

| Universal gas constant $\mathrm{R}=8.314 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}$ | Avogadro constant $\mathrm{N}_{\mathrm{A}}=6.022 \times 10^{23} \mathrm{~mol}^{-1}$ |
| :--- | :--- |
| Planck's constant $\mathrm{h}=6.626 \times 10^{-34} \mathrm{Js}$ | Velocity of light $\mathrm{C}=3 \times 10^{8} \mathrm{~ms}^{-1}$ |

1. Select the element which is having the highest second ionization energy,
2. $C$
3. $N$
4. 0
5. $F$
6. $S$
7. Select the molecule which is having the largest number of lone pairs in it.
8. $S F_{4}$
9. $\mathrm{CCl}_{4}$
10. $\mathrm{HClO}_{4}$
11. $\mathrm{H}_{3} \mathrm{PO}_{4}$
12. $O F_{2}$
13. The Order of increasing bond angle of the species, $\mathrm{N}^{+} \mathrm{O}_{2}, \mathrm{NO}_{2}, \mathrm{NO}_{2}^{-}, \mathrm{NO}_{3}^{-}$
14. $\mathrm{NO}_{3}^{-}<\mathrm{NO}_{2}<\mathrm{NO}_{2}^{-}<\mathrm{N}^{+} \mathrm{O}_{2}$
15. $\mathrm{NO}_{2}^{-}<\mathrm{NO}_{3}^{-}<\mathrm{N}^{+} \mathrm{O}_{2}<\mathrm{NO}_{2}$
16. $\mathrm{NO}_{2}^{-}<\mathrm{NO}_{2}<\mathrm{NO}_{3}^{-}<\mathrm{N}^{+} \mathrm{O}_{2}$
17. $\mathrm{NO}_{2}^{-}<\mathrm{NO}_{3}^{-}<\mathrm{NO}_{2}<\mathrm{N}^{+} \mathrm{O}_{2}$
18. $\mathrm{NO}_{2}<\mathrm{NO}_{2}^{-}<\mathrm{NO}_{3}^{-}<\mathrm{N}^{+} \mathrm{O}_{2}$
19. IUPAC name of the compound,

20. ethyl-2-amino - 3 - methylhex - 3 -en - 5 - oneoate
21. ethyl 2 -amino - 3 - methyl-5-oxohex - 3 - enoate
22. ethyl-3-methyl-2-amino - 5 - oxohex -3 - enoate
23. ethyl 2 -ammine -3 -methylhex -3 -en -5 - onoate
24. ethyl 2 -ammine -3 -methyl -5 -oxohex -3 -enoate
25. Select the statement which is not true?
26. Temperature, concentration, pressure, physical state and catalysts affect for the reaction rate.
27. For a given reaction, the rate of removal of each of the reactant is equal to the rate of formation of each of the product.
28. The rate with respect to a reactant depends on the stoichiometric coefficient of the relevant substance.
29. The rate of a reaction is the change of concentration occurred within a unit time.
30. The time taken to occur a given definite change can be used to measure the rate.

31. The product / products obtained when $\mathrm{CH}_{3}-\mathrm{C}-\mathrm{CH}_{2} \mathrm{CH}_{3}$ undergoes the self condenzation at the presence of aqueous NaOH is / are?
A. $\mathrm{CH}_{3}-\stackrel{\stackrel{\mathrm{OH}}{\mathrm{C}} \underset{\mathrm{C}}{\mathrm{C} \mathrm{H}_{2} \mathrm{CH}_{3}} \mathrm{CH}_{2} \mathrm{CH}_{2}-\stackrel{\text { I }}{\mathrm{O}} \mathrm{C}-\mathrm{CH}_{3}}{ }$



32. A only
33. A and B only
34. B and D only
35. C and D only
36. D only
37. Consider the following reaction, taking place at 298 K ,

$$
A_{2}(\mathrm{~g})+3 B_{2}(\mathrm{~g}) \rightarrow 2 A B_{3}(\mathrm{~g}) ; \Delta H^{\theta}<0
$$

Which of the following statements is true regarding the above reaction?

1. For all temperatures, $\Delta G^{\theta}$ of the reaction takes a negative value.
2. At the low temperatures the reaction is spontaneous.
3. When the reaction is taking place, the entropy of the surroundings decreases.
4. When the reaction is taking place, the entropy of the system increases.
5. For all the temperatures the reaction is not spontaneous.
6. At 298 K the ratio of $\frac{\left[\mathrm{Pb}^{2+}(a q)\right]}{\left[\mathrm{Ca}^{2+}(a q)\right]}$ in a saturated solution of $\mathrm{PbCO}_{3}$ and $\mathrm{CaCO}_{3}$ is,

At $298 \mathrm{~K} \quad K_{s p}\left(\mathrm{PbCO}_{3}(s)\right)=6 \times 10^{-14} \mathrm{~mol}^{2} \mathrm{dm}^{-6}$

$$
K_{s p}\left(\mathrm{CaCO}_{3}(s)\right)=3 \times 10^{-9} \mathrm{~mol}^{2} \mathrm{dm}^{-6}
$$

1. $\frac{\sqrt{6}}{\sqrt{3}}$
2. $\frac{1 \times 10^{-2}}{\sqrt{5}}$
$32 \times 10^{-4}$
3. $0.5 \times 10^{-5}$
4. $2 \times 10^{-5}$
5. The compound can be used to distinguish both solutions of $\mathrm{MgCl}_{2}$ and $\mathrm{MgSO}_{4}$ is,
6. $\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}(\mathrm{aq})$
7. $\mathrm{Na}_{2} \mathrm{CO}_{3}(\mathrm{aq})$
8. $\mathrm{Ba}\left(\mathrm{NO}_{3}\right)_{2}(\mathrm{aq})$
9. $\mathrm{Na}_{3} \mathrm{PO}_{4}(\mathrm{aq})$
10. $\mathrm{NaOH}(\mathrm{aq})$
11. A mixture of $N_{2^{(g)}}$ and $O_{2^{(g)}}$ is obtained when nitrous oxide $\left(N_{2} O\right)(g)$ is undergoing the thermal decomposition. Here $56 \%$ of $\mathrm{N}_{2} \mathrm{O}(\mathrm{g})$ is, under $1.2 \times 10^{5} \mathrm{~Pa}$ pressure. The partial pressure of the $N_{2}(g)$ formed is, (Consider that the temperature and volume are constants
12. $0.41 \times 10^{5} \mathrm{~Pa}$
13. $0.53 \times 10^{5} \mathrm{~Pa}$
14. $0.26 \times 10^{5} \mathrm{~Pa}$
15. $0.8 \times 10^{5} \mathrm{~Pa}$
16. $0.4 \times 10^{5} \mathrm{~Pa}$
17. The increasing order of melting points of the following substances is,
$\mathrm{H}_{2} \mathrm{O}, \mathrm{NH}_{3}, \mathrm{CH}_{4}, \mathrm{HF}, \mathrm{SbH}_{3}$
18. $\mathrm{CH}_{4}<\mathrm{NH}_{3}<\mathrm{HF}<\mathrm{H}_{2} \mathrm{O}<\mathrm{SbH}_{3}$
19. $\mathrm{CH}_{4}<\mathrm{SbH}_{3}<\mathrm{NH}_{3}<\mathrm{H}_{2} \mathrm{O}<\mathrm{HF}$
20. $\mathrm{SbH}_{3}<\mathrm{CH}_{4}<\mathrm{NH}_{3}<\mathrm{H}_{2} \mathrm{O}<\mathrm{HF}$
21. $\mathrm{CH}_{4}<\mathrm{SbH}_{3}<\mathrm{NH}_{3}<\mathrm{HF}<\mathrm{H}_{2} \mathrm{O}$
22. $\mathrm{SbH}_{3}<\mathrm{CH}_{4}<\mathrm{NH}_{3}<\mathrm{HF}<\mathrm{H}_{2} \mathrm{O}$
23. Which of the following statements is false regarding the colours of the complex ions, formed by the cations of 3d block?
24. $\left[\mathrm{CuCl}_{4}\right]^{2-}(a q)$ is yellow in colour.
25. $\left[\mathrm{NiCl}_{4}\right]^{2-}(a q)$ is yellow in colour.
26. $\left[\mathrm{CoCl}_{4}\right]^{2-}(a q)$ is blue in colour.
27. $\left[\mathrm{Ni}\left(\mathrm{NH}_{3}\right)_{6}\right]^{2+}(\mathrm{aq})$ is dark blue in colour.
28. $\left[\mathrm{Co}\left(\mathrm{NH}_{3}\right)_{6}\right]^{2+}(\mathrm{aq})$ is dark blue in colour.
29. $A$ and $B$ liquids mix together to form an ideal solution. This solution mixture exists in equilibrium with it's vapour inside a closed vessel at a constant temperature. At that temperature the saturated vapour pressure of $A$ and $B$ are $P_{A}^{0}$ and $P_{B}^{0}$ respectively. If the mole fraction of $A$ in the liquid phase is $X_{A}$, then the mole fraction of A in vapour phase is,
30. $\frac{P_{A}^{0} \times A}{\left(P_{A}^{0}-P_{B}^{0}\right) X_{A}+P_{B}^{0}}$
31. $\frac{P_{B}^{0}\left(1-X_{A}\right)}{P_{A}^{0}\left(1-X_{A}\right)+P_{B}^{0}}$
32. $\frac{P_{B}^{0} \mathrm{X} A}{P_{A}^{0} \times A+P_{B}^{0}\left(1-X_{A}\right)}$
33. $\frac{P_{A}^{0} \mathrm{X} A}{P_{A}^{0}+P_{B}^{0}}$
34. $\frac{P_{A}^{0}\left(1-X_{A}\right)}{P_{A}^{0} \times A+P_{B}^{0}\left(1-X_{A}\right)}$
35. The correct statement from the followings is,
36. In the emission spectrum of atomic hydrogen, the distance between spectral lines is decreasing to the direction of increasing wave Length.
37. Among the species $\mathrm{N}^{+} \mathrm{O}_{2}, \mathrm{NO}_{3}^{-}$and $\mathrm{NO}_{2}^{-}$the electronegativity of the central $\mathrm{N}_{1}$ atiom is highest in $\mathrm{NO}_{2}^{-}$.
38. The highest reducing property is shown by $K$ among $A l, N a, M g, K$
39. The ability of hydrolysis is maximum for $\mathrm{AsCl}_{3}$ among $\mathrm{NCl}_{3}, \mathrm{PCl}_{3}, \mathrm{AsCl}_{3}$
40. Only H bonds exist among the molecules of $\mathrm{CH}_{3} \mathrm{COOH}$
41. At 298 K Calculate the $p H$ of the buffer solution which is formed by mixing $50 \mathrm{~cm}^{3}$ of $0.2 \mathrm{~mol} \mathrm{dm}{ }^{-3} \mathrm{CH}_{3} \mathrm{COOH}$ acid solution and $50 \mathrm{~cm}^{3}$ of $0.1 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{KOH}$ solution.
At $298 \mathrm{~K} \quad K_{a_{\left(\mathrm{CH}_{3} \mathrm{COOH}\right)}}=1.8 \times 10^{-5} \mathrm{~mol} \mathrm{dm}{ }^{-3}$
42. 4.74
43. 5.74
44. 5.26
45. 4.26
46. 5.32
47. The standard electrode potentials of two standard electrodes formed by $A$ and $B$ are given below.
$E^{\theta}\left(A_{(a q)}^{+} / A_{(s)}\right)=+0.52 \mathrm{~V}$

$$
E_{\left(B^{3+}(a q) / B(s)\right)}^{\theta}=-0.74 \mathrm{~V}
$$

Which is true regarding the electro chemical cell formed using the above half cells? ,

