

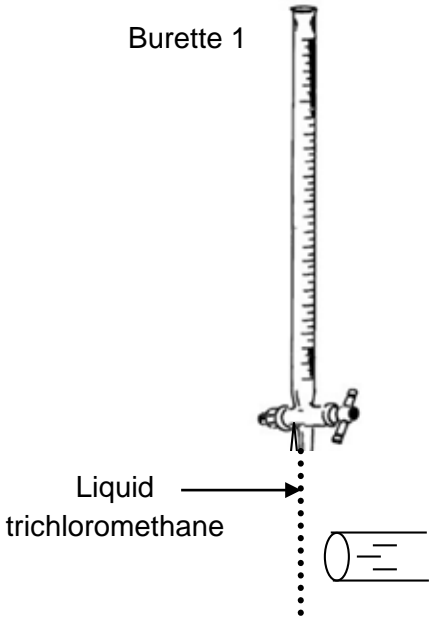
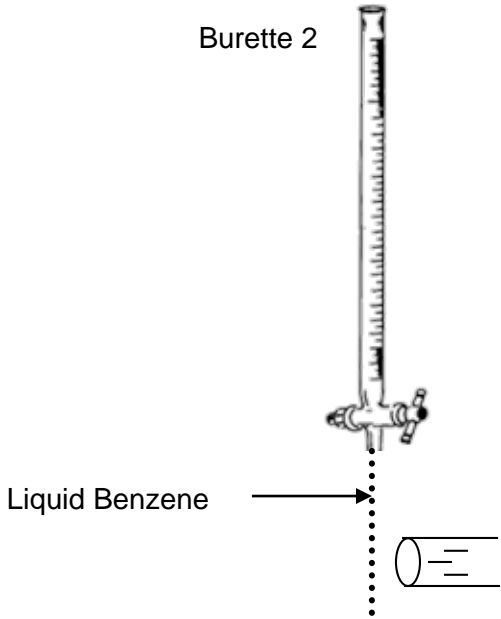
Section A

Answer all questions

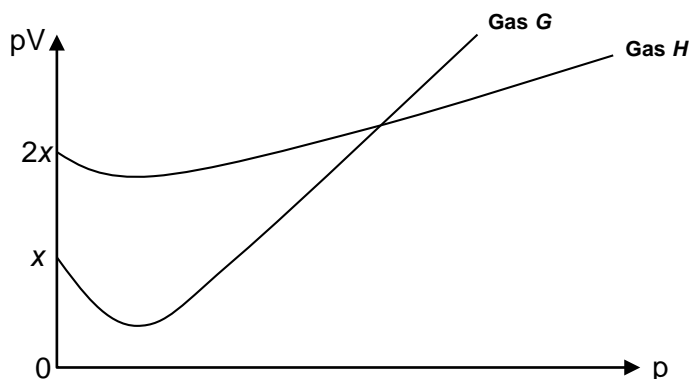
| | | | | | | | | |
|---|--|------|---|------|---|------|---|------|
| 1 | <p>To determine the percentage of nitrogen present in a snack, 1.0 g of the snack was boiled with concentrated sulphuric acid to convert all the nitrogen into ammonium sulphate. The ammonium salt obtained was then boiled with excess aqueous sodium hydroxide to liberate the ammonia, which was passed into 25.0 cm³ of 0.20 mol dm⁻³ hydrochloric acid. The unreacted hydrochloric acid required 20.0 cm³ of 0.10 mol dm⁻³ aqueous sodium hydroxide for complete neutralisation.</p> <p>What is the percentage by mass of nitrogen in the snack?</p> | | | | | | | |
| | A | 2.8% | B | 4.2% | C | 7.2% | D | 8.4% |
| <p>Answer: B</p> <p>Amt of NH₃ = $(\frac{25}{1000} \times 0.20) - (\frac{20}{1000} \times 0.10) = 0.003 \text{ mol}$</p> <p>$2\text{NH}_3 \equiv 1(\text{NH}_4)_2\text{SO}_4 \equiv 2\text{N}$</p> <p>Mass of nitrogen in snack = $0.003 \times 14.0 = 0.042 \text{ g}$</p> <p>% by mass = $\frac{0.042}{1.0} \times 100 = \underline{\underline{4.2\%}}$</p> | | | | | | | | |

| | | | | | | | | |
|---|--|----|----------|----|----------|----|----------|----|
| 2 | <p>In an experiment, 25.0 cm³ of 0.20 mol dm⁻³ solution of K₂AO₄ reacted exactly with 25.0 cm³ of 0.10 mol dm⁻³ aqueous sodium sulfate(IV). The half-equation for the oxidation of the sulfate(IV) ion is shown below.</p> $\text{SO}_3^{2-} (\text{aq}) + \text{H}_2\text{O} (\text{l}) \rightarrow \text{SO}_4^{2-} (\text{aq}) + 2\text{H}^+ (\text{aq}) + 2\text{e}^-$ <p>Calculate the final oxidation state of A.</p> | | | | | | | |
| | A | +2 | B | +3 | C | +4 | D | +5 |
| | <p>Answer: D</p> <p>$2\text{K}_2\text{AO}_4 \equiv \text{SO}_3^{2-}$</p> <p>No. of e gained by A in K₂AO₄ = No. of e lost by S in SO₃²⁻ = 2e</p> <p>$2\text{A}^{6+} + 2\text{e} \rightarrow 2\text{A}^{n+}$</p> <p>$12 - 2 = 2n$</p> <p>$n = \underline{\underline{+5}}$</p> | | | | | | | |

| | | | | |
|------------------|---|-------------------------|-------------------------|--|
| 3 | Two elements D and E have the following properties. | | | |
| | <ul style="list-style-type: none"> D and E form ionic compounds Na_2D and Na_2E respectively. Element E forms EF_6 molecules whereas D is not able to do so. <p>Which pair of electronic configurations for D and E is correct?</p> | | | |
| | | D | E | |
| | A | $[\text{He}] 2s^2 2p^2$ | $[\text{Ne}] 3s^2 3p^4$ | |
| | B | $[\text{He}] 2s^2 2p^2$ | $[\text{Ne}] 3s^2 3p^2$ | |
| | C | $[\text{He}] 2s^2 2p^4$ | $[\text{Ne}] 3s^2 3p^2$ | |
| | D | $[\text{He}] 2s^2 2p^4$ | $[\text{Ne}] 3s^2 3p^4$ | |
| Answer: D | | | | |

| | | | | |
|---|---|--------------------------------|---------------------------|--|
| 4 | <p>The diagram below shows liquid trichloromethane and liquid benzene flowing from burettes 1 and 2 respectively.</p> <div style="display: flex; justify-content: space-around; align-items: center;"> <div style="text-align: center;"> <p>Burette 1</p>  <p>Liquid trichloromethane</p> </div> <div style="text-align: center;"> <p>Burette 2</p>  <p>Liquid Benzene</p> </div> </div> <p>What would happen to the flow of the liquids trichloromethane and benzene when a negatively-charged rod is brought near to each of them?</p> | | | |
| | | Liquid trichloromethane | Liquid benzene | |
| | A | Deflected towards the rod | Deflected towards the rod | |
| | B | Undeflected | Deflected towards the rod | |
| | C | Deflected towards the rod | Undeflected | |
| | D | Undeflected | Undeflected | |
| <p>Answer: C</p> <p>Liquid trichloromethane is polar. The partial positive charge can be attracted by the negatively charged rod.</p> <p>Benzene is non-polar and does not have partial charges.</p> | | | | |

- 5** The value of pV is plotted against p for two gases, **G** and **H**, where p is the pressure and V is the volume of the gas.



