

1. Assign oxidation number to the underlined elements in each of the following species:

(a)  $NaH_2PO_4$ 

(b)  $NaHSO_4$ 

(c)  $H_4 \underline{P}_2 O_7$ 

(d)  $K_2 \underline{Mn} O_4$ 

(e) Ca<u>O</u>2

(f) Na<u>B</u>H<sub>4</sub>

(g)  $H_2 S_2 O_7$ 

(h)	$KAl(\underline{S}O_4)_2.12H_2O_4$
An	swer:

(a)  $NaH_2\underline{P}O_4$ 

Let x be the oxidation no. of P.

Oxidation no. of Na = +1

Oxidation no. of H = +1

Oxidation no. of O = -2

 $\operatorname{Na}^{+1}_{2}\operatorname{P}^{+1}_{0}\operatorname{P}^{x}_{4}$ 

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Then,

1(+1) + 2(+1) + 1(x) + 4(-2) = 01 + 2 + x -8 = 0 x = +5 Therefore, oxidation no. of P is +5.

(b)  $NaH\underline{S}O_4$ 

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Let x be the oxidation no. of S.

Oxidation no. of Na = +1

Oxidation no. of H = +1

Oxidation no. of O = -2

Then,

1(+1) + 1(+1) + 1(x) + 4(-2) = 0

1 + 1 + x -8 = 0

х = +б

Therefore, oxidation no. of S is +6.

(c)  $H_4 \underline{P}_2 O_7$ 

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 $H_{4}^{+1} P_{2}^{x} O_{7}^{-2}$ 

Let x be the oxidation no. of P.

Oxidation no. of H = +1

Oxidation no. of O = -2

Then,

4(+1) + 2(x) + 7(-2) = 0



4 + 2x - 14 = 0

2x = +10

x = +5

Therefore, oxidation no. of P is +5.

(d)  $K_2 \underline{Mn} O_4$ 

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 $K_{2}^{+1} MnO_{4}^{-2}$ 

Let x be the oxidation no. of Mn.

Oxidation no. of K = +1

Oxidation no. of O = -2

Then,

2(+1) + x + 4(-2) = 0

2 + x - 8 = 0

х = +6

Therefore, oxidation no. of Mn is +6.

(e) Ca<u>O</u><sub>2</sub>

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# $CaO_2^*$

Let x be the oxidation no. of O.

Oxidation no. of Ca = +2

Then,

(+2) + 2(x) = 0

2 + 2x = 0

2x = -2

x = -1

Therefore, oxidation no. of O is -1.



(f)  $Na\underline{B}H_4$ 

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# <sup>+1</sup> X -1 Na BH<sub>4</sub>

Let x be the oxidation no. of B.

Oxidation no. of Na = +1

Oxidation no. of H = -1

Then,

1(+1) + 1(x) + 4(-1) = 0

1 + x -4 = 0

x = +3

Therefore, oxidation no. of B is +3.

(g)  $H_2 S_2 O_7$ 

 $\overset{\textbf{+1}}{\textbf{H}_2}\overset{\textbf{x}}{\textbf{S}_2}\overset{\textbf{-2}}{\textbf{O}_7}$ 

Let x be the oxidation no. of S.

Oxidation no. of H = +1

Oxidation no. of O = -2

Then,

2(+1) + 2(x) + 7(-2) = 0

2 + 2x - 14 = 0

2x = +12

x = +6

Therefore, oxidation no. of S is +6.

(h)  $KAl(\underline{S}O_4)_2.12H_2O$ 





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+1 3+ x 2-+1 -2 K AI (SO<sub>4</sub>)<sub>2</sub> .12H<sub>2</sub>O

Let x be the oxidation no. of S.

Oxidation no. of K = +1

Oxidation no. of AI = +3

Oxidation no. of O= -2

Oxidation no. of H = +1

Then,

1(+1) + 1(+3) + 2(x) + 8(-2) + 24(+1) + 12(-2) = 0

1 + 3 + 2x - 16 + 24 - 24 = 0

2x = +12

x = +6

Therefore, oxidation no. of S is +6.

OR

Ignore the water molecules because it is neutral. Then, the summation of the oxidation no. of all atoms of water molecules can be taken as 0. Hence, ignore the water molecule.

2x = 12

х = +б

Therefore, oxidation no. of S is +6.

2.What are the oxidation number of the underlined elements in each of the following and how do you rationalise your results?

(a) K<u>I</u>3

(b)  $H_2 \underline{S}_4 O_6$ 

(c)  $\underline{Fe}_{3}O_{4}$ 



(d)  $\underline{C}H_3\underline{C}H_2OH$ 

# (e) <u>C</u>H<sub>3</sub>COOH

Answer:

(a) K<u>I</u>3

Let x be the oxidation no. of I.

Oxidation no. of K = +1

Then,

1(+1) + 3(x) = 0

1 + 3x = 0

 $x = -\frac{1}{3}$ 

Oxidation no. cannot be fractional. Hence, consider the structure of  $\,KI_3$  .

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In  $KI_3\,$  molecule, an iodine atom forms coordinate covalent bond with an iodine molecule.

+1 K⁺[I-I←I]

Therefore, in  $KI_3$  molecule, the oxidation no. of I atoms forming the molecule  $I_2$  is 0, while the oxidation no.

of I atom which is forming coordinate bond is -1.

(b)  $H_2 \underline{S}_4 O_6$ 

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Let x be the oxidation no. of S.

Oxidation no. of H = +1 Oxidation no. of O = -2



Then,

$$2(+1) + 4(x) + 6(-2) = 0$$
$$2 + 4x - 12 = 0$$

$$x = +2\frac{1}{2}$$

Oxidation no. cannot be fractional. Therefore, S would be present with different oxidation state in molecule.

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The oxidation no. of two out of the four S atoms is +5 while that of other two atoms is 0.

# (c) $\underline{Fe}_{3}O_{4}$

Let x be the oxidation no. of Fe.

Oxidation no. of O = -2

Then,

3(x) + 4(-2) = 0

3x -8 = 0

$$x = \frac{8}{3}$$

Oxidation no. cannot be fractional.

One of the three atoms of Fe has oxidation no. +2 and other two atoms of Fe has oxidation no. +3.

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$$\dot{Fe}_{2}^{2}O, \dot{Fe}_{2}O_{3}$$





(d)  $\underline{C}H_3\underline{C}H_2OH$ 

 $\dot{\mathbf{C}}_{2}\dot{\mathbf{H}}_{6}\dot{\mathbf{O}}$ 

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Let x be the oxidation no. of C.

Oxidation no. of O= -2

Oxidation no. of H = +1

Then,

2(x) + 4(+1) + 1(-2) = 0

2x + 4 - 2 = 0

x = -2

Therefore, oxidation no. of C is -2.

(e)  $\underline{C}H_3\underline{C}OOH$ 

# $\overset{\textbf{x}}{\textbf{C}}_{2}\overset{\textbf{+1}}{\textbf{H}}_{4}\overset{\textbf{-2}}{\textbf{O}}_{2}$

Let x be the oxidation no. of C. Oxidation no. of O = -2Oxidation no. of H = +1Then,

2(x) + 4(+1) + 2(-2) = 0

2x + 4 - 4 = 0

x = 0

Therefore, average oxidation no. of C is 0. Both the carbon atoms are present in different environments so they

cannot have same oxidation no. Therefore, carbon has oxidation no. of +2 and \_2 in  $CH_3COOH$  .



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3. Justify that the following reactions are redox reactions: (a)  $CuO_{(s)} + H_{2(g)} \rightarrow Cu_{(s)} + H_2O_{(g)}$ 

(b)  $Fe_2O_{3\,(s)} + 3 CO_{(g)} \rightarrow 2 Fe_{(s)} + 3 CO_{2\,(g)}$ 

(c)  $4 BCl_{3(g)} + 3 LiAlH_{4(s)} \rightarrow 2 B_2H_{6(g)} + 3 LiCl_{(s)} + 3 AlCl_{3(s)}$ 

(d)  $2 K_{(s)} + F_{2 (g)} \rightarrow 2 K + F_{(s)}$ 

(e)  $4 \ NH_{3 \ (g)} + 5 \ O_{2 \ (g)} \rightarrow 4 \ NO_{(g)} + 6 \ H_2O_{(g)}$ Answer:

(a) 
$$CuO_{(s)} + H_{2(g)} \rightarrow Cu_{(s)} + H_2O_{(g)}$$

Oxidation no. of Cu and O in CuO is +2 and -2 respectively.

Oxidation no. of  $H_2$  is 0.

Oxidation no. of Cu is 0.

Oxidation no. of H and O in  $H_2O$  is +1 and -2 respectively.

The oxidation no. of Cu decreased from +2 in CuO to 0 in Cu. That is CuO is reduced to Cu.

The oxidation no. of H increased from 0 to +1 in  $H_2$  . That is  $H_2$  is oxidized to  $H_2O$  .



Therefore, the reaction is redox reaction.

(b)  $Fe_2O_{3\ (s)}$  + 3  $CO_{(g)}$  ightarrow 2  $Fe_{(s)}$  + 3  $CO_{2\ (g)}$ 

In the above reaction,

Oxidation no. of Fe and O in  $Fe_2O_3$  is +3 and -2 respectively.

