

## \*General chem\*

Ch. 1

Luminous intensity  $\xrightarrow{\text{units}}$  Candela (cd)

exa	$10^{18}$
peta	$10^{15}$
terra	$10^{12}$
hecta	$10^2$
deka	$10^1$
atto	$10^{-18}$

جوابات مختبری S.F کی  
1.00 تریکوں کی وجہ سے اسیلی  
S.F = infinite number of SF

\*Rules for counting Significant figures :-

1. Non zero integers count as significant figures ( $45 \times 10^{-3} \rightarrow 2 \text{ S.F}$ )
2. Zeros:-
  - a. leading zeros  $\rightarrow$  not count (zeros at the left side from the non zeros)
  - b. captive zeros  $\rightarrow$  count (between non zero)  $6.0048 \rightarrow 2 \text{ S.F}$
  - c. Trailing zeros  $\rightarrow$  count if the number contains decimal point (at the right end of the number)  
 $9000. \rightarrow 4 \text{ S.F}$  /  $0.300 \rightarrow 3 \text{ S.F}$  /  $150 \rightarrow 2 \text{ S.F}$

\* Accuracy:- related to how much your figures closed to correct value  
\* precision :- " " " " " " to each other

\* Temperature :-

$$T_K = T_C + 273.15 \quad / \quad T_C = \frac{5}{9} (T_F - 32)$$

## ch.2

- atom : electrons + protons + neutrons

- nucleus :- small / extremely dense : account for almost all of the atom's mass

- isotopes :- same number of protons but different numbers of neutrons

↳ identical chemical properties because the chemistry of an atom is due to its valence ( $e^-$ )

النواة في الجدول الدوري لا يغير وزنها بل يغير وزن الذرة

average atomic mass =  $(\text{atomic mass of isotope}_1 \times \text{percentage}) + \dots + (\text{atomic mass of isotope}_n \times \text{percentage}_n)$

- Atomic Number  $\Rightarrow Z = P$

- Mass Number / atomic mass  $\Rightarrow A = P + N$

- إذا طلب السؤال إنه أوجد الـ  $Z$  فهو العنصر من الجدول الدوري فتبينت عـ

$Z$  ليس عدد



- Chemical Bonds :-

1. covalent bond :- between atoms by sharing electrons to form molecules (none of atoms should be metal) ( $H_2O$ )

2. ionic bond :- force of attraction between oppositely charged ions ( $NaCl$ )

$N_2 / Cl_2 / Co \rightarrow$  diatomic molecule.

$NO / CO \rightarrow$  heterogeneous diatomic molecule.

$Cl_2 / O_2 / N_2 \rightarrow$  homogeneous " "

$H_2O \rightarrow$  heterogeneous molecule

metal  
cation : positive ion  
non-metal  
anion : negative ion

## \*Periodic table :-

-Groups/Families :- elements in the same vertical columns ; have similar chemical properties

e.g. → Alkaline metals / Alkaline earth metals / Halogens / noble gases

- periods :- horizontal rows of elements

## metals :

(representative + transition + lanthanides + actinides) elements

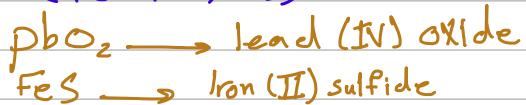
# \* Naming compounds :-

- Binary compound :- Composed of two elements

1. Binary ionic compounds :- metal + non-metal → Type I and II  
 - Ionic compounds with polyatomic ions

\* Type I :-  $\text{NaCl} \rightarrow \text{Sodium chloride}$

\* Type II :- metals in these compounds form more than one positive charge ( $\text{Fe}^{+2}, +3$ )



