

# ARJUNA

## JEE AIR 2024

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**Physical Chemistry**

### Ionic Equilibrium



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Lecture No. - 09



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# Topics to be Covered

**Topic**

✓ Solubility (S)

**Topic**✓ Solubility product ( $K_{SP}$ )

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Solubility : "Amount of substance dissolved in  
per L solution"

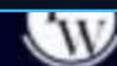
If represent as "S"

# If amount is taken in terms of mole,

Unit of Solubility "M"

# If amount is taken in terms of "g",

Unit of Solubility : "g/L"



# Types of salt (on the basis of solubility)

1. Soluble salt :

$$S > 0.1 \text{ M}$$

2. Partial Soluble Salt :

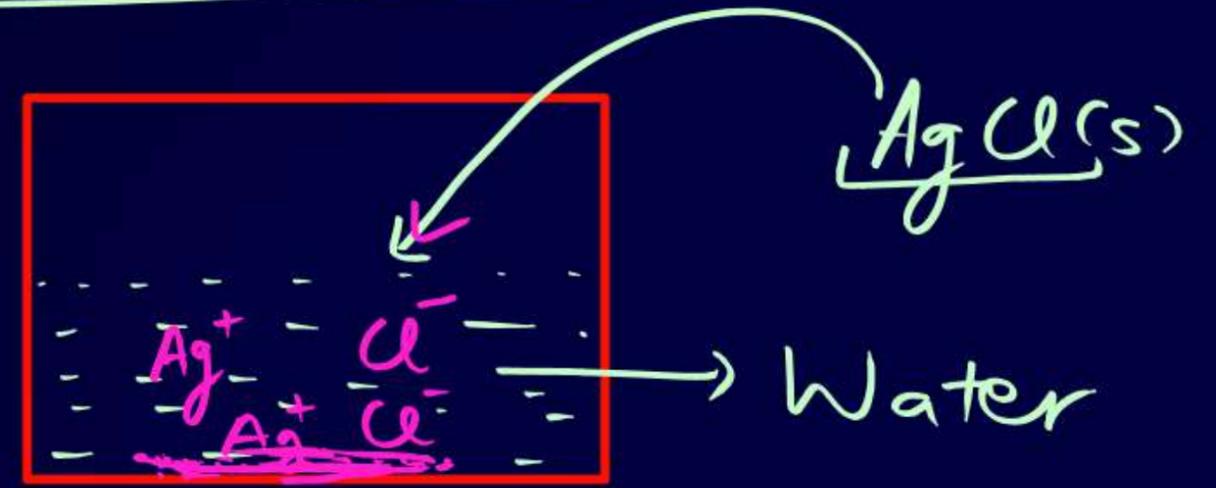
$$S = (0.1 \leftrightarrow 0.01) \text{ M}$$

3. Sparingly Soluble Salt (SSS) :

$$S < 0.01 \text{ M}$$



# Sparingly Soluble Salt:



$$K_{sp} = [Ag^+(aq)] [Cl^-(aq)]$$



At equilibrium state

Rate of forward direction = Rate of backward direction

Precipitate: PPT

formation of ppt



$$Q = \text{Reaction Quotient} = \underbrace{[Ag^+(aq)] [Cl^-(aq)]}$$

Equilibrium Constant  $\leftarrow K_{sp} = \text{Solubility product.}$

(i) If  $Q < K_{sp}$ , Reaction shift to forward,

Unsaturated Solution

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(ii) If  $Q = K_{sp}$ , Reaction is in equilibrium state  
ppt formation will start.

(iii) If  $Q > K_{sp}$ , Reaction shift in backward direction  
Appearance of ppt in solution



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$$Q = [A^{y+}(aq)]^x [B^{x-}(aq)]^y$$

$$K_{sp} = [A^{y+}(aq)]^x [B^{x-}(aq)]^y \rightarrow \text{Eqm state.}$$

(A<sub>x</sub>B<sub>y</sub>)



Q. If 0.1 mole of a SSS is dissolved in water.

Only 40% of salt dissolved in the solution (in 1L)

Calculate: [Provided Cation : Anion = 1 : 2]  
Assuming no hydrolysis of cation & ion.

(1) [Cation]

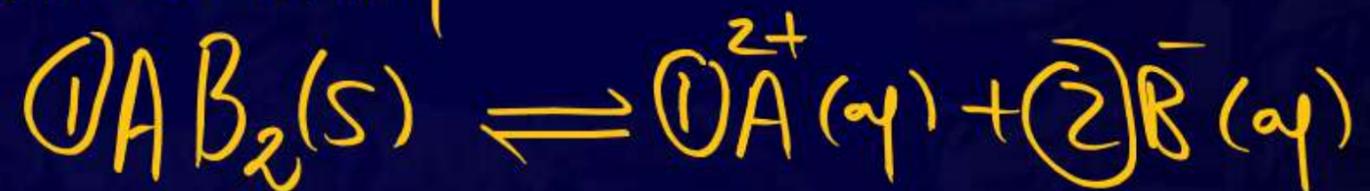
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(2) [Anion]

(3)  $K_{sp}$  of SSS

Sol<sup>n</sup>  $S = \frac{0.1 \times 40}{100}$

Let Solubility = S



$$[A^{2+}(aq)] = \frac{0.1 \times 40}{100}, \quad [B^{-}(aq)] = 2 \times \frac{0.1 \times 40}{100}$$

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$$\begin{aligned}K_{sp}(AB_2) &= (A^{+2})(B^{-1})^2 \\ &= (4 \times 10^{-2})(8 \times 10^{-2})^2 \\ &= 64 \times 4 \times 10^{-6}\end{aligned}$$

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## Relationship between Solubility and Solubility Product)

(Assuming no hydrolysis of ion)