Oxidation no. of C and O in CO is +2 and -2 respectively. Oxidation no. of Fe is 0.

Oxidation no. of C and O in  $CO_2$  is +4 and -2 respectively.

The oxidation no. of Fe decreased from +3 in  $Fe_2O_3$  to 0 in Fe. That is  $Fe_2O_3$  is reduced to Fe.

The oxidation no. of C increased from 0 to +2 in CO to +4 in  $CO_2$  . That is CO is oxidized to  $CO_2$  .

Therefore, the reaction is redox reaction.

(c)  $4 BCl_{3(g)} + 3 LiAlH_{4(s)} \rightarrow 2 B_2H_{6(g)} + 3 LiCl_{(s)} + 3 AlCl_{3(s)}$ 

the above reaction,

Oxidation no. of B and Cl in  $BCl_3$  is +3 and -1 respectively.

Oxidation no. of Li, Al and H in  $LiAlH_4$  is +1, +3 and -1 respectively.

Oxidation no. of B and H in  $B_2H_6$  is -3 and +1 respectively.

Oxidation no. of Li and Cl in LiCl is +1 and -1 respectively.

Oxidation no. of Al and Cl in  $AlCl_3$  is +3 and -1 respectively.



The oxidation no. of B decreased from +3 in  $BCl_3$  to -3 in  $B_2H_6$ . That is  $BCl_3$  is reduced to  $B_2H_6$ .

The oxidation no. of H increased from -1 in  $LiAlH_4$  to +1 in  $B_2H_6$  . That is  $LiAlH_4$  is oxidized to  $B_2H_6$ 

Therefore, the reaction is redox reaction.

(d) 
$$2 K_{(s)} + F_{2(g)} \rightarrow 2 K + F_{(s)}$$

In the above reaction,

Oxidation no. of K is 0.

Oxidation no. of F is 0.

Oxidation no. of K and F in KF is +1 and -1 respectively.

The oxidation no. of K increased from 0 in K to +1 in KF. That is K is oxidized to KF.

The oxidation no. of F decreased from 0 in  $\,F_2\,$  to -1 in KF. That is  $\,F_2\,$  is reduced to KF.

Therefore, the reaction is a redox reaction.

(e) 
$$4 \ NH_{3 \ (g)} + 5 \ O_{2 \ (g)} \rightarrow 4 \ NO_{(g)} + 6 \ H_2O_{(g)}$$

In the above reaction,

Oxidation no. of N and H in  $NH_3$  is -3 and +1 respectively.

Oxidation no. of  $O_2$  is 0.

Oxidation no. of N and O in NO is +2 and -2 respectively.

Oxidation no. of H and O in  $H_2O$  is +1 and -2 respectively.

The oxidation no. of N increased from -3 in  $\,NH_3$  to +2 in NO.

The oxidation no. of  $O_2$  decreased from 0 in  $O_2$  to -2 in NO and  $H_2O$  . That is  $O_2$  is reduced.



Therefore, the reaction is a redox reaction.

4. Fluorine reacts with ice and results in the change:

 $H_2O_{(s)} + F_{2\,(g)} \rightarrow HF_{(g)} + HOF_{(g)}$ 

Justify that this reaction is a redox reaction Answer:

 $H_2O_{(s)} + F_{2(g)} \rightarrow HF_{(g)} + HOF_{(g)}$ 

In the above reaction,

Oxidation no. of H and O in  $H_2O$  is +1 and -2 respectively.

Oxidation no. of  $F_2$  is 0.

Oxidation no. of H and F in HF is +1 and -1 respectively.

Oxidation no. of H, O and F in HOF is +1, -2 and +1 respectively.

The oxidation no. of F increased from 0 in  $\,F_2\,$  to +1 in HOF.

The oxidation no. of F decreased from 0 in  $O_2$  to -1 in HF.

Therefore, F is both reduced as well as oxidized. So, it is redox reaction.

5. Calculate the oxidation no. of sulphur, chromium and nitrogen in  $H_2SO_5$ ,  $Cr_2O_7^{2-}$  and  $NO_3^-$ . Suggest structure of these compounds. Count for the fallacy.

#### Answer:

For  $H_2SO_5$ 

Let x be the oxidation no. of S.



Oxidation no. of O= -2

Oxidation no. of H = +1

 $H^{+1} = O^{-2} = U^{-2} = O^{-1} = O$ 

Then,

2(+1) + 1(x) + 5(-2) = 0

2 + x - 10 = 0

x = +8

But the oxidation no. of S cannot be +8 as S has 6 valence electrons. Therefore, the oxidation no. of S cannot be more than +6.

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The structure of  $H_2SO_5$  is as given below:

Now,

$$2(+1) + 1(x) + 3(-2) + 2(-1) = 0$$
$$2 + x - 6 - 2 = 0$$

x = +6

Therefore, the oxidation no. of S is +6.

For  $Cr_2O_7^{2-}$ 

Let x be the oxidation no. of Cr.

Oxidation no. of O= -2

Then,

2(x) + 7(-2) = -2

2x -14 = -2

х = +б



There is no fallacy about the oxidation no. of Cr in  $\, Cr_2 O_7^{2-}$  .

The structure of  $\ Cr_2O_7^{2-}$  is as given below.

Each of the two Cr atoms has the oxidation no. of +6.



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For  $NO_3^-$ 

Let x be the oxidation no. of N. Oxidation no. of O= -2

Then,

1(x) + 3(-2) = -1

x = +5

There is no fallacy about the oxidation no. of N in  $\,NO_3^-$  .

The structure of  $\,NO_3^-\,$  is as given below.



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Nitrogen atom has the oxidation no. of +5.

6.	Write	formulas	for the	following	compounds:
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- (a) Mercury (II) chloride (b) Nickel (II) sulphate
- (c) Tin (IV) oxide (d) Thallium (l) sulphate
- (e) Iron (III) sulphate (f) Chromium (III) oxide

#### Answer:

(a) Mercury (II) chloride

# $HgCl_2$

(b) Nickel (II) sulphate

# $NiSO_4$

(c) Tin (IV) oxide

# $SnO_2$

(d) Thallium (I) sulphate

# $Tl_2SO_4$

(e) Iron (III) sulphate

# $Fe_2(SO_4)_3$

(f) Chromium (III) oxide

# $Cr_2O_3$



7. Suggest a list of the substances where carbon can exhibit oxidation states from -4 to +4 and nitrogen from -3 to +5.

#### Answer:

The compound where carbon has oxidation no. from -4 to +4 is as given below in the table:

Compounds	Oxidation no. of carbon	
$CH_2Cl_2$	0	
$HC\equiv\ CH$	-1	
$ClC\equiv\ CCl$	+1	
$CH_3Cl$	-2	
$CHCl_3$ , CO	+2	
$H_3C-\ CH_3$	-3	
$Cl_3C CCl_3$	+3	
$CH_4$	-4	
$CCl_4$ , $CO_2$	+4	



Compounds	Oxidation no. of nitrogen
$N_2$	0
$N_2H_2$	-1
$N_2O$	+1
$N_2H_4$	-2
NO	+2
$NH_3$	-3
$N_2O_3$	+3
$NO_2$	+4
$N_{2}O_{5}$	+5

# 8. While sulphur dioxide and hydrogen peroxide can act as oxidising as well as reducing agents in their reactions, ozone and nitric acid act only as oxidants. Why?

#### Answer:

In sulphur dioxide (  $SO_2$  ) the oxidation no. of S is +4 and the range of oxidation no. of sulphur is from +6 to -2.



Hence,  $SO_2$  can act as a reducing and oxidising agent.

In hydrogen peroxide (  $H_2O_2$  ) the oxidation no. of O is -1 and the range of the oxidation no. of oxygen is from 0

to -2. Oxygen can sometimes attain the oxidation no. +1 and +2.

Therefore,  $H_2O_2$  can act as a reducing and oxidising agent.

In ozone ( $O_3$ ) the oxidation no. of O is O and the range of the oxidation no. of oxygen is from 0 to -2. Hence,

the oxidation no. of oxygen only decreases in this case.

Therefore,  $O_3$  acts only as an oxidant.

In nitric acid ( $HNO_3$ ) the oxidation no. of nitrogen is +5 and the range of the oxidation no. that nitrogen can have is from +5 to -3. Hence, the oxidation no. of nitrogen can only decrease in this case.

Therefore,  $HNO_3$  acts only as an oxidant.

#### 9.Consider the reactions:

(a) 
$$6 \ CO_{2 \ (g)} + 6 \ H_2O_{(l)} \rightarrow C_6H_{12}O_{6 \ (aq)} + 6 \ O_{2 \ (g)}$$

(b)  $O_{3(g)} + H_2 O_{2(l)} \rightarrow H_2 O_{(l)} + 2 O_{2(g)}$ 

Why it is more appropriate to write these reactions as :

(a) 
$$6 CO_{2(q)} + 12 H_2O_{(l)} \rightarrow C_6H_{12}O_{6(aq)} + 6 H_2O_{(l)} + 6 O_{2(q)}$$

(b) 
$$O_{3(g)} + H_2 O_{2(l)} \rightarrow H_2 O_{(l)} + O_{2(g)} + O_{2(g)}$$

Also suggest a technique to investigate the path of the above (a) and (b) redox reactions



#### Answer:

(a)

Step 1 :

 $H_2O$  breaks to give  $H_2$  and  $O_2$ .

$$2 H_2 O_{(l)} \rightarrow 2 H_{2(g)} + O_{2(g)}$$

Step 2 :

The  $H_2$  produced in earlier step reduces  $CO_2$  , thus produce glucose and water.