\* Ionic compounds with polyatomic ions

Name	Formula	Name	Formula
ammonium	$\text{NH}_4^+$	phosphite	$\text{PO}_3^{3-}$
nitrite	$\text{NO}_2^-$	phosphate	$\text{PO}_4^{3-}$
nitrate	$\text{NO}_3^-$	hydrogen phosphate	$\text{HPO}_4^{2-}$
sulfite	$\text{SO}_3^{2-}$	dihydrogen phosphate	$\text{H}_2\text{PO}_4^-$
sulfate	$\text{SO}_4^{2-}$	perchlorate	$\text{ClO}_4^-$
hydrogen sulfate (aka: bisulfate)	$\text{HSO}_4^-$	chlorate	$\text{ClO}_3^-$
thiosulfate	$\text{S}_2\text{O}_3^{2-}$	chlorite	$\text{ClO}_2^-$
oxalate	$\text{C}_2\text{O}_4^{2-}$	hypochlorite	$\text{ClO}^-$
hydroxide	$\text{OH}^-$	bromate	$\text{BrO}_3^-$
IUPAC Name	older name	Name	Formula
copper(I)	cuprous	acetate	$\text{CH}_3\text{COO}^-$
copper(II)	cupric	acetato	$\text{C}_2\text{H}_3\text{O}_2^-$
iron(II)	ferrous	carbonate	$\text{CO}_3^{2-}$
iron(III)	ferric	hydrogen carbonate (aka: bicarbonate)	$\text{HCO}_3^-$
lead(II)	plumbous	chromate	$\text{CrO}_4^{2-}$
lead(IV)	plumbeous	dichromate	$\text{Cr}_2\text{O}_7^{2-}$
mercury(I)	mercurous	permanganate	$\text{MnO}_4^-$
mercury(II)	mercuric	peroxide	$\text{O}_2^{2-}$
tin(II)	stannous	cyanide	$\text{CN}^-$
tin(IV)	stannic	cyanate	$\text{OCN}^-$
		thiocyanate	$\text{SCN}^-$

\* Binary covalent compounds (Type III) :- formed between 2 non-metals

## Ch. 3

- the reference for atomic masses is  $^{12}\text{C}$  because its atomic mass is exactly 12 a.u

- the ratio of the mass of an isotope to  $^{12}\text{C}$  by mass spectrometer

$$\text{1 mole of substance} = 6.022 \times 10^{23} \text{ atoms}$$

(an arrow points from the 6.022 to the text "avogadro's number")

$$1 \text{ mole (C)} = 6.022 \times 10^{23} \text{ atoms} = 12.01 \text{ g (C)}$$

mass for 1 mole of substance = atomic mass

$$1 \text{ mole of an element} = 6.022 \times 10^{23} \text{ atoms} = \text{atomic mass}$$

for elements  $\rightarrow n = \frac{\text{mass}}{\text{atomic mass}}$

$$1 \text{ mole of a compound} = 6.022 \times 10^{23} \text{ molecules} = M.M$$

For molecule  $\rightarrow n = \frac{\text{mass}}{M.M}$  Molar Mass

$$\% \text{ mass} = \frac{\text{mass of element in compound}}{\text{mass of compound}} \times 100\%$$

**Exercise:** calculate the mass percent of ethanol ( $\text{C}_2\text{H}_5\text{OH}$ )?



$$M.M (\text{C}_2\text{H}_5\text{OH}) = 2(12.01) + 6(1) + 16 = 46.022$$

$$\text{mass} (\text{C}_2\text{H}_5\text{OH}) = 46.022 \times 1 = 46.022 \text{ g}$$

$$M.M (\text{O}_2) = 16 \times 2 = 32 \rightarrow \text{mass} = 32 \times 3 = 96$$

$$\text{mass of compound} = 46.022 + 96 = 142.022$$

$$\% \text{ mass} (\text{C}_2\text{H}_5\text{OH}) = \frac{\text{mass} (\text{C}_2\text{H}_5\text{OH})}{\text{mass of compound}} \times 100\%$$

$$= \frac{46.022}{142.022} \times 100\% = 32.405\%$$

imp. note :- coefficients can not be fractions , although they are usually given as lowest integer multiples.

mass of reactants = mass of products

- Stoichiometric mixing :- all reactants are consumed and converted into products
- non - stoichiometric mixing :- limiting reactant

when the reaction is 1:1 then the lower number of moles is the L.R  
but when the reaction is not 1:1 then simple calculation is needed to find out the L.R



percentage yield :- an important indicator of the efficiency of a particular laboratory or industrial reaction.