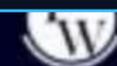


Let solubility of  $A_x B_y$  =  $S$

$$[A^{y+} (aq)] = xS, \quad [B^{x-} (aq)] = yS$$

$$K_{sp} = [A^{y+}]^x [B^{x-}]^y = (xS)^x (yS)^y = x^x y^y S^{x+y}$$

( $A_x B_y$ )



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$$\underline{K_{sp}(A_x B_y)} = x^x y^y \equiv S^{x+y} M^{x+y}$$

$$S = \left[ \frac{K_{sp}(A_x B_y)}{x^x y^y} \right]^{\frac{1}{x+y}}$$

(1) Cation : Anion  
1 : 2  
or  
2 : 1

$$K_{sp}(AB_2) = 4S^3$$

or

$$K_{sp}(A_2B)$$



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$$(2) \quad 2 : 3$$

or

$$3 : 2$$

$$K_{sp}(A_2B_3) = 108 \text{ S}^5 \text{ M}^5$$

or

$$K_{sp}(A_3B_2)$$

$$(3) \quad 3 : 1$$

or

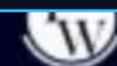
$$1 : 3$$

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$$K_{sp}(A_3B) = 27 \text{ S}^4 \text{ M}^4$$

or

$$K_{sp}(AB_3)$$



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Relation bet<sup>n</sup>  $K_{sp}$  &  $S$

Ans.

$3:1$

$$K_{sp} = 27S^4$$

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Q. If  $10^{-4} \text{ M}$   $\text{KCl(aq)}$  solution is mixed with  $10^{-6} \text{ M}$   $\text{AgNO}_3(\text{aq})$  in such a way that their volume ratio is  $\boxed{3:1}$  respectively

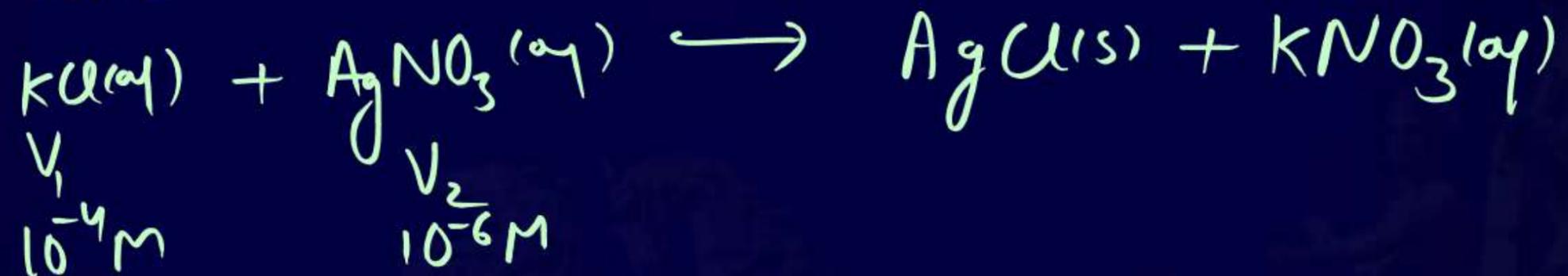
Predict whether precipitation of  $\text{AgCl}$  will occur or not

Given:  $K_{sp}(\text{AgCl}) = 10^{-10} \text{ M}^2$

$25^\circ\text{C}$

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Sol<sup>n</sup>



$$V_f = V_1 + V_2 \rightarrow \left[ \begin{aligned} [\text{Ag}^+(\text{aq})] &= \frac{10^{-6} \times V_2}{V_1 + V_2}, & [\text{Cl}^-(\text{aq})] &= \frac{V_1 \times 10^{-4}}{V_1 + V_2} \\ &= 10^{-6} \times \frac{1}{4} & &= \frac{3}{4} \times 10^{-4} \end{aligned} \right], \quad \frac{V_1}{V_2} = \frac{3}{1}$$



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1:1

$$Q = [\text{Ag}^+(\text{aq})] [\text{Cl}^-(\text{aq})] = \frac{1}{4} \times 10^{-6} \times \frac{3}{4} \times 10^{-9}$$
$$= 10^{-10} \times \frac{3}{16} < 1 \times 10^{-10}$$

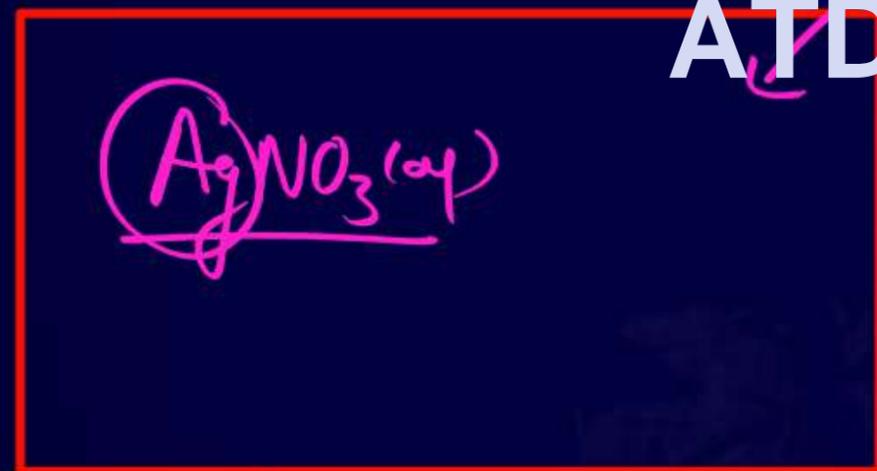
ppt will not occur

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# Calculation of Solubility in presence of Common ion: GURUKUL

Q. Calculate solubility of  $\text{AgCl}$  in  $0.1\text{M}$   $\text{AgNO}_3(\text{aq})$  solution at  $25^\circ\text{C}$ . Given  $K_{sp}(\text{AgCl}) = 10^{-10} \text{M}^2$

Sol<sup>n</sup>.