1. Electrode $A$ is the anode and electrode $B$ is the cathode.
2. Electrode $A$ is the cathode and electrode $B$ is the anode and the electromotive force of that cell is to +0.22 V .
3. When the cell is functioning electrons are following from the electrode $A$ towards the electrode $B$ through the external circuit.
4. Electrode $B$ is the anode and the electro motive force of the cell is +1.26 V .
5. Electrode $B$ is the anode and when the cell is functioning the mass of the electrode $B$ is increasing.
6. Which of the following statements is false?
7. The fractional distillation can be used to obtain distilled water from a salt solution.
8. The fractional distillation can be used to separate the volatile compounds in a liquid mixture.
9. The simple distillation is used to separate the components in a solution, which is formed by dissolving a non volatile substance in a volatile solvent.
10. In order to separate two liquids by the fractional distillation there should be a considerable difference between their boiling points.
11. In the fractional distillation the mole fraction of the substance in the solution which has the high boiling point increases gradually.
12. Which of the following statements is true regarding the given compound ?
13. More acidic than phenol.
14. Less acidic than phenol.

15. reacting with $\mathrm{CH}_{3} \mathrm{COOH}$ to form esters.
16. reacting with $\mathrm{Na}_{2} \mathrm{CO}_{3}$ to liberate $\mathrm{CO}_{2}(\mathrm{~g})$
17. Undergoing nucleophilic substitution reaction with $\mathrm{CH}_{3} \mathrm{COCl}$ to form esters.
18. Consider the reaction, $A_{(g)}+B_{(g)}+$ energy $\rightleftharpoons C(g)$

At 298 K the equal number of moles of $A_{(g)}$ and $B_{(g)}$ are mixed inside a rigid vessel and allowed it to reach the equilibrium. Select the method which can not be used to increase the yield of $C_{(g)}$

1. Increasing the temperature of the system.
2. Addition of a certain amount of $B_{(g)}$ to the system.
3. Removal of $C_{(g)}$ from the system by liquifying.
4. Increasing the pressure of the system by adding an inert gas at constant temperature.
5. Addition of a catalysts.
6. A metal mixture in which Na and Cu are mixed, contains $64.25 \%$ of Na by mass. When 3.58 g of that metal mixture is added to $250 \mathrm{~cm}^{3}$ of water. What is the pH value of that solution at 298 K . (At $298 K K_{w}=1 \times 10^{-14} \mathrm{~mol}^{2} \mathrm{dm}^{-6}$ and $\mathrm{Cu}=64, \mathrm{Na}=23$ )
7. 13.4
8. 13.6
9. 1.6
10. 0.6
11. 0.4
12. The products of the reaction,

excess
(i) $\mathrm{CH}_{3} \mathrm{MgBr}$
$\xrightarrow[\text { (ii) } \mathrm{H}_{2} \mathrm{O}]{ }$
13. 


$\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$
2.

3.

4.

5.

22. At 298 K when $0.1 \mathrm{moldm}^{-3}$ mono basic weak with $p H=4$ is diluted hundred times, $p H$ value of the resultant solutions is,

1. 4.5
2. 5
3. 6
4. 6.5
5. 5.5
6. For the electroplating of Ni metal 0.5 A constant current is passed through a $\mathrm{NiCl}_{2}$ solution, during 2 hours. The maximum mass of Ni that can be coated is, $\left(1 F=96500 \mathrm{C} \mathrm{mol}^{-1}, \mathrm{Ni}=58.7\right)$
7. 2.56 g
8. 5.87 g
9. 1.09 g
10. 58.7 g
11. 1.17 g
12. Consider the following reaction series.

structures obtained for $\mathrm{A}, \mathrm{B}, \mathrm{C}$, and D respectively.
(1)

13. Which of the followings is not true regarding the reactions of Diazonium Salt.
(1)

(4)

(2)

(3)

(5)

14. Which of the following compounds can be answered for all the following observations?

- Shows the geometrical isomerism.
- The product obtained in the addition of $H C N$ shows enantiomerism.
- Gives a dark yellow colour precipitate with Bready's reagent ( $2,4-$ DNP)
- Does not give a brick red colour precipitate with Fehling's solution.
(1)

(2)

(3)

(4)

(5)


27. The chemical formula of hexaaquairon(iii) sulfate is,
28. $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right] \mathrm{SO}_{4}$
29. $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]_{2}\left(\mathrm{SO}_{4}\right)_{3}$
30. $\left[\mathrm{Fe}\left(\mathrm{OH}_{2}\right)_{6}\right]_{2}\left(\mathrm{SO}_{4}\right)_{3}$
31. $\left[\mathrm{Fe}\left(\mathrm{OH}_{2}\right)_{6}\right] \mathrm{SO}_{4}$
32. $\left[\mathrm{Fe}\left(\mathrm{OH}_{2}\right)_{6}\right]\left(\mathrm{SO}_{4}\right)_{3}$
33. Which of the following statements is false regarding S- block elements?
34. Only $L i$ of the first group reacts with $N_{2}$ gas.
35. $L i$ forms only $L i_{2} O(s)$ with excess $O_{2}(g)$.
36. Na reacts with dil. $\mathrm{H}_{2} \mathrm{SO}_{4}$ and liberates $\mathrm{H}_{2}(\mathrm{~g})$ rapidly.
37. All $S$ block elements react with $H_{2}(\mathrm{~g})$ and form metal hydrides.
38. Except $B e$ rest of all the metals of the second group react with $N_{2}$ gas.
39. Which of the following statements is false regarding the $d$ block elements.
40. Unlike $s$ block elements, $d$ block elements can show variable oxidation states.
41. The electronegativity of $d$-block elements is less than the electronegativity of $s$-block elements of the same period.
42. Metallic properties $d$-block elements is greater than the metallic properties of $s$-block elements.
43. $d$-block elements act as catalysts.
44. $d$ - block elements can form acidic / basic / amphoteric oxides.
45. Consider the following reactions.


Which of the following responses shows a step of the mechanism of the above reaction correctly?

1. $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}-\mathrm{Cl}+\mathrm{AlCl}_{3} \rightarrow\left(\mathrm{CH}_{3}\right)_{3} \mathrm{C}-\mathrm{Cl}^{+}-\overline{\mathrm{Al}} \mathrm{Cl}_{3}$
2. 



4.

5.


- For each of the questions 31 to 40 , one or more responses out of the four responses (a), (b) , (c) and (d) given is /are correct. Select the correct response/responses in accordance with the instructions given on your answer sheet, mark
(1) If only (a) and (b) are correct.
(2) If only (b) and (c) are correct.
(3) If only (c) and (d) are correct.
(4) If only (d) and (a) are correct.
(5) If any other number or combination of responses is correct.

Summary of above Instructions,

| 1 | 2 | 3 | 4 | 5 |
| :---: | :---: | :---: | :---: | :---: |
| Only (a) and (b) <br> are correct | Only (b) and (c) <br> are correct | Only (c) and (d) <br> are correct | Only (a) and (d) <br> are correct | Any other number or <br> combination of <br> responses is correct |

31. Consider the titration of $0.1 \mathrm{moldm}^{-3} \mathrm{HCl}$ in burette and $25 \mathrm{~cm}^{3}$ of $0.1 \mathrm{~mol} \mathrm{dm}{ }^{-3} \mathrm{NH}_{4} \mathrm{OH}$ taken in to a titration flask at 298 K .
a) $\quad \mathrm{pH}$ value at the equivalence point is less than 7.
b) Consumed volume of HCl at the equivalence point is less than $25 \mathrm{~cm}^{3}$.
c) Phenolphthalein is a suitable indicator to determine the equivalence point.
d) Methyl arrange is a suitable indicator to determine the equivalence point.
32. Select the pairs of ions which show the same colour in the aqueous solution.
(a) $\mathrm{Mn}^{2+}, \mathrm{Ni}^{2+}$
(b) $\mathrm{V}^{3+}, \mathrm{Cr}^{3+}$
(c) $\mathrm{Cu}^{2+}, \mathrm{Co}^{2+}$
(d) $\mathrm{Mn}^{2+}, \mathrm{Co}^{2+}$
33. At 298 K , some amount of $\mathrm{CaCl}_{2}$ is add to a saturated solution of $\mathrm{Ca}(\mathrm{OH})_{2}$. Which of the followings is / are occured in this solution.
(a) $\left[\mathrm{Ca}^{2+}(a q)\right]$ increasing
(b) $\left[\mathrm{OH}^{-}(\mathrm{aq})\right]$ increasing
(c) $\left[H^{+}(a q)\right]$ does not change.
(d) $p H$ value is decreasing.
34. At 800 K , for the equilibrium, $\mathrm{N}_{2}(g)+3 \mathrm{H}_{2}(g) \rightleftharpoons 2 \mathrm{NH}_{3}(g)$ the equilibrium constant $K_{c}=0.08 \mathrm{dm}^{6} \mathrm{~mol}^{-2}$ At $800 \mathrm{~K}, 4 \mathrm{~mol}$ of $\mathrm{N}_{2}(\mathrm{~g}), 5 \mathrm{~mol}$ of $\mathrm{H}_{2}(\mathrm{~g})$ and 2.5 mol of $\mathrm{NH}_{3}(\mathrm{~g})$ are placed in a rigid vessel of $5 \mathrm{dm}^{3}$. Which of the followings is / are true regarding the above equilibrium.
(a) Initially $Q_{C}<K_{C}$ and the reaction shifts to left.
(b) Initially $Q_{C}>K_{C}$ and the reaction shifts to right.
(c) Initially, $Q_{C}<K_{C}$ and the reaction shifts to right by forming excess $\mathrm{NH}_{3}$.
(d) Initially $Q_{C}<K_{C}$ and the reaction shifts to right consuming $H_{2}(g)$ and $N_{2}(g)$.

(a) All the $C$ atoms of the molecules exist in the same plane.
(b) $\mathrm{p}, \mathrm{q}$ and r carbon atoms exist in a same line.
(c) All $H$ atoms of the molecule exist in the same plane.
(d) Above compound shows diasteriomerism.
35. Consider the following two compounds.

