



**Catholic Junior College**  
**JC 2 Preliminary Examinations**  
**Higher 2**

CANDIDATE  
NAME

CLASS

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**CHEMISTRY**

Paper 1 Multiple Choice

**9729/01**

**15 September 2022**  
**1 hour**

Additional Materials: Multiple Choice Answer Sheet  
Data Booklet

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**READ THESE INSTRUCTIONS FIRST**

Write in soft pencil.

Do not use staples, paper clips, glue or correction fluid.

Write your name, class and NRIC/FIN number on the Answer Sheet in the spaces provided.

There are **thirty** questions on this paper. Answer **all** questions. For each question there are four possible answers **A, B, C** and **D**.

Choose the **one** you consider correct and record your choice in **soft pencil** on the separate Answer Sheet.

**Read the instructions on the Answer Sheet very carefully.**

Each correct answer will score one mark. A mark will not be deducted for a wrong answer.

Any rough working should be done in this booklet.

The use of an approved scientific calculator is expected, where appropriate.

**WORKED**  
**SOLUTIONS**

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This document consists of **31** printed pages.

1 Which statement about 27 g of Al is always correct?

- A It contains the same number of atoms as  $\frac{1}{12}$  g of  $^{12}\text{C}$ .  
**B It contains the same number of atoms as 24 dm<sup>3</sup> of krypton gas at room temperature and pressure.**  
 C It contains the same number of hydrogen ions as 1 dm<sup>3</sup> of 1 mol dm<sup>-3</sup> aqueous sulfuric acid.  
 D It contains the same number of atoms as 28 g of nitrogen gas.

**Topic: Mole Concept**

No of Al atoms in 27 g =  $\frac{27}{27.0} \times 6.02 \times 10^{23} = \underline{6.02 \times 10^{23}}$

A No of C atoms in  $\frac{1}{12.0}$  g of  $^{12}\text{C} = \frac{1}{12.0} \times 6.02 \times 10^{23} = 4.18 \times 10^{21}$

**B No of Kr atoms in 24 dm<sup>3</sup> of Kr =  $\frac{24}{24.0} \times 6.02 \times 10^{23} = 6.02 \times 10^{23}$**

C  $\text{H}_2\text{SO}_4 \equiv 2\text{H}^+$

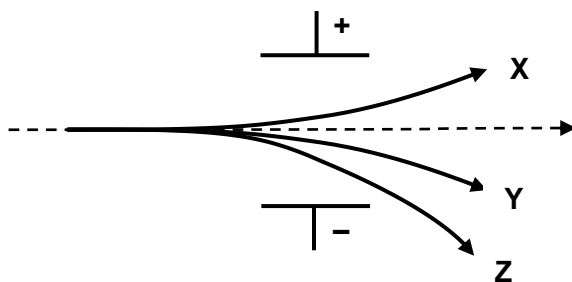
No of  $\text{H}^+$  ions in 1 dm<sup>3</sup> of aq  $\text{H}_2\text{SO}_4 = 1 \times 1 \times 2 \times 6.02 \times 10^{23}$   
 $= 1.20 \times 10^{24}$

D  $\text{N}_2 \equiv 2\text{N}$

No of N atoms in 28 g of  $\text{N}_2 = \frac{28}{28.0} \times 2 \times 6.02 \times 10^{23} = 1.20 \times 10^{24}$

**Answer: B**

2 The following are flight paths of charged particles as they pass through an electric field at the same speed.



Which of the following correctly identifies X, Y and Z?

	X	Y	Z
A	$^{14}\text{N}^-$	$^{16}\text{O}^{2+}$	$^{28}\text{Si}^{2+}$
<b>B</b>	<b><math>^{14}\text{N}^-</math></b>	<b><math>^{14}\text{C}^+</math></b>	<b><math>^{28}\text{Si}^{4+}</math></b>
C	$^{15}\text{O}^+$	$^{14}\text{C}^+$	$^{14}\text{N}^+$
D	$^{15}\text{O}^-$	$^{14}\text{C}^+$	$^{28}\text{Si}^+$

**Concept: Angle of deflection**

particles	$^{14}\text{N}^-$	$^{14}\text{C}^+$	$^{28}\text{Si}^{4+}$
m/z	1/14	1/14	4/28 = 1/7

**angle of deflection**  $\propto \frac{\text{charge}}{\text{mass}}$  of particle

$^{14}\text{N}^-$  is negatively charged, so it is attracted to anode while  $^{14}\text{C}^+$  and  $^{28}\text{Si}^{4+}$  are attracted to cathode. Since angle of deflection is charge/mass ratio,  $^{14}\text{N}^-$  and  $^{14}\text{C}^+$  have roughly the same angle of deflection (paths X and Y), while  $^{28}\text{Si}^{4+}$  has almost double the angle of deflection (path Z).

**Answer: B**

3 *Use of the Data Booklet is relevant to this question.*

Species containing one or more unpaired electrons can be attracted by an external magnetic field and are said to be paramagnetic.

Which of the following species is paramagnetic?

- 1  $\text{Cr}^{3+}$
- 2  $\text{Cu}^+$
- 3  $\text{Ni}^{2+}$

**A** 3

**B** 1 and 2

**C** 1 and 3

**D** 1, 2 and 3

**Concept: Electronic Configuration**

$\text{Cr}^{3+}$ :  $[\text{Ar}]3\text{d}^3$

$\text{Cu}^+$ :  $[\text{Ar}]3\text{d}^{10}$  (Does not have any unpaired electrons)

$\text{Ni}^{2+}$ :  $[\text{Ar}]3\text{d}^8$

**Answer: C**

- 4 In which of the following sequences are the species quoted in order of decreasing boiling points?

- A RbCl, KCl  
**B HF, HCl**  
 C K, Ca  
 D CO, CO<sub>2</sub>

**Concept: Boiling points**

**Option A:** Both of them have giant ionic lattice structure with strong electrostatic forces of attractions between cations and anions. The strength of ionic bonding  $\propto \frac{q_+ \cdot q_-}{r_+ + r_-}$

Rb<sup>+</sup> has larger cationic size than K<sup>+</sup>, hence RbCl has weaker ionic bonding than KCl.

**Option B:** HF has a higher boiling point than HCl as it has stronger intermolecular hydrogen bonding while HCl has weaker permanent dipole-permanent dipole between its molecules.

**Option C:** Both K and Ca have giant metallic lattice structure with strong electrostatic forces of attractions between cations and sea of delocalised electrons. The strength of metallic bonding  $\propto$  the number of valence electrons involved in the delocalisation.

K: 4s<sup>1</sup>

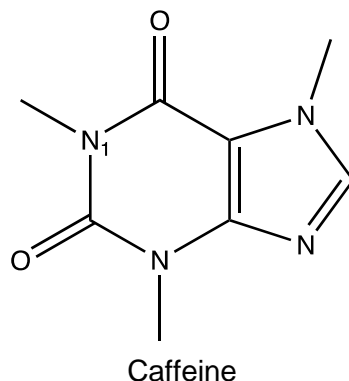
Ca: 4s<sup>2</sup>

Ca has higher boiling point than K because it has more valence electrons than K involved in the delocalisation.

**Option D:** Both of them have simple molecular structure. CO<sub>2</sub> has larger electron cloud size compared to CO. CO<sub>2</sub> has higher boiling point as it has stronger intermolecular instantaneous dipole-induced dipole attractions between its molecules compared to the weaker intermolecular permanent dipole-permanent dipole between CO molecules.

**Answer: B**

- 5 People drink beverages containing caffeine to relieve or prevent drowsiness and to improve cognitive performance.



Which statement about caffeine is true?

- A Caffeine molecule has a planar structure.
- B The  $\pi$  bond in  $C=C$  is formed by sideways overlap of 2p orbitals.**
- C The nitrogen atom in caffeine,  $N_1$  is basic.
- D There are 16 sigma bonds in a caffeine molecule.

