A little different than normal... We will do the first part of the lecture in our desks, and then the rest of this **PowerPoint will be done as** you do the lab activity.

## **N50 – Titrations** Our last lecture of the year!

**Target:** I can set up and perform a titration.

Link to YouTube Presentation: <u>https://youtu.be/6owm822vyhl</u>

# What is titration?

A way to determine the concentration of an unknown substance.

- Uses the fact that acids and bases react with each other in "neutralization reactions"
- At the point where the neutralization reaction is finished # moles Acid = # moles Base

# Key Terms

#### **Titrand**

The unknown solution you are interested in

#### **Titrant**

The solution with the known concentration

#### **Equivalence Point**

The point at which all the titrand has reacted with the titrant. # Moles Acid = # Moles Base

#### End Point

The point at which your titration *seems* finished during the lab

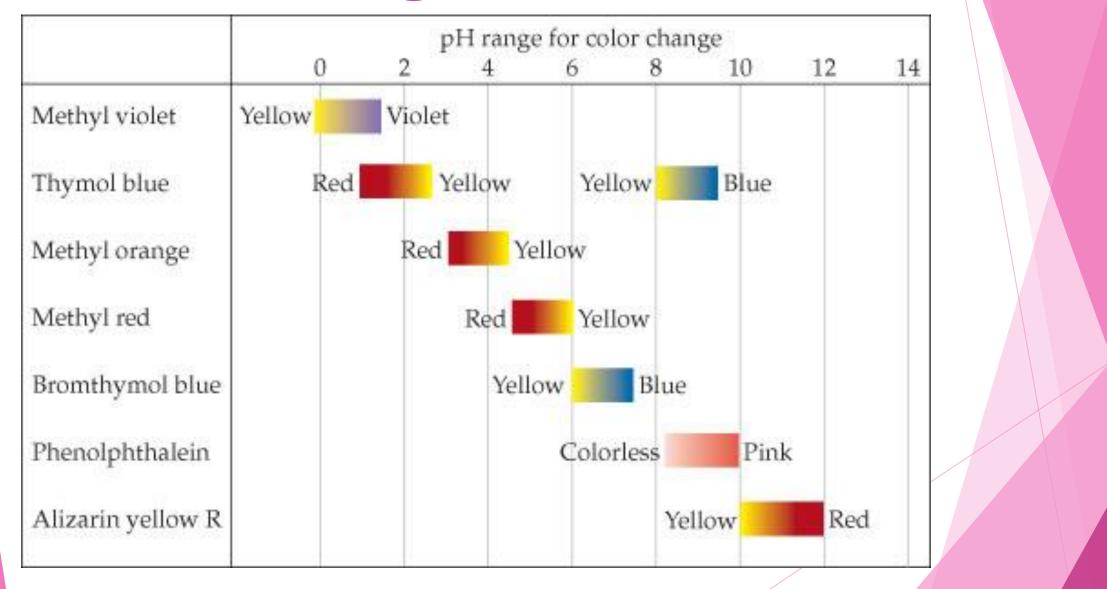
- a color change happens for example

# How do you know you reached the end point?

#### Use an **INDICATOR**

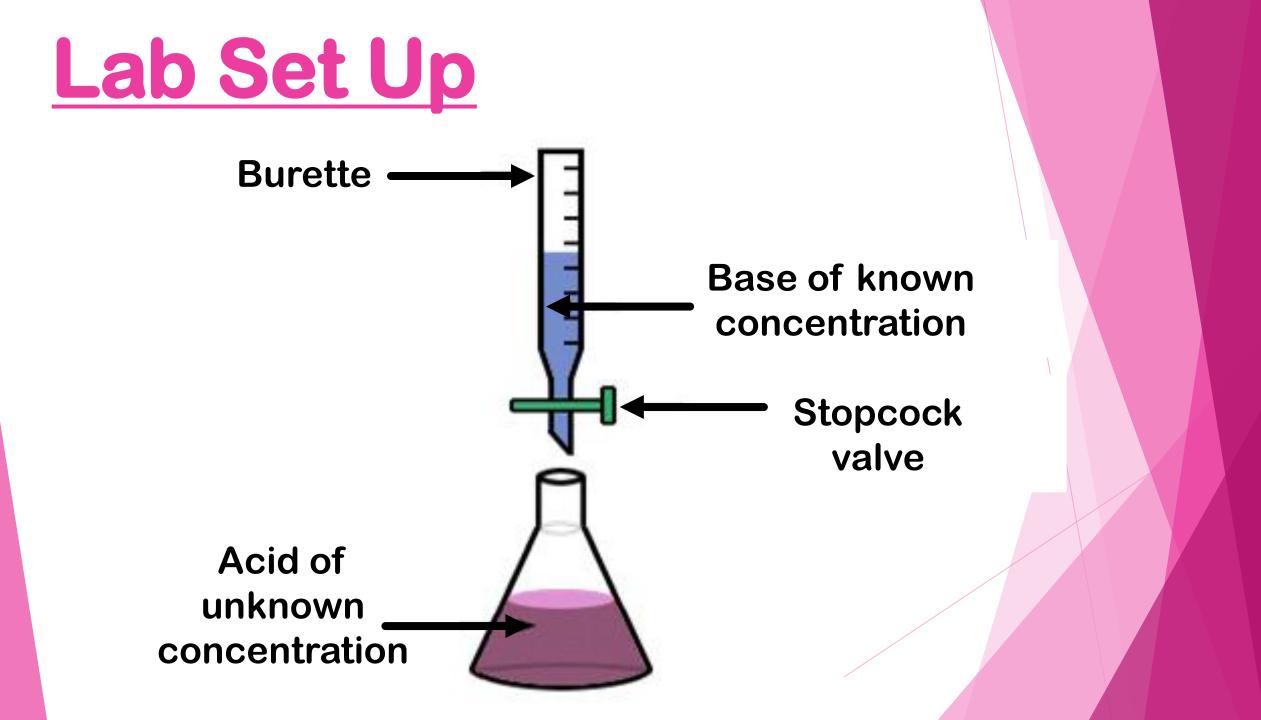
Turns colors based on pH – can show you visually when you have reached the end point.