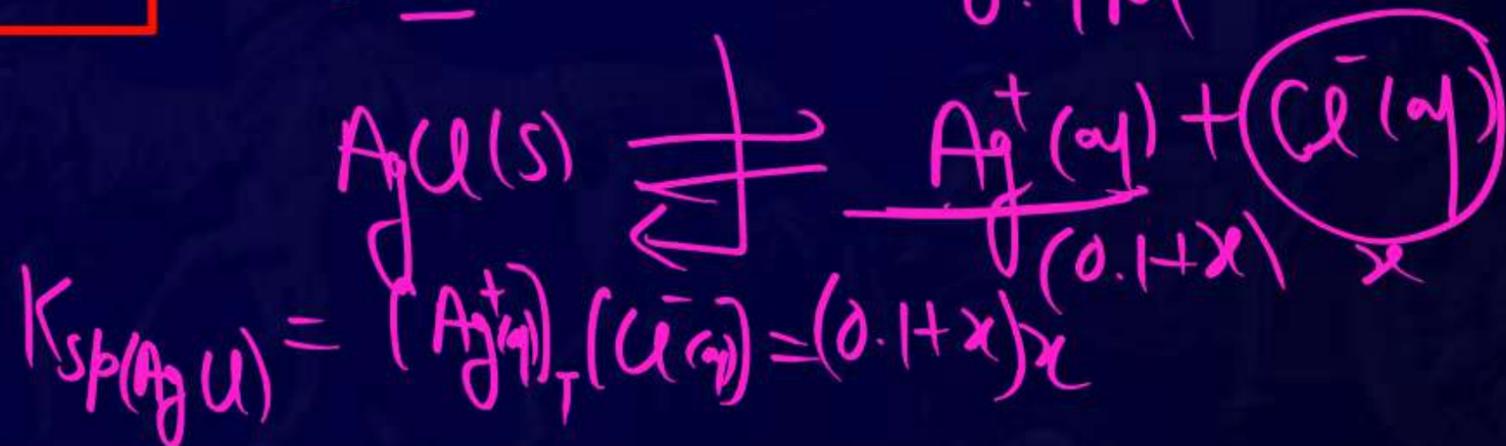


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Let solubility of  $\text{AgCl}$  in  $0.1\text{M}$   $\text{AgNO}_3 = x$



Due to pr. of C.I of  $\text{Ag}^+$ ,  
Solubility of  $\text{AgCl} \downarrow$





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$$K_{sp}(\text{AgCl}) = 10^{-10} = (0.1+x)x$$

Solubility of AgCl,  $K_{sp} = S^2$ ,  $S = 10^{-5}$   
in water

$\downarrow$   
 $10^{-10}$

$$x < S$$

due to C.I. effect.

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$$0.1+x \approx 0.1$$

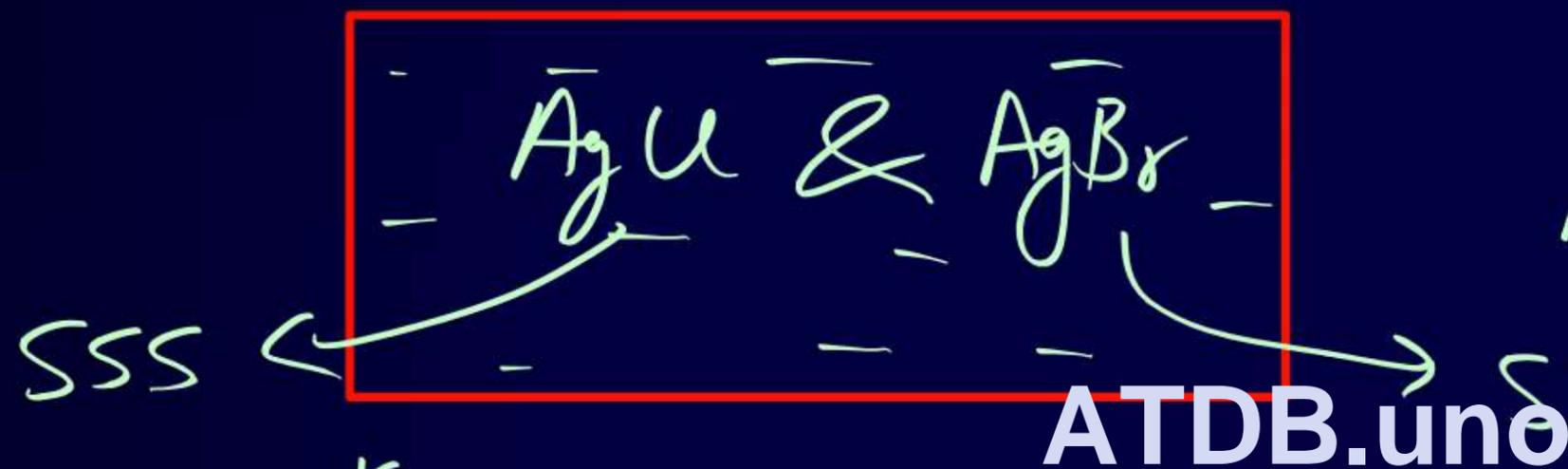
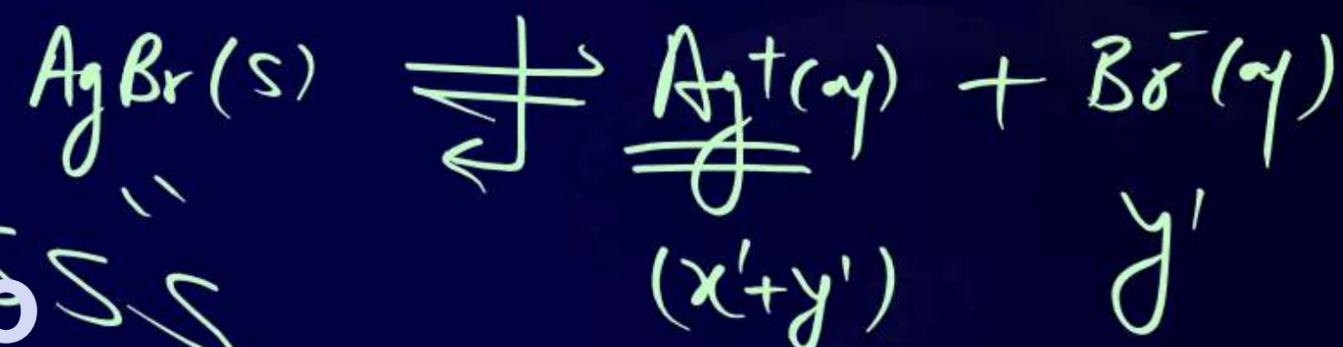
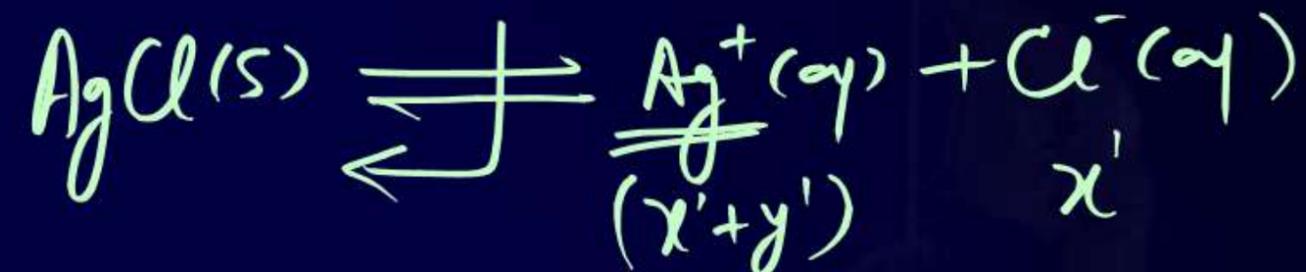
$$10^{-10} = (0.1)x$$