**Concept: Chemical Bonding**

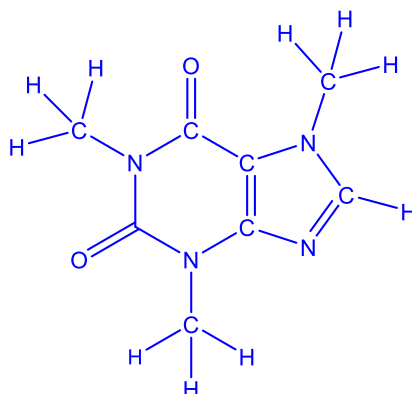
**Option A is wrong. There are  $sp^3$  carbon atoms which have tetrahedral geometry.**

**Option B is correct. The pi bond is formed by sideways overlap of parallel unhybridised p orbitals.**

**Option C is wrong. The lone pair of electrons in the unhybridised p orbital of  $N_1$  has overlapped sideways with the adjacent  $\pi$  electron cloud of  $C=O$  bond resulting in the delocalisation of lone pair of electrons. Hence, lone pair of electrons is unavailable for protonation.**

**Option D is wrong. There are 25  $\sigma$  bonds in a caffeine molecule. There is one  $\sigma$  bond and one  $\pi$  bond in a double bond.**

**Answer: D**



6 Which of the following substances conduct electricity due to delocalised electrons?

- 1 Graphite
- 2 Solid magnesium
- 3 Molten sodium chloride

- A 1 only
- B 1 and 2 only**
- C 2 and 3 only
- D 1, 2 and 3

**Concept: Physical properties of compounds**

**Answer: B**

**Molten sodium chloride conducts electricity due to the presence of free mobile ions.**

7 Which of the following statements is **incorrect**?

- A When methane gas is subjected to low pressure, it liquefies.**
- B Tyre pressure readings are higher on a hot day.
- C The boiling point of water is lower than 100°C at a higher altitude.
- D The density of an ideal gas at constant pressure is inversely proportional to the temperature.

**Concept: Gaseous State**

**A: Gas should be subjected to high pressure for it to be liquified.**

**B:  $P \propto T$ , when tyre is hot, T increases hence pressure is higher.**

**C: At a higher altitude than sea level, the atmospheric pressure is lower than atmospheric pressure at sea level, thus boiling point will be lowered as  $P \propto T$ .**

**D:  $\rho = \frac{PM}{RT}$  density is inversely proportional to temperature.**

**Answer: A**

- 8 For the oxides of Period 3 elements (Na to P), which property decreases from  $\text{Na}_2\text{O}$  to  $\text{P}_4\text{O}_{10}$ ?

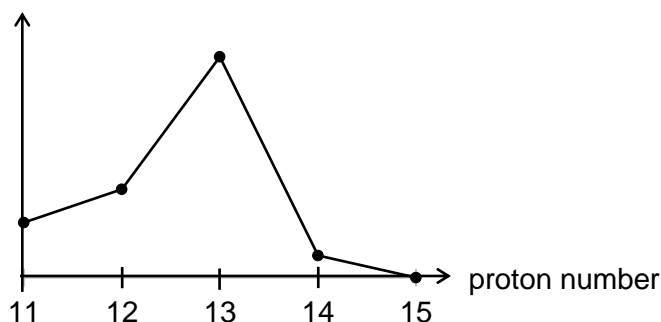
- A melting point  
 B covalent character  
 C pH when mixed with water  
 D solubility in aqueous alkali

Concept: Properties of oxides of Period 3

Answer: C

Oxide formula		$\text{Na}_2\text{O}$	$\text{MgO}$	$\text{Al}_2\text{O}_3$	$\text{SiO}_2$	$\text{P}_4\text{O}_6$ and $\text{P}_4\text{O}_{10}$
Structure		Giant ionic			Giant covalent	Simple covalent (molecular)
Melting Point / °C		1280	2900	2040	1610	24 580
Acid/base behaviour		Basic: $\text{Na}_2\text{O} + 2\text{H}^+ \rightarrow 2\text{Na}^+ + \text{H}_2\text{O}$	Basic: $\text{MgO} + 2\text{H}^+ \rightarrow \text{Mg}^{2+} + \text{H}_2\text{O}$	Amphoteric: $\text{Al}_2\text{O}_3 + 6\text{H}^+ \rightarrow 2\text{Al}^{3+} + 3\text{H}_2\text{O}$  $\text{Al}_2\text{O}_3 + 2\text{OH}^- + 3\text{H}_2\text{O} \rightarrow 2\text{Al}(\text{OH})_4^-$	Acidic: forms $\text{SiO}_3^{2-}$ with base $\text{SiO}_2 + 2\text{NaOH} \rightarrow \text{Na}_2\text{SiO}_3 + \text{H}_2\text{O}$ $\text{SiO}_2 + \text{CaO} \rightarrow \text{CaSiO}_3$	Acidic $\text{P}_4\text{O}_6 + 8\text{NaOH} \rightarrow 4\text{Na}_2\text{HPO}_3 + 2\text{H}_2\text{O}$  $\text{P}_4\text{O}_{10} + 12\text{NaOH} \rightarrow 4\text{Na}_3\text{PO}_4 + 6\text{H}_2\text{O}$
Rxn w/ $\text{H}_2\text{O}$	Eqn	$\text{Na}_2\text{O} + \text{H}_2\text{O} \rightarrow 2\text{NaOH}$	$\text{MgO} + \text{H}_2\text{O} \rightleftharpoons \text{Mg}(\text{OH})_2$ (slightly alkaline as only mildly soluble)	<u>Giant ionic</u> with high lattice energy. Will not dissolve in water.	<u>Giant covalent.</u> Will not dissolve.	$\text{P}_4\text{O}_6 + 6\text{H}_2\text{O} \rightarrow 4\text{H}_3\text{PO}_3$ $\text{P}_4\text{O}_{10} + 6\text{H}_2\text{O} \rightarrow 4\text{H}_3\text{PO}_4$
	pH	13	9	<u>7 (the pH of water)</u>	<u>7 (the pH of water)</u>	2

- 9 The following graph shows how a property of the elements in Period 3, from Na to P, or their compounds, varies with proton number.



What property is shown by the graph?

- A ionisation energies of elements
- B melting point of element
- C pH of aqueous chloride
- D electrical conductivity of element**

**Concept: Trends of different physical properties of Period 3 elements.**

**Electrical conductivity increases from Na to Al due to the increase in number of mobile delocalised electrons. Electrical conductivity then decrease to Si as Si is a semi-conductor and has poor conductivity under normal conditions. P does not conduct electricity as it neither have delocalised electrons nor mobile charge carriers.**

**Answer: D**

- 10 *Use of Data Booklet is relevant to this question.*

The  $\Delta G^\ominus_{\text{solution}}$  and  $\Delta S^\ominus_{\text{solution}}$  for silver chloride, AgCl are  $+55.6 \text{ kJ mol}^{-1}$  and  $+33.2 \text{ J mol}^{-1} \text{ K}^{-1}$  respectively.

What is the standard enthalpy change ( $\Delta H^\ominus$ ) when 287 g of AgCl is precipitated under the same conditions?

- A  $+65.5 \text{ kJ}$
- B  $-65.5 \text{ kJ}$
- C  $+131 \text{ kJ}$
- D  $-131 \text{ kJ}$**

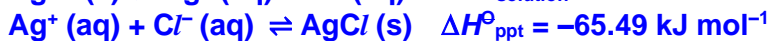


**Concept: Gibbs' Free Energy**

$$\text{Amount of AgCl precipitated} = \frac{287}{107.9 + 35.5} = 2.00 \text{ mol}$$

$$\Delta G = \Delta H - T\Delta S$$

$$55.6 = \Delta H_{\text{solution}}^{\circ} - (298)(0.0332)$$



For 2 mol of AgCl precipitated, standard enthalpy change is -131 kJ.

**Answer: D**

11 The rate equation for a reaction between **A** and **B** is given by: rate =  $k[\text{A}]$

Which of the following statements about the reaction is true?

