

## Tips & Trick

Remember: That For Strong Acid  $K_a_1 \gg \gg K_{a_2}$   
So it is always 100% dissociation of first  $H^+$ .

Remember these formulas like this for salt hydrolysis:

$$S \cdot A + W \cdot B \Rightarrow \left[ pH = 7 - \frac{1}{2} pK_b - \frac{\log c}{2} \right]$$

it is S. Acid so  $[pH < 7] \Rightarrow$  we use '-' sign in all terms  
& here the basic mol. is weaker & so we use  
 $pK_b$  as base. Hydrolysis is alone.

Similarly for  $W \cdot A + S \cdot B \quad pH > 7$

$$\left( pH = 7 + \frac{1}{2} pK_a + \frac{\log c}{2} \right)$$

Here Acidic part hydrolysis as it is weak  
so we use  $pK_a$ .

for  $S \cdot A + S \cdot B \Rightarrow$  It's True we all know.

for  $(W \cdot A + W \cdot B)$  it will be ~~diss.~~ of  $pK_a + pK_b$   
added to  $\textcircled{7}$ . It depends on  $pK_a + pK_b$   
for soln to be Acidic / Basic

$$\left( pH = 7 + \frac{1}{2} (pK_a - pK_b) \right)$$

Always Remember  $\left( K_h = \frac{10^{-14}}{K_a} \right)$  & H is subs. for

& in salt hydrolysis & rest procedure  
is same.

Remember  $1 - \alpha \approx 1$  only if  $(\alpha < 6.4 \times 10^{-4})$  i.e.  $\textcircled{0.1}$

$$\left( pH + pOH = 14 \right)$$