$$x = 10^{-9}$$

# Calculation of Solubility of SSS in presence of other SSS



Simultaneous Solubility

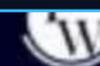


$K_{sp}(\text{AgCl})$

$K_{sp}(\text{AgBr})$

Let Solubility of AgCl in pr. of AgBr =  $x'$

Solubility of AgBr in pr. of AgCl =  $y'$



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$$\frac{K_{sp}(\text{AgCl})}{K_{sp}(\text{AgBr})} = \frac{[\text{Ag}^+]_T [\text{Cl}^-]_{aq}}{[\text{Ag}^+]_T [\text{Br}^-]_{aq}} ;$$

$$\frac{K_{sp}(\text{AgCl})}{K_{sp}(\text{AgBr})} = \frac{[\text{Cl}^-]_{aq}}{[\text{Br}^-]_{aq}} = \frac{x'}{y'} \quad \text{--- (1)}$$

$$K_{sp}(\text{AgCl}) = [\text{Ag}^+]_T [\text{Cl}^-]_{aq} = (x' + y') x' \quad \text{--- (2)}$$

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$$K_{sp}(\text{AgBr}) = [\text{Ag}^+]_T [\text{Br}^-]_{aq} = (x' + y') y' \quad \text{--- (3)}$$

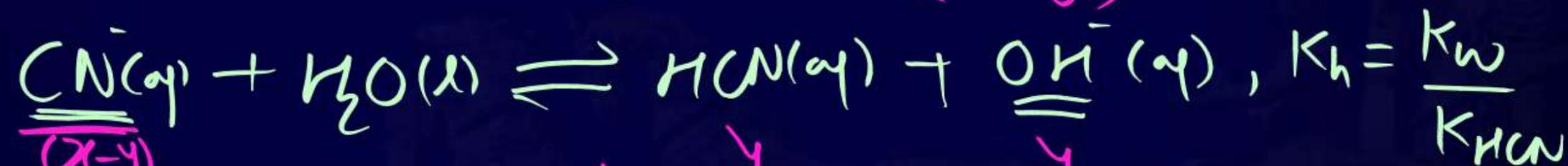
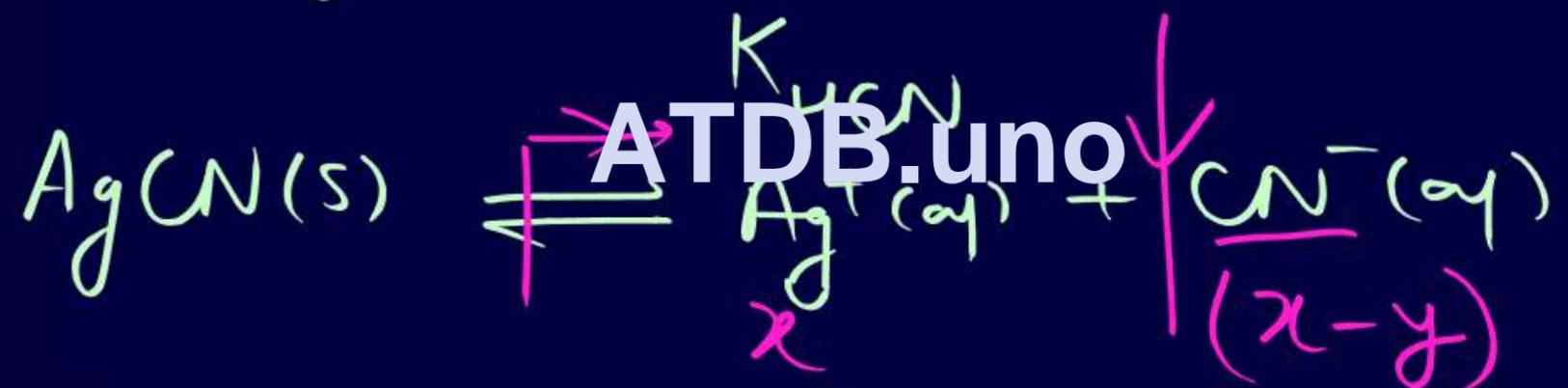
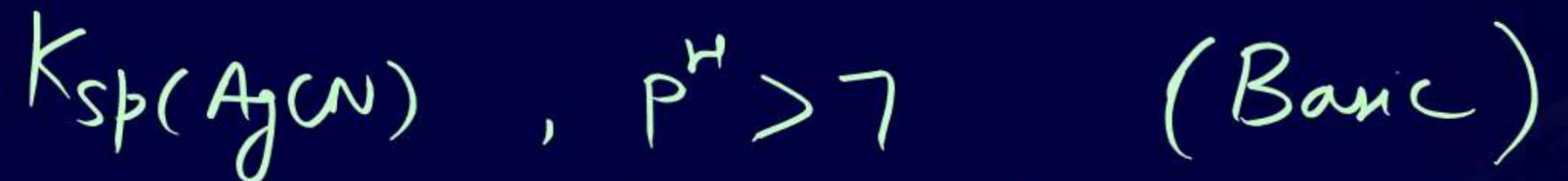
$$\text{Eqn (2) + Eqn (3), we get } K_{sp}(\text{AgCl}) + K_{sp}(\text{AgBr}) = (x' + y')^2 \quad \text{--- (4)}$$