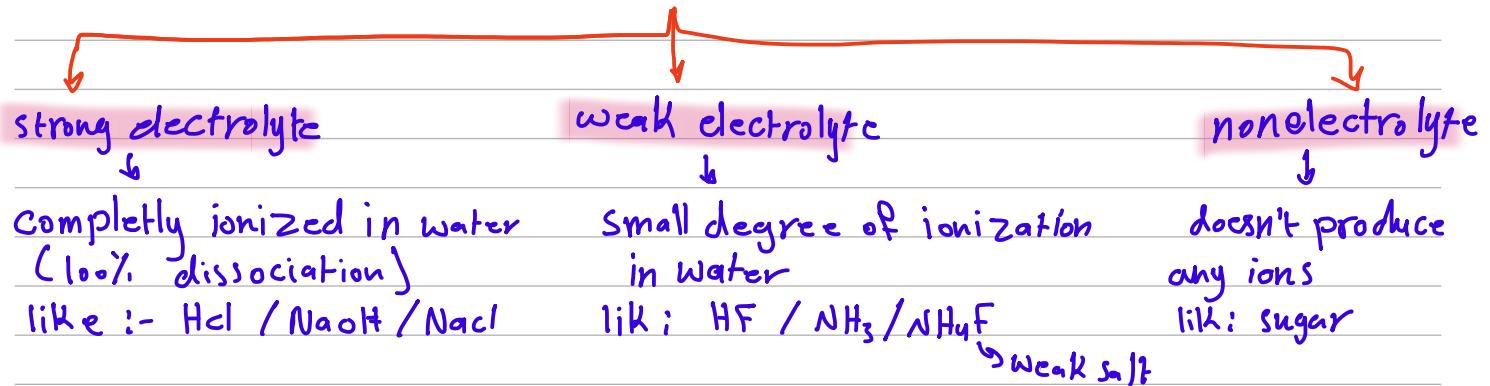
→ الفعل العملي

$$\text{Percent yield} = \frac{\text{actual yield}}{\text{theoretical yield}} \times 100\%$$

↓ من التجربة (المختبر)

## Ch 4

\* **electrolyte :-** Substance that when dissolved in water produces a solution that conducts electricity ,  
↳ depends on the amount of ions



Dilution :- adding water to a concentrated or stock solution to achieve the molarity desired for a particular solution .

$$M = \frac{n}{V}$$

∴ dilution of water does not alter the number of moles of solute .

Moles of solute before dilution = moles of solute after dilution

$$\begin{aligned} n &= n \\ M_i V_i &= M_f V_f \end{aligned}$$

- Types of chemical reactions :-

1. Precipitation reactions :- one of the products is insoluble
2. Acid - Base reactions :- neutralization reactions ( अम्ल + स्लॉ गुणी)
3. Oxidation - Reduction reactions (Redox reactions) :- involve electron transfer

- ∵ the number of ions ↗

number of ions = number of moles × avogadro's number

## \* Rules for Solubility :-

1.  $\text{NO}_3^-$  /  $\text{NH}_4^+$  / alkaline metal (group 1A)  $\rightarrow$  salts are soluble.
2.  $\text{Cl}^-$  /  $\text{Br}^-$  /  $\text{I}^-$   $\rightarrow$  salts are soluble.  
except:  $(\text{Ag}^+, \text{Pb}^{+2}, \text{Hg}^{+2})$
3. Most sulfate ( $\text{SO}_4^{-2}$ ) salts are soluble.  
except:  $\text{BaSO}_4, \text{PbSO}_4, \text{Hg}_2\text{SO}_4, \text{CaSO}_4$
4.  $\text{BaCrO}_4$  insoluble
5. Most  $\text{OH}^-, \text{S}^{+2}, \text{CO}_3^{-2}, \text{CrO}_4^{-2}, \text{PO}_4^{-3}$  salts are insoluble  
except:- (group 1A,  $\text{NH}_4^+$ )  $\rightarrow$  in rule number ①

Solid + liquid  $\rightarrow$  ionic equation  $\downarrow$  (in case of  
معزل عن الماء)

