

## TEXTBOOK QUESTIONS SOLVED

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

- (a) NaH<sub>2</sub><u>P</u>O<sub>4</sub>
- (b) NaH<u>S</u>O<sub>4</sub>
- (c)  $H_4\underline{P}_2O_7$
- (d)  $K_2MnO_4$

- (e) Ca<u>O</u><sub>2</sub>
- (f) NaBH<sub>4</sub>
- (g)  $H_{2}S_{2}O_{7}$
- (h)  $KAl(\underline{SO}_4)_2.12H_2O$

Answer:

(a) 
$$\cdot$$
 +1 +1 x -2  
P in Na H<sub>2</sub> P O<sub>4</sub>  
(+1) + 2(+1) + x + 4 (-2) = 0  
 $x + 3 - 8$  or  $x = +5$ 

(b) S in NaHSO₄

$$\begin{array}{l}
+1 & +1 & x - 2 \\
\text{Na H S O}_4 \\
(+1) + (+1) + x + 4 (-2) &= 0 \\
x - 6 &= 0 \\
x &= +6
\end{array}$$

- (c) P in H<sub>4</sub> P<sub>2</sub> O<sub>7</sub>  $^{+1} x \sim ^{-2}$ H<sub>4</sub> P<sub>2</sub> O<sub>7</sub>  $^{4}$  (+1) + 2x + 7 (-2) = 0  $^{2x} - 10 = 0$  $^{x} = +5$
- (d) Mn in  $K_2MnO_4$ +1 x -2  $K_2MnO_4$ 2 (+1) + x + 4 (-2) = 0

$$x - 6 = 0$$

$$x = +6 \text{ oxygen.}$$

(e) Let the oxidation number of  $CaO_2$  be x.

$$2 + 2x = 0$$
 (: oxy No. of  $a = +2$ )  
 $x = -1$ 

Thus, oxidation number of O in  $CaO_2 = -1$ .

(f) In NaBH $_4$ , H is present as hydride ion. Therefore, its oxidation number is -1. Thus,

+1 
$$x$$
 -1  
Na B H<sub>4</sub>  $\therefore$  1 (+1) +  $x$  + 4 (-1) = 0 or  $x$  = +3  
Thus, the oxidation number of B in NaBH<sub>4</sub> = +3.

(g) +1 x -2  $H_2 S_2 O_7$   $\therefore$  2 (+1) + 2 (x) + 7 (-2) = 0 or x = +6 Thus, the oxidation number of S in  $H_2 S_2 O_7 = + 6$ .

(h) +1+3 x -2 +1-2 K Al (S O<sub>4</sub>)<sub>2</sub> 12 (H<sub>2</sub> O) or +1+3+2x+8 (-2) +12 (2 × 1 - 2) or x = +6 Alternatively, since H<sub>2</sub>O is a neutral molecule, therefore, sum of oxidation numbers of all the atoms in H<sub>2</sub>O may be taken as zero. As such water molecules may be ignored white computing the oxidation number of S.

$$\therefore$$
 + 1 + 3 + 2x - 16 = 0 or x = +6

Thus, the oxidation number of S in  $KAl(SO_4)_2.12H_2O = +6$ .

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

(a) 
$$KI_3$$
 (b)  $H_2S_4O_6$  (c)  $\underline{Fe}_3O_4$  (d)  $\underline{CH}_3\underline{CH}_2OH$  (e)  $\underline{CH}_3\underline{COOH}$ .  
Answer: (a) In  $KI_3$ , since the oxidation number of K is +1, therefore,

the average oxidation number of iodine = -1/3. But the oxidation number cannot be fractional. Therefore, we must consider its structure,  $K^+[I-I < -I]$ . Here, a coordinate bond is formed between  $I_2$  molecule and  $I^-$  ion. The oxidation number of two iodine atoms forming the  $I_2$  molecule is zero while that of iodine forming the coordinate bond is -1. Thus, the O.N. of three I atoms, atoms in  $KI_3$  are 0, 0 and -1 respectively.

(b) By conventional method. O.N. of S in 
$$H_2S_4O_6 = H_2 S_4 O_6^{-2}$$
  
or  $2(+1) + 4x + 6(-2) = 0$  or  $x = +2.5$  (wrong)

But it is wrong because all the four S atoms cannot be in the same oxidation state.