Which of the following could be the identities of the gases?

| | | Gas G | Gas H |
|----------|--|----------------------------|----------------------------|
| A | | 0.5 mol of H_2 at 25 °C | 0.5 mol of H_2 at 50 °C |
| B | | 0.5 mol of H_2 at 25 °C | 1 mol of SO_2 at 25 °C |
| C | | 0.5 mol of SO_2 at 25 °C | 0.5 mol of SO_2 at 50 °C |
| D | | 0.5 mol of SO_2 at 25 °C | 1 mol of H_2 at 25 °C |

Answer: D

Amount of gas **H** should be twice the amount of gas **G**.

According to the shape of the curves, gas **G** should be a less ideal gas than gas **H**.

- 6** During an inspection, a small spacecraft of capacity 20 m³ was connected to another of capacity 50 m³. Before connection, the pressure inside the smaller craft was 40 atm and that inside the larger one was 150 atm.

Given that all measurements were made at the same temperature, What is the pressure of the system after the connection?

| | | | | | | | |
|----------|--------|----------|--------|----------|---------|----------|---------|
| A | 78 atm | B | 95 atm | C | 119 atm | D | 190 atm |
|----------|--------|----------|--------|----------|---------|----------|---------|

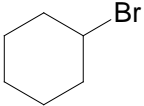
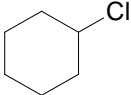
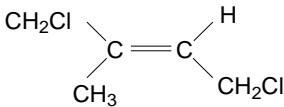
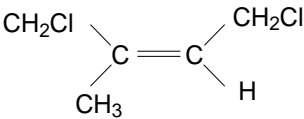
Answer: C

$$\text{Amt of gas in small spacecraft} = \frac{pV}{RT} = \frac{(40)(20)}{RT} = \frac{800}{RT}$$

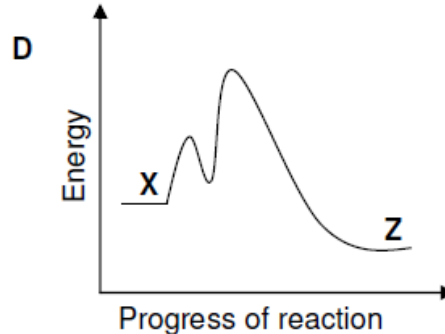
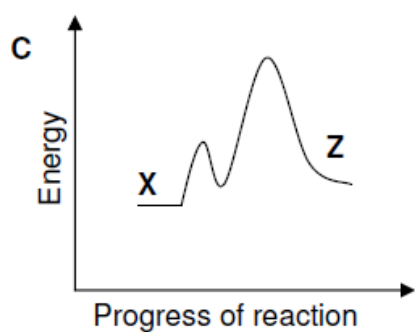
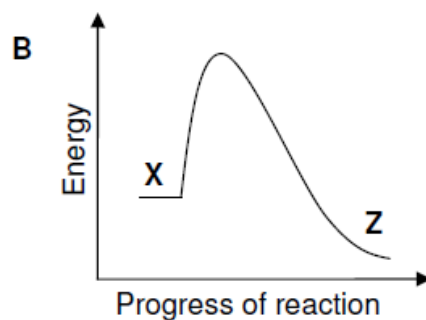
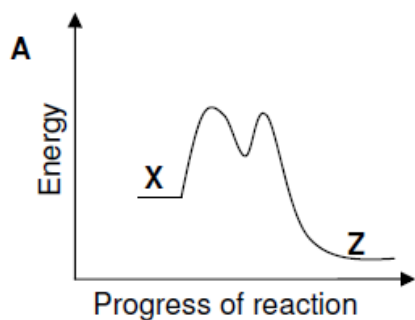
$$\text{Amt of gas in large spacecraft} = \frac{pV}{RT} = \frac{(150)(50)}{RT} = \frac{7500}{RT}$$

$$\text{total amt of moles of gas} = \frac{800 + 7500}{RT} = \frac{8300}{RT}$$

$$\text{Pressure in the combined arrangement} = \frac{nRT}{V} = \frac{\left(\frac{8300}{RT}\right)RT}{(20 + 50)} = 119 \text{ atm}$$

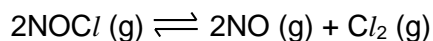
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| 7 | In which of the following pairs of compounds will compound II have a higher boiling point than compound I ? | | | | |
| | | | I | II | |
| A | | |  |  | |
| B | | | $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ | $\text{C}(\text{CH}_3)_4$ | |
| C | | | $\text{CH}_3\text{CH}_2\text{CH}_2\text{COOH}$ | $\text{CH}_3\text{CH}_2\text{CH}_2\text{OH}$ | |
| D | | |  |  | |
| <p>Answer: D</p> <p>A: I has higher Mr than II and hence has more extensive intermolecular VDW forces of attraction and a higher boiling point.</p> <p>B: I is linear and II is branched and hence has more extensive intermolecular VDW forces of attraction and a higher boiling point.</p> <p>C: I has more extensive intermolecular hydrogen bonding than II and hence a higher boiling point.</p> <p>D: I is non-polar and has temporary dipole-induced dipole interactions while II is polar and has permanent dipole-dipole interactions. Hence II has a higher boiling point.</p> | | | | | |

| | |
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| 8 | <p>The conversion of compound X into Z was exothermic and proceeded by two steps, where Y was the intermediate. The steps involved were:</p> <p>Step 1: X → Y</p> <p>Step 2: Y → Z</p> <p>It was found that Step 1 is the rate-determining step. Which diagram represents the energy level diagram for the reaction?</p> |
|---|---|



Answer: **A**

| | |
|---|--|
| 9 | Pure nitrosyl chloride, NOCl gas, was heated at 320°C in a 2.0 dm ³ vessel. At equilibrium, 30% of the NOCl gas had dissociated according to the equation below and the total pressure was P atm. |
|---|--|



What is value of the equilibrium constant, K_p ?

| | | | | | | | |
|----------|------------------|----------|------------------|----------|---------|----------|---------|
| A | $\frac{17.9}{p}$ | B | $\frac{41.7}{p}$ | C | 0.0120p | D | 0.0130p |
|----------|------------------|----------|------------------|----------|---------|----------|---------|