 $6 \ CO_{2 \ (g)} \ + \ 12 \ H_{2 \ (g)} \ o \ C_{6} H_{12} O_{6 \ (s)} \ + \ 6 \ H_{2} O_{(l)}$ 

The net reaction is as given below:

$$[2 \ H_2 O_{(l)} \ 
ightarrow \ 2 \ H_{2 \ (g)} \ + \ O_{2 \ (g)}] imes 6$$

$$6 \ CO_{2 \ (g)} + 12 \ H_{2 \ (g)} \rightarrow C_{6}H_{12}O_{6 \ (s)} + 6 \ H_{2}O_{(l)}$$

$$6 \; CO_{2\;(g)} \; + \; \; 12 \; H_2O_{(l)} \; o \; \; C_6H_{12}O_{6\;(g)} \; + \; \; 6 \; H_2O_{(l)} \; + \; \; 6 \; O_{2\;(g)}$$

This is the suitable way to write the reaction as the reaction also produce water molecules in the photosynthesis process.

The path can be found with the help of radioactive  $\,H_2O^{18}\,$  instead of  $\,H_2O\,$  .

(b)

Step 1 :

 $O_2$  is produced from each of the reactants  $O_3$  and  $H_2O_2$ . That is the reason  $O_2$  is written two times.



 $O_3$  breaks to form  $O_2$  and 0.

Step 2 :

 $H_2O_2\,$  reacts with 0 produced in the earlier step, thus produce  $\,H_2O\,$  and  $\,O_2$  .

 $O_{3\,(g)} \rightarrow O_{2\,(g)} + O_{(g)}$ 

 $H_2O_2_{(l)} + O_{(g)} \rightarrow H_2O_{(l)} + O_2_{(g)}$ 

 $H_2 O_2 \,_{(l)} \,\, + \,\, O_3 \,_{(g)} \,\, 
ightarrow \,\, H_2 O_{(l)} \,\, + \,\, O_2 \,_{(g)} \,\, + \,\, O_2 \,_{(g)}$ 



The path can be found with the help of  $\,H_2O_2^{18}\,$  or  $\,O_3^{18}$  .

# 10. The compound $AgF_2$ is unstable compound. However, if formed, the compound acts as a very strong oxidising agent. Why?

Answer:

The oxidation no. of Ag in  $AgF_2$  is +2. But, +2 is very unstable oxidation no. of Ag. Hence, when  $AgF_2$  is

formed, silver accepts an electron and forms  $Ag^+$  . This decreases the oxidation no. of Ag from +2 to +1. +1

state is more stable. Therefore,  $AgF_2$  acts as a very strong oxidizing agent.

11. Whenever a reaction between an oxidising agent and a reducing agent is carried out, a compound of lower oxidation state is formed if the reducing agent is in excess and a compound of higher oxidation state is formed if the oxidising agent is in excess. Justify this statement giving three illustrations.

Justify the above statement with three examples.



#### Answer:

When there is a reaction between reducing agent and oxidizing agent, a compound is formed which has lower oxidation number if the reducing agent is in excess and a compound is formed which has higher oxidation number if the oxidizing agent is in excess.

(i)  $P_4$  and  $F_2$  are reducing and oxidizing agent respectively.

In an excess amount of  $P_4$  is reacted with  $F_2$  , then  $PF_3$  would be produced, where the oxidation no. of P is

+3.

 $P_{4\ (excess)}$  +  $F_2$  ightarrow  $PF_3$ 

If  $P_4$  is reacted with excess of  $F_2$ , then  $PF_5$  would be produced, where the oxidation no. of P is +5.

$$P_4 + F_2 (excess) \rightarrow PF_5$$

(ii) K and  $\,O_2\,$  acts as a reducing agent and oxidizing agent respectively.

If an excess of K reacts with  $O_2$  , it produces  $K_2O$  . Here, the oxidation number of 0 is -2.

$$4 \ K_{\ (excess)} \ + \ O_2 \ o \ 2 \ K_2 O^{-2}$$

If K reacts with an excess of  $O_2$  , it produces  $K_2O_2$  , where the oxidation number of 0 is –1.

 $2 K + O_2 (excess) \rightarrow K_2 O_2^{-1}$ 

(iii) C and  $O_2$  acts as a reducing agent and oxidizing agent respectively.



If an excess amount of C is reacted with insufficient amount of  $O_2$ , then it produces CO, where the oxidation

number of C is +2.

 $C_{(excess)} + O_2 \rightarrow CO$ 

If C is burnt in excess amount of  $O_2$ , then  $CO_2$  is produced, where the oxidation number of C is +4.

 $C + O_2 (excess) \rightarrow CO_2$ 

#### 12. How do you count for the following observations?

(a) Though alkaline potassium permanganate and acidic potassium permanganate both are used as oxidants, yet in the manufacture of benzoic acid from toluene we use alcoholic potassium permanganate as an oxidant. Why? Write a balanced redox equation for the reaction.

(b) When concentrated sulphuric acid is added to an inorganic mixture containing chloride, we get colourless pungent smelling gas HCI, but if the mixture contains bromide then we get red vapour of bromine. Why?

#### Answer:

(a) While manufacturing benzoic acid from toluene, alcoholic potassium permanganate is used as an oxidant due to the given reasons.

(i) In a neutral medium,  $OH^-$  ions are produced in the reaction. Due to that, the cost of adding an acid or a

base can be reduced.

(ii)  $KMnO_4$  and alcohol are homogeneous to each other as they are polar. Alcohol and toluene are

homogeneous to each other because both are organic compounds. Reactions can proceed at a faster rate in a

homogeneous medium compared to heterogeneous medium. Therefore, in alcohol,  $KMnO_4\,$  and toluene can

react at a faster rate.

The redox reaction is as given below:





(b) When concentrated  $H_2SO_4$  is added to an inorganic mixture containing bromide, firstly HBr is produced.

HBr, a strong reducing agent, reduces  $H_2SO_4$  to  $SO_2$  with the evolution of bromine's red vapour.

 $2 \ NaBr \ + \ 2 \ H_2SO_4 \ o \ 2 \ NaHSO_4 \ + \ 2 \ HBr$ 

 $2 HBr + H_2SO_4 \rightarrow Br_2 + SO_2 + 2 H_2O$ 

When concentrated  $H_2SO_4$  I added to an inorganic mixture containing chloride, a pungent smelling gas (HCl)

is evolved. HCl, a weak reducing agent, cannot reduce  $\,H_2SO_4\,$  to  $\,SO_2$  .

 $2 NaCl + 2 H_2SO_4 \rightarrow 2 NaHSO_4 + 2 HCl$ 

13. Identify the substance oxidised, reduced, oxidising agent and reducing agent for each of the following reactions:

(a) 
$$2 AgBr_{(s)} + C_6 H_6 O_{2 (aq)} \rightarrow 2 Ag_{(s)} + 2 HBr_{(aq)} + C_6 H_4 O_{2 (aq)}$$

(b) 
$$HCHO_{(l)} + 2 \left[ Ag(NH_3)_2 \right]^+_{(aq)} + 3 OH^-_{(aq)} \rightarrow 2 Ag_{(s)} + HCOO^-_{(aq)} +$$

$$4 NH_{3 (aq)} + 2 H_2 O_{(l)}$$

(c)  $HCHO_{(l)} + 2 Cu_{(aq)}^{2+} + 5 OH_{(aq)}^{-} \rightarrow Cu_2O_{(s)} + HCOO_{(aq)}^{-} + 3 H_2O_{(l)}$ 



(d)  $N_2 H_{4\,(l)} + 2 H_2 O_{2\,(l)} \rightarrow N_{2\,(g)} + 4 H_2 O_{(l)}$ 

(e) 
$$Pb_{(s)} + PbO_{2(s)} + 2 H_2SO_{4(aq)} \rightarrow 2 PbSO_{4(aq)} + 2 H_2O_{(l)}$$

Answer:

(a) 
$$2 \ AgBr_{(s)} + C_6 H_6 O_{2 \ (aq)} \rightarrow 2 \ Ag_{(s)} + 2 \ HBr_{(aq)} + C_6 H_4 O_{2 \ (aq)}$$

 $C_6H_6O_2 \Rightarrow$  Oxidized substance

AgBr => Reduced substance AgBr =>Oxidizing agent

 $C_6H_6O_2 \Rightarrow$  Reducing agent

(b)  $HCHO_{(l)} + 2 \left[ Ag(NH_3)_2 \right]_{(aq)}^+ + 3 OH_{(aq)}^- \rightarrow 2 Ag_{(s)} + HCOO_{(aq)}^- +$ 

 $4 NH_{3 (aq)} + 2 H_2O_{(l)}$ 

HCHO => Oxidized substance

 $[Ag(NH_3)_2]^+$  => Reduced substance

 $[Ag(NH_3)_2]^+ \Rightarrow$  Oxidizing agent

HCHO=> Reducing agent

(c) 
$$HCHO_{(l)} + 2 Cu_{(aq)}^{2+} + 5 OH_{(aq)}^{-} \rightarrow Cu_2O_{(s)} + HCOO_{(aq)}^{-} + 3 H_2O_{(l)}$$