- 1 **A** is involved in the rate-determining step in the reaction mechanism.
- 2 The rate constant,  $k$ , increases with increasing concentration of **A**.
- 3 A graph of rate against  $[\text{A}]$  gives a straight line that passes through the origin.

**A** 1,2 and 3      **B** 1 and 3 only      **C** 2 only      **D** 1 only

**Concept: Reaction mechanism**

1. True statement. From the rate equation, 1 molecule of A is involved in the rate determining step of the reaction.
2. False. Rate constant,  $k$ , is only affected by temperature and catalyst.
3. True statement. From the rate equation, rate is directly proportional to  $[\text{A}]$  hence is reflected as a straight line passing through the origin.

**Answer: B**

- 12 What is the pH of the final solution when  $V \text{ cm}^3$  of dilute nitric acid of pH 2.0 is mixed with  $V \text{ cm}^3$  of dilute nitric acid of pH 4.0 followed by the addition of  $2V \text{ cm}^3$  of water?

A 2.3      **B 2.6**      C 3.0      D 3.6

**Topic: Chemistry of aqueous solutions**

**Concept: Calculation of pH of acidic solutions**

**Total volume of final solution =  $V + V + 2V = 4V \text{ cm}^3$**

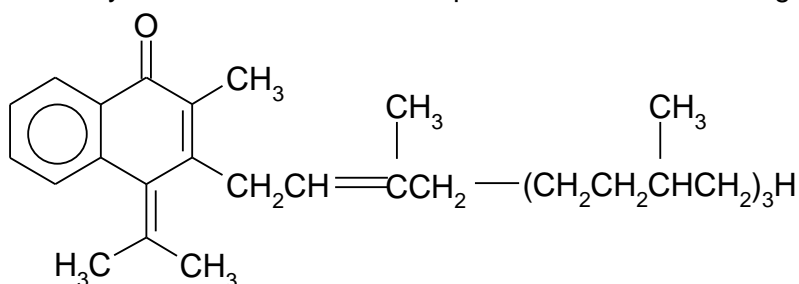
**Total amount of  $\text{H}^+$  ions in final solution =  $(10^{-2} \times \frac{V}{1000}) + (10^{-4} \times \frac{V}{1000}) \text{ mol}$**

$$[\text{H}^+] \text{ in final solution} = \frac{(10^{-2} \times \frac{V}{1000}) + (10^{-4} \times \frac{V}{1000})}{\frac{4V}{1000}} = 0.002525 \text{ mol dm}^{-3}$$

**pH of final solution =  $-\log(0.002525) = 2.6$**

**Answer: B**

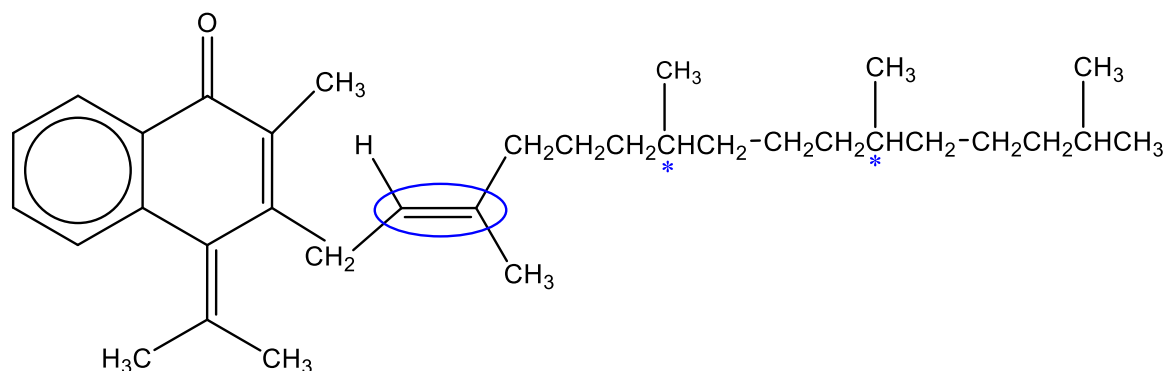
- 13 Compound **K** is a yellow viscous oil found in plants. It has the following structure.



What is the total number of stereoisomers for compound **K**?

A 4      **B 8**      C 16      D 32

**Topic: Stereoisomerism**



There are 2 chiral carbons and 1 carbon-carbon double bond that can display cis-trans isomerism. Therefore  $n = 3$

$$\text{No. of stereoisomers} = 2^3 \\ = 8$$

**Answer: B**

- 14 In the free radical substitution reaction of methane,  $\text{CH}_4$ , one of the side-products formed is ethane,  $\text{CH}_3\text{CH}_3$ , which is formed when two  $\bullet\text{CH}_3$  radicals combine. Upon careful heating, a sample of butane,  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$ , reacted with chlorine gas in a free radical substitution reaction to give only mono-substituted products.

How many possible organic side-products would be obtained in this reaction when the radicals produced combine with each other?

- A 2                      **B 3**                      C 4                      D 5

**Topic: Free Radical substitution**

**Only 2 organic radicals are produced in the mono-substitution of  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3$  :**

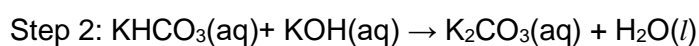


**Therefore, there can be 3 different products:**

**$\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$  ,  $\text{CH}_3\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$ , and  $\text{CH}_3\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_3$ .**

**Answer: B**

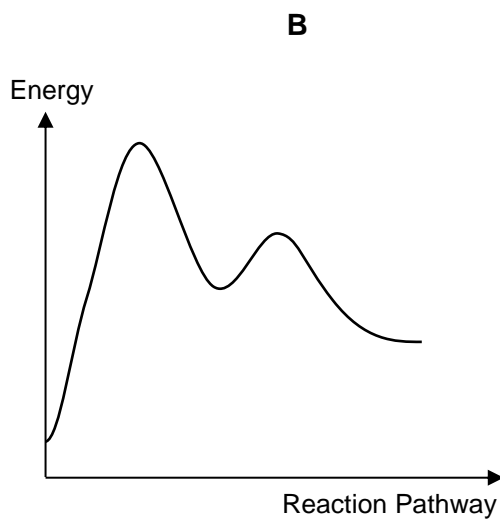
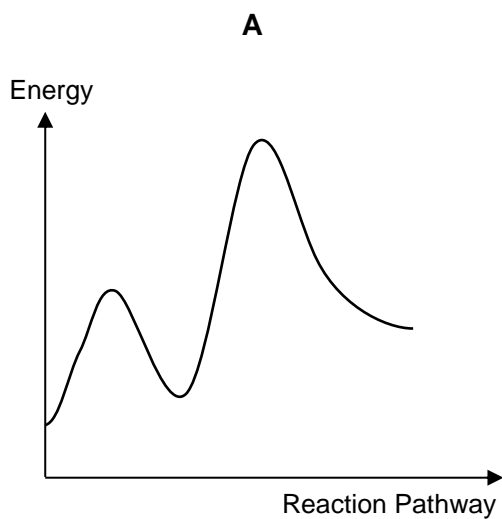
- 15 The reaction between carbon dioxide and potassium hydroxide is exothermic. The proposed two-step mechanism of the reaction is shown below:



Experiments were carried out to study the rate of the reaction above.

