

## Speed Test-36

1. (a)  $[\text{Fe}(\text{H}_2\text{O})_6]\text{NO}]\text{SO}_4$   
Let O.N. of Fe be  $x$  then,  
 $1 \times (x) + 5 \times (0) + 1 \times (+1) + 1 \times (-2) = 0 \quad \therefore x = +1$
2. (c) In  $\text{BaO}_2 + \text{H}_2\text{SO}_4 \rightarrow \text{BaSO}_4 + \text{H}_2\text{O}_2$  all atoms are present in the same O.S. in reactants and products.
3. (a)  $\text{Cr}_2\text{O}_7^{2-} + 6\text{I}^- + 14\text{H}^+ \longrightarrow 2\text{Cr}^{3+} + 3\text{I}_2 + 7\text{H}_2\text{O}$   
+ 3 oxidation state of Cr.
4. (d) O.N. of N in  $\text{NO}_2$  and  $\text{N}_2\text{O}_4$  is +4  
 $\therefore$  difference is zero.  
O.N. of P in  $\text{P}_2\text{O}_3$  and  $\text{P}_4\text{O}_{10}$  is +5  
 $\therefore$  difference is zero  
O.N. of N in  $\text{N}_2\text{O}$  is +1 and in NO is +2. The difference is 1  
O.N. of S in  $\text{SO}_2$  is +4 and in  $\text{SO}_3$  is +6. The difference is +2.
5. (d)  $\text{Xe} = 53.5\% \quad \therefore \text{F} = 46.5\%$   
Relative number of atoms Xe  
$$= \frac{53.5}{131.2} = 0.4 \text{ and } \text{F} = \frac{46.5}{19} = 2.4$$
  
Simple ratio  $\text{Xe} = 1$  and  $\text{F} = 6$   
 $\therefore$  Molecular formula is  $\text{XeF}_6$   
O.N. of Xe is +6.
6. (d) The element is Ti (At. no. 22). Electronic configuration is  $1s^2, 2s^2p^6, 3s^2p^6d^2, 4s^2$ , the energy level of 3d and 4s is very close. It can exhibit +4 oxidation state ( $\text{Ti}^{4+}$ ).
7. (d) In  $\text{Ag}_2\text{O}$ , O.N. of Ag is +1 and in Ag the O.N. is 0. There is gain of electrons, hence  $\text{H}_2\text{O}_2$  act as a reducing agent.
8. (d) In redox reaction oxidation and reduction take place simultaneously.  $\text{Cu}_2\text{S} + 2\text{FeO} \rightarrow 2\text{Cu} + 2\text{Fe} + \text{SO}_2$ .  
O.N. of Cu changes from +1 to 0 (reduction) and O.N. of S changes from -2 to +4 (oxidation).
9. (d) Order of decreasing electrode potentials of Mg, K, Ba and Ca is:  
 $\text{Mg} > \text{Ca} > \text{Ba} > \text{K}$   
It can be explained by their standard reduction potentials.  
$$E_{\text{K}^+/\text{K}}^\circ = -2.925$$
  
$$E_{\text{Ba}^{2+}/\text{Ba}}^\circ = -2.90$$
  
$$E_{\text{Ca}^{2+}/\text{Ca}}^\circ = -2.87$$
  
$$E_{\text{Mg}^{2+}/\text{Mg}}^\circ = -2.37$$
  
Highly negative value of  $E_{\text{red}}^\circ$  shows the least value of electrode potential.

10. (a) (i)  $\text{Mn}^{n+} + n\text{e}^- \rightleftharpoons \text{M}$ , for this reaction, high negative value of  $E^\circ$  indicates lower reduction potential, that means M will be a good reducing agent.

Stronger reducing agent  $\Rightarrow$  Easy to oxidise  
 $\Downarrow$   
Lower reduction potential  $\Leftarrow$  higher oxidation potential

(ii) Element	F	Cl	Br	I
Reduction potential ( $E^\circ$ volt)	+2.87	+1.36	+1.06	+0.54

As reduction potential decreases from fluorine to iodine, oxidising nature also decreases from fluorine to iodine.

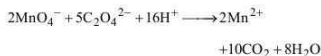
(iii) The size of halide ions increases from  $\text{F}^-$  to  $\text{I}^-$ . The bigger ion can loose electron easily. Hence the reducing nature increases from HF to HI.