HCHO => Oxidized substance

 $Cu^{2+} \Rightarrow$  Reduced substance

 $Cu^{2+}\,$  => Oxidizing agent



HCHO => Reducing agent

(d)  $N_2 H_{4\,(l)} + 2 H_2 O_{2\,(l)} \rightarrow N_{2\,(g)} + 4 H_2 O_{(l)}$ 

 $N_2H_4\,$  => Oxidized substance

 $H_2O_2 \Rightarrow$  Reduced substance

 $H_2O_2 \Rightarrow$  Oxidizing agent

 $N_2H_4\;$  => Reducing agent

$$(e) \ Pb_{(s)} \ + \ \ PbO_{2 \ (s)} \ + \ \ 2 \ H_2SO_{4 \ (aq)} \ \rightarrow \ \ 2 \ PbSO_{4 \ (aq)} \ + \ \ 2 \ H_2O_{(l)}$$

Pb=> Oxidized substance

 $PbO_2 \Rightarrow$  Reduced substance

 $PbO_2 \Rightarrow$  Oxidizing agent

Pb => Reducing agent

14. Consider the reactions :

 $2 \,\, S_2 O^{2-}_{3 \,\,(aq)} \,\, + \,\, I_{2 \,\,(s)} \,\, 
ightarrow \,\, S_4 O^{2-}_{6 \,\,(aq)} \,\, + \,\, 2 \,\, I^-_{(aq)}$ 

$$S_2 O^{2-}_{3\,(aq)} \ + \ 2 \ Br_{2\,(l)} \ + \ 5 \ H_2 O_{(l)} \ o \ 2 \ SO^{2-}_{4\,(aq)} \ + \ 4 \ Br^-_{(aq)} \ + \ 10 \ H^+_{(aq)}$$

Why does the same reductant, thiosulphate react differently with iodine and bromine ? Answer:



The average oxidation no. of S in  $S_2 O_3^{2-}$  is +2.

The average oxidation no. of S in  $S_4 O_6^{2-}$  is +2.5.

The oxidation no. of S in  $S_2 O_3^{2-}$  is +2.

The oxidation no. of S in  $SO_4^{2-}$  is +6.

As  $Br_2$  is a stronger oxidizing agent than  $I_2$  , it oxidizes S of  $S_2 O_3^{2-}$  to a higher oxidation no. of +6 in

 $SO_4^{2-}$  .

As  $I_2$  is a weaker oxidizing agent so it oxidizes S of  $S_2O_3^{2-}$  ion to a lower oxidation no. that is 2.5 in

 $S_4 O_6^{2-}$  ions.

Thus, thiosulphate react differently with  $I_2$  and  $Br_2$  .

15. Justify giving reactions that among halogens, fluorine is the best oxidant and among hydrohalic compounds, hydroiodic acid is the best reductant.

Answer:

 $F_2$  can oxidize  $Cl^-$  to  $Cl_2$  ,  $Br^-$  to  $Br_2$  , and  $I^-$  to  $I_2$  as:

$$F_{2 \; (aq)} \; + \; 2 \; Cl^{-}_{(s)} \; \rightarrow \; 2 \; F^{-}_{(aq)} \; + \; Cl_{2 \; (g)}$$

 $F_{2\;(aq)} \;+\;\; 2\;Br^-_{(aq)} \; o\;\; 2\;F^-_{(aq)} \;+\;\; Br_{2\;(l)}$ 



$$F_{2\;(aq)} \;+\;\; 2\; I^-_{(aq)} \; o \;\; 2\; F^-_{(aq)} \;+\;\; I_{2\;(s)}$$

But,  $Cl_2$ ,  $Br_2$ , and  $I_2$  cannot oxidize  $F^-$  to  $F_2$ . The oxidizing power of halogens increases in the order as given below:

 $I_2 < Br_2 < Cl_2 < F_2$ 

Therefore, fluorine is the best oxidant among halogens.

HI and HBr can reduce  $H_2SO_4$  to  $SO_2$  , but HCl and HF cannot. Hence, HI and HBr are

stronger reductants compared to HCl and HF.

 $2 \; HI \; + \; H_2 SO_4 \; o \; I_2 \; + \; SO_2 \; + \; 2 \; H_2 O$ 

 $2 HBr + H_2SO_4 \rightarrow Br_2 + SO_2 + 2 H_2O$ 

 $I^-$  can reduce  $Cu^{2+}$  to  $Cu^+$  , but  $Br^-$  cannot.

 $4 \,\, I^-_{(aq)} \ + \ 2 \,\, Cu^{2+}_{(aq)} \ \rightarrow \ Cu_2 I_{2 \,\,(s)} \ + \ I_{2 \,\,(aq)}$ 

Therefore, hydrochloric acid is the best reductant among hydrohalic compounds. Hence, the reducing power of hydrohalic acids increases as given below:

HF < HCl < HBr < HI

#### 16. Why does the following reaction occur ?

 $XeO_{6\;(aq)}^{4-} \ + \ 2\;F_{(aq)}^{-} \ + \ 6\;H_{(aq)}^{+} \ \rightarrow \ XeO_{3\;(g)} \ + \ F_{2\;(g)} \ + \ 3\;H_2O_{(l)}$ 



What conclusion about the compound  $Na_4 XeO_6$  ( of which  $XeO_6^{4-}$  is a part) can be drawn from the

reaction?

Answer:

$$XeO^{4-}_{6\,(aq)} \ + \ 2\ F^-_{(aq)} \ + \ 6\ H^+_{(aq)} \ o \ XeO_{3\,(g)} \ + \ F_{2\,(g)} \ + \ 3\ H_2O_{(l)}$$

The oxidation no. of Xe reduces from +8 in  $\,XeO_6^{4-}\,$  to +6 in  $\,XeO_3$  .

The oxidation no. of F increases from -1 in  $\,F^-\,$  to 0 in  $\,F_2$  .

Hence,  $XeO_6^{4-}$  is reduced on the other hand  $F^-$  is oxidized. As  $Na_2XeO_6^{4-}$  (or  $XeO_6^{4-}$  ) is a stronger

oxidizing agent compared to  $\,F_2$  , this reaction occurs.

#### 17. Consider the reactions:

(a) 
$$H_3PO_{2(aq)} + 4 AgNO_{3(aq)} + 2 H_2O_{(l)} \rightarrow H_3PO_{4(aq)} + 4 Ag_{(s)} +$$

 $4 HNO_{3 (aq)}$ 

**(b)** 
$$H_3PO_{2 (aq)} + 2 CuSO_{4 (aq)} + 2 H_2O_{(l)} \rightarrow H_3PO_{4 (aq)} + 2 Cu_{(s)} +$$

 $H_2SO_4$  (aq)

(c) 
$$C_6H_5CHO_{(l)} + 2 \left[Ag(NH_3)_2\right]^+_{(aq)} + 3 OH^-_{(aq)} \rightarrow C_6H_5COO^-_{(aq)} + 2 Ag_{(s)} +$$

 $4 NH_{3 (aq)} + 2 H_2 O_{(l)}$ 



(d)  $C_6H_5CHO_{(l)}~+~2~Cu^{2+}_{(aq)}~+~5~OH^-_{(aq)}~
ightarrow$  No change is observed

What inference do you draw about the behavior of  $Ag^+$  and  $Cu^{2+}$  from these reactions?

#### Answer:

 $Ag^+$  and  $Cu^{2+}$  behaves as oxidizing agent in reactions (i) and (ii) respectively.

In reaction (iii),  $Ag^+$  oxidizes  $C_6H_5CHO$  to  $C_6H_5COO^-$ 

In reaction (iv),  $Cu^{2+}$  cannot oxidize  $C_6H_5CHO$  .

Therefore,  $Ag^+$  is a stronger oxidizing agent compared to  $\,Cu^{2+}$  .

18. Balance the following redox reactions by ion - electron method :

- (a)  $MnO^-_{4~(aq)} + I^-_{(aq)} 
  ightarrow MnO_{2~(s)} + I_{2~(s)}$  (Basic medium)
- (b)  $MnO^-_{4\,(aq)} + SO_{2\,(g)} \rightarrow Mn^{2+}_{(aq)} + H_2SO^-_4$  (Acidic medium)
- (c)  $H_2O_{2\ (aq)} + Fe^{2+}_{(aq)} o Fe^{3+}_{(aq)} + H_2O_{(l)}$  (Acidic medium)

(d)  $Cr_2^{2-}O_{7\,(aq)} + SO_{2\,(g)} \rightarrow Cr_{(aq)}^{3+} + SO_{(aq)}^{2-}$  (Acidic medium) Answer:

(a) 
$$MnO_{4\ (aq)}^{-} + I_{(aq)}^{-} \rightarrow MnO_{2\ (s)} + I_{2\ (s)}$$

Step 1



The two half reactions are given below:

Oxidation half reaction:  $I_{(aq)} 
ightarrow I_{2\ (s)}$ 

Reduction half reaction:  $MnO_4^- 
ightarrow MnO_2$ 

Step 2

Balance I in oxidation half reaction:

 $2 \ I^-_{(aq)} \ o \ I_{2 \ (s)}$ 

Add 2  $e^-$  to the right hand side of the reaction to balance the charge:

$$2I^{-}_{(aq)} \ 
ightarrow \ I_{2 \ (s)} \ + \ 2 \ e^{-}$$

Step 3

The oxidation no. of Mn has decreased from +7 to +4 in the reduction half reaction. Therefore, 3 electrons are added to the left hand side of the reaction.