$$x' + y' = \sqrt{K_{sp}(\text{AgCl}) + K_{sp}(\text{AgBr})}$$

# Calculation of solubility of SSS in presence of hydrolysis: GURUKUL

eg Solubility of AgCN at 25°C

Given:



∴ Solubility of AgCN ↑

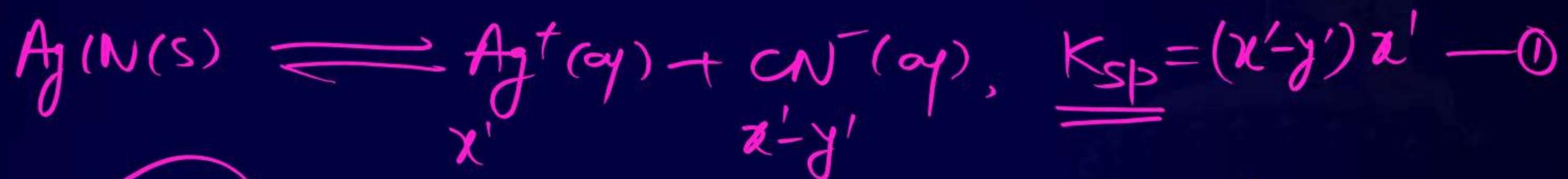


Let solubility of AgCN in basic medium  
 $= x$

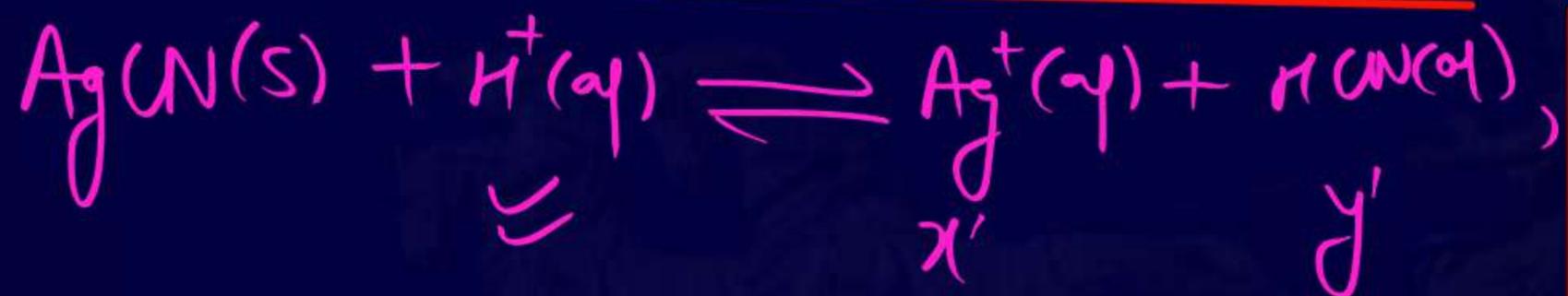
$$K_{sp}(\text{AgCN}) = x(x-y) \quad \text{--- ①}$$

$$\frac{K_w}{K_{\text{HCN}}} = \frac{[\text{HCN}][\text{OH}^-]}{[\text{CN}^-]} = \frac{y \cdot x \cdot y}{x-y} \quad \text{--- ②}$$

# In presence of Acidic medium  
 $p^H < 7$



$$\therefore \frac{1}{K_{HCN}} \gg K_{sp}$$



$$\frac{K_{sp}}{K_{HCN}} = \frac{[Ag^+][HCN]}{[H^+]}$$



# Solubility of SSS in Complex formation :-

e.g. Solubility of AgCl in 0.1M NH<sub>3</sub>

Given:

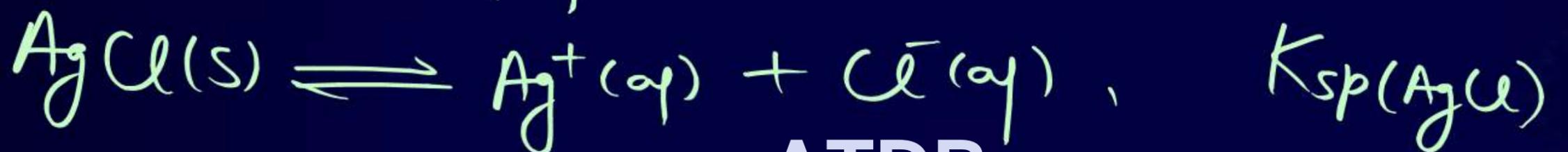
$K_f [Ag(NH_3)_2]^+$  = Complex formation Constant  
or  
Stability Constant