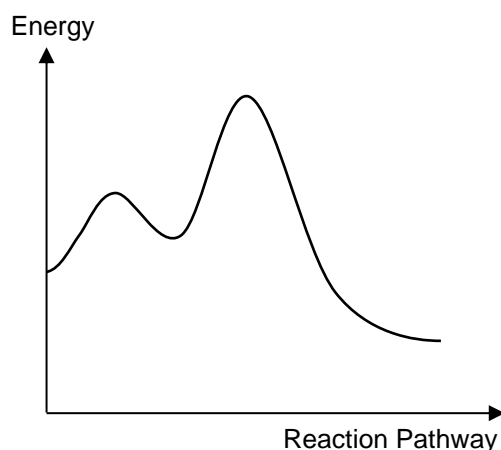
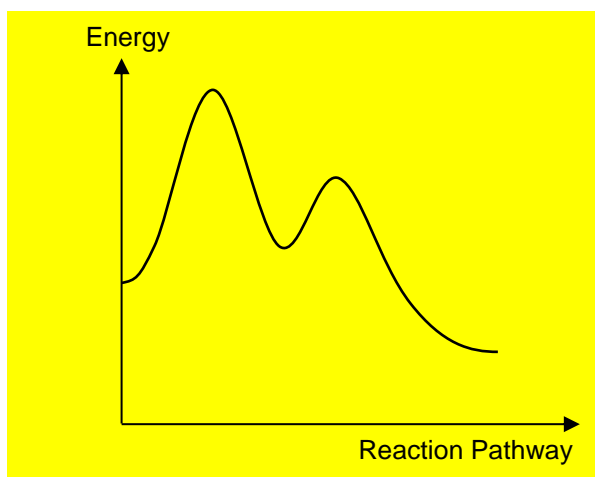
Experiment Number	Initial concentration of $\text{CO}_2$ / $\text{mol dm}^{-3}$	Initial concentration of $\text{KOH}$ / $\text{mol dm}^{-3}$	Initial reaction rate / $\text{mol dm}^{-3} \text{ s}^{-1}$
1	0.2	0.2	0.0034
2	0.4	0.2	0.0068
3	0.2	0.1	0.0017

Which of the following graphs most likely describes the energy profile of the reaction above?



**C**

**D**



**Concept: Kinetics (Reaction mechanism and order of reactions)**

**Answer: C**

Comparing Expts 1 and 2, using the inspection method, by keeping concentrations of KOH constant, while doubling the concentration of  $\text{CO}_2$ , the rate doubles. Hence, the rate of reaction is directly proportional to  $[\text{CO}_2]$ . Therefore the reaction is first order wrt  $\text{CO}_2$ .

Comparing Expts 1 and 3, using the inspection method, by keeping concentrations of  $\text{CO}_2$  constant, while halving the concentration of KOH, the rate is halved. Hence, the rate of reaction is directly proportional to  $[\text{KOH}]$ . Therefore the reaction is first order wrt KOH.

$$\text{Rate} = k[\text{CO}_2][\text{KOH}]$$

Based on the rate equation, it can be inferred that 1 molecule of  $\text{CO}_2$  reacts with 1 molecule of KOH in the rate determining step, i.e. slow step which corresponds to step 1 of the reaction mechanism proposed.

Since the slow step has a high activation energy and is an overall exothermic reaction, therefore the graph reflected corresponds to option C.

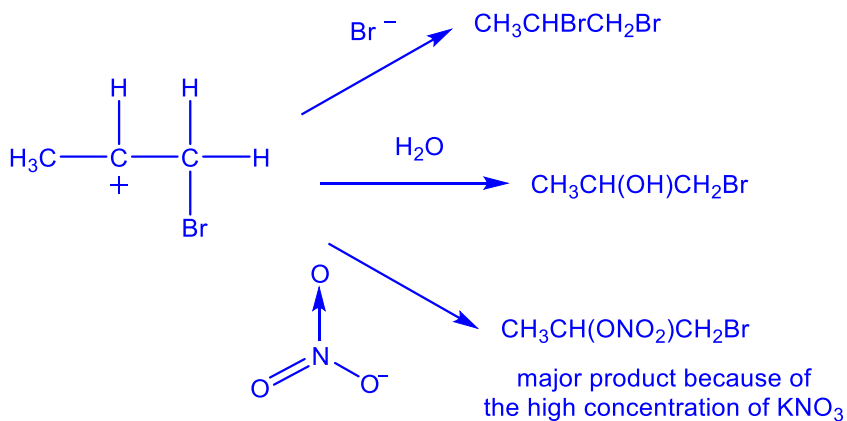
- 16 Aqueous bromine can react with propene in the presence of concentrated potassium nitrate solution.

Which of the following is the major product formed in the reaction?

- A  $\text{CH}_3\text{CHBrCH}_2\text{ONO}_2$
- B  $\text{CH}_3\text{CHBrCH}_2\text{Br}$
- C  $\text{CH}_3\text{CHBrCH}_2\text{OH}$
- D  $\text{CH}_3\text{CH}(\text{ONO}_2)\text{CH}_2\text{Br}$

**Topic: Alkenes**

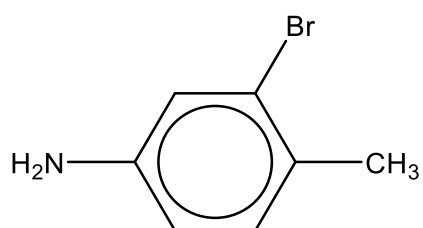
**Concept: Electrophilic addition, Markovnikov's rule**



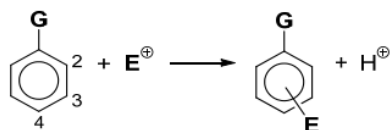
**In the second step of the reaction, the nitrate ion is preferred relative to water owing to its negative charge.**

**Answer: D**

- 17 Which synthetic route is most likely to lead to the most successful synthesis of the following product from benzene?



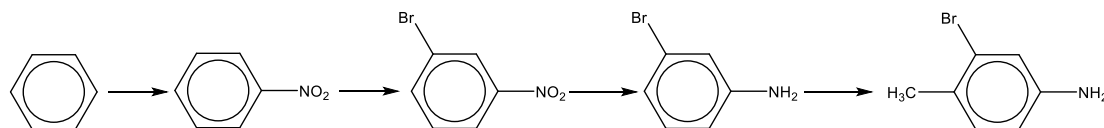
- A nitration, bromination, alkylation, reduction
- B nitration, bromination, reduction, alkylation**
- C nitration, alkylation, reduction, bromination
- D alkylation, bromination, nitration, reduction

**Topic: Arenes****Concept: Orientating effect of groups in aromatic substitution reactions**

G	-alkyl -OH or -OR -NH <sub>2</sub> , -NHR or -NR <sub>2</sub> -NHCOR	-Cl, -Br, -I	-CHO, -COR -CO <sub>2</sub> H, -CO <sub>2</sub> R -NH <sub>3</sub> <sup>+</sup> -NO <sub>2</sub> , -CN
Reactivity of ring (compared to benzene)	Activated	Deactivated	Deactivated
Position of E (relative to position of G)	2- and/or 4-	2- and/or 4-	3-

**Option A results in a contest between the -NO<sub>2</sub> and -Br groups to determine the position of the -CH<sub>3</sub> group**

**Option B will result in the best yield because the -NH<sub>2</sub> and -Br groups are both -2,4-directing and will reinforce each other's activating effects on the incoming -CH<sub>3</sub>.**



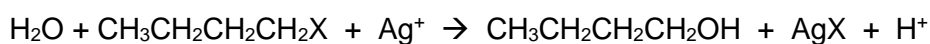
**Option C results in a contest between the -NO<sub>2</sub> and -CH<sub>3</sub> groups to determine the position of the Br group.**

**Option D results in a contest between the -Br and -CH<sub>3</sub> groups to determine the position of the NO<sub>2</sub> group.**

**Answer: B**

- 18** A few drops of 1-chlorobutane, 1-bromobutane, 1-iodobutane were placed separately into three test-tubes each, containing 1.0 cm<sup>3</sup> of aqueous silver nitrate at 60 °C.

A hydrolysis reaction occurred. (X is the halogen atom)



Which of the following would be the best explanation for the rate of the reaction?

- A** The bond energy of C-X bond decreases from C-Cl to C-I.  
**B** The bond polarity of C-X bond decreases from C-Cl to C-I.  
**C** The electron deficiency of the carbon atom bonded to X decreases from C-Cl to C-I.  
**D** The solubility of AgX decreases from AgCl to AgI.