 $MnO^-_{4 \ (aq)} + 3 \ e^- \rightarrow MnO_{2 \ (aq)}$ 

Add 4  $OH^-$  ions to right hand side of the reaction to balance the charge.

$$MnO^{-}_{4\;(aq)} + 3 e^{-} \rightarrow MnO_{2\;(aq)} + 4 OH^{-}$$

Step 4

There are 6 oxygen atoms on the right hand side and 4 oxygen atoms on the left hand side. Hence, 2 water molecules are added to the left hand side.

$$MnO^{-}_{4\;(aq)} + 2 H_2O + 3 e^{-} \rightarrow MnO_{2\;(aq)} + 4OH^{-}$$

Step 5



Equal the no. of electrons on both the sides by multiplying oxidation half reaction by 3 and reduction half reaction by 2:

$$6 \,\, I^-_{(aq)} \,\, 
ightarrow \,\, 3 \,\, I_{2 \,\, (s)} \,\, + \,\, 6 \,\, e^-$$

$$2 \,\, MnO^-_{4 \,\, (aq)} \,\, + \,\, 4 \,\, H_2O \,\, + \,\, 6 \,\, e^- \,\, \rightarrow \,\, 2 \,\, MnO_{2 \,\, (s)} \,\, + \,\, 8 \,\, OH^-_{(aq)}$$

Step 6

After adding both the half reactions, we get the balanced reaction as given below:

$$6 \,\, I^-_{(aq)} \,\, + \,\, 2 \,\, Mn O^-_{4 \,\, (aq)} \,\, + \,\, 4 \,\, H_2 O_{(l)} \,\, 
ightarrow \,\, 3 \,\, I_{2 \,\, (s)} \,\, + \,\, 2 \,\, Mn O_{2 \,\, (s)} \,\, + \,\, 8 \,\, OH^-_{(aq)}$$

(b) 
$$MnO_{4\ (aq)}^{-} + SO_{2\ (g)} \rightarrow Mn_{(aq)}^{2+} + H_2SO_4^{-}$$

Step 1

Similar to (i), oxidation half reaction is:

$$SO_{2\ (g)} \ + \ 2\ H_2O_{(l)} \ o \ HSO_4^-\ (aq) \ + \ 3\ H^+_{(aq)} \ + \ 2\ e^-_{(aq)}$$

Step 2

Reduction half reaction is:

$$MnO^{-}_{4\;(aq)} + 8 H^{+}_{(aq)} + 5 e^{-} \rightarrow Mn^{2+}_{(aq)} + 4 H_2O_{(l)}$$

Step 3

Multiply the oxidation half reaction with 5 and the reduction half reaction with 2, then add them. We get the balanced reaction as given below:

Step 1

Similar to (i), oxidation half reaction is:



$$Fe^{2+}_{(aq)} \rightarrow Fe^{3+}_{(aq)} + e^{-}$$

Step 2

Reduction half reaction is:

$$H_2 O_{2 \; (aq)} \; + \; 2 \; H^+_{(aq)} \; + \; 2 \; e^- \; 
ightarrow \; 2 \; H_2 O_{(l)}$$

Step 3

Multiply the oxidation half reaction with 2 then add it to the reduction half reaction. We get the balanced reaction as given below:

$$H_2O_{2\;(aq)} \;+\;\; 2\;Fe^{2+}_{(aq)}\;2\;H^+_{(aq)}\;
ightarrow\; 2\;Fe^{3+}_{(aq)}2\;H_2O_{(l)}$$

(d) 
$$Cr_2^{2-}O_{7(aq)} + SO_{2(g)} \rightarrow Cr_{(aq)}^{3+} + SO_{(aq)}^{2-}$$

Step 1

Similar to (i), oxidation half reaction is:

$$SO_{2\;(g)} \;+\; 2\; H_2O_{(l)} \;
ightarrow \; SO_{4\;(aq)}^{2-} \;+\; 4\; H_{(aq)}^+ \;+\; 2\; e^-$$

Step 2

Reduction half reaction is:

$$Cr_2 O^{2-}_{7\,(aq)} \ + \ 14 \ H^+_{(aq)} \ + \ 6 \ e^- \ o \ 2 \ Cr^{3+}_{(aq)} \ + \ 7 \ H_2 O_{(l)}$$

Step 3

Multiply the oxidation half reaction with 2 then add it to the reduction half reaction. We get the balanced reaction as given below:

$$Cr_2^{2-}O_{7\,(aq)} + 3 \ SO_{2\,(g)} + 2 \ H^+_{(aq)} \rightarrow 2 \ Cr^{3+}_{(aq)} + 3 \ SO^{2-}_{4\,(aq)} + H_2O_{(l)}$$

19. Balance the following equations in basic medium by ion-electron method and oxidation number methods and identify the oxidising agent and the reducing agent.

(a) 
$$P_{4(s)} + OH^{-}_{(aq)} \rightarrow PH_{3(g)} + HPO^{-}_{2(aq)}$$



**(b)** 
$$N_2H_4_{(l)} + ClO_3^-_{(aq)} \rightarrow NO_{(g)} + Cl_{(g)}^-$$

(c) 
$$Cl_2O_{7(g)} + H_2O_{2(aq)} \rightarrow ClO_{2(aq)}^- + O_{2(g)} + H_{(aq)}^+$$

#### Answer:

(a) The Oxidation no. of P reduces from 0 in  $\,P_4\,$  to – 3 in  $\,PH_3\,$ 

The oxidation no. of P increases from 0 in  $P_4$  to + 2 in  $HPO_2^-$  . Therefore,  $P_4$  behaves both as areducing

agent as well asoxidizing agent in the reaction.

Ion - electron method:

- The oxidation half reaction:

$$P_{4\,(s)} \rightarrow HPO^{-}_{2\,(aq)}$$

- Balance atom P:

$$P_{4(s)} \rightarrow 4 HPO_{2(aq)}^{-}$$

- Add 8 electrons to balance oxidation no.

$$P_{4\,(s)} \rightarrow 4 \; HPO^{-}_{2\,(aq)} + 8 \; e^{-}$$

– Add  $12 \ OH^-$  to balance the charge:

$$P_{4\,(s)} \ + \ 12 \ OH^{-}_{(aq)} \ o \ 4 \ HPO^{-}_{2\,(aq)} \ + \ 8 \ e^{-}$$

– Add 4  $\,H_2O\,$  to balance H and O atoms:

$$P_{4\,(s)} + 12 OH^{-}_{(aq)} \rightarrow 4 HPO^{-}_{2\,(aq)} + 4 H_2O_{(l)} + 8 e^{-} ----(1)$$

The reduction half reaction:



$$P_{4\ (s)} \ 
ightarrow \ PH_{3\ (g)}$$

- Balance atom P:

$$P^0_{4\,(s)} \ o \ 4 \ P^{-3} H_{3\,(g)}$$

- Add 12 electrons to balance oxidation no.

$$P_{4\,(s)} + 12 e^- \rightarrow 4 P H_{3\,(g)}$$

– Add  $12 \ OH^-$  to balance the charge:

$$P_{4\,(s)} \ + \ 12 \ e^- \ o \ 4 \ PH_{3\,(g)} \ + \ 12 \ OH_{(aq)}^-$$

– Add 12  $H_2O$  to balance H and O atoms:

$$P_{4\,(s)} + 12 H_2 O_{(l)} + 12 e^- \rightarrow 4 P H_{3\,(g)} + 12 O H^-_{(aq)} ----(2)$$

- Now, multiply the equation (1) by 3 and equation (2) by 2. Then, after adding them, we get the balanced redox reaction as given below:

$$5 P_{4 (s)} + 12 H_2 O_{(l)} + 12 HO_{(aq)} \rightarrow 8 PH_{3 (g)} + 12 HPO_{2 (aq)}^-$$
  
(b)





The Oxidation no. of N increases from -2 in  $\,N_2H_4\,$  to –+2 in NO.

The oxidation no. of CI reduces from +5 in  $\,ClO_3^-\,$  to +-1 in  $\,Cl^-$  .

Therefore,  $N_2H_4$  behaves as a reducing agent while  $ClO_3^-$  behaves as an oxidizing agent in the reaction.

