

## 4E : A Course on the Quantum Universe

- Quantum physics is the most exciting advance in the history of science. Its firestorm like birth and development makes it an excellent example of the symbiosis between theory and experimentation
- It is the fountainhead of Modern Chemistry, Biology and many fields of Engineering
- What to expect in this course:
- You will see Quantum mechanics a few times as UCSD UG
- For Example, $130 \mathrm{~A}, \mathrm{~B}, \mathrm{C}$ series will be a formal and mathematical account of the methods of quantum Mechanics
- This course will be a more conceptual and "intuitive" introduction to quantum physics
- The last part of this course will be a survey of some interesting examples of the Quantum Universe:
- Particle Physics
- Astrophysics and Cosmology


## Some Bookkeeping Issues Related to This Course <br> -

- Course text: Modern Physics by Tipler, Llewellyn
- $4^{\text {th }}$ edition, Published by WH Freeman
- Instructor :
- Vivek Sharma: modphys@ucsd.edu
- 3314 Mayer Hall, Ph: (858) 5341943
- Office Hours:
- Mon: 1:30-2:30pm, Tue: 2:30-3:30pm
- Teaching Assistant:
- Jason Wright: jwright@physics.ucsd.edu
- 5116 Mayer Hall

- Office Hour: Thursday 3:00-4:00pm in Mayer 2101 (Tutorial Center)
- Class Web Site: http://modphys.ucsd.edu/4es05
- Web page is important tool for this class, make sure you can access it


## 4E Website: http://modphys.ucsd.edu/4es05/

## UCSD Physics Department

Welcome to the Spring '05 Modern Physics (PHYS 4E) Web Site

| Announcements | $\underline{\text { Course Handouts }}$ |
| :---: | :---: |
| Office Hours $\underline{\text { Weekly Schedule }}$ <br> Homework Solutions $\underline{\text { Lecture Slides }}$ <br> Quiz \& Finals Solutions [Absolute] Grading Scale <br> Quiz \& Final Grades $\underline{\text { Scientific American }}$ <br> Material on E-Reserve (Free From UCSD location or Proxy) |  |

E-Mail: Prof. Sharma Jason Wright
Free Stuff You Need: Acrobat Reader
Pl. try to access this website and let me know if you have problems viewing any content.

## Weekly Schedule

Pl. attend discussion session on Wednesday and problem session on Thursday if you plan to do well in this course

General Class Schedule

| Lecture | Monday | 11:00-11:50 am | CNTR 217A | Prof. Sharma |
| :---: | :---: | :---: | :---: | :---: |
| Prof. Office Hour | Monday | 1:30-2:30 pm | Mayer 3314 | Prof. Sharma |
| Lecture | Tuesday | 9:00-9:50 am | SOLIS 109 | Prof. Sharma |
| Prof. Office Hour | Tuesday | 2:30-3:30 pm | Mayer 3314 | Prof. Sharma |
| Lecture | Wednesday | 11:00-11:50 am | CNTR 217A | Prof. Sharma |
| Discussion | Wednesday | 12:00-12:50 pm | SOLIS 109 | Prof. Sharma/Jason Wright |
| TA Office Hour | Thursday | 3:00-4:00 pm | Mayer 2101 | Jason Wright |
| Problem Solving | Thursday | 7:00-8:50 pm | SOLIS 111 | Jason Wright |
| Lecture | Friday | 11:00-11:50 am | CNTR 217A | Announce Quiz/Midterm Dates |
| Prof. Office Hour | Weekend | By Appointment | Mayer 3314 | Prof. Sharma |

Check the announcements page for important schedule changes

## Week 1 Schedule \& HW

| Date | Time | Read | Topic | HV problems for the week | Location |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Monday | $\begin{aligned} & \text { 11:00 } \\ & \text { am } \end{aligned}$ | $\begin{gathered} \text { Ch. } \\ 3 \end{gathered}$ | ```Quantization of Charge, Light & Energy``` | Read Section 3.1 | $\begin{aligned} & \text { CNTR } \\ & 217 \mathrm{~A} \end{aligned}$ |
| Tuesday | $\begin{aligned} & \text { 9:00 } \\ & \text { am } \end{aligned}$ | $\begin{gathered} \text { Ch. } \\ 3 \end{gathered}$ | Quantization of Charge, Light \& Energy | Ch 3: $2,6,11,15,21,22$ | $\begin{aligned} & \text { SOLIS } \\ & 109 \end{aligned}$ |
| Wednesday | $\begin{gathered} \text { 11:00 } \\ \text { am } \end{gathered}$ | $\begin{gathered} \text { Ch. } \\ 3 \end{gathered}$ | Quantization of Charge, Light \& Energy | $\begin{gathered} \text { Ch 3: } \\ 25,32,34,38,41 \end{gathered}$ | $\begin{aligned} & \text { CNTR } \\ & 217 A \end{aligned}$ |
| Wednesday | $\begin{aligned} & \text { 12:00 } \\ & \mathrm{pm} \end{aligned}$ | ----- | Discussion | - | $\begin{gathered} \text { SOLIS } \\ 109 \end{gathered}$ |
| Thursday | $\begin{gathered} 7: 00- \\ 8: 50 \mathrm{pm} \end{gathered}$ | - | Problem Session | Attempt problems before PS | $\begin{gathered} \text { SOLIS } \\ 111 \end{gathered}$ |
| Friday | 11:00am | Ch.. 3 | ```Quantization of Charge, Light & Energy``` | Ch. 3: 42,46 | $\begin{aligned} & \text { CNTR } \\ & 217 \mathrm{~A} \end{aligned}$ |

Check the announcements page for important schedule changes

## Quizzes, Final and Grades

- Course score = 60\% Quiz + 40\% Final Exam
- 5 quizzes if I can schedule them, best $4(=\mathrm{n}-1)$ scores used
- Two problems in each quiz, 45 minutes to do it
- One problem HW like, other more interesting
- Closed book exam, but you can bring one page "CHEAT SHEET"
- Blue Book required, Code numbers will be given at the 1st quiz. Bring calculator, check battery !
- No makeup quizzes
- See handout for Quiz regrade protocol
- Final Exam : TBA, but in Week of June 6-10
- Inform me of possible conflict within 2 weeks of course
- Don't plan travel/vacation before finals schedule is confirmed !
- No makeup finals for any reason


## All Quizzes During My Research Related Travel

Tentative Schedule, TBC next week

- Quiz 1 on Monday April 11
- Quiz 2 on Friday April 29
- Quiz 3 Friday May 13
- Quiz 4 Friday 20 or $27^{\text {th }}$ (TBC)
- Quiz 5 Friday June 3


## Course Grade

- Our wish is that every body gets an A!...So no curve
- Grading is on an absolute scale. Roughly it looks like this :

| Total Score | Grade |
| :---: | :---: |
| $>85$ | $\mathrm{~A}+$ |
| $>75$ | A |
| $>60$ | B |
| $>45$ | C |
| $<30$ | F |

## How To Do Well In This Course

- Read the assigned text BEFORE lecture to get a feel of the topic
- Don't rely on your intuition ! The concepts are quite abstract.
- Attend lecture (ask questions during/before/after lecture) and discussion.
- Do not just accept a concept without understanding the logic
- Attempt all homework problems yourself
- Before looking at the problem solutions (available on web by Tuesday afternoon) \& before attending Problem Solving session
- The textbook, the lectures and the discussions are all integral to this course. Just following lectures is not sufficient (I won't cover every thing)
- Quarter goes fast, don't leave every thing for the week before exam !!
- Don't hesitate to show up at Prof. or TA office hour (they don't bite !)


Constituents of Nature: The Ancient View
Every civilization has speculated about the constitution of the Universe. The Greek philosophers thought that the universe was made up of just four elements: Earth, air, Fire and Water


This was a great "scientific" theory because it was simple but it had one drawback: It was wrong! There was no experimental proof for it.

## Concept of An Atom

- Around $6^{\text {th }}-5^{\text {th }}$ century BC, Indians and more famously the Greeks speculated on "indivisible" constituents of matter
- In $5^{\text {th }} \mathrm{BC}$, Leucippus and his follower Democritus set the scene for modern physics by asking " what would happen if you chopped up matter into ever smaller pieces. There would be a limit beyond which you could chop no more!"
- They called this indivisible piece an Atom (or Anu in Sanskrit)



## Some Highlights in Understanding Matter

- Lavosier's measurement of conservation of matter in chemical reactions
- Faraday's Electrolysis experiment (1833) : Same amount of charge F is required to decompose 1 gram-ionic weight of monovalent ions
- 1 F passed thru NaCl leads to 23 gm of Na at cathode and 35.5 gm Cl at anode but it takes 2F to disassociate $\mathrm{CuSO}_{4}$
$-\Rightarrow$ Mass of element liberated at an electrode is directly proportional to charge transferred and inversely prop. to the valence of the freed element
- Avagadro postulated that pure gases at same temprature and pressure have same number of molecules per unit volume.
$-\Rightarrow \mathrm{N}_{\mathrm{A}}=6.023 \times 10^{23}$
- Dalton \& Mendeleev's theory that all elementary atoms differing in mass and chemical properties
- Discovery of cathode rays and measurement of their properties


## Quantum Nature of Matter

- Fundamental Characteristics of different forms of matter
- Mass
- Charge
- Experimentally measurable
- using some combination of E \& B

$$
\vec{F}=q(\vec{E}+\vec{v} \times \vec{B})
$$

-Or E/B and some other macroscopic force
e.g. Drag Force

## Thomson's Determination of e/m of Electron



- In E Field alone, electron lands at D
- In B field alone, electron lands at E
- When E and B field adjusted to cancel each other's force $\rightarrow$ electron lands at F $\rightarrow \mathrm{e} / \mathrm{m}=1.7588 \times 10^{11} \mathrm{C} / \mathrm{Kg}$



## Necessary Homework Reading

- Pl. read Section 3.1, including the discussion detailing the Millikan's oil drop experiment (download from www.freeman.com/modphys4e)
- This is straightforward reading. HW problems are assigned on this and the material may show up in the quiz


## Ch 3 : Quantum Theory Of Light

- What is the nature of light ?
- When it propagates?
- When it interacts with Matter?
- What is Nature of Matter ?
- When it interacts with light?
- As it propagates ?
- Revolution in Scientific Thought
- A firestorm of new ideas (NOT steady dragged out progress)
- Old concepts violently demolished, new ideas born
- Rich interplay of experimental findings \& scientific reason
- One such revolution happened at the turn of $20^{\text {th }}$ Century
- Led to the birth of Quantum Theory \& Modern Physics


Hertz \& Experimental Demonstration of Light as EM Wave


## Properties of EM Waves: Maxwell's Equations



$$
\text { Intensity of Radiation } \mathrm{I}=\frac{1}{2 \mu_{0} c} E_{0}^{2}
$$

Larger the amplitude of Oscillation
More intense is the radiation

## Disasters in Classical Physics (1899-1922)

Disaster $\rightarrow$ Experimental observation that could not be explained by Classical theory

- Disaster \# 1 : Nature of Blackbody Radiation from your BBQ grill
- Disaster \# 2: Photo Electric Effect
- Disaster \# 3: Scattering light off electrons (Compton Effect)

Resolution of Experimental Observation will require radical changes in how we think about nature
$-\rightarrow$ QUANTUM PHYSICS: The Art of Conversation with Subatomic Particles

Nature of Radiation: An Expt with BBQ Grill


(a) Intensity of Radiation $\mathrm{I}=\int R(\lambda) d \lambda \propto T^{4}$
$I=\sigma T^{4}$ (Area under curve)
Stephan-Boltzmann Constant $\sigma=5.6710^{-8} \mathrm{~W} / \mathrm{m}^{2} \mathrm{~K}^{4}$

(b) Higher the temperature of BBQ, Lower is the $\lambda$ of PEAK intensity

$$
\mathrm{I}_{\operatorname{MAX}} \propto 1 / \mathrm{T}
$$

$\lambda_{\text {MAX }} \mathrm{T}=$ const

$$
=2.89810^{-3} \mathrm{mK}
$$

As a body gets hotter it gets more RED then White : Wein's Law

Reason for different shape of $R(\lambda) \vee s \lambda$ for different temperature? Can one explain in on basis of Classical Physics ??

## Blackbody Radiator: An Idealization



Classical Thought:

- Box is filled with EM standing waves
- Radiation reflected back-and-forth between walls
- Radiation in thermal equilibrium with walls of Box
- How may waves of wavelength $\lambda$ can fit inside the box?

Blackbody Absorbs everything
Reflects nothing
All light entering opening gets absorbed (ultimately) by the cavity wall

Cavity in equilibrium T
w.r.t. surrounding. So it
radiates everything It absorbs
Emerging radiation is a sample
of radiation inside box at temp T
Predict nature of radiation inside Box ?


Standing Waves

(c)


## The Beginning of The End ! How BBQ Broke Physics

## Classical Calculation

\# of standing waves between Wavelengths $\lambda$ and $\lambda+\mathrm{d} \lambda$ are

$$
\mathrm{N}(\lambda) \mathrm{d} \lambda=\frac{8 \pi \mathrm{~V}}{\lambda^{4}} \bullet d \lambda ; \mathrm{V}=\text { Volume of box }=\mathrm{L}^{3}
$$

Each standing wave contributes energy $\mathrm{E}=\mathrm{kT}$ to radiation in Box
Energy density $u(\lambda)=[\#$ of standing waves $/$ volume $] \times$ Energy $/$ Standing Wave

$$
=\frac{8 \pi \mathrm{~V}}{\lambda^{4}} \times \frac{1}{\mathrm{~V}} \times \mathrm{kT}=\frac{8 \pi}{\lambda^{4}} \mathrm{kT}
$$

$\operatorname{Radiancy} \mathrm{R}(\lambda)=\frac{\mathrm{c}}{4} \mathrm{u}(\lambda)=\frac{\mathrm{c}}{4} \frac{8 \pi}{\lambda^{4}} \mathrm{kT}=\frac{2 \pi \mathrm{c}}{\lambda^{4}} \mathrm{kT}$
Radiancy is Radiation intensity per unit $\lambda$ interval: Lets plot it
Prediction : as $\lambda \rightarrow 0$ (high frequency f$), \mathbf{R}(\lambda) \rightarrow$ Infinity ! Oops !

## Ultra Violet (Frequency) Catastrophe



## That was a Disaster !

(\#1)

## Disaster \# 2 : Photo-Electric Effect

Light of intensity I, wavelength $\lambda$ and frequency fincident on a photo-cathode


Measure characteristics of current in the circuit as a fn of $I, f, \lambda$

## Photo Electric Effect: Measurable Properties

- Rate of electron emission from cathode
- From current $i$ seen in ammeter in the circuit. More photoelectrons $\rightarrow$ more current registered in ammeter
- Maximum kinetic energy of emitted electron
- By applying retarding potential on electron moving left to tright towards Collector plate
- $\mathrm{K}_{\mathrm{MAX}}=\mathrm{eV}_{0}$ ( $\mathrm{V}_{0}=$ Stopping voltage)
- Stopping potential $\rightarrow$ no current flows
- Photoelectric Effect on different types of photo-cathode metal surface
- Time between shining light and first sign of photo-current in the circuit

Observations:PhotoCurrent Vs Intensity of Incident Light


Observations: Photocurrent Vs frequency of incident light


Stopping Voltage ( $V_{0}$ ) Vs Incident Light Frequency ( $f$ )
Try different photocathode materials.....see what happens


## Conclusions from the Experimental Observations

- Max Kinetic energy $\mathrm{K}_{\mathrm{MAX}}$ independent of Intensity I for light of same frequency
- No photoelectric effect occurs if light frequency f is below a threshold no matter how high the intensity of light
- For a particular metal, light with $\mathrm{f}>\mathrm{f}_{\mathrm{t}}$ causes photoelectric effect IRRESPECTIVE of light intensity.
- $f_{t}$ is characteristic of that metal
- Photoelectric effect is instantaneous !...not time delay
Can one Explain all this Classically !


## Classical Explanation of Photo Electric Effect

- As light Intensity increased $\Rightarrow \vec{E}$ field amplitude larger
- E field and electrical force seen by the "charged subatomic oscillators" Larger
- $\vec{F}=e \vec{E}$
- More force acting on the subatomic charged oscillator
- $\Rightarrow$ More (work done) $\rightarrow$ more energy transferred to it
- $\Rightarrow$ Charged particle "hooked to the atom" should leave the surface with more Kinetic Energy KE !! The intensity of light (EM Wave) shining rules !
- As long as light is intense enough, light of ANY frequency f should cause photoelectric effect
- Because the Energy in a Wave is uniformly distributed over the Spherical wavefront incident on cathode, should be a noticeable time lag $\Delta \mathrm{T}$ between time is incident \& the time a photo-electron is ejected : Energy absorption time
- How much time for electron ejection? Lets calculate it classically


## Classical Physics: Time Lag in Photo-Electric Effect ?

- Electron absorbs energy incident on a surface area where the electron is confined $\cong$ size of atom in cathode metal
- Electron is "bound" by attractive Coulomb force in the atom, so it must absorb a minimum amount of radiation before its stripped off
- Example : Laser light Intensity I = $120 \mathrm{~W} / \mathrm{m}^{2}$ on Na metal
- Binding energy $=2.3 \mathrm{eV}=$ "Work Function $\Phi$ "
- Electron confined in Na atom, size $\cong 0.1 \mathrm{~nm}$; how long before ejection?
- Average Power Delivered $\mathrm{P}_{\mathrm{AV}}=\mathbf{I} . \mathbf{A}, \mathrm{A}=\pi \mathrm{r}^{2} \cong 3.1 \times 10^{-20} \mathrm{~m}^{2}$
- If all energy absorbed then $\Delta \mathrm{E}=\mathrm{P}_{\mathrm{AV}} \cdot \Delta \mathrm{T} \Rightarrow \Delta \mathrm{T}=\Delta \mathrm{E} / \mathrm{P}_{\mathrm{AV}}$

$$
\Delta T=\frac{(2.3 \mathrm{eV})\left(1.6 \times 10^{-19} \mathrm{~J} / \mathrm{eV}\right)}{\left(120 \mathrm{~W} / \mathrm{m}^{2}\right)\left(3.1 \times 10^{-20} \mathrm{~m}^{2}\right)}=0.10 \mathrm{~S}
$$

- Classical Physics predicts measurable delay even by the primitive clocks of 1900
- But in experiment, the effect was observed to be instantaneous !!
- Classical Physics fails in explaining all results


## That was a Disaster !

> (\# 2)

Beginning of a search for a new hero or an explanation or both !

## Max Planck \& Birth of Quantum Physics

## Back to Blackbody Radiation Discrepancy

Planck noted the Ultraviolet catastrophe at high frequency
"Cooked" calculation with new "ideas" so as bring:

$$
\begin{aligned}
\mathrm{R}(\lambda) \rightarrow 0 \text { as } \lambda & \rightarrow 0 \\
\mathrm{f} & \rightarrow \infty
\end{aligned}
$$

- Cavity radiation as equilibrium exchange of energy between EM radiation \& "atomic" oscillators present on walls of cavity
- Oscillators can have any frequency f
- But the Energy exchange between radiation and oscillator NOT continuous, it is discrete ...in packets of same amount

$$
\mathrm{E}=\mathrm{n} \text { hf, with } \mathrm{n}=1,2,3,4, \ldots \infty
$$

$\mathrm{h}=$ constant he invented, a number he made up !

Planck did not know about electrons, Nucleus etc: They had not been discovered then


## Planck, Quantization of Energy \& BlackBody Radiation

- Keep the rule of counting how many waves fit in a BB Volume
- Radiation Energy in cavity is quantized
- EM standing waves of frequency f have energy

$$
\mathrm{E}=\mathrm{n} \text { hf }(\mathrm{n}=1,2,3 \ldots 10 \ldots .1000 \ldots)
$$

$\qquad$

- Probability Distribution: At an equilibrium temp T, possible energy of oscillators is distributed over a spectrum of states: $\mathrm{P}(\mathrm{E})=\mathrm{e}^{(-\mathrm{E} / \mathrm{kT})}$
- Modes of Oscillation with :
-Less energy $\mathrm{E}=\mathrm{hf}$ = favored
- More energy $\mathrm{E}=\mathrm{hf}=$ disfavored

P(E)


By this discrete statistics, large energy = high f modes of EM disfavored

Planck's Calculation: A preview to keep the story going

$$
R(\lambda)=\left(\frac{c}{4}\right)\left(\frac{8 \pi}{\lambda^{4}}\right)\left[\frac{h c}{\lambda}\left(\frac{1}{e^{\frac{h c}{\lambda k T}}-1}\right)\right]
$$

Odd looking form
When $\lambda \rightarrow$ large $\Rightarrow \frac{\mathrm{hc}}{\lambda \mathrm{kT}} \rightarrow$ small
Recall $\mathrm{e}^{\mathrm{x}}=1+x+\frac{x^{2}}{2!}+\frac{x^{3}}{3!}+\ldots$.

$$
\Rightarrow e^{\frac{h c}{\lambda k T}}-1=\left(1+\frac{h c}{\lambda k T}+\frac{1}{2}\left(\frac{h c}{\lambda k T}\right)^{2}+\ldots . .\right]-1
$$

$=\frac{h c}{\lambda k T} \quad$ plugging this in $\mathrm{R}(\lambda)$ eq:
$R(\lambda)=\left(\frac{c}{4}\right)\left(\frac{8 \pi}{\lambda^{4}}\right) \frac{h c}{\lambda k T}$
Graph \& Compare With BBQ data

## Planck's Formula and Small $\lambda$

When $\lambda$ is small (large f)

$$
\frac{1}{e^{\frac{h c}{\lambda k T}}-1} \cong \frac{1}{e^{\frac{h c}{\lambda k T}}}=e^{-\frac{h c}{\lambda k T}}
$$

Substituting in $R(\lambda)$ eqn:

$$
\begin{aligned}
& R(\lambda)=\left(\frac{c}{4}\right)\left(\frac{8 \pi}{\lambda^{4}}\right) e^{-\frac{h c}{\lambda k T}} \\
& \text { As } \lambda \rightarrow 0, e^{-\frac{h c}{\lambda k T}} \rightarrow 0
\end{aligned}
$$

$$
\Rightarrow R(\lambda) \rightarrow 0
$$

Just as seen in the experimental data !

## Planck's Explanation of Black Body Radiation



Major Consequence of Planck's Energy Postulate
nipatirationof Enarg!


Judging Planck's Postulate : Visionary or just a Wonk?

"IT'S AN EXCELLENT PROOF, BUT IT LACKS WARMTH AND FEELING,"

## Einstein's Explanation of Photoelectric Effect



## Einstein's Explanation of Photoelectric Effect

- Energy associated with EM waves not uniformly distributed over wavefront, rather is contained in packets of energy $\Rightarrow$ PHOTON
- $\mathrm{E}=\mathrm{hf}=\mathrm{hc} / \lambda$ [but is it the same h as in Planck's th.?]
- Light shining on metal emitter/cathode is a stream of photons of energy E which depends on frequency $f$
- Photons knock off electron from metal instantaneously
- Transfer all energy to electron
- Energy gets used up to pay for Work Function Ф. Remaining energy shows up as KE of electron $\mathrm{KE}=\mathrm{hf}-\Phi$
- Cutoff Frequency hf $=\Phi$ (pops an electron, $\mathrm{KE}=0$ )
- Larger intensity I $\rightarrow$ more photons incident
- Low frequency light $f \rightarrow$ not energetic enough to overcome work function of electron in atom

Einstein's Interpretation of Photoelectric Effect (1905)

$$
\begin{aligned}
& E=h f=\varphi+K E_{\text {electron }} \\
& e V_{0}=K E=h f-\varphi
\end{aligned}
$$

Now interpret the experimental data Under the "single bullet" theory

## Makes Sense!

(b)


Modern View of Photoelectric Effect



Reinterpreting Photoelectric Effect With Light as Photons
Photoelectric Effect on An Iron Surface
Light of Intensity $\mathrm{I}=1.0 \mu \mathrm{~W} / \mathrm{cm}^{2}$ incident on $1.0 \mathrm{~cm}^{2}$ surface of Fe
Assume Fe reflects $96 \%$ of light
further only $3 \%$ of incident light is Violet region ( $\lambda=250 \mathrm{~nm}$ )
barely above threshold frequency for Photoelectric effect
(a) Intensity available for Ph. El effect $\mathrm{I}=3 \% \times 4 \% \times\left(1.0 \mu \mathrm{~W} / \mathrm{cm}^{2}\right)$
(b) how many photo-electrons emitted per second?

$$
\# \text { of photoelectrons }=\frac{\text { Power }}{\mathrm{hf}}=\frac{3 \% \times 4 \% \times\left(1.0 \mu \mathrm{~W} / \mathrm{cm}^{2}\right) \lambda}{\mathrm{hc}}
$$

$$
=\frac{\left(250 \times 10^{-9} \mathrm{~m}\right)\left(1.2 \times 10^{-9} \mathrm{~J} / \mathrm{s}\right)}{\left(6.6 \times 10^{-34} \mathrm{~J} \cdot \mathrm{~s}\right)\left(3.0 \times 10^{8} \mathrm{~m} / \mathrm{s}\right)}
$$

$$
1.5 \times 10^{9}
$$

(c) Current in Ammeter : $\mathrm{i}=\left(1.6 \times 10^{-19} \mathrm{C}\right)\left(1.5 \times 10^{9}\right)=2.4 \times 10^{-10} \mathrm{~A}$
(d) W ork Function $\Phi=\mathrm{hf}_{0}=\left(4.14 \times 10^{-15} \mathrm{eV} \cdot \mathrm{s}\right)\left(1.1 \times 10^{15} \mathrm{~s}^{-1}\right)$

$$
=4.5 \mathrm{eV}
$$

## Facts about Light Quantum

- The human eye is a sensitive photon detector at visible wavelengths: Need $>5$ photons of $\cong 550 \mathrm{~nm}$ to register on your optical sensor
- The Photographic process :
- Energy to Dissociate an AgBr molecule $=0.6 \mathrm{eV}$
- Photosynthesis Process : 9 sunlight photon per reaction cycle of converting $\mathrm{CO}_{2}$ and water to carbohydrate \& $\mathrm{O}_{2}$ - chlorophyll absorbs best at $\lambda \cong 650-700 \mathrm{~nm}$
- Designing Space Shuttle "skin": Why Platinum is a good thing
- designing Solar cells : picking your metal cathode


## Photon \& Relativity: Wave or a Particle ?

- Photon associated with EM waves, travel with speed $=\mathrm{c}$
- For light $(\mathrm{m}=0)$ : Relativity says $\mathrm{E}^{2}=(\mathrm{pc})^{2}+\left(\mathrm{mc}^{2}\right)^{2}$
- $\Rightarrow \mathrm{E}=\mathrm{pc}$
- But Planck tells us : $\mathrm{E}=\mathrm{hf}=\mathrm{h}(\mathrm{c} / \lambda)$
- Put them together : hc $/ \lambda=\mathrm{pc}$
$-\Rightarrow \mathrm{p}=\mathrm{h} / \lambda$
- Momentum of the photon (light) is inversely proportional to $\lambda$
- But we associate $\lambda$ with waves $\& \mathrm{p}$ with particles .... what is going on??
-Quantum Physics !


## X Rays "Bremsstrahlung": The Braking Radiation

- EM radiation, produced by bombarding a metal target with energetic electrons.
- Produced in general by ALL decelerating charged particles
- X rays : very short $\lambda \cong 60-100 \mathrm{pm}\left(10^{-12} \mathrm{~m}\right)$, large frequency f
- Very penetrating because very energetic $E=h f!!$


Useful for probing structure of sub-atomic Particles (and your teeth !)

X Ray Production Mechanism


When electron passes near a positively charged target nucleus contained in target material, its deflected from its path because of Coulomb attraction, experiences acceleration.