**Topic: Halogen Derivatives**

**Concept: Hydrolysis in RX**

**Answer: A**

**Bond energy of C–X (kJ mol<sup>-1</sup>) :** C–Cl (340) > C–Br (280) > C–I (240).

The rate of hydrolysis increases from CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Cl to CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>I. The rate of hydrolysis is determined by the bond energy of C–X bond.

The bond energy of C–X bond decreases from C–Cl to C–I.

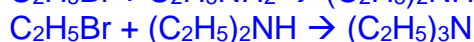
Ease to break the C–X bond increases from C–Cl to C–I.

- 19 Concentrated ammonia was heated in a sealed tube with excess bromoethane. Which of the following product will **not** be formed?

- A** C<sub>4</sub>H<sub>10</sub>N  
**B** C<sub>4</sub>H<sub>11</sub>N  
**C** C<sub>6</sub>H<sub>15</sub>N  
**D** C<sub>8</sub>H<sub>20</sub>NBr

**Topic: Halogen derivatives**

**Concept: Alkylation of nitrogen compounds**



**Answer: A**

- 20 An alcohol **A** with molecular formula C<sub>4</sub>H<sub>10</sub>O is oxidised by acidified potassium dichromate(VI) under certain conditions to give **B**. The following shows some properties of **B**:

- 1 **B** does not produce a yellow precipitate with aqueous alkaline iodine.
- 2 **B** gives a brick – red precipitate when reacted with Fehling’s solution.

How many isomers of alcohol **A** could result in the observations for **B**?

- A** 1                      **B** 2                      **C** 3                      **D** 4

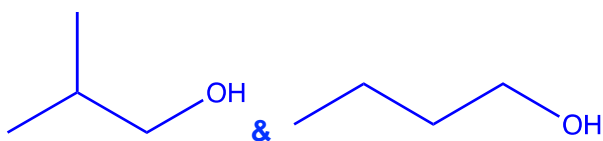
**Concept: Reactions of alcohols**

**Answer: B**

Since product B does not produce a yellow precipitate, it can be concluded that B does not contain a secondary alcohol with a methyl group.

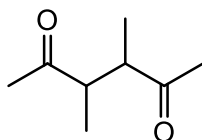
Since product B gives a brick-red precipitate when reacted with Fehling's solution, an aliphatic aldehyde functional group is present.

Since aldehydes are formed from controlled oxidation of primary alcohol, the possible structures of primary alcohol from  $C_4H_{10}O$  are:



Therefore, there are 2 isomers.

- 21 Compound **Z** shown below is an intermediate used to generate pyrroles which are essential to the production of many different chemicals in the pharmaceutical industry.



compound **Z**

Which sentence is correct for compound **Z**?

- A It produces a silver mirror with Tollens' reagent.
- B It decolourises acidified potassium manganate(VII).
- C It produces a yellow precipitate with aqueous alkaline iodine.**
- D It does not produce an orange precipitate with 2,4-dinitrophenylhydrazine.

**Concept: reactions and distinguishing tests relating to carbonyl compounds**

**Compound Z is a diketone.**

**Option A: Incorrect since there is no aldehyde functional group in compound Z and thus cannot give a silver mirror with Tollens' reagent.**

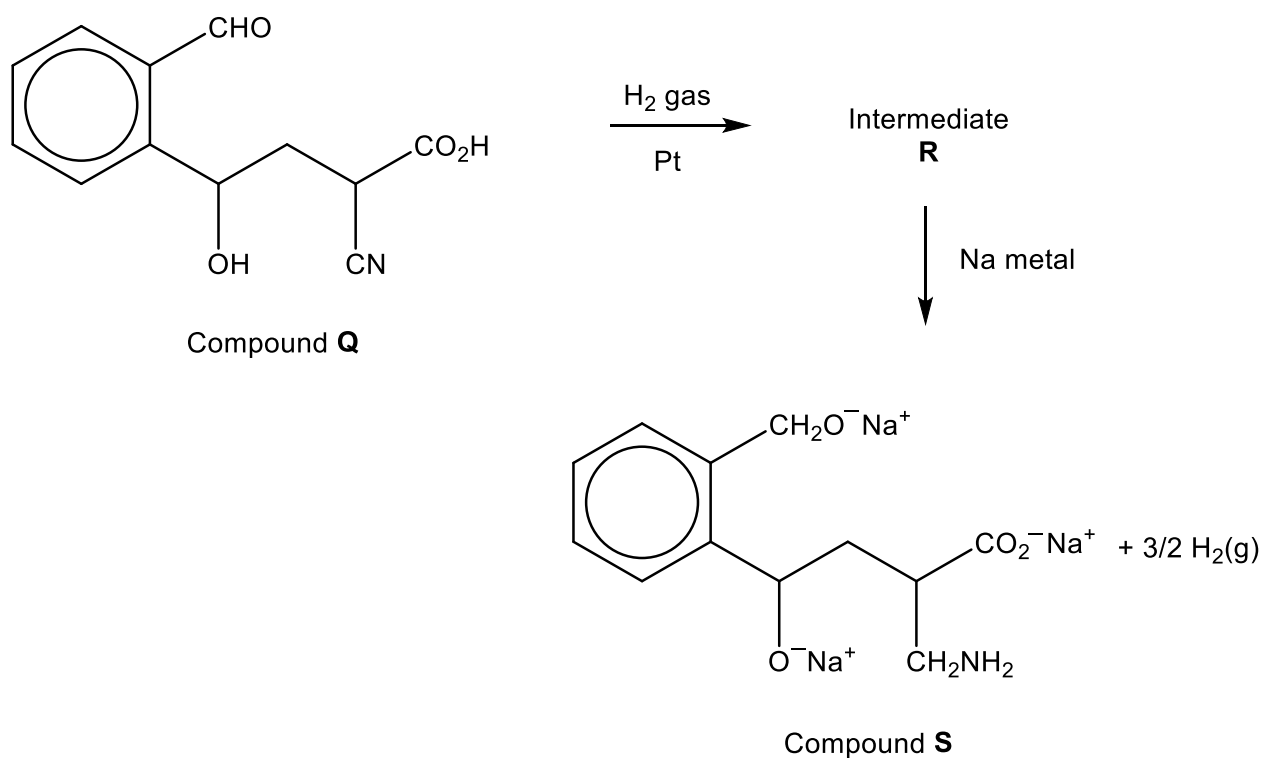
**Option B: Incorrect since ketone cannot undergo further oxidation.**

**Option C: Correct since it has  $\text{—}\overset{\text{O}}{\parallel}{\text{C}}\text{—CH}_3$  group and thus able to give a yellow precipitate with alkaline aqueous iodine.**

**Option D: Incorrect as ketone can undergo condensation with 2,4-DNPH to give an orange precipitate.**

**Answer: C**

- 22** Compound **S** can be obtained via the following 2-step synthesis from compound **Q**.

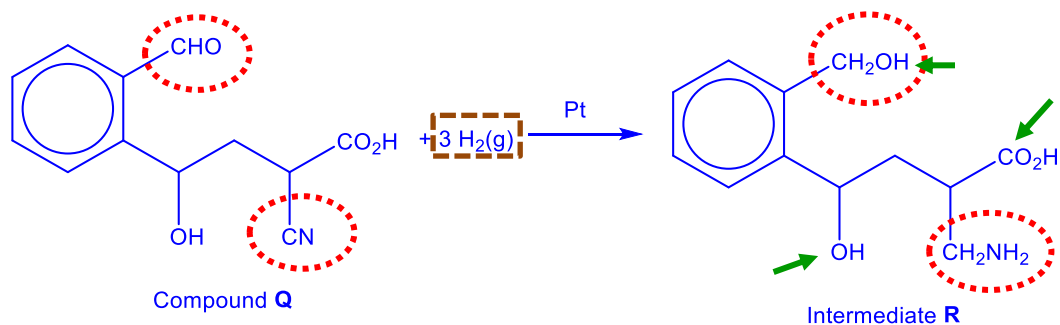


If 96 dm<sup>3</sup> of hydrogen gas was reacted with **one mole** of compound **Q**, followed by the addition of sodium metal at room temperature and pressure, what is the final gas volume when the reaction was completed? (Given  $V_m = 24 \text{ dm}^3 \text{ mol}^{-1}$  at r.t.p.)