Ion - electron method:

- The oxidation half reaction:

$$N_2 H_{4(l)} \rightarrow NO_{(g)}$$

- Balance atom N:

 $N_2 H_{4\,(l)} ~
ightarrow ~2~NO_{(g)}$ 

- Add 8 electrons to balance oxidation no:

 $N_2 H_{4\,(l)} \rightarrow 2 \ NO_{(q)} + 8 \ e^-$ 

- Add 8 OH<sup>-</sup> to balance the charge:

 $N_2 H_{4\;(l)} \;+\;\; 8\; OH^-_{(aq)} \; o \;\; 2\; NO_{(g)} \;+\;\; 8\; e^-$ 

– Add 6  $H_2O$  to balance O atoms:

$$N_2 H_{4\,(l)} + 8 OH^-_{(aq)} \rightarrow 2 NO_{(g)} + 6 H_2 O_{(l)} + 8 e^- --- (1)$$

- The reduction half reaction:

 $ClO^-_{3\,(aq)} \rightarrow Cl^-_{(aq)}$ 



Add 6 electrons to balance oxidation no.

$$ClO^{-}_{3\,(aq)} ~+~~ 6~e^- ~\rightarrow~~ Cl^{-}_{(aq)}$$

– Add  $6 OH^-$  ions to balance the charge:

$$ClO^{-}_{3 \; (aq)} \; + \; \; 6 \; e^{-} \; 
ightarrow \; Cl^{-}_{(aq)} \; + \; \; 6 \; OH^{-}_{(aq)}$$

– Add 3  $H_2O$  to balance O atoms:

$$ClO_{3(aq)}^{-} + 3 H_2O_{(l)} + 6 e^- \rightarrow Cl_{(aq)}^{-} + 6 OH_{(aq)}^{-} --- (2)$$

Now, multiply the equation (1) by 3 and equation (2) by 4. Then, after adding them, we get the balanced redox reaction as given below:

$$3 N_2 H_{4 \ (l)} + 4 \ ClO_{3 \ (aq)}^- \rightarrow 6 \ NO_{(g)} + 4 \ Cl_{(aq)}^- + 6 \ H_2 O_{(l)}$$

Oxidation number method:

- Reduction in the oxidation no. of N = 2 × 4 = 8
- Increment in the oxidation no. of Cl = 1 × 6 = 6

Multiply  $\,N_2H_4\,$  by 3 and  $\,ClO_3^-\,$  by 4 to balance the reduction and increment of the oxidation no. :

$$3 N_2 H_{4 \ (l)} + 4 C l O_{3 \ (aq)}^- \rightarrow N O_{(g)} + C l_{(aq)}^-$$

- Balance Cl and n atoms:

$$3 N_2 H_{4 \ (l)} + 4 C l O_{3 \ (aq)}^- \rightarrow 6 N O_{(g)} + 4 C l_{(aq)}^-$$

– Add 6  $H_2O$  to balance O atoms:



$$3 N_2 H_{4 \ (l)} + 4 C l O_{3 \ (aq)}^- \rightarrow 6 N O_{(g)} + 4 C l_{(aq)}^-$$

– Add 6  $H_2O$  to balance O atoms:

$$3 N_2 H_{4(l)} + 4 C l O_{3(aq)}^- \rightarrow 6 N O_{(g)} + 4 C l_{(aq)}^- + 6 H_2 O_{(l)}$$

This is the required reaction equation.

(c)



The Oxidation no. of Cldecreases from +7 in  $\ Cl_2O_7$  to +3 in  $\ ClO_2^-$  .

The oxidation no. of Oincreases from -1 in  $\,H_2O_2\,$  to 0 in  $\,O_2$  .

Therefore,  $H_2O_2$  behaves as a reducing agent while  $Cl_2O_7$  behaves as an oxidizing agent in the reaction.

Ion - electron method:

- The oxidation half reaction:

 $H_2O_2(aq) \rightarrow O_2(g)$ 

- Add 2 electrons to balance oxidation no:

$$H_2 O_2 \,_{(aq)} \ o \ O_2 \,_{(g)} \ + \ 2 \ e^-$$



– Add  $2 \ OH^-$  to balance the charge:

$$H_2 O_{2\;(aq)} \;+\;\; 2\; OH^-_{(aq)} o \;\; O_{2\;(g)} \;+\;\; 2\; e^-$$

– Add 2  $H_2O$  to balance O atoms:

$$H_2O_2_{(aq)} + 2 OH^-_{(aq)} \rightarrow O_2_{(g)} + 2 H_2O_{(l)} + 2 e^- --- (1)$$

- The reduction half reaction:

$$Cl_2O_7(g) \rightarrow ClO_2^-(aq)$$

- Balance Cl atoms:

$$Cl_2O_7_{\ (g)} \ 
ightarrow \ 2 \ ClO_2^-_{\ (aq)}$$

- Add 8 electrons to balance oxidation no.

 $Cl_{2}O_{7\,(g)} + 8 \ e^{-} \rightarrow 2 \ ClO_{2\,(aq)}^{-}$ 

– Add  $6 OH^-$  ions to balance the charge:

 $Cl_2O_{7\,(g)} ~+~ 8~e^- ~\rightarrow~ 2~ClO^-_{2\,(aq)} ~+~ 6~OH^-_{(aq)}$ 

– Add 3  $H_2O$  to balance O atoms:

 $Cl_2O_{7\,(g)} ~+~ 3~H_2O_{(l)} ~+~ 8~e^- ~\rightarrow~ 2~ClO_{2\,(aq)}^- ~+~ 6~OH_{(aq)}^-$ 

Now, multiply the equation (1) by 4. Then, adding equation (1) and (2), we get the balanced redox reaction as given below:



$$Cl_2O_{7(g)} + 4 H_2O_{2(aq)} + 2 OH^-_{(aq)} \rightarrow 2 ClO^-_{2(aq)} + 4 O_{2(g)} + 5 H_2O_{(l)}$$

Oxidation number method:

- Reduction in the oxidation no. of  $\ Cl_2O_7$  = 4× 2 = 8
- Increment in the oxidation no. of  $H_2O_2$  = 2×1 = 2

Multiply  $H_2O_2$  by 4 and  $O_2$  by 4 to balance the reduction and increment of the oxidation no. :

$$3 N_2 H_{4(l)} + 4 C l O_{3(aq)}^- \rightarrow N O_{(g)} + C l_{(aq)}^-$$

- Balance Cl and n atoms:

 $Cl_2O_{7\,(g)} + 4 H_2O_{2\,(aq)} \rightarrow 2 ClO_{2\,(aq)}^- + 4 O_{2\,(g)}$ 

– Add 3  $H_2O$  to balance O atoms:

$$Cl_2O_{7(g)} + 4 H_2O_{2(aq)} \rightarrow 2 ClO_{2(aq)}^- + 4 O_{2(g)} + 3 H_2O_{(l)}$$

– Add  $2 \ OH^-$  and  $2 \ H_2O$  to balance H atoms:

$$Cl_2O_{7\,(g)} + 4 H_2O_{2\,(aq)} 2 OH^-_{(aq)} \rightarrow 2 ClO^-_{2\,(aq)} + 4 O_{2\,(g)} + 5 H_2O_{(l)}$$

This is the required reaction equation.

20. What sorts of informations can you draw from the following reaction ?

 $(CN)_{2\,(g)} \ + \ 2 \ OH^-_{(aq)} \ o \ CN^-_{(aq)} \ + \ CNO^-_{(aq)} \ + \ H_2O_{(l)}$ 



#### Answer:

The oxidation no. of C in  $(CN)_2$ ,  $CN^-$  and  $CNO^-$  are +3, +2 and +4 respectively.

Let the oxidation no. of C be y.

 $(CN)_2$ 

2(y - 3) = 0

Therefore, y = 3

 $CN^{-}$ 

y – 3 = -1

Therefore, y = 2

 $CNO^{-}$ 

y - 3 -2 = -1

Therefore, y = 4

The oxidation no. of C in the reaction is:

Oxidation no. of C in  $(CN)_2$  is +3

Oxidation no. of C in  $\,CN^{-}\,$  is +2

Oxidation no. of C in  $CNO^-$  is +4

We can see that the same compound is oxidized and reduced simultaneously in the reaction.

The reactions in which the same compound is oxidized and reduced is known as disproportionation reaction. Then, we can say that alkaline decomposition of cyanogens is a disproportionation reaction.

21. The  $Mn^{3+}$  ion is unstable in solution and undergoes disproportionation to give Mn<sup>2+</sup>, MnO<sub>2</sub>, and H<sup>+</sup> ion.



Write a balanced ionic equation for the reaction.

#### Answer:

The reaction is as given below:

$$Mn^{3+}_{(aq)} \rightarrow Mn^{2+}_{(aq)} + MnO_{2} \, _{(s)} + H^+_{(aq)}$$

The oxidation half reaction:

 $Mn^{3+}_{(aq)} 
ightarrow MnO_{2\ (s)}$ 

Add 1 electron to balance the oxidation no. :

$$Mn^{3+}_{(aq)} \rightarrow MnO_{2 \ (s)} + e^{-}$$

Add  $4 H^+$  ions to balance the charge:

$$Mn^{3+}_{(aq)} 
ightarrow MnO_{2~(s)} + e^- + 4 H^+_{(aq)}$$

Add 2  $\,H_2O\,$  to balance O atoms and  $\,H^+\,$  ions:

$$Mn^{3+}_{(aq)} + 2 H_2 O_{(l)} \rightarrow MnO_{2 (s)} + e^- + 4 H^+_{(aq)} ---- (1)$$

The reduction half reaction:

 $Mn^{3+}_{(aq)} 
ightarrow Mn^{2+}_{(aq)}$ 

Add 1 electron to balance the oxidation no. :

$$Mn^{3+}_{(aq)} \ + \ e^- \ o \ Mn^{2+}_{(aq)} \ ---$$
 (2)

Add equation (1) and (2) to get the balanced chemical equation:



# $2 \ Mn^{3+}_{(aq)} \ + \ 2 \ H_2O_{(l)} \ o \ MnO_{2 \ (s)} \ + \ 2 \ Mn^{2+}_{(aq)} \ + \ 4 \ H^+_{(aq)}$

22.Consider the elements:

Cs, Ne, I and F

(a) Identify the element that exhibits only negative oxidation.