E\&M $\rightarrow$ that any charged particle will emit radiation when accelerated. This EM radiation "appears" as photons. Since photo carries energy and momentum, the electron must lose same amount. If all of electron's energy is lost in just one single collision then:
$e \Delta V=\mathrm{hf}_{\max }=\frac{h c}{\lambda_{\text {min }}}$ or $\lambda_{\text {min }}=\frac{h c}{e \Delta V}$

## X Ray Spectrum in Molybdenum (Mo)



## X rays As Subatomic Probes

X rays are EM waves of low wavelength, high frequency (and energy) and demonstrate characteristic features of a wave

- Interference \&Diffraction
- To probe into a structure size $\Delta \mathrm{X}$ you need a light source with wavelength much smaller than the features of the object being probed
- Good Resolution $\rightarrow \lambda_{\text {SOURCE }} \ll \Delta X$
- X rays allows one probe at atomic size $\left(10^{-10}\right) \mathrm{m}$



## Compton Scattering : Quantum Pool!

- Arthur Compton (USA) proves that X-rays (EM Waves) have particle like properties (acts like photons)
- Showed that classical theory failed to explain the scattering effect of X rays on to free (not bound, barely bound electrons)
- Experiment : shine X ray on to a surface with "almost" free electrons
- Watch the scattering of light off electron : measure time $+\lambda$ of scattered X-ray



## Compton Effect: what should Happen Classically?

- Plane wave [ $\mathrm{f}, \lambda$ ] incident on a surface with loosely bound electrons $\rightarrow$ interaction of E field of EM wave with electron: $\mathbf{F}=\mathrm{eE}$
- Electron oscillates with
$\mathrm{f}=\mathrm{f}_{\text {incident }}$
- Eventually radiates spherical waves with $\mathrm{f}_{\text {radiated }}=\mathrm{f}_{\text {incident }}$
- At all scattering angles, $\Delta \mathrm{f}$ \&
$\Delta \lambda$ must be zero
- Time delay while the
 electron gets a "tan": soaks in radiation




## Compton Scattering : Summary of Observations



How does one explain this startling anisotropy?
67

Compton Effect : Quantum (Relativistic) Pool


Compton Scattering: The Quantum Picture


Compton Scattering: The Quantum Picture


Rules of Quantum Pool between Photon and Electron


$$
\left(\lambda^{\prime}-\lambda\right)=\left(\frac{\boxed{h}}{m_{e} c}\right)(1-\cos \theta)
$$

## Checking for " $h$ " in Compton Scattering

From scattered photon $\lambda$, plot $\Delta \lambda$, calculate slope and measure "h"


Other Forms of Energy Exchange between Radiation and Matter
$\mathrm{E} \rightleftharpoons \mathrm{mc}^{2}+\mathrm{mc}^{2}$ same kind of matter \& antimatter produced or destroyed in pairs

Particle creation

gamma-ray photon

Particle annihilation
antielectron gamma-ray photon


Constructive Interference depends on Path (or phase) diff. Traversed
Two Identical waves $y_{i}(x, t)=y_{\max } \sin \left(k_{i} x-\omega_{i} t+\phi_{i}\right)$ travel along +x and interefere to give a resulting wave $\mathrm{y}^{\prime}(x, t)$. The resulting wave form depends on relative phase difference between 2 waves. Shown for $\Delta \phi=0, \pi, \frac{2}{3} \pi$

(a)
$\downarrow$


(b)
$\downarrow$


(c)
$\downarrow$



Bragg Scattering: Probing Atoms With X-Rays Incident X-ray detector


Constructive Interference when net phase difference is $0,2 \pi$ etc This implied path difference traveled by two waves must be integral multiple of wavelength : $\mathbf{n} \boldsymbol{\lambda}=\mathbf{2 d} \sin \vartheta$

From X Ray (EM Wave) Scattering data,
size of atoms was known to be about $10^{-10} \mathrm{~m}$


## Where are the electrons inside the atom?

Early Thought: "Plum pudding" model $\rightarrow$ Atom has a homogenous distribution of Positive charge with electrons embedded in them


- How to test these hypotheses? $\rightarrow$ Shoot "bullets" at the atom and watch their trajectory. What Kind of bullets ?
$\cdot$ Indestructible charged bullets $\rightarrow$ Ionized $\mathrm{He}^{++}$atom $=\alpha^{++}$particles $\cdot \mathrm{Q}=+2 \mathrm{e}$, Mass $\mathrm{M}_{\alpha}=4 \mathrm{amu} \gg \mathrm{m}_{\mathrm{e}}, \mathrm{V}_{\alpha}=2 \times 10^{7} \mathrm{~m} / \mathrm{s}$ (non-relavistic) [charged to probe charge \& mass distribution inside atom]


## Plum Pudding Model of Atom

- Non-relativistic mechanics $\left(\mathrm{V}_{\alpha} / \mathrm{c}=0.1\right)$
- In Plum-pudding model, $\alpha$-rays hardly scatter because
- Positive charge distributed over size of atom $\left(10^{-10} \mathrm{~m}\right)$
$-\mathrm{M}_{\alpha} \gg \mathrm{M}_{\mathrm{e}}$ (like moving truck hits a bicycle)
$-\rightarrow$ predict $\alpha$-rays will pass thru array of atoms with little scatter


Need to test this hypothesis $\rightarrow$ Ernest Rutherford

## "Rutherford Scatterina" discovered by his PhD Student (Marsden)



Force on $\alpha$-particle due to heavy Nucleus

$\alpha$ particle trajectory is hyperbolic
Scattering angle is related to impact par.
-Outside radius $\mathrm{r}=\mathrm{R}, \mathrm{F} \propto \mathrm{Q} / \mathrm{r}^{2}$
$\cdot$ Inside radius $\mathrm{r}<\mathrm{R}, \mathrm{F} \propto \mathrm{q} / \mathrm{r}^{2=} \mathrm{Qr} / \mathrm{R}^{2}$
-Maximum force at radius $\mathrm{r}=\mathrm{R}$
Impact Parameter $b=\left(\frac{k q_{\alpha} Q}{m_{\alpha} v_{\alpha}^{2}}\right)\left(\cot \frac{\theta}{2}\right)$

Rutherford Scattering: Prediction and Experimental Result

\# scattered Vs $\phi$ depends on:
$\mathrm{n}=\#$ of incident alpha particles
$\mathrm{N}=$ \# of nuclei/area of foil
$\mathrm{Ze}=$ Nuclear charge
$\mathrm{K}_{\alpha}$ of incident alpha beam
$\mathrm{A}=$ detector area



## Dimension Matters !

## Size of Nucleus $=10^{-15} \mathrm{~m}$ Size of Atom $=10^{-10} \mathrm{~m}$

How are the electrons located inside an atom? How are they held in a stable fashion ? necessary condition for us to exist !

All these discoveries will require new experiments and observations

Where are the Electrons in an Atom ?


## Clues: Continuous \& Discrete spectra of Elements

Hot blackbody

a Continuous spectrum

c Emission line spectrum


Emission \& Absorption Line Spectra of Elements



## Kirchhoff’ Experiment : "D" Lines in Na



D lines darken noticeably when Sodium vapor introduced Between slit and prism

Emission \& Absorption Line Spectrum of Elements

Emission line appear dark because of photographic exposure


Spectral Observations : series of lines with a pattern


- Empirical observation (by trial \& error)
- All these series can be summarized in a simple formula

$$
\frac{1}{\lambda}=R\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right), n_{f}>n_{i}, n_{i}=1,2,3,4 . .
$$

Fitting to spectral line series data

$$
\mathrm{R}=1.09737 \times 10^{7} \mathrm{~m}^{-1}
$$

How does one explain this ?

The Rapidly Vanishing Atom: A Classical Disaster ! Not too hard to draw analogy with dynamics under another Central Force

Think of the Gravitational Force between two objects and their circular orbits.
Perhaps the electron rotates around the Nucleus and is bound by their electrical charge

$$
\mathrm{F}=\mathrm{G} \frac{\mathrm{M}_{1} \mathrm{M}_{2}}{\mathrm{r}^{2}} \Rightarrow \mathrm{k} \frac{Q_{1} Q_{2}}{\mathrm{r}^{2}}
$$

Laws of E\&M destroy this equivalent picture : Why?


## Classical Trajectory of The Orbiting Electron

Classical model of Hydrogenic Atom (Z protons) is mechanically stable but is electrically unstable !
Mechanically balanced : $\mathrm{F}=\frac{\mathrm{kZe}^{2}}{r^{2}}=\frac{m v^{2}}{r} \quad$ (Coulomb force $=$ Centripetal force)
But electron is always accelerating towards center of circle. Laws of classical electrodynamics predict that accelerating charge will radiate light of frequency $f=$ freq. of periodic motion
$f=\frac{v}{2 \pi r}=\left(\frac{k Z e^{2}}{r m}\right)^{\frac{1}{2}} \frac{1}{2 \pi r}=\left(\frac{k Z e^{2}}{4 \pi^{2} m}\right)^{\frac{1}{2}} \frac{1}{r^{\frac{3}{2}}} \sim \frac{1}{r^{\frac{3}{2}}}$
And Total energy $\mathrm{E}=\mathrm{KE}+\mathrm{U}=\frac{\mathrm{mv}^{2}}{2}+\left(-\frac{k Z e^{2}}{r}\right)$, but since $\frac{\mathrm{kZe} \mathrm{e}^{2}}{2 r}=\frac{m \nu^{2}}{2}$
$\Rightarrow E=\frac{k Z e^{2}}{2 r}-\frac{k Z e^{2}}{r}=\frac{k Z e^{2}}{2 r} \sim-\frac{1}{r}$
Thus Classical physics predicts that as energy is lost to radiation, electron's orbit will become smaller and smaller while frequency of radiation will become larger and larger!
The electron will reach the Nucleus in $\sim 1 \mu$ s !!
In reality, this does not occur. Unless excited by external means, atoms do not radiate AT ALL!

Bohr's Bold Model of Atom: Semi Quantum/Classical


1. Electron in circular orbit around proton with vel=v
2. Only stationary orbits allowed Electron does not radiate when in these stable (stationary) orbits
3. Orbits quantized:
$-\mathrm{M}_{\mathrm{e}} \mathrm{v} \mathrm{r}=\mathrm{nh} / 2 \pi(\mathrm{n}=1,2,3 \ldots)$
4. Radiation emitted when electron "jumps" from a stable orbit of higher energy $\rightarrow$ stable orbit of lower energy $\mathrm{E}_{\mathrm{f}}-\mathrm{E}_{\mathrm{i}}=\mathrm{hf}=\mathrm{hc} / \lambda$
5. Energy change quantized

- $f=$ frequency of radiation
- Both Nucleus \& e e revolve around their common center of mass (CM)
- Such a system is equivalent to single particle of "reduced mass" $\mu$ that revolves around position of Nucleus at a distance of $\left(\mathrm{e}^{-}-\mathrm{N}\right)$ separation
$\square \mu=\left(m_{\mathrm{e}} \mathrm{M}\right) /\left(\mathrm{m}_{\mathrm{e}}+\mathrm{M}\right)$, when $\mathrm{M} \gg \mathrm{m}, \mu=\mathrm{m}$ (Hydrogen atom)
$\square$ Not so when calculating Muon $\left(\mathrm{m}_{\mu}=207 \mathrm{~m}_{\mathrm{e}}\right)$ or equal mass charges rotating around each other (similar to what you saw in gravitation)


## Allowed Energy Levels \& Orbit Radii in Bohr Model

$\mathrm{E}=\mathrm{KE}+\mathrm{U}=\frac{1}{2} m_{e} v^{2}-k \frac{e^{2}}{r}$
Force Equality for Stable Orbit
$\Rightarrow$ Coulomb attraction $=$ CP Force

$$
\begin{aligned}
k \frac{e^{2}}{r^{2}}=\frac{m_{e} v^{2}}{r} \\
\Rightarrow K E=\frac{m_{e} v^{2}}{2}=k \frac{e^{2}}{2 r}
\end{aligned}
$$

Total Energy $\mathrm{E}=\mathrm{KE}+\mathrm{U}=-k \frac{e^{2}}{2 r}$
Negative $\mathrm{E} \Rightarrow$ Bound system
This much energy must be added to
the system to break up the bound atom

Radius of Electron Orbit :
$m v r=n \hbar$
$\Rightarrow v=\frac{n \hbar}{m r}$,
substitute in $\mathrm{KE}=\frac{1}{2} m_{e} v^{2}=\frac{k e^{2}}{2 r}$
$\Rightarrow r_{n}=\frac{n^{2} \hbar^{2}}{m k e^{2}}, n=1,2, \ldots \infty$
$n=1 \Rightarrow$ Bohr Radius $a_{0}$
$a_{0}=\frac{1^{2} \hbar^{2}}{m k e^{2}}=0.529 \times 10^{-10} \mathrm{~m}$
In general $r_{n}=n^{2} a_{0} ; n=1,2, \ldots \infty$
Quantized orbits of rotation

## Energy Level Diagram and Atomic Transitions

$$
\begin{aligned}
& E_{n}=K+U=\frac{-k e^{2}}{2 r} \\
& \text { since } r_{n}=a_{0} n^{2}, \mathrm{n}=\text { quantum number } \\
& E_{n}=\frac{-k e^{2}}{2 a_{0} n^{2}}=-\frac{13.6}{n^{2}} e V, n=1,2,3 . . \infty \\
& \text { Interstate transition: } \mathrm{n}_{\mathrm{i}} \rightarrow n_{f} \\
& \Delta E=h f=E_{i}-E_{f} \\
& =\frac{-k e^{2}}{2 a_{0}}\left(\frac{1}{n_{i}^{2}}-\frac{1}{n_{f}^{2}}\right) \\
& f=\frac{k e^{2}}{2 h a_{0}}\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right) \\
& \begin{array}{r}
\frac{1}{\lambda}=\frac{f}{c}=\frac{k e^{2}}{2 h c a_{0}}\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right) \\
=\mathrm{R})\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right)
\end{array}
\end{aligned}
$$



## Hydrogen Spectrum: as explained by Bohr


$E_{n}=-\left(\frac{k e^{2}}{2 a_{0}}\right) \frac{Z^{2}}{n^{2}}$
Bohr's " $R$ " same as Rydberg Constant $\mathbf{R}$ derived emperically from spectral series

A Look Back at the Spectral Lines With Bohr's Optic


$$
E_{n}=-\left(\frac{k e^{2}}{2 a_{0}}\right) \frac{Z^{2}}{n^{2}}
$$

Rydberg Constant


Bohr's Atom: Emission \& Absorption Spectra


## Some Notes About Bohr Like Atoms

- Ground state of Hydrogen atom $(\mathrm{n}=1) \mathrm{E}_{0}=-13.6 \mathrm{eV}$
- Method for calculating energy levels etc applies to all Hydrogenlike atoms $\rightarrow-1 \mathrm{e}$ around +Ze
- Examples : $\mathrm{He}^{+}, \mathrm{Li}^{++}$
- Energy levels would be different if replace electron with Muons
- Reduced Mass
- Necessity of Reduced Mass calculation enhanced for "positronium" like systems
- Bohr's method can be applied in general to all systems under a central force (e.g. gravitational instead of Coulombic)

$$
\begin{array}{|l}
\text { If change } U(r)=k \frac{Q_{1} Q_{2}}{r} \rightarrow G \frac{M_{1} M_{2}}{r} \\
\text { Changes every thing: } \mathrm{E}, \mathrm{r}, \mathrm{f} \text { etc } \\
\text { "Importance of constants in your life" } \\
\hline
\end{array}
$$

## Bohr's Correspondence Principle

- It now appears that there are two different worlds with different laws of physics governing them
- The macroscopic world
- The microscopic world
- How does one transcend from one world to the other?
- Bohr's correspondence Principle
- predictions of quantum theory must correspond to predictions of the classical physics in the regime of sizes where classical physics is known to hold.

```
when n}->\infty\mathrm{ [Quantum Physics] = [Classical Physics]
```


## Correspondence Principle for Bohr Atom

- When $\mathrm{n} \gg 1$, quantization should have little effect, classical and quantum calculations should give same result: Check this $\rightarrow$
Compare frequency of transition between level $n_{i}=n$ and $n_{f}=n-1$
In Bohr Model : $f=\frac{c}{\lambda}=\frac{Z^{2} m k^{2} e^{4}}{4 \pi \hbar^{3}}\left(\frac{1}{(n-1)^{2}}-\frac{1}{n^{2}}\right)$
$=\frac{Z^{2} m k^{2} e^{4}}{4 \pi \hbar^{3}} \frac{2 n-1}{n^{2}(n-1)^{2}} \approx \frac{Z^{2} m k^{2} e^{4}}{4 \pi \hbar^{3}} \frac{1}{n^{3}} \quad$ (since when $\mathrm{n} \gg 1, \mathrm{n}-1 \approx \mathrm{n}$ )

And Classically: $f_{\text {rev }}=\frac{v}{2 \pi r}$; using $v=\frac{n \hbar}{m r}$ and $r=\frac{n^{2} \hbar^{2}}{m k Z e^{2}}$
$\Rightarrow f_{\text {rev }}=\frac{n \hbar / m r}{2 \pi r}=\frac{n \hbar}{2 \pi m r^{2}}=\frac{n \hbar}{2 \pi m\left(n^{2} \hbar^{2} / m k Z e^{2}\right)^{2}}=\frac{m k^{2} Z^{2} e^{4}}{2 \pi \hbar^{3} n^{3}}$
Same!

## Atomic Excitation by Electrons: Franck-Hertz Expt

Other ways of Energy exchange are also quantized ! Example:

- Transfer energy to atom by colliding electrons on it -Elastic and inelastic collisions with a heavy atom (Hg)
- Accelerate electrons, collide with Hg atoms, measure energy transfer in inelastic collision (by applying retarding voltage)
- Count how many electrons get thru and arrive at plate P
(a)


Atomic Excitation by Electrons: Franck-Hertz Expt
Plot \# of electrons/time (current) overcoming the retarding potential (V)


Atoms accept only discrete amount of Energy, no matter the fashion in which energy is transfered

## Bohr's Explanation of Hydrogen like atoms

- Bohr's Semiclassical theory explained some spectroscopic data $\rightarrow$ Nobel Prize : 1922
- The "hotch-potch" of clasical \& quantum attributes left many (Einstein) unconvinced
- "appeared to me to be a miracle - and appears to me to be a miracle today One ought to be ashamed of the successes of the theory"
- Problems with Bohr's theory:
- Failed to predict INTENSITY of spectral lines
- Limited success in predicting spectra of multi-electron atoms (He)
- Failed to provide "time evolution" of system from some initial state
- Overemphasized Particle nature of matter-could not explain the wave-particle duality of light
- No general scheme applicable to non-periodic motion in subatomic systems
- "Condemned" as a one trick pony! Without fundamental insight ...raised the question : Why was Bohr successful?


## Prince Louise de Broglie \& Matter Waves

- Key to Bohr atom was Angular momentum quantization
- Why this Quantization: $\mathrm{mvr}=|\mathrm{L}|=\mathrm{nh} / 2 \pi$ ?
- Invoking symmetry in nature, Prince Louise de Broglie conjectured:

Because photons have wave and particle like nature $\rightarrow$ particles may have wave like properties !!

Electrons have accompanying "pilot" wave (not EM) which guide particles thru spacetime.


## A PhD Thesis Fit For a Prince!

- Matter Wave !
- "Pilot wave" of $\lambda=\mathrm{h} / \mathrm{p}=\mathrm{h} /(\gamma \mathrm{mv})$
- Frequency of pilot wave f=E/h
- Consequence:
- If matter has wave like properties then there would be interference (destructive \& constructive) of some kind!
- Analogy of standing waves on a plucked string to explain the quantization condition of Bohr orbits


## Matter Waves : How big, how small ?

$$
\begin{aligned}
& \text { 1. Wavelength of baseball, } \mathrm{m}=140 \mathrm{~g}, \mathrm{v}=27 \mathrm{~m} / \mathrm{s} \\
& \lambda=\frac{\mathrm{h}}{\mathrm{p}}=\frac{h}{m v}=\frac{6.63 \times 10^{-34} \mathrm{~J} . \mathrm{s}}{(.14 \mathrm{~kg})(27 \mathrm{~m} / \mathrm{s})}=1.75 \times 10^{-34} \mathrm{~m} \\
& \Rightarrow \quad \lambda_{\text {baseball }} \lll \text { size of nucleus }
\end{aligned}
$$

Baseball "looks" like a particle
2. Wavelength of electron $K=120 \mathrm{eV}$ (assume NR)
$\mathrm{K}=\frac{\mathrm{p}^{2}}{2 \mathrm{~m}} \Rightarrow p=\sqrt{2 m K}$

$$
=\sqrt{2\left(9.11 \times 10^{-31}\right)(120 \mathrm{eV})\left(1.6 \times 10^{-19}\right)}
$$

$$
=5.91 \times 10^{-24} \mathrm{Kg} . \mathrm{m} / \mathrm{s}
$$

$$
\lambda_{e}=\frac{h}{p}=\frac{6.63 \times 10^{-34} \mathrm{~J} . \mathrm{s}}{5.91 \times 10^{-24} \mathrm{~kg} . \mathrm{m} / \mathrm{s}}=1.12 \times 10^{-10} \mathrm{~m}
$$

$$
\Rightarrow \quad \lambda_{e} \simeq \text { Size of atom }!!
$$

Models of Vibrations on a Loop: Model of e in atom



Reminder: Light as a Wave : Bragg Scattering Expt


## Verification of Matter Waves: Davisson \& Germer Expt

If electrons have associated wave like properties $\rightarrow$ expect interference pattern when incident on a layer of atoms (reflection diffraction grating) with inter-atomic separation $d$ such that

$$
\text { path } \operatorname{diff} \mathrm{AB}=\mathrm{d} \sin \vartheta=\mathrm{n} \lambda
$$



Electrons Diffract in Crystal, just like X-rays


Diffraction pattern produced by 600 eV electrons incident on a Al foil target

Notice the waxing
and waning of scattered electron

Intensity.

What to expect if
electron had no wave
like attribute

Davisson-Germer Experiment: 54 eV electron Beam


Polar graphs of DG expt with different electron accelerating potential when incident on same crystal ( $d=$ const $)$


Peak at $\Phi=50^{\circ}$
when $\mathrm{V}_{\mathrm{acc}}=54 \mathrm{~V}$

Analyzing Davisson-Germer Expt with de Broglie idea
de Broglie $\lambda$ for electron accelerated thru $\mathrm{V}_{\text {acc }}=54 \mathrm{~V}$
$\frac{1}{2} m v^{2}=K=\frac{p^{2}}{2 m}=e V \Rightarrow v=\sqrt{\frac{2 e V}{m}} ; p=m v=m \sqrt{\frac{2 e V}{m}}$
If you believe de Broglie
$\lambda=\frac{h}{p}=\frac{h}{m v}=\frac{h}{m \sqrt{\frac{2 e V}{m}}=\frac{h}{\sqrt{2 m e V}}=\lambda^{\text {predict }}}$


For $\mathrm{V}_{\text {acc }}=54$ Volts $\Rightarrow \lambda=1.67 \times 10^{-10} \mathrm{~m}$ (de Broglie) Exptal data from Davisson-Germer Observation: $\mathrm{d}_{\text {nickel }}=2.15 \dot{\mathrm{~A}}=2.15 \times 10^{-10} \mathrm{~m}$ (from Bragg Scattering)
$\theta_{\text {diff }}^{\max }=50^{\circ}$ (observation from scattering intensity plot)
Diffraction Rule : $\mathrm{d} \sin \phi=\mathrm{n} \lambda$
For Principal Maxima $(\mathrm{n}=1) ; \lambda^{\text {meas }}=(2.15 \dot{\mathrm{~A}})\left(\sin 50^{\circ}\right)$




## Just What is Waving in Matter Waves?

For waves in an ocean, it's the water that "waves"
For sound waves, it's the molecules in medium For light it's the $\mathbf{E} \& \mathbf{B}$ vectors that oscillate

Just What's "waving" in matter waves ?

- It's the PROBABLILITY OF FINDING THE PARTICLE that waves!
- Particle can be represented by a wave packet
- At a certain location (x)
- At a certain time ( t )
- Made by superposition of many sinusoidal waves of different amplitudes, wavelengths $\lambda$ and frequency $f$
- It's a "pulse" of probability in spacetime


## What Wave Does Not Describe a Particle



- What wave form can be associated with particle's pilot wave?
- A traveling sinusoidal wave? $y=A \cos (k x-\omega t+\Phi)$
- Since de Broglie "pilot wave" represents particle, it must travel with same speed as particle ......(like me and my shadow)

Phase velocity $\left(\mathrm{v}_{\mathrm{p}}\right)$ of sinusoidal wave: $\mathrm{v}_{\mathrm{p}}=\lambda f$ In Matter:
(a) $\lambda=\frac{\mathrm{h}}{p}=\frac{h}{\gamma m v}$
(b) $\mathrm{f}=\frac{\mathrm{E}}{\mathrm{h}}=\frac{\gamma m c^{2}}{\mathrm{~h}}$
$\Rightarrow \mathrm{v}_{\mathrm{p}}=\lambda f=\frac{E}{p}=\frac{\gamma m c^{2}}{\gamma m v}=\frac{c^{2}}{v}>c!$

Single sinusoidal wave of infinite extent does not represent particle localized in space

Need "wave packets" localized Spatially (x) and Temporally (t)

## How To Make Wave Packets : Just Beat it !

Superposition of two sound waves of slightly different frequencies $f_{1}$ and $f_{2}, f_{1} \cong f_{2}$

Pattern of beats is a series of wave packets
Beat frequency $\mathrm{f}_{\text {beat }}=\mathrm{f}_{2}-\mathrm{f}_{1}=\Delta \mathrm{f}$
$\Delta f=$ range of frequencies that are superimposed to form
the wave packet


Addition of 2 Waves with slightly different $\lambda$ and slightly different $\omega$ Resulting wave's "displacement " $\mathrm{y}=\mathrm{y}_{1}+y_{2}$ :
$y=A\left[\cos \left(k_{1} x-w_{1} t\right)+\cos \left(k_{2} x-w_{2} t\right)\right]$
Trignometry : $\cos \mathrm{A}+\cos \mathrm{B}=2 \cos \left(\frac{\mathrm{~A}+\mathrm{B}}{2}\right) \cos \left(\frac{\mathrm{A}-\mathrm{B}}{2}\right)$
$\therefore y=2 A\left[\left(\cos \left(\frac{k_{2}-k_{1}}{2} x-\frac{w_{2}-w_{1}}{2} t\right)\right)\left(\cos \left(\frac{k_{2}+k_{1}}{2} x-\frac{w_{2}+w_{1}}{2} t\right)\right)\right]$
since $\mathrm{k}_{2} \cong k_{1} \cong k_{\text {ave }}, w_{2} \cong w_{1} \cong w_{\text {ave }}, \Delta k \ll k, \Delta w \ll w$
$y=2 A\left[\left(\cos \left(\frac{\Delta k}{2} x-\frac{\Delta w}{2} t\right)\right) \cos (k x-w t)\right] \equiv \mathrm{y}=\mathrm{A}^{\prime} \cos (k x-w t)$,
$\mathrm{A}^{\prime}$ oscillates in $\mathrm{x}, \mathrm{t} ; A^{\prime}=2 A\left(\cos \left(\frac{\Delta k}{2} x-\frac{\Delta w}{2} t\right)\right)=$ modulated amplitude
$w$ High-frequency wave
Phase Vel $\mathrm{V}_{p}=\frac{W_{\text {ave }}}{k_{\text {ave }}}$
Group Vel $\quad \mathrm{V}_{\mathrm{g}}=\frac{\Delta w}{\Delta k}$
$V_{g}:$ Vel of envelope $=\frac{d w}{d k}$
Wave Group Or packet

Non-repeating wave packet can be created thru superposition Of many waves of similar (but different) frequencies and wavelengths


The superposition of the many waves spanning a range of frequencies is a wave packet.