Answer: **C**

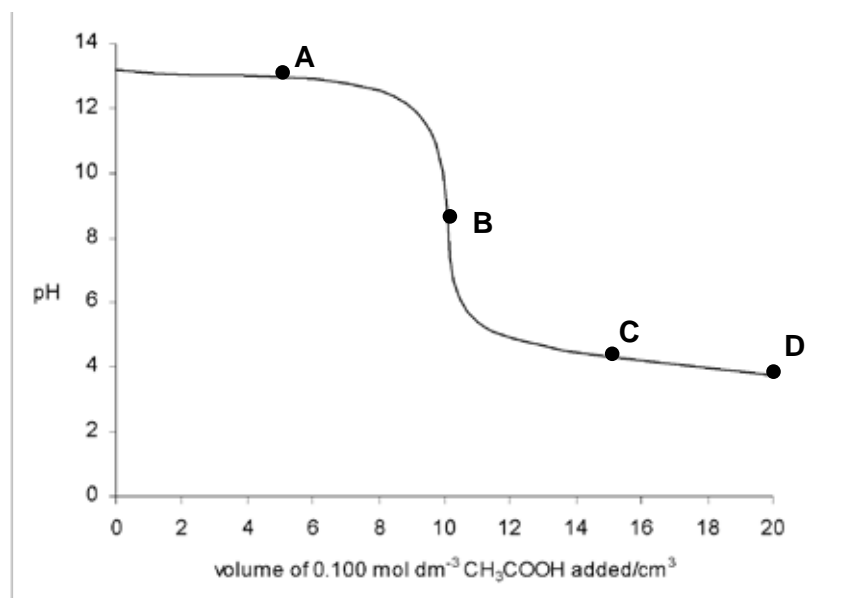
| | | | |
|---|---|----------|----------|
| | $2\text{NOCl (g)} \rightleftharpoons 2\text{NO (g)} + \text{Cl}_2 \text{(g)}$ | | |
| Initial partial pressure/atm | x | 0 | 0 |
| Change in partial pressure / atm | -0.3x | +0.3x | +0.15x |
| Equilibrium partial pressure / atm | 0.7x | 0.3x | 0.15x |
| | 0.6087p | 0.2609p | 0.1304p |

$$0.7x + 0.3x + 0.15x = p$$
$$x = 0.8696p$$

$$\begin{aligned} K_p &= \frac{(0.2609p)(0.1304p)^2}{(0.6087p)^2} \\ &= 0.01197 \\ &= 0.0120p \end{aligned}$$

- 10** The pH change when $0.100 \text{ mol dm}^{-3} \text{ CH}_3\text{COOH}$ is added drop-wise to 10.0 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ NaOH (aq)}$ is shown below.

At which point on the graph does $\text{pH} = \text{p}K_a$, where K_a is the acid dissociation constant of the weak acid?



Answer: **D**

Regions **C** and **D** are where there is an excess of weak acid CH_3COOH as well as the salt $\text{CH}_3\text{COO}^- \text{Na}^+$ that is formed. Hence, buffer region.

At Region **D**,

$$\text{Amt of } \text{CH}_3\text{COO}^- \text{Na}^+ \text{ formed} = \frac{10}{1000} \times 0.100 = 0.001 \text{ mol}$$

$$\text{Amt of excess } \text{CH}_3\text{COOH} \text{ added} = \frac{(20 - 10)}{1000} \times 0.100 = 0.001 \text{ mol}$$

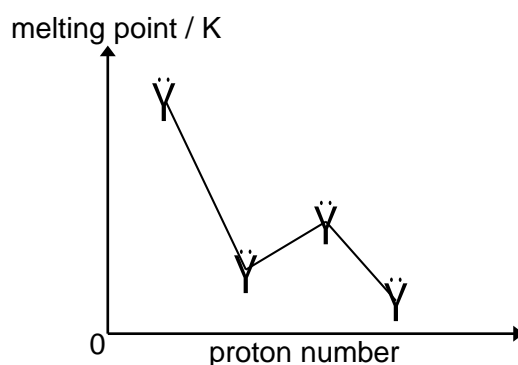
Amt of $\text{CH}_3\text{COO}^- \text{Na}^+$ and CH_3COOH is the same, hence this is the point where there is maximum buffering capacity and $\text{pH} = \text{p}K_a$.

| | | | | | | | | |
|--|---|------------------------|----------|-----------------------|----------|-----------------------|----------|-----------------------|
| 11 | <p>In an experiment, 70 cm³ of water at 25°C was brought to boiling point by burning butane in excess oxygen. Given that the standard enthalpy change of combustion of butane is -2877 kJ mol⁻¹, calculate the volume of butane needed if this process is only 85% efficient.</p> <p>Assume that the specific heat capacity of water is 4.2 J g⁻¹ K⁻¹ and that 1 mole of gas occupies 24 dm³ under the given conditions.</p> | | | | | | | |
| | A | 0.0721 dm ³ | B | 0.156 dm ³ | C | 0.184 dm ³ | D | 0.216 dm ³ |
| <p>Answer: D</p> <p>Since $Q' = mc\Delta T = 70 \times 4.2 \times (100.0 - 25.0) = 22050 \text{ J}$</p> <p>Apparent amount of heat absorbed by water, Q'</p> $= \frac{85}{100} Q \text{ (Actual amount of heat evolved by burning butane)}$ <p>Actual amount of heat evolved, $Q = \frac{100}{85} \times 22050 = 25941 \text{ J}$</p> $\Delta H_c^\ominus(\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3) = -\frac{25941}{n} = -2877 \times 10^3$ <p>Amount of butane = $9.017 \times 10^{-3} \text{ mol}$</p> <p>Volume of butane = $9.017 \times 10^{-3} \times 24 = \underline{\underline{0.216 \text{ dm}^3}}$</p> | | | | | | | | |

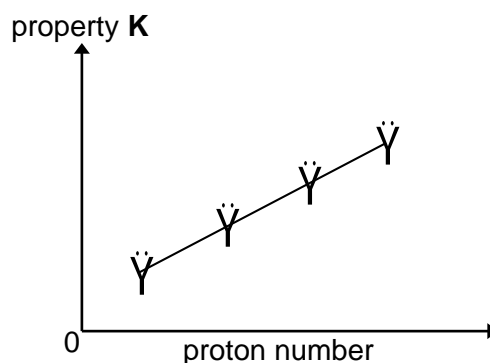
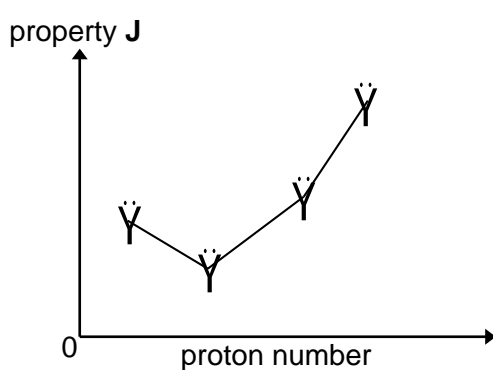
| | | |
|---|---|---|
| 12 | Which of the following changes does not alter the E^\ominus value measured for a Cl_2/Cl^- half-cell that is under standard conditions? | |
| | A | Adding water into the half-cell. |
| | B | Placing the half-cell in an ice bath. |
| | C | Adding copper(II) ions into the half-cell. |
| | D | Introducing an inert gas into the half-cell at a pressure of 1 atm through a separate inlet from the Cl_2 gas inlet. |
| <p>Answer: D</p> <p>A: Dilution causes the concentration of Cl^- ions to be lower than 1 mol dm^{-3}.</p> <p>B: The ice bath lowers the temperature of the half-cell to less than 298 K.</p> <p>C: Cu^{2+} ions will form a complex with Cl^- ions and lower the concentration of Cl^- ions to less than 1 mol dm^{-3}.</p> <p>D: Introducing an inert gas through a separate inlet does not affect the pressure of Cl_2 gas hence doing so does not affect the E^\ominus value of the half-cell.</p> | | |