A

B

C

The more correct statement/s regarding the above compound is / are.
(a) Rate of nucleophilic substitution reaction of $B$ is less than $C$.
(b) A undergoes nucleophilic substitution reactions.
(c) Rate of nucleophilic substitution reaction of $A$ is greater than $B$.
(d) While $B$ undergoes nucleophilic substitution reactions of a single step $C$ undergoes nucleophilic substitution reactions of two steps.
37. Which of the following solutions turns / turn red litmus blue.
(a) $\mathrm{NH}_{4} \mathrm{Cl}$
(b) $L i F$
(c) $\mathrm{CH}_{3} \mathrm{COOK}$
(d) $\mathrm{CH}_{3} \mathrm{COONa}$
38. Select the factor / factors which affects / affect the electromotive forces of an electro chemical cell.
(a) temperature
(b) Cross section area of the electrodes
(c) the distance between the electrodes
(d) nature of the electrode used.
39. Which of the followings can be used to distinguish $\mathrm{SO}_{2}(g)$ and $\mathrm{H}_{2} \mathrm{~S}(\mathrm{~g})$.
(a) $\mathrm{H}^{+} / \mathrm{KMnO}_{4}(\mathrm{aq})$
(b) $\mathrm{H}^{+} / \mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}$
(c) wet colourful petal of a flower.
(d) nature of the electrode used
40. Which of the following statement $/ \mathrm{s}$ is / are false regarding an aliquot of an ideal gas.
(a) Total energy of the molecules is changed, in the molecular collisions occurred at constant temperature
(b) There are not attraction forces among gas molecules.
(c) At constant temperatures mean kinetic energy of gas molecules is a constant.
(d) Gas molecules travel in all directions in the same speed.

In question numbers 41 to 50 , two statements are given in respect of each question. From the table given below, select the response out of the responses (1), (2), (3), (4) and (5) that best fits the two statements and mark appropriately on your answer sheet.

| $\mathbf{1}^{\text {st }}$ Statement | $\mathbf{2}^{\text {nd }}$ Statement | Response |
| :--- | :--- | :--- |
| True | True and $1^{\text {stststatement is explained correctly }}$ | 1 |
| True | True and $1^{\text {st }}$ statement is not explained correctly | 2 |
| True | False | 3 |
| False | True | 4 |
| False | False | 5 |
|  |  |  |


|  | $1{ }^{\text {st }}$ Statement | $2^{\text {nd }}$ Statement |
| :---: | :---: | :---: |
| 41. | The reaction rate is increased by the catalyzed. | The activation energy of the reaction is decreased by the catalysts. |
| 42. | The basicity of amides is less than that of amine. | The electron pair on nitrogen in the amide group is de localized on to the carbonyl group by the resonance. |
| 43. | The order of an elementary reaction is equal to the molecularity. | The number of molecules participated for the balanced chemical equation is the molecularity. |
| 44. | When the concentration of a monobasic weak acid is decreasing pH value is decreasing. | When the concentration of a monobasic weak acid is decreasing then its degree of dissociation is increasing. |
| 45. | Alkanes are not reactive towards the non polar reagents. | $C-C$ bond of the alkane is non polar and $C-H$ bond is polar. |
| 46. | At high temperature He gas reaches the ideal behaviour. | At higher temperatures the speed of the molecules is increasing, therefore the strength of the attractive forces is decreasing. |
| 47. | In acidic medium when $H_{2} S$ is passed, not only $\mathrm{Cu}^{2+}$ but also $\mathrm{Ni}^{2+}$ can be precipitated as its sulphide. | Because of the high $\mathrm{H}^{+}$concentration of the acidic medium concentration of $S^{2-}$ ions decreases relatively. |
| 48. | Acidity of the benzoic acid is greater than the acidity of phenol. | The stability of the phenate ion is greater than the stability of the benzoate ion. |
| 49. | The melting point of $\mathrm{MgCO}_{3}$ is less than the melting point of $\mathrm{BaCO}_{3}$. | The polarizing power of $\mathrm{Mg}^{2+}$ ion is less than the polarizing power of $\mathrm{Ba}^{2+}$ ion. |
| 50. | The conductivity of $0.1 \mathrm{moldm}^{-3} \mathrm{NaCl}$ is less than the conductivity of $0.1 \mathrm{~mol} \mathrm{dm}{ }^{-3} \mathrm{NaCl}$. | The electric conductivity of the solution depends only on the temperature on the nature of the solute. |

## ๔లిరిธిమ อథరి <br>  <br> Periodic Table



| 57 | 58 | 59 | 60 | 61 | 62 | 63 | 64 | 65 | 66 | 67 | 68 | 69 | 70 | 71 |
| :--- | :--- | :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| La | Ce | Pr | Nd | Pm | Sm | En | Gd | Tb | Dy | Ho | Er | Tm | Yb | Lu |
| 89 | 90 | 91 | 92 | 93 | 94 | 95 | 96 | 97 | 98 | 99 | IOO | 101 | 102 | 103 |
| Ac | Th | Pa | U | Np | Pu | Am | Cm | Br | Cr | Es | Fm | Md | No | Lr |



Index No :

* A Periodic Table is provided on page 16.
* Use of calculators is not allowed.
* Universal gas constant, $R=8.314 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$
* Avogadro constant, $N_{A}=6.022 \times 10^{23} \mathrm{~mol}^{-1}$

㫧 In answering this paper, you may represent alkyl groups in a condensed manner.

Example:


PART A - Structured Essay (pages 2-8)
Answer all the questions on the question paper itself.

* Write your answer in the space provided for each question. Please note that the space provided is sufficient for the answer and that extensive answers are not expected.

PART B and PART C - Essay (pages 9-15)

* Answer four questions selecting two questions from each part. Use the papers supplied for this purpose.
* At the end of the time allotted for this paper, tie the answers to the three Parts $\mathbf{A}, \mathbf{B}$ and $\mathbf{C}$ together so that Part $\mathbf{A}$ is on top and hand them over to the Supervisor.
* You are permitted to remove only Parts $\mathbf{B}$ and $\mathbf{C}$ of the question paper from the Examination Hall.

For Examiner's Use Only

| Part | Question No. | Marks |
| :---: | :---: | :---: |
| A | 1 |  |
|  | 2 |  |
|  | 3 |  |
|  | 4 |  |
| B | 5 |  |
|  | 6 |  |
|  | 9 |  |
|  | 9 |  |
|  | 10 |  |
| Total |  |  |
| Percentage |  |  |

Final Mark

| In Numbers |  |
| :--- | :--- |
| In Letters |  |

Code Numbers

| Marking Examiner 1 |  |
| :--- | :--- |
| Marking Examiner 2 |  |
| Checked by : |  |
| Supervised by : |  |

## Part A - Structured Essay

Answer all four questions on this paper itself. ( Each question carries 10 marks)
(01) (a) (i) Mention the oxidation state and hybridization shown by the central N atom in the following compounds and ions.

| Compound | $\mathrm{NO}_{2}$ | $\mathrm{NO}_{2} \mathrm{~F}$ | $\mathrm{NH}_{2} \mathrm{OH}$ | $\mathrm{N}_{2} \mathrm{O}$ | $\mathrm{NH}_{4}^{+}$ |
| :--- | :--- | :--- | :--- | :--- | :--- |
| 1. Oxidation number |  |  |  |  |  |
| 2. Hybridization |  |  |  |  |  |

(ii) Draw the acceptable Lewis structure for the molecule $\mathrm{N}_{2} \mathrm{O}$.
(iii) Draw the resonance structures for the above molecules expect the drawn in part II above. Mention the stabilities by giving reasons
(b) $\mathrm{KHF}_{2}$ is an ionic compound with a cation of one type and an anion of one type.
i. Write IUPAC name of $\mathrm{KHF}_{2}$.
$\qquad$
$\qquad$
ii. Write the chemical of the cation and the anion?
cation
anion
iii. Draw the structure of the anion and name the bond types of it.
$\qquad$
$\qquad$
iv. Answer the following questions regarding the compounds of $\mathrm{H}_{2} \mathrm{~S}$ and $\mathrm{CS}_{2}$.

1. Mention the secondary interaction type / types existing among each of the species. $\mathrm{H}_{2} \mathrm{~S}$ $\qquad$ $\mathrm{CS}_{2}$ $\qquad$
2. Which of the above molecule has the strongest secondary interactions?
$\qquad$
3. Which of the above compounds has the highest boiling point.?
4. Consider the combustion of each of the above compounds at the presence of excess $\mathrm{O}_{2}$. Write the balanced chemical equations for the above reactions.
(C) Mention whether the following statements are 'True" or 'False'
i. Unit of the rate constant depends on the order of the reactions.
( ................... )
ii. All the molecules which are having polar covalent bonds have dipole moments.(................... )
iii. Energy of an electron in a 3d orbital is greater than the energy of an electron in 4s orbital. ( ................... )
iv. pH value of $10^{-10} \mathrm{~mol} \mathrm{dm}^{-3}$ aqueous HCl solution at $25^{0} \mathrm{C}$ is 10 ( .................... )
v. Heat of the reaction of a chemical reaction depends on the temperature. ( ................... )
vi. Atomic radius of Cl is greater than that of Li . (................... )
vii. In the order of $\mathrm{NaCl}<\mathrm{MgCl}_{2}<\mathrm{AlCl}_{3}$, their melting points are increasing.( ................... )
(02) (a) The element M belongs to s block. Although the sulphate of M is more soluble be in water, the carbonate is insoluble in water. Although M does not react with cold water, it reacts with hot water.
(i) What is the element M ? $\qquad$
(ii) When the element $M$ is combusted in air it forms two compounds $Q_{1}$ and $Q_{2}$. Only $Q_{2}$ of them reacts with water. Write the chemical formulae of the compounds $Q_{1}$ and $Q_{2}$.
$Q_{1}$ $\qquad$ $Q_{2}$ $\qquad$
(iii) Write the balanced chemical equations for the followings.
5. Reaction of M with hot water.
6. Reaction of $\mathrm{Q}_{2}$ with water.
(iv) In a certain experiment the above element M is combusted in air resulting a mixture of $\mathrm{Q}_{1}$ and $\mathrm{Q}_{2}$. 1.0 g of that mixture is allowed to react with excess $\mathrm{H}_{2} \mathrm{O}$. Then a gas is liberated and that gas is allowed react completely with the $0.5 \mathrm{~mol} \mathrm{dm}{ }^{-3} \mathrm{HCl}$. The percentage of $\mathrm{Q}_{2}$ contained in 1.0 g of the above sample is $30.3 \%$ Calculate the volume of HCl reacted above.
(b) When 2.48 g of a hydrated inorganic salt X is heated, the anhydrous salt Y and 0.9 g of water is obtained as products. The following observations are given by an aqueous solution of Y .
7. Decolourises the solution of $\mathrm{I}_{2}$ dissolved in aqueous KI.
8. At the presence of HCl , forms a light yellow colour precipitate, liberating a colourlessgas with a pungent smell.
9. At the presence of aqueous $\mathrm{AgNO}_{3}$,forms a white colour, precipitate and it turns black within a short period of time.
10. A conc. HCl solution of Y imparts yellow colour to the flame in the flame test.
(i) What is the salt of Y?
$\qquad$
(ii) Write the balanced equations for the reactions relevant to observations 1, 2 and 3.
(iii) Derive the chemical formula of the hydrated compound X .
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) Write the balanced chemical equations for the following reactions.
(i) $\quad \mathrm{NaNH}_{2}+\mathrm{NH}_{4} \mathrm{Cl} \xrightarrow{-}$
(ii) $\mathrm{H}_{2} \mathrm{~S}+\mathrm{H}_{2} \mathrm{O}_{2} \rightarrow$
(iii) $\mathrm{Al}+\mathrm{OH}^{-}+\mathrm{NO}_{3}^{-} \rightarrow$
(iv) $\mathrm{FeCl}_{3}+\mathrm{H}_{2} \mathrm{~S} \rightarrow$
(03) (a) At $25^{0} \mathrm{Cp}^{\mathrm{K}_{\mathrm{b}}}$ of $\mathrm{NH}_{3}(\mathrm{aq})$ is 4.8 and $\mathrm{Kw}=10^{-14} \mathrm{~mol}^{2} \mathrm{dm}^{-6}$
(i) What do you mean by $\mathrm{p}^{\mathrm{K}_{\mathrm{b}}}$ ?
$\qquad$
$\qquad$
$\qquad$
(ii) Write the equilibrium attained when the aqueous $\mathrm{NH}_{3}$ is decomposed.
$\qquad$
$\qquad$
(iii) Write an expression for the dissociation constant ( $\mathrm{K}_{\mathrm{b}}$ ) of $\mathrm{NH}_{3}$ (aq).
(iv) According to the reaction $\mathrm{NH}_{4(\mathrm{aq})}^{+}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})} \rightleftharpoons \mathrm{NH}_{3(\mathrm{aq})}+\mathrm{H}_{3} \mathrm{O}_{(\mathrm{aq})}^{+}$. Write an expression for $\left(\mathrm{K}_{\mathrm{a}}\right)$ of $\mathrm{NH}_{4}^{+}(\mathrm{aq})$.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(v) Accordingly derive the relationship among $\mathrm{k}_{\mathrm{a}}, \mathrm{k}_{\mathrm{b}}$ and $\mathrm{k}_{\mathrm{w}}$.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(vi) At $25^{\circ} \mathrm{C}$ Calculate $\mathrm{p}^{\mathrm{K}_{\mathrm{b}}}$ of $\mathrm{NH}_{4}^{+}$(aq).
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(b) An aqueous solution contain NaOH and $\mathrm{Na}_{2} \mathrm{CO}_{3}$ is provided. Two portions of $50 \mathrm{~cm}^{3}$ from that solution are seperated and labelled as A and B. Those solutions are titrated as follows.