## Wave Packet : Localization

Finite \# of diff. Monochromatic waves always produce INFINTE sequence of repeating wave groups $\rightarrow$ can't describe (localized) particle To make localized wave packet, add " infinite" \# of waves with
Well chosen Amplitude A, Wave number k and ang. f requency $\omega$


## Group Velocity, Phase Velocity and Dispersion



Example: Water Wave packet With $V_{g}=V_{p} / 2$


Wave packet for which the group velocity=1/2 phase velocity

The $\uparrow$, representing a point of constant phase for the dominant $\lambda$, travels with $\mathrm{V}_{\mathrm{p}}$

The $\oplus$ at center of group travels with group velocity $\left(\mathrm{V}_{\mathrm{g}}\right)$

A Dispersive Wave Packet Moving Along X Axis


The O indicates position of the classical particle. The Wave packet spreads out in x \& y directions since $\mathrm{V}_{\mathrm{p}}$ of constituent waves depends on wavelength $\lambda$ of the wave

## Group Velocity $V_{g}$ of Matter Wave Packets

Consider An Electron:
mass $=\mathrm{m} \quad$ velocity $=\mathrm{v}, \quad$ momentum $=\mathrm{p}$
Energy $\mathrm{E}=\mathrm{hf}=\gamma \mathrm{mc}^{2} ; \quad \omega=2 \pi f=\frac{2 \pi}{\mathrm{~h}} \gamma \mathrm{mc}^{2}$
Wavelength $\lambda=\frac{\mathrm{h}}{\mathrm{p}} ; \mathrm{k}=\frac{2 \pi}{\lambda} \Rightarrow k=\frac{2 \pi}{h} \gamma m \nu$
Group Velocity : $\mathrm{V}_{\mathrm{g}}=\frac{d w}{d k}=\frac{d w / d v}{d k / d v}$
(b)
$\frac{d w}{d v}=\frac{d}{d v}\left[\frac{\frac{2 \pi}{\mathrm{~h}} \mathrm{mc}^{2}}{\left[1-\left(\frac{\mathrm{v}}{\mathrm{c}}\right)^{2}\right]^{1 / 2}}\right]=\frac{2 \pi \mathrm{mv}}{\mathrm{h}\left[1-\left(\frac{\mathrm{v}}{\mathrm{c}}\right)^{2}\right]^{3 / 2}} \& \frac{d k}{d \nu}=\frac{d}{d v}\left[\frac{2 \pi}{h\left[1-\left(\frac{\mathrm{v}}{\mathrm{c}}\right)^{2}\right]^{1 / 2}} m v\right]=\frac{2 \pi \mathrm{~m}}{\mathrm{~h}\left[1-\left(\frac{\mathrm{v}}{\mathrm{c}}\right)^{2}\right]^{3 / 2}}$
$\mathrm{V}_{\mathrm{g}}=\frac{d w}{d k}=\frac{d w / d v}{d k / d v}=v \Rightarrow$ Group velocity of electron Wave packet "pilot wave"
is same as electron's physical velocity
But velocity of individual waves making up the wave packet $\mathrm{V}_{\mathrm{p}}=\frac{w}{k}=\frac{c^{2}}{v}>c$ ! (not physical)

## Wave Packets \& Uncertainty Principles



What can we learn from this simple model?

- Distance $\Delta \mathrm{X}$ between adjacent minima $=\left(\mathrm{X}_{2}\right)_{\text {node }}-\left(\mathrm{X}_{1}\right)_{\text {node }}$
- Define $\mathrm{X}_{1}=0$ then phase diff from $\mathrm{X}_{1} \rightarrow \mathrm{X}_{2}=\pi \quad$ (similarly for $\mathrm{t}_{1} \rightarrow \mathrm{t}_{2}$ )

Node at $\mathrm{y}=0=2 \mathrm{~A} \cos \left(\frac{\Delta \mathrm{w}}{2} t-\frac{\Delta k}{2} x\right)$, Examine x or t behavior
$\Rightarrow$ in $\mathrm{x}: \Delta k \cdot \Delta x=\pi \Rightarrow$ Need to combine many waves of diff. k to make small $\Delta x$ pulse
$\Delta \mathrm{x}=\frac{\pi}{\Delta \mathrm{k}}$, for small $\Delta \mathrm{x} \rightarrow 0 \Rightarrow \Delta \mathrm{k} \rightarrow \infty$ \& Vice Verca
and In $\mathrm{t}: \Delta \mathrm{w} \cdot \Delta t=\pi \Rightarrow$ Need to combine many waves of diff $\omega$ to make small $\Delta t$ pulse

$$
\Delta t=\frac{\pi}{\Delta \omega} \text {, for small } \Delta t \rightarrow 0 \Rightarrow \Delta \omega \rightarrow \infty \text { \& Vice Verca }
$$

## Signal Transmission and Bandwidth Theory

- Short duration pulses are used to transmit digital info
- Over phone line as brief tone pulses
- Over satellite link as brief radio pulses
- Over optical fiber as brief laser light pulses
- Ragardless of type of wave or medium, any wave pulse must obey the fundamental relation
- $\Delta \omega \Delta t \cong \pi$
- Range of frequencies that can be transmitted are called bandwidth of the medium
- Shortest possible pulse that can be transmitted thru a medium is $\Delta \mathrm{t}_{\text {min }} \cong \pi / \Delta \omega$
- Higher bandwidths transmits shorter pulses \& allows high data rate

Wave Packets \& The Uncertainty Principles of Subatomic Physics
in space $\mathrm{x}: \Delta \mathrm{k} \cdot \Delta x=\pi \Rightarrow$ since $\mathrm{k}=\frac{2 \pi}{\lambda}, \mathrm{p}=\frac{\mathrm{h}}{\lambda}$

$$
\Rightarrow \quad \Delta p . \Delta x=h / 2
$$

usually one writes $\Delta p . \Delta x \geq \hbar / 2$ approximate relation

In time $\mathrm{t}: \Delta w \cdot \Delta t=\pi \Rightarrow$ since $\omega=2 \pi f, E=h f$

$$
\Rightarrow \Delta E . \Delta t=h / 2
$$

usually one writes $\Delta E . \Delta t \geq \hbar / 2$ approximate relation
What do these inequalities mean physically?

Know the Error of Thy Ways: Measurement Error $\rightarrow \Delta$

- Measurements are made by observing something : length, time, momentum, energy
- All measurements have some (limited) precision....no matter the instrument used
- Examples:
- How long is a desk? $\mathrm{L}=(5 \pm 0.1) \mathrm{m}=\mathrm{L} \pm \Delta \mathrm{L}$ (depends on ruler used)
- How long was this lecture ? $\mathrm{T}=(50 \pm 1)$ minutes $=\mathrm{T} \pm \Delta \mathrm{T}$ (depends on the accuracy of your watch)
- How much does Prof. Sharma weigh ? $\mathrm{M}=(1000 \pm 700) \mathrm{kg}=\mathrm{m} \pm \Delta \mathrm{m}$ - Is this a correct measure of my weight ?
- Correct (because of large error reported) but imprecise
- My correct weight is covered by the (large) error in observation


Best Estimate Length: 36 mm
Best Estimate Length: 36 mm
Probable Range: 35.5 to 36.5 mm
Length Measure


Voltage (or time) Measure

## Measurement Error : $x \pm \Delta x$

- Measurement errors are unavoidable since the measurement procedure is an experimental one
- True value of an measurable quantity is an abstract concept
- In a set of repeated measurements with random errors, the distribution of measurements
resembles a Gaussian distribution characterized by the parameter $\sigma$ or $\Delta$ characterizing the width of the distribution


The Gauss, or Normal, Distribution
$G_{X, \sigma}(x)=\frac{1}{\sigma \sqrt{2 \pi}} e^{(x-x)^{2} / 2 \sigma^{2}}$.


135

Measurement Error : $x \pm \Delta x$



## Interpreting Measurements with random Error : $\Delta$



Figure 5.12. The shaded area between $X \pm t \sigma$ is the probability of a measurement within $t$ standard deviations of $X$.


## Where in the World is Carmen San Diego?

Carmen San Diego hidden inside a big box of length L
Suppose you can't see thru the (blue) box, what is you best estimate
of her location inside box (she could be anywhere inside the box)


Your best unbiased measure would be $\mathrm{x}=\mathrm{L} / 2 \pm \mathrm{L} / 2$
There is no perfect measurement, there are always measurement error

## Wave Packets \& Matter Waves



- What is the Wave Length of this wave packet?
- made of waves with $\lambda-\Delta \lambda<\lambda<\lambda+\Delta \lambda$
- De Broglie wavelength $\lambda=\mathrm{h} / \mathrm{p}$
- $\rightarrow$ Momentum Uncertainty: $p-\Delta p<p<p+\Delta p$
- Similarly for frequency $\omega$ or f
- made of waves with $\omega-\Delta \omega<\omega<\omega+\Delta \omega$

Planck's condition $\mathrm{E}=\mathrm{hf}=\mathrm{h} \omega / 2$
$\rightarrow$ Energy Uncertainty: $\mathrm{E}-\Delta \mathrm{E}<\mathrm{E}<\mathrm{E}+\Delta \mathrm{E}$

## Back to Heisenberg's Uncertainty Principle

$\Delta x . \Delta p \geq h / 4 \pi \Rightarrow$ If the measurement of the position of a particle is made with a precision $\Delta x$ and a SIMULTANEOUS measurement of its momentum $p_{x}$ in the $X$ direction, then the product of the two uncertainties (measurement errors) can never be smaller than $\cong h / 4 \pi$ irrespective of how precise the measurement tools

- $\Delta \mathrm{E} . \Delta \mathrm{t} \geq \mathrm{h} / 4 \pi \Rightarrow$ If the measurement of the energy E of a particle is made with a precision $\Delta \mathrm{E}$ and it took time $\Delta \mathrm{t}$ to make that measurement, then the product of the two uncertainties (measurement errors) can never be smaller than $\cong h / 4 \pi$ irrespective of how precise the measurement tools

These rules arise from the way we constructed the Perhaps these rules wave packets describing Matter "pilot" waves are bogus, can we verify this with some physical picture ??

## Are You Experienced ?

- What you experience is what you observe
- What you observe is what you measure
- No measurement is perfect, they all have measurement error: question is of the degree
- Small or large $\Delta$
- Uncertainty Principle and Breaking of Conservation Rules
- Energy Conservation
- Momentum Conservation


## The Act of Observation (Compton Scattering)

Observations of particle motion by means of scattered illumination. When the incident wavelength is reduced to accommodate the size of the particle, the momentum transferred by the photon becomes large enough to disturb the observed motion.


## Act of Observation Tells All



Compton Scattering: Shining light to observe electron
Photon scattering off an electron, Seeing $\rightarrow$ the photon enters my eye


The act of Observation DISTURBS
the object being watched, here the electron moves away from where it was originally



## Diffraction By a Circular Aperture (Lens)

See Resnick, Halliday Walker 6 ${ }^{\text {th }}$ Ed (on S.Reserve), Ch 37, pages 898-900


Fig. 37-9 The diffraction pattern of a circular aperture. Note the central maximum and the circular secondary maxima. The figure has been overexposed to bring out these secondary maxima, which are much less intense than the central maximum.

Diffracted image of a point source of light thru a lens ( circular aperture of size d )

First minimum of diffraction pattern is located by


See previous picture for definitions of $\vartheta, \lambda, \mathrm{d}$

## Resolving Power of Light Thru a Lens

Image of 2 separate point sources formed by a converging lens of diameter d , ability to resolve them depends on $\lambda \& d$ because of the Inherent diffraction in image formation


## Aftermath of Uncertainty Principle

- Deterministic (Newtonian) physics topples over
- Newton's laws told you all you needed to know about trajectory of a particle
- Apply a force, watch the particle go!
- Know every thing! X, v, p, F, a

- Can predict exact trajectory of particle if you had perfect device
- No so in the subatomic world !
- Of small momenta, forces, energies
- Can't predict anything exactly
- Can only predict probabilities
- There is so much chance that the particle landed here or there
- Cant be sure !....cognizant of the errors of thy observations


## All Measurements Have Associated Errors

- If your measuring apparatus has an intrinsic inaccuracy (error) of amount $\Delta \mathrm{p}$
- Then results of measurement of momentum $p$ of an object at rest can easily yield a range of values accommodated by the measurement imprecision :
$--\Delta \mathrm{p} \leq \mathrm{p} \leq \Delta \mathrm{p}$ : you will measure any of these values for the momentum of the particle
- Similarly for all measurable quantities like X, t, Energy !



## Implications of Uncertainty Principles

A bound "particle" is one that is confined in some finite region of space


One of the cornerstones of Quantum mechanics is that bound particles can not be stationary - even at Zero absolute temperature !

There is a non-zero limit on the kinetic energy of a bound particle


## Fluctuations In The Vacuum : Breaking Energy Conservation Rules

Vacuum, at any energy, is bubbling with particle creation and annihilation
AE . At a $h 2 \pi$ implies that you can (in principle) pull out an clephant t antecephant from NOTHING (Vacemm) but for a very very shor fime At!

How Much Time : $\Delta t=\frac{\hbar}{2 M c^{2}}$
How cool is that!


Strong Force Within Nucleus $\rightarrow$ Exchange Force and Virtual Particles


Repulsive force
Strong Nuclear force can be modeled as exchange of virtual particles called $\pi^{ \pm}$mesons by nucleons (protons neutrons)
$\pi^{ \pm}$mesons are emitted by proton and reabsorbed by a neutron

The short range of the Nuclear force is due to the "large mass of the exchanged meson
$\mathrm{M}_{\pi}=140 \mathrm{MeV} / \mathrm{c}^{2}$


## Range of Nuclear Exchange Force

How long can the emitted virtual particle last?
$\Delta E \times \Delta \mathrm{t} \geq \hbar$
The virtual particle has rest mass + kinetic energy
$\Rightarrow$ Its energy $\Delta E \geq M c^{2}$

$\Rightarrow$ Particle can not live for more than $\square$
Range R of the meson (and thus the exchange force)
$\mathrm{R}=\mathrm{c} \Delta \mathrm{t}=\mathrm{c} \hbar / M c^{2}=\hbar / M c$

For $\mathrm{M}=140 \mathrm{MeV} / \mathrm{c}^{2} \Rightarrow R \simeq \frac{1.06 \times 10^{-34} \mathrm{~J} . \mathrm{s}}{\left(140 \mathrm{MeV} / \mathrm{c}^{2}\right) \times \mathrm{c}^{2} \times\left(1.60 \times 10^{-13} \mathrm{~J} / \mathrm{MeV}\right)}$
$R \simeq 1.4 \times 10^{-15} \mathrm{~m}=1.4 \mathrm{fm}$

## Subatomic Cinderella Act

- Neutron emits a charged pion for a time $\Delta t$ and becomes a (charged) proton
- After time $\Delta \mathrm{t}$, the proton reabsorbs charged pion particle ( $\pi^{-}$) to become neutron again
- But in the time $\Delta t$ that the positive proton and $\pi$ - particle exist, they can interact with other charged particles
- After time $\Delta t$ strikes, the Cinderella act is over!




Wave Phenomena $\rightarrow$ Interference and Diffraction



## Interference Pattern of Electrons When Both slits open

Growth of 2-slit Interference pattern thru different exposure periods photographic plate (screen) struck by


Watching Which Hole Electron Went Thru By Shining Intense Light


Watching electrons with dim light: See light flash \& hear detector clicks
Low intensity light $\rightarrow$ Not many photons incident
Maybe a photon hits the electron (See flash, hear click)
Or Maybe the photon misses the electron (no flash, only click)
Probability $\mathrm{P}_{12}$ when both holes open and I see the flash and hear the detector click


Watching electrons in dim light: don't see flash but hear detector clicks


Compton Scattering: Shining light to observe electron



## Summary of Experiments So Far

1. Probability of an event is given by the square of amplitude of a complex \# $\Psi$ : Probability Amplitude
2. When an event occurs in several alternate ways, probability amplitude for the event is sum of probability amplitudes for each way considered seperately. There is interference:

$$
\begin{aligned}
& \Psi=\Psi_{1}+\Psi_{2} \\
& P_{12}=\left|\Psi_{1}+\Psi_{2}\right|^{2}
\end{aligned}
$$

3. If an experiment is done which is capable of determining whether one or other alternative is actually taken, probability for event is just sum of each alternative - Interference pattern is LOST !

## Is There No Way to Beat Uncertainty Principle?

- How about NOT watching the electrons!
- Let's be a bit crafty !!
- Since this is a Thought experiment $\rightarrow$ ideal conditions
- Mount the wall on rollers, put a lot of grease $\rightarrow$ frictionless
- Wall will move when electron hits it
- Watch recoil of the wall containing the slits when the electron hits it
- By watching whether wall moved up or down I can tell
- Will my ingenious plot succeed?

Measuring The Recoil of The Wall $\Rightarrow$ Not Watching Electron!


## Losing Out To Uncertainty Principle

- To measure the RECOIL of the wall $\Rightarrow$
- must know the initial momentum of the wall before electron hit it
- Final momentum after electron hits the wall
- Calculate vector sum $\rightarrow$ recoil
- Uncertainty principle :
- To do this $\Rightarrow \Delta \mathrm{P}=0 \rightarrow \Delta \mathrm{X}=\infty$ [can not know the position of wall exactly]
- If don't know the wall location, then down know where the holes are
- Holes will be in different place for every electron that goes thru
$\rightarrow$ The center of interference pattern will have different (random) location for each electron
- Such random shift is just enough to Smear out the pattern so that no interference is observed !


## Summary

- Probability of an event in an ideal experiment is given by the square of the absolute value of a complex number $\Psi$ which is call probability amplitude
- $\mathrm{P}=$ probability
- $\Psi=$ probability amplitude, $\mathrm{P}=|\Psi|^{2}$
- When an even can occur in several alternative ways, the probability amplitude for the event is the sum of the probability amplitudes for each way considered separately. There is interference:

$$
\begin{aligned}
& \Psi=\Psi_{1}+\Psi_{2} \\
& \mathrm{P}=\left|\Psi_{1}+\Psi_{2}\right|^{2}
\end{aligned}
$$

- If an experiment is performed which is capable of determining whether one or other alternative is actually taken, the probability of the event is the sum of probabilities for each alternative. The interference is lost: $\mathrm{P}=\mathrm{P}_{1}+\mathrm{P}_{2}$


## The Lesson Learnt

- In trying to determine which slit the particle went through, we are examining particle-like behavior
- In examining the interference pattern of electron, we are using wave like behavior of electron

Bohr's Principle of Complementarity:
It is not possible to simultaneously determine physical observables in terms of both particles and waves

The Bullet Vs The Electron: Each Behaves the Same Way


## Quantum Mechanics of Subatomic Particles

- Act of Observation destroys the system (No watching!)
- If can't watch then all conversations can only be in terms of Probability P
- Every particle under the influence of a force is described by a Complex wave function $\Psi(x, y, z, t)$
- $\Psi$ is the ultimate DNA of particle: contains all info about the particle under the force (in a potential e.g Hydrogen )
- Probability of per unit volume of finding the particle at some point ( $\mathrm{x}, \mathrm{y}, \mathrm{z}$ ) and time t is given by
- $\mathrm{P}(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})=\Psi(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t}) . \Psi^{*}(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})=|\Psi(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})|^{2}$
- When there are more than one path to reach a final location then the probability of the event is
$-\Psi=\Psi_{1}+\Psi_{2}$
$-\mathrm{P}=\left|\Psi^{*} \Psi\right|=\left|\Psi_{1}\right|^{2}+\left|\Psi_{2}\right|^{2}+2\left|\Psi_{1}\right| \Psi_{2} \mid \cos \phi$


## Quantum Mechanics of Subatomic Particles

- Act of Observation destroys the system (No watching!)
- If can't watch then all conversations can only be in terms of Probability P
- Every particle under the influence of a force is described by a Complex wave function $\Psi(x, y, z, t)$
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$-\mathrm{P}(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})=\Psi(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t}) . \Psi^{*}(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})=|\Psi(\mathrm{x}, \mathrm{y}, \mathrm{z}, \mathrm{t})|^{2}$
- When there are more than one path to reach a final location then the probability of the event is
$-\Psi=\Psi_{1}+\Psi_{2}$
$-\mathrm{P}=\left|\Psi^{*} \Psi\right|=\left|\Psi_{1}\right|^{2}+\left|\Psi_{2}\right|^{2}+2\left|\Psi_{1}\right| \Psi_{2} \mid \cos \phi$

- Although not possible to specify with certainty the location of particle, its possible to assign probability $\mathrm{P}(\mathrm{x}) \mathrm{dx}$ of finding particle between $x$ and $x+d x$
- $\mathrm{P}(\mathrm{x}) \mathrm{dx}=|\Psi(\mathrm{x}, \mathrm{t})|^{2} \mathrm{dx}$
- E.g intensity distribution in light diffraction pattern is a measure of the probability that a photon will strike a given point within the pattern



## Y: The Wave function Of A Particle

- The particle must be some where

$$
\int_{-\infty}^{+\infty}|\psi(x, t)|^{2} d x=1
$$

- Any $\Psi$ satisfying this condition is

NORMALIZED

- Prob of finding particle in finite interval
$P(a \leq x \leq b)=\int_{a} \psi^{*}(x, t) \psi(x, t) d x$
- Fundamental aim of Quantum Mechanics
- Given the wavefunction at some instant (say $\mathrm{t}=0$ ) find $\Psi$ at some subsequent time t
- $\Psi(\mathrm{x}, \mathrm{t}=0) \rightarrow \Psi(\mathrm{x}, \mathrm{t})$...evolution
- Think of a probabilistic view of particle's "newtonian trajectory"
- We are replacing Newton's $2^{\text {nd }}$ law for subatomic systems

The Wave Function is a mathematical function that describes a physical object $\rightarrow$ Wave function must have some rigorous properties :

- $\Psi$ must be finite
- $\Psi$ must be continuous fn of $\mathrm{x}, \mathrm{t}$
- $\Psi$ must be single-valued
- $\Psi$ must be smooth $\mathrm{fn} \rightarrow$ $\frac{d \psi}{d x}$ must be continuous
WHY?

Bad Wave Functions Of Physical Systems : You Decide Why


## A Simple Wave Function : Free Particle

- Imagine a free particle of mass $m$, momentum $p$ and $K=p^{2} / 2 m$
- Under no force, no attractive or repulsive potential to influence it
- Particle is where it wants : can be any where $[-\infty \leq x \leq+\infty]$
- Has No relationship, no mortgage, no quiz, no final exam....its essentially a bum!
- how to describe a quantum mechanical bum?
- $\Psi(\mathrm{x}, \mathrm{t})=\mathrm{Ae}^{\mathrm{i}(\mathrm{kx}-\omega \mathrm{t})}=\mathrm{A}(\operatorname{Cos}(\mathrm{kx}-\omega \mathrm{t})+\mathrm{i} \sin (\mathrm{kx}-\omega \mathrm{t}))$
$k=\frac{p}{\hbar} ; \quad \omega=\frac{\mathrm{E}}{\hbar}$
For non-relativistic particles
Has definite momentum
and energy but location unknown!
$\mathrm{E}=\frac{\mathrm{p}^{2}}{2 \mathrm{~m}} \Rightarrow \omega(\mathrm{k})=\frac{\hbar \mathrm{k}^{2}}{2 \mathrm{~m}}$

Wave Function of Different Kind of Free Particle : Wave Packet
Sum of Plane Waves:
$\Psi(x, 0)=\int^{+\infty} a(k) e^{i k x} d k$
Combine many free waves to create a localized wave packet (group)
$\Psi(x, t)=\int a(k) e^{i(k x-\omega t)} d k$
Wave Packet initially localized
in $\Delta X, \Delta t$ undergoes dispersion
The more you know now, The less you will know later

Why?

Spreading is due to DISPERSION resulting
from the fact that phase velocity of individual
waves making up the packet depends on $\lambda(\mathrm{k})$


## Normalization Condition: Particle Must be Somewhere

$\xrightarrow{\square}$
Example: $\psi(x, 0)=C e^{\left|x_{0}\right|}, \quad C \& x_{0}$ are constants
This is a symmetric wavefunction with diminishing amplitude
The Amplitude is maximum at $\mathrm{x}=0 \Rightarrow$ Probability is max too
Normalization Condition: How to figure out C ?

$$
\begin{aligned}
& \text { A real particle must be somewhere: Probability of finding particle is finite } \\
& \mathrm{P}(-\infty \leq \mathrm{x} \leq+\infty)=\int_{-\infty}^{+\infty}|\psi(x, 0)|^{2} d x=\int_{-\infty}^{+\infty} C^{2} e^{-2 \frac{\mid x}{x_{0}}} d x=1 \\
& \Rightarrow 1=2 C^{2} \int_{0}^{\infty} e^{-2-\frac{\mid x}{x_{0}}} d x=2 C^{2}\left[\frac{x_{0}}{2}\right]=C^{2} x_{0} \\
& \Rightarrow \psi(x, 0)=\frac{1}{\sqrt{x_{0}}} e^{-\frac{x}{x_{0}}}
\end{aligned}
$$

Probability of finding particle within a certain location $x \pm \Delta x$


$$
\begin{aligned}
& \mathrm{P}\left(-\mathrm{x}_{0} \leq \mathrm{x} \leq+\mathrm{x}_{0}\right)=\int_{-x_{0}}^{+\mathrm{x}_{0}}|\psi(x, 0)|^{2} d x=\int_{-\mathrm{x}_{0}}^{+\mathrm{x}_{0}} C^{2} e^{-2 \left\lvert\, \frac{x}{x_{0}}\right.} d x \\
& =2 C^{2}\left[\frac{x_{0}}{2}\right]\left[1-e^{-2}\right]=\left[1-e^{-2}\right]=0.865 \Rightarrow 87 \%
\end{aligned}
$$

## Where Do Wave Functions Come From ?

- Are solutions of the time dependent Schrödinger Differential Equation (inspired by Wave Equation seen in 2C)


## $-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \Psi(x, t)}{\partial x^{2}}+U(x) \Psi(x, t)=i \hbar \frac{\partial \Psi(x, t)}{\partial t}$

- Given a potential $\mathrm{U}(\mathrm{x}) \rightarrow$ particle under certain force
$-\mathrm{F}(\mathrm{x})=-\frac{\partial U(x)}{\partial x}$



## Introducing the Schrodinger Equation

Consider for simplicity just a one-dimensional system

$$
\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \Psi(x, t)}{\partial x^{2}}+U(x) \Psi(x, t)=i \hbar \frac{\partial \Psi(x, t)}{\partial t}
$$

- $\mathrm{U}(\mathrm{x})=$ characteristic Potential of the system
- Different potential for different types of forces
- Hence different solutions for the S eqn.
- $\rightarrow$ characteristic wavefunctions for a particular $\mathrm{U}(\mathrm{x})$

Schrodinger Equation in 1, 2, 3 dimensional systems
1-dimension

$$
-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \Psi(x, t)}{\partial x^{2}}+U(x) \Psi(x, t)=i \hbar \frac{\partial \Psi(x, t)}{\partial t}
$$

2-dimension

$$
-\frac{\hbar^{2}}{2 m}\left[\frac{\partial^{2} \Psi(x, y, t)}{\partial x^{2}}+\frac{\partial^{2} \Psi(x, y, t)}{\partial y^{2}}\right]+U(x, y) \Psi(x, y, t)=i \hbar \frac{\partial \Psi(x, y, t)}{\partial t}
$$

3-dimension
$-\frac{\hbar^{2}}{2 m}\left[\frac{\partial^{2} \Psi(x, y, z, t)}{\partial x^{2}}+\frac{\partial^{2} \Psi(x, y, z, t)}{\partial y^{2}}+\frac{\partial^{2} \Psi(x, y, z, t)}{\partial z^{2}}\right]+U(x, y, z) \Psi(x, y, z, t)=i \hbar \frac{\partial \Psi(x, y, z, t)}{\partial t}$

Schrodinger Wave Equation in Quantum Mechanics
Wavefunction $\psi$ which is a sol. of the Sch. Equation embodies all modern physics experienced/learnt so far:
$\mathrm{E}=\mathrm{hf}, \quad \mathrm{p}=\frac{\mathrm{h}}{\lambda}, \quad \Delta x . \Delta p \sim \hbar, \Delta E . \Delta t \sim \hbar$, quantization etc
Schrodinger Equation is a Dynamical Equation much like Newton's Equation $\overrightarrow{\mathrm{F}}=\mathrm{m} \overrightarrow{\mathrm{a}}$
$\psi(\mathrm{x}, 0) \rightarrow$ Force (potential) $\rightarrow \psi(\mathrm{x}, \mathrm{t})$
Evolves the System as a function of space-time
The Schrodinger Eq. propogates the system
forward \& backward in time:
$\psi(\mathrm{x}, \delta \mathrm{t})=\psi(\mathrm{x}, 0) \pm\left[\frac{d \psi}{d t}\right]_{t=0} \delta t$
Where does it come from ?? ..."First Principles"
......no real "derivation" exists.............