| | | |
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| 13 | Which statement concerning the chlorine-containing compounds of elements in the third period, sodium to argon, is correct? | |
| | A | NaCl dissolves easily in water due to favourable ion-dipole interactions and the compound with the highest electrical conductivity in molten state is AlCl_3 . |
| | B | PCl_3 and Cl_2O_7 are both acidic in nature due to hydration of the compounds in water. |
| | C | The low boiling points of PCl_3 and Cl_2O_7 are due to the weak bond energies involved in the P-Cl bonds and the Cl-O bonds. |
| | D | The different oxidation states of chlorine in NaCl and Cl_2O_7 are due to the relative electronegativities of the pairing element and chlorine. |
| <p>Answer: D</p> <p>A: AlCl_3 is a simple molecular structure, it does not conduct electricity.</p> <p>B: PCl_3 and Cl_2O_7 are both acidic in nature due to <u>hydrolysis</u> of the compounds in water.</p> <p>C: The low boiling points of PCl_3 and Cl_2O_7 are due to the weak <u>intermolecular</u> forces of attraction present and not related to bond energy.</p> | | |

- 14** The diagram represents the melting points of four consecutive elements in the third period of the Periodic Table.



The sketches below represent another two properties of the elements.



What are properties **J** and **K**?

| | | property J | property K |
|------------------|----------|-----------------------------|-------------------|
| | A | third ionisation energy | electronegativity |
| | B | number of valence electrons | boiling point |
| | C | ionic radius | nuclear charge |
| | D | electrical conductivity | atomic radius |
| Answer: A | | | |

| | | |
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| 15 | Which of the following is true about the thermal decomposition of magnesium nitrate? | |
| | A | Sodium nitrate is thermally unstable as compared to magnesium nitrate. |
| | B | Every 1 mole of magnesium nitrate burnt gives 2.5 moles of gaseous products. |
| | C | The decomposition temperature of magnesium nitrate is higher than that of barium nitrate. |
| | D | The solid product of the thermal decomposition of magnesium nitrate readily dissolves in water to give an alkaline solution. |
| <p>Answer: B</p> <p>A: Sodium nitrate is more thermally stable than magnesium nitrate due to the lower charge density, and hence lower polarising power, of Na^+ as compared to Mg^{2+}, leading to a smaller polarising effect on the NO_3^- electron cloud.</p> <p>B: $2\text{Mg}(\text{NO}_3)_2 (\text{s}) \rightarrow 2\text{MgO} (\text{s}) + 4\text{NO}_2 (\text{g}) + \text{O}_2 (\text{g})$</p> <p>C: The decomposition temperature of magnesium nitrate is lower than that of barium nitrate due to the higher charge density, and hence greater polarising power, of Mg^{2+} as compared to Ba^{2+}, leading to a greater polarising effect on the NO_3^- electron cloud.</p> <p>D: $\text{MgO} (\text{s})$ dissolves very slowly, if at all, in water.</p> | | |

| | | |
|--|--|---|
| 16 | Which of the following gives the best description of the reactions of Group II metals and their compounds? | |
| | A | All Group II metals react with steam to give hydrogen gas. |
| | B | Barium oxide is stored in oil due to its explosive reaction with oxygen gas in air. |
| | C | All Group II oxides undergo neutralisation with hot acids to give a salt and water. |
| | D | Beryllium hydroxide is amphoteric due to the high charge density of the Be^{2+} ion. |
| <p>Answer: C</p> <p>A: $\text{Be} (\text{s})$ does not react with steam.</p> <p>B: $\text{Ba} (\text{s})$, not barium oxide, is explosive in air.</p> <p>D: The acidic nature of $\text{Be}(\text{OH})_2$ can be explained by its high charge density but not its basic nature.</p> | | |

17

The table below shows the results of experiments in which the halogens, P_2 , Q_2 and R_2 were added to separate aqueous solutions containing P^- , Q^- and R^- ions.

| | P^- (aq) | Q^- (aq) | R^- (aq) |
|-------|--------------|-------------|--------------|
| P_2 | no reaction | no reaction | R_2 formed |
| Q_2 | P_2 formed | no reaction | R_2 formed |
| R_2 | no reaction | no reaction | no reaction |

In which sequence is the solubility of the silver halides in aqueous ammonia arranged in **increasing** order?

| | |
|----------|-------------------|
| A | $AgP < AgQ < AgR$ |
| B | $AgQ < AgP < AgR$ |
| C | $AgQ < AgR < AgP$ |
| D | $AgR < AgP < AgQ$ |

Answer: **D**


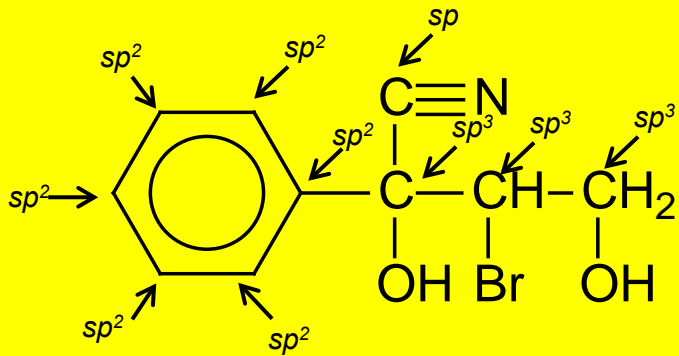
From the table R_2 is the weakest oxidising agent while Q_2 is the strongest oxidising agent.

A logical deduction means Q_2 is chlorine, and R_2 is iodine.