$K_{sp}$

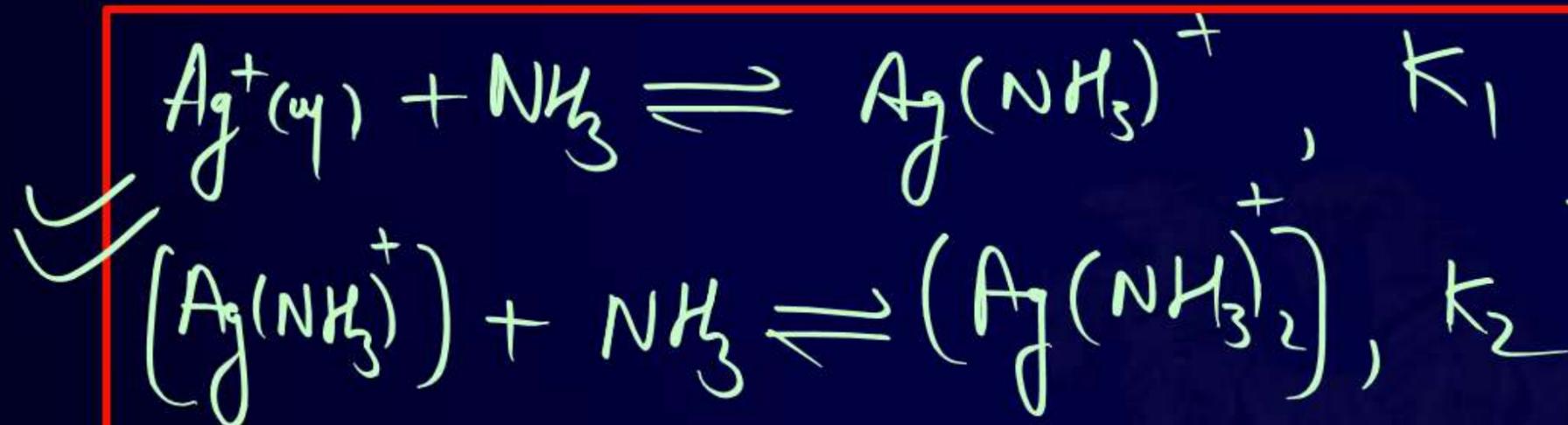


$K_{eq} \uparrow, \Delta G^\circ = -RT \ln K, \text{ Stability} \uparrow$

$\downarrow$   $K_d$  or  $K_{inst.}$  = Complex dissociation Constant or Instability Constant.

$$= \left( \frac{1}{K_f} \right)$$


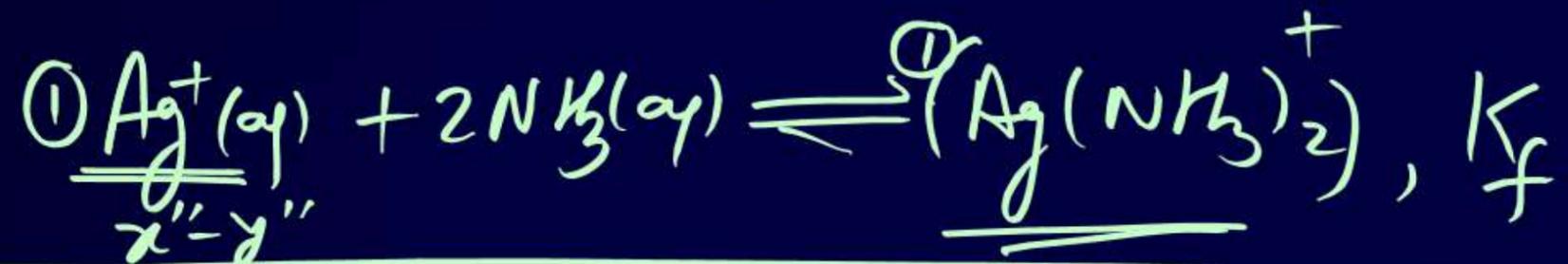
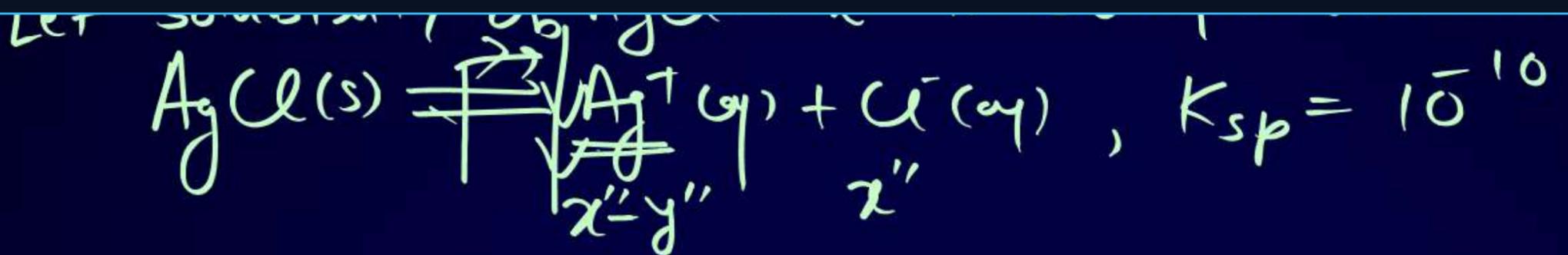
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Net reaction