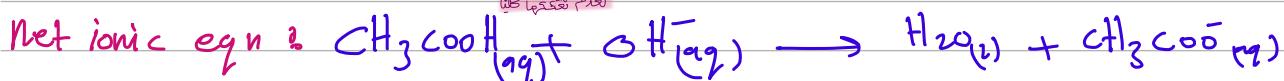
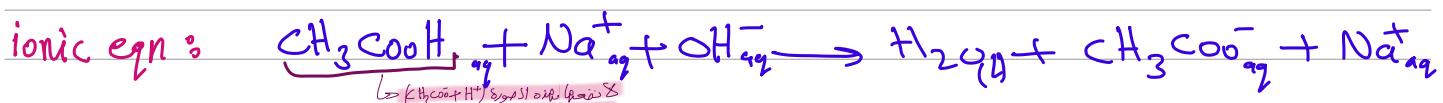
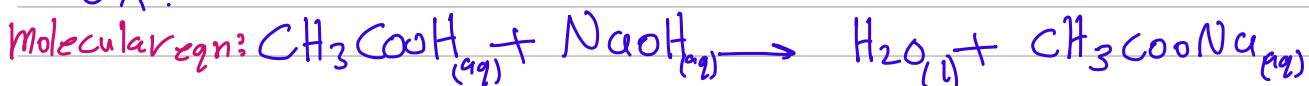
(الذريون الغير مشارك) spectator ions  $\downarrow$  الباقي Net equation  $\downarrow$  (in  
ionic)

\* For a strong acid / strong base reaction the net ionic equation  
is always:



\* Weak acid/strong base or weak base/strong acid reactions :-

Ex.



## Rules for Assigning Oxidation States

Oxidation state of an atom in an element = 0

Oxidation state of monatomic ion = charge of the ion

Oxygen = -2 in covalent compounds (except in peroxides)

where it = -1  $\text{H}_2\text{O}_2$

Hydrogen = +1 in covalent compounds (except in hydrides, -1)

Fluorine = -1 in compounds

Sum of oxidation states = 0 in neutral compounds

Sum of oxidation states = charge of the ion in ions

$\boxed{\text{LiH, NaH}}$ ,  $\text{H}_2\text{CO}_3$ ,  $\text{MnO}_4^-$



- \* Oxidation :- increase in oxidation state , it happens by loss of electrons  
(Reducing agent)
- \* Reduction :- decrease " " " , " " " gain of electron  
(Oxidizing agent)

## Ch.5

- the atmospheric pressure is measured by barometer
- the pressure of a gas confined in a container is measured by manometer (car tire , home gas cylinder)

### \* Properties of gases :-

- 1- Uniformly fill any container and take its shape.
- 2- easily compressed .
- 3- Mixes completely with any other gas .
- 4- Exerts pressure on its surroundings .

$$P \rightarrow N/m^2 = \text{pascal (pa)}$$

$$P = \frac{\text{Force}}{\text{area}}$$

$$\begin{aligned} 1 \text{ atm} &= 101.325 \text{ kPa} = 101.325 \text{ Pa} = 1.01325 \text{ bar} = 760 \text{ mmHg} \\ &= 760 \text{ torr} = 14.7 \text{ lb/in}^2 \end{aligned}$$

### \* Ideal gas law :-

$$PV = nRT, R = 0.08206 \text{ L.atm/mole.K}$$

(constant)

معكث تختبر عن صحة المعرفة  
وهي بمعنى كل وحدة (P) هي

$$\frac{P_i V_i}{T_i} = \frac{P_f V_f}{T_f}$$

### - Standard Molar volume of an ideal gas (SMV) :-

for 1 mole of an ideal gas at (0)°C and 1 atm , the volume of the gas is 22.42 L

- STP :- 0°C , 1 atm

$$\begin{aligned} \text{density} &= d = \frac{\text{mass}}{\text{volume}} \\ d &= \frac{m}{V} = \frac{P}{RT} \end{aligned}$$

\* the total pressure exerted is the sum of the pressure that each gas would exert if it were alone under the same conditions of  $N, T, n$ .

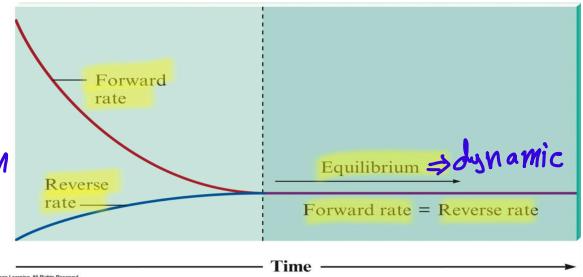
$$P_{\text{total}} = P_1 + P_2 + P_3 \quad \begin{array}{l} \text{في حالات مجموعات الغازات} \\ \text{في نفس المكان} \end{array}$$

## Ch. 13

\* After the equilibrium is reached, none of the reactants or products has a concentration of zero