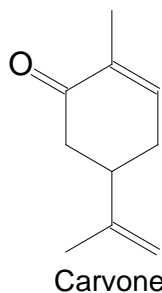
Solubility of silver halides in NH_3 (aq) in increasing order is AgR , AgP and AgQ .

| | | | | |
|-----------|--|------------------------------|---------------------------|---------------------------|
| 18 | Aqueous sodium hydroxide was added to a pale green solution of a mixture of two metal cations. The resulting precipitate was treated with excess ammonia solution giving an intense deep blue mixture. The mixture was filtered giving a reddish-brown residue and a deep blue filtrate. The residue was washed with deionised water and treated with excess acidified ammonium thiocyanate giving a blood red solution. | | | |
| | Which of the following substances best explains these observations? | | | |
| | | <i>reddish-brown residue</i> | <i>deep blue filtrate</i> | <i>blood-red solution</i> |
| A | $Fe(OH)_2$ | $Cu(OH)_2$ | $[Fe(H_2O)_6]^{2+}$ | |
| B | $Fe(OH)_3$ | $[Cu(NH_3)_4]^{2+}$ | $[Fe(H_2O)_6]^{3+}$ | |
| C | $Fe(OH)_2$ | $[Cu(H_2O)_6]^{2+}$ | $[Fe(H_2O)_5SCN]^{2+}$ | |
| D | $Fe(OH)_3$ | $[Cu(NH_3)_4]^{2+}$ | $[Fe(H_2O)_5SCN]^{2+}$ | |
| Answer: D | | | | |

| | | |
|---|---|---|
| 19 | Rhodium and its compounds are used as catalysts in many important reactions. Which of the following properties makes rhodium a suitable heterogeneous catalyst? | |
| | A | Rhodium has vacant d-orbitals of suitable energy. |
| | B | Rhodium exhibits variable oxidation states. |
| | C | Rhodium is able to form stable complexes. |
| | D | Rhodium is able to form coloured compounds. |
| <p>Answer: A</p> <p>The only answer to explain the feature of transition elements which allow them to function as heterogeneous catalyst.</p> <p>Statement B is more for homogeneous catalyst.</p> | | |

| | | |
|---|---|---------|
| 20 | <p>Compound S was used in the following synthesis route.</p> <div style="text-align: center;">  <p>Compound S Compound T Compound U</p> </div> <p>What are the numbers of sp, sp^2 and sp^3 hybridised carbon atoms in compound U?</p> | |
| | A | 0, 6, 4 |
| | B | 0, 7, 3 |
| | C | 1, 6, 3 |
| | D | 1, 7, 2 |
| <p>Answer: C</p> <p>Compound U is</p> <div style="text-align: center;">  </div> | | |

- 21** Carvone is the main active ingredient found in spearmint and it has the following structure:



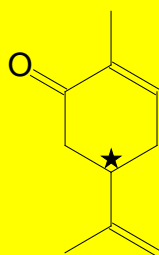
Carvone can be reduced to compound **V** by heating with hydrogen gas in the presence of nickel.

How many stereoisomers do the molecules of carvone and **V** each have?

| | | | Carvone | Compound V |
|----------|--|--|---------|------------|
| A | | | 0 | 4 |
| B | | | 0 | 8 |
| C | | | 2 | 4 |
| D | | | 2 | 8 |

Answer: **D**

Carvone has one chiral carbon as shown:



hence it will have $2^1 = 2$ optical isomers

Compound **V** has 3 chiral carbon as shown:



hence it will have $2^3 = 8$ optical isomers

| | | |
|--|---|---|
| 22 | How many alcohols (including both structural and stereoisomers) can have the molecular formula $C_4H_{10}O$? | |
| | A | 3 |
| | B | 4 |
| | C | 5 |
| | D | 6 |
| <p>Answer: C</p> <p>$CH_3CH_2CH_2CH_2OH$ $CH_3CH_2CH(OH)CH_3$ (presence of chiral carbon \Rightarrow 2 isomers)</p> <p>$CH_3C(CH_3)CH_2OH$ $CH_3C(CH_3)(OH)CH_3$</p> | | |

| | | | | |
|-----------|--|------------------|----------|------------------|
| 23 | A sample of ethene was added to a solution of Br_2 (aq) and $NaCl$ (aq). Which of the following products is not likely to be found in the reaction mixture? | | | |
| | A | $CH_2(OH)CH_2Br$ | C | $CH_2(OH)CH_2Cl$ |
| | B | CH_2BrCH_2Cl | D | CH_2BrCH_2Br |

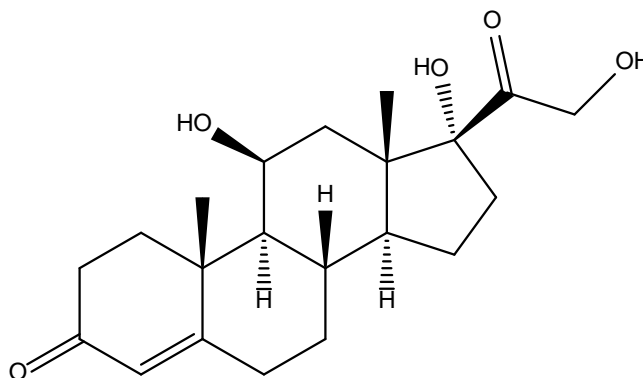
Answer: **C**

In the electrophilic substitution of ethene and $Br_2(aq)$, the first step is:

carbocation intermediate

Hence, the product must contain at least one Br.

- 24** Hydrocortisone is a steroid hormone produced by the adrenal gland and is released in response to stress. It is commonly used as an active ingredient in anti-inflammatory creams.



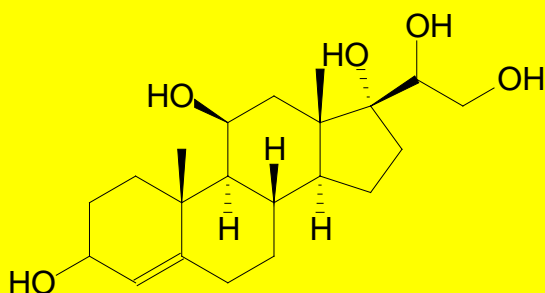
Hydrocortisone

Which of the following statement about hydrocortisone is true?

- | | |
|----------|---|
| A | When treated with an excess of hot concentrated acidified KMnO_4 , it forms a compound containing 7 carbonyl groups. |
| B | When warmed with aqueous alkaline iodine, a yellow precipitate is observed. |
| C | When treated with cold dilute KMnO_4 , it forms a compound containing 2 hydroxy groups. |
| D | When treated with NaBH_4 in the presence of methanol, it forms a compound containing 5 hydroxy groups. |

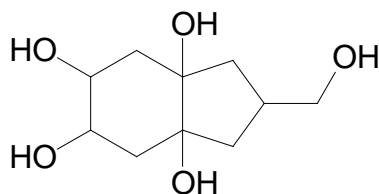
Answer: **D**

Ketone undergoes reduction to form:

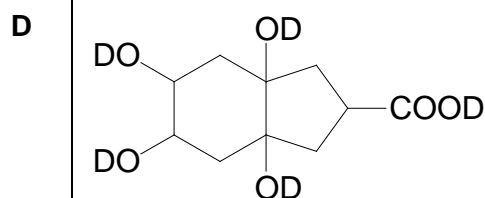
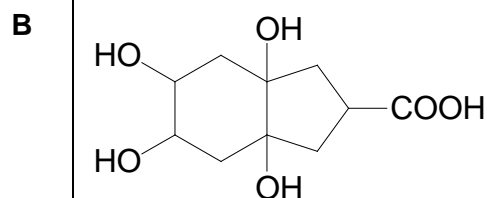
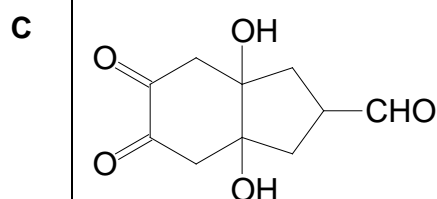
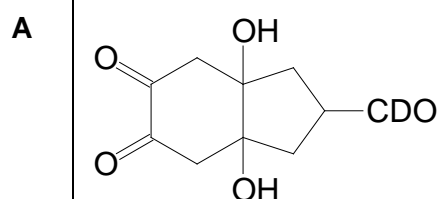