$$K_f = K_1 \times K_2$$



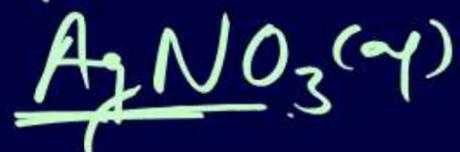
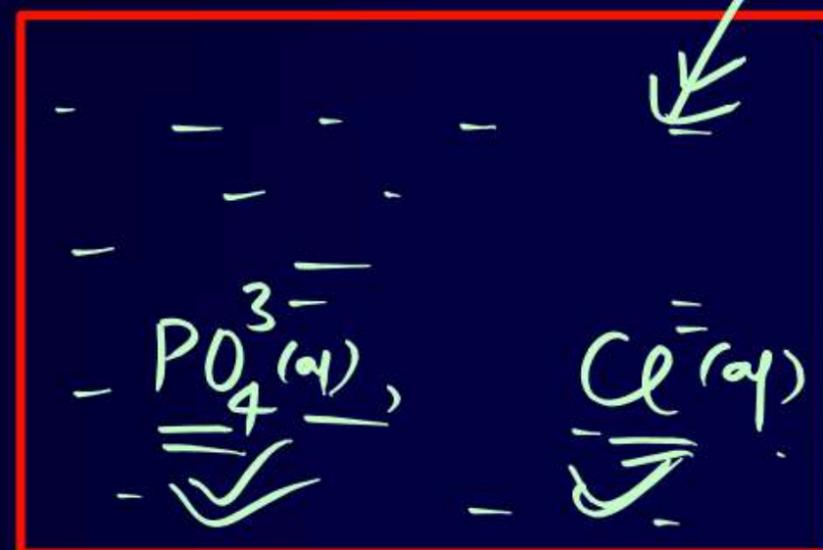
$$y'' \approx x''$$



$$K_{sp} \times K_f = \frac{[\text{Ag}(\text{NH}_3)_2^+][\text{Cl}^-]}{(\text{NH}_3)^2}; \quad [\text{Ag}(\text{NH}_3)_2^+] = (\text{Cl}^-)$$

$y'' \approx x''$

# Selective precipitation :-



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$$K_{sp}(\text{Ag}_3\text{PO}_4) = [\text{Ag}^+(\text{aq})]_{\text{req}}^3 [\text{PO}_4^{3-}(\text{aq})]$$

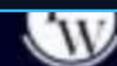
$$K_{sp}(\text{AgCl}) = [\text{Ag}^+(\text{aq})]_{\text{req}} [\text{Cl}^-(\text{aq})]$$

$$[\text{Ag}^+(\text{aq})]_{\text{req}} = \left[ \frac{K_{sp}(\text{Ag}_3\text{PO}_4)}{[\text{PO}_4^{3-}(\text{aq})]} \right]^{1/3}$$

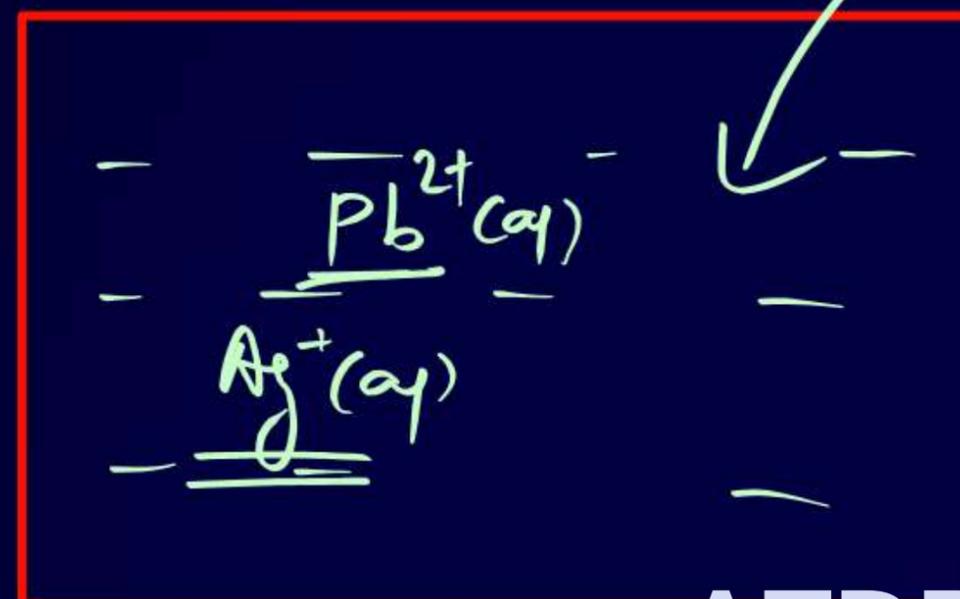
for  $\text{Ag}_3\text{PO}_4$

$$[\text{Ag}^+(\text{aq})]_{\text{req}} = \frac{K_{sp}(\text{AgCl})}{[\text{Cl}^-(\text{aq})]}$$

for  $\text{AgCl}$



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$$K_{sp}(\text{Ag}_2\text{CrO}_4) = [\text{Ag}^+]^2 [\text{CrO}_4^{2-}]$$

$$K_{sp}(\text{PbCrO}_4) = [\text{Pb}^{2+}] [\text{CrO}_4^{2-}]$$

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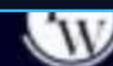
$$[\text{CrO}_4^{2-}]_{\text{req.}} = \frac{K_{sp}(\text{Ag}_2\text{CrO}_4)}{[\text{Ag}^+]^2}$$

$\text{Ag}_2\text{CrO}_4$

$$[\text{CrO}_4^{2-}]_{\text{req.}} = \frac{K_{sp}(\text{PbCrO}_4)}{[\text{Pb}^{2+}]}$$

for  $\text{PbCrO}_4$

## Question



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#Q. The values of  $K_{sp}$  for the slightly soluble salts MX and  $QX_2$  are each equal to  $4.0 \times 10^{-18}$ . Which salt is more soluble? Explain your answer fully.