## Method 01

Portion A is titrated with $1.0 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{HCl}$ at the presence of phenolphthalein. The consumed volume of the acid at the endpoint is 10.0 ml .

## Method 02

A few drops of phenolphthalein is added to part B and $\mathrm{CO}_{2}(\mathrm{~g})$ is bubbled through the solution until the obtained red colour turns colourless. After that at this solution is titrated with $1.0 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{HCl}$ at the presence of methyl orange. The acid volume consumed at the endpoint is 15.0 ml .
(i) Write the balanced equations for the reactions taken place in the first titration.
(ii) Write the balanced equations for the reaction taken place, when $\mathrm{CO}_{2}(\mathrm{~g})$ is bubbled through the solution B in the method 02 .
$\qquad$
$\qquad$
$\qquad$
(iii) Write the balanced equation for the reaction taken place in the second titration?
$\qquad$
$\qquad$
$\qquad$
(iv) If the number of moles of NaOH in the initial mixture is x and the number of moles of $\mathrm{Na}_{2} \mathrm{CO}_{3}$ in it is y . Calculate the values of x and y .
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(04) (a) Write the structures of the major organic products formed by adding of the following reagents to an aqueous solution containing phenol and aniline.
i. $\mathrm{Br}_{2}(\mathrm{l})$


Product formed by Phenol
ii $\mathrm{CH}_{3} \mathrm{COCl}$


Product formed by Phenol


Product formed by aniline


Product formed by aniline
iii. Nmae the reactions carried out in (i) and (ii) above considering the type of the mechanism.
(i)
(ii) $\qquad$
iv. Write the mechanism for the reaction between phenol and $\mathrm{CH}_{3} \mathrm{COCl}$.
(b) Show how would you synthesis the organic compound material in not more than seven steps.

using

as the starting
(c) i Write the structures of the major organic products obtained in the following reactions.
1.

2. $\mathrm{C}_{6} \mathrm{H}_{5}-\mathrm{CH}=\mathrm{CH}-\mathrm{CH}_{3} \xrightarrow{\mathrm{HBr}}$
3. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOH} \xrightarrow{\mathrm{CH}_{3} \mathrm{MgBr}}$
4. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CONH}_{2} \xrightarrow{\mathrm{P}_{2} \mathrm{O}_{5} / \Delta}$
5.


## Second Term Test - 2019 <br> Chemisty 13 - II - PART B

## - Answer two question only ( Each question carries 15 mark)

((05) (a) (i) The following Born-Harbor cycle of $\mathrm{Na}_{2} \mathrm{O}$ can be used to calculate the lattice enthalpy of $\mathrm{Na}_{2} \mathrm{O}$.


1. Write the relevant balanced chemical equation to represent the standard lattice enthalpy of $\mathrm{Na}_{2} \mathrm{O}$.
2. Identify the species A and B by giving the physical states of them.
3. Introduce the enthalpy changes mentioned as (1) - (5).
(ii) Using the Born-Habour cycle mentioned and the following data calculate the lattice enthalpy of $\mathrm{Na}_{2} \mathrm{O}$.
The enthalpy of atomization of $\mathrm{Na}(\mathrm{s})=+107 \mathrm{kJmol}^{-1}$
The enthalpy of first electron affinity of Oxygen $=+141 \mathrm{kJmol}^{-1}$
The enthalpy of second electron affinity of oxygen $=+798 \mathrm{kJmol}^{-1}$
The enthalpy of formation of $\mathrm{Na}_{2} \mathrm{O}(\mathrm{s})=-414 \mathrm{kJmol}^{-1}$
The enthalpy of first ionization energy of $\mathrm{Na}=+494 \mathrm{kJmol}^{-1}$
The enthalpy of bond dissociation of $\mathrm{O}_{2}=+496 \mathrm{kJmol}^{-1}$
(iii) How to compare the lattice enthalpy of MgO (s) relative to the lattice enthalpy of $\mathrm{Na}_{2} \mathrm{O}$ (s) by giving reasons.
(b) Under certain conditions, partial pressures in equilibrium of $\mathrm{N}_{2}, \mathrm{O}_{2}$ and $\mathrm{NH}_{3}$ are, $\mathrm{N}_{2(\mathrm{~g})} 44.8 \mathrm{~atm}, \mathrm{H}_{2(\mathrm{~g})} 105.6 \mathrm{~atm}, \mathrm{NH}_{3(\mathrm{~g})} 37.2 \mathrm{~atm}$
$\mathrm{N}_{2(\mathrm{~g})}+3 \mathrm{H}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NH}_{3(\mathrm{~g})}$
(i) Write an expression for $\mathrm{K}_{\mathrm{p}}$ in the production of $\mathrm{NH}_{3}$ in the Haber process.
(ii) Using above data calculate the value of $\mathrm{K}_{\mathrm{p}}$ with the units.
(c) At a certain temperature the equilibrium constant $\mathrm{K}_{\mathrm{c}}$ for the following reaction is 16
$\mathrm{SO}_{2(\mathrm{~g})}+\mathrm{NO}_{2(\mathrm{~g})} \rightleftharpoons \mathrm{SO}_{3(\mathrm{~g})}+\mathrm{NO}_{(\mathrm{g})}$
One mole of each of all the above gases are placed inside a closed vessel of $1 \mathrm{dm}^{3}$ Calculate the equilibrium concentrations in mol dm ${ }^{-3}$ of $\mathrm{NO}_{(\mathrm{g})}$ and $\mathrm{NO}_{2(\mathrm{~g})}$ at the equilibrium.
(06) (a) The marble and limestone monuments situated in the places such as Taj Mahal in India are undergoing corrosion due to the acid rain. The carbonate of these monuments is converted to the relatively more soluble sulphate by the acid rain. The relevant reaction is given below.

$$
\mathrm{CaCO}_{3(\mathrm{~s})}+\mathrm{H}_{2} \mathrm{SO}_{4(\mathrm{aq})} \rightarrow \mathrm{CaSO}_{4(\mathrm{~s})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}+\mathrm{CO}_{2(\mathrm{~g})}
$$

(i) Write an expression for the solubility product Ksp of $\mathrm{CaSO}_{4}$. Give the units of $\mathrm{K}_{\text {sp }}$.
(ii) At the temperature of occurring the above reaction the value of $\mathrm{K}_{\mathrm{sp}}$ is $3 \times 10^{-5}$. Calculate the concentration of $\mathrm{SO}_{4}^{2-}$ ions in the saturated $\mathrm{CaSO}_{4}$ solution, using the above expression.
(iii) At the above temperature when $100 \mathrm{dm}^{3}$ of the acid rain is fallen on a small monument, calculate the mass loss that occurred. Assume that the monument is made up of pure $\mathrm{CaCO}_{3}$ and the acid rain is saturated with $\mathrm{CaSO}_{4}$ due to the corrosion.
(Ca - 40, S - 32, O-16)
(b) These monuments are treated with a mixture of urea and $\mathrm{Ba}(\mathrm{OH})_{2}$ to extend the life time. When the mixture is seeping through the small holes of the carbonate stone, urea dissociate in to $\mathrm{NH}_{3}$ and $\mathrm{CO}_{2}$. This $\mathrm{CO}_{2}$ reacts with, $\mathrm{Ba}(\mathrm{OH})_{2}$ to form $\mathrm{BaCO}_{3}$.
The reaction occurred are given below.
$\left(\mathrm{NH}_{2}\right)_{2} \mathrm{CO}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g})$
$\mathrm{Ba}(\mathrm{OH})_{2}(\mathrm{aq})+\mathrm{CO}_{2}(\mathrm{~g}) \rightarrow \mathrm{BaCO}_{3}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$
Then this $\mathrm{BaCO}_{3(\mathrm{~s})}$ is converted to its sulphate by the acid rain.
$\mathrm{BaCO}_{3(\mathrm{~s})}+\mathrm{H}_{2} \mathrm{SO}_{4(\mathrm{aq})} \rightarrow \mathrm{BaSO}_{4(\mathrm{~s})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}+\mathrm{CO}_{2(\mathrm{~g})}$
The solubility of $\mathrm{CaSO}_{4}$ is too small relative $\mathrm{BaSO}_{4}$. At that temperature in a saturated solution of pure $\mathrm{BaSO}_{4}$, $\left[\mathrm{Ba}^{2+}\right]=9 \times 10^{-6} \mathrm{~mol} \mathrm{dm}^{-3}$
(i) The solubility of $\mathrm{BaSO}_{4}$ is too smaller than $\mathrm{CaSO}_{4}$
(ii) Write the expression for the $\mathrm{K}_{\text {sp }}$ of $\mathrm{BaSO}_{4}$. Calculate the value of it. Using above data. Mention the units.
(c) (i) Methonic acid ( HCOOH ) is a weak acid with $\mathrm{K}_{\mathrm{a}}=1.77 \times 10^{-4} \mathrm{~mol} \mathrm{dm}^{-3}$