\* Chemical Equilibrium :-

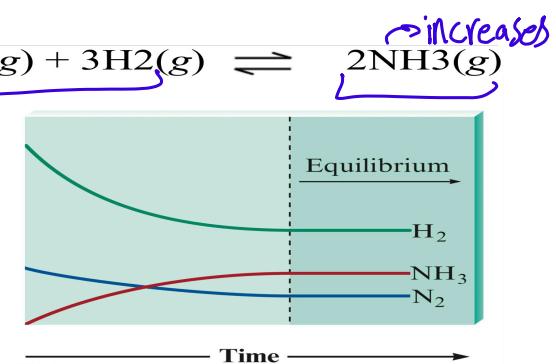
A state of the system where the concentration of all products and reactants remain constant with time.



concentration decreases

$$K = \frac{[C]^m [D]^n}{[A]^j [B]^k}$$

equilibrium expression



$$K' = \frac{1}{K} \quad \text{عند جمعيتي المقادير} \quad \Rightarrow \quad *$$

-When a balanced equation for a reaction is multiplied by a factor of ( $n$ )

$$K' = K^n$$



$$K_p = \frac{P_z^z \cdot P_w^w}{P_x^x \cdot P_y^y} = K (RT)^{\Delta n_g}, \quad \Delta n_g = n_p - n_r$$

بس في حالات الظروف الغازية

$$\Delta n_g = 0 \Rightarrow K_p = K$$

$$\Delta n_g > 0 \Rightarrow K_p > K$$

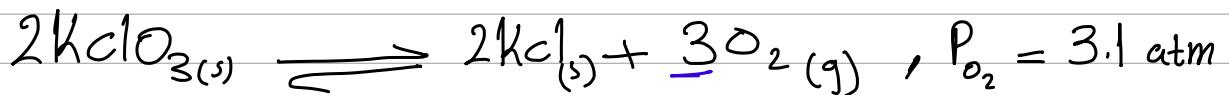
$$\Delta n_g < 0 \Rightarrow K_p < K$$

- Homogeneous equilibria :- involve the same phase.

(s, g, aq) تكون المكونات في نفس الحالة

- Heterogeneous equilibria :- involve more than one phase.

\* the position of a heterogeneous equilibrium doesn't depend on the amount of pure solid or liquids present because their concentrations are constant.



$$K_p = P_{\text{O}_2}^3, \Delta n_g = 3, K_p > K, K = [\text{O}_2]^3$$

\*  $K > 1$  means that at equilibrium the reaction system consist of mostly products or the equilibrium lies to the right.

\*  $K < 1$  means that " " " " " " mostly reactants or " " " " to the left.

- Reaction Quotient ( $Q$ ) :- used when all of the initial concentration are non-zero.

\*  $Q = K$  (just at equilibrium), no shift will occur

\*  $Q > K \rightarrow$  the system shifts to the left. After equilibrium consuming products and forming reactants, until equilibrium is achieved

\*  $Q < K \rightarrow$  " " " " right. Before " " reactants " " products " " "

-  $K$  will change depending upon the temperature.

\* Endothermic reaction :- energy is a reactant,  $K$  increase, shift to right

\* Exothermic reaction :- " " " " a product,  $K$  decrease, shift to left  
at least

- Pressure effect  $K$  when one of the reactants is gas

- Addition of inert gas doesn't affect the equilibrium position.

## Ch. 14

### - Arrhenius :-

- \* acids :- produce  $H^+$
- \* bases :- produce  $OH^-$

### - Bronsted - Lowry :-

- \* acids :- proton ( $H^+$ ) donor
- \* bases :- proton ( $H^+$ ) acceptor

$\begin{matrix} HA & A^- \end{matrix}$   
 acid / base conjugate pair  
 Conjugate acid / base pair are related  
 by one proton transfer

### - Lewis :-

- \* acids :- electron pair acceptor
- \* bases :- electron pair donor

### - Strong acid :-

Ionization equilibrium lies far to the right.  
 Yields a weak conjugate base.

### - Strong base :-

يُنْسَى جَدِيداً جَدِيداً  
 يُنْسَى الْفَكِيرَةُ  
 yield a weak conjugate acid.

### - Weak acid :-

يُنْسَى الْعَنْتَرَةُ

Ionization equilibrium lies far to the left.