## Time Independent S. Equation

$-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \Psi(x, t)}{\partial x^{2}}+U(x) \Psi(x, t)=i \hbar \frac{\partial \Psi(x, t)}{\partial t}$
Sometimes (depending on the character of the Potential $\mathrm{U}(\mathrm{x}, \mathrm{t})$ )
The Wave function is factorizable: can be broken up

$$
\Psi(\mathrm{x}, \mathrm{t})=\psi(x) \phi(t)
$$

Example: Plane Wave $\Psi(x, t)=e^{i(k x-c t)}=e^{i(k x)} e^{-i(\omega t)}$
In such cases, use seperation of variables to get :
$\frac{-\hbar^{2}}{2 \mathrm{~m}} \phi(t) \cdot \frac{\partial^{2} \psi(x)}{\partial^{2} x}+U(x) \psi(x) \phi(t)=i \hbar \psi(x) \frac{\partial \phi(t)}{\partial t}$
Divide throughout by $\Psi(\mathrm{x}, \mathrm{t})=\psi(\mathrm{x}) \phi(\mathrm{t})$
$\Rightarrow \frac{-\hbar^{2}}{2 \mathrm{~m}} \frac{1}{\psi(x)} \cdot \frac{\partial^{2} \psi(x)}{\partial^{2} x}+U(x)=i \hbar \frac{1}{\phi(t)} \frac{\partial \phi(t)}{\partial t}$
LHS is a function of $x$; RHS is fn of $t$
x and t are independent variables, hence :
$\Rightarrow$ RHS $=$ LHS $=$ Constant $=\mathrm{E}$

Factorization Condition For Wave Function Leads to:
$\square$

$$
\frac{-\hbar^{2}}{2 \mathrm{~m}} \frac{\partial^{2} \psi(x)}{\partial^{2} x}+U(x) \psi(x)=E \psi(x)
$$

$$
i \hbar \frac{\partial \phi(t)}{\partial t}=E \phi(t)
$$

What is the Constant E ? How to Interpret it?
Back to a Free particle :

$$
\begin{aligned}
& \Psi(\mathrm{x}, \mathrm{t})=A \mathrm{e}^{\mathrm{ikx}} \mathrm{e}^{-\mathrm{i} \omega \mathrm{t}}, \psi(\mathrm{x})=A \mathrm{e}^{\mathrm{ikx}} \\
& \mathrm{U}(\mathrm{x}, \mathrm{t})=0
\end{aligned}
$$

Plug it into the Time Independent Schrodinger Equation (TISE) $\Rightarrow$
$\frac{-\hbar^{2}}{2 m} \frac{d^{2}\left(A e^{(i k x)}\right)}{d x^{2}}+0=E A e^{(i k x)} \Rightarrow E=\frac{\hbar^{2} k^{2}}{2 m}=\frac{p^{2}}{2 m}=$ (NR Energy)
Stationary states of the free particle: $\Psi(\mathrm{x}, \mathrm{t})=\psi(\mathrm{x}) \mathrm{e}^{-\mathrm{i} \omega \mathrm{t}}$
$\Rightarrow|\Psi(x, t)|^{2}=|\psi(x)|^{2}$
Probability is static in time t , character of wave function depends on $\psi(x)$

## Schrodinger Eqn: Stationary State Form

- Recall $\rightarrow$ when potential does not depend on time explicitly
- $\mathrm{U}(\mathrm{x}, \mathrm{t})=\mathrm{U}(\mathrm{x})$ only... we used separation of $\mathrm{x}, \mathrm{t}$ variables to simplify
- $\Psi(x, t)=\psi(x) \phi(t)$
- broke S. Eq. into two: one with $x$ only and another with $t$ only
$\frac{-\hbar^{2}}{2 m} \frac{\partial^{2} \psi(x)}{\partial^{2} x}+U(x) \psi(x)=E \psi(x)$

$$
i \hbar \frac{\partial \phi(t)}{\partial t}=E \phi(t)
$$

How to put Humpty-Dumpty back together ? e.g to say how to go from an expression of $\psi(\mathrm{x}) \rightarrow \Psi(\mathrm{x}, \mathrm{t})$ which describes time-evolution of the overall wave function

## Schrodinger Eqn: Stationary State Form

Since $\frac{\mathrm{d}}{\mathrm{dt}}[\ln f(t)]=\frac{1}{f(t)} \frac{\mathrm{d} f(t)}{\mathrm{dt}}$
In i $\hbar \frac{\partial \phi(t)}{\partial \mathrm{t}}=E \phi(t)$, rewrite as $\frac{1}{\phi(t)} \frac{\partial \phi(t)}{\partial \mathrm{t}}=\frac{E}{i \hbar}=-\frac{i E}{\hbar}$
and integrate both sides w.r.t. time
$\int_{t=0}^{t=t} \frac{1}{\phi(t)} \frac{\partial \phi(t)}{\partial \mathrm{t}} d t=\int_{0}^{t}-\frac{i E}{\hbar} d t \Rightarrow \int_{0}^{t} \frac{1}{\phi(t)} \frac{\mathrm{d} \phi(t)}{\mathrm{dt}} d t=-\frac{i E}{\hbar}$
$\therefore \ln \phi(t)-\ln \phi(0)=-\frac{i E}{\hbar} t$, now exponentiate both sides
$\Rightarrow \phi(t)=\phi(0) e^{-\frac{i E}{\hbar} t} \quad ; \phi(0)=$ constant $=$ initial condition $=1(\mathrm{e} . \mathrm{g})$
$\Rightarrow \phi(t)=e^{-\frac{i E}{\hbar} t} \quad \&$ Thus $\Psi(\mathrm{x}, \mathrm{t})=\psi(\mathrm{x}) e^{-\frac{i E}{\hbar} t}$ where $\mathrm{E}=$ energy of system


## Example of a Particle Inside a Box With Infinite Potential



## $\Psi(x)$ for Particle Inside 1D Box with Infinite Potential Walls

Why can't the
particle exist
Outside the box ?
$\rightarrow$ E Conservation


## $\Psi(x)$ for Particle Inside 1D Box with Infinite Potential Walls



Need to figure out values of A, B : How to do that?
Apply BOUNDARY Conditions on the Wavefunction
Since $\psi(x)$ must be continuous everywhere
$\Rightarrow$ match the wavefunction just outside box with
the wavefunction value just inside the box
$\Rightarrow$ At $\mathrm{x}=0 \Rightarrow \psi(x=0)=0 \quad \& \quad$ At $\mathrm{x}=\mathrm{L} \Rightarrow \psi(x=L)=0$
$\mathrm{X}=0 \quad \mathrm{X}=\mathrm{L} \therefore \psi(x=0)=B=0$ (Continuity condition at $\mathrm{x}=0)$
$\& \psi(x=L)=0 \Rightarrow \mathrm{~A} \mathrm{Sin} \mathrm{kL}=0$ (Continuity condition at $\mathrm{x}=\mathrm{L}$ )

$$
\Rightarrow \mathrm{kL}=\mathrm{n} \pi \Rightarrow \mathrm{k}=\frac{\mathrm{n} \pi}{\mathrm{~L}}, n=1,2,3, \ldots \infty
$$

So what does this say about Energy E ? : $\mathrm{E}_{\mathrm{n}}=\frac{\mathrm{n}^{2} \pi^{2} \hbar^{2}}{2 m L^{2}}$
Quantized (not Continuous)!

Quantized Energy levels of Particle in a Box


## What About the Wave Function Normalization ?

The particle's Energy and Wavefunction are determined by a number $n$ We will call $n \rightarrow$ Quantum Number, just like in Bohr's Hydrogen atom What about the wave functions corresponding to each of these energy states?

$$
\begin{aligned}
\psi_{\mathrm{n}} & =A \sin (k x)=A \sin \left(\frac{n \pi x}{L}\right) & & \text { for } 0<\mathrm{x}<\mathrm{L} \\
& =0 & & \text { for } \mathrm{x} \geq 0, \mathrm{x} \geq \mathrm{L}
\end{aligned}
$$

Normalized Condition :
$1=\int_{0}^{\mathrm{L}} \psi_{\mathrm{n}}^{*} \psi_{\mathrm{n}} d x=A^{2} \int_{0}^{L} \operatorname{Sin}^{2}\left(\frac{n \pi x}{L}\right) \quad$ Use $2 \operatorname{Sin}^{2} \theta=1-2 \operatorname{Cos} 2 \theta$
$1=\frac{A^{2}}{2} \int_{0}^{L}\left(1-\cos \left(\frac{2 n \pi x}{L}\right)\right)$ and since $\int \cos \theta=\sin \theta$
$1=\frac{A^{2}}{2} L \Rightarrow A=\sqrt{\frac{2}{L}}$
So $\psi_{\mathrm{n}}=\sqrt{\frac{2}{L}} \sin (k x)=\sqrt{\frac{2}{L}} \sin \left(\frac{n \pi x}{L}\right) \quad$...What does this look like?

Wave Functions : Shapes Depend on Quantum \# n


## Where in The World is Carmen San Diego?

- We can only guess the probability of finding the particle somewhere in x
- For $\mathrm{n}=1$ (ground state) particle most likely at $\mathrm{x}=\mathrm{L} / 2$
- For n=2 (first excited state) particle most likely at L/4, 3L/4
- Prob. Vanishes at $\mathrm{x}=\mathrm{L} / 2$ \& L
- How does the particle get from just before $\mathrm{x}=\mathrm{L} / 2$ to just after?
- QUIT thinking this way, particles don't have trajectories
- Just probabilities of beir somewhere



Classically, where is the particle most Likely to be : Equal prob of being anywhere inside the Box NOT SO says Quantum Mechanics!

## Remember Sesame Street?



This particle in the box is brought to you by the letter

## The QM Prob. of Finding Particle in Some Region in Space

Consider $\mathrm{n}=1$ state of the particle
Ask : What is $\mathrm{P}\left(\frac{\mathrm{L}}{4} \leq x \leq \frac{3 L}{4}\right)$ ?
$\mathrm{P}=\int_{\frac{L}{4}}^{\frac{3 L}{4}}\left|\psi_{1}\right|^{2} d x=\frac{2}{L} \int_{\frac{L}{4}}^{\frac{3 L}{4}} \sin ^{2} \frac{\pi x}{L} d x=\left(\frac{2}{L}\right) \cdot \frac{1}{2} \int_{\frac{L}{4}}^{\frac{3 L}{4}}\left(1-\cos \frac{2 \pi x}{L}\right) d x$
$P=\frac{1}{L}\left[\frac{L}{2}-\right]\left[\frac{L}{2 \pi} \sin \frac{2 \pi x}{L}\right]_{L / 4}^{3 L / 4}=\frac{1}{2}-\frac{1}{2 \pi}\left(\sin \frac{2 \pi}{L} \cdot \frac{3 L}{4}-\sin \frac{2 \pi}{L} \cdot \frac{L}{4}\right)$
$P=\frac{1}{2}-\frac{1}{2 \pi}(-1-1)=0.818 \Rightarrow 81.8 \%$

Classically $\Rightarrow 50 \%$ (equal prob over half the box size)
$\Rightarrow$ Substantial difference between Classical \& Quantum predictions

## When The Classical \& Quantum Pictures Merge: $n \rightarrow \infty$



But one issue is irreconcilable:
Quantum Mechanically the particle can not have $\mathrm{E}=0$
This is a direct consequence of the Uncertainty Principle
The particle moves around with KE inversely proportional to the length Of the (1D) Box

## Finite Potential Barrier

- There are no Infinite Potentials in the real world
- Imagine the cost of as battery with infinite potential diff
- Will cost infinite \$ sum + not available at Radio Shack
- Imagine a realistic potential : Large U compared to KE but not infinite


Classical Picture : A bound particle (no escape) in $0<x<L$ Quantum Mechanical Picture : Use $\Delta \mathrm{E} . \Delta \mathrm{t} \leq \mathrm{h} / 2 \pi$ Particle can leak out of the Box of finite potential $\mathrm{P}(|\mathrm{x}|>\mathrm{L}) \neq 0$

## Finite Potential Well

$$
\begin{aligned}
& \quad \frac{-\hbar^{2}}{2 m} \frac{d^{2} \psi(x)}{d x^{2}}+U \psi(x)=E \psi(x) \\
& \Rightarrow \quad \frac{d^{2} \psi(x)}{d x^{2}}=\frac{2 m}{\hbar^{2}}(U-E) \psi(x) \\
& =\alpha^{2} \psi(x) ; \alpha=\sqrt{\frac{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}{\hbar^{2}}} \\
& \Rightarrow \text { General Solutions : } \psi(x)=A e^{+\alpha x}+B e^{-\alpha x} \\
& \text { Require finiteness of } \psi(x) \\
& \Rightarrow \psi(x)=A e^{+\alpha x} \ldots . . \mathrm{x}<0 \quad \text { (region I) } \\
& \psi(x)=A e^{-\alpha x} \ldots . . \mathrm{x}>\mathrm{L} \quad \text { (region III) }
\end{aligned}
$$

Again, coefficients A \& B come from matching conditions at the edge of the walls $(x=0, L)$
But note that wave fn at $\psi(x)$ at $(\mathrm{x}=0, \mathrm{~L}) \neq 0!$ ! (why?)
Further require Continuity of $\psi(x)$ and $\frac{d \psi(x)}{d x}$
These lead to rather different wave functions

Finite Potential Well: Particle can Burrow Outside Box!
$\square$


Finite Potential Well: Particle can Burrow Outside Box




Particle can be outside the box but only for a time $\Delta t \approx h / \Delta E$
$\Delta \mathrm{E}=$ Energy particle needs to borrow to Get outside $\Delta \mathrm{E}=\mathrm{U}-\mathrm{E}+\mathrm{KE}$
The Cinderella act (of violating E conservation cant last very long

Particle must hurry back (cant be caught with its hand inside the cookie-jar)
Penetration Length $\delta=\frac{1}{\alpha}=\frac{\hbar}{\sqrt{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}}$
If $\mathrm{U} \gg \mathrm{E} \Rightarrow$ Tiny penetration
If $\mathrm{U} \rightarrow \infty \Rightarrow \delta \rightarrow 0$

Finite Potential Well: Particle can Burrow Outside Box

# ! 

$$
\begin{aligned}
& \text { Penetration Length } \delta=\frac{1}{\alpha}=\frac{\hbar}{\sqrt{2 m(U-E)}} \\
& \text { If } \mathrm{U}>\mathrm{E} \Rightarrow \text { Tiny penetration } \\
& \text { If } \mathrm{U} \rightarrow \infty \Rightarrow \delta \rightarrow 0
\end{aligned}
$$

$$
\mathrm{E}_{\mathrm{n}}=\frac{\mathrm{n}^{2} \pi^{2} \hbar^{2}}{2 m(L+2 \delta)^{2}}, n=1,2,3,4 \ldots
$$

When $\mathrm{E}=\mathrm{U}$ then solutions blow up
$\Rightarrow$ Limits to number of bound states $\left(\mathrm{E}_{\mathrm{n}}<U\right)$
When $\mathrm{E}>\mathrm{U}$, particle is not bound and can get either reflected or transmitted across the potential "barrier"

## Measurement Expectation: Statistics Lesson

- Ensemble \& probable outcome of a single measurement or the average outcome of a large \# of measurements
$<x>=\frac{n_{1} x_{1}+n_{2} x_{2}+n_{3} x_{3}+\ldots n_{i} x_{i}}{n_{1}+n_{2}+n_{3}+\ldots n_{i}}=\frac{\sum_{i=1}^{n} n_{i} x_{i}}{N}=\frac{\int_{-\infty}^{\infty} x P(x) d x}{\int_{-\infty}^{\infty} P(x) d x}$

For a general Fn $f(x)$
$<f(x)>=\frac{\sum_{i=1}^{n} n_{i} f\left(x_{i}\right)}{N}=\frac{\int_{-\infty}^{\infty} \psi^{*}(x) f(x) \psi(x) d x}{\int_{-\infty}^{\infty} P(x) d x}$

Sharpness of A Distr:
Scatter around average
$\sigma=\sqrt{\frac{\sum\left(\mathrm{x}_{\mathrm{i}}-\bar{x}\right)^{2}}{N}}$
$\sigma=\sqrt{\left(\overline{x^{2}}\right)-(\bar{x})^{2}}$
$\sigma=$ small $\rightarrow$ Sharp distr.
Uncertainty $\Delta \mathrm{X}=\sigma$

## Particle in the Box, $n=1$, find $<x\rangle$ \& $\Delta x$ ?

$\psi(\mathrm{x})=\sqrt{\frac{2}{\mathrm{~L}}} \sin \left(\frac{\pi}{L} x\right)$
$<\mathrm{x}>=\int_{-\infty}^{\infty} \sqrt{\frac{2}{\mathrm{~L}}} \sin \left(\frac{\pi}{L} x\right) x \sqrt{\frac{2}{\mathrm{~L}}} \sin \left(\frac{\pi}{L} x\right) d x$
$=\frac{2}{L} \int_{0}^{L} x \sin ^{2}\left(\frac{\pi}{L} x\right) d x$, change variable $\theta=\left(\frac{\pi}{L} x\right)$
$\Rightarrow\langle\mathrm{x}\rangle=\frac{2}{\mathrm{~L} \pi^{2}} \int_{0}^{\pi} \theta \sin ^{2} \theta$, use $\sin ^{2} \theta=\frac{1}{2}(1-\cos 2 \theta)$
$\Rightarrow\langle\mathrm{x}\rangle=\frac{2 \mathrm{~L}}{2 \pi^{2}}\left[\int_{0}^{\pi} \theta \mathrm{d} \theta-\int_{0}^{\pi} \theta \cos 2 \theta d \theta\right]$ use $\int u d v=u v-\int \mathrm{vdu}$
$\Rightarrow\langle\mathrm{x}\rangle=\frac{\mathrm{L}}{\pi^{2}}\left(\frac{\pi^{2}}{2}\right)=\frac{L}{2} \quad$ (same result as from graphing $\left.\psi^{2}(x)\right)$
Similarly $\left\langle\mathrm{x}^{2}\right\rangle=\int_{0}^{L} \mathrm{x}^{2} \sin ^{2}\left(\frac{\pi}{L} x\right) d x=\frac{L^{2}}{3}-\frac{L^{2}}{2 \pi^{2}}$
and $\Delta \mathrm{X}=\sqrt{\left\langle\mathrm{x}^{2}\right\rangle-\langle x\rangle^{2}}=\sqrt{\frac{L^{2}}{3}-\frac{L^{2}}{2 \pi^{2}}-\frac{L^{2}}{4}}=0.18 L$

$\Delta X=20 \%$ of L, Particle not sharply confined in Box


## Expectation Values \& Operators: More Formally

- Observable: Any particle property that can be measured
- X,P, KE, E or some combination of them, e,g: $\mathrm{x}^{2}$
- How to calculate the probable value of these quantities for a QM state ?
- Operator: Associates an operator with each observable
- Using these Operators, one calculates the average value of that Observable
- The Operator acts on the Wavefunction (Operand) \& extracts info about the Observable in a straightforward way $\rightarrow$ gets Expectation value for that observable

$$
Q>=\int^{+\infty} \Psi^{*}(x, t)[\hat{Q}] \Psi^{*}(x, t) d x
$$

$Q$ is the observable, $[\hat{Q}]$ is the operator
\& $\langle Q\rangle$ is the Expectation value

$$
\begin{array}{ll}
\text { Examples: }[\mathrm{X}]=\mathrm{x}, & {[\mathrm{P}]=\frac{\hbar}{\mathrm{i}} \frac{d}{d x}} \\
{[\mathrm{~K}]=\frac{[\mathrm{P}]^{2}}{2 \mathrm{~m}}=\frac{-\hbar^{2}}{2 \mathrm{~m}} \frac{\partial^{2}}{\partial x^{2}}} & {[\mathrm{E}]=\mathrm{i} \hbar \frac{\partial}{\partial t}}
\end{array}
$$

## Table 5.2 Common Observables and Associated Operators

| Observable | Symbol | Associated <br> Operator |
| :--- | :---: | :---: |
| position | $x$ | $x$ |
| momentum | $p$ | $\frac{\hbar}{i} \frac{\partial}{\partial x}$ |
| potential energy | $U$ | $U(x)$ |
| kinetic energy | $K$ | $-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial x^{2}}$ |
| hamiltonian | $H$ | $-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial x^{2}}+U(x)$ |
| total energy | $E$ | $i \hbar \frac{\partial}{\partial t}$ |

## Operators $\rightarrow$ Information Extractors

[p] or $\hat{\mathrm{p}}=\frac{\hbar}{\mathrm{i}} \frac{d}{d x}$
Momentum Operator
gives the value of average mometum in the following way:
$<\mathrm{p}>=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})[p] \psi(x) d x=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})\left(\frac{\hbar}{\mathrm{i}}\right) \frac{d \psi}{d x} d x$
Plug \& play form
Similerly :
$[\mathrm{K}]$ or $\hat{\mathrm{K}}=-\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{d^{2}}{d x^{2}}$ gives the value of average KE
$<\mathrm{K}>=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})[K] \psi(x) d x=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})\left(-\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{d^{2} \psi(x)}{d x^{2}}\right) d x$
Similerly
$<\mathrm{U}>=\int^{+\infty} \psi^{*}(\mathrm{x})[U(x)] \psi(x) d x \quad$ : plug in the $\mathrm{U}(\mathrm{x})$ fn for that case
and $<\mathrm{E}\rangle=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})[K+U(x)] \psi(x) d x=\int_{-\infty}^{+\infty} \psi^{*}(\mathrm{x})\left(-\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{d^{2} \psi(x)}{d x^{2}}+U(x)\right) d x$
Hamiltonian Operator $[\mathrm{H}]=[\mathrm{K}]+[\mathrm{U}]$
The Energy Operator $[\mathrm{E}]=\mathrm{i} \hbar \frac{\partial}{\partial t}$ informs you of the average energy

## [H] \& [E] Operators

- $[\mathrm{H}]$ is a function of x
- [E] is a function of t .......they are really different operators
- But they produce identical results when applied to any solution of the time-dependent Schrodinger Eq.
- $[\mathrm{H}] \Psi(\mathrm{x}, \mathrm{t})=[\mathrm{E}] \Psi(\mathrm{x}, \mathrm{t})$

$$
\left[-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial x^{2}}+U(x, t)\right] \Psi(x, t)=\left[i \hbar \frac{\partial}{\partial t}\right] \Psi(x, t)
$$

- Think of S. Eq as an expression for Energy conservation for a Quantum system


## Where do Operators come from ? A touchy-feely answer

Example:[p] The momentum Extractor (operator):
Consider as an example: Free Particle Wavefunction
$\Psi(\mathrm{x}, \mathrm{t})=\mathrm{Ae}^{\mathrm{i}(\mathrm{kx}-\mathrm{wt})} \quad ; \mathrm{k}=\frac{2 \pi}{\lambda}, \lambda=\frac{h}{p} \Rightarrow k=\frac{p}{\hbar}$
rewrite $\Psi(\mathrm{x}, \mathrm{t})=\mathrm{Ae} \mathrm{e}^{\mathrm{i}\left(\frac{p}{\hbar} \mathrm{x}-\mathrm{wt}\right)} ; \frac{\partial \Psi(\mathrm{x}, \mathrm{t})}{\partial \mathrm{x}}=i \frac{p}{\hbar} \mathrm{Ae} \mathrm{e}^{\mathrm{i}\left(\frac{p}{\hbar} \mathrm{x}-\mathrm{wt}\right)}=i \frac{p}{\hbar} \Psi(\mathrm{x}, \mathrm{t})$
$\Rightarrow\left[\frac{\hbar}{\mathrm{i}} \frac{\partial}{\partial x}\right] \Psi(\mathrm{x}, \mathrm{t})=\mathrm{p} \Psi(\mathrm{x}, \mathrm{t})$

So it is not unreasonable to associate $[\mathrm{p}]=\left[\frac{\hbar}{\mathrm{i}} \frac{\partial}{\partial x}\right]$ with observable p

## Example : Average Momentum of particle in box

- Given the symmetry of the 1 D box, we argued last time that $<\mathrm{p}>=0$
: now some inglorious math to prove it !
- Be lazy, when you can get away with a symmetry argument to solve a problem..do it \& avoid the evil integration \& algebra......but be sure!
$\psi_{n}(x)=\sqrt{\frac{2}{L}} \sin \left(\frac{n \pi}{L} x\right) \quad \& \quad \psi^{*}{ }_{n}(x)=\sqrt{\frac{2}{L}} \sin \left(\frac{n \pi}{L} x\right)$
$\langle p\rangle=\int_{-\infty}^{+\infty} \psi^{*}[p] \psi d x=\int_{-\infty}^{\infty} \psi^{*}\left[\frac{\hbar}{i} \frac{d}{d x}\right] \psi d x$
$\langle p\rangle=\frac{\hbar}{i} \frac{2}{L} \frac{n \pi}{L} \int_{-\infty}^{\infty} \sin \left(\frac{n \pi}{L} x\right) \cos \left(\frac{n \pi}{L} x\right) d x$
Since $\int \sin \cos \operatorname{cosax} \mathrm{dx}=\frac{1}{2 \mathrm{a}} \sin ^{2} a x \quad \ldots$ here $\mathrm{a}=\frac{\mathrm{n} \pi}{\mathrm{L}}$
$\Rightarrow\langle p\rangle=\frac{\hbar}{i L}\left[\sin ^{2}\left(\frac{n \pi}{L} x\right]_{x=0}^{x=L}=0\right.$ since $\operatorname{Sin}^{2}(0)=\operatorname{Sin}^{2}(n \pi)=0$
We knew THAT before doing any math !

Quiz 1: What is the <p> for the Quantum Oscillator in its symmetric ground state
Quiz 2: What is the $<\mathrm{p}>$ for the Quantum Oscillator in its asymmetric first excited state

## But what about the $<K E>$ of the Particle in Box?

$<p>=0$ so what about expectation value of $\mathrm{K}=\frac{\mathrm{p}^{2}}{2 \mathrm{~m}}$ ?
$<K\rangle=0$ because $\langle p\rangle=0$; clearly not, since we showed $\mathrm{E}=\mathrm{KE} \neq 0$
Why? What gives?
Because $\mathrm{p}_{\mathrm{n}}= \pm \sqrt{2 m E_{n}}= \pm \frac{n \pi \hbar}{L} ; \quad$ " $\pm$ " is the key!
The AVERAGE $\mathrm{p}=0$, since particle is moving back \& forth

$$
<\mathrm{KE}>=<\frac{\mathrm{p}^{2}}{2 \mathrm{~m}}>\neq 0 ; \quad \operatorname{not} \frac{\left\langle\mathrm{p}^{2}\right\rangle}{2 m}!
$$

Be careful when being "lazy"

Quiz: what about <KE> of a quantum Oscillator?
Does similar logic apply??

## Schrodinger Eqn: Stationary State Form

$P(x, t)=\Psi^{*} \Psi=\psi^{*}(x) e^{+\frac{i E}{\hbar} t} \psi(x) e^{-\frac{i E}{\hbar} t}=\psi^{*}(x) \psi(x) e^{\frac{i E}{\hbar} t-\frac{i E}{\hbar} t}=|\psi(x)|^{2}$
In such cases, $\mathrm{P}(\mathrm{x}, \mathrm{t})$ is INDEPENDENT of time.
These are called "stationary" states because Prob is independent of time Examples : Particle in a box (why?)

Quantum Oscillator (why?)
Total energy of the system depends on the spatial orientation of the system : charteristic of the potential situation!

Simple Harmonic Oscillator:Quantum and Classical
?