11. (d) Negative  $E^\circ \Rightarrow$  Stronger reducing agent or weaker oxidising agent  
Positive  $E^\circ \Rightarrow$  Weaker reducing agent or stronger oxidising agent.
12. (a) Both Fe(II) and S(IV) in  $\text{SO}_3^{2-}$  can be oxidised to Fe(III) and  $(\text{SO}_4)^{2-}$  respectively hence  $(3/5) \times 0.5 = 0.3$  mol /litre.  
$$\left[ \text{MnO}_4^- + 5\text{e}^- + 8\text{H}^+ \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O} \right] \times \frac{3}{5}$$
  
$$\text{Fe}^{2+} \longrightarrow \text{Fe}^{3+} + \text{e}^-$$
  
$$\text{SO}_3^{2-} \longrightarrow \text{SO}_4^{2-} + 2\text{e}^-$$
  
$$\frac{3}{5}\text{MnO}_4^- + \frac{24}{5}\text{H}^+ + \text{Fe}^{2+} + \text{SO}_3^{2-} \longrightarrow \frac{3}{5}\text{Mn}^{2+} + 12/5\text{H}_2\text{O} + \text{Fe}^{3+} + \text{SO}_4^{2-}$$
13. (d) Reduction potential of Cu(II) is greater than that of Zn(II) and Al(III) thus can be easily replaced by these ions. Moreover solution of copper is blue in color.
14. (b)  $3\text{SO}_2 + \text{O}_3 \rightarrow 3\text{SO}_3$   
O.N. of S changes from +4 to +6. Two electron change  
 $\therefore$  Eq. Wt = M/2. (molecular wt. = M)
15. (a) 
$$2\text{KMnO}_4 + 3\text{H}_2\text{SO}_4 \xrightarrow{+7} \text{K}_2\text{SO}_4 + 2\text{MnSO}_4 + 3\text{H}_2\text{O} + 5\text{O}$$
  
$$2\text{FeSO}_4 + \text{H}_2\text{SO}_4 + \text{O} \xrightarrow{+2} \text{Fe}_2(\text{SO}_4)_3 + \text{H}_2\text{O}$$

O.N. of Mn changes from +7 to +2 (Reduction)

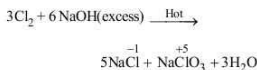
O.N. of Fe changes from +2 to +3 (Oxidation)

16. (c) On balancing the given equations, we get

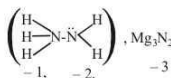


So,  $x = 2, y = 5$  &  $z = 16$

17. (b) On reaction with hot and concentrated alkali a mixture of chloride and chlorate is formed



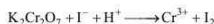
18. (a)  $[\text{Co}(\text{NH}_3)_5\text{Cl}]\text{Cl}_2$ ,  $\begin{array}{c} +1 \\ \text{H} \\ -1 \\ \text{H} \end{array} \text{N} \begin{array}{c} -1 \\ -2 \\ -1 \end{array} \text{H}$



19. (c) The redox couple with maximum reduction potential will be best oxidising agent and with minimum reduction potential will be best reducing agent.

20. (a)  $3\text{MnO}_4^{2-} + 2\text{H}_2\text{O} \rightarrow \text{MnO}_2 + 2\text{MnO}_4^- + 4\text{OH}^-$   
or  $\text{MnO}_4^{2-} + \frac{2}{3}\text{H}_2\text{O} \rightarrow \frac{1}{3}\text{MnO}_2 + \frac{2}{3}\text{MnO}_4^- + \frac{4}{3}\text{OH}^-$

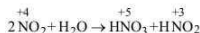
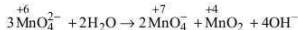
21. (b) In iodometry,  $\text{K}_2\text{Cr}_2\text{O}_7$  liberates  $\text{I}_2$  from iodides (NaI or KI) which is titrated with  $\text{Na}_2\text{S}_2\text{O}_3$  solution.



Here, one mole of  $\text{K}_2\text{Cr}_2\text{O}_7$  accepts 6 mole of electrons.

$$\therefore \text{Equivalent weight} = \frac{\text{molecular weight}}{6}$$

22. (d)  $\text{Cl}_2 + 2\text{NaOH} \rightarrow \text{NaCl} + \text{NaClO} + \text{H}_2\text{O}$



All undergo disproportionation.

23. (c)  $\begin{array}{c} -4 & +4 \\ \text{N}_2 & \text{H}_4 \end{array} \xrightarrow{\text{loss of } 10e^-} \text{N}_2^{+6}(\text{y})$

O.N. of N changes from -2 to +3

24. (b) Phosphorus, sulphur and chlorine disproportionate in the alkaline medium.