The weaker the acid - the stronger its conjugate base

\* Water is amphoteric (auto ionization) :- behaves either as an acid or as a base.

$$K_w = [H^+][OH^-] = 1 \times 10^{-14} \text{ at } 25^\circ C$$

- \*  $[H^+] = [OH^-] \Rightarrow pH = 7$  (neutral)
- \*  $[H^+] > [OH^-] \Rightarrow pH < 7$  (acidic)
- \*  $[H^+] < [OH^-] \Rightarrow pH > 7$  (basic)

- If the equilibrium lies to the right ,  $K_a / K_b > 1$

إذَا تَكُونُ النَّوَاطِقُ بِحُوْجَنِ اعْلَى

- ↪ ↪ ↪ ↪ ↪ left ,  $K_a / K_b < 1$

إذَا تَكُونُ النَّوَاطِقُ بِعُوْجَنِ اعْلَى

$$K_a = \frac{[H_3O^+][A^-]}{[HA]}, \quad K_b = \frac{[OH^-][BH^+]}{[B]}$$

- The number of decimal places in the log is equal to the number of Significant figures in the original number.

$$[\text{H}^+] = 1.0 \times 10^{-9} \text{ M}$$

2 S.F

$$\text{pH} = 9.00$$

$$\text{pH} = -\log [\text{H}^+] , \text{ pOH} = -\log [\text{OH}^-] , \text{ pH} + \text{pOH} = 14$$

2 decimal places

\* إذا سأله عن مكونات محلول لحمض قوي / قاعدة قوية يبعاون

- في حالة الحمض القوي :- القاعدة المعرفة / (المحض)  $\text{H}^+ / \text{H}_2\text{O}$  - في حالة القاعدة القوية: (الحمض)  $\text{OH}^- / \text{H}_2\text{O}$

\* Two factors for acidity in binary compounds :-

1- Bond polarity : كل ما زادت الخطبية (الفرق في الكهروستاتية) بزيادة قوة الحمض

2. Bond strength : كل ما زادت بتعل قوة الحمض لمlein الحمض  
العوبي مثل HCl يتكون الرابطة ما بين  $\text{H}^+$  و  $\text{Cl}^-$  ضعيفة جداً فبسهل من تفكيكه

\* Oxyacids :-  $\text{H}-\text{O}-\text{X}$  ( $\text{HClO}$ ,  $\text{HBrO}$ )

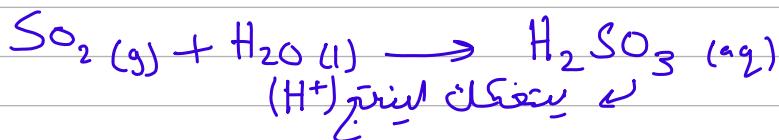
- كلما زادت عدد ذرات O بزيادة قوة الحمض (لأنه O<sub>2</sub> بتعل على زيادة الخطبية)  
- كلما زادت قدرة X على انتهاج ذب ازداد نحوها بزيادة قوة الحمض (لأنه يودي إلى زيادة الفرق في الكهروستاتية)

\* Oxides :-

فقط من دخ

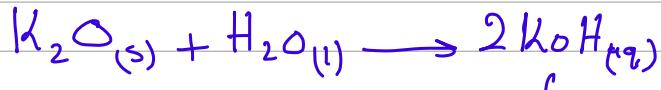
- Acidic oxides (acid Anhydrides) :-  $\text{O}-\text{X}$  is strong and covalent. ( $\text{SO}_2$ ,  $\text{NO}_2$ ,  $\text{CO}_2$ )

- يتفاعلوا مع  $\text{H}_2\text{O}$  فبنيكون حمض و هو الذي يجعل العامل حمسي



- Basic oxides (basic Anhydrides) :- O-X bond is ionic has a very low electronegativity (K<sub>2</sub>O, CaO)

- يتفاعل مع H<sub>2</sub>O وينتج OH<sup>-</sup> و يجعل المحلول قاعدي



يُنتج OH<sup>-</sup>



- Buffer Solutions resist a change in pH

Buffer :- weak acid and its conjugate base  
weak base and its conjugate acid

- Henderson-Hasselbatch equation :-

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$pOH = pK_b + \log \frac{[BH^+]}{[B]}$$

