

(s) (3s)  

$$K_{sp} = s \times (3s)^{3}$$
  
 $= 27s^{4}$   
 $2.7 \times 10^{-31} = 27s^{4}$   
 $\therefore$   $s = \sqrt[4]{\frac{K_{sp}}{27}} = \sqrt[4]{\frac{2.7 \times 10^{-31}}{27}}$   
 $= \sqrt[4]{10^{-32}}$   
 $= 10^{-8} \text{ mol/L}$   
(d)  
 $N_{2}O_{4} \neq 2NO_{2}$   
 $\frac{1}{1-\alpha} \qquad 0$   
 $X_{p} = \frac{(n_{NO_{2}})^{2}}{n_{N_{2}O_{4}}} \times \left[\frac{P}{2\pi}\right]^{1}$   
For 33% dissociation:  $K_{p} = \frac{(2 \times 0.33)^{2}}{0.67} \times \left[\frac{P}{1.33}\right]$   
For 40% dissociation:  $K_{p} = \frac{(2 \times 0.40)^{2}}{0.60} \times \left[\frac{P}{1.40}\right]$   
 $\therefore \frac{P_{1}}{P_{2}} = 1.56 \approx 1.60 = \frac{8}{5}$   
(a)  
 $A + 2B \approx 2C$   
 $2 \quad 3 \quad 2 \quad \text{initial moles}$   
 $(2 - 0.5)(3 - 0.5)(2 \times 0.5)$  at equilibrium  
Molar concentration of  $A = \frac{1.5}{2}$   
Molar concentration of  $B = \frac{2.5}{2}$   
Molar concentration of  $C = \frac{1}{2}$   
 $K = \frac{[C]^{2}}{[A][B]^{2}} = \frac{1 \times 1 \times 2 \times 2 \times 2 \times 2 \times 2}{1.5 \times 2.5 \times 2.5} = \frac{2}{1.5 \times 2.5 \times 2.5} = 0.21$   
(d)  
In  $\frac{N}{1000}$  KOH solution,  $[OH^{-}] = 10^{-3}$  M  
 $pOH = -\log [OH^{-}] = -\log [10^{-3}]$   
 $= +3\log 10 = 3$   
 $pH + pOH = 14$   
 $pH = 14 - pOH$ 

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$$= 14 - 3$$
  
= 11

## 12 **(a)**

**(b)** 

Tears are alkaline in nature.

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$$MX_{2} = M^{2+} + 2X^{-}$$

$$s \qquad 2s$$

$$K_{sp} = (2s)^{2}(s) = 4s^{3}$$

$$\Rightarrow s = \sqrt[3]{\frac{K_{sp}}{4}} = \sqrt[3]{\frac{4 \times 10^{-12}}{4}} = 1.0 \times 10^{-4} M$$
(d)
Let colubility of PeCO = mod L<sup>-1</sup>

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Let solubility of BaSO<sub>4</sub> = mol L<sup>-1</sup>  
BaSO<sub>4</sub>
$$\Rightarrow$$
Ba<sup>2+</sup> +SO<sub>4</sub><sup>2-</sup>  
 $x$   $x$   $x$   
 $\therefore$  Ions at equilibrium  
 $K_{sp} = [Ba^{2+}][SO_4^{2-}]$   
 $= x \times x$   
 $x^2$   
Given,  $K_{sp} = 1.5 \times 10^{-9}$   
 $(1.5 \times 10^{-9}) = x^2$   
or  $\sqrt{1.5 \times 10^{-9}} = x$   
 $x = 3.9 \times 10^{-5} \text{ mol/L}$ 

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(d)

Strongest Bronsted base is that which has weakest conjugate acid.

Base	Conjugate				
	acid (base				
	+ H <sup>+</sup> )				
ClO <sup>-</sup>	HClO				
$ClO_2^-$	HClO <sub>2</sub>				
$ClO_3^-$	HClO <sub>3</sub>				
$ClO_4^-$	HClO <sub>4</sub>				

• HClO is weak conjugate acid.

 $\therefore$  ClO<sup>-</sup> is strongest Bronsted base.

## 17 **(b)**

At equilibrium  $Q = K_c$  (or  $Q = K_p$ )

## 18 **(a)**

Oxalic acid =  $x \mod/L$ Oxalic acid KMnO<sub>4</sub>

$$M_1 V_1 = M_2 V_2$$
  
40 mL × x = 16 mL × 0.05  
$$x = \frac{16 \times 0.05}{40} = \frac{1}{50}$$
$$x = \frac{1}{50} M$$

Now, convert molarity into normality  $N \times eq.wt = M \times mol.wt.of$  oxalic acid

$$N \times 45 = \frac{1}{50} \times 90$$
$$N = \frac{1}{25}$$

This normality represents the hydrogen ion concentration.

So, 
$$[H^+] = \frac{1}{25}$$
  
 $pH = \log \frac{1}{[H^+]}$   
 $= \log 25 = 1.3$ 

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(c)

Simple cations such as Ag<sup>+</sup>, Cu<sup>2+</sup>,Fe<sup>3+</sup> etc. can accept pairs of electrons and hence are Lewis acids.

20 (d)

Aspirin is a weak acid. Due to common ion effect, it is unionised in acid medium but completely ionised in alkaline medium

ANSWER-KEY											
Q.	1	2	3	4	5	6	7	8	9	10	
<b>A.</b>	В	D	Α	В	С	Α	В	Α	D	Α	
<b>Q</b> .	11	12	13	14	15	16	17	18	19	20	
A.	D	Α	D	В	D	D	B	Α	С	D	