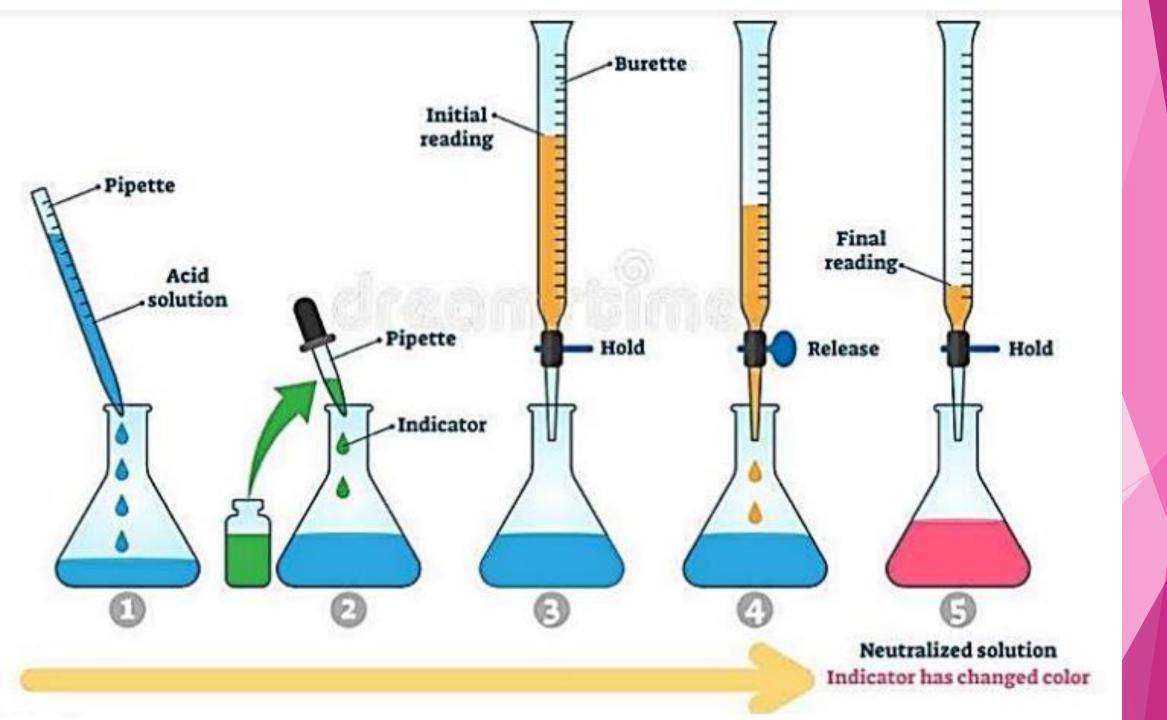
# **Pick the right indicator!**



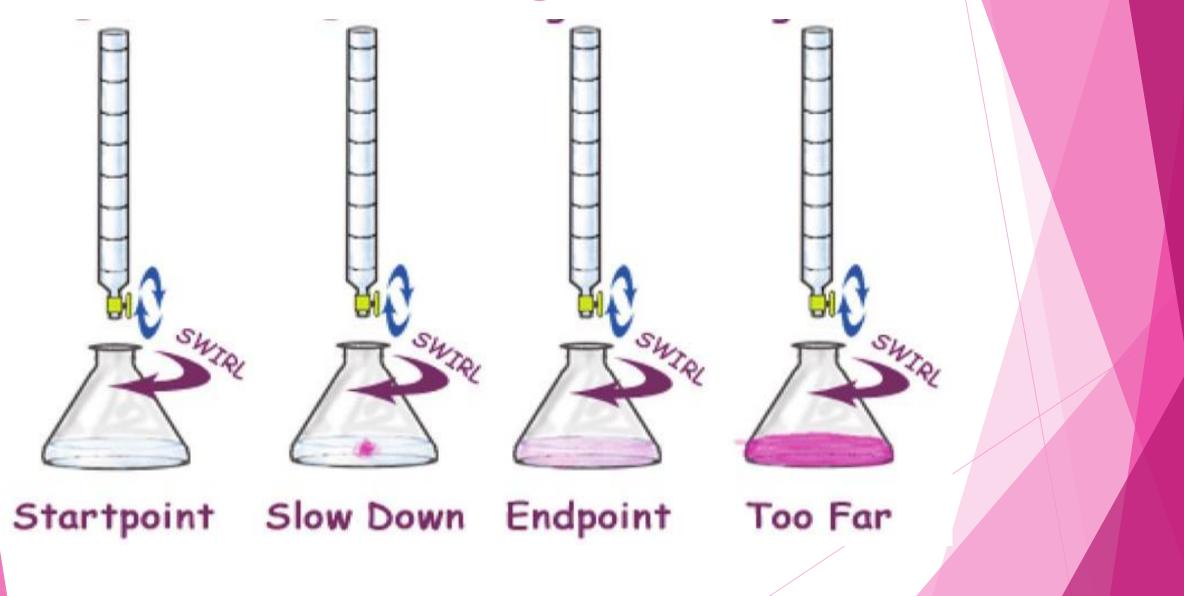
## **Pick the right indicator!**

Crystal Violet	0		2	3	4	5	6	7	8	9	10	11	12	13	1
Cresol Red			_			-									
Cresolphthalein (meta)	-		_	_		-					1	_			
-	-			-		-		-	1		-	-		_	
Cresol Purple	-			-	-	_						_		-	
Thymol Blue	-	-	-			-	-	-	-	-				_	
Methyl Orange - Xylene Cyanol	-	_					_	_	_	_		_	_	_	
Bromophenol Blue				_				_		_					
Congo Red															
Methyl Orange															
Alizarin Red S															
Bromocresol Green															
Dichlorofluorescein															
Methyl Red											1				
Bromocresol Green/Methyl Red	I I														
Bromocresol Purple															
Chlorophenol Red							10								
Bromothymol Blue															
Phenol Red															
Naphtholphthalein (alpha)															
Phenolphthalein															
Cresolphthalein (ortho)									1						
Thymolphthalein															
Indigo Carmine															
Universal Indicator														04	