1. Write an expression for $\mathrm{K}_{\mathrm{a}}$ of HCOOH .
2. Calculate $\left[\mathrm{H}^{+}(\mathrm{aq})\right]$ of $0.05 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{HCOOH}$ solution using the above expression.
3. Calculate the percentage of ionized molecule of HCOOH in the solution.
4. Calculate the pH of the solution.
(ii) Both HCOOH and HCl acid reacts with Mg metal powder.
5. Write the balanced chemical equation for the reaction between HCOOH and Mg reacted.
6. $20 \mathrm{~cm}^{3}$ volumes of $0.05 \mathrm{~mol} \mathrm{dm}^{-3}$ of these two acid solutions are taken and reacted with excess Mg powder. Then the equal volumes of $\mathrm{H}_{2}(\mathrm{~g})$ are liberated. But relative to HCl acid, the reaction rate of HCOOH is very low. Calculate the volume of liberated $\mathrm{H}_{2}(\mathrm{~g})$.
(At STP, $\mathrm{V}_{\mathrm{m}}=22.4 \mathrm{dm}^{3} \mathrm{~mol}^{-1}$ )
7. The reaction rate of HCOOH is too low relative to HCl acid. Explain the reasons.
8. Explain the reason for the liberation of the same gas volume as HCl by HCOOH , at the end of the reaction.
(07) (a) A low quality sample of an alloy which is used in welding's is analyzed for Pb . A mass of 0.759 g of that sample is dissolved is an acid to from a solution of $\mathrm{Pb}^{2+}$. An excess $\mathrm{K}_{2} \mathrm{CrO}_{4}$ solution is added to the above solution to precipitate all $\mathrm{Pb}^{2+}$ present in it. After that the precipitate is filtered and washed. Then the precipitate is again dissolved in an acid and treated it with excess KI. The liberated $\mathrm{I}_{2}$ during the reaction of $\mathrm{CrO}_{4}^{2-}$ with KI is titrated with $0.051 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ solution. The consumed volume of $\mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ solution for the titration is $11.22 \mathrm{~cm}^{3}$. $\quad\left(\mathrm{Pb}=207 \mathrm{~g} \mathrm{~mol}^{-1}\right)$
(i) Write the balanced chemical equations for all the reactions above.
(ii) Calculate the mass percentage of Pb in this alloy.
(b) (i) Explain that "what is referred as standard hydrogen electrode"
(ii) The standard hydrogen electrode is connected to the standard Zn electrode through a salt bridge, to from an electro chemical cell. The following questions are based on the formed cell.
9. Mention the anode and the cathode.
10. When the two terminals are not connected write the balanced chemical equations for the reactions.
11. Write the balanced chemical equation for the cell reaction when a current flows through the cell.
12. Write the standard cell notation for the above cell.
13. If $\mathrm{E}^{\theta}$ value of the standard Zn electrode is $\mathrm{Zn}-0.76 \mathrm{v}$. Calculate the electromotive force of the above cell.
14. Mention a method that can be used to increase the e.m.f. of the above electric cell.
(c) At $27^{\circ} \mathrm{C}$ an ideal solution, formed by mixing $\mathrm{A}(\mathrm{l})$ and $\mathrm{B}(\mathrm{l})$ is in equilibrium with its vapour. At that temperature the standard vapour pressure of A is $4 \times 10^{5} \mathrm{~Pa}$ and the satarated vapour pressure of $B$ is $2.5 \times 10^{5} \mathrm{~Pa}$. The total pressure of the gas phase in $3 \times 10^{5} \mathrm{~Pa}$.
(i) Explain the ideal behaviour of the above solution considering the intermolecular interactions.
(ii) Calculate the composition of A and B in the liquid phase in equilibrium. Mention the used assumptions.
(iii) Calculate the composition of the vapour phase in equilibrium.
(iv) draw the phase diagram of the variation of the temperature and the composition at the constant pressure, relevant to the above ideal solution.
(v) Mention the method that is used to separate a mixture A and B.

## Part - C essay

## - Answer two question only ( Each question carries 15 marks)

(08) (a) Mention the structures of the compounds $A$ to $G$ and the reagents $R_{1}$ to $R_{8}$ given $n$ the following conversion.

(b) Show how would you synthesis the organic compound $\mathrm{CH}_{3} \mathrm{CH}_{2}-\mathrm{C}-\mathrm{CH}_{3}$ using $\mathrm{C}_{2} \mathrm{H}_{4}$ using as
the starting matrial, in not more than six steps.
(c) How to distinguish the following compounds.
(i) $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ and $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{CH}_{2} \mathrm{OH}$
(ii) $\mathrm{CH}_{3} \mathrm{NH}_{2}$ and $\mathrm{CH}_{3} \mathrm{CONH}_{2}$
(iii) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CHO}$ and $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COCH}_{3}$
(09) (a) The compomds A and B are solids and soluble in water. The observations obtained for the experiments carried out to identify the two cations present in those compounds, are given below.

| Experiment | Observations |
| :---: | :---: |
| (1) Added a small amount of NaOH to an aqueous solution of $A$. | A brownish black precipitate is obtained and it is soluble in $\mathrm{NH}_{4} \mathrm{OH}$ |
| (2) Added diluted HCl to a portion of an aqueous solution A | A white precipitate which is insoluble in hot water and soluble in dil. $\mathrm{NH}_{4} \mathrm{OH}$ |
| (3) added a small amount of NaOH to an aq. solution of B | A white precipitate which is soluble in excess NaOH |
| (4) Added a small amount of $\mathrm{NH}_{4} \mathrm{OH}$ to an aqueous solution of B | A white precipitate which is insoluble in excess $\mathrm{NH}_{4} \mathrm{OH}$ |
| (5) Added dil HCl to an aq. solution of B and passed $\mathrm{H}_{2} \mathrm{~S}(\mathrm{~g})$. | No observation. |
| (6) Boiled the solution obtained in above (5) and added | A gelatinous white precipitate is obtained. |

(i) Identify the cations present in A and B.
(ii) Write the chemical formulae of the precipitates obtained in 1 and 3 above.
(b) Write the answers considering the elements $\mathrm{P}, \mathrm{S}$ and Cl .
(i) Mention the naturally existing molecular type in each of the above elements.
(ii) Arrange the above elements in the order of increasing their melting points.
(iii) Mention the reason for the answer in above (ii).
(iv) Write the chemical formulae of the hydrides formed by the above their elements. Mention their chemical properties in front of them as, weak acidic / weak basic / strong acidic / strong bases.
(v) Write the balanced chemical equations for the reactions of Sulphur with the followings.

1. $\mathrm{NaOH}(\mathrm{aq})$
2. hot concentrated $\mathrm{H}_{2} \mathrm{SO}_{4}$ (g)
(vi) Write the balanced chemical equations for the reactions of chlorine with the followings.
3. hot concentrated KOH
4. excess $\mathrm{NH}_{3}$ (g)
(c) (i) Mention the order of decreasing water solubilities of the group II elements. Explain the reasons for that.
(ii) Mention the order of increasing melting points of $\mathrm{LiF}, \mathrm{LiCl}, \mathrm{LiBr}, \mathrm{LiI}$ and explain the reasons.
(10) (a) Community of the area complained to the central environmental authority that the waste water drained out from an industry of manufacturing $\mathrm{H}_{2} \mathrm{SO}_{4}$ is polluted with the acid. To test that waste water $50 \mathrm{~cm}^{3}$ of it measured and diluted up to $500 \mathrm{~cm}^{3}$ by adding distilled water. To react with all $\mathrm{H}^{+}$present in $25 \mathrm{~cm}^{3}$ of the above prepared solution, 0.2 g of excess $\mathrm{KIO}_{3}$ and KI was added. To react completely with the formed iodine $20 \mathrm{~cm}^{3}$ of $0.1 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ solution was required.
(i) Write the balanced chemical equations for the reactions taken place.
(ii) Calculate the concentration of $\mathrm{H}^{+}$present in the waste water.
(b) Mention the chemical formulae of the complex ions formed by the given cations below with the relevant ligand and mention the colours of those complex ions.

| cation | ligand | complex ion | colour |
| :---: | :---: | :---: | :---: |
| $(1) \mathrm{Cr}^{3+}$ | $\mathrm{Cl}^{-}$ |  |  |
| $(2) \mathrm{Fe}^{3+}$ | $\mathrm{H}_{2} \mathrm{O}$ |  |  |
| $(3) \mathrm{Co}^{3+}$ | $\mathrm{NH}_{3}$ |  |  |
| (4) $\mathrm{Co}^{2+}$ | $\mathrm{Cl}^{-}$ |  |  |
| $(5) \mathrm{Cu}^{2+}$ | $\mathrm{NH}_{3}$ |  |  |

(c) A is an inorganic mix salt, of two compounds (a dualsalt) having a single anion and two cations. The tests carried out to identify the anion and two cation and their observations are given below.

(i) Identify the two cations and the anion contain in A Write the chemical formula.
(ii) Write the chemical formulae of the species C, J and K
(iii) Write the formulae of coordinated complexes which cause to the colour of $B$ and $H$
(iv) Write the chemical formulae of the precipitates of F and G.
(v) Write the balanced chemical equations for, the reactions between B and NaOH the reactions between H and Zn dust.



| 57 | 58 | 59 | 60 | 61 | 62 | 63 | 64 | 65 | 66 | 67 | 68 | $\mathbf{G 9}$ | 70 | 71 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| La | Ce | Pr | Nd | Pm | Sm | Eu | Gd | Tb | Dy | Ho | Er | Tm | Yb | Lu |
| 89 | 90 | 91 | 92 | 93 | 94 | 95 | 96 | 97 | 98 | 99 | 100 | 101 | 102 | 103 |
| Ac | Th | Pa | U | Np | Pu | Am | Cm | Bk | Cr | Es | Fm | Md | No | Lr |


| $(1)$ | 3 | $(11)$ | 4 | $(21)$ | 4 | $(31)$ | 4 | $(41)$ | 3 |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| $(2)$ | 1 | $(12)$ | 5 | $(22)$ | 2 | $(32)$ | 5 | $(42)$ | 1 |
| $(3)$ | 4 | $(13)$ | 1 | $(23)$ | 3 | $(33)$ | 4 | $(43)$ | 1 |
| $(4)$ | 2 | $(14)$ | 3 | $(24)$ | 3 | $(34)$ | 3 | $(44)$ | 4 |
| $(5)$ | 2 | $(15)$ | 1 | $(25)$ | 4 | $(35)$ | 5 | $(45)$ | 5 |
| $(6)$ | 3 | $(16)$ | 4 | $(26)$ | 1 | $(36)$ | 4 | $(46)$ | 1 |
| $(7)$ | 2 | $(17)$ | 1 | $(27)$ | 3 | $(37)$ | 5 | $(47)$ | 4 |
| $(8)$ | 5 | $(18)$ | 5 | $(28)$ | 5 | $(38)$ | 4 | $(48)$ | 3 |
| $(9)$ | 3 | $(19)$ | 5 | $(29)$ | 2 | $(39)$ | 5 | $(49)$ | 3 |
| $(10)$ | 2 | $(20)$ | 2 | $(30)$ | 3 | $(40)$ | 4 | $(50)$ | 5 |