HW

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Ans. (QX<sub>2</sub> is more soluble)



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## Question

#Q. The solubility of  $\text{PbSO}_4$  water is 0.038 g/L. Calculate the solubility product constant of  $\text{PbSO}_4$ .

HW

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Ans.  $1.6 \times 10^{-8}$



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## Question



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#Q. Calculate the solubility of  $\text{Mg}(\text{OH})_2$  in water.  $K_{sp} = 1.2 \times 10^{-11}$ .

HW

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Ans.  $1.4 \times 10^{-4}$



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## Question

#Q. How many mol  $\text{CuI}$  ( $K_{sp} = 5 \times 10^{-12}$ ) will dissolve in 1.0 L of 0.10 M  $\text{NaI}$  solution?

HW

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Ans.  $[\text{Cu}^+] = 5 \times 10^{-11} \text{ M}$



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## Question



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#Q. A solution of saturated  $\text{CaF}_2$  is found to contain  $4.1 \times 10^{-4}$  M fluoride ion.  
Calculate the  $K_{sp}$  of  $\text{CaF}_2$ . Neglect hydrolysis.

HW

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Ans.  $3.4 \times 10^{-11}$



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## Question



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#Q. The solubility of  $ML_2$  (formula weight, 60 g/mol) in water is  $2.4 \times 10^{-5}$  g/100 mL solution. Calculate the solubility product constant for  $ML_2$ .

HW

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Ans.  $2.6 \times 10^{-16}$



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## Question

#Q. What is the solubility (in mol/L) of  $\text{Fe}(\text{OH})_3$  in a solution of  $\text{pH} = 8.0$ ? [ $K_{\text{sp}}$  for  $\text{Fe}(\text{OH})_3 = 1.0 \times 10^{-36}$ ]

HW

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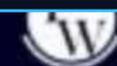


Ans.  $1.0 \times 10^{-18} \text{ M}$



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## Question



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#Q. Calculate the solubility of  $A_2X_3$  in pure water, assuming that neither kind of ion reacts with water. For  $A_2X_3$ ,  $[K_{sp} = 1.1 \times 10^{-23}]$

HW

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Ans.  $1.0 \times 10^{-5}$  mol/lit



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## Question



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#Q. Determine the solubility of AgCl in 0.1 M BaCl<sub>2</sub>. [ $K_{sp}$  for AgCl =  $1 \times 10^{-10}$ ]

HW

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Ans.  $5 \times 10^{-10} \text{ M}$



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## Question



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#Q. A solution has a  $\text{Mg}^{2+}$  concentration of  $0.0010 \text{ mol/L}$ . Will  $\text{Mg}(\text{OH})_2$  precipitate if the  $\text{OH}^-$  concentration of the solution is  $[\text{K}_{\text{sp}} = 1.2 \times 10^{-11}]$

- (a)  $10^{-5} \text{ mol/L}$
- (b)  $10^{-3} \text{ mol/L} ?$

HW

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Ans. (a) no precipitation will occur,  
(b) a precipitate will form

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## Question

#Q. Calculate solubility of  $\text{PbI}_2$  ( $K_{\text{sp}} = 1.4 \times 10^{-8}$ ) in water at  $25^\circ$ , which is 90% dissociated.

MW

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Ans.  $1.6 \times 10^{-3}$



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## Question



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#Q. Calculate solubility of AgCN ( $K_{sp} = 4 \times 10^{-16}$ ) in a buffer solution of PH = 3.

$$[K_{a(\text{HCN})} = 4 \times 10^{-10}]$$

HW

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Ans.  $3.16 \times 10^{-5}$



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## Question

#Q. Calculate the Simultaneous solubility of AgSCN and AgBr.  $K_{sp}$  (AgSCN) =  $1.1 \times 10^{-12}$ ,  $K_{sp}$ (AgBr) =  $5 \times 10^{-13}$

HW

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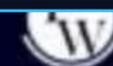
Ans.  $4 \times 10^{-7}$  mol/L AgBr,  $9 \times 10^{-7}$  mol/L AgSCN



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## Question



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#Q. Calculate  $F^-$  in a solution saturated with respect of both  $MgF_2$  and  $SrF_2$

$$K_{sp}(MgF_2) = 9.5 \times 10^{-9}, K_{sp}(SrF_2) = 4 \times 10^{-9}$$

HW

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Ans.  $[F^-] = 3 \times 10^{-3}M$



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## Question



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#Q. Assuming no change in volume, calculate the minimum mass of NaCl necessary to dissolve 0.010 mol AgCl in 100 L solution.

$$[K_f(\text{AgCl}_2^-) = 3 \times 10^5, K_{sp}(\text{AgCl}) = 1 \times 10^{-10}]$$

HW

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Ans. 19.5 kg



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## Question

#Q. A recent investigation of the complexation of  $\text{SCN}^-$  with  $\text{Fe}^{3+}$  represented by constant  $K_1$ ,  $K_2$  and  $K_3$  as 130, 16 and 1.0 respectively. What is the overall formation constant of  $\text{Fe}(\text{SCN})_3$  from its component ions, and what is the dissociation constant of  $\text{Fe}(\text{SCN})_3$  into its simplest ions on the basis of these data?

HW

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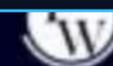
Ans.  $K_d = 1/K_f = 4.8 \times 10^{-4}$



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## Question



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#Q. How much AgBr could dissolve in 1.0 L of 0.40 M NH<sub>3</sub>? Assume that Ag(NH<sub>3</sub>)<sub>2</sub><sup>+</sup> is the only complex formed.

$$[K_f(\text{Ag}(\text{NH}_3)_2^+) = 1 \times 10^8 ; K_{sp}(\text{AgBr}) = 5 \times 10^{-13}]$$

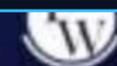
HW

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Ans.  $2.8 \times 10^{-3} \text{ M}$



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# Thank You ATDB.uno