By chemical bonding method. The structure of H<sub>2</sub>S<sub>4</sub>O<sub>6</sub> is shown below:

$$H - O \xrightarrow{+5} S - S - S - S - OH$$

The O.N. of each of the S-atoms linked with each other in the middle is zero while that of each of the remaining two S-atoms is +5.

- (c) By conventional method. O.N. of Fe in  $\operatorname{Fe}_3 \operatorname{O}_4^2$  or 3x + 4 (-2) = 0 or x = 8/3. By stoichiometry.  $\operatorname{Fe}_3 \operatorname{O}_4 \equiv \operatorname{Fe}_3 \operatorname{O} \cdot \operatorname{Fe}_2 \operatorname{O}_3^2$ .  $\therefore$  O.N. of Fe in  $\operatorname{Fe}_3 \operatorname{O}_4$  is + 2 and + 3
- (d) By conventional method. O.N. of C in  $CH_3CH_2OH = C_2 H_6 O$ or 2x + 6 (+ 1) + 1 (- 2) = 0 or x = -2.
- (e) By conventional method.  $CH_3COOH = C_2^x + H_4 C_2^2$  or 2x + 4 4 = 0 or x = 0By chemical bonding method,  $C_2$  is attached to three H-atoms (less electronegative than carbon) and one – COOH group (more electronegative than carbon).

therefore, O.N. of  $C_2 = 3 (+1) + x + 1 (-1) = 0$  or x = -2

 $C_1$  is, however, attached to one oxygen atom by a double bond, one OH (O.N. = -1) and one CH<sub>3</sub> (O.N. = +1) group, therefore, O.N. of  $C_1$  = + 1 + x + 1 (-2) + 1 (-1) = 0 or x = +2

Question 3. Justify that the following reactions are redox reactions:

- (a)  $CuO(s) + H_2(g) \rightarrow Cu(s) + H_2O(g)$
- (b)  $Fe_2O_3(s) + 3CO(g) \rightarrow 2Fe(s) + 3CO_2(g)$
- (c)  $4BCl_3(g) + 3LiAlH_4(s) \rightarrow 2B_2H_6(g) + 3LiCl(s) + 3AlCl_3(s)$
- (d)  $2K(s) + F_2(g) \rightarrow 2K + F^-(s)$

Answer:

(a) 
$$C_{uO(s)}^{+2-2} + H_{2}(g) \longrightarrow C_{u(s)}^{0} + H_{2O(g)}^{+1-2}$$

Here, O is removed from CuO, therefore, it is reduced to Cu while O is added to  $H_2$  to form  $H_2O$ , therefore, it is oxidised. Further, O.N. of Cu decreases from + 2 in CuO to 0 in Cu but that of H increases from 0 in  $H_2$  to +1 in  $H_2O$ . Therefore, CuO is reduced to Cu but  $H_2$  is oxidised to  $H_2O$ . Thus, this is a redox reaction.

(b) 
$$\text{Fe}_2 \overset{+3}{\text{O}_3} \overset{-2}{\text{O}_3} (s) + 3 \overset{+2}{\text{CO}} (g) \longrightarrow 2 \overset{0}{\text{Fe}} (s) + 3 \overset{+4}{\text{CO}_2} (g)$$

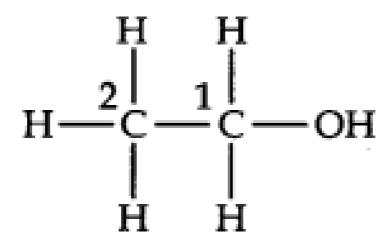
Here O.N. of Fe decreases from +3 if  $Fe_2O_3$  to 0 in Fe while that of C increases from +2 in CO to +4 in  $CO_2$ . Further, oxygen is removed from  $Fe_2O_3$  and added to CO, therefore,  $Fe_2O_3$  is reduced while CO is oxidised. Thus, this is a redox reaction.