## Part II

Structured Essay - Part II

$$
\begin{aligned}
& \text { (gl) (a) I } \quad \mathrm{NO}_{2} \quad \mathrm{NO}_{2} \mathrm{NH} \mathrm{NH}_{2} \mathrm{~N}_{2} \mathrm{O} \quad \mathrm{WH}_{4}^{+} \\
& \text {Oxidation number }+4 \quad \text { in } \quad-1 \cdots+2 \quad-3 \\
& \text { Hybridization. } s p^{2} S p^{2} S p^{3} \quad S p p^{s} \quad 10 \times 1 \text { marks } \neq(10) \\
& \text { II: } \quad: N \equiv N^{+}-0_{0}^{-}: \quad(10 \text { marks }) \\
& \text { II } \quad: \dot{N}=\stackrel{+}{N}=\ddot{O}: \longleftrightarrow \quad \ddot{N}-\dot{N}^{+} \equiv \ddot{\theta^{+}}(\operatorname{mink} 5 \times 2=10) \\
& \text { * not stable. } \\
& \text { * not stable. } \\
& \text { * charge distribution } \\
& \text { *. }(\rightarrow \text { charge is on } \\
& \text { *. ( } t \text { ) charge is on the } \\
& \text { oxygen atom } \\
& \text { (O2 meres). } \\
& \text { (O3morles) } \\
& \text { (b). i. potassium hydrogen difluorde. ( } 10 \mathrm{men} / \mathrm{lg} \text { ) } \\
& \text { ii. cation } \begin{array}{ll}
\text { anion } & \mathrm{HF}_{2}-
\end{array} \\
& \text { ( } 02 \times 2 \text { marks) } \\
& \text { II. }
\end{aligned}
$$

iv: 1: $\mathrm{H}_{2} \mathrm{~S}$ *. dipole-dipole interactions.

* dispersion forces. ( $0_{2} \times 2$ mark $=44$ )
$\mathrm{CS}_{2}$ dispersion forces (o 2mark)

2. $C_{2} \quad(05$ marks
$3, C S_{2}, \quad(05$ marks $)$

$$
\begin{aligned}
& \text { 4. } 2 \mathrm{H}_{2} \mathrm{~S}+3 \mathrm{O}_{2} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{SO}_{2} \\
& \mathrm{CS}_{2}+3 \mathrm{O}_{2} \rightarrow \mathrm{CO}_{2}+2 \mathrm{SO}_{2} \\
&(\mathrm{OS} \times 2 \text { 2 } 10 \mathrm{merkg})
\end{aligned}
$$

(c). $i$. True. iii True. $v$. True.
ii. False ir. False.

$$
r_{i} \text { False. }
$$

vii.

$$
\text { (0 } 2 \times 8=16 \text { monks })
$$

(a2) $(a)$ i. $M=M g$ (vine

$$
\text { ii. } Q_{1}=M g O \quad Q_{2}=M_{3} N_{2} .(10 \text { marks) }
$$

iii. i. $\left.\mathrm{Mg}+2 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{MgCOH}\right)_{2}+\mathrm{H}_{2}$
2. $\mathrm{Mg}_{8} \mathrm{~N}_{2}+6 \mathrm{H}_{2} \mathrm{O} \longrightarrow 3 \mathrm{Mg}(\mathrm{OH})_{2}+2 \mathrm{NH}_{3}$ ( 10 marls)

$$
\text { Pr. Mass of } Q_{2} \text { in irreg }=\frac{30.3}{100} \times 1=0.303 \mathrm{~g}
$$

$\left.\begin{array}{c}\text { number of moles } \\ O F \\ Q_{2}\end{array}\right\}=\frac{0.303}{100}=0.003 \mathrm{~mol}$

$$
\mathrm{Mg}_{3} \mathrm{~N}_{2}+\mathrm{CH}_{2} \mathrm{O} \rightarrow 3 \mathrm{Mg}(\mathrm{OH})_{2}+2 \mathrm{NH}_{3}
$$

number of moles ${ }^{\text {of } \mathrm{NH}_{3}}$ liberated $=0.006 \mathrm{~mol}$.

$$
\mathrm{NH}_{3}+\mathrm{HCl} \longrightarrow \mathrm{NH}_{4} \mathrm{Cl}
$$

$\therefore$ number OF moles of ACl treated. $\}=0.006 \mathrm{~mol}^{6}$.

$$
\begin{aligned}
\text { Volume of } \mathrm{HCl} & =\frac{\left.1000 \mathrm{~cm}^{3} \times 0.00606 \mathrm{mc}\right)}{0.5 \mathrm{~mol}} \\
& =\frac{12.12 \mathrm{~cm}^{3}}{} \quad(01 \times 8=8 \mathrm{marks})
\end{aligned}
$$

(b). i. $Y=\mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}$ ( 10 marls)
ii. $2 \mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{8}+I_{2} \longrightarrow 2 \mathrm{NaI}+\mathrm{Na}_{2} \mathrm{SO}_{4}$
$\mathrm{Na}_{2} \otimes_{2} \mathrm{O}_{3}+2 \mathrm{HCl} \longrightarrow 2 \mathrm{NaCl}+\mathrm{SO}_{2}+\mathrm{H}_{2} \mathrm{O}+\mathrm{S}$

$$
\mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}+2 \mathrm{AgNO}_{3} \rightarrow \mathrm{Ag}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}+2 \mathrm{NaNO}_{3}
$$

$$
\mathrm{Ag}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{Ag}_{2} \mathrm{~S}+\mathrm{H}_{2} \mathrm{SO}_{4}\left(\begin{array}{l}
5 \times 4=20 \mathrm{mon} / \mathrm{lg})
\end{array}\right.
$$

M


$\therefore \quad \therefore \quad \begin{array}{ll}\quad \therefore \quad \mathrm{Na}_{2} \mathrm{O}_{3}-5 \mathrm{H}_{2} \mathrm{O} \\ (2 \times 6=12 \text { marts })^{3}\end{array}$
(C) i. $\mathrm{NaNH}_{2}+\mathrm{NH}_{4} \mathrm{Cl} \longrightarrow 2 \mathrm{NH}_{3}+\mathrm{NaCl}$

$$
\text { ii. } \mathrm{PbS}+4 \mathrm{H}_{2} \mathrm{O}_{2} \rightarrow \mathrm{PbSO}_{4}+4 \mathrm{H}_{2} \mathrm{O}
$$

$$
\text { iii } \quad 8 \mathrm{Al}+5 \mathrm{OH}^{-}+3 \mathrm{NO}_{3}^{-} \rightarrow 8 \mathrm{AlO}_{2}^{-}+3 \mathrm{NH}_{3}
$$

$$
\text { iv. } 2 \mathrm{FeCl}_{3}+\mathrm{H}_{2} \mathrm{~S} \rightarrow 2 \mathrm{FeCl}_{2}+2 \mathrm{HCl}+\mathrm{S}
$$

$$
\text { (03). (a) i. } P k_{b}=-\log k_{b} \quad \text { ( } 4 \times 5=20 \mathrm{marls} \text { ) }
$$

$$
\text { ii. } \mathrm{NH}_{3} \text { aq }+\mathrm{Hi}_{2}(l) \rightleftarrows \mathrm{NH}_{4} \mathrm{NCan}_{4}^{+\mathrm{OH}_{\text {(aq }}^{-}}
$$

$$
\text { Mi: } \quad K_{b}=\left[\mathrm{NH}_{4}^{+(a q)}\right]\left[\mathrm{OH}_{4}^{-}\right]
$$

$$
\frac{\left.\mathrm{NH}_{4}^{\mathrm{H}} \text { (ag) }\right]\left[\mathrm{OH}_{\text {cog, }}^{-}\right]}{\left[\mathrm{NH}_{\mathrm{H}} \text { (ag) }\right]} \mathrm{V}
$$

$$
v-K_{a} \times K_{b}=\left[\mathrm{NH}_{4}^{\top}{ }_{\text {cap }}\right]
$$

$$
\begin{gathered}
=\left[\mathrm{OH}_{3}^{-}\right]\left[\mathrm{H}_{3} \mathrm{O}^{+}\right] \\
{\left[\mathrm{OH}^{-}\right]=\mathrm{KW}^{2} .}
\end{gathered}
$$

$$
\therefore k_{a} \cdot k_{b}=k_{w}
$$

$$
\text { vi. } \quad k_{a}=\frac{k_{w}}{k_{b}}
$$

$$
\begin{aligned}
-\log _{10} k_{a} & =-\log _{10} \frac{k_{w}}{k_{b}} \quad(11 \times 5=55 \text { marta }) \\
p k_{a} & =-\log _{10} k_{w}-\log _{10} k_{b}=14-4.8=9.2 \mathrm{~V}
\end{aligned}
$$

3. 

(b)

$$
\text { ri. } \mathrm{NaHCO} 3+\mathrm{HCl} \longrightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{CO}_{3}
$$

ir. In the mitial mixture, if there are $x$ mol of NaOH and $y \mathrm{~mol}$ of $\mathrm{Na}_{2} \mathrm{CO}_{3}$

$$
\begin{align*}
x+y & =\frac{1 \times 10}{1000}-(1)  \tag{1}\\
x+2 y & =\frac{1 \times 15}{1000}  \tag{2}\\
x-1) & =0.015-0.01 \\
x & =0.01-0.005
\end{align*}
$$

(ii) $\mathrm{C}_{6} \mathrm{H}_{5}-\mathrm{O}-\mathrm{C}^{\mathrm{N}} \mathrm{CH}_{3}$
III. i. electrophilic substitution. vercleophilic substitution.
iv:

(b)


$$
\mathrm{C}_{6} \mathrm{H}_{5}^{\circ}+\mathrm{HCl}_{2}
$$

$$
\begin{gathered}
(2 \times 8=16 \mathrm{monk})
\end{gathered}
$$


dill
$\mathrm{H}^{+}$


$$
\begin{aligned}
& \mathrm{NaOH}+\mathrm{HCl} \longrightarrow \mathrm{NaCl}_{2}+\mathrm{H}_{2} \\
& \mathrm{Na}_{2} \mathrm{CO}_{3}+\mathrm{HCl} \longrightarrow \mathrm{NaHCO}_{3}+\mathrm{NaCl}- \\
& \text { it. } 2 \mathrm{NaOH}+\mathrm{CO}_{2} \longrightarrow \mathrm{NaCO}_{2}+4 \mathrm{H}_{2}- \\
& \mathrm{Na}_{2} \mathrm{CO}_{3}+\mathrm{CO}_{2}+\mathrm{H}_{2} \mathrm{O} \rightarrow \underset{2}{\mathrm{Na}_{3}}+\mathrm{HOO}_{2} \sim
\end{aligned}
$$

(c) i. 0 , $\mathrm{C}-\mathrm{O}-a\rangle-\mathrm{NO}_{2}$
ii $\mathrm{O}-\underset{\mathrm{Br}}{\mathrm{CH}}-\mathrm{CH}_{2} \mathrm{CH}_{3}$
iii. $\quad \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{COOMgBr}$
iv. $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CN}$.
$r$.