25. (c) Oxidation number of oxygen in  $\text{OF}_2 = +2$  and

$$\text{in } \text{K}_2\text{O}_2 = \frac{-1}{2}$$

26. (d) In  $\text{H}_2\text{SO}_4$ , sulphur is in highest oxidation state (+6). Hence  $\text{H}_2\text{SO}_4$  will be strongest oxidising agent.

27. (c)  $\text{K}_2\text{Cr}_2\text{O}_7 + 3\text{SO}_2 + 4\text{H}_2\text{SO}_4 \rightarrow$

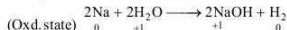


O.N. of chromium changes from +6 to +3

28. (a) Higher the value of reduction potential higher will be the oxidising power whereas the lower the value of reduction potential higher will be the reducing power.

29. (a) More the reduction potential, more will be the oxidising power.

30. (c) The violent reaction between sodium and water is an example of redox reaction:



(Oxd. state)  $\begin{array}{c} 0 \\ \text{Na} \end{array} + \begin{array}{c} 0 \\ +1 \\ \text{H}_2\text{O} \end{array} \longrightarrow \begin{array}{c} +1 \\ \text{NaOH} \end{array} + \begin{array}{c} 0 \\ \text{H}_2 \end{array}$

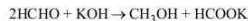
In this reaction, sodium (Na) is oxidised to NaOH while  $\text{H}_2\text{O}$  is reduced to  $\text{H}_2$ .

31. (a)  $\text{FeSO}_4$  is oxidised to  $\text{Fe}_2(\text{SO}_4)_3$ , change in O.N. of Fe is by 1. Hence equivalent weight of Mohr's salt is  $M/1 = M$ .

32. (d)  $2\text{K}_2\text{CrO}_4 + 2\text{HCl} \rightarrow \text{K}_2\text{Cr}_2\text{O}_7 + 2\text{KCl} + \text{H}_2\text{O}$   
Coefficients are 2, 2, 1, 2, 1

33. (a)

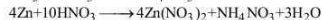
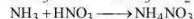
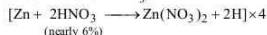
34. (c) In Cannizzaro's reaction



formaldehyde is getting reduced as well as oxidised.

35. (a)

36. (d) Zinc gives  $\text{H}_2$  gas with dil  $\text{H}_2\text{SO}_4/\text{HCl}$  but not with  $\text{HNO}_3$  because in  $\text{HNO}_3$ ,  $\text{NO}_3^-$  ion is reduced and give  $\text{NH}_4\text{NO}_3$ ,  $\text{N}_2\text{O}$ ,  $\text{NO}$  and  $\text{NO}_2$  (based upon the concentration of  $\text{HNO}_3$ )



Zn is on the top position of hydrogen in electrochemical series. So Zn displaces  $\text{H}_2$  from dilute  $\text{H}_2\text{SO}_4$  and  $\text{HCl}$  with liberation of  $\text{H}_2$ .



37. (c)

38. (a) ON of S in  $\text{S}_8 = 0$

ON of S in  $\text{S}_2\text{F}_2 = +1$

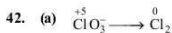
ON of S in  $\text{H}_2\text{S} = -2$

39. (a) More is  $E_{\text{RP}}^\circ$ , more is the tendency to get itself reduced or more is oxidising power.

40. (a) For statement (iii),  $\text{HgCl}_2$  is reduced to  $\text{Hg}_2\text{Cl}_2$

41. (d) More the negative reduction potential, more is the tendency to lose electron. The reducing power increases as the standard reduction potential becomes more and more negative.

Thus, Li is the strongest reducing agent as the standard reduction potential of  $\text{Li}^+/\text{Li}$  is most negative,  $-3.05\text{ V}$ .



$$x - 6 = -1 \quad x = 0$$

$$x = +5 \quad x = 0 \quad (x = \text{oxidation number})$$

$$\text{Equivalent mass} = \frac{\text{Molecular mass}}{\text{Oxidation number}} = \frac{84.45}{5} = 16.89$$

43. (b) In  $\text{SO}_3$ , the O.N. of S can increase and decrease. Hence can behave as reducing and oxidising agent. Oxidation state of S varies from  $-2$  to  $6$ .

44. (a)  $-(4/3)$  is the average oxidation state of C in  $\text{C}_3\text{H}_4$ .

45. (c) The redox reaction involve loss or gain of electron(s) i.e. change in oxidation state. Given reaction is not a redox reaction as this reaction involves no change in oxidation state of reactant or product.