

I(1):

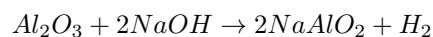
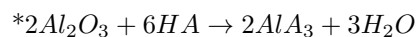
Element with a greater number of protons and a smaller atomic radius has larger electronegativity. Among the four elements, H and F have the smallest atomic radius. Although the atomic radius of F is slightly larger than that of H, it has many more protons than H. Therefore, F  $\boxed{(2)}$  has the largest electronegativity.

Note: When compared with period 2 elements, the electronegativity of H is slightly greater than that of B (Boron) but less than other elements with larger atomic numbers.

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I(2):

$Al_2O_3$   $\boxed{(2)}$  can undergo reaction with both acid and base\*. Therefore, it is amphoteric.



I(3):

1) H-Cl bond is polar.

2) H-O bonds are polar.  $H_2O$  molecules are in V-shape, hence  $H_2O$  molecules are also polar as the vector sum of dipole moments is not equal to 0.

3) C=O bonds are polar. However,  $CO_2$  molecules are in linear shape, the polarities cancelled out. Therefore,  $CO_2$  molecules are nonpolar.

4) N-H bonds are polar.  $NH_3$  molecules are in the shape of a trigonal pyramid, hence  $NH_3$  molecules are also polar as the vector sum of dipole moments is not equal to 0.

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I(4):

1) He is a gas (He is used to fill up balloons)

2) S is a solid (crystal)

3) Hg is a liquid (Hg is the liquid inside thermometer)

4) Pb is a solid

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I(5):

1) As Cl has a larger number of electron shells, the atomic radius of Cl is larger than that of F.

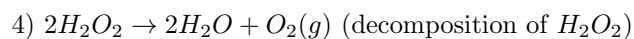
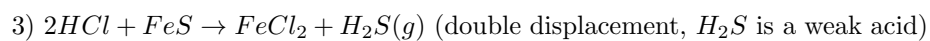
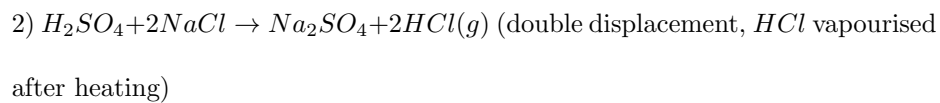
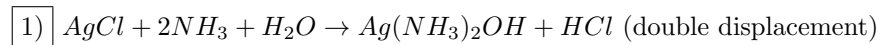
2) True.

3) As the atomic radius of Cl is larger than that of F, its electrons are attracted less strongly and can more easily be given out. Therefore, Cl is a strong reducing agent (i.e. a weaker oxidising agent).

4) Both of them can form oxides ( $OF_2$  and  $OCl_2$ ).

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I(6):



I(7):

1) The predominant bonding force of ionic crystals is ionic bond.

2) False.

3)  $\boxed{\text{Those dissociated ions can attract (or repulse) by charges and hence conduct electricity.}}$

4) Ionic crystals are brittle, as the repulsive force due to the same charge will break the crystal easily in case there are any displacements of layers.

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II:

(1): There are  $H^+$  ions,  $OH^-$  ions and  $Na^+$  ions in the aqueous solution.

As  $OH^-$  has the highest reducing power among them, it is oxidised to  $O_2$ .

As  $H^+$  has the highest oxidising power among them, it is reduced to  $H_2$ .

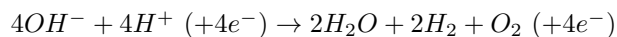
Therefore,  $\boxed{H_2}$  is formed at the cathode.

(2):

The half equation at the anode:  $4OH^- \rightarrow 2H_2O + O_2 + 4e^-$

The half equation at the cathode:  $2H^+ + 2e^- \rightarrow H_2$

Therefore, the full equation of the reaction is



When a current of 9.65A flowed for 2 hours, a charge of

$$9.65 \cdot 2 \cdot 60 \cdot 60 = 19.3 \cdot 3600 \text{ C passed through the circuit, i.e. } \frac{19.3 \cdot 3600}{96500} = 0.72 \text{ mol}$$

of electrons passed through the circuit.

By the equation, the molar ratio  $e^- : O_2 = 4 : 1$ . Therefore,  $\frac{0.72}{4} = \boxed{0.18} \text{ mol}$

of substance formed at the anode.

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III:

(1): Referring to the table:

Chemical	$N_2O_4$	$NO_2$
Number of moles before equilibrium	1.0	0
Number of moles after equilibrium	$1.0 - \frac{1.2}{2} = 0.4$	1.2

As partial pressure is proportional to molar fraction, we have the partial pres-

$$\text{sure of } N_2O_2 = 1.00 \times 10^5 \cdot \frac{0.4}{0.4+1.2} = 2.5 \times 10^4 \text{ Pa.}$$

(2): Based on (1), we have the partial pressure of

$$NO_2 = 1.0 \times 10^5 - 2.5 \times 10^4 = 7.5 \times 10^4 \text{ Pa.}$$

$$\text{Therefore, the pressure equilibrium constant} = \frac{(7.5 \times 10^4)^2}{2.5 \times 10^4} \approx \boxed{2.3 \times 10^5} \text{ Pa.}$$

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IV:

(1): Under heating,  $H_2O_2$  oxidises  $Cr^{3+}$  to  $CrO_4^{2-}$  and it itself becomes  $OH^-$ .

Therefore,  $Fe^{3+}$  is precipitated as  $Fe(OH)_3$ .

By adding acetic acid,  $OH^-$  ions are completely neutralised and  $H_2CrO_4$  formed so that the solution becomes weak acid.

When  $Pb(CH_3COO)_2$  is added, precipitation reaction occurs which gives  $PbCrO_4$  precipitate.