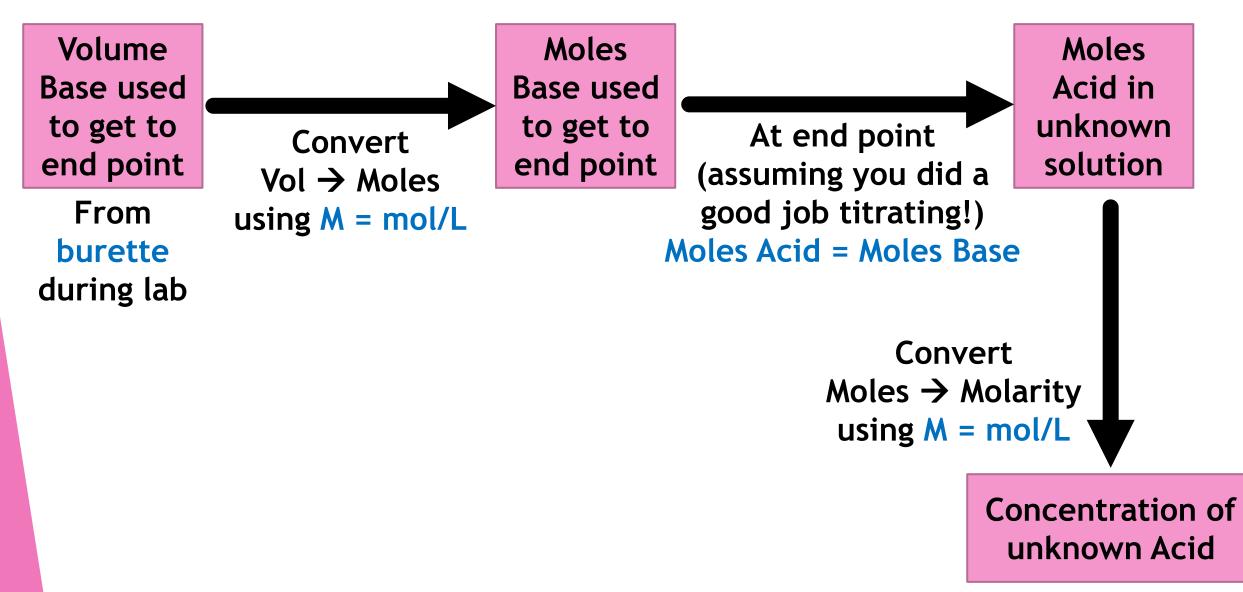




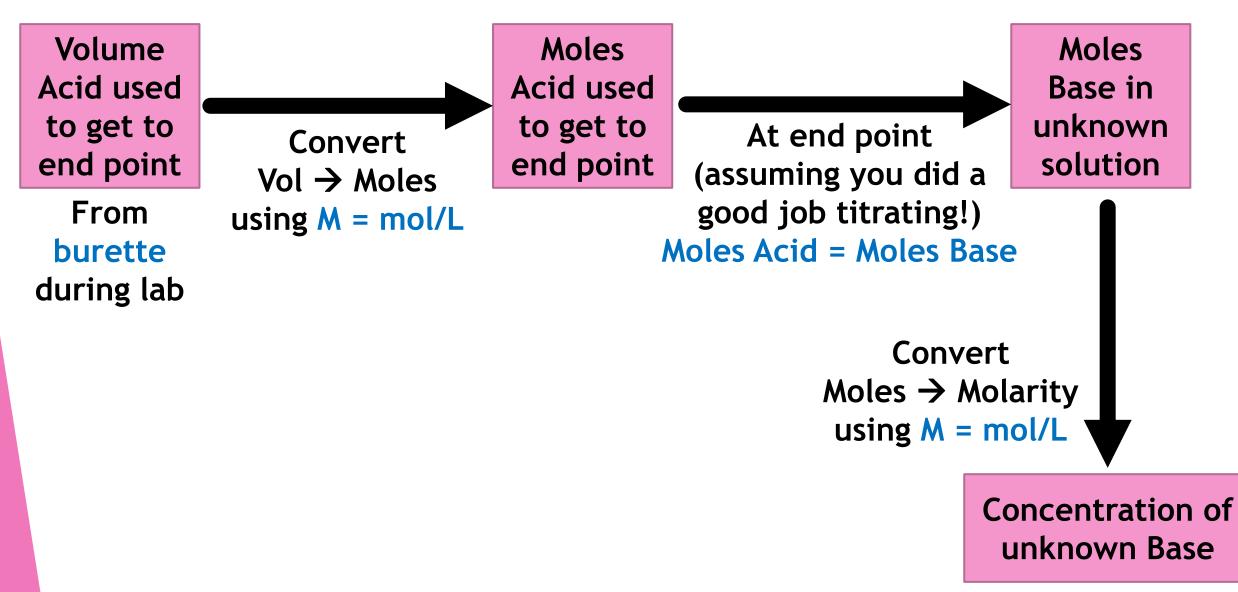
## Careful! Don't go too fast!