(b) Identify the element that exhibits only positive oxidation.

(c) Identify the element that exhibits both negative and positive oxidation states.

(d) Identify the element that exhibits neither negative nor positive oxidation state?

#### Answer:

(a) F exhibits only negative oxidation no. That is -1.

(b) Cs exhibits only positive oxidation no. That is +1.

- (c) I exhibits both negative and positive oxidation no. That is -1, +1, +3, +5 and +7.
- (d) Ne exhibits neither negative nor positive oxidation no. That is 0.

23. Chlorine is used to purify drinking water. Excess of chlorine is harmful. The excess of chlorine is removed by treating with sulphur dioxide. Present a balanced equation for this redox change taking place in water.

#### Answer:

The redox reaction is as given below:

$$Cl_{2(s)} + SO_{2(aq)} + H_2O_{(l)} \rightarrow Cl^{-}_{(aq)} + SO^{2-}_{4(aq)}$$

The oxidation half reaction:

$$SO_{2 \ (aq)} \ 
ightarrow \ SO_{2 \ (aq)}^{2-}$$

Add 2 electrons to balance the oxidation no. :

$$SO_{2\;(aq)} \ 
ightarrow \ SO_{4\;(aq)}^{2-} \ + \ 2 \ e^{-}$$

Add  $4 H^+$  ions to balance the charge:

$$SO_{2\;(aq)} \ 
ightarrow \ SO_{4\;(aq)}^{2-} \ + \ 4 \ H^+_{(aq)} \ + \ 2 \ e^-$$



Add 2  $H_2O$  to balance O atoms and  $H^+$  ions:

$$SO_{2 \ (aq)} \ + \ 2 \ H_2O \ o \ SO_{4 \ (aq)}^{2-} \ + \ 4 \ H_{(aq)}^+ \ + \ 2 \ e^- \ --- \ (1)$$

The reduction half reaction:

 $Cl_{2(s)} \rightarrow Cl^{-}_{(aq)}$ 

Balance Cl atoms:

 $Cl_{2(s)} \rightarrow 2 Cl^{-}_{(aq)}$ 

Add 2 electrons to balance the oxidation no. :

 $Cl_{2 \ (s)} \ + \ 2 \ e^- \rightarrow \ 2 \ Cl^-_{(aq)} \ ---$  (2)

Add equation (1) and (2) to get the balanced chemical equation:

 $Cl_{2\ (s)} + SO_{2\ (aq)} + 2 H_2O_{(l)} \rightarrow 2 Cl^-_{(aq)} + SO^{2-}_{4\ (aq)} + 4 H^+_{(aq)}$ 

24. Refer to the periodic table given in your book and now answer the following questions:

(a) Select the possible non – metals that can show disproportionation reaction?

(b) Select three metals that show disproportionation reaction?

#### Answer:

One of the reacting elements always has an element that can exist in at least 3 oxidation numbers.

- (i) The non metals which can show disproportionation reactions are P, Cl and S.
- (ii) The three metals which can show disproportionation reactions are Mn, Ga and Cu.

25. In Ostwald's process for the manufacture of nitric acid, the first step involves the oxidation of ammonia gas by oxygen gas to give nitric oxide gas and steam. What is the maximum weight of nitric oxide that can be obtained starting only with 10.00 g. of ammonia and 20.00 g of oxygen?

#### Answer:

The balanced reaction is as given below:



 $4 \, NH_{3 \, (g)} \ + \ 5 \, O_{2 \, (g)} \ o \ 4 \, NO_{(g)} \ + \ 6 \, H_2O_{(g)}$ 

 $4 NH_3 = 4 \times 17 g = 68 g$ 

5 O<sub>2</sub> = 5 × 32 g = 160 g

 $4 NO = 4 \times 30 g = 120 g$ 

 $6 H_2O = 6 \times 18 g = 108 g$ 

Thus,  $NH_3$  (68 g) reacts with  $O_2$  (20 g)

Therefore, 10 g of  $\,NH_3\,$  reacts with  $\,rac{160 imes 10}{68}\,$  g = 23.53 g of  $\,O_2\,$ 

But only 20 g of  $O_2$  is available.

Hence,  $O_2$  is a limiting reagent.

Now, 160 g of  $O_2$  gives  $rac{120 imes 20}{160}$  g of N = 15 g of NO.

Therefore, max of 15 g of nitric oxide can be obtained.

26. Using the standard electrode potentials given in Table 8.1, predict if the reaction between the following is feasible:

(a) 
$$Fe^{3+}_{(aq)}$$
 and  $I^-_{(aq)}$ 

(b)  $Ag^+_{(aq)}$  and  $Cu_{(s)}$ 



(c) 
$$Fe^{3+}_{(aq)}$$
 and  $Cu_{(s)}$ 

(d)  $Ag_{(s)}$  and  $Fe^{3+}_{(aq)}$ 

(e)  $Br_{2\;(aq)}$  and  $Fe^{2+}_{(aq)}$ 

Answer:

(a)  $Fe^{3+}_{(aq)}$  and  $I^-_{(aq)}$ 

 $2 \, \, Fe^{3+}_{(aq)} \ + \ 2 \, \, I^-_{(aq)} \ \rightarrow \ 2 \, \, Fe^{2+}_{(aq)} \ + \ I_{2 \ (s)}$ 

Oxidation half reaction:  $2~I^-_{(aq)}~
ightarrow~I_{2~(s)}~+~2~e^-$  ;  $E^\circ~=~-0.54V$ 

Reduction half reaction:  $[Fe^{3+}_{(aq)} + e^- 
ightarrow Fe^{2+}_{(aq)}] imes 2;$  ;  $E^\circ = +0.77V$ 

 $2\;Fe^{3+}_{(aq)}\;+\;\;2\;I^-\;
ightarrow\;2\;Fe^{2+}_{(aq)}\;+\;\;I_{2\;(s)};;\qquad\qquad E^\circ\;=\;+0.23V$ 

 $E^\circ$  for the overall reaction is positive. Therefore, the reaction between  $Fe^{3+}_{(aq)}$  and  $I^-_{(aq)}$  is feasible.

(b)  $Ag^+_{(aq)}$  and  $Cu_{(s)}$ 

 $2 \; Ag^+_{(aq)} \; + \; Cu_{(s)} \; o \; 2 \; Ag_{(s)} \; + \; Cu^{2+}_{(aq)}$ 

Oxidation half reaction:  $Cu_{(s)} \ 
ightarrow \ Cu_{(aq)}^{2+} \ + \ 2 \ e^-$  ;  $E^\circ = -0.34 V$ 



Reduction half reaction:  $[Ag^+_{(aq)}~+~e^ightarrow~Ag_{(s)}]~ imes~2$  ;  $E^\circ~=~+0.80V$ 

 $2 \; Ag^+_{(ag)} \; + \; Cu_{(s)} \; o \; 2 \; Ag_{(s)} \; + \; Cu^{2+}$  ;  $E^\circ \; = \; +0.46 V$ 

 $E^{\circ}$  for the overall reaction is positive. Therefore, the reaction between  $Ag^+_{(aq)}$  and  $Cu_{(s)}$  is feasible.

- (c)  $Fe^{3+}_{(aq)} \ {\rm and} \ Cu_{(s)}$
- $2 \; Fe^{3+}_{(aq)} \; + \; Cu_{(s)} \; o \; 2 \; Fe^{2+}_{(s)} \; + \; Cu^{2+}_{(aq)}$

Oxidation half reaction:  $Cu_{(s)} \ o \ Cu_{(aq)}^{2+} \ + \ 2 \ e^-$  ;  $E^\circ = -0.34 V$ 

Reduction half reaction:  $[Fe^{3+}_{(aq)} + e^- 
ightarrow Fe^{2+}_{(s)}] imes 2$  ;  $E^\circ = +0.77V$ 

 $2 \; F e^{3+}_{(aq)} \; + \; C u_{(s)} \; o \; 2 \; F e^{2+}_{(s)} \; + \; C u^{2+}_{(aq)}; \qquad E^\circ \; = \; +0.43 V$ 

 $E^\circ$  for the overall reaction is positive. Therefore, the reaction between  $Fe^{3+}_{(aq)}$  and  $Cu_{(s)}$  is feasible.

(d)  $Ag_{(s)}$  and  $Fe_{(aq)}^{3+}$ 

 $Ag_{(s)} + 2 F e^{3+}_{(aq)} \rightarrow Ag^{+}_{(aq)} + F e^{2+}_{(aq)}$ 

Oxidation half reaction:  $Ag_{(s)} ~
ightarrow~ Ag^+_{(aq)} ~+~ e^-$  ;  $E^\circ = -0.80 V$ 

Reduction half reaction:  $Fe^{3+}_{(aq)}~+~e^ightarrow~Fe^{2+}_{(aq)}$  ;  $E^\circ~=~+0.77V$ 



$$Ag_{(s)} + Fe^{3+}_{(aq)} \rightarrow Ag^{+}_{(aq)} + Fe^{2+}_{(aq)}; \qquad E^{\circ} = -0.03V$$

 $E^{\circ}$  for the overall reaction is positive. Therefore, the reaction between  $Ag_{(s)}$  and  $Fe_{(aq)}^{3+}$  is feasible.