Particle of mass $m$ within a potential $U(x)$ $\overrightarrow{\mathrm{F}}(\mathrm{x})=-\frac{d U(x)}{d x}$
$\left.\overrightarrow{\mathrm{F}}(\mathrm{x}=\mathrm{a})=-\frac{d U(x)}{d x} \right\rvert\,=0$,
$\overrightarrow{\mathrm{F}}(\mathrm{x}=\mathrm{b})=0, \overrightarrow{\mathrm{~F}}(\mathrm{x}=\mathrm{c})=0 \ldots$...But $\ldots$
look at the Curvature:

Stable Equilibrium: General Form :
$\mathrm{U}(\mathrm{x})=\mathrm{U}(\mathrm{a})+\frac{1}{2} k(x-a)^{2}$
Rescale $\Rightarrow U(x)=\frac{1}{2} k(x-a)^{2}$
Motion of a Classical Oscillator (ideal)
Ball originally displaced from its equilibirium position, motion confined between $\mathrm{x}=0$ \& $\mathrm{x}=\mathrm{A}$
$\mathrm{U}(\mathrm{x})=\frac{1}{2} k x^{2}=\frac{1}{2} m \omega^{2} x^{2} ; \omega=\sqrt{\frac{k}{m}}=$ Ang. Freq $E=\frac{1}{2} k A^{2} \Rightarrow$ Changing A changes E
E can take any value \& if $\mathrm{A} \rightarrow 0, \mathrm{E} \rightarrow 0$
Max. KE at $\mathrm{x}=0, \mathrm{KE}=0$ at $\mathrm{x}= \pm \mathrm{A}$

## Quantum Picture: Harmonic Oscillator

Find the Ground state Wave Function $\psi(\mathrm{x})$
Find the Ground state Energy E when $\mathrm{U}(\mathrm{x})=\frac{1}{2} m \omega^{2} x^{2}$
Time Dependent Schrodinger Eqn: $\frac{-\hbar^{2}}{2 m} \frac{\partial^{2} \psi(x)}{\partial^{2} x}+\frac{1}{2} m \omega^{2} x^{2} \psi(x)=E \psi(x)$
$\Rightarrow \frac{d^{2} \psi(x)}{d x^{2}}=\frac{2 m}{\hbar^{2}}\left(E-\frac{1}{2} m \omega^{2} x^{2}\right) \psi(x)=0$ What $\psi(\mathrm{x})$ solves this?

Two guesses about the simplest Wavefunction:

1. $\psi(\mathrm{x})$ should be symmetric about x 2. $\psi(\mathrm{x}) \rightarrow 0$ as $\mathrm{x} \rightarrow \infty$
$+\psi(x)$ should be continuous $\& \frac{d \psi(x)}{d x}=$ continuous

My guess: $\psi(\mathrm{x})=\mathrm{C}_{0} e^{-\alpha x^{2}} ;$ Need to find $\mathrm{C}_{0} \& \alpha$ :

What does this wavefunction \& PDF look like?

How to Get $\mathrm{C}_{0} \& \alpha$ ?? ...Try plugging in the wave-function into the time-independent Schr. Eqn.

Time Independent Sch. Eqn \& The Harmonic Oscillator
Master Equation is : $\frac{\partial^{2} \psi(x)}{\partial x^{2}}=\frac{2 m}{\hbar^{2}}\left[\frac{1}{2} m \omega^{2} x^{2}-E\right] \psi(x)$
Since $\psi(x)=C_{0} e^{-\alpha x^{2}}, \frac{d \psi(x)}{d x}=C_{0}(-2 \alpha x) e^{-\alpha x^{2}}$,
$\frac{d^{2} \psi(x)}{d x^{2}}=C_{0} \frac{d(-2 \alpha x)}{d x} e^{-\alpha x^{2}}+C_{0}(-2 \alpha x)^{2} e^{-\alpha x^{2}}=C_{0}\left[4 \alpha^{2} x^{2}-2 \alpha\right] e^{-\alpha x^{2}}$
$\left.\left.\Rightarrow C_{0}\left[4 \alpha^{2} x^{2}\right]-2 \alpha\right] e^{-\alpha x^{2}}=\frac{2 m}{\hbar^{2}}\left[\frac{1}{2} m \omega^{2} x^{2}\right]-E\right] C_{0} e^{-\alpha x^{2}}$
Match the coeff of $x^{2}$ and the Constant terms on LHS \& RHS
$\Rightarrow 4 \alpha^{2}=\frac{2 m}{\hbar^{2}} \frac{1}{2} m \omega^{2}$ or $\alpha=\frac{\mathrm{m} \omega}{2 \hbar}$
\& the other match gives $2 \alpha=\frac{2 m}{\hbar^{2}} E$, substituing $\alpha \Rightarrow$

$$
\mathrm{E}=\frac{1}{2} \hbar \omega=\mathrm{hf} \quad!!!!. . . . .(\text { Planck's Oscillators) }
$$

What about $C_{0}$ ? We learn about that from the Normalization cond.

## SHO: Normalization Condition

$\int_{-\infty}^{+\infty}\left|\psi_{0}(x)\right|^{2} d x=1=\int_{-\infty}^{+\infty} C_{0}^{2} e^{\frac{-m \omega x^{2}}{\hbar}} d x$
Since $\int_{-\infty}^{+\infty} e^{-a x^{2}} d x=\sqrt{\frac{\pi}{a}}$
(dont memorize this)
Identifying $\mathrm{a}=\frac{m \omega}{\hbar}$ and using the identity above
$\Rightarrow \quad C_{0}=\left[\frac{m \omega}{\pi \hbar}\right]^{\frac{1}{4}}$
Hence the Complete NORMALIZED wave function is :

$$
\psi_{0}(\mathrm{x})=\left[\frac{m \omega}{\pi \hbar}\right]^{\frac{1}{4}} e^{\frac{-m \omega \alpha^{2}}{2 \hbar}} \text { Ground State Wavefunction }
$$

has energy $\mathrm{E}=\mathrm{hf}$

Planck's Oscillators were electrons tied by the "spring" of the mutually attractive Coulomb Force

## Quantum Oscillator In Pictures

$E=K E+U(x)>0$ for $\mathrm{n}=0$
Quantum Mechanical prob for particle To live outside classical turning points


Classically particle most likely to be at the turning point (velocity $=0$ ) Quantum Mechanically, particle most likely to be at $\mathrm{x}=\mathrm{x}_{0}$ for $\mathrm{n}=0$

## Classical \& Quantum Pictures of SHO compared

- Limits of classical vibration : Turning Points (do on Board)
- Quantum Probability for particle outside classical turning points $\mathrm{P}(|\mathrm{x}|>\mathrm{A})=16 \%$ !!
- Do it on the board (see Example problems in book)


## Excited States of The Quantum Oscillator

$$
\begin{aligned}
& \psi_{n}(x)=C_{n} H_{n}(x) e^{-\frac{m \omega x^{2}}{2 \hbar}} ; \\
& H_{n}(x)=\text { Hermite Polynomials } \\
& \text { with } \\
& \mathrm{H}_{0}(\mathrm{x})=1 \\
& \mathrm{H}_{1}(\mathrm{x})=2 \mathrm{x} \\
& \mathrm{H}_{2}(\mathrm{x})=4 \mathrm{x}^{2}-2 \\
& \mathrm{H}_{3}(\mathrm{x})=8 \mathrm{x}^{3}-12 x \\
& \mathrm{H}_{\mathrm{n}}(\mathrm{x})=(-1)^{\mathrm{n}} e^{x^{2}} \frac{d^{n} e^{-x^{2}}}{d x^{n}} \\
& \text { and } \\
& E_{n}=\left(n+\frac{1}{2}\right) \hbar \omega=\left(n+\frac{1}{2}\right) h f
\end{aligned}
$$



Again $\mathrm{n}=0,1,2,3 \ldots \infty$ Quantum \#


The Case of a Rusty "Twisted Pair" of Naked Wires \& How Quantum Mechanics Saved ECE Majors!


- Twisted pair of Cu Wire (metâl) in virgin form
- Does not stay that way for long in the atmosphere
-Gets oxidized in dry air quickly $\mathrm{Cu} \rightarrow \mathrm{Cu}_{2} \mathrm{O}$
-In wet air $\mathrm{Cu} \rightarrow \mathrm{Cu}(\mathrm{OH})_{2}$ (the green stuff on wires)
- Oxides or Hydride are non-conducting ..so no current can flow across the junction between two metal wires
- No current means no circuits $\rightarrow$ no EE, no ECE !!
- All ECE majors must now switch to Chemistry instead \& play with benzene !!! Bad news !


Tunneling Through A Potential Barrier

-Classical \& Quantum Pictures compared: When $\mathrm{E}>\mathrm{U}$ \& when $\mathrm{E}<\mathrm{U}$
-Classically, an particle or a beam of particles incident from left encounters barrier:
-when $\mathrm{E}>\mathrm{U} \rightarrow$ Particle just goes over the barrier (gets transmitted )
-When $\mathrm{E}<\mathrm{U} \rightarrow$ particle is stuck in region I, gets entirely reflected, no transmission (T)
-What happens in a Quantum Mechanical barrier? No region is inaccessible for particle since the potential is (sometimes small) but finite

Beam Of Particles With $E<U$ Incident On Barrier From Left


Description Of WaveFunctions in Various regions: Simple Ones first
In Region I : $\quad \Psi_{\mathrm{I}}(x, t)=A e^{i(k x-\omega t)}+B e^{i(-k x-\omega t)}=$ incident + reflected Waves with $\mathrm{E}=\hbar \omega=\frac{\hbar^{2} k^{2}}{2 m}$
define Reflection Coefficient : $\mathrm{R}=\frac{|\mathrm{B}|^{2}}{|\mathrm{~A}|^{2}}=$ frac of incident wave intensity reflected back
In Region III: $\Psi_{\mathrm{III}}(x, t)=F e^{i(k x-\omega t)}+G e^{i(-k x-\omega t)}=$ transmitted
Note : $G e^{i(-h x-\omega t)}$ corresponds to wave incident from right!
This piece does not exist in the scattering picture we are thinking of now $(G=0)$
So $\Psi_{\text {III }}(x, t)=F e^{i(k x-\omega t)}$ represents transmitted beam. Define $T=\frac{|F|^{2}}{|A|^{2}}$
Unitarity Condition $\Rightarrow \mathrm{R}+\mathrm{T}=1$ (particle is either reflected or transmitted)

Wave Function Across The Potential Barrier
In Region II of Potential U
TISE: $-\frac{\hbar^{2}}{2 m} \frac{d^{2} \psi(x)}{d x^{2}}+U \psi(x)=E \psi(x)$
$\Rightarrow \frac{d^{2} \psi(x)}{d x^{2}}=\frac{2 m}{\hbar^{2}}(U-E) \psi(x)$

$$
=\alpha^{2} \psi(x)
$$

with $\alpha^{2}=\frac{\sqrt{2 m(U-E)}}{\hbar} ; \quad U>E \Rightarrow \alpha^{2}>0$
Solutions are of form $\psi(x) \propto e^{ \pm \alpha x}$

$$
\Psi_{I I}(x, t)=C e^{+\alpha x-i \omega t}+D e^{-\alpha x-i \omega t} 0<\mathrm{x}<\mathrm{L}
$$

To determine $\mathrm{C} \& \mathrm{D} \Rightarrow$ apply matching cond. $\Psi_{I I}(x, t)=$ continuous across barrier $(\mathrm{x}=0, \mathrm{~L})$

$\frac{d \Psi_{I I}(x, t)}{d x}=$ continuous across barrier $(\mathrm{x}=0, \mathrm{~L})$


## Continuity Conditions Across Barrier

At $\mathrm{x}=0$, continuity of $\psi(\mathrm{x}) \Rightarrow$
$A+B=C+D \quad$ (1)
At $\mathrm{x}=0$, continuity of $\frac{d \psi(\mathrm{x})}{d x} \Rightarrow$

$$
i k A-i k B=\alpha C-\alpha D
$$

Similarly at $\mathrm{x}=\mathrm{L}$ continuity of $\psi(\mathrm{x}) \Rightarrow$

$$
\begin{equation*}
C e^{-\alpha L}+D e^{+\alpha L}=F e^{i k L} \tag{3}
\end{equation*}
$$

at $\mathrm{x}=\mathrm{L}$, continuity of $\frac{d \psi(\mathrm{x})}{d x} \Rightarrow$

$$
\begin{equation*}
-(\alpha \mathrm{C}) e^{-\alpha L}+(\alpha \mathrm{D}) e^{\alpha L}=i k F e^{i k L} \tag{4}
\end{equation*}
$$

Four equations \& four unknowns
Cant determine A,B,C,D but if you
Divide thruout by A in all 4 equations :
$\Rightarrow$ ratio of amplitudes $\rightarrow$ relations for $\mathrm{R} \& \mathrm{~T}$
That's what we need any way


## Potential Barrier when $E<U$

Expression for Transmission Coeff T=T(E)
Depends on barrier Height U, barrier Width L and particle Energy E

and $\quad R(E)=1-T(E) \ldots \ldots .$. what's not transmitted is reflected


Above equation holds only for $\mathrm{E}<\mathrm{U}$
For $\mathrm{E}>\mathrm{U}, \alpha=$ imaginary $\#$
$\operatorname{Sinh}(\alpha \mathrm{L})$ becomes oscillatory This leads to an Oscillatory T(E) and Transmission resonances occur where For some specific energy ONLY, $\mathrm{T}(\mathrm{E})=1$ At other values of E , some particles are reflected back ..even though $\mathrm{E}>\mathrm{U}$ !! [do the derivation on blackboard] That's the Wave nature of the Quantum particle

## A Special Case That is Instructive \& Useful: U>>E

Given the 4 equations from Continuity Conditions: Solve for $\frac{\mathrm{A}}{\mathrm{F}}$
$\frac{\mathrm{A}}{\mathrm{F}}=\left[\frac{1}{2}+\frac{i}{4}\left(\frac{\alpha}{k}-\frac{k}{\alpha}\right)\right] e^{(i k+\alpha) L}+\left[\frac{1}{2}-\frac{i}{4}\left(\frac{\alpha}{k}-\frac{k}{\alpha}\right)\right] e^{(i k-\alpha) L}$
Remember $\alpha=\frac{\sqrt{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}}{\hbar}, k=\frac{\sqrt{2 \mathrm{mE}}}{\hbar}$,
when $\mathrm{U} \gg \mathrm{E}, \alpha \gg \mathrm{k} \& \frac{\alpha}{k} \gg \frac{k}{\alpha}$ So $\frac{\alpha}{k}-\frac{k}{\alpha} \approx \frac{\alpha}{k}$; For large Barrier $\mathrm{L}, \alpha \mathrm{L} \gg 1$
$\frac{\mathrm{A}}{\mathrm{F}}=\left[\frac{1}{2}+\frac{i}{4}\left(\frac{\alpha}{k}\right)\right] e^{(i k+\alpha) L} ;\left(\frac{\mathrm{A}}{\mathrm{F}}\right)^{*}=\left[\frac{1}{2}-\frac{i}{4}\left(\frac{\alpha}{k}\right)\right] e^{(-i k+\alpha) L}$
$\mathrm{T}^{-1}=\left(\frac{\mathrm{A}}{\mathrm{F}}\right)^{*} \frac{\mathrm{~A}}{\mathrm{~F}}=\left[\frac{1}{4}+\frac{1}{16}\left(\frac{\alpha^{2}}{k^{2}}\right)\right] e^{(2 \alpha) L}=\frac{1}{T(E)} ;$ now invert \& consolidate
$\mathrm{T}=\left|\frac{\mathrm{F}^{*} \mathrm{~F}}{\mathrm{~A}^{*} \mathrm{~A}}\right|=\left(\frac{16}{4+\left(\frac{\alpha}{k}\right)^{2}}\right) e^{-2 \alpha L} ;$ now watch the variables employed

## A Special Case That is Instructive \& Useful: $U \gg E$

$\alpha=\frac{\sqrt{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}}{\hbar}, k=\frac{\sqrt{2 \mathrm{mE}}}{\hbar}=\frac{\mathrm{p}}{\hbar}=\frac{2 \pi}{\lambda}$
$\left(\frac{\alpha}{k}\right)^{2}=\frac{2 m(U-E) / \hbar^{2}}{2 m E / \hbar^{2}}=\frac{U}{E}-1 \simeq \frac{U}{E}$
$\Rightarrow\left(\frac{16}{4+\left(\frac{\alpha}{k}\right)^{2}}\right)=\left(\frac{16}{4+\left(\frac{U}{E}-1\right)}\right) \rightarrow$ varies slowly compared with $\mathrm{e}^{-2 \alpha \mathrm{~L}}$ term
keeing in mind only the Order of magnitude, I suggest
$\left(\frac{16}{4+\left(\frac{U}{E}-1\right)}\right) \approx 1$; back to $\mathrm{T}=\left(\frac{16}{4+\left(\frac{\alpha}{k}\right)^{2}}\right) e^{-2 \alpha L}$ substituting
So approximately
$T \approx \mathrm{e}^{-2 \alpha \mathrm{~L}}$ Transmission Prob is fn of U,E,L

Why subject you to this TORTURE? $\rightarrow$ Estimate T for complicated Potentials
See next (example of Cu oxide layer, radioactivity, blackhole blowup etc)


Story involves an Allied plan for a massive breakout from a Nazi P.O.W. camp,
during World War Two. The Nazis had created a high-security, escape -proof prisoner of war camp for those annoying detainees who have attempted escape from their other prison of war camps. These prisoners are not discouraged at all, as they plan a huge escape of 100 men.

## Ceparated in Coppertino



Q: 2 Cu wires are seperated by insulating Oxide layer. Modeling the Oxide layer as a square barrier of height $\mathrm{U}=10.0 \mathrm{eV}$, estimate the transmission coeff for an incident beam of electrons of $E=7.0 \mathrm{eV}$ when the layer thickness is
(a) 5.0 nm (b) 1.0 nm

Q: If a 1.0 mA current in one of the intwined wires is incident on Oxide layer, how much of this current passes thru the Oxide layer on to the adjacent wire if the layer thickness is 1.0 nm ? What becomes of the remaining current?



## A Complicated Potential Barrier Can Be Broken Down

| Multiplicative Transmission prob, ever decreasing but not 0 |
| :--- | :--- |

$\mathrm{U}(\mathrm{x})$


Can be broken down into a sum of successive Rectangular potential barriers for which we learnt to find the Transmission probability $\mathrm{T}_{\mathrm{i}}$
The Transmitted beam intensity thru one small barrier becomes incident beam intensity for the following one So on \& so forth ...till the particle escapes with final Transmission prob T

$$
\left.T=\int T_{i} d x=e^{-2\left[\frac{\sqrt{2 m}}{\hbar} \int \sqrt{U(x)-E} d x\right.}\right]
$$

Radioactivity: The $\alpha$-particle \& Steve McQueen Compared


- In a Nucleus such as Ra, Uranium etc $\alpha$ particle rattles around parent nucleus, "hitting"the nuclear walls with a very high frequency (probing the "fence"), if the Transmission prob $\mathrm{T}>0$, then eventually particle escapes
- Within nucleus, $\alpha$ particle is virtually free but is trapped by the Strong nuclear force
- Once outside nucleus, the particle "sees" only the repulsive ( + ) columbic force (nuclear force too faint outside) which keeps it within nucleus
- Nuclear radius $\mathrm{R}=10^{-14} \mathrm{~m}, \mathrm{E}_{\alpha}=9 \mathrm{MeV}$
- Coulomb barrier $\mathrm{U}(\mathrm{r})=\mathrm{kq}_{1} \mathrm{q}_{2} / \mathrm{r}$ - At $\mathrm{r}=\mathrm{R}, \mathrm{U}(\mathrm{R}) \approx 30 \mathrm{MeV}$ barrier
- $\alpha$-particle, due to QM , tunnels thru
- It's the sensitivity of T on $\mathrm{E}_{\alpha}$ that accounts for the wide range in half-lives of radioactive nuclei


## Radioactivity Explained Roughly (..is enough!)

- Protons and neutrons rattling freely inside radioactive nucleus $\left(\mathrm{R} \simeq 10^{-15} \mathrm{~m}\right)$
- Constantly morphing into clusters of protons and neutrons
- Proto-alpha particle $=(2 p+2 n)$ of $\approx 9 \mathrm{MeV}$ prevented from getting out by the imposing Coulombic repulsion of remaining charge ( $\approx 30 \mathrm{MeV}$ )
- Escapes by tunneling thru Coloumb potential...but some puzzling features:
- a particles emitted from all types of radioactive nuclei have roughly same KE $\cong 4-9 \mathrm{MeV}$

- In contrast, the half live $\mathrm{T}\left(\mathrm{N} \rightarrow \mathrm{e}^{-1} \mathrm{~N}\right)$ differ by more than 20 orders of magnitude !

| Element | KE of emitted $\alpha$ | Half Life |
| :--- | :--- | :--- |
| ${ }^{212} \mathrm{Po}$ | 8.95 MeV | $3 \times 10^{-7} \mathrm{~s}$ |
| ${ }^{240} \mathrm{Cm}$ | 6.40 MeV | 27 days |
| ${ }^{226} \mathrm{Ra}$ | 4.90 MeV | $1.60 \times 10^{3} \mathrm{Yr}$ |
| ${ }^{232} \mathrm{Th}$ | 4.05 MeV | $1.41 \times 10^{10} \mathrm{yr}$ |



## Radioactivity Explained Crudely

$$
\begin{aligned}
& T(E) \approx e^{\left[\frac{-2}{\hbar} \sqrt{2 m} \int \sqrt{U(x)-E} d x\right]}, U(x)=\frac{2 e^{2} Z}{4 \pi \varepsilon_{0} r} \\
& \ln T=\frac{-2}{\hbar} \int_{0}^{\mathrm{b}} \sqrt{2 m_{\alpha}\left(\frac{2 e^{2} Z}{4 \pi \varepsilon_{0} r}-E_{\alpha}\right)} d r,
\end{aligned}
$$

limits of integration correspond to values of $r$ when $E=U$
$\Rightarrow \frac{2 e^{2} Z}{4 \pi \varepsilon_{0} b}=E \Rightarrow b=\frac{2 e^{2} Z}{4 \pi \varepsilon_{0} E_{\alpha}}$
Define $\xi=\frac{\mathrm{r}}{\mathrm{b}} ;=\frac{r}{2 e^{2} Z / 4 \pi \varepsilon_{0} E} \Rightarrow \ln T \cong \frac{-2\left(\sqrt{2 m_{\alpha} E}\right) b}{\hbar} \int_{0}^{1} \sqrt{\frac{1}{\xi}-1} d \xi$
Substitute $\xi=\sin ^{2} \theta$ in integration, change limits $\Rightarrow$
$\ln T \cong \frac{-4\left(\sqrt{2 m_{\alpha} E_{\alpha}}\right) b}{\hbar} \int_{0}^{\pi / 2} \cos ^{2} \theta d \theta ;$ use $\int_{0}^{\pi / 2} \cos ^{2} \theta d \theta=\pi / 4 \& E_{\alpha}=\frac{m_{\alpha} V_{\alpha}^{2}}{2}$
$\ln T \cong \frac{-2 \pi}{\hbar} \frac{Z e^{2}}{4 \pi \varepsilon_{0}} \sqrt{\frac{2 m_{\alpha}}{E_{\alpha}}}=\frac{-4 \pi}{\hbar} \frac{Z e^{2}}{4 \pi \varepsilon_{0}} \frac{1}{V_{\alpha}} \Rightarrow T \cong e^{\left[\frac{-4 \pi Z e^{2}}{\hbar 4 \pi \varepsilon_{0} V_{\alpha}}\right]}$
$T \propto e^{\frac{-1}{V_{\alpha}}}$ and $T \propto e^{-z}$

## Radioactivity

A more eloberate calculation (Bohm) yields

where $\mathrm{r}_{0}=\frac{\hbar^{2}}{m_{\alpha} k e^{2}} \simeq 8 \mathrm{fm}$ is the "Bohr Radius" of alpha particles
and $\mathrm{E}_{0}=\frac{k e^{2}}{2 r_{0}}=0.0993 \mathrm{MeV}=$ Nuclear "Rydberg"
To obtain decay rates, need to multiply $\mathrm{T}(\mathrm{E})$ by the number of collisions
$\alpha$ particle makes with the "walls" of the nuclear barrier. This collission frequency
$\mathrm{f}=\frac{\mathrm{V}_{\alpha}}{2 \mathrm{R}}=$ transit time for $\alpha$ particle crossing the nuclear barrier (rattle time)
Typically $\mathrm{f}=10^{21}$ collissions/second

Decay rate (prob. of $\alpha$ emission per unit time) $\lambda=\mathrm{f} \bullet \mathrm{T}(\mathrm{E})$
$\lambda=10^{21} e^{\left\{-4 \pi z \sqrt{\frac{E_{0}}{E_{\alpha}}}+8 \sqrt{\frac{Z Z}{r_{0}}}\right\}}$
Definition : Half life $\mathrm{t}_{1 / 2}=\frac{\ln 2}{\lambda}$

## Half Lives Compared: Sharp dependence on $E_{\alpha}$

$\alpha$ particles emerge with (a) $\mathrm{E}=4.05 \mathrm{MeV}$ in Thorium (b) $\mathrm{E}=8.95 \mathrm{MeV}$ in Polonium. The Nuclear size $\mathrm{R}=9 \mathrm{fm}$ in both cases. Which one will outlive you?

Thorium ( $\mathrm{Z}=90$ ) decays into Radium ( $\mathrm{Z}=88$ )
$\mathrm{T}(\mathrm{E})=\exp \{-4 \pi(\sqrt{88}) \sqrt{(0.0993 / 4.05)}+8 \sqrt{88}(9.00 / 7.25)\}$

$$
=1.3 \times 10^{-39}
$$

Taking $\mathrm{f}=10^{21} \mathrm{~Hz} \Rightarrow \lambda=1.3 \times 10^{-18} \alpha$ emission $\Rightarrow \mathrm{t}_{1 / 2}=\frac{0.693}{1.3 \times 10^{-18}}=1.7 \times 10^{10} y r!!$ !
Polonium ( $Z=84$ ) decays into Lead $(Z=82)$
$\mathrm{T}(\mathrm{E})=\exp \{-4 \pi(\underline{82}) \sqrt{(0.0993 / 8.95)}+8 \sqrt{82(9.00 / 7.25)}\}$

$$
=8.2 \times 10^{-13}
$$

Taking $\mathrm{f}=10^{21} \mathrm{~Hz} \Rightarrow \lambda=8.2 \times 10^{-8} \alpha$ emission $\Rightarrow \mathrm{t}_{1 / 2}=\frac{0.693}{8.2 \times 10^{-8}}=8.4 \times 10^{-10} \mathrm{~s}!!!$ !