- A:** $\text{C}=\text{C}$, primary and secondary alcohol undergo oxidation to form a compound with 4 carbonyl groups.
- B:** No $\text{CH}_3\text{CO}-$ or $-\text{CH}(\text{CH}_3)(\text{OH})$ group observed.
- C:** Only alkene undergoes mild oxidation to form a diol, a compound with 5 hydroxy groups is formed.

| | |
|----|---|
| 25 | Deuterium, D, is an isotope of hydrogen, ${}^2_1\text{H}$. |
|----|---|

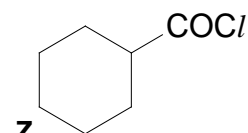
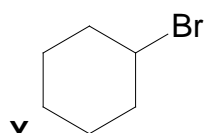
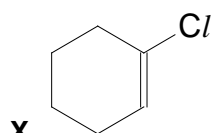
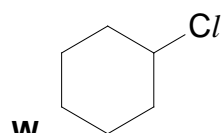


Which of the following is the product formed when the above compound reacts with hot $\text{K}_2\text{Cr}_2\text{O}_7$ in aqueous D_2SO_4 under controlled conditions?



Answer: **C**

26 Which of the following shows the correct sequence in order of increasing ease of hydrolysis for the compounds below?



| | |
|---|-----------------|
| A | $X < W < Y < Z$ |
|---|-----------------|

| | |
|----------|--|
| B | $Z < X < W < Y$ |
|----------|--|

| | |
|----------|--|
| C | $Z < Y < X < W$ |
|----------|--|

| | |
|---|-----------------|
| D | $W < X < Y < Z$ |
|---|-----------------|

Answer: **A**

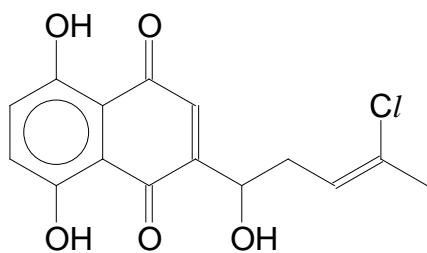
Acyl chloride, Z undergoes hydrolysis readily at rtp.

Strength of C-X bond: $\text{C-Cl} > \text{C-Br}$

Ease of hydrolysis: $R-Cl < R-Br$

For X, presence of double bond strengthen C-Br bond making nucleophilic substitution less readily to happen.

27

Which of the following statement about compound **A** is true?Compound **A**

- | | |
|----------|--|
| A | When treated with alkaline Tollens' reagent, it forms a compound with molecular formula, $C_{15}H_{13}O_5Cl$. |
| B | When reacted with PCl_5 , 1 mole of HCl (g) are formed. |
| C | When heated with H_2 (g) in the presence of Ni catalyst, it forms a compound containing 3 hydroxy groups. |
| D | When treated with sodium hydroxide, hydrolysis occurred. |

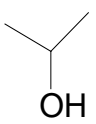
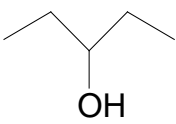
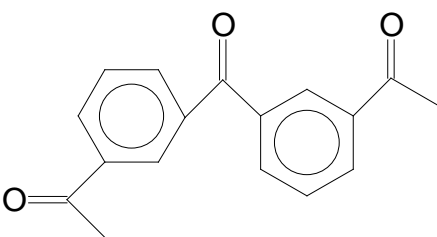
Answer: **B**

A: Compound with formula, $C_{15}H_{11}O_5Cl$ is formed as phenol undergoes neutralisation with alkaline Tollens' reagent.

B: Only 1 mole of $HCl(g)$ is formed as phenol do not react with PCl_5

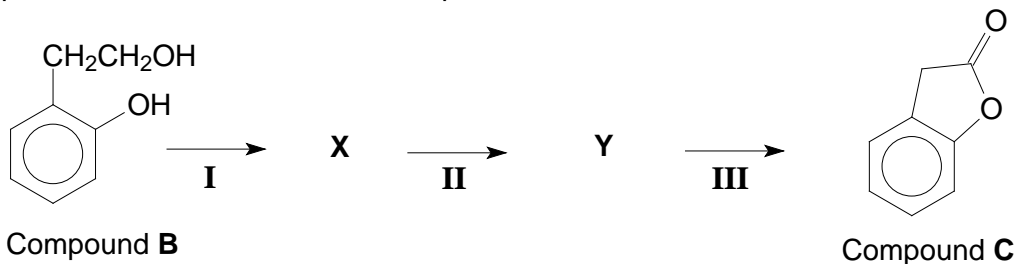
C: After reduction of ketone and $C=C$, a compound containing 5 hydroxy groups is formed.

D: Neutralisation occurred not hydrolysis.

| | | |
|----|--|-------|
| 28 | <p>What is the ratio of sodium iodide formed when each of the three compounds reacts with alkaline aqueous iodine?</p> <div style="display: flex; justify-content: space-around; align-items: center; margin-top: 20px;"> <div style="text-align: center;">  <p>OH</p> </div> <div style="text-align: center;">  <p>OH</p> </div> <div style="text-align: center;">  </div> </div> | |
| | A | 1:1:1 |
| | B | 5:0:6 |
| | C | 5:0:3 |
| | D | 1:1:2 |
| | <p>Answer: B</p> | |

29

Compound **B** can be converted to compound **C** as shown below.



Which of the following statements is correct with regards to the given reaction scheme?

A Step **I** may involve the use of PCl_5 .

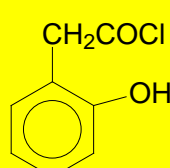
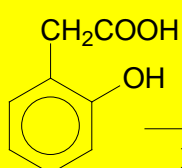
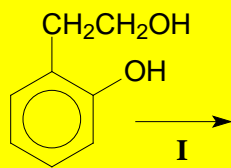
B Step **I** may involve the use of hot acidified potassium manganate(VII).

C Step **III** may involve the use of aqueous sodium hydroxide.

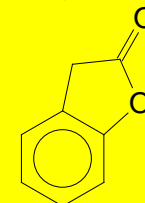
D Step **III** may involve the use of hot concentrated sulfuric acid.