(c) 
$$4 \stackrel{+3-1}{BCl_3}(g) + \stackrel{+1+3-1}{LiAlH_4}(s) \longrightarrow 2 \stackrel{-3}{B_2} \stackrel{+1}{H_6}(g) + 3 \stackrel{+1}{LiCl}(s) + 3 \stackrel{+3}{Al} \stackrel{-1}{Cl_3}(s)$$

Here, O.N. of B decreases from +3 in  $BrCl_3$  to -3 in  $B_2H_6$  while that of H increases from -1 in  $LiAlH_4$  to +1 in  $B_2H_6$ . Therefore,  $BCl_3$  is reduced while  $LiAlH_4$  is oxidised. Further, H is added to  $BCl_3$  but is removed from  $LiAlH_4$ , therefore,  $BCl_3$  is reduced while  $LiAlH_4$  is oxidised. Thus, it is a redox reaction.

Here, each K atom as lost one electron to form  $K^+$  while  $F_2$  has gained two electrons to form two  $F^-$  ions. Therefore, K is oxidised while  $F_2$  is reduced. Thus, it is a redox reaction.

By chemical bonding,  $C_2$  is attached to three H-atoms (less electronegative than carbon) and one  $CH_2OH$  group (more electronegative than carbon), therefore,



O.N. of  $C_2$  = 3 (+1) + x + 1 (-1) = 0 or x = -2  $G_2$  is, however, attached to one OH (O.N. = -1) and one CH<sub>3</sub> (O.N. = +1) group, therefore, O.N. of  $C_4$  = +1 + 2 (+1) + x + 1 (-1) = 0 or x = -2

Question 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: Writing the O.N. of each atom above its symbol, we have,

$$\overset{+1}{\text{H}_2}\overset{-2}{\text{O}} + \overset{0}{\text{F}_2} \longrightarrow \overset{+1}{\text{H}}\overset{-1}{\text{F}} + \overset{+1}{\text{H}}\overset{-2}{\text{O}}\overset{+1}{\text{F}}$$

Here, the O.N. of F decreases from 0 in  $F_2$  to -1 in HF and increases from 0 in  $F_2$  to +1 in HOF. Therefore,  $F_2$  is both reduced as well as oxidised. Thus, it is a redox reaction and more specifically, it is a disproportionation reaction.

Question 5. Calculate the oxidation number of sulphur, chromium and nitrogen in  $H_2SO_5$ ,  $Cr_2O_2$  and NOT. Suggest structure of these compounds. Count for the fallacy.

Answer: O.N. of S in  $H_2SO_5$ . By conventional method, the O.N. of S in  $H_2SO_5$  is 2 (+1) + x + 5 (-2) = 0 or x = +8 This is impossible because the maximum O.N. of S cannot be more than six since it has only six electrons in the valence shell. This fallacy is overcome if we calculate the O.N. of S by chemical bonding method. The structure of  $H_2SO_5$  is

$$H - O - S - O - O - H$$

$$2 \times (+1) + x + 2 (-1) + 3 \times (-2) = 0 \text{ or } x = +6$$

$$(\text{for H}) \quad (\text{for S}) \quad \text{for } (O - O) \text{ (for other O)}$$

$$atoms$$

$$Cr \text{ in } Cr_2O_7^{-2}$$

$$2x + (-2 \times 7) = -2$$

$$2x - 14 = -2$$

$$2x = -2 + 14 \qquad x = +6$$

$$x + 1 (-1) + 1 (-2) + 1 (-2) = 0 \text{ or } x + 5$$

$$(\text{for O}) \quad (\text{for = O)} \quad \text{for } \rightarrow O$$

Thus, there is no fallacy about the O.N. of N in NO<sub>3</sub><sup>-</sup>whether one calculates by conventional method or by chemical bonding method.

Question 6. Write formulas for the following compounds:

- (a) Mercury (II) chloride,
- (b) Nickel (II) sulphate,
- (c) Tin (IV) oxide,
- (d) Thallium (I) sulphate,
- (e) Iron (III) sulphate,
- (f) Chromium (III) oxide.

Answer:

- (a)  $Hg(II) Cl_2$ ,
- (b) Ni(II) SO<sub>4</sub>,
- (c)  $S_n(IV) O_2$
- (4) T (1) CO
- (d)  $T_{12}(I) SO_4$ ,
- (e)  $Fe_2(III) (SO_4)_3$ ,
- (f)  $Cr_2(III) O_3$ .

