of the electrons.
- The “position” of the electron is spread over space and is not well defined.
- We may use the radial wave function  $R(r)$  to calculate radial probability distributions of the electron.
- The probability of finding the electron in a differential volume element  $d\tau$  is

$$dP = \psi^*(r, \theta, \phi) \psi(r, \theta, \phi) d\tau$$



# Probability Distribution Functions

- The differential volume element in spherical polar coordinates is

$$d\tau = r^2 \sin\theta dr d\theta d\phi$$

Therefore,

$$P(r)dr = r^2 R^*(r)R(r)dr \int_0^\pi |f(\theta)|^2 \sin\theta d\theta \int_0^{2\pi} g(\phi)d\phi$$

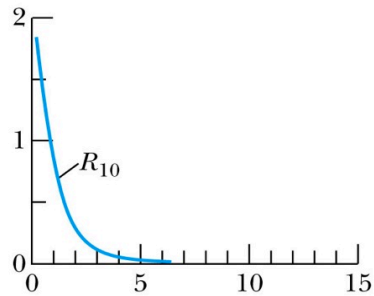
- We are only interested in the radial dependence.

$$P(r)dr = r^2 |R(r)|^2 dr$$

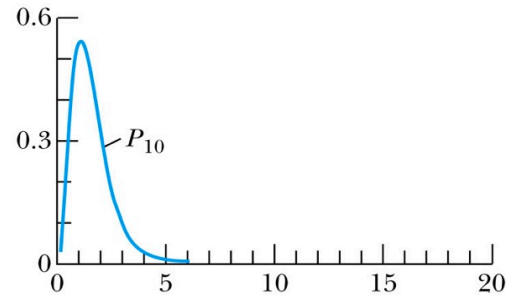
- The radial probability density is  $P(r) = r^2 |R(r)|^2$  and it depends only on  $n$  and  $l$ .

# Probability Distribution Functions

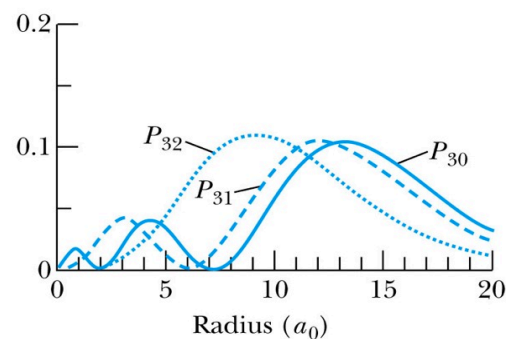
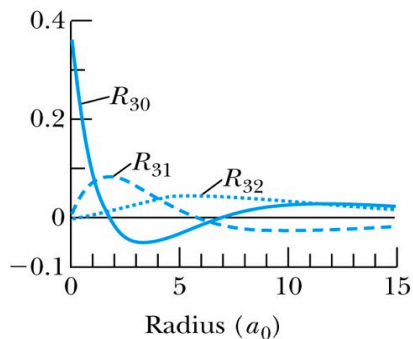
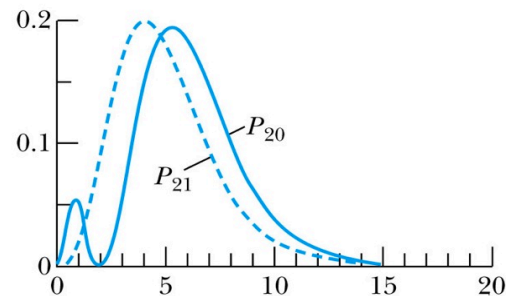
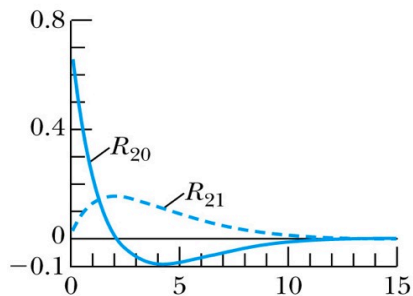
Radial wave functions ( $R_{n\ell}$ )



Radial probability distribution ( $P_{n\ell}$ )



- $R(r)$  and  $P(r)$  for the lowest-lying states of the hydrogen atom



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# Probability Distribution Functions

- The probability density for the hydrogen atom for three different electron states