- A** Gas volume decreases by 36 dm<sup>3</sup>.
- B** Gas volume decreases by 60 dm<sup>3</sup>.
- C** Gas volume increases by 36 dm<sup>3</sup>.
- D** No change in gas volume.

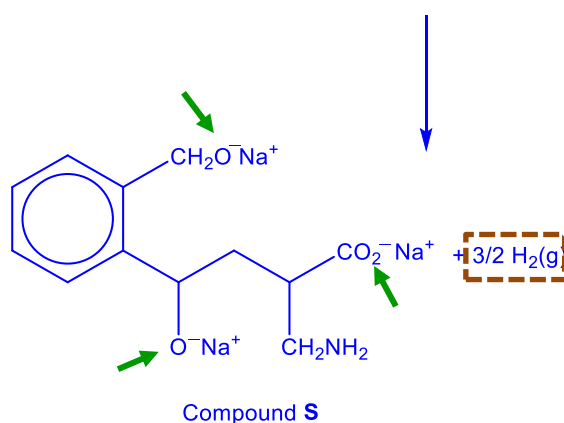
**Topic: Carbonyl Compounds, Alcohols and Carboxylic Acids**

**Concept: Reduction of carbonyl compounds and nitriles, redox reaction of –OH group**



$\text{H}_2$  gas with Pt will only reduce alkenes, carbonyl compounds and nitriles (note:  $\circ\circ\circ$ )

Na metal will react with the –OH group in alcohols and carboxylic acids. (note:  $\rightarrow$ )

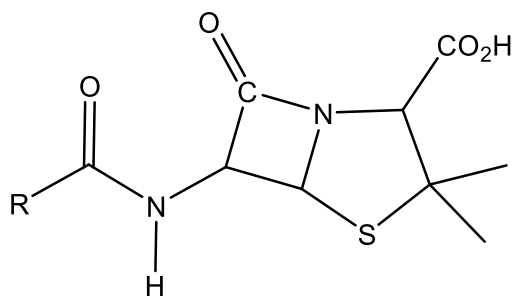


3 mol of  $\text{H}_2$  gas was used up for reduction while 1.5 mol of  $\text{H}_2$  gas was produced from the redox reaction, hence more  $\text{H}_2$  gas was consumed and there will be a decrease in gas volume.

Change in gas volume =  $1.5 \times 24 = 36 \text{ dm}^3$

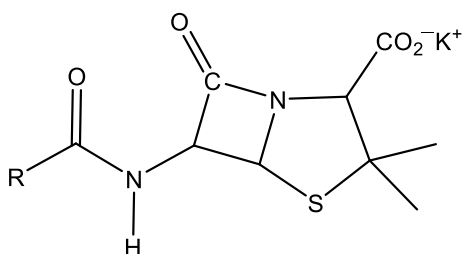
**Answer: A**

- 23** Penicillin is an antibiotic commonly used to treat a number of bacterial infections. The general structure of a penicillin molecule is given below.

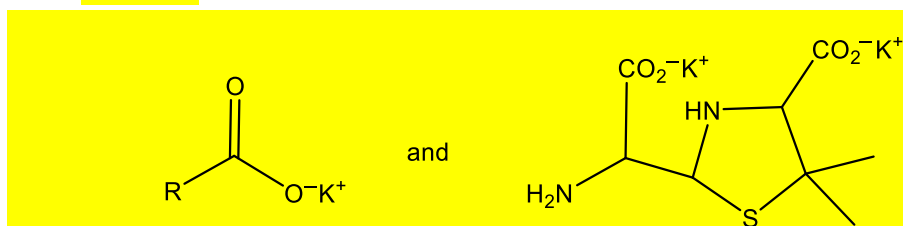


What are the products formed when penicillin is boiled with excess aqueous potassium hydroxide?

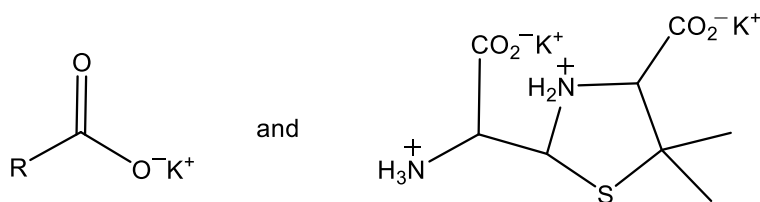
**A**



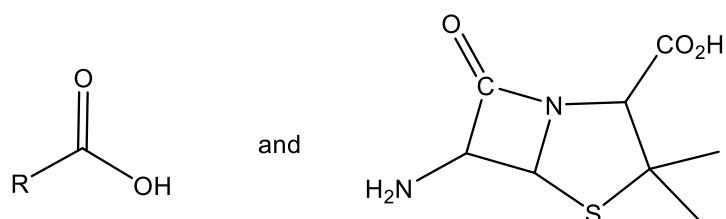
**B**



**C**

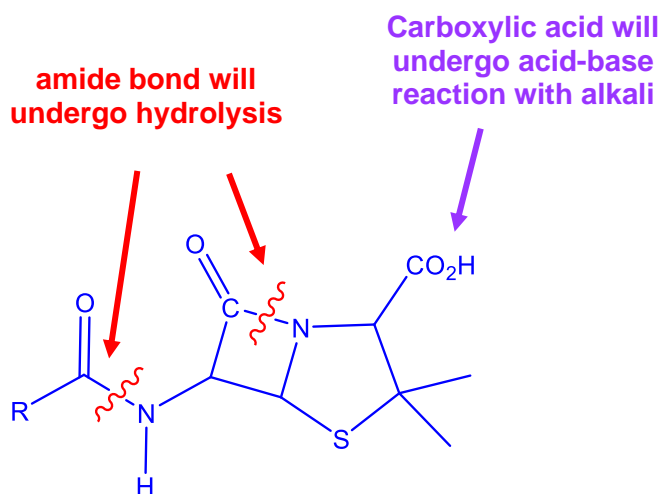


**D**

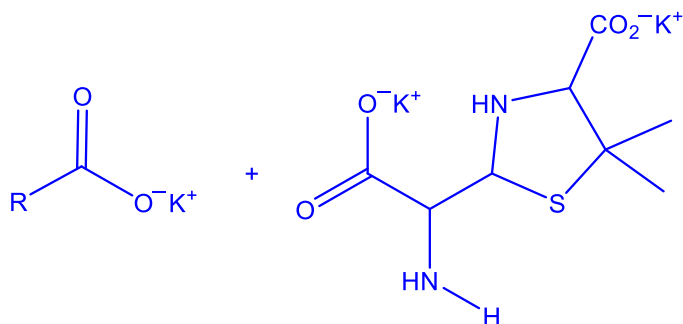


Topic: Nitrogen Compounds

Concept: Alkaline hydrolysis of amides

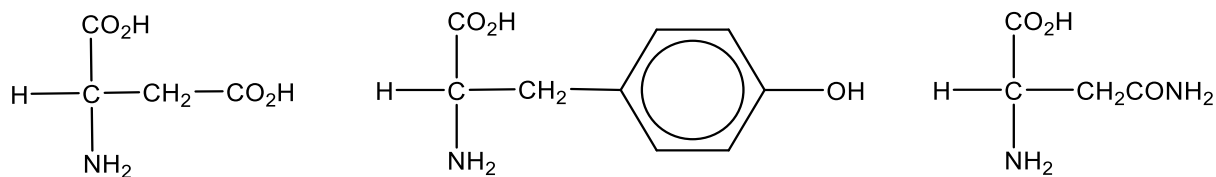


Hence, the hydrolysed products are:



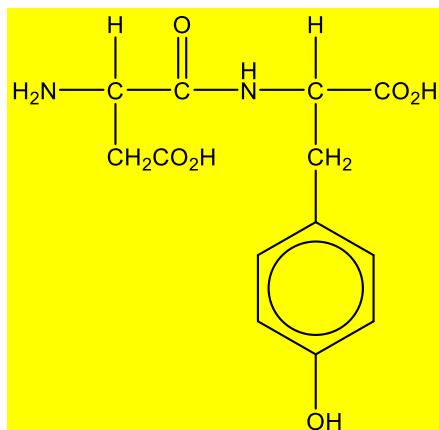
Answer: B

24 The following shows the structures of three amino acids.

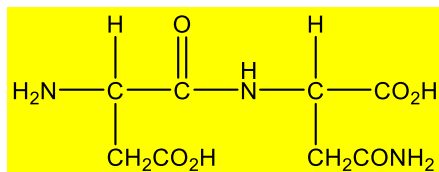


Which of the following represent the dipeptides formed from these amino acids?