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

7 (110 17 01)			
Compound	O.N. of Carbon	Compound	O.N. of Nitrogen
CH <sub>4</sub>	-4	$NH_3$	-3
CH <sub>3</sub> CH <sub>3</sub>	, -3	$NH_2 - NH_2$	-2
CH <sub>2</sub> =CH <sub>2</sub> or CH <sub>3</sub> Cl	-2	NH=NH	-1
CH = CH	-1	N=N	0
CH <sub>2</sub> Cl <sub>2</sub> or C <sub>6</sub> H <sub>12</sub> O <sub>6</sub>	0	$N_2O$	+1
C2Cl2 or C6Cl6	+1	NO	+2
CO or CHCl <sub>3</sub>	+2	$N_2O_3$	+3
C2Cl6 or (COOH)2	+3	$N_2O_4$	+4
CO <sub>2</sub> or CCl <sub>4</sub>	+4	$N_2O_5$	+5

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

Answer:

- (i) In  $SO_2$ , O.N. of S is +4. In principle, S can have a minimum O.N. of 2 and maximum of +6. Therefore, S in  $SO_2$  can either decrease or increase its O.N. and hence can act both as an oxidising as well as a reducing agent.
- (ii) In  $H_2O_2$ , the O.N. of O is -1. In principle, O can have a minimum O.N. of -2 and maximum of zero (+1 is possible in  $O_2F_2$  and +2 in  $OF_2$ ). Therefore, O in  $H_2O_2$  can either decrease its O.N. from -1 to -2 or can increase its O.N. from -1 to zero. Therefore,  $H_2O_2$  acts both as an oxidising as well as a reducing agent.
- (iii) In  $O_3$ , the O.N. of O is zero. It can only decrease its O.N. from zero to -1 or -2, but cannot increase to +2. Therefore,  $O_3$  acts only as an oxidant.
- (iv) In  ${\rm HNO_3}$ , O.N. of N is +5 which is maximum. Therefore, it can only decrease its O.N. and hence it acts as an oxidant only.

Question 9. Consider the reactions: (a)  $6CO_2(g) + 6H_2O(I) \rightarrow C_6H_{12}O_6(s) + 6O_6(g)$  (b)  $O_3(g) + H_2O_2(l) H_2O(l) + 2O_2(g)$ 

Why it is more appropriate to write these reactions as:

(a) 
$$6CO_2(g) + 12H_2O(l) \rightarrow C_6H_{12}O_6(s) + 6H_2O(l) + 6O_2(g)$$

(b) 
$$O_3(g) + H_2O_2(I) \rightarrow H_2O(I) + O_2(g) + O_2(g)$$

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

Answer:

- (a) Therefore, it is more appropriate to write the equation for photosynthesis as (iii) because it emphasises that  $12H_2O$  are used per molecule of carbohydrate formed and  $6H_2O$  are produced during the process.
- (b) The purpose of writing  $O_2$  two times suggests that  $O_2$  is being obtained from each of the two reactants.

$$\begin{array}{ccc}
O_3(g) & \longrightarrow & O_2(g) + O(g) \\
H_2O_2 + O(g) & \longrightarrow & H_2O(l) + O_2(g) \\
\hline
O_3(g) + H_2O_2(l) & \longrightarrow & H_2O(l) + O_2(g) + O_2(g)
\end{array}$$

The path of reactions (a) and (b) can be determined by using  $\rm\,H_2O_2^{18}$  or  $\rm\,D_2O$  in reaction

(a) or by using  $H_2O_2^{18}$  or  $O_3^{18}$  in reaction (b).

Question 10. The compound  ${\rm AgF_2}$  is unstable. However, if formed, the compound acts as a very strong oxidising agent. Why? Answer:

In  ${\rm AgF_2}$  oxidation state of Ag is +2 which is very very unstable. Therefore, it quickly accepts an electron to form the more stable +1 oxidation state.

$$Ag^{2+} + e^{-} \rightarrow Ag^{+}$$

Therefore, AgF<sub>2</sub>, if formed, will act as a strong oxidising agent.

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