## **So...** Known [base] & unknown [acid]



## **So...** Known [acid] & unknown [base]



## Some things to be careful of...

- mL versus L
- Stoichiometry is it a 1:1 ratio H<sup>+</sup> : OH<sup>-</sup> ? Or is it 1:2, or 2:1, or 2:3, etc 1 mol NaOH = 1 mol OH<sup>-</sup> 1 mole Ca(OH)<sub>2</sub> = 2 mol OH<sup>-</sup>
- End point and equivalence point are only identical if your titration is absolutely perfect. It never is, there are lab errors!

# Lab Activity Portion of Lecture

## Instead of a practice problem just on paper, we are going to have our practice problem be an actual titration! On WS #14

#### **Question #1:** What is our titrand?

### **Question #2:** What is our titrant?

## Titrand = HCI – unknown [] Titrant = NaOH – known [], 0.10 M



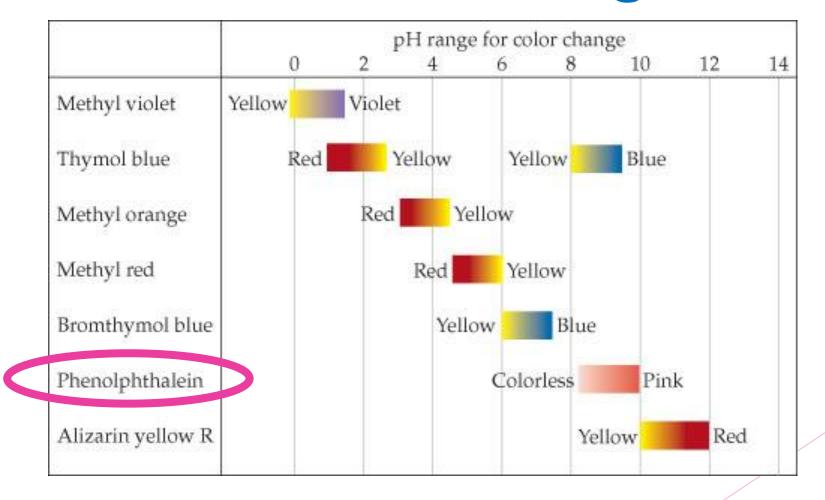
- 1) Make sure everything is rinsed with distilled water, and then rinse the burette with the titrant (NaOH).
- 2) Clamp burette into burette clamp onto a ring stand
- 3) Fill burette with NaOH with a known concentration (0.10 M)
- 4) Put a beaker under burette and slowly open valve, letting some NaOH out until bottom of meniscus is reading at an easy to read value. Careful! Make sure that the entire tip of the burette is filled with NaOH.

# Lab Set Up continued...

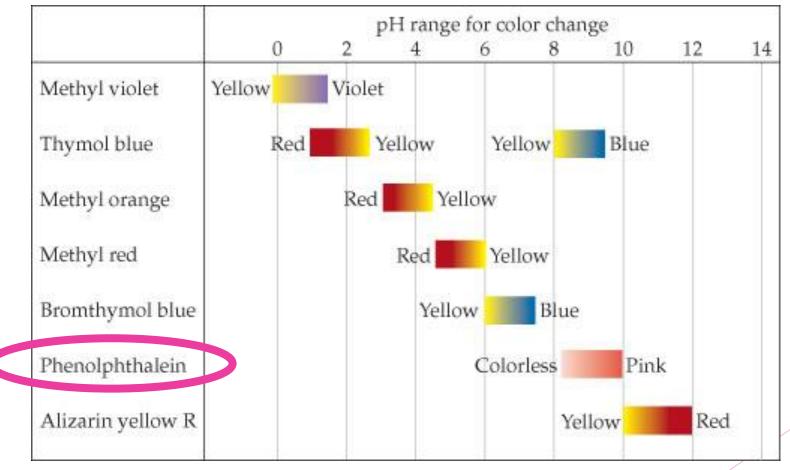
- 5) Using a graduated cylinder, put a known volume of your titrand (unknown concentration HCL) into an Erlenmeyer flask.
- 6) Add a small amount of appropriate indicator to the flask (phenolphthalein).



# **Question #3:** Which indicator should we pick? Our unknown will be in the 7-10 range.



## <u>Question #4:</u> What color shift do you expect to see for this indicator? From $\rightarrow$ \_\_\_\_.



# How many trials to do?

Do **FOUR** trials (typically)

1<sup>st</sup> Trial – Rough trial – "Quick and dirty" Just a rough estimate so you have an idea of when you need to start slowing down. DON'T include this trial when averaging your data!

