## **CBSE Test Paper-04**

## **Class - 12 Chemistry (The p - Block Elements)**

- 1. Which of the following hydrides is the most acidic?
  - a. H<sub>2</sub>S
  - b. H<sub>2</sub>Se
  - c. H<sub>2</sub>Te
  - d. H<sub>2</sub>O
- 2. Which among the following forms basic oxide?
  - a. Nitrogen
  - b. Antimony
  - c. Phosphorous
  - d. Bismuth
- 3. Of the following hydrides which is the strongest reducing agent?
  - a. PH<sub>3</sub>
  - b. AsH<sub>3</sub>
  - c. BiH<sub>3</sub>
  - d. NH<sub>3</sub>
- 4. Fluorine reacts with H<sub>2</sub>S to produce
  - a. SF<sub>6</sub> and HF
  - b. SF<sub>4</sub> and HF
  - c.  $SF_6$ , S and HF
  - d.  $SF_2$  and  $HF_4$
- 5. Bleaching powder is treated with  $CO_2$ 
  - a. It absorbs the gas
  - b. CaO is formed
  - c. CaCl<sub>2</sub> is formed
  - d.  $CaCO_3$  and  $Cl_2$  is formed
- 6. Balance the following equation:  $XeF_6 + H_2O 
  ightarrow XeO_2F_2 + HF$  .

- 7. Which is the strongest oxidizing agent among  $ClO_4^-$ ,  $\ BrO_4^-$  and  $IO_4^-$ ? Given Reduction potentials  $E^\circ$  for  $ClO_4^-$ ,  $BrO_4^-$  and  $IO_4^-$  are 1.19, 1.74 and 1.65 V respectively.
- 8. Give he disproportionation reaction of  $H_3PO_3$ .
- 9. Halogens have maximum negative electron gain enthalpy in the respective periods of the periodic table. Why?
- 10. Complete the following reactions:

i. 
$$C_2H_4+O_2
ightarrow$$

ii. 
$$4Al+3O_2 
ightarrow$$

- 11. What happens when sulphur dioxide is passed through an aqueous solution of Fe (III) salt?
- 12. Draw the structure of XeF<sub>4</sub>.
- 13. Draw the structure of  $H_3PO_2$ .
- 14. Why is  $K_{a2} \le K_{a1}$  for  $H_2SO_4$  in water?
- 15. How would you account for the following:
  - i.  $NH_3$  is a stronger base than  $PH_3$
  - ii. Sulphur has a greater tendency for catenation than oxygen.
  - iii.  $\,F_2$  is a stronger oxidizing agent than  $\text{\rm Cl}_2$

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## Class - 12 Chemistry (The p - Block Elements) Solutions

1. c. H<sub>2</sub>Te

**Explanation:**  $H_2$ Te. Acidic strength increases from  $H_2$ O to  $H_2$ Te. This is because the dissociation energies decreases as bond length of M-H increases from oxygen and tellurium.

2. d. Bismuth

**Explanation:** As we move down the group in periodic table, metallic character increases so Bi is a metal thus its oxide is basic.

3. c. BiH<sub>3</sub>

**Explanation:** The reducing character of the hydrides of Group 15 elements increases from  $NH_3$  to  $BiH_3$ (Bismuthine) because the reducing character depends upon the stability of the hydride. The greater the unstability of hydride, the greater is its reducing character. Since the  $BiH_3$  is least stable (because the size of a central atom is greatest & therefore its tendency to form stable covalent bond with small hydrogen atom decreases, as a result, the bond strength decreases) in this series,  $BiH_3$  is a strongest reducing agent.

4. a.  $SF_6$  and HF

Explanation:  $4F_2 + \ H_2S 
ightarrow SF_6 + \ 2HF$ 

5. d.  $CaCO_3$  and  $Cl_2$  is formed

**Explanation:**  $CaCO_3$  and  $Cl_2$  is produced when Bleaching powder is treated with  $CO_2$ .

$$Ca(OCl)_2 \ + \ CO_2 
ightarrow CaCO_3 \ + \ Cl_2 \uparrow$$

- 6. Balanced equation:  $XeF_6 + H_2O 
  ightarrow XeO_2F_2 + 4HF$
- 7. Higher the value of reduction potentials, stronger is the oxidising behaviour. Therefore,  $BrO_4^-$  is the strongest oxidizing agent.

8.  $H_3PO_3$  on heating undergoes self-oxidation-reduction

$$4H_3\overset{+3}{P}O_3 o P\overset{-3}{H_3} + 3H_3\overset{+5}{P}O_4 \ phosphorons \ acid \ acid$$

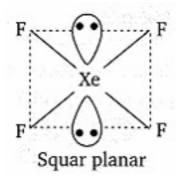
9. Halogens have the smallest size in their respective periods and therefore high effective nuclear charge. As a consequence, they readily accept one electron to an acquine noble gas electronic configuration.

10. i. 
$$C_2H_4 + 3O_2 o 2CO_2 + 2H_2O_2$$
 $Ethane Oxygen Carbondioxide Water$ 
ii.  $4Al + 3O_2 o 2Al_2O_3$ 
Aluminium  $Oxygen Alumina$ 

11. It reduces Fe (III)salt to Fe (II) salt.

$$2\mathrm{Fe^{3^+}}$$
 +  $\mathrm{SO_2}$  +  $2\mathrm{H_2O} \rightarrow 2\mathrm{Fe^{2^+}}$  +  $SO_4^{2^-}$  +  $4\mathrm{H^+}$ 

12. The square planar structure of  $XeF_4$  is shown below:



13. The phosphorus acid is a dibasic acid having an oxidation state of P = +3 as shown below;

$$H \xrightarrow{\begin{array}{c} O \\ \parallel \\ P \\ \end{array}} OH$$

14. 
$$H_2SO_{4(aq)} + H_2O_{(l)} \rightarrow H_3O_{(aq)}^+ + HSO_{4(aq)}^-; \ K_{a_1} > 10$$
 $HSO_4^- + H_2O \rightarrow H_3O^+ + SO_4^{2-}, \ K_{a_2} = 1.2 \times 10^2$ 
It can be noticed that  $K_{a_1} >> K_{a_2}$ .

This is because a neutral  ${\rm H_2SO_4}$  has a much higher tendency to lose a proton than the negatively charged  $HSO_4^-$ . Thus, the former is a much stronger acid than the latter.

15. i. Due to the presence of lone pair of electrons on the centre atom both  $NH_3$  and  $PH_3$  are Lewis Bases. When  $NH_3$  or  $PH_3$  accepts a proton, an additional N - H or P - H bond is formed.

$$H_3N + H^+ \rightarrow NH_4^+, H_3P + H^+ \rightarrow PH_4^+$$

Due to smaller size of N than P, N - H bond thus formed is much stronger than P - H bond. As a result  $NH_3$  has more tendency than  $PH_3$  to accept a proton. Therefore,  $NH_3$  is a stronger base than  $PH_3$ .

- ii. The property of catenation depends upon the strength of the element element bond. Since sulphur S S bond strength is much more than O O bond strength. So sulphur has greater tendency for catenation than oxygen.
- iii. Since  $F_2$  has smaller size than  $Cl_2$  and there is absence of d-orbital in fluorine, that's why  $F_2$  is stronger oxidizing agent than  $Cl_2$ .