Answer: **C**

Compound **B**



Compound **C**



Step **I**: hot, acidified $\text{K}_2\text{Cr}_2\text{O}_7$

Step **II**: PCl_5 , rtp

Step **III**: NaOH (aq), rtp (to form phenoxide)

| | | | |
|----------|--|----------|---|
| 30 | <p>A polypeptide, was partially hydrolysed using two different enzymes and the fragments were separated. Analysis of the fragments gave the following results:</p> <p>Using enzyme D: glu-val-phe glu-asp-leu ala-ser val-ala</p> <p>Using enzyme E: val-phe ala-glu-asp val ser-glu leu-ala</p> <p>What is the amino acid sequence of the polypeptide?</p> | | |
| | <table border="1"> <tr> <td data-bbox="260 763 331 831">A</td><td data-bbox="331 763 1426 831">leu-ala-ser-glu-val-phe-val-ala-glu-asp</td></tr> </table> | A | leu-ala-ser-glu-val-phe-val-ala-glu-asp |
| A | leu-ala-ser-glu-val-phe-val-ala-glu-asp | | |
| | <table border="1"> <tr> <td data-bbox="260 831 331 898">B</td><td data-bbox="331 831 1426 898">ala-glu-asp-leu-ala-ser-glu-val-phe-val</td></tr> </table> | B | ala-glu-asp-leu-ala-ser-glu-val-phe-val |
| B | ala-glu-asp-leu-ala-ser-glu-val-phe-val | | |
| | <table border="1"> <tr> <td data-bbox="260 898 331 965">C</td><td data-bbox="331 898 1426 965">glu-val-phe-ala-glu-asp-leu-ala-ser-val</td></tr> </table> | C | glu-val-phe-ala-glu-asp-leu-ala-ser-val |
| C | glu-val-phe-ala-glu-asp-leu-ala-ser-val | | |
| | <table border="1"> <tr> <td data-bbox="260 965 331 1032">D</td><td data-bbox="331 965 1426 1032">val-ala-glu-asp-leu-ala-ser-glu-val-phe</td></tr> </table> | D | val-ala-glu-asp-leu-ala-ser-glu-val-phe |
| D | val-ala-glu-asp-leu-ala-ser-glu-val-phe | | |
| | <p>Answer: D</p> | | |

Section B

For **questions 31-40**, one or more of the numbered statements **1** to **3** may be correct. Decide whether each of the statements is or is not correct. The responses **A** to **D** should be selected on the basis of:

| A | B | C | D |
|----------------------------------|------------------------------------|------------------------------------|-----------------------------|
| 1, 2 and 3 are correct | 1 and 2 only are correct | 2 and 3 only are correct | 1 only is correct |

No other combination of statements is to be used as a correct response.

| | | |
|--|--|-------------|
| 31 | The compound ammonium sulfate is primarily used as a fertiliser for alkaline soils. Which type(s) of bonding is/are found in the compound? | |
| | 1 | ionic |
| | 2 | covalent |
| | 3 | dative bond |
| <p>Answer: A (1, 2 and 3)</p> <div style="text-align: center;"> $2 \left[\begin{array}{c} \text{H} \\ \\ \text{H}-\text{N}-\text{H} \\ \\ \text{H} \end{array} \right]^+ \left[\begin{array}{c} \text{O} \\ \\ \text{O}-\text{S}-\text{O} \\ \\ \text{O} \end{array} \right]^{2-}$ </div> | | |

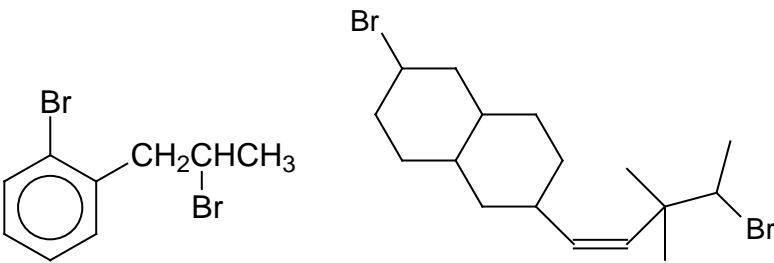
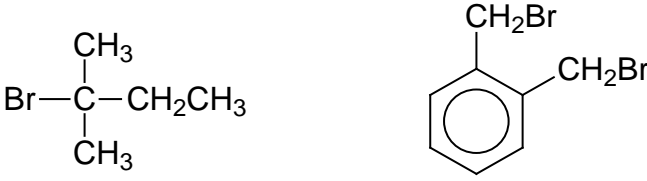
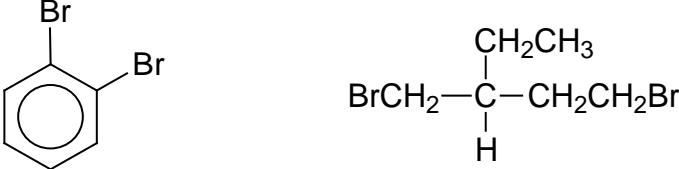
| | | |
|---|--|---|
| 32 | The rate of reaction of a strip of magnesium ribbon and 45 cm ³ of 1.5 mol dm ⁻³ HNO ₃ is determined at 25°C. Which of the following cases would both conditions contribute to an increase in the rate of reaction? | |
| | 1 | Mg powder and 90 cm ³ of 1.5 mol dm ⁻³ HNO ₃ |
| | 2 | Mg powder and 45 cm ³ of 2.0 mol dm ⁻³ HNO ₃ |
| | 3 | 45 cm ³ of 2.0 mol dm ⁻³ HNO ₃ at 35°C |
| <p>Answer: C (2 and 3 only)</p> <p>1: Use of Mg powder increases rate of reaction due to increase in surface area of reaction but increase in volume does not increase rate of reaction.</p> <p>2: Use of Mg powder increases rate of reaction due to increase in surface area of reaction and increase in concentration of HNO₃ increases rate of reaction due to increase in effective collisions.</p> <p>3: Increase in concentration and temperature increases rate of reaction due to increase in effective collisions.</p> | | |

| | | | | | |
|-----------|--|-------------------------|--------------------------------|----------------------|--|
| 33 | Which of the following statements is/are correct for the following equilibrium? $3\text{H}_2(\text{g}) + \text{N}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g}) \quad \Delta H < 0$ | | | | |
| | | Condition | Position of equilibrium | K_p | Rate of formation of NH₃ |
| | 1 | Increase in pressure | Right | No change | Increase |
| | 2 | Decrease in temperature | Right | Increase | Decrease |
| | 3 | Addition of catalyst | Left | No change | Increase |

Answer: **B** (1 and 2 only)