(2): Similar to a diamond, the number of nearby silicon atoms to one silicon atom is 4.

The thermochemical equation of formation of  $SiH_4$  is  $Si + 2H_2 \rightarrow SiH_4$ .

On the product side, 4 Si-H bonds are formed, hence  $4 \cdot 318 = 1272 \text{ kJ/mol}$  of energy is released.

On the reactant side, Si crystal is converted to Si atoms, which absorbs 439 kJ/mol of energy. Also, two H-H bonds are broken, which absorbs

$2 \cdot 436 = 872 \text{ kJ/mol}$  of energy.

Given the above, during the formation of  $SiH_4$ ,  $439+872=1311 \text{ kJ/mol}$  of energy is absorbed and  $1272 \text{ kJ/mol}$  of energy is released. Therefore, the heat of formation is  $1272-1311=-39 \text{ kJ/mol}$ .

Note: "Heat of formation" here should refer to the heat released during formation, but not the enthalpy change. Therefore, it should take the negative

sign.

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V:

(1): Consider the mass of C atom in  $CO_2$  formed, we have the mass of C atom in ester B  $= 7.92 \cdot \frac{12}{12+2 \cdot 16} = 2.16 \text{ mg}$ .

Consider the mass of H atom in  $H_2O$  formed, we have the mass of H atom in ester B  $= 3.24 \cdot \frac{2 \cdot 1.0}{2 \cdot 1.0 + 16.0} = 0.36 \text{ mg}$ .

Therefore, the mass of the O atom in ester B  $= 3.48 - 2.16 - 0.36 = 0.96 \text{ mg}$ . The

molar ratio C:H:O in ester B  $= \frac{2.16}{12.0} : \frac{0.36}{1.0} : \frac{0.96}{16.0} = 0.18 : 0.36 : 0.06 = 3:6:1$ .

Therefore, the empirical formula of B is  $C_3H_6O$  (1).

(2): Let the molecular formula of B be  $C_{3n}H_{6n}O_n$ .

Consider the molecular weight of B, we have  $110 < n(3 \cdot 12.0 + 6 \cdot 1.0 + 16.0) < 118$ .

Therefore, we have  $n = 2$ , i.e. the molecular formula of B is  $C_6H_{12}O_2$  (4).

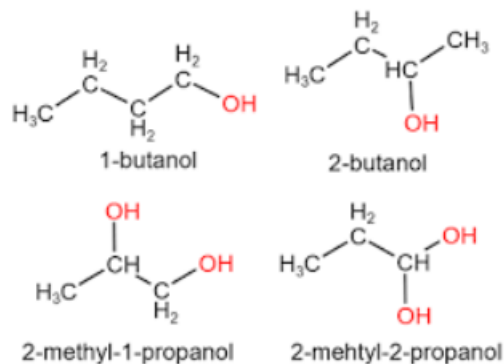
(3): As A undergoes esterification with acetic acid, it is an alcohol (4).

(4): The carboxylic acid used in the esterification is acetic acid,  $H_3CCOOH$ .

Therefore, the condensed formula of B is  $H_3CCOOC_4H_9$ , which suggests the alcohol, A, has a molecular formula of  $C_4H_{10}O$  (4).

(5): The number of structural isomers exist for B is equal to the number of

structural isomers exist for A. For A, we have 4 structural isomers:



Therefore, there are 4 structural isomers for B.

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VI:

With  $H_2SO_4$  catalyst,  $-NO_2$  substituted  $-H$  in benzene and gave nitrobenzene

(9). Reduced by  $Sn/HCl$ , it becomes aniline (1). Undergoing diasotiasation

with  $NaNO_2/HCl$ , it becomes (15). Heat it with water, we get phenol (11).

With  $Fe$  catalyst,  $-Cl$  substituted  $-H$  in benzene and gives chlorobenzene (7).

Under high temperature and high pressure,  $-OH$  substituted  $-Cl$  and on the other hand  $Na^+$  displaced  $H^+$ , which turns out giving (14).

Treating benzene with conc.  $H_2SO_4$ , we get benzene sulfuric acid (8). Neu-

tralise it with  $NaOH$ , we get (12). Moreover, undergoing alkali fusion, it becomes (14).

For the above two paths, (14) becomes phenol after being treated with acid ( $CO_2 + H_2O$ ). Or it becomes phthalic acid (5) after  $-COOH$  group is substituted  $-H$  under high temperature and pressure.

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VII:

(1):

☐ True

2) False

3)  $CaCO_3 + 2H_2O \rightarrow Ca(OH)_2 + H_2CO_3$ . No acetylene is formed.

4) False

☐  $C_2H_2 + H_2O \rightarrow H_2C = CHOH$ . As  $H_2C = CHOH$  is very unstable, it turns to  $H_3CCHO$  (acetaldehyde) immediately.

6)  $C_2H_2 + Br_2 \rightarrow C_2H_2Br_2$ . An Additional reaction occurs.

(2):

1) Ketones have a C=O double bond.

2) Aniline contains the benzene ring, which contains C=C double bonds.

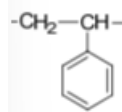
3) Carboxylic acids contain a C=O double bond.

4) Alkenes contain C=C double bond

☐ Alkanes contain only single bonds.

(3):

The right-hand side of the repeating unit:





Which is a styrene whose double bond is opened.

Consider the left-hand side of the repeating unit:  $-CH_2-CH=CH-CH_2-$ ,

before it opens the double bonds, it should be  $CH_2=CH-CH=CH_2$ , i.e.

1,3-butadiene (5).

Note: 1,3-butadiene is involved in the synthesis of most of the synthetic gums.