2<sup>nd</sup> – 4<sup>th</sup> Trials – Real ones – Be careful!

## Set up your Data Table

Titration of Unknown HCl Solution with Phenolphthale						
[ ] of Titrant:	Vo	lume of Titra	nd Used:			

		Rough Trial	Trial #1	Trial #2	Trial #3
	Name				
	Burette Starting Volume (mL)				
	Burette Ending Volume (mL)				
	Volume of Titrant Used (mL)				

**Every group** needs to do one rough trial, and each person will do a real trial. Four people in a group = Four trials. Make the right number of columns in your table!

## Set up your Data Table

	Titration of Unknown HCl Solution with Phenolphthalein						
	[ ] of Titrant: 0.10 M			Volume of Titrand Used: 5 mL			
	Rough Trial Trial #		#1	Trial #2	Trial #3		
Name							
Burette Starting Volume (mL)							
Burette Ending Volume (mL)							
Volume of Titrant Used (mL)							

## **Titration Lecture Videos**

- 1) What is a titration? FuseSchool https://youtu.be/tlbD8MG1qMM
- 2) Setting up and Performing a Titration. CarolinaBiological https://youtu.be/sFpFCPTDv2w
- 3) How to Prepare a Burette for a Titration. Wits University https://youtu.be/Lr1nLTCqZvM
- 4) How to Read the Volume off a Burette. Wits University https://youtu.be/qdmp4\_Nwd-Q
- 5) What is a Titration and how is it performed? Wits University https://www.youtube.com/watch?v=YqfvRBJ-iPg
- 6) Acid Base Equilibrium. Bozeman Science <u>https://youtu.be/l5fk7HPmo5g</u>

# Set up your lab station

### Burette's are already clamped in for you, and filled with NaOH – not super safe to fill them, so I did it for you.

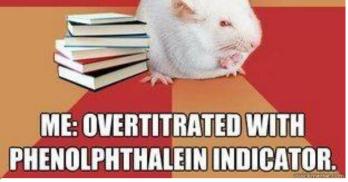
## **Perform Rough Trial**

		Titration of Unknown HCl Solution with Phenolphthalein						
		[ ] of Titrant: 0.20 M			Volume of Titrand Used: 10 mL			
		Rough Trial Trial #			Trial #2	Trial #3		
	Burette Starting Volume (mL)							
	Burette Ending Volume (mL)							
	Volume of Titrant Used (mL)							

## Each person does their trial

	Titration of Unknown HCl Solution with Phenolphthalein						
	[ ] of Titrant: 0.20 M			Volume of Titrand Used: 10 mL			
	Rough Trial	#1	Trial #2	Trial #3			
Burette Starting Volume (mL)							
Burette Ending Volume (mL)							
Volume of Titrant Used (mL)							

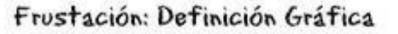
#### NORMAL PERSON: WOW PRETTY PINK COLOR

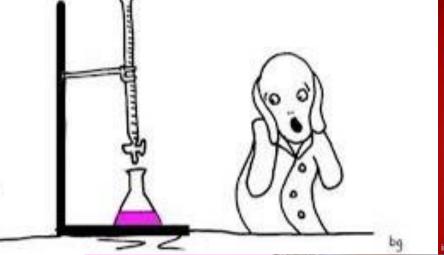


When your titration turns bright pink, and the professor starts walking towards you...

r,











The perfect game doesn't exiCalculate how many moles of **NaOH** you used **From burette** Molarity = Moles / Liters in the lab Moles NaOH used = Volume used x Molarity NaOH Known []  $Moles = L \times mol$ given to you Moles NaOH = x mL 11 Y mol 1000 mL 1 L

Calculate the unknown concentration of the acid

At End Point  $\rightarrow$  Moles NaOH = Moles HCI

Molarity = Moles / Liter

Same as mol NaOH used!

Molarity Acid = Moles Acid Liters Acid Used

> The amount in your Erlenmeyer flask!

Come check what your unknown concentration of Acid was!

Calculate the % error for each person's acid – let's see which person had the most accurate titration per group!

Average your group member's answers together – report % error of averaged data on the whiteboard – let's see which group had the best titration skills!

#### YouTube Link to Presentation

https://youtu.be/6owm822vyhl