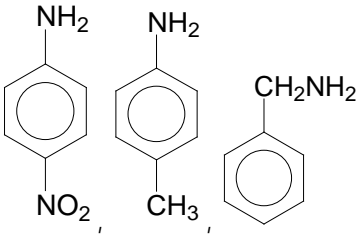
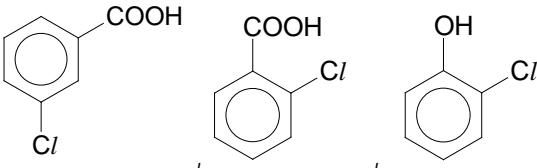
| | | | | |
|----------|-------------------------|---|--|---|
| | Condition | Position of equilibrium | K_p | Rate of forward reaction |
| 1 | Increase in pressure | Right (✓) Position of eqm will shift to decrease amt of gases. | No change (✓) K _p is independent of pressure | Increase (✓) Increase in pressure results in increase number of effective collisions. |
| 2 | Decrease in temperature | Right (✓) Position of eqm will shift to forward exothermic reaction. | Increase (✓) Forward rate constant will decrease less than backward rate constant, hence K _p increases | Decrease (✓) Decrease in temperature leads to lower kinetic energy and decrease in number of effective collisions. |
| 3 | Addition of catalyst | Left (x) Catalyst does not affect eqm position | Increase (x) K _p is independent of catalyst | Increase (✓) Catalyst increases rate of reaction. |

| | | | | |
|---|---|--|-------------------|----------------------------|
| 34 | Which of the following indicators can be used for the titration between ethylamine and hydrochloric acid? | | | |
| | | | Indicator | pH transition range |
| | 1 | | Naphtholphthalein | 7.3 – 8.7 |
| | 2 | | Congo red | 3.0 – 5.0 |
| | 3 | | Azolitmin | 4.5 – 8.3 |
| <p>Answer: C (2 and 3 only)</p> <p>For the titration between ethylamine and hydrochloric acid, it is a strong acid/weak base titration. Hence, at equivalence point of titration, pH increases sharply from » 3 to 7.</p> <p>Only Congo red and Azolitmin pH transition range) lies within the rapid pH change (» 3 to 7) over the equivalence point</p> | | | | |

| | | |
|--|--|---|
| 35 | In which of the following pairs will compound I and compound II give the same number of cis-trans isomers after reaction with hot ethanolic potassium hydroxide? | |
| | I | II |
| | 1 |  |
| | 2 |  |
| | 3 |  |
| <p>Answer: A (1, 2 and 3)</p> <p>Pair 1 both give 2 cis-trans isomers. Pair 2 and 3 both give 0 cis-trans isomers.</p> | | |

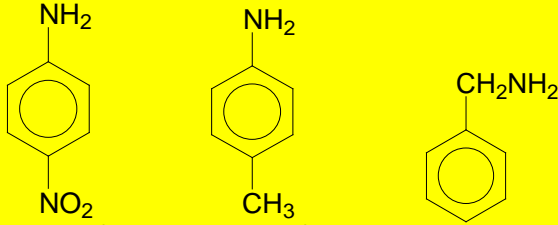
| | | |
|---|---|--|
| 36 | Which of the following processes lead(s) to an increase in entropy? | |
| | 1 | Diffusion of CFCs into the stratosphere. |
| | 2 | Combustion of a piece of charcoal to form CO ₂ (g) and H ₂ O (g). |
| | 3 | Desalination of sea water by reverse osmosis (solvent passes from a more concentrated solution to a more dilute solution). |
| <p>Answer: B (1 and 2 only)</p> <p>For option 3, there is an increase in orderliness as the solvent passes from a more concentrated solution to a more diluted solution. Hence, entropy will decrease.</p> | | |


| | |
|----|--|
| 37 | <p>The diagram shows the structure of salicylic acid:</p> <div data-bbox="758 257 917 414" data-label="Chemical-Block"> </div> <p>salicylic acid</p> <p>Which compound(s) give(s) salicylic acid on acidic hydrolysis?</p> |
| 1 | <div data-bbox="343 560 566 705" data-label="Chemical-Block"> </div> |
| 2 | <div data-bbox="343 750 486 896" data-label="Chemical-Block"> </div> |
| 3 | <div data-bbox="343 940 502 1086" data-label="Chemical-Block"> </div> |
| | <p>Answer: D (1 only)</p> <p>2: Halogenoarenes do not undergo nucleophilic substitution.</p> <div data-bbox="295 1332 494 1478" data-label="Chemical-Block"> </div> <p>3: will be formed instead.</p> |


| | | |
|-----------|---|--|
| 38 | In which of the following sequences does the value of pK_b decrease? | |
| 1 |  | |
| 2 | $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2$, $\text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2$, $\text{CH}_3\text{CH}_2\text{CH}_2\text{NH}_2$ | |
| 3 |  | |

Answer: **B (1 and 2)**

pK_b decreases means weakest base to strongest base or strongest acid to weakest acid.



1: 

- The **electron donating alkyl group increases** the **availability** of the **lone pair of electrons on N atom to accept a proton via a dative bond**. Hence,  is the strongest base.

4-methylphenylamine is a **stronger** base than 4-nitrophenylamine.

- The presence of the **electron-donating methyl group** in 4-methylphenylamine **reduces** the **delocalization** of the **lone pair of electrons on N into benzene ring**.
- This **increases** the **availability** of lone pair of electrons **to accept a proton via dative bond**.

| | |
|--|---|
| | <p>2: $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2$, $\text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2$, $\text{CH}_3\text{CH}_2\text{CH}_2\text{NH}_2$</p> <ul style="list-style-type: none"> The electron donating alkyl group increases the availability of the lone pair of electrons on N atom to accept a proton via a dative bond. Hence, $\text{CH}_3\text{CH}_2\text{CH}_2\text{NH}_2$ is the strongest base. Proximity of withdrawing $-\text{OH}$ substituent to NH_2 group: $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2 > \text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2$ <p>Electron withdrawing effect: $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2 > \text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2$</p> <p>Availability of the lone pair of electrons on N atom to accept a proton via a dative bond: $\text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2 > \text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2$</p> <p>Strength of base: $\text{CH}_2\text{CH}_2\text{CH}_2\text{NH}_2 > \text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{NH}_2$</p> <p>3: The order of the acidity is already incorrect to begin.</p> |
|--|---|

| | |
|----|--|
| 39 | <p><i>Use of the Data Booklet is relevant to this question.</i></p> <p>A student set up an electrolytic cell for the purpose of purifying copper metal. However, the set-up did not lead to successful purification of copper.</p> <p>Which of the following could explain the failure of the set-up?</p> |
| 1 | The electrolyte used was aqueous chromium(III) chloride. |
| 2 | The impure copper was used as the cathode and the anode was made of pure copper metal. |
| 3 | The E°_{cell} for the reaction is a negative value. |
| | <p>Answer: B (1 and 2 only)</p> <p>1: Chromium metal would be deposited on the cathode before copper metal, hence leading to the failure of the purification of copper.</p> <p>2: Impure copper should be the anode and the cathode should be pure copper metal. When the polarities are switched, the pure copper anode would be oxidised and copper metal would be deposited on the impure copper cathode, leading to the failure of the purification of copper.</p> <p>3: Even if E°_{cell} is a negative value, the reaction can still occur because electricity is the source of energy for a chemical reaction to take place in an electrolytic cell.</p> |

| | | |
|--|--|-------------|
| 40 | Vanadium has the electronic structure $1s^2 2s^2 2p^6 3s^2 3p^6 3d^3 4s^2$. Which of the following vanadium compounds is/are likely to exist? | |
| | 1 | V_2O_5 |
| | 2 | $VOCl$ |
| | 3 | $K_2V_2O_7$ |
| <p>Answer: B (1 and 2 only)</p> <p>Oxidation of Vanadium in $V_2O_7^{2-}$ is expected to be 6 which is not possible.</p> | | |