## Potential Barrier : An Unintuitive Result When $E>U$



Description Of WaveFunctions in Various regions: Simple Ones first
In Region I : $\Psi_{\mathrm{I}}(x, t)=A e^{i(k x-\omega t)}+B e^{i(-k x-\omega t)}$; In Region III: $\Psi_{\text {III }}(x, t)=F e^{i(k x-\omega t)}$ In Region II of Potential U :
TISE: $-\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{d^{2} \psi(x)}{d x^{2}}+U \psi(x)=E \psi(x) \Rightarrow \frac{d^{2} \psi(x)}{d x^{2}}=\frac{2 m}{\hbar^{2}}(U-E) \psi(x)=\alpha^{2} \psi(x)$
with $\alpha^{2}=\frac{\sqrt{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}}{\hbar} ; \quad \mathrm{U}<\mathrm{E} \Rightarrow \alpha^{2}<0$
Define $\alpha=i k^{\prime} ; \alpha^{2}=-\left(k^{\prime}\right)^{2} ; k^{\prime}=\sqrt{\frac{2 m(E-U)}{\hbar^{2}}}$
$\Rightarrow \Psi_{\text {II }}=C e^{i\left(-k^{\prime} x-\omega t\right)}+D e^{i\left(k^{\prime} x-\omega t\right)} \Rightarrow$ Oscillatory W avefunction
Apply continuity condition at $x=0 \& x=L$
$\mathrm{A}+\mathrm{B}=\mathrm{C}+\mathrm{D} ; \quad k A-k B=k^{\prime} D-k^{\prime} C ; \quad C e^{-i k^{\prime} L}+D e^{i k^{\prime} L}=F e^{i k L} ; k^{\prime} D e^{i k^{\prime} L}-k^{\prime} C e^{-i k^{\prime} L}=k F e^{i k L}$
Eliminate $\mathrm{B}, \mathrm{C}, \mathrm{D}$ and write every thing in terms of A and $\mathrm{F} \Rightarrow$
$\mathrm{A}=\frac{1}{4} F e^{i k L}\left\{\left[2-\left(\frac{k^{\prime}}{k}+\frac{k}{k^{\prime}}\right)\right] e^{i k^{\prime} L}+\left[2+\left(\frac{k^{\prime}}{k}+\frac{k}{k^{\prime}}\right)\right] e^{-i k^{\prime} L}\right\}$

## Potential Barrier : An Unintuitive Result When $E>U$



Only when $\sin k^{\prime} L=0, T=1$; this happens when $k^{\prime} L=n \pi$
Since $k^{\prime}=\sqrt{\frac{2 m(E-U)}{\hbar^{2}}} \Rightarrow \sqrt{\frac{2 m(E-U)}{\hbar^{2}}}=n \pi$
$\Rightarrow E_{n}=U+n^{2}\left(\frac{\pi^{2} \hbar^{2}}{2 m L^{2}}\right)$ is the condition
for particle to be completely transmitted

For all other energies, $\mathrm{T}<1$ and $\mathrm{R}>0$ !!!
This is Quantum Mechanics in your face


## Special Case: A Potential Step



## Transmission Probability in A Potential Step

$$
\mathrm{U}=0 \quad \text { for } \mathrm{x}<0
$$

$$
\mathrm{U}=\mathrm{U} \quad \text { for } \mathrm{x} \geq 0
$$

$$
\Psi_{I I}=C e^{-\alpha x-i \omega t}
$$

$$
\text { Since } \Psi_{I}(x, t)=A e^{i k x-i \omega t}+B e^{-i k x-i \omega t} ; \Psi_{\mathrm{II}}(x, t)=C e^{-\alpha x-i \omega t}
$$

Applying Continuity conditions of $\Psi$ and $\frac{d \Psi}{d x}$ at $\mathrm{x}=0$ :
$\mathrm{A}+\mathrm{B}=\mathrm{C} \Rightarrow \frac{\mathrm{C}}{\mathrm{A}}=1+\frac{\mathrm{B}}{\mathrm{A}}=1-\frac{(1+\mathrm{ik} \delta)}{(1-\mathrm{ik} \delta)}$
$\Rightarrow \frac{\mathrm{C}}{\mathrm{A}}=-\frac{2 i k \delta}{1-i k \delta} \neq 0 \Rightarrow T=\left(\frac{\mathrm{C}}{\mathrm{A}}\right)\left(\frac{\mathrm{C}}{\mathrm{A}}\right)^{*}>0!!!$
The particle burrows into the skin of the step barrier. If one has a barrier of width $\mathrm{L}=\delta$, particle escapes thru the barrier.
penetration distance $\Delta x=$ distance for which prob. drops by $1 / \mathrm{e}$.
$|\psi(\mathrm{x}=\Delta \mathrm{x})|^{2}=\mathrm{C}^{2} \mathrm{e}^{-2 \alpha \Delta \mathrm{x}}=\mathrm{C}^{2} \mathrm{e}^{-1}$; happens when $2 \alpha \Delta \mathrm{x}=1$ or $\Delta \mathrm{x}=\frac{1}{2} \frac{\hbar}{\sqrt{2 \mathrm{~m}(\mathrm{U}-\mathrm{E})}}$

## Particle Beams and Flux Conservation



If we write the particle wavefunction for incident as $\psi_{\mathrm{I}^{+}}=A e^{i k_{I} x}$ and reflected as $\psi_{\mathrm{I}-}=B e^{-i k_{I} x}$
The particle flux arriving at the barrier, defined as number of particles per unit length per unit time
$\mathrm{S}_{\mathrm{I}^{+}}=\left|\psi_{\mathrm{I}^{+}}^{*} \psi_{\mathrm{I}^{+}}\right| V_{I+}=\left|\psi_{\mathrm{I}^{+}} \psi^{*} \psi_{\mathrm{I}+}\right|\left(\frac{p}{m}\right)$ and $\mathrm{S}_{\mathrm{I}-}=\left|\psi_{\mathrm{I}^{-}}{ }^{*} \psi_{\mathrm{I}}\right|-\left(\frac{p}{m}\right) ; \quad$ (for non-relativistic case)
Since the wavefunction in region III $\psi_{\mathrm{III}+}=\psi_{\mathrm{III}}=F e^{i k_{I I} x}$ and $\mathrm{S}_{\mathrm{III}+}=\left|\psi_{\mathrm{III}+}^{*} \psi_{\mathrm{III}+}\right|\left(\frac{p_{I I I}}{m}\right)$
The general expression for flux probabilities : number of particles passing by any point per unit time:
Transmission Probability $\mathrm{T}=\frac{\left|\psi_{\mathrm{III}+}^{*} \psi_{\mathrm{III+}}\right| V_{I I++}}{\left|\psi_{\mathrm{I}+} \psi_{\mathrm{I}+}\right| V_{I+}}=\left(\frac{F}{A}\right)^{*}\left(\frac{F}{A}\right)\left(\frac{V_{I I I+}}{V_{I+}}\right)$
Reflection Probability $\mathrm{R}=\frac{\left|\psi_{\mathrm{I}-}{ }^{*} \psi_{\mathrm{I}-}\right| V_{I-}}{\left|\psi_{\mathrm{I}+} \psi_{\mathrm{I}+}\right| V_{I+}}=\left(\frac{B}{A}\right)^{*}\left(\frac{B}{A}\right)\left(\frac{V_{I-}}{V_{I+}}\right)$

The general expression for conservation of particle flux remains: $1=\mathrm{T}+\mathrm{R}$

## Where does this generalization become important?

$\longrightarrow$
(a) Energy

- Particle with energy E incident from left on a potential step U, with $\mathrm{E}>\mathrm{U}$
- Particle momentum, wavelength and velocities are different in region I and II
- Is the reflection probability $=0$ ?

Where does this generalization become important?
■


## QM in 3 Dimensions

- Learn to extend S. Eq and its solutions from "toy" examples in 1-Dimension (x) $\rightarrow$ three orthogonal dimensions
( $\mathrm{r} \equiv \mathrm{x}, \mathrm{y}, \mathrm{z}$ )

$$
\vec{r}=\hat{i} x+\hat{j} y+\hat{k} z
$$

- Then transform the systems
- Particle in 1D rigid box $\rightarrow 3 \mathrm{D}$ rigid box
- 1D Harmonic Oscillator $\rightarrow$ 3D

Harmonic Oscillator

- Keep an eye on the number of different integers needed to specify system $1 \rightarrow 3$ (corresponding to 3 available degrees of freedom $\mathrm{x}, \mathrm{y}, \mathrm{z}$ )


The Schrodinger Equation in 3 Dimensions: Cartesian Coordinates
Time Dependent Schrodinger Eqn:

$\nabla^{2}=\frac{\partial^{2}}{\partial x^{2}}+\frac{\partial^{2}}{\partial y^{2}}+\frac{\partial^{2}}{\partial z^{2}}$
So $-\frac{\hbar^{2}}{2 m} \nabla^{2}=\left(-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial x^{2}}\right)+\left(-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial y^{2}}\right)+\left(-\frac{\hbar^{2}}{2 m} \frac{\partial^{2}}{\partial z^{2}}\right)=[K]$
$=\left[\mathrm{K}_{\mathrm{x}}\right]$
$\left[\mathrm{K}_{\mathrm{x}}\right]+\left[\mathrm{K}_{\mathrm{x}}\right]$
so $[H] \Psi(x, t)=[E] \Psi(x, t)$ is still the Energy Conservation Eq

Stationary states are those for which all probabilities are constant in time and are given by the solution of the TDSE in seperable form:

$$
\Psi(x, y, z, t)=\Psi(\vec{r}, t)=\psi(\overrightarrow{\mathrm{r}}) \mathrm{e}^{-i \omega t}
$$

This statement is simply an extension of what we derived in case of 1D time-independent potential

Particle in 3D Rigid Box : Separation of Orthogonal Spatial (x,y,z) Variables
TISE in 3D: $-\frac{\hbar^{2}}{2 \mathrm{~m}} \nabla^{2} \psi(x, y, z)+U(x, y, z) \psi(x, y, z)=E \psi(x, y, z)$
$\mathrm{x}, \mathrm{y}, \mathrm{z}$ independent of each other, write $\psi(x, y, z)=\psi_{1}(x) \psi_{2}(y) \psi_{3}(z)$
and substitute in the master TISE, after dividing thruout by $\psi=\psi_{1}(x) \psi_{2}(y) \psi_{3}(z)$
and noting that $U(r)=0$ for $(0<x, y, z,<L) \Rightarrow$
$\left(-\frac{\hbar^{2}}{2 m} \frac{1}{\psi_{1}(x)} \frac{\partial^{2} \psi_{1}(x)}{\partial x^{2}}\right)+\left(-\frac{\hbar^{2}}{2 m} \frac{1}{\psi_{2}(y)} \frac{\partial^{2} \psi_{2}(y)}{\partial y^{2}}\right)+\left(-\frac{\hbar^{2}}{2 m} \frac{1}{\psi_{3}(z)} \frac{\partial^{2} \psi_{3}(z)}{\partial z^{2}}\right)=E=$ Const
This can only be true if each term is constant for all $x, y, z \Rightarrow$
$-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \psi_{1}(x)}{\partial x^{2}}=E_{1} \psi_{1}(x) ;-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \psi_{2}(y)}{\partial y^{2}}=E_{2} \psi_{2}(y) ;-\frac{\hbar^{2}}{2 m} \frac{\partial^{2} \psi_{3}(z)}{\partial z^{2}}=E_{3} \psi_{3}(z)$
With $E_{1}+E_{2}+E_{3}=E=$ Constant (Total Energy of 3 D system)
Each term looks like particle in 1D box (just a different dimension)
So wavefunctions must be like $\psi_{1}(x) \propto \sin k_{1} \mathrm{x}, \psi_{2}(y) \propto \sin k_{2} \mathrm{y}, \psi_{3}(z) \propto \sin k_{3} z$

## Particle in 3D Rigid Box : Separation of Orthogonal Variables

Wavefunctions are like $\psi_{1}(x) \propto \sin k_{1} x, \psi_{2}(y) \propto \sin k_{2} y, \psi_{3}(z) \propto \sin k_{3} z$ Continuity Conditions for $\psi_{\mathrm{i}}$ and its first spatial derivatives $\Rightarrow n_{i} \pi=k_{i} L$
Leads to usual Quantization of Linear Momentum $\overrightarrow{\mathrm{p}}=\hbar \overrightarrow{\mathrm{k}}$.....in 3D
$p_{x}=\left(\frac{\pi \hbar}{L}\right) n_{1} ; p_{y}=\left(\frac{\pi \hbar}{L}\right) n_{2} ; p_{z}=\left(\frac{\pi \hbar}{L}\right) n_{3}\left(\mathrm{n}_{1}, \mathrm{n}_{2}, \mathrm{n}_{3}=1,2,3, . . \infty\right)$
Note: by usual Uncertainty Principle argument neither of $\mathrm{n}_{1}, \mathrm{n}_{2}, \mathrm{n}_{3}=0$ ! (why?)
Particle Energy $\mathrm{E}=\mathrm{K}+\mathrm{U}=\mathrm{K}+0=\frac{1}{2 \mathrm{~m}}\left(p_{x}^{2}+p_{y}^{2}+p_{z}^{2}\right)=\frac{\pi^{2} \hbar^{2}}{2 m L^{2}}\left(n_{1}^{2}+n_{2}^{2}+n_{3}^{2}\right)$
Energy is again quantized and brought to you by integers $\mathrm{n}_{1}, \mathrm{n}_{2}, \mathrm{n}_{3}$ (independent) and $\psi(\overrightarrow{\mathrm{r}})=\mathrm{A} \sin k_{1} \mathrm{x} \sin k_{2} \mathrm{y} \sin k_{3} z \quad(\mathrm{~A}=\mathrm{Overall}$ Normalization Constant $)$
$\Psi(\overrightarrow{\mathrm{r}}, \mathrm{t})=\psi(\overrightarrow{\mathrm{r}}) \mathrm{e}^{-\mathrm{i} \frac{\mathrm{E}}{\hbar} t}=A\left[\sin k_{1} \mathrm{x} \sin k_{2} \mathrm{y} \sin k_{3} \mathrm{z}\right] \mathrm{e}^{-\mathrm{i} \frac{\mathrm{E}}{\hbar} t}$

## Particle in 3D Box :Wave function Normalization Condition

$\Psi(\overrightarrow{\mathrm{r}}, \mathrm{t})=\psi(\mathrm{r}) \mathrm{e}^{-\mathrm{E} \mathrm{E} t}{ }^{\frac{\mathrm{E}}{}}=A\left[\sin k_{1} \mathrm{x} \sin k_{2} \mathrm{y} \sin k_{3} z\right] \mathrm{e}^{-\frac{\mathrm{E}}{} \mathrm{E}^{\mathrm{E}} t}$
$\Psi^{*}(\overrightarrow{\mathrm{r}}, \mathrm{t})=\psi^{*}(\overrightarrow{\mathrm{r}}) \mathrm{e}^{\mathrm{i} \frac{\mathrm{E}}{\hbar} t}=A\left[\sin k_{1} \mathrm{x} \sin k_{2} \mathrm{y} \sin k_{3} \mathrm{z}\right] \mathrm{e}^{\mathrm{i} \frac{\mathrm{E}}{\hbar} t}$
$\Psi^{*}(\overrightarrow{\mathrm{r}}, \mathrm{t}) \Psi(\overrightarrow{\mathrm{r}}, \mathrm{t})=A^{2}\left[\sin ^{2} k_{1} \mathrm{x} \sin ^{2} k_{2} \mathrm{y} \sin ^{2} k_{3} z\right]$
Normalization Condition : $1=\iiint_{\mathrm{x}, \mathrm{y}, \mathrm{z}} \mathrm{P}(\mathrm{r}) \mathrm{dx} \mathrm{dydz} \Rightarrow$ $1=A^{2} \int_{x=0}^{\mathrm{L}} \sin ^{2} k_{1} \mathrm{x} d \mathrm{x} \int_{\mathrm{y}=0}^{\mathrm{L}} \sin ^{2} k_{2} \mathrm{y} d \mathrm{y} \int_{\mathrm{z}=0}^{\mathrm{L}} \sin ^{2} k_{3} \mathrm{z} d \mathrm{~d}=A^{2}\left(\sqrt{\frac{L}{2}}\right)\left(\sqrt{\frac{L}{2}}\right)\left(\sqrt{\frac{L}{2}}\right)$
$\Rightarrow A=\left[\frac{2}{L}\right]^{\frac{3}{2}}$ and $\Psi(\overrightarrow{\mathrm{r}}, \mathrm{t})=\left[\frac{2}{L}\right]^{\frac{3}{2}}\left[\sin k_{1} \mathrm{x} \sin k_{2} \mathrm{y} \sin k_{3} \mathrm{z}\right] \mathrm{e}^{-\mathrm{i} \frac{\mathrm{E}}{\hbar} t}$

## Particle in 3D Box : Energy Spectrum \& Degeneracy

$\mathrm{E}_{\mathrm{n}_{1}, \mathrm{n}_{2}, \mathrm{n}_{3}}=\frac{\pi^{2} \hbar^{2}}{2 m L^{2}}\left(n_{1}^{2}+n_{2}^{2}+n_{3}^{2}\right) ; \mathrm{n}_{\mathrm{i}}=1,2,3 \ldots \infty, n_{i} \neq 0$
Ground State Energy $\mathrm{E}_{111}=\frac{3 \pi^{2} \hbar^{2}}{2 m L^{2}}$
Next level $\Rightarrow 3$ Excited states $\mathrm{E}_{211}=\mathrm{E}_{121}=\mathrm{E}_{112}=\frac{6 \pi^{2} \hbar^{2}}{2 m L^{2}}$
Different configurations of $\psi(\mathrm{r})=\psi(\mathrm{x}, \mathrm{y}, \mathrm{z})$ have same energy $\Rightarrow$ degeneracy



Probability Density Functions for Particle in 3D Box


Source of Degeneracy: How to "Lift" Degeneracy

- Degeneracy came from the threefold symmetry of a
CUBICAL Box $\left(\mathrm{L}_{\mathrm{x}}=\mathrm{L}_{\mathrm{y}}=\mathrm{L}_{\mathrm{z}}=\mathrm{L}\right)$
- To Lift (remove) degeneracy $\rightarrow$ change each dimension such that CUBICAL box $\rightarrow$ Rectangular Box
- $\left(\mathrm{L}_{\mathrm{x}} \neq \mathrm{L}_{\mathrm{y}} \neq \mathrm{L}_{\mathrm{z}}\right)$
- Then
$E=\left(\frac{n_{1}^{2} \pi^{2}}{2 m L_{x}^{2}}\right)+\left(\frac{n_{2}^{2} \pi^{2}}{2 m L_{y}^{2}}\right)+\left(\frac{n_{3}^{2} \pi^{2}}{2 m L_{z}^{2}}\right)$


## Did 0 포

$$
L_{1}<L_{2}<L_{3}
$$

$$
=E_{221}
$$

$E_{122}$
$\qquad$ $E_{211}$ $E_{121}$

$$
\square
$$

$E_{112}$

$$
112
$$

## The Coulomb Attractive Potential That Binds the electron and Nucleus (charge +Ze) into a Hydrogenic atom



The Hydrogen Atom In Its Full Quantum Mechanical Glory

$U(r) \propto \frac{1}{r}=\frac{1}{\sqrt{x^{2}+y^{2}+z^{2}}} \Rightarrow \mathrm{x}, \mathrm{y}, \mathrm{z}$ all mixed up !
As in case of particle in 3D box, we should use seperation of variables ( $x, y, z$ ??) to derive 3 independent differential. eqns.
This approach will get very ugly since we have a "conjoined triplet"
To simplify the situation, choose more appropriate variables Cartesian coordinates $(\mathrm{x}, \mathrm{y}, \mathrm{z}) \rightarrow$ Spherical Polar $(\mathrm{r}, \theta, \phi)$ coordinates


$$
\begin{aligned}
& x=r \sin \theta \cos \phi \\
& y=r \sin \theta \sin \phi \\
& z=r \cos \theta \\
& r=\sqrt{x^{2}+y^{2}+z^{2}} \\
& \theta=\cos ^{-1} \frac{z}{r} \text { (Polar angle) } \\
& \phi=\tan ^{-1} \frac{y}{x} \text { (Azimuthal angle) }
\end{aligned}
$$



The Hydrogen Atom In Its Full Quantum Mechanical Glory


The Schrodinger Equation in Spherical Polar Coordinates (is bit of a mess!)

The TISE is :
$\frac{1}{\mathrm{r}^{2}} \frac{\partial}{\partial r}\left(r^{2} \frac{\partial \psi}{\partial r}\right)+\frac{1}{r^{2} \sin \theta} \frac{\partial}{\partial \theta}\left(\sin \theta \frac{\partial \psi}{\partial \theta}\right)+\frac{1}{r^{2} \sin ^{2} \theta} \frac{\partial^{2} \psi}{\partial^{2} \phi}+\frac{2 \mathrm{~m}}{\hbar^{2}}(\mathrm{E}-\mathrm{U}(\mathrm{r})) \psi(\mathrm{r}, \theta, \phi)=0$

Try to free up second last term from all except $\phi$

This requires multiplying thruout by $r^{2} \sin ^{2} \theta \Rightarrow$
$\sin ^{2} \theta \frac{\partial}{\partial r}\left(r^{2} \frac{\partial \psi}{\partial r}\right)+\sin \theta \frac{\partial}{\partial \theta}\left(\sin \theta \frac{\partial \psi}{\partial \theta}\right)+\frac{\partial^{2} \psi}{\partial^{2} \phi}+\frac{2 \mathrm{mr}^{2} \sin ^{2} \theta}{\hbar^{2}}\left(\mathrm{E}+\frac{\mathrm{ke}^{2}}{\mathrm{r}}\right) \psi=0$

For Seperation of Variables, Write $\psi(\mathrm{r}, \theta, \phi)=\mathrm{R}(\mathrm{r}) . \Theta(\theta) . \Phi(\phi)$
Plug it into the TISE above \& divide thruout by $\psi(\mathrm{r}, \theta, \phi)=\mathrm{R}(\mathrm{r}) \cdot \Theta(\theta) . \Phi(\phi)$

Note that

$$
\begin{aligned}
& \frac{\partial \Psi(r, \theta, \phi)}{\partial \mathrm{r}}=\Theta(\theta) \cdot \Phi(\phi) \frac{\partial \mathrm{R}(\mathrm{r})}{\partial \mathrm{r}} \\
& \frac{\partial \Psi(r, \theta, \phi)}{\partial \theta}=R(r) \Phi(\phi) \frac{\partial \Theta(\theta)}{\partial \theta} \\
& \frac{\partial \Psi(r, \theta, \phi)}{\partial \theta}=R(r) \Theta(\theta) \frac{\partial \Phi(\phi)}{\partial \phi}
\end{aligned} \Rightarrow \text { when substituted in TISE }
$$

## Solving For the Hydrogen Atom: Separation of Variables

LHS is fn. of $r, \theta$ \& RHS is fn of $\phi$ only, for equality to be true for all $r, \theta, \phi$

$$
\Rightarrow \text { LHS }=\text { constant }=\mathrm{RHS}=\mathrm{m}_{1}^{2}
$$

## Deconstructing The Schrodinger Equation for Hydrogen

Now go break up LHS to seperate the $\mathrm{r} \& \theta$ terms.....
LHS: $\frac{\sin ^{2} \theta}{R} \frac{\partial}{\partial r}\left(r^{2} \frac{\partial R}{\partial r}\right)+\frac{\sin \theta}{\Theta} \frac{\partial}{\partial \theta}\left(\sin \theta \frac{\partial \Theta}{\partial \theta}\right)+\frac{2 \mathrm{~m} r^{2} \sin ^{2} \theta}{\hbar^{2}}\left(\mathrm{E}+\frac{\mathrm{ke}}{\mathrm{r}}\right)=\mathrm{m}_{l}^{2}$

Divide Thruout by $\sin ^{2} \theta$ and arrange all terms with r away from $\theta \Rightarrow$
$\frac{1}{R} \frac{\partial}{\partial r}\left(r^{2} \frac{\partial R}{\partial r}\right)+\frac{2 m r^{2}}{\hbar^{2}}\left(\mathrm{E}+\frac{\mathrm{ke}^{2}}{\mathrm{r}}\right)=\frac{\mathrm{m}_{l}^{2}}{\sin ^{2} \theta}-\frac{1}{\Theta \sin \theta} \frac{\partial}{\partial \theta}\left(\sin \theta \frac{\partial \Theta}{\partial \theta}\right)$

Same argument : LHS is fn of r, RHS is fn of $\theta$;

For them to be equal for all $\mathrm{r}, \theta \Rightarrow$ LHS $=$ const $=$ RHS $=l(l+1)$

What is the mysterious $l(l+1)$ ? Just a number like $2(2+1)$

So What do we have after all the shuffling!

$$
\begin{align*}
& \frac{\mathrm{d}^{2} \Phi}{d \phi^{2}}+\mathrm{m}_{l}^{2} \Phi=0 \ldots \ldots \ldots \ldots \ldots . .(1) \\
& \frac{1}{\sin \theta} \frac{d}{d \theta}\left(\sin \theta \frac{d \Theta}{d \theta}\right)+\left[l(l+1)-\frac{\mathrm{m}_{l}^{2}}{\sin ^{2} \theta}\right] \Theta(\theta)=0 \ldots .  \tag{2}\\
& \frac{1}{r^{2}} \frac{d}{d r}\left(r^{2} \frac{\partial R}{\partial r}\right)+\left[\frac{2 m r^{2}}{\hbar^{2}}\left(\mathrm{E}+\frac{\mathrm{ke}^{2}}{\mathrm{r}}\right)-\frac{l(l+1)}{r^{2}}\right] R(r)=0 . . \tag{3}
\end{align*}
$$

These 3 "simple" diff. eqn describe the physics of the Hydrogen atom.

All we need to do now is guess the solutions of the diff. equations

Each of them, clearly, has a different functional form

## And Now the Solutions of The S. Eqns for Hydrogen Atom

The Azimuthal Diff. Equation : | $\frac{\mathrm{d}^{2} \Phi}{d \phi^{2}}+\mathrm{m}_{l}^{2} \Phi=0$ |
| :--- |

Solution: $\Phi(\phi)=\mathrm{A} \mathrm{e}^{\mathrm{im} \|_{\phi}}$ but need to check "Good Wavefunction Condition" Wave Function must be Single Valued for all $\phi \Rightarrow \Phi(\phi)=\Phi(\phi+2 \pi)$
$\Rightarrow \Phi(\phi)=\mathrm{A}^{\mathrm{im} m_{l} \phi}=\mathrm{A}^{\mathrm{i} m_{l}(\phi+2 \pi)} \Rightarrow m_{l}=0, \pm 1, \pm 2, \pm 3 \ldots .($ Magnetic Quantum \#)

The Polar Diff. Eq: $\frac{1}{\sin \theta} \frac{d}{d \theta}\left(\sin \theta \frac{d \Theta}{d \theta}\right)+\left[l(l+1)-\frac{\mathrm{m}_{l}^{2}}{\sin ^{2} \theta}\right] \Theta(\theta)=0$

Solutions : go by the name of "Associated Legendre Functions" only exist when the integers $l$ and $m_{l}$ are related as follows $m_{l}=0, \pm 1, \pm 2, \pm 3 \ldots . \pm l ; l=$ positive number

1: Orbital Quantum Number

Wavefunction Along Azimuthal Angle $\phi$ and Polar Angle $\theta$

For $l=0, m_{l}=0 \Rightarrow \Theta(\theta)=\frac{1}{\sqrt{2}}$


For $l=1, m_{l}=0, \pm 1 \Rightarrow$ Three Possibilities for the Orbital part of wavefunction

$$
\left[l=1, m_{l}=0\right] \Rightarrow \Theta(\theta)=\frac{\sqrt{6}}{2} \cos \theta \quad\left[l=1, m_{l}= \pm 1\right] \Rightarrow \Theta(\theta)=\frac{\sqrt{3}}{2} \sin \theta
$$

$\left[l=2, m_{l}=0\right] \Rightarrow \Theta(\theta)=\frac{\sqrt{10}}{4}\left(3 \cos ^{2} \theta-1\right)$
and so on and so forth (see book for more Functions)

Radial Differential Equations and Its Solutions

The Radial Diff. Eqn: $\frac{1}{r^{2}} \frac{d}{d r}\left(r^{2} \frac{\partial R}{\partial r}\right)+\left[\frac{2 \mathrm{mr}}{}{ }^{2}\left(\mathrm{E}+\frac{\mathrm{ke}^{2}}{\mathrm{r}}\right)-\frac{l(l+1)}{r^{2}}\right] R(r)=0$
Solutions : Associated Laguerre Functions R(r), Solutions exist only if:

1. $\mathrm{E}>0$ or has negtive values given by

$$
\mathrm{E}=-\frac{\mathrm{ke}^{2}}{2 \mathrm{a}_{0}}\left(\frac{1}{n^{2}}\right) ; \text { with } a_{0}=\frac{\hbar^{2}}{m k e^{2}}=\text { Bohr Radius }
$$

2. And when $\mathrm{n}=$ integer such that $l=0,1,2,3,4, \ldots \ldots . .(n-1)$
n = principal Quantum \# or the "big daddy" quantum \#

The Hydrogen Wavefunction: $\psi(r, \theta, \phi)$ and $\Psi(r, \theta, \phi, t)$
To Summarize : The hydrogen atom is brought to you by the letters:

```
n=1,2,3,4,5,\ldots.\infty
l=0,1,2,3,,4...(n-1)
m}=0,\pm1,\pm2,\pm3,\ldots\pm
```

The Spatial part of the Hydrogen Atom Wave Function is:

$$
\psi(r, \theta, \phi)=R_{n l}(r) . \Theta_{l m_{l}}(\theta) \quad . \Phi_{\mathrm{m}_{1}}(\phi)=R_{n \mid} \mathrm{Y}_{l}^{m_{l}}
$$

$\mathrm{Y}_{1}^{m_{l}}$ are known as Spherical Harmonics. They define the angular structure in the Hydrogen-like atoms.