(e)  $Br_{2\;(aq)}$  and  $Fe^{2+}_{(aq)}$ 

 $Br_{2\,(s)} \ + \ 2 \ Fe^{2+}_{(aq)} \ o \ 2 \ Br^-_{(aq)} \ + \ 2 \ Fe^{3+}_{(aq)}$ 

Oxidation half reaction:  $[Fe^{2+}_{(aq)} 
ightarrow Fe^{3+}_{(aq)} + e^-] imes 2$  ;  $E^\circ = -0.77 V$ 

Reduction half reaction:  $Br_{2~(aq)}~+~2~e^ightarrow~2~Br^-_{(aq)}$  ;  $E^\circ~=~+1.09V$ 

 $Br_{2\,(s)} \ + \ 2 \ Fe^{2+}_{(aq)} \ o \ 2 \ Br^-_{(aq)} \ + \ 2 \ Fe^{3+}_{(aq)}; \hspace{1cm} E^\circ \ = \ -0.32V$ 

 $E^{\circ}$  for the overall reaction is positive. Therefore, the reaction between  $Br_{2~(aq)}$  and  $Fe^{2+}_{(aq)}$  is feasible.

27. Predict the products of electrolysis in each of the following:

(i) An aqueous solution of  $AgNO_3$  with silver electrodes

(ii) An aqueous solution AgNO<sub>3</sub> with platinum electrodes

(iii) A dilute solution of  $H_2SO_4$  with platinum electrodes

(iv) An aqueous solution of  $CuCl_2$  with platinum electrodes.



#### Answer:

(i)  $AgNO_3$  ionizes in aqueous solution to form  $Ag^+$  and  $NO_3^-$  ions.

On electrolysis, either  $Ag^+$  ion or  $H_2O$  molecule can be decreased at cathode. But the reduction potential of

 $Ag^+$  ions is higher than that of  $\,H_2O$  .

$$Ag^+_{(aq)}$$
 +  $e^ 
ightarrow$   $Ag_{(s)}$  ;  $E^\circ$  = +0.80V

 $2 \; H_2 O_{(l)} \; + \; 2 \; e^- \; 
ightarrow \; H_{2 \; (g)} \; + \; 2 \; O H^-_{(aq)}$  ;  $E^\circ \; = \; -0.83 V$ 

Therefore,  $Ag^+$  ions are decreased at cathode. Same way, Ag metal or  $H_2O$  molecules can be oxidized at

anode. But the oxidation potential of Ag is greater than that of  $H_2O$  molecules.

$$Ag_{(s)} \rightarrow Ag^+_{(aq)} + e^-$$
;  $E^\circ = -0.80V$ 

$$2 \; H_2 O_{(l)} \; 
ightarrow \; O_{2 \; (g)} \; + \; 4 \; H^+_{(aq)} \; + \; 4 \; e^-$$
 ;  $E^\circ \; = \; -1.23 V$ 

Hence, Ag metal gets oxidized at anode.

(ii) Pt cannot be oxidized very easily. Therefore, at anode, oxidation of water occurs to liberate  $\,O_2$  . At the

cathode,  $Ag^+$  ions are decreased and get deposited.

(iii)  $H_2SO_4$  ionizes in aqueous solutions to give  $H^+$  and  $SO_4^{2-}$  ions.

$$H_2 SO_4 \,_{(aq)} \ o \ 2 \, H^+_{(aq)} \ + \ SO^{2-}_{4(aq)}$$



On electrolysis, either of  $H_2O$  molecules or  $H^+$  ions can get decreased at cathode. But the decreased

potential of  $H^+$  ions is higher than that of  $H_2O$  molecules.

 $2 \; H^+_{(ag)} \; + \; 2 \; e^- \; o \; H_{2 \; (g)}$  ;  $E^\circ \; = \; 0.0 V$ 

 $2 \; H_2 O_{(aq)} \; + \; 2 \; e^- \; 
ightarrow \; H_{2 \; (g)} \; + \; 2 \; O H^-_{(aq)}$  ;  $E^\circ \; = \; -0.83 V$ 

Therefore, at cathode,  $H^+$  ions are decreased to free  $H_2$  gas.

On the other hand, at anode, either of  $H_2O$  molecules or  $SO_4^{2-}$  ions can be oxidized. But the oxidation of  $SO_4^{2-}$  involves breaking of more bonds than that of  $H_2O$  molecules. Therefore,  $SO_4^{2-}$  ions have lower oxidation potential than  $H_2O$ . Hence,  $H_2O$  is oxidized at anode to free  $O_2$  molecules.

(iv) In aqueous solutions,  $CuCl_2$  ionizes to give  $\,Cu^{2+}\,$  and  $\,Cl^-\,$  ions as:

$$CuCl_{2 \ (aq)} \ 
ightarrow \ Cu^{2+}_{(aq)} \ + \ 2 \ Cl^{-}_{(aq)}$$

 $CuCl_{2\;(aq)} \ 
ightarrow \ Cu^{2+}_{(aq)} \ + \ 2 \ Cl^{-}_{(aq)}$ 

On electrolysis, either of  $Cu^{2+}$  ions or  $H_2O$  molecules can get decreased at cathode. But the decreased potential of  $Cu^{2+}$  is more than that of  $H_2O$  molecules.

$$Cu^{2+}_{(aq)}$$
 +  $2~e^ 
ightarrow$   $Cu_{(aq)}$  ;  $E^\circ$  =  $+0.34V$ 



 $H_2O_{(l)} ~+~ 2~e^- ~
ightarrow ~H_{2~(q)} ~+~ 2~OH^-$  ;  $E^\circ ~=~ -0.83V$ 

Therefore,  $\,Cu^{2+}\,$  ions are decreased at cathode and get deposited. In the same way, at anode, either of  $\,Cl^-\,$ 

or  $H_2O$  is oxidized. The oxidation potential of  $H_2O$  is higher than that of  $Cl^-$  .

 $2 \; Cl^{-}_{(aq)} \; 
ightarrow \; Cl_{2 \; (g)} \; + \; \; 2 \; e^{-}$  ;  $E^{\circ} \; = \; +0.34 V$ 

 $2 \; H_2 O_{(l)} \; 
ightarrow \; O_{2 \; (g)} \; + \; 4 \; H^+_{(aq)} \; + \; 4 \; e^-$  ;  $E^\circ \; = \; -1.23 V$ 

But oxidation of  $H_2O$  molecules occurs at a lower electrode potential compared to that of  $Cl^-$  ions because

of over-voltage (extra voltage required to liberate gas). As a result,  $Cl^-$  ions are oxidized at the anode to

liberate  $Cl_2$  gas.

28. Arrange the given metals in the order in which they displace each other from the solution of their salts.

#### Al, Fe, Cu, Zn, Mg

Answer:

A metal with stronger reducing power displaces another metal with weaker reducing power from its solution of salt.

The order of the increasing reducing power of the given metals is as given below:

Cu < Fe < Zn < AI < Mg

Therefore, Mg can displace AI from its salt solution, but AI cannot displace Mg. Thus, the order in which the given metals displace each other from the solution of their salts is as given below: Mg >AI>Zn> Fe >Cu

29. Given the standard electrode potentials,

K<sup>+</sup> /K = −2.93V



Ag<sup>+</sup> /Ag = 0.80V

 $Hg^{2+}$  /Hg = 0.79V

 $Mg^{2+}$  /Mg = -2.37V

 $Cr^{3+}$  /Cr = -0.74V

#### Arrange these metals in their increasing order of reducing power.

#### Answer:

The reducing agent is stronger as the electrode potential decreases. Hence, the increasing order of the reducing power of the given metals is as given below:

Ag < Hg < Cr < Mg < K

30. Depict the galvanic cell in which the reaction is:

$$Zn_{(s)} \ + \ 2 \ Ag^+_{(aq)} \ o \ Zn^{2+}_{(aq)} \ + \ 2 \ Ag_{(s)}$$

#### Further show:

- (i) which of the electrode is negatively charged?
- (ii) the carriers of the current in the cell.
- (iii) individual reaction at each electrode.

#### Answer:

The galvanic cell corresponding to the given redox reaction can be shown as:

Zn| 
$$Zn^{2+}_{(aq)}$$
 ||  $Ag^+_{(aq)}$  |Ag

(i) Zn electrode is negatively charged because at this electrode, Zn oxidizes to  $\,Zn^{2+}\,$  and the leaving electrons

accumulate on this electrode.

(ii) The carriers of current are ions in the cell.



(iii) Reaction at Zn electrode is shown as:

 $Zn_{(s)} \ o \ Zn_{(aq)}^{2+} \ + \ 2 \ e^-$ 

Reaction at Ag electrode is shown as:

 $Ag^+_{(aq)}$  +  $e^-$  ightarrow  $Ag_{(s)}$ 