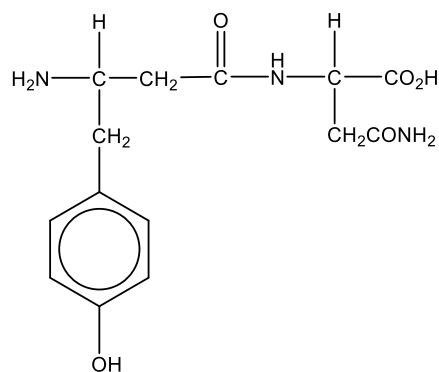
1



2



3



- A 1, 2 and 3  
 B 1 and 2  
 C 1 and 3  
 D 2 only

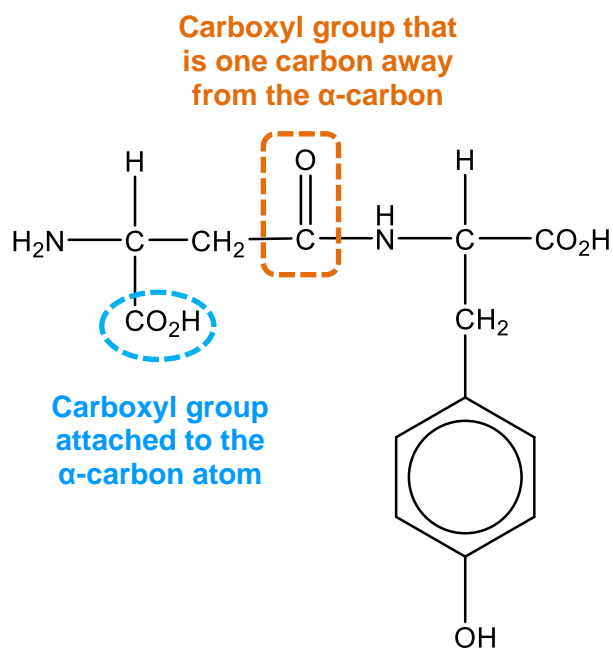


**Topic: Nitrogen Compounds**

**Concept: Formation of peptide bonds**

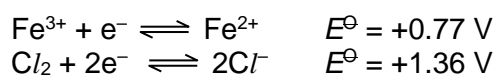
Peptide bonds are formed between the carboxyl group ( $-\text{CO}_2\text{H}$ ) of one amino acid with the amino ( $-\text{NH}_2$ ) group of another amino acid. Both the carboxyl and amino groups are attached to the  $\alpha$ -carbon atoms in the amino acids.

Option 3 is wrong because the carboxyl group in the first amino acid is not attached to the  $\alpha$ -carbon atom.



**Answer: B**

- 25 Two electrode potentials are given.



Which species is the strongest reducing agent?

- A  $\text{Fe}^{3+}$       B  $\text{Fe}^{2+}$       C  $\text{Cl}_2$       D  $\text{Cl}^{-}$

**Topic: Electrochemistry**

**Concept: Using  $E^{\ominus}$  to determine strength of oxidising power.**

**A reducing agent reduces others and it gets oxidised (donates electrons).**

**The only 2 species undergoing oxidation are  $\text{Fe}^{2+}$  and  $\text{Cl}^{-}$ .**

**For the species ions to donate electrons, it must be more readily oxidised, i.e. less positive  $E^{\ominus}$  value in the data booklet. Thus the only possible answer is B.**

**Answer: B**

- 26 Use of the Data Booklet is relevant to this question.

An electrochemical cell is set up using a  $\text{Fe}^{2+}(\text{aq})|\text{Fe}(\text{s})$  half-cell and a  $\text{VO}_2^{+}(\text{aq}), \text{VO}^{2+}(\text{aq})|\text{Pt}(\text{s})$  half-cell.

Which of the following gives a correct effect on the  $E_{\text{cell}}$  and a correct explanation for the effect when each of the changes is made to the cell separately?

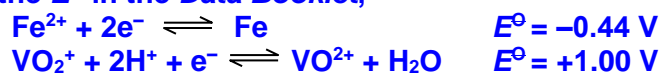
	change	effect on $E_{\text{cell}}$	explanation
1	add KCN(aq) to the $\text{Fe}^{2+}(\text{aq}) \text{Fe}(\text{s})$ half-cell	increases	concentration of $\text{Fe}^{2+}(\text{aq})$ decreases
2	add water to the $\text{VO}_2^{+}(\text{aq}), \text{VO}^{2+}(\text{aq}) \text{Pt}(\text{s})$ half-cell	decreases	concentration of water increases
3	increase temperature of the $\text{Fe}^{2+}(\text{aq}) \text{Fe}(\text{s})$ half-cell	no change	temperature change does not affect $E_{\text{cell}}$

- A 1, 2 and 3      B 1 and 2      C 2 and 3      D 1 only

Topic: Electrochemistry

Concept: Finding value of  $E_{\text{cell}}$  under non standard conditions

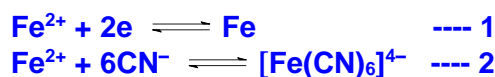
From the  $E^\ominus$  in the *Data Booklet*,



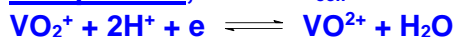
$\text{Fe}^{2+}/\text{Fe}$  is the anode ( $-0.44\text{V}$ ) while  $\text{VO}_2^+/\text{VO}^{2+}$  is the cathode ( $+1.00\text{V}$ ).

$$\begin{aligned} E_{\text{cell}}^\ominus &= E_{\text{red}}^\ominus - E_{\text{oxid}}^\ominus \\ &= (+1.00) - (-0.44) \\ &= +1.44 \text{ V} \end{aligned}$$

- 1: Adding KCN to  $\text{Fe}^{2+}/\text{Fe}$  half-cell will result in formation of  $[\text{Fe}(\text{CN})_6]^{4-}$  and hence cause  $[\text{Fe}^{2+}]$  to decrease.  $E_{\text{Fe}^{2+}/\text{Fe}}$  will become more negative as its p.o.e. in equation 1 shifts to the left, hence  $E_{\text{cell}}$  will become more positive (increases).



- 2: Adding water to  $\text{VO}_2^+/\text{VO}^{2+}$  half-cell will decrease both  $[\text{VO}_2^+]$  and  $[\text{VO}^{2+}]$ . Its p.o.e. shifts to the left as there are more ions on the left.  $E_{\text{VO}_2^+/\text{VO}^{2+}}$  becomes less positive, hence  $E_{\text{cell}}$  will become less positive (decreases).



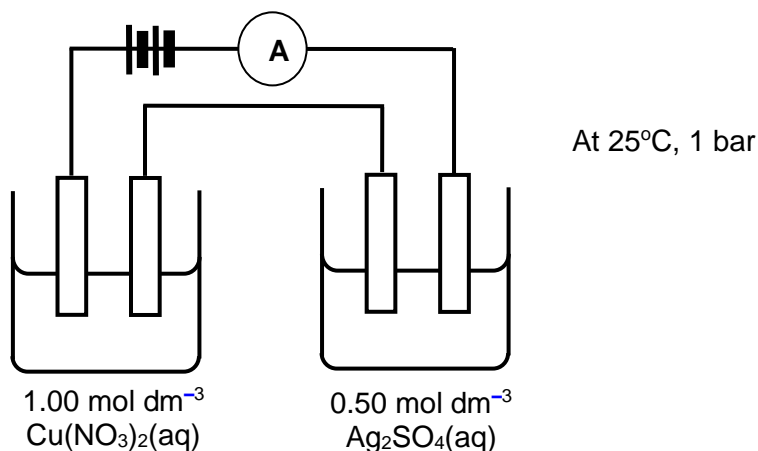
OR concentration of water remains unchanged.