The Full wavefunction is $\Psi(\mathrm{r}, \theta, \varphi, t)=\psi(r, \theta, \phi) e^{-\frac{i E}{\hbar} t}$

Radial Wave Functions For $n=1,2,3$


| $\mathrm{n}=1 \rightarrow$ K shell |  |  |
| :--- | :--- | :--- |
| $\mathrm{n}=2 \rightarrow$ L Shell |  |  |
| $\mathrm{n}=3 \rightarrow$ M shell |  |  |
| $\mathrm{n}=4 \rightarrow$ N Shell |  | $l=0 \rightarrow s($ harp sub shell <br> $l=1 \rightarrow p($ rincipal) sub shell <br>  <br> $\ldots . .$. |
| $l=2 \rightarrow d$ (iffuse) sub shell |  |  |
| $l=3 \rightarrow f($ undamental) ss |  |  |
| $l=4 \rightarrow g$ sub shell |  |  |
| $\ldots \ldots .$. | 279 |  |

## Symbolic Notation of Atomic States in Hydrogen

| $l \rightarrow$ | $s(l=0)$ | $p(l=1)$ | $d(l=2)$ | $f(l=3)$ | $g(l=4)$ | $\ldots \ldots$ |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| $n$ |  |  |  |  |  |  |
| $\downarrow$ |  |  |  |  |  |  |
| 1 | $1 s$ |  |  |  |  |  |
| 2 | $2 s$ | $2 p$ | $3 d$ |  |  |  |
| 3 | $3 s$ | $3 p$ | $4 d$ | $4 f$ |  |  |
| 4 | $4 s$ | $4 p$ | $5 d$ | $5 f$ | $5 g$ |  |
| 5 | $5 s$ | $5 p$ |  |  |  |  |

Note that:
$\cdot \mathrm{n}=1$ is a non-degenerate system
$\cdot \mathrm{n}>1$ are all degenerate in $l$ and $m_{l}$.
All states have same energy
But different angular configuration


Energy States, Degeneracy \& Transitions


## Facts About Ground State of H Atom



We need structure, we need variety, we need some curves!

Cross Sectional View of Hydrogen Atom prob. densities in r, $\theta, \phi$ Birth of Chemistry (Can make Fancy Bonds $\rightarrow$ Overlapping electron "clouds")

What's the electron "cloud" : Its the Probability Density in r, $\theta, \phi$ space!


## Interpreting Orbital Quantum Number (l)

Radial part of S.Eqn: $\frac{1}{r^{2}} \frac{d}{d r}\left(r^{2} \frac{d R}{d r}\right)+\left[\frac{2 \mathrm{~m}}{\hbar^{2}}\left(\mathrm{E}+\frac{\mathrm{ke}^{2}}{\mathrm{r}}\right)-\frac{l(l+1)}{r^{2}}\right] R(r)=0$
For H Atom: $\mathrm{E}=\mathrm{K}+\mathrm{U}=\mathrm{K}_{\text {RADIAL }}+\mathrm{K}_{\text {ORBITAL }}-\frac{k e^{2}}{r}$; substitute this in E
$\frac{1}{r^{2}} \frac{d}{d r}\left(r^{2} \frac{d R}{d r}\right)+\frac{2 \mathrm{~m}}{\hbar^{2}}\left[\mathrm{~K}_{\text {RADIAL }}+\mathrm{K}_{\text {ORBITAL }}-\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{l(l+1)}{r^{2}}\right] R(r)=0$
Examine the equation, if we set $\mathrm{K}_{\text {orbital }}=\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{l(l+1)}{r^{2}}$ then

what remains is a differential equation in $r$
$\frac{1}{r^{2}} \frac{d}{d r}\left(r^{2} \frac{d R}{d r}\right)+\frac{2 \mathrm{~m}}{\hbar^{2}}\left[\mathrm{~K}_{\text {RADIAL }}\right] R(r)=0$ which depends only on radius $r$ of orbit
Further, we also know that $\mathrm{K}_{\text {ORBTIAL }}=\frac{1}{2} m v_{\text {orbit }}^{2} ; \quad \overrightarrow{\mathrm{L}}=\overrightarrow{\mathrm{r}} \times \overrightarrow{\mathrm{p}} ; \quad|\mathrm{L}|=\mathrm{mv}_{\text {ort }} \mathrm{r} \Rightarrow \mathrm{K}_{\text {ORBTIAL }}=\frac{L^{2}}{2 m r^{2}}$
Putting it all togather: $\mathrm{K}_{\text {ORBTIAL }}=\frac{\hbar^{2}}{2 \mathrm{~m}} \frac{l(l+1)}{r^{2}}=\frac{L^{2}}{2 m r^{2}} \Rightarrow$ magnitude of Ang. Mom $|L|=\sqrt{l(l+1)} \hbar$
Since $l=$ positive integer $=0,1,2,3 \ldots(\mathrm{n}-1) \Rightarrow$ angular momentum $|L|=\sqrt{l(l+1)} t=$ discrete values
$|L|=\sqrt{l(l+1)} \hbar: \quad$ QUANTIZATION OF Electron's Angular Momentum

## Magnetic Quantum Number : $m_{l}$


$\vec{L}=\vec{r} \times \vec{p}$ (Right Hand Rule)
Classically, direction \& Magnitude of $\overrightarrow{\mathrm{L}}$ always well defined
QM: Can/Does $\overrightarrow{\mathrm{L}}$ have a definite direction? Proof by Negation: Suppose $\overrightarrow{\mathrm{L}}$ was precisely known/defined ( $\overrightarrow{\mathrm{L}} \| \hat{\mathrm{z}})$
Since $\vec{L}=\vec{r} \times \vec{p} \Rightarrow$ Electron MUST be in $\mathrm{x}-\mathrm{y}$ orbit plane
$\Rightarrow \Delta \mathrm{z}=0 ; \Delta \mathrm{p}_{z} \Delta \mathrm{z} \sim \hbar \Rightarrow \Delta \mathrm{p}_{\mathrm{z}} \sim \infty ; E=\frac{p^{2}}{2 m} \sim \infty!!!$
So, in Hydrogen atom, $\overrightarrow{\mathrm{L}}$ can not have precise measurable value Uncertainty Principle \& Angular Momentum : $\Delta \mathrm{L}_{z} \Delta \phi \sim \hbar$



## Where is it likely to be ? $\rightarrow$ Radial Probability Densities

$\Psi(r, \theta, \phi)=R_{n l}(r) \cdot \Theta_{l m_{l}}(\theta) \quad \Phi_{\mathrm{m}_{l}}(\phi)=R_{n l} \mathrm{Y}_{l}^{m_{l}}$
Probability Density Function in 3D:
$\mathrm{P}(\mathrm{r}, \theta, \phi)=\Psi^{*} \Psi=|\Psi(r, \theta, \phi)|^{2}=\left|R_{n \mid}\right|^{2} \cdot\left|\mathrm{Y}_{l}^{m_{1}}\right|^{2}$
Note : 3D Volume element $\mathrm{dV}=\mathrm{r}^{2} \cdot \sin \theta \cdot d r \cdot d \theta \cdot d \phi$ Prob. of finding particle in a tiny volume dV is P.dV $=\left|R_{n \mid}\right|^{2} .\left|\mathrm{Y}_{l}^{m_{1}}\right|^{2} . \mathrm{r}^{2} . \sin \theta . d r . d \theta \cdot d \phi$

The Radial part of Prob. distribution: $\mathrm{P}(\mathrm{r}) \mathrm{dr}$
$\mathrm{P}(\mathrm{r}) \mathrm{dr}=\left|R_{n l}\right|^{2} \cdot r^{2} d r \int_{0}^{\pi}\left|\Theta_{l m_{l}}(\theta)\right|^{2} d \theta \int_{0}^{2 \pi}\left|\Phi_{\mathrm{m}_{1}}(\phi)\right|^{2} d \phi$
When $\Theta_{l m_{I}}(\theta) \& \Phi_{\mathrm{m}_{1}}(\phi)$ are auto-normalized then $\mathrm{P}(\mathrm{r}) \mathrm{dr}=\left|R_{n l}\right|^{2} \cdot r^{2}{ }^{2} d r$; in other words $\mathrm{P}(\mathrm{r})=\mathrm{r}^{2}\left|R_{n l}\right|^{2}$
Normalization Condition: $1=\int_{0}^{\infty} \mathrm{r}^{2}\left|\mathrm{R}_{\mathrm{n} 1}\right|^{2} \mathrm{dr}$

Expectation Values


## Ground State: Radial Probability Density

$P(r) d r=|\psi(r)|^{2} .4 \pi r^{2} d r$
$\Rightarrow P(r) d r=\frac{4}{a_{0}^{3}} r^{2} e^{-2 \frac{r}{a_{0}}}$
Probability of finding Electron for $\mathrm{r}>\mathrm{a}_{0}$
$P_{r>a_{0}}=\int_{a}^{\infty} \frac{4}{a_{0}^{3}} r^{2} e^{-2 \frac{r}{a_{0}}} d r$
To solve, employ change of variable
Define $\mathrm{z}=\left[\frac{2 \mathrm{r}}{\mathrm{a}_{0}}\right]$;change limits of integration
$P_{r>a_{0}}=\frac{1}{2} \int_{2}^{\infty} z^{2} e^{-z} d z \quad$ (such integrals called Error. Fn)
$=-\left.\frac{1}{2}\left[z^{2}+2 z+2\right] e^{-z}\right|_{2} ^{\infty}=5 e^{2}=0.667 \Rightarrow 66.7 \%$


## Most Probable \& Average Distance of Electron from Nucleus

Most Probable Distance:
In the ground state $\left(n=1, l=0, m_{l}=0\right)$ $\square$
Most probable distance $r$ from Nucleus $\Rightarrow$ What value of $r$ is $P(r) \max$ ?
$\Rightarrow \frac{\mathrm{dP}}{\mathrm{dr}}=0 \Rightarrow \frac{4}{a_{0}^{3}} \cdot \frac{d}{d r}\left[r^{2} e^{-2 \frac{r}{a_{0}}}\right]=0 \Rightarrow\left[\frac{-2 r^{2}}{a_{0}}+2 r\right] e^{-2 \frac{r}{a_{0}}}=0$
$\Rightarrow \frac{2 r^{2}}{a_{0}}+2 r=0 \Rightarrow r=0$ or $r=a_{0} \ldots$ which solution is correct?
(see past quiz) : Can the electron BE at the center of Nucleus $(\mathrm{r}=0)$ ?
$P(r=0)=\frac{4}{a_{0}^{3}} 0^{2} e^{-2 \frac{0}{a_{0}}}=0!\Rightarrow$ Most Probable distance $r=a_{0} \quad$ (Bohr guessed right)
What about the AVERAGE location $<\mathrm{r}>$ of the electron in Ground state?
$<\mathrm{r}>=\int_{\mathrm{r}=0}^{\infty} \mathrm{rP}(\mathrm{r}) \mathrm{dr}=\frac{4}{a_{0}^{3}} \int_{0}^{\infty} r r^{2} e^{-2 \frac{r}{a_{0}}} d r \quad \ldots$ change of variable $\mathrm{z}=\frac{2 \mathrm{r}}{\mathrm{a}_{0}}$
$\Rightarrow\langle r\rangle=\frac{a_{0}}{4} \int_{z=0}^{\infty} z^{3} e^{-z} d z \ldots \ldots . \quad$ Use general form $\int_{0}^{\infty} z^{n} e^{-z} d z=n!=n(n-1)(n-2) \ldots(1)$
$\Rightarrow \quad<r>=\frac{a_{0}}{4} 3!=\frac{3 a_{0}}{2} \neq a_{0}!$ Average $\&$ most likely distance is not same. Why?
Asnwer is in the form of the radial Prob. Density: Not symmetric

Radial Probability Distribution $P(r)=r^{2} R(r)$


Normalized Spherical Harmonics \& Structure in H Atom
TABLE 7-1 Spherical harmonics

| $l=0$ | $m=0$ | $Y_{\infty 0}=\sqrt{\frac{1}{4 \pi}}$ |
| :---: | :---: | :---: |
| $l=1$ | $m=1$ | $Y_{11}=-\sqrt{\frac{3}{8 \pi}} \sin \theta e^{i \phi}$ |
|  | $m=0$ | $Y_{10}=\sqrt{\frac{3}{4 \pi}} \cos \theta$ |
|  | $m=-1$ | $Y_{1-1}=\sqrt{\frac{3}{8 \pi}} \sin \theta e^{-i \phi}$ |
| $l=2$ | $m=2$ | $Y_{22}=\sqrt{\frac{15}{32 \pi}} \sin ^{2} \theta e^{2 / \phi}$ |
|  | $m=1$ | $Y_{21}=-\sqrt{\frac{15}{8 \pi}} \sin \theta \cos \theta e^{i \phi}$ |
|  | $m=0$ | $Y_{20}=\sqrt{\frac{5}{16 \pi}}\left(3 \cos ^{2} \theta-1\right)$ |
|  | $m=-1$ | $Y_{2-1}=\sqrt{\frac{15}{8 \pi}} \sin \theta \cos \theta e^{-i \phi}$ |
|  | $m=-2$ | $Y_{2-2}=\sqrt{\frac{15}{32 \pi}} \sin ^{2} \theta e^{-2 i \phi}$ |

## Excited States ( $n>1$ ) of Hydrogen Atom : Birth of Chemistry !

| Features of Wavefunction in $\theta \& \phi$ : |  |
| :---: | :---: |
| Consider $n=2, l=0 \Rightarrow \psi_{200}=$ Spherically Symmetric (last slide) |  |
| Excited States (3 \& each with same $\mathrm{E}_{\mathrm{n}}$ ) : |  |
| $\psi_{211}, \psi_{210}, \psi_{21-1}$ are all $2 p$ states |  |
| $\psi_{211}=\mathrm{R}_{21} \mathrm{Y}_{1}^{1}=\left(\frac{1}{\pi}\right)\left(\frac{\mathrm{Z}}{\mathrm{a}_{0}}\right)^{3 / 2}\left(\frac{Z}{8}\right)\left(\frac{r}{a_{0}}\right) e^{\frac{-\mathrm{Zr}}{a_{0}}} \cdot \sin \theta \cdot e^{i \phi}$ |  |
| $\left\|\psi_{211}\right\|^{2}=\left\|\psi_{211}^{*} \psi_{211}\right\| \propto \sin ^{2} \theta$ Max at $\theta=\frac{\pi}{2}$, min at $\theta=0$; Symm in $\phi$ |  |
| What about ( $\mathrm{n}=2, \ell=1, \mathrm{~m}_{1}=0$ ) $\psi_{210}=R_{21}(\mathrm{r}) \mathrm{Y}_{1}^{0}(\theta, \phi) ;$ | $z \quad 2 p_{z}$ |
| $\mathrm{Y}_{1}^{0}(\theta, \phi) \propto \frac{1}{2} \sqrt{\frac{3}{\pi}} \cos \theta ;$ | $y$ |
| Function is max at $\theta=0$, min at $\theta=\frac{\pi}{2}$ |  |
| We call this $2 p_{z}$ state because of its extent in z |  |

## Excited States $(n>1)$ of Hydrogen Atom : Birth of Chemistry!

Remember Principle of Linear Superposition
for the TISE which is basically a simple differential equation:
$-\frac{\hbar^{2}}{2 \mathrm{~m}} \nabla^{2} \psi+U \psi=E \psi$

Principle of Linear Superposition $\Rightarrow$ If $\psi_{1}$ and $\psi_{2}$ are sol. of TISE then a "designer" wavefunction made of linear sum $\psi=a \psi_{1}+b \psi_{2}$ is also a sol. of the diff. equation !
To check this, just substitute $\psi$ ' in place of $\psi$
\& convince yourself that

$$
-\frac{\hbar^{2}}{2 \mathrm{~m}} \nabla^{2} \psi^{\prime}+U \psi^{\prime}=E \psi^{\prime}
$$

The diversity in Chemistry and Biology DEPENDS on this superposition rule

## Designer Wave Functions: Solutions of S. Eq !

Linear Superposition Principle means allows me to "cook up" wavefunctions $\psi_{2 \mathrm{p}_{\mathrm{x}}}=\frac{1}{\sqrt{2}}\left[\psi_{211}+\psi_{21-1}\right] \ldots .$. has electron "cloud" oriented along x axis $\psi_{2 \mathrm{p}_{\mathrm{y}}}=\frac{1}{\sqrt{2}}\left[\psi_{211}-\psi_{21-1}\right] \ldots .$. has electron "cloud" oriented along y axis So from 4 solutions $\psi_{200}, \psi_{210}, \psi_{211}, \psi_{21-1} \rightarrow 2 s, 2 p_{x}, 2 p_{y}, 2 p_{z}$

Similarly for $\mathrm{n}=3$ states ... and so on ...can get very complicated structure in $\theta \& \phi \ldots . .$. which I can then mix \& match to make electrons "most likely" to be where I want them to be !


Designer Wave Functions: Solutions of S. Eq !



## More Radial Probabilities P(r) Vs. $r / a_{0}$



Net Prob. densities for $\mathrm{n}=2$ states


## Transition Between States In Quantum Systems

In formulating the Hydrogen Atom, Bohr was obliged to postulate that the frequency
of radiation emitted by an atom dropping from energy level $\mathrm{E}_{\mathrm{m}}$ to a lower level $\mathrm{E}_{\mathrm{n}}$ is:

$$
\mathrm{f}=\frac{\mathrm{E}_{\mathrm{m}}-\mathrm{E}_{\mathrm{n}}}{h}
$$

This relationship rises naturally in Quantum Mechanics, consider for simplicity a system in which an electron only in the x direction:The time-dependent Wavefunction $\Psi_{n}(\mathrm{x}, \mathrm{t})=\psi_{n}(\mathrm{x}) \mathrm{e}^{-\mathrm{i} \frac{\mathrm{E}_{\mathrm{n}}}{\hbar} t}$;
$<\mathrm{x}\rangle=\int^{\infty} \mathrm{x} \psi_{\mathrm{n}}^{*} \psi_{n} d x=$ constant in time, does not oscillate, no radiation occurs
But, due to an external perturbation lasting some time, electron shifts from one state (m) to another(n) In this period wavefunction of electron is a linear superposition of two possible states $\Psi=a \Psi_{n}+b \Psi_{m} ; \mathrm{a}^{*} \mathrm{a}=$ prob. of electron in state n and $\mathrm{b}^{*} \mathrm{~b}=$ prob. of electron in state $\mathrm{m} ; \mathrm{a}^{*} \mathrm{a}+\mathrm{b}^{*} \mathrm{~b}=1$ Initially $a=1, b=0$ and finally $a=0, b=1$. While the electron is in either state there is no radiation but when it is in the midst of transition from $\mathrm{m} \rightarrow \mathrm{n}$, both a and b have non-vanishing values and radiation is produced. Expectation value for compostive wavefunction $\langle\mathrm{x}\rangle=\int \mathrm{x} \Psi \Psi^{*} \Psi d x$;
$<\mathrm{x}>=\int^{\infty} \mathrm{x}\left(\mathrm{a}^{2} \Psi_{\mathrm{n}}^{*} \Psi_{n}+b^{*} a \Psi_{\mathrm{m}}^{*} \Psi_{n}+\mathrm{a}^{*} \mathrm{~b} \Psi_{n}^{*} \Psi_{m}+\mathrm{b}^{2} \Psi_{\mathrm{m}}^{*} \Psi_{m}\right) d x$

## Transition Between States In Quantum Systems

$<\mathrm{x}>=\int_{-\infty} \mathrm{x}\left(\mathrm{a}^{2} \Psi_{\mathrm{n}}^{*} \Psi_{n}+b^{*} a \Psi_{\mathrm{m}}^{*} \Psi_{n}+\mathrm{a}^{*} \mathrm{~b} \Psi_{n}^{*} \Psi_{m}+\mathrm{b}^{2} \Psi_{\mathrm{m}}^{*} \Psi_{m}\right) d x$
$<x>=a^{2} \int x \psi_{n}^{*} \psi_{n} d x+b^{2} \int x \psi_{m}^{*} \psi_{m} d x$

$$
+a b^{*} \int x \psi_{m}^{*} e^{+i\left(E_{m} / \hbar\right) t} \psi_{n} e^{-i\left(E_{n} / \hbar\right) t} d x+a^{*} b \int x \psi_{n}^{*} e^{+i\left(E_{n} / \hbar\right) t} \psi_{m} e^{-i\left(E_{m} / \hbar\right) t} d x
$$

Use $\mathrm{e}^{\mathrm{i} \theta}=\cos \theta+i \sin \theta$ and $\mathrm{e}^{-\mathrm{i} \theta}=\cos \theta-i \sin \theta$ in the above and consider just the REAL part of expression for the last two terms, it varies with time as $\cos \left(\frac{\mathrm{E}_{\mathrm{m}}-\mathrm{E}_{\mathrm{n}}}{\hbar}\right) t=\cos 2 \pi\left(\frac{\mathrm{E}_{\mathrm{m}}-\mathrm{E}_{\mathrm{n}}}{h}\right) t=\cos 2 \pi \mathrm{ft}$

So the $\langle\mathrm{x}\rangle$ of the electron oscillates with frequency f and one has a nice electric dipole analogy $\Rightarrow$ Hence radiative transtions !

Similarly for particle in an infinite well or harmonic oscillator

## What's So "Magnetic" about $m_{l}$ ?

Precessing electron $\rightarrow$ Current in loop $\rightarrow$ Magnetic Dipole moment $\mu$


The electron's motion $\rightarrow$ hydrogen atom is a dipole magnet

## The "Magnetism" of an Orbiting Electron

Precessing electron $\rightarrow$ Current in loop $\rightarrow$ Magnetic Dipole moment $\mu$


Electron in motion around nucleus $\Rightarrow$ circulating charge $\Rightarrow$ curent $i$
$i=\frac{-e}{T}=\frac{-e}{\frac{2 \pi r}{v}}=\frac{-e p}{2 \pi m r} ;$ Area of current loop $\mathrm{A}=\pi \mathrm{r}^{2}$
Magnetic Moment $|\mu|=\mathrm{iA}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) r p ; \quad \vec{\mu}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) \vec{r} \times \vec{p}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) \vec{L}$
Like the $\overrightarrow{\mathrm{L}}$, magnetic moment $\vec{\mu}$ also precesses about " z " axis

$$
\text { z component, } \mu_{\mathrm{z}}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) L_{z}=\left(\frac{-\mathrm{e} \hbar}{2 \mathrm{~m}}\right) m_{l}=-\mu_{B} m_{l}=\text { quantized! }
$$

## Quantized Magnetic Moment



Why all this ? Need to find a way to break the Energy Degeneracy \& get electron in each $\left(n, l, m_{l}\right)$ state to identify itself, so we can "talk" to it and make it do our bidding:
" Walk this way, talk this way!"

## The "Magnetism" of an Orbiting Electron

Precessing electron $\rightarrow$ Current in loop $\rightarrow$ Magnetic Dipole moment $\mu$


Electron in motion around nucleus $\Rightarrow$ circulating charge $\Rightarrow$ curent $i$
$i=\frac{-e}{T}=\frac{-e}{\frac{2 \pi r}{v}}=\frac{-e p}{2 \pi m r} ;$ Area of current loop $\mathrm{A}=\pi \mathrm{r}^{2}$
Magnetic Moment $|\mu|=\mathrm{iA}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) r p ; \vec{\mu}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) \vec{r} \times \vec{p}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) \vec{L}$
Like the $\overrightarrow{\mathrm{L}}$, magnetic moment $\vec{\mu}$ also precesses about " z " axis

$$
\text { z component, } \mu_{\mathrm{z}}=\left(\frac{-\mathrm{e}}{2 \mathrm{~m}}\right) L_{\mathrm{z}}=\left(\frac{-\mathrm{e} \hbar}{2 \mathrm{~m}}\right) m_{l}=-\mu_{\mathrm{B}} m_{l}=\text { quantized! }
$$

## Bar Magnet Model of Magnetic Moment

In external B field, magnet experiences a torque which tends to align it with the field direction


If the magnet is spinning, torque causes magnet to precess around the ext. B field with a constant frequency: Larmor frequency



In presence of External B Field, Total energy of H atom changes to

$$
\mathrm{E}=\mathrm{E}_{0}+\hbar \omega_{L} m_{l}
$$

So the Ext. B field can break the E degeneracy "organically" inherent in the H atom. The Energy now depends not just on $n$ but also $m_{l}$

## 4E : The Quantum Universe

Lecture 29, May 24
Vivek Sharma
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## Electron has "Spin": An additional degree of freedom

Electron possesses additional "hidden" degree of freedom : "Spinning around itself" !
Spin Quantum \# $s=\frac{1}{2}$ (either Up or Down)
How do we know this ? $\Rightarrow$ Stern-Gerlach expt Spin Vector $\vec{S}$ (a form of angular momentum) is also Quantized

$$
|\overrightarrow{\mathrm{S}}|=\sqrt{s(s+1)} \hbar=\sqrt{\frac{3}{4}} \hbar
$$

$\& \mathrm{~S}_{\mathrm{z}}=m_{\mathrm{s}} \hbar ; m_{\mathrm{s}}= \pm \frac{1}{2}$
Spinning electron is an entitity defying any simple classical description. ....hidden D.O.F
$|\overrightarrow{\mathrm{S}}|=\sqrt{s(s+1)} \hbar$


Spin angular momentum S also exhibits Space quantization


In an inhomogeneous field perpendicular to beam direction, magnetic moment $\mu$ experiences a force $F_{z}$ whose direction depends on $Z$ component of the net magnetic moment \& inhomogeneity $\mathrm{dB} / \mathrm{dz}$. The force deflects magnetic moment up or down. Space Quantization means expect $(21+1)$ deflections. For $l=0$, expect all electrons to arrive on the screen at the center (no deflection)


This was a big surprise for Stern-Gerlach ! They had accidentally Discovered a new degree of freedom for electron : "spin" which Can take only two orientations for angular momentum S : up or down Leads to a new quantum number $s=1 / 2$. As a result:

> Z Component of Spin Angular Momentum $S_{z}=m_{s} \hbar$
> The magnitude $|S|=\sqrt{s(s+1)} \hbar$ is FIXED, never changes !
> Allowed orientations are $s(s+1)=2$
> $\overrightarrow{\mathrm{~S}} \Rightarrow \vec{\mu}_{s}$; The corresponding Spin Magnetic Moment

## What Stern\&Gerlach Saw in l=0 Silver Atoms <br> $\square$



Picture changes instantaneously as the external Field is switched off or on....discovery !

## Four (not 3) Numbers Describe Hydrogen Atom $\rightarrow n, l, m_{l}, m_{s}$

## "Spinning" charge gives rise to a dipole moment $\vec{\mu}_{\mathrm{s}}$

Imagine (semi-clasically, incorrectly!) electron as sphere: charge q , radius r Total charge uniformly distributed: $q=\sum \Delta q_{i}$;
as electron spins, each "chargelet" rotates $\Rightarrow$ current $\Rightarrow$ dipole moment $\vec{\mu}_{\mathrm{s}_{\mathrm{i}}}$

$$
\vec{\mu}_{s}=\left(\frac{q}{2 m_{e}}\right) \sum_{i} \vec{\mu}_{s_{\mathrm{i}}}=g\left(\frac{q}{2 m_{e}}\right) \vec{S} ; g=2
$$

In a Magnetic Field $\overrightarrow{\mathrm{B}} \Rightarrow$ magnetic energy due to spin $\mathrm{U}_{\mathrm{S}}=\vec{\mu}_{\mathrm{s}} \cdot \vec{B}$
Net Angular Momentum in H Atom $\overrightarrow{\mathrm{J}}=\overrightarrow{\mathrm{L}}+\overrightarrow{\mathrm{S}}$
Net Magnetic Moment of H atom: $\vec{\mu}=\vec{\mu}_{0}+\vec{\mu}_{s}=\left(\frac{-e}{2 m_{e}}\right)(\vec{L}+g \vec{S})$


Notice that since $g=2$, net dipole moment vector $\vec{\mu}$ is not $\|$ to $\vec{J}$
(There are many such "ubiquitous" quantum numbers for elementary particles!)