(05 $\times 5=25 \mathrm{manls}$ ?
$05(a)$.
(i) $1.2 \mathrm{Na}_{(g)}^{+}+\mathrm{O}_{(g)}^{2-} \longrightarrow \mathrm{Na}_{2} \mathrm{O}_{\text {cs }}$ (10)
i) 2.: $A=N a(g)$
$\cos$
$(05)$
3. (1) - The first ionigation enthalpg of Ma 105)
(2) - The addition of first and second Stendard enthalpres of electron affimitres (05) of oxygen.
(3) - The stendard. lattice enthalpy of $\mathrm{Na}_{2} \mathrm{O}_{\mathrm{cs}}$, (as)
(4) - The standard enthalpy of formation op $\mathrm{N}_{2} \mathrm{O}_{\mathrm{e}}$ (ar)

C if the symbals of the enthalpies mentroned, give ( 2 manks on(y)
(II)

$$
\begin{aligned}
& +\Delta H_{a+m}^{\theta} O_{\text {Cg }}+\left(\Delta H_{E A_{1}}^{\theta}+\Delta H_{E A_{2}}^{Q}\right)_{O_{C D}}
\end{aligned}
$$

$$
\begin{align*}
& -414 \mathrm{kjmol}^{-1}=\left(2 \times 107 \mathrm{kjmal}^{-1}\right)+\left(2 \times 494 \mathrm{kjmal}^{1}\right)+  \tag{10}\\
& \left(\frac{1}{2} \times 496 \mathrm{kjmol}\right)+\left(-141+798 \mathrm{kjmol}^{\mathrm{K}}\right) \\
& +\Delta H_{L}^{8} \text { (NaOCS) } \\
& \Delta H_{L\left(\mathrm{Na}_{2}\right.}^{\theta}=-2521 \mathrm{kjmol} \tag{10}
\end{align*}
$$

(III) The lattice enthalpy of AgO is a s mower than the lattice enthalpy of Nra. (more negative)

Reason:- $\mathrm{Mg}^{2 t}$ is corbie. Charged relative to $\mathrm{Nat}^{2}$ and the charge density of $\mathrm{Mg}^{2}$ is high and its radius is small
(b) (i) $K_{P}=\frac{P^{2} N_{3}(g)}{P_{N_{2}(g)} \times P_{H_{2}(g)}^{3}}$

(8) (ii)

$$
\begin{aligned}
K_{p} & =\frac{(37.2)^{2} \mathrm{~atm}^{2}}{44.8 \times(105.6)^{3} \mathrm{~atm}} \\
& =2.62 \times 10^{-5} \mathrm{~atm}^{2}
\end{aligned}
$$

 moles


$$
\begin{equation*}
k_{s p}=\left[\mathrm{ca}_{\mathrm{cq},}^{2+}\right]\left[\mathrm{sch}_{\mathrm{coq}}^{2}\right] \quad \text { mol }{ }^{2} d m^{-8} \text {. (os) } \tag{0}
\end{equation*}
$$

(ii)

$$
\begin{aligned}
{[\text { cast } 4] } & =\sqrt{k_{S P}}=\sqrt{3 \times 10^{-5} \mathrm{~mol}^{2} \mathrm{dm}^{-6}} \\
& =5.5 \times 10^{-3} \mathrm{moldm} \mathrm{~m}^{-3}
\end{aligned}
$$

(III) The amount of $\mathrm{CaSO}_{4}$ in $100 \mathrm{dm}^{3}=5.5 \times 10^{-3} \times 100$

$$
=0.55 \mathrm{~mol} \quad(05)
$$

dissolved amount of $\mathrm{CaCO}_{3}=$ amount of $\mathrm{CaSO}_{4}$ in the solution
molar mass of $\mathrm{CaCl}_{3}=100 \mathrm{gmol}$ mass $\operatorname{loss}\left(\mathrm{CaCO}_{3}\right)$.

$$
=\frac{0.55 \times 100=55 \mathrm{~g} \quad(5)}{a \rightarrow 30}
$$

(b) (i). When the ionic solids are dissolved in water, to for the difference between lattice enthalpy and solution enthalpy, felerent energy should be absorbed.
*. When going down the group, the solution enthalpy

* takes a very low $(-)$ value relative to $\mathrm{CaSO}_{4}$ ? since the earl cationic radius is large Therefore the solution enthalpy of $\mathrm{BaSO}_{4}$ toes takes a more $(t)$ value than the solution. enthalpy of $\mathrm{CaSO}_{4}$.
$\therefore$ The Solubility of $\mathrm{BaSO}_{4}$ is too low relative to $\mathrm{CaSO}_{4}$.
(ii)

$$
\begin{aligned}
K_{s p} & =\left[\mathrm{Ba}_{\mathrm{cag}}^{2+}\right]\left[\mathrm{SO}_{4}^{2} \mathrm{caq}^{2}\right] \\
& =\left(9 \times 10^{-6} \mathrm{moldm}^{-3}\right)^{2} \\
& =8.1 \times 10^{-11} \mathrm{mal}^{2} \mathrm{dma}^{-6} \\
\therefore K_{a} & =\frac{\left[\mathrm{HCOO}^{-} \mathrm{caql}\right]\left[\mathrm{H}_{\mathrm{caq} 2}^{+}\right]}{\left[\mathrm{HCOOH}_{\mathrm{av}}\right]} \text { moldm-s }
\end{aligned}
$$

$$
\begin{align*}
{\left[4_{\text {aq }}^{+}\right] } & =\left[H_{\text {(oo }}^{-a g)}\right]  \tag{05}\\
K_{a} & =\frac{\left[H_{(a q)}\right]^{2}}{\left[H\left(a 0 H_{\text {(ag) }}\right]\right.}
\end{align*}
$$

(05)

$$
\left[H^{+}(a q)\right]=\sqrt{K_{a} \times\left[H_{(O O H}^{(a g)}\right)}
$$

Since the dissociation of the acid is two small, it is considered that the equilibrium concentration or the acid is equal to the initial concentration.

$$
\begin{aligned}
& =\sqrt{1.77 \times 10^{-4} \times 0.05} \\
& =2.97 \times 10^{3} \quad\left(\simeq 3 \times 10^{-3}\right) \mathrm{mol} \mathrm{dmin}^{-3}
\end{aligned}
$$

(3)

$$
\begin{aligned}
\frac{\left[H_{\text {cog }}\right]}{\left[H\left(00 H_{\text {cog } 3}\right]\right.} \times 100 & =\frac{297 \times 10^{-3} \times 100}{0.05} \\
& =5.94 \% \quad(\simeq 6 \%)
\end{aligned}
$$

Co (4)

$$
\begin{align*}
p^{H} & =-\log _{10}\left[H_{(09)}^{+}\right] \\
& =-\log _{10}\left(2.97 \times 10^{-3}\right) \\
& =2.52 \tag{05}
\end{align*}
$$

(II) i)

$$
\begin{aligned}
& \mathrm{H} \text { (00H (ag) }+M g(s) \longrightarrow\left(\mathrm{H}_{(000)_{2} \mathrm{Mg}_{(s)}+\mathrm{H}_{2(g)}} \longrightarrow \mathrm{Mg}_{\text {(aq) }}+\mathrm{H}_{2 \text { (g) }}\right.
\end{aligned}
$$

2) $H^{+}$aqs amount $=0.05 \mathrm{mold} \mathrm{me}^{-3} \times \frac{20}{10^{3}} \mathrm{dm}^{3}$

$$
=1 \times 10^{-3} \mathrm{~mol}
$$

(05)
amount of

$$
\mathrm{H}_{2}(g)
$$

$$
\begin{aligned}
& =\frac{1}{2} \times 1 \times 10^{-3} \mathrm{~mol} \\
& =0.5 \times 10^{-3} \mathrm{~mol}
\end{aligned}
$$

volume of $\mathrm{H}_{2}(\mathrm{~g})$

$$
\begin{align*}
& =0.5 \times 10^{-3} \times 22400 \\
& =11.2 \mathrm{~cm}^{3} \tag{0,5}
\end{align*}
$$

(3) Rate $\propto\left[\mathrm{H}^{+}(a q)\right]$
$\mathrm{HCOOH}^{-1}(a q)$ acid partially decomposes slowly. (03)
(4) The same amount of $\mathrm{H}^{+}\left(\mathrm{aq}_{q}\right)$ is consumed as HCl , because $\mathrm{H}_{(a q)}^{+}$is consumed to form $\mathrm{H}_{2}(g)$. Reason is, the equal volumes of acids are used.
$\frac{\left(\begin{array}{c}(05) \\ e \rightarrow(8) \\ \end{array} \frac{150}{=}\right.}{}$

$$
\begin{align*}
& \text { (07) (a). (i) } \left.2 \mathrm{~S}_{2} \mathrm{O}_{3}{ }^{2} \longrightarrow \mathrm{~S}_{4} \mathrm{O}_{6}{ }^{2-}+2 e \cdot\right\} \text { This step } \\
& \left.2 \mathrm{e}+\mathrm{I}_{2} \longrightarrow 2 I \mathrm{I} \quad\right\} \text { is not essential } \\
& 2 \mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}+\mathrm{I}_{2} \longrightarrow \mathrm{Na}_{2} \mathrm{~S}_{4} \mathrm{O}_{6}+2 \mathrm{NaI}  \tag{05}\\
& 6 \mathrm{I}^{-}+16 \mathrm{H}^{+}+2 \mathrm{CrO}_{4}^{2-} \rightarrow 3 \mathrm{I}_{2}+2 \mathrm{Cr}^{3+}+8 \mathrm{H}_{2} \mathrm{O}  \tag{10}\\
& \mathrm{~Pb}^{2+}+\mathrm{CrO}_{4}^{2} \longrightarrow \mathrm{PbCrO}_{4}(s) . \\
& \text { (a) }
\end{align*}
$$

$\therefore$ (I). The amount of $\mathrm{Ng}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}=0.051 \mathrm{NoD} 1 \mathrm{dm}^{-3} \times \frac{11.22}{1000} \mathrm{dm}^{3}$ (05)
The amount of $I_{2} \quad=0.051 \times \frac{11.22}{1000} \times \frac{1}{2} \mathrm{~mol}$ (a5) $\begin{aligned} \mathrm{Cro}_{4}{ }^{2} \text { cmount. } & =0.051 \times \frac{11.22}{1000} \times \frac{1}{2} \times \frac{2}{3} \mathrm{~mol}(05) \\ \therefore \text { amount of } \mathrm{Pb} & =0.051 \times 11.22 \times 1 \times 0)\end{aligned}$ $\therefore$ amount of $\mathrm{Pb}=0.051 \frac{\times 11.22}{1000} \times \frac{1}{2} \times \frac{2}{3} \mathrm{~mol}$
The mass of $\mathrm{Pb}=0.05(\times 11.22$ (05)
$\begin{aligned} \text { The mass percentage } & =0.051 \times \frac{11.22}{1000} \times \frac{1}{2} \times \frac{2}{3} \times \frac{207 \times 100}{0.759}\end{aligned}$
(07) (b) (1). At 298 K