- 3:  $E_{\text{cell}}$  changes with temperature unless  $\Delta H$  of half-cell = 0

Answer: D

27 Use of the Data Booklet is relevant to this question.

Using inert electrodes, a current was passed through two beakers containing aqueous silver sulfate and aqueous copper(II) nitrate, connected in series under standard conditions.



What is the ratio of the mass of silver to copper deposited after the current was passed for  $t$  minutes?

- |          |      |          |      |
|----------|------|----------|------|
| <b>A</b> | 0.59 | <b>B</b> | 0.85 |
| <b>C</b> | 1.70 | <b>D</b> | 3.40 |

**Topic: Electrochemistry**

**Concept: Electrolysis**

**Relevant mole ratio: Cu  $\equiv$  2e (from Cu<sup>2+</sup>) and Ag  $\equiv$  e (from Ag<sup>+</sup>)**

**Since same current pass through in  $t$  minutes, same amount of electrons passed (notice this is INDEPENDENT of the initial concentrations of the solutions) to find mole ratio between Cu and Ag formed,  
Cu  $\equiv$  2e and 2Ag  $\equiv$  2e so Cu  $\equiv$  2Ag**

**Ratio of mass of Ag to mass of Cu = 2 x 107.9 : 63.5 = 3.40**

**Answer: D**

- 28 A current of 10 A is passed for 150 minutes through molten aluminium oxide using inert electrodes.

What will be the approximate volume of gas liberated, measured at s.t.p.?

- A 0.089 dm<sup>3</sup>    **B 5.3 dm<sup>3</sup>**    C 5.6 dm<sup>3</sup>    D 11.2 dm<sup>3</sup>

**Topic: Electrochemistry**

**Concept: Electrolysis**

**Using  $Q = It$ ,**

$$Q = 10 \times 150 \times 60$$

$$= 90\,000\text{C}$$

$$\text{No. of Faradays} = \frac{90\,000}{9.65 \times 10^4} = 0.9326\text{ F}$$

**In molten aluminium oxide, no water is present, so the oxygen gas liberated is from  $\text{Al}_2\text{O}_3$ . The oxide ion in  $\text{Al}_2\text{O}_3$  is  $\text{O}^{2-}$ .**



**Since  $\text{O}_2 \equiv 4\text{e}^-$ ,**

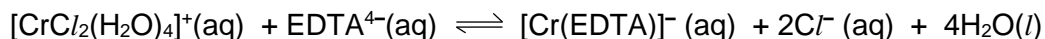
**4 F liberates 1 mol of  $\text{O}_2$  gas**

**0.9326 F liberates 0.2332 mol of  $\text{O}_2$  gas**

$$\text{Vol of } \text{O}_2 \text{ gas} = 0.2332 \times 22.7 \text{ dm}^3 = 5.29 \text{ dm}^3$$

**Answer: B**

- 29  $\text{EDTA}^{4-}(\text{aq})$  solution is added dropwise until in excess to a solution of  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$ . The equilibrium constant for this reaction is greater than 1 and the equation for the reaction is as shown below.



Which one of the following statements about the above reaction is correct?

- A There is no change in colour of the solution after addition of  $\text{EDTA}^{4-}$ .  
 B  $[\text{Cr}(\text{EDTA})]^{-}$  is a less stable complex ion than  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$ .  
**C Both  $[\text{Cr}(\text{EDTA})]^{-}$  and  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$  are octahedral complexes.**  
 D The above is a redox reaction.

**Topic: Transition Metals**

**Option A is incorrect:**

The colour of a transition metal complex depends on the energy gap,  $\Delta E$ . The energy gap,  $\Delta E$ , in turn depends on the nature of the ligands.

Different ligands split the energy level of d orbitals to different extent.

⇒ Amount of energy,  $\Delta E$ , absorbed by d electron in d-d transition differ.

⇒ Colour of transition metal complex solution would change with different ligands used.

⇒ There would be color of the solution will lighten after adding  $\text{EDTA}^{4-}$ .

**Option B is incorrect:**

It is given that the equilibrium constant for the above reaction is more than 1,

⇒ Concentration of  $[\text{Cr}(\text{EDTA})]^- >$  concentration of  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$

⇒  $[\text{Cr}(\text{EDTA})]^-$  is more stable than  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$ .

**Option C is correct:**

For  $[\text{CrCl}_2(\text{H}_2\text{O})_4]^+$ , the central  $\text{Cr}^{3+}$  ion has 6 bond pairs. Hence, it is an octahedral complex.

$\text{EDTA}^{4-}$  is a hexadentate ligand that forms only six dative bonds with the central metal cation. Hence, the central  $\text{Cr}^{3+}$  ion in  $[\text{Cr}(\text{EDTA})]^-$  has 6 bond pairs. It is an octahedral complex.

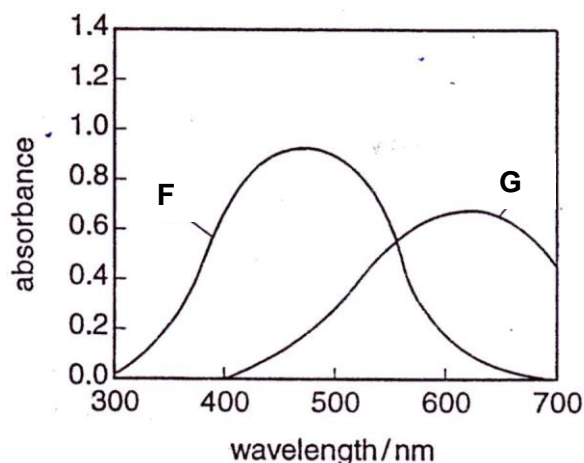
**Option D is incorrect:**

The above reaction does not involve any change in the oxidation no of the elements involved. Hence, it is not a redox reaction. Instead, it is a ligand exchange reaction.

**Answer: C**

- 30** The absorbance of a solution at a particular wavelength is proportional to the concentration of ion responsible for the absorption.

The visible spectra of solutions of two transition metal complexes **F** and **G** are shown in the diagram below. Both complexes contain the same transition metal ion.



Given that energy is inversely proportional to wavelength and the visible region of the electromagnetic spectrum is as follows:

violet	blue	green	yellow	orange	red
400		500		600	700
Wavelength (nm)					

Which of following statements can be deduced from the spectra?

- 1 Complex **F** is likely to be red while complex **G** is likely to be blue.
  - 2 The energy gap in complex **F** is greater than that in complex **G**.
  - 3 The  $K_c$  value for the formation of complex **F** is higher than the  $K_c$  value for formation of complex **G**.
- A** 1, 2 and 3 are correct
- B** 1 and 2 only are correct
- C** 2 and 3 only are correct
- D** 1 only is correct

**Topic: Transition Metals**

Option 1 is correct as Complex F absorbed light of lower wavelength (green) and the complementary colour (red) is observed. Complex G absorbed light of higher wavelength (orange) and the complementary colour (blue) is observed.

Option 2 is correct as Complex F absorbed light of lower wavelength and thus  $\Delta E$  is larger (since  $\Delta E \propto 1/\lambda$ ) than that of Complex G. This means that the ligand in F is a stronger field ligand than that in complex G.

Option 3 is wrong as we cannot determine  $K_c$  value base on individual absorption curve. We can determine the magnitude of  $K_c$  value only if we are looking at a mixture absorption curve whereby there is equal amount of both ligands.

**Ans: B**