## Magnetic Energy in an External B Field

Contributions from Orbital and Spin motions. Defining Z axis to be the orientation of the B field:

$$
\mathrm{U}=-\vec{\mu} \cdot \vec{B}=\frac{e}{2 m} B\left\{L_{z}+g S_{z}\right\}=\frac{e \hbar}{2 m} B\left\{m_{l}+g m_{s}\right\}
$$

Example: Zeeman spectrum in $B=1 T$ produced by Hydrogen initially in $n=2$ state
after taking spin into account: $\mathrm{n}=2 \Rightarrow \mathrm{E}_{2}=-13.6 \mathrm{eV} / 2^{2}=-3.40 \mathrm{eV}$
Since $m_{1}=0, \pm 1$, orbital contribution to Magnetic energy $\mathrm{U}_{0}=m_{l} \hbar \omega_{L}$
This splits energy levels to $E=E_{2} \pm \hbar \omega_{L}$; for $m_{1}= \pm 1$ states
These states get further split in pairs due to spin magnetic moment
Since $\mathrm{g}=2$ and $\mathrm{m}_{\mathrm{s}}= \pm \frac{1}{2}$; spin energy is again Zeeman energy $=\hbar \omega_{L}$
As a result electrons in this shell have one of the following energies

$$
\begin{array}{|l|}
\hline \mathrm{E}_{2} \\
\mathrm{E}_{2} \pm \hbar \omega_{L} \\
\mathrm{E}_{2} \pm 2 \hbar \omega_{L} \\
\hline
\end{array}
$$

This leads to a variety of allowed $\left(\Delta\left(m_{1}+\mathrm{m}_{\mathrm{s}}\right)=0, \pm 1\right)$ energy transitions with different intensities (Principal and satellites) which are visible when B field is large (ignore LS couplinॄ


Spin-Orbit Interaction: L and S Momenta are Linked Magnetically
.


Electron revolving around Nucleus finds itself in a "internal" B field because in its frame of reference, the nucleus is orbiting around it

This B field, due to orbital motion, interacts with electron's spin dipole moment $\vec{\mu}_{s}$ $U_{m}=-\vec{\mu} \cdot \vec{B} \Rightarrow$ Energy larger when $\overrightarrow{\mathrm{S}} \| \overrightarrow{\mathrm{B}}$, smaller when anti-parallel
$\Rightarrow$ States with same $\left(n, l, m_{l}\right)$ but diff. spins $\Rightarrow$ energy level splitting/doubling due to $\overrightarrow{\mathrm{S}}$


Vector Model For Total Angular Momentum J


## Vector Model For Total Angular Momentum J

Coupling of Orbital \& Spin magnetic moments $\Rightarrow$
Neither Orbital nor Spin angular Momentum are conserved seperately!
$\overrightarrow{\mathrm{J}}=\overrightarrow{\mathrm{L}}+\overrightarrow{\mathrm{S}}$ is conserved so long as there are no external torques present Rules for Total Angular Momentum Quantization :
$|J|=\sqrt{j(j+1)} \hbar \quad$ with $j=|l+s|, l+s-1, l+s-2 \ldots \ldots, \ldots,|l-s|$
$\mathrm{J}_{z}=m_{j} \hbar \quad$ with $m_{j}=j, j-1, j-2 \ldots \ldots,-j$
Example: state with $\left(l=1, s=\frac{1}{2}\right)$
$j=3 / 2 \Rightarrow m_{j}=-3 / 2,-1 / 2,1 / 2,3 / 2$
$\mathrm{j}=1 / 2 \Rightarrow m_{j}= \pm 1 / 2$
In general $m_{j}$ takes $(2 j+1)$ values
$\Rightarrow$ Even \# of orientations

## Addition of Orbital and Spin Angular Momenta



## Complete Description of Hydrogen Atom

Full description
of the Hydrogen atom:
$\left\{n, l, m_{l}, m_{s}\right\}$
$\Downarrow$


LS Coupling
$\Downarrow$
$\left\{n, l, j, m_{s}\right\} \quad$ How to describe multi-electrons atoms like $\mathrm{He}, \mathrm{Li}$ etc?
corresponding How to order the Periodic table?
to 4 D.O.F.

- Four guiding principles:
- Indistinguishable particle \& Pauli Exclusion Principle
-Independent particle model (ignore inter-electron repulsion)
-Minimum Energy Principle for atom
-Hund's "rule" for order of filling vacant orbitals in an atom


## Multi-Electron Atoms : >1 electron in orbit around Nucleus

In Hydrogen Atom $\psi(\mathrm{r}, \theta, \phi)=\mathrm{R}(\mathrm{r}) \cdot \Theta(\theta) . \Phi(\phi) \equiv\left\{n, l, j, m_{j}\right\}$

In n-electron atom, to simplify, ignore electron-electron interactions complete wavefunction, in "independent"particle approx" :

$$
\psi(1,2,3, . . n)=\psi(1) \cdot \psi(2) \cdot \psi(3) \ldots \psi(n) \quad ? ? ?
$$

Complication $\rightarrow$ Electrons are identical particles, labeling meaningless! Question: How many electrons can have same set of quantum \#s?
Answer: No two electrons in an atom can have SAME set of quantum\#s (if not, all electrons would occupy 1s state (least energy)... no structure!!

Example of Indistinguishability: electron-electron scattering


If we cant follow electron path, don't know between which of the two scattering events actually happened

## Helium Atom: Two electrons around a Nucleus

In Helium, each electron has : kinetic energy + electrostatic potential energy
If electron " 1 " is located at $\mathrm{r}_{1}$ \& electron " 2 "is located at $\mathrm{r}_{2}$ then TISE has terms like:
$H_{1}=-\frac{\hbar^{2}}{2 \mathrm{~m}} \nabla_{1}^{2}+k \frac{(2 e)(-e)}{r_{1}} ; \mathrm{H}_{2}=-\frac{\hbar^{2}}{2 \mathrm{~m}} \nabla_{2}^{2}+k \frac{(2 e)(-e)}{r_{2}}$ such that
$\mathrm{H}_{1} \psi+\mathrm{H}_{2} \psi=\mathrm{E} \psi ; \mathrm{H}_{1} \& \mathrm{H}_{2}$ are same except for "label"
Independent Particle Approx $\Rightarrow$ ignore repulsive $U=k \frac{e^{2}}{\left|r_{2}-r_{1}\right|}$ term


Helium WaveFunction: $\psi=\psi\left(\mathrm{r}_{1}, r_{2}\right)$; Probability $P=\psi^{*}\left(\mathrm{r}_{1}, r_{2}\right) \psi\left(\mathrm{r}_{1}, r_{2}\right)$
But if we exchange location of (identical, indistinguishable) electrons $\Rightarrow\left|\psi\left(\mathrm{r}_{1}, r_{2}\right)\right|=\left|\psi\left(\mathrm{r}_{2}, r_{1}\right)\right|$
 when $\psi\left(\mathrm{r}_{1}, r_{2}\right)=-\psi\left(\mathrm{r}_{2}, r_{1}\right) \ldots \ldots \ldots \ldots \ldots \ldots .$. fermionic System (made of electron, proton e.g)
$\Rightarrow$ Helium wavefunction must be ODD; if electron " 1 " is in state a \& electron " 2 " is in state b
Then the net wavefunction $\psi_{a b}\left(\mathrm{r}_{1}, \mathrm{r}_{2}\right)=\psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)$ satisfies
$\mathrm{H}_{1} \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)=E_{a} \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)$
$\mathrm{H}_{2} \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)=E_{b} \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)$
and the sum
$\left[\mathrm{H}_{1}+\mathrm{H}_{2}\right] \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)=\left(E_{a}+E_{b}\right) \psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)$
Total Helium Energy $\mathrm{E} \simeq \mathrm{E}_{\mathrm{a}}+\mathrm{E}_{\mathrm{b}}=$ sum of Hydrogen atom like E

Helium wavefunction must be ODD $\Rightarrow$ anti-symmetric: $\psi_{a b}\left(\mathrm{r}_{1}, \mathrm{r}_{2}\right)=-\psi_{a b}\left(\mathrm{r}_{2}, \mathrm{r}_{1}\right)$


So it must be that $\psi_{a b}\left(\mathrm{r}_{1}, \mathrm{r}_{2}\right)=\psi_{\mathrm{a}}\left(r_{1}\right) \cdot \psi_{\mathrm{b}}\left(r_{2}\right)-\psi_{\mathrm{a}}\left(r_{2}\right) \cdot \psi_{\mathrm{b}}\left(r_{1}\right)$
It is impossible to tell, by looking at probability or energy which particular electron is in which state
If both are in the same quantum state $\Rightarrow \mathrm{a}=\mathrm{b} \& \psi_{a a}\left(\mathrm{r}_{1}, \mathrm{r}_{2}\right)=\psi_{b b}\left(\mathrm{r}_{1}, \mathrm{r}_{2}\right)=0 \ldots$ Pauli Exclusion principle
General Principles for Atomic Structure for n-electron system:

1. n-electron system is stable when its total energy is minimum
2.Only one electron can exist in a particular quantum state in an atom...not 2 or more!
2. Shells \& SubShells In Atomic Structure:
(a) ignore inter-electron repulsion (crude approx.)
(b) think of each electron in a constant "effective" mean Electric field
(Effective field: "Seen" Nuclear charge (+Ze) reduced by partial screening due to other electrons "buzzing" closer (in r) to Nucleus)
Electrons in a SHELL: have same n, are at similar <r> from nucleus, have similar energies
Electons in a SubShell: have same principal quantum number n

- Energy depends on $l$, those with lower $l$ closer to nucleus, more tightly bound
- all electrons in sub-shell have same energy, with minor dependence on $m_{l}, m_{s}$


## Shell \& Sub-Shell Energies \& Capacity



1. Shell \& subshell capacity limited due to Pauli Exclusion principle
2. Shell is made of sub-shells (of same principal quantum \# n)
3. Subshell $\Leftarrow(n, l)$, given $n \Rightarrow l=0,1,2,3, . .(n-1)$,

$$
\text { for any } l \Rightarrow m_{l}=0, \pm 1, \pm 2, . . \Rightarrow(2 l+1), m_{s}= \pm \frac{1}{2}
$$

$\qquad$ $5 p$
$-4 d$
$\Rightarrow$ Max. \# of electrons in a shell $=\sum$ subshell capacity $\mathrm{N}_{\mathrm{MAX}}=\sum_{l=0}^{n-1} 2 .(2 l+1)=2[1+3+5+. .2(n-1)+1]=2(n)\left[\frac{1}{2}(1+(2 n-1))\right]=2 n^{2}$ 4. The "K" Shell ( $n=1$ ) holds 2 electrons, "L" Shell ( $n=1$ ) holds 8 electrons, M shell ( $n=3$ ) holds 18 electrons ......
5. Shell is closed when fully occupied
6. Sub-Shell closed when
(a) $\sum \overrightarrow{\mathrm{L}}_{\mathrm{i}}=0, \sum \vec{S}_{\mathrm{i}}=0 \Rightarrow$ Effective charge distribution= symmetric
(b) Electrons are tightly bound since they "see" large nuclear charge
(c) Because $\sum \overrightarrow{\mathrm{L}}_{\mathrm{i}}=0 \Rightarrow$ No dipole moment $\Rightarrow$ No ability to attract electrons $\Rightarrow$ Inert! Noble gas!
6.Alkali Atoms: have a single "s" electron in outer orbit; nuclear charge heavily shielded by inner shell electrons
enrgy of "valence" electron
large orbital radius of valence electron


## Topics In Particle Physics

- Cosmic Messengers!
- Dirac, Anderson and the Positron !
- antimatter
- Fundamental forces in nature
- How elementary particles are produced: Accelerators
- Classification of Particle and How we know this
- Conservation laws
- Colored Quarks and Quantum Chromodynamics!
- Electroweak theory and Standard model
- The Higgs Particle and Large Hadron Collider
- Beyond the Standard model : Supersymmetry \& Strings


## Fundamental Particle Physics

What are the elementary constituents of matter ?
What are the forces that control their behaviour at the most basic level?


Size of Things


## Probing The Cosmic Onion: Experimentally



Rutherford (1909): Nuclear atom (proton)
Chadwick (1932): Discovers neutron
SLAC (1968): Quarks in neutrons and protons

Power of Microscope


Wavelength of probe radiation should be smaller than the object to be resolved

$$
\lambda \ll \frac{h}{p}=\frac{h c}{E}
$$

| Object | Size | Energy of Radiation |
| :--- | :--- | :--- |
| Atom | $10^{-10} \mathrm{~m}$ | 0.00001 GeV (electrons) |
| Nucleus | $10^{-14} \mathrm{~m}$ | 0.01 GeV (alphas) |
| Nucleon | $10^{-15} \mathrm{~m}$ | 0.1 GeV (electrons) |
| Quarks | $?$ | $>1 \mathrm{GeV}$ (electrons) |

Radioactive sources give energies in the range of MeV
Need accelerators for higher energies.



## Relativity, Dirac and Anti-matter

$$
E^{2}=(p c)^{2}+\left(m c^{2}\right)^{2} \Rightarrow E= \pm \sqrt{(p c)^{2}+\left(m c^{2}\right)^{2}}
$$

What does the negative energy solution imply ??!

- Dirac postulated that all negative energy states were filled with electrons. They exert no net force on any thing and thus are unobservable
- Used Pauli Excl. principle to claim that only "holes" in this infinite sea of negative energy states observable
- Holes would act as positive charge with positive energy
- Anderson's discovery of positron !


Pair Production: Photoelectric effect with a negative energy electron!


Pair Production Photographed in B field: Note Curvature


All particles have an anti-matter partner !




## The Four Fundamental Forces



## Forces in Nature

Table 15.1 Particle Interactions

| Interaction (Force) | Particles Acted on by Force | Relative <br> Strength ${ }^{\text {a }}$ | Typical Lifetimes for Decays via a Given Interaction | Range of <br> Force | Force- <br> Carrying <br> Particle <br> Exchanged |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Strong | Quarks, hadrons | 1 | $\leq 10^{-20} \mathrm{~s}$ | Short ( $\approx 1 \mathrm{fm}$ ) | Gluon |
| Electromagnetic | Charged particles | $\approx 10^{-2}$ | $\approx 10^{-16} \mathrm{~s}$ | Long ( $x$ ) | Photon |
| Wcak | Quarks, leptons | $\approx 10^{-6}$ | $\geq 10^{-10} \mathrm{~s}$ | Short ( $\approx 10^{-3} \mathrm{fm}$ ) | $\mathrm{W}^{ \pm}, \mathrm{Z}^{0}$ bosons |
| Gravitational | All particles | $\approx 10^{-43}$ | ? | Long ( $\infty$ ) | Graviton ${ }^{\text {b }}$ |

${ }^{\text {a }}$ For two $u$ quarks at $3 \times 10^{-17} \mathrm{~m}$.
${ }^{\mathrm{b}}$ Not experimentally detected.

Compton Scattering, Pair Production and Annihilation
!


Strong Interaction Between Protons and Neutrons

Uncertainty Principle and range of strong force

(a)


Weak Force of Beta Decay: Uncertainty Principle

(a)

(b)


## Unification of Physical Laws



## Unification of Forces



Analogy: Are steam, water and ice manifestation of the same thing ?

## Cosmic Ray Smashing Through Nucleus



## Particle Categories: Hadrons \& Leptons

| Category | Particle Name | Symbol | Antiparticle | $\begin{gathered} \text { Mass } \\ \left(\mathrm{MeV} / \mathrm{c}^{2}\right) \end{gathered}$ | B | $L_{\text {e }}$ | $\boldsymbol{L}_{\mu}$ | $L_{\text {r }}$ | S | Lifetime <br> (s) | Principal <br> Decay <br> Modes ${ }^{\text {a }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Leptons | Electron | $\mathrm{e}^{-}$ | $\mathrm{e}^{+}$ | 0.511 | 0 | +1 | 0 | 0 | 0 | Stable |  |
|  | Electron-neutrino | $\nu_{\mathrm{c}}$ | $\bar{v}_{\text {c }}$ | $<2.8 \times 10^{-6}$ | 0 | +1 | 0 | 0 | 0 | Stable |  |
|  | Muon | $\mu^{-}$ | $\mu^{+}$ | 105.7 | 0 | 0 | +1 | 0 | 0 | $2.19 \times 10^{-6}$ | $\mathrm{e}^{-\bar{\nu}_{\mathrm{c}} \nu_{\mu}}$ |
|  | Muon-neutrino | $\nu_{\mu}$ | $\bar{v}_{\mu}$ | $<3.5 \times 10^{-6}$ | 0 | 0 | +1 | 0 | 0 | Stable |  |
|  | Tau | $\tau^{-}$ | $\tau^{+}$ | 1784 | 0 | 0 | 0 | +1 | 0 | $3.3 \times 10^{-13}$ | $\mu^{-} \bar{\nu}_{\mu} \nu_{\gamma}$ |
|  | Tau-neutrino | $\nu_{\boldsymbol{r}}$ | $\bar{\nu}_{\sim}$ | $<8.4 \times 10^{-6}$ | 0 | 0 | 0 | +1 | 0 | Stable |  |
| Hadrons Mesons |  |  |  |  |  |  |  |  |  |  |  |
|  | Pion |  | $\pi^{-}$ | 139.6 | 0 | 0 | 0 | 0 | 0 | $2.60 \times 10^{-8}$ | $\mu^{+} \nu_{\mu}$ |
|  |  | $\pi^{0}$ | Self | 135.0 | 0 | 0 | 0 | 0 | 0 | $0.83 \times 10^{-16}$ | ${ }_{2 \gamma}{ }^{\mu}$ |
|  | Kaon | $\mathrm{K}^{+}$ | $\mathrm{K}^{-}$ | 493.7 | 0 | 0 | 0 | 0 | +1 | $1.24 \times 10^{-8}$ | $\begin{aligned} & \mu^{+} v_{\mu} \\ & n^{+} \end{aligned}$ |
|  |  | $\mathrm{K}_{5}{ }^{\text {¢ }}$ | $\overline{\mathbf{K}}_{s}^{0}$ | 497.7 | 0 | 0 | 0 | 0 | $+1$ | $0.89 \times 10^{-10}$ | ${ }^{\pi^{+} \pi^{+}}$ |
|  |  |  |  |  |  |  |  |  |  |  | $2 \pi^{0}$ |
|  |  | $\mathrm{K}_{L}^{0}$ | $\bar{\kappa}_{L}^{0}$ | 497.7 | 0 | 0 | 0 | 0 | $+1$ | $5.2 \times 10^{-8}$ | $\begin{aligned} & \pi^{*} \mathrm{e}^{\mp} \bar{\nu}_{\mathrm{c}} \\ & 3 \pi^{0} \end{aligned}$ |
|  |  |  |  |  |  |  |  |  |  |  | $\pi^{ \pm} \mu^{+} \bar{\nu}_{\mu}$ |
|  | Eta | $\eta$ | Self | 548.8 | 0 | 0 | 0 | 0 | 0 | $<10^{-18}$ | $2 \gamma, 3 \pi^{0}$ |
|  |  | $\eta^{\prime}$ | Self | 958 | 0 | 0 | 0 | 0 | 0 | $2.2 \times 10^{-21}$ | $\eta \pi^{+} \pi^{-}$ |
| Baryons | Proton Neutron <br> Lambda Sigma | P | $\overline{\text { p }}$ | 938.3 | +1 | 0 | 0 | 0 | 0 | Stable |  |
|  |  | n | $\bar{\square}$ | 939.6 | +1 | 0 | 0 | 0 | 0 | 624 | $\mathrm{pe}^{-\bar{\nu}_{\mathrm{c}}}$ |
|  |  | $\Lambda^{0}$ | $\overline{\bar{\Lambda}}^{0}$ | 1115.6 | +1 | 0 | 0 | 0 | -1 | $2.6 \times 10^{-10}$ | $\mathrm{p} \mathrm{z}^{-}, \mathrm{n} \pi^{0}$ |
|  |  | $\Sigma^{+}$ | $\overline{\Sigma^{-}}$ | 1189.4 | +1 | 0 | 0 | 0 | -1 | $0.80 \times 10^{-10}$ | $\mathrm{p} \pi^{0}, \mathrm{n} \pi^{+}$ |
|  |  | $\Sigma^{0}$ | $\stackrel{\bar{\Sigma}}{ }{ }^{0}$ | 1192.5 | +1 | 0 | 0 | 0 | -1 | $6 \times 10^{-20}$ | $\Lambda^{0} \gamma$ |
|  |  | $\mathrm{\Sigma}^{-}$ | $\bar{\Sigma}^{+}$ | 1197.3 | $+1$ | 0 | 0 | 0 | -1 | $1.5 \times 10^{-10}$ | $\mathrm{n} \pi^{-}$ |
|  | Delta | $\Delta^{++}$ | $\triangle$ | 1230 | +1 | 0 | 0 | 0 | 0 | $6 \times 10^{-24}$ |  |
|  |  | $\Delta^{\text {a }}$ | $\bar{\Delta}^{-}$ | 1231 | +1 | 0 | 0 | 0 | 0 | $6 \times 10^{-24}$ | $\mathrm{p} \pi^{0}, \mathrm{n} \pi^{+}$ |
|  |  | $\Delta^{0}$ | $\stackrel{\Delta^{0}}{ }{ }^{\text {a }}$ | 1232 | +1 | 0 | 0 | 0 | 0 | $6 \times 10^{-24}$ | $\mathrm{n} \pi^{0} \cdot \mathrm{p} \pi^{-}$ |
|  |  | $\Delta^{-}$ | $\bar{\Delta}^{+}$ | 1234 | +1 | 0 | 0 | 0 | 0 | $6 \times 10^{-24}$ | $\mathrm{n} \pi^{-}$ |
|  | Xi | $\Xi^{0}$ | $\bar{\Xi}^{\circ}$ | 1315 | +1 | 0 | 0 | 0 | -2 | $2.9 \times 10^{-10}$ | $\Lambda^{0} \pi^{0}$ |
|  |  | $\Xi^{-}$ | $\Xi^{+}$ | 1321 | +1 | 0 | 0 | 0 | -2 | $1.64 \times 10^{-10}$ | $\Lambda^{0} \pi^{-}$ |
|  | Omega | $\Omega^{-}$ | $\Omega^{+}$ | 1672 | +1 | 0 | 0 | 0 | -3 | $0.82 \times 10^{-10}$ | $\begin{aligned} & \Xi_{\Xi^{-}}^{\Lambda^{0} \pi^{0}}, \Xi^{0} \pi^{-}, \end{aligned}$ |



## Relativistic Force \& Acceleration

$\vec{p}=\frac{m \vec{u}}{\sqrt{1-(u / c)^{2}}}=\gamma m \vec{u}$

Relativistic
Force
And Acceleration
$F=\left[\frac{m}{\left(1-(u / c)^{2}\right)^{3 / 2}}\right] \frac{d u}{d t}:$ Relativistic Force
Since Acceleration $\overrightarrow{\mathrm{a}}=\frac{d \vec{u}}{d t}$, [rate of change of velocity]

$$
\overrightarrow{\mathrm{a}}=\frac{\overrightarrow{\mathrm{F}}}{\mathrm{~m}}\left[1-(u / c)^{2}\right]^{3 / 2}
$$

Reason why you cant quite get up to the speed of light no matter how hard you try!

Linear Particle Accelerator : Parallel Plates With Potential Difference


Charged particle q moves in straight line in a uniform electric field $\vec{E}$ with speed $\overrightarrow{\mathrm{u}}$ accelarates under force $\overrightarrow{\mathrm{F}}=\mathrm{q} \overrightarrow{\mathrm{E}}$
$|\vec{a}|=\left|\frac{d \vec{u}}{d t}\right|=\left|\frac{\vec{F}}{m}\right|\left(1-\frac{u^{2}}{c^{2}}\right)^{3 / 2}=\left|\frac{q \vec{E}}{m}\right|\left(1-\frac{u^{2}}{c^{2}}\right)^{3 / 2}$
Under force, work is done on the particle, it gains Kinetic energy

New Unit of Energy
$1 \mathrm{eV}=1.6 \times 10^{-19} \mathrm{Joules}$
larger the potential difference V across
$1 \mathrm{MeV}=1.6 \times 10^{-13}$ Joules
$1 \mathrm{GeV}=1.6 \times 10-10$ Joules

Linear Accelerator : 50 Billion Volts Accelerating Potential


Discovery of Quarks: Constituents of Proton


High Energy Proton smashing into a Proton Rich Target


Magnetic Confinement \& Circular Particle Accelerator


## Charged Form of Matter \& Anti-Matter in a B Field

Antimatter form of electron = Positron ( $\mathrm{e}^{+}$)
Same Mass but opposite Charge
Positron curls the other way from electron ir a B Field


Accelerating Electrons Thru RF Cavities



Circular Particle Accelerator: LEP @ CERN, Geneve
Accelerated electron through an effective voltage of 100 Billion Volts !

circular track for accelerating electron




Sequence of Events Following $e^{+} e^{-}$Annihilation


Collider Detector: Concentric Array of Specialized Particle Detectors



## DNA of Fundamental Particles: Vital Statistics



Some Quantum Numbers of Quarks

| Properties of Quarks and Antiquarks |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Flavor | Spin | Charge | Baryon <br> Number | Strangeness | Charm | Topness | Bottomness |
| Quarks |  |  |  |  |  |  |  |
| $u$ (up) | $\frac{1}{2} \hbar$ | $+\frac{2}{3} e$ | $+\frac{1}{3}$ | 0 | 0 | 0 | 0 |
| $d$ (down) | $\frac{1}{2} \hbar$ | $-\frac{1}{3} e$ | $+\frac{1}{3}$ | 0 | 0 | 0 | 0 |
| $s$ (strange) | $\frac{1}{2} \hbar$ | $-\frac{1}{3} e$ | $+\frac{1}{3}$ | -1 | 0 | 0 | 0 |
| $c$ (charmed) | $\frac{1}{2} \hbar$ | $+\frac{2}{3} e$ | $+\frac{1}{3}$ | 0 | +1 | 0 | 0 |
| $t$ (top) | $\frac{1}{2} h$ | $+\frac{2}{3} e$ | $+\frac{1}{3}$ | 0 | 0 | +1 | 0 |
| $b$ (bottom) | $\frac{1}{2} \hbar$ | $-\frac{1}{3} e$ | $+\frac{1}{3}$ | 0 | 0 | 0 | +1 |
| Antiquarks |  |  |  |  |  |  |  |
| $\bar{u}$ | $\frac{1}{2} \hbar$ | $-\frac{2}{3} e$ | $-\frac{1}{3}$ | 0 | 0 | 0 | 0 |
| $\bar{d}$ | $\frac{1}{2} \hbar$ | $+\frac{1}{3} e$ | $-\frac{1}{3}$ | 0 | 0 | 0 | 0 |
| $\bar{s}$ | $\frac{1}{2} \hbar$ | $+\frac{1}{3} e$ | $-\frac{1}{3}$ | +1 | 0 | 0 | 0 |
| $\bar{c}$ | $\frac{1}{2} \hbar$ | $-\frac{2}{3} e$ | $-\frac{1}{3}$ | 0 | -1 | 0 | 0 |
| $\bar{t}$ | $\frac{1}{2} \hbar$ | $-\frac{2}{3} e$ | $-\frac{1}{3}$ | 0 | 0 | -1 | 0 |
| $\bar{b}$ | $\frac{1}{2} \hbar$ | $+\frac{1}{3} e$ | $-\frac{1}{3}$ | 0 | 0 | 0 | -1 |

Composite Particles are made of Quarks held by "glue"
Proton = (uud); Neutron = (udd), Pion+ = (u dbar), K+ = (sbar d)

## Some Open Questions In Particle Physics

- How do particles get the masses they have?
- Physicists believe particle masses are generated by interaction with a mysterious field that permeates the entire universe
- Stronger the particle interacts with the field, the more massive it is
- It could be a new fundamental field called HIGGS field
- Or it may be a composite object made of new particles (techniquarks) tightly bound together by a new force (technicolor!)
- Whatever the nature of this mass mechanism, odds are solid that it will be produced when beams of protons with energy of 7 TRILLION eV collide at the LHC accelerator
- Could be seen as one or many new Higgs particle
- If the Universe is made of $>4$ dimensions, some of the extra dimensions could "pop" out in these violent collisions
- Little blackholes could also be produced in these high energy interactions....and the detector will catch them in action !!


## Hunting for Higgs Particle With CMS Detector