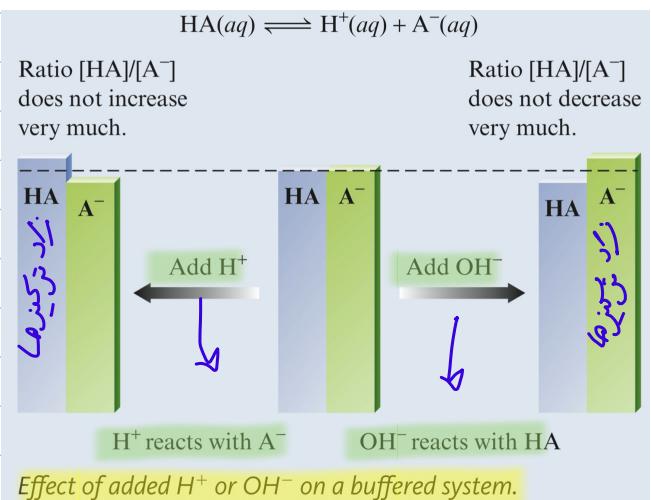
\* تتحدد pH buffer على النسبة بين الحمض الضعيف وقاعدته المرفقة وبين قيم pH ثابتة ماء الأمونيوم هذه النسبة ثابتة (إذا كان تركيز مكونات buffer من ترتكب H<sup>+</sup> أو OH<sup>-</sup> أكثاف)

- Titration Curve

\* Equivalence (Stoichiometric) point :-

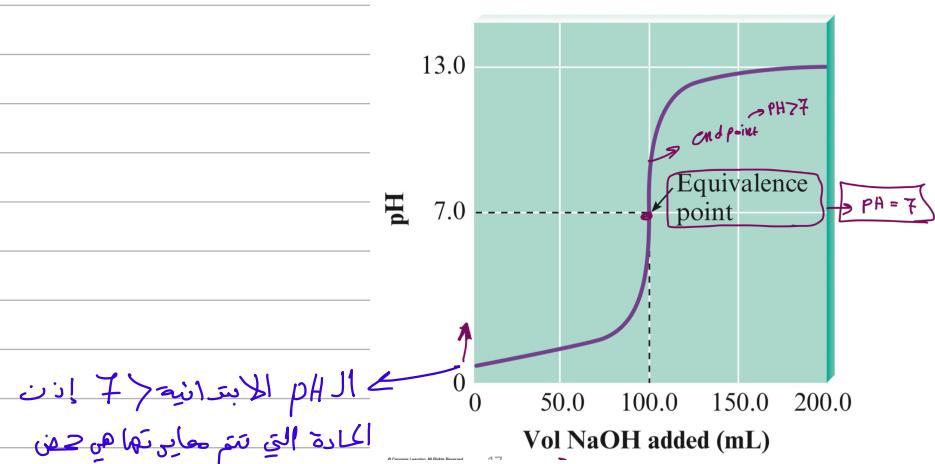
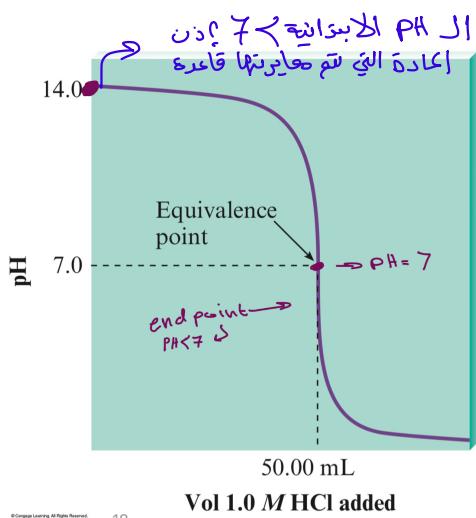
النقطة التي تتعادل فيها جمجمة مول من إلكترافاف (سوار حمض / قاعدة) مع الحمض / القاعدة التي بدنا نفاخوها و تكون pH = 7 (في حال كانت المعايرة بين حمض قوي و قاعدة قوية)

\* end point :- هي التي يحدد انتهاء المعايرة وهي تتحدد من قربها خصوبة من نقطة التكافؤ و تقع على المعايرة



- معايرة إضافية :-

- عند معايرة حمض ضعيف بقاعدة قوية تكون الـ pH عن نقطة التكافؤ > 7
- عند معايرة قاعدة ضعيفة بحمض قوي تكون الـ pH عن نقطة التكافؤ < 7



phenolphthalein →

\* Methyl orange: yellow in basic  
red in acidic