## (06)

(2) 1. anode $2 n$ (2) cathock - $H$ elestrode (2) llat anode $\mathrm{Zn}^{2+}$ (q) t2e $\rightleftharpoons \quad$ Zncs, (s)

$$
2 n^{\top}(9)+20<H_{2}(9) \text { (5) }
$$

iii

$$
Z_{n}(s)+2 \mathrm{H}_{(\text {eq })}^{+} \longrightarrow \text { 生 }^{+g}+Z_{n}^{2+}(5)
$$


$v . E^{\theta}=E_{C}^{\theta}-E_{A}^{\theta}$

$$
=0-(0.76 \mathrm{~V})
$$

$$
=+0.76 \mathrm{~V}
$$

$$
\begin{aligned}
& \text { +. Addition of con } \mathrm{FCl} \text { to the solutica, } \cos 3 \\
& \text { contams } \mathrm{H}^{+}
\end{aligned}
$$

(c) 1. In the solution

$$
f_{(A \ldots A)}=f_{(B \ldots B)}=F_{(A \ldots B)^{2} .}
$$

then the ability to vaporize does not depend on each other.
(10)
(ii) By assuming the Ideal nature of the rapor
$P_{\tau}=P_{A}+P_{B}$
$3 \times 10^{5} \mathrm{~Pa}=4 \times 10^{5} X_{A}+2.5 \times 10^{5}\left(1-x_{A}\right) 2$

$$
x_{A}=\frac{1}{3}
$$

Then $X_{B}=1-\frac{1}{3}=2 / 3$
(ii)

$$
\begin{aligned}
P_{A} & =P_{A}^{0} \times{ }_{A} V \\
P_{A} & =4 \times 10^{50} P_{a} \times \frac{1}{3}=4 / 3 \times 10^{5} P a . \\
P_{A} & =P_{T} \times Y_{A} V \\
Y_{A} & =\frac{P_{A}}{P_{T}} V=\frac{4 \times 10^{5}}{3 \times 3 \times 10^{5}}=\frac{4}{9} / / / 2
\end{aligned}
$$

(08) (a) $\mathrm{CH}_{3} \mathrm{COCH}_{3} \xrightarrow[\mathrm{R}_{1}]{\mathrm{NaBH}_{4}}$

$\mathrm{CH}_{3}-\mathrm{CH}=\mathrm{CH}_{2}$
$\mathrm{CH}_{3} \mathrm{COCH}_{3}$
R5



$$
\mathrm{CH}_{3}-\underset{\substack{\mathrm{C} \\
1 \\
\mathrm{CH}_{3}}}{\substack{\left.\mathrm{CH}_{2} \\
\mathrm{CH}_{3}\right)_{2}}} \stackrel{\mathrm{NH}_{3}}{\mathrm{~N}_{8}}\left[\begin{array}{c}
\mathrm{CH}_{3}-\mathrm{C}_{1}-\mathrm{CH}\left(\mathrm{CH}_{3}\right)_{2} \\
\mathrm{CH}_{3}
\end{array}\right]
$$

$$
\begin{aligned}
& A, B, G, D, E, F, G, \rightarrow(7 \times 5=35 \text { marks }) \\
& R_{1} \rightarrow R_{8} \quad(8 \times 3=24 \text { manks) }
\end{aligned}
$$


(c) $\mathrm{C} . \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$


Turbidity is


## Give turbardmy take abound

one hour to
give the turbidity,
shown within 5 minutes.
$\mathrm{CH}_{4} \mathrm{CHO} \underset{\text { decane }}{\mathrm{CH}_{3} \mathrm{COCH}_{3}}$
( $2 \times 9^{2}=18$ mors:)
Ii. $\mathrm{CH}_{3} \mathrm{NH}_{2}, \mathrm{CH}_{3} \mathrm{CONH}_{2}$
" $\mathrm{NaOH} / \Delta$ is tested with rustlers reagent.


$$
\begin{aligned}
& \text { decolours. } \\
& \text { gas } \\
& \text { otters reagent. }
\end{aligned}
$$


$\begin{array}{lll}\text { (G) I. } \mathrm{A} \rightarrow \mathrm{Ag}^{+} & \mathrm{B} \rightarrow \mathrm{Al}^{3+} \quad \text { (1 0marls) } \\ \text { II } 19 \mathrm{Ag}_{2} \mathrm{O} & 3 \rightarrow \mathrm{Al}(\mathrm{OH})_{3} \text { (lomrls) }\end{array}$
(b). I phospechous - $P_{4}$ Sulphur $S_{8}$ (hiorine $\left(C_{2}\right)$ II belting point $\mathrm{Cl}_{2}<p_{4}<s_{8} \quad(\circ 9 \mathrm{mmlk})$ III The number of electrons present within the molecule is increasing as $\mathrm{Cl}_{2} \leftharpoonup p_{4}<S_{8}$, Then the ability of forming London fores among the non polar molecules is increasing as $C_{2}<P_{4}<S_{8}$.
$\therefore$ The melting point is increasing as above.
Pr. $\mathrm{P} \rightarrow \mathrm{PH}_{3}$ weak basic: $\quad(6 \times 2=12$ mums)

$$
s \rightarrow \vec{H}_{2} s \backsim \text { weak acidic } \vdots
$$

$$
\mathrm{el} \rightarrow \mathrm{HCl} \leadsto \sim \text { strong acid } \cdot(06 \times 3=18 \mathrm{mols})
$$

V. $\quad 6 \cdot \mathrm{NaOH}+4 \mathrm{~S} \longrightarrow 2 \mathrm{Na}_{2} \mathrm{~S}+\mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}+3 \mathrm{H}_{2} \mathrm{O}$
$2 \mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{S} \rightarrow 3 \mathrm{SNO}_{2}+2 \mathrm{H}_{2} \mathrm{O}$ (20 marls)
vi. $6 \mathrm{KOH}+3 \mathrm{Cl}_{2} \rightarrow 5 \mathrm{KCl}+\mathrm{KClO}_{3}+3 \mathrm{H}_{2} \mathrm{O}$ $8 \mathrm{NH}_{3}+3 \mathrm{Cl}_{2} \longrightarrow \mathrm{~N}_{2}+6 \mathrm{NH}_{4}^{\mathrm{Cl}}$ (20morm)
 the solubility ( 10 mols )
Down the group, $*_{n}$ The cationic radius is the same lattice energy is decreasing. *. Down the radius is increasing, hydration
*. When the cationic radius is increasing, energy is decreasing.
*. When going down the group, the hydration energy of ion decreases in a higheramount than decreasing of lattice energy.

Down the group, the solution enthalpy of the sulphates is $(t) v e . \quad(02 \times 6=12 \mathrm{mols})$
$\pi$. Melrira point LiI $<$ Li Br LLiCl<LiF (moors) *. Cation is the same, $\checkmark$

* The charge of the anion is the same.
* radius $\mathrm{F}^{-}<\mathrm{CI}^{-}<\mathrm{Br}^{-}<\mathrm{I}^{-}$
* polarizability $\mathrm{F}^{-}<\mathrm{Cl}^{-}<\mathrm{Br}^{-}<\mathrm{I}^{-}$
*, court property of the th Life $<$ LiB $<L I C I<L i f$ $\therefore$ melting point is increarmg

$$
\text { (5 } 52=10 \mathrm{monly})
$$

(10)
(a)

$$
\text { (a) i. } \begin{aligned}
& \mathrm{IO}_{3}^{-}+5 \mathrm{I}^{-}+6 \mathrm{H}^{+} \rightarrow 3 \mathrm{I}_{2}+3 \mathrm{H}_{2} \mathrm{O} \\
& \mathrm{I}_{2}+2 \mathrm{Na}_{2} \mathrm{~S}_{2} \mathrm{O}_{3} \rightarrow 2 \mathrm{NaI}+\mathrm{Na}_{2} \mathrm{~S}_{4} \mathrm{O}_{6} \\
&(10 \times 2=20 \mathrm{mals})
\end{aligned}
$$

$$
n_{\mathrm{Nq}_{2} \mathrm{~S}_{2} \mathrm{O}_{3}}=\frac{0.1 \times 20}{1000}=2 \times 10^{-3} \mathrm{~mol}
$$

$$
n I_{2} \quad=1 \times 10^{-3} \mathrm{~mol}
$$

$$
n_{H^{+}}=2 \times 10^{-3} \mathrm{~mol}
$$

$$
\begin{aligned}
n_{H}+\text { present in } 500 \mathrm{~cm}^{3} & \left.=\frac{2 \times 10^{-3} \times 500}{25}=0.04 \mathrm{mo}\right) \\
n_{H^{+}} \text {of } 50 \mathrm{~cm}^{3} & =0.04 \mathrm{~mol} \\
{\left[\mathrm{H}^{+}\right] \text {in waste watè } } & =\frac{0.04 \times 10^{3}}{50} \mathrm{~V} \\
& =0.8 \text { mold ī }
\end{aligned}
$$

(b)
(e). i. $\mathrm{Fe}^{2+}, \mathrm{iNH}_{\mathrm{A}}{ }^{+}, \mathrm{SO}_{4}{ }^{2-} \quad\left(\mathrm{O}_{3 \times 3} 3\right.$ 2 9males)

$$
\begin{aligned}
& \text { ii. } \mathrm{C} \rightarrow \mathrm{NH}_{3} \text {, } \mathrm{J} \rightarrow \mathrm{NH}_{4} \mathrm{Cl}, \quad \mathrm{~K}=\mathrm{BaSO}_{4} \\
& \text { ( } 03 \times 3=9 \text { ) } \\
& \text { pipp. }\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+},\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}_{6}\right]^{3+} \quad(03 \times 2=06)\right. \\
& \text { iv. } \left.\mathrm{F} \rightarrow \mathrm{Fe}(\mathrm{OH})_{2} \quad G=\mathrm{Fe}_{2} \mathrm{OH}\right)_{3}(03 \times 2=6) \\
& \text { V. }\left(\mathrm{NH}_{4}\right)_{2} \mathrm{SO}_{4}+2 \mathrm{NaOH} \longrightarrow \mathrm{NCSO}_{4}+2 \mathrm{NH}_{3}+42 \\
& 2\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{8}\right]^{8+}+\mathrm{Zn} \rightarrow 2\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}_{6}\right]^{2+}+\mathrm{Zn}^{2+}\right. \\
& \text { (10) }
\end{aligned}
$$

$$
\begin{aligned}
& \text { iii }\left[\mathrm{CO}\left(\mathrm{NH}_{3}\right)_{6}\right]^{3+} \text { - dark brown } \\
& \text { ir }\left[\mathrm{COCl}_{4}\right]^{2} \quad \text {-blue. } \\
& \text { r. }\left[\mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{4}\right]^{2+} \quad \text { - blue. } \\
& \text { ( } 10 \times 5=50 \text { marlas) }
\end{aligned}
$$

