## Section I <br> Total Marks (75)

## Part A

Total marks (15)
Attempt Questions 1-15
Allow about 30 minutes for this part

Use the multiple choice answer sheet.
Select the alternative A, B, C or D that best answers the question. Fill in the response square completely.

Sample 2+4=(A) $2 \quad$ (B) $6 \quad$ (C) $8 \quad$ (D) 9
A Å
B $\zeta$
C Å
D $\AA$

If you think you have made a mistake, put a cross through the incorrect answer and fill in the new answer.
$A \zeta$
B $\zeta$
C Å
D Å

If you change your mind and have crossed out what you consider to be the correct answer, then indicate this by writing the word correct and drawing an arrow as follows:
correct
$A \zeta$
B $\zeta$
C Å
D $\AA$

## SECTION I <br> Part A <br> Multiple Choice

1. The extraction of copper from copper(I) sulfide produces sulfur dioxide as a by-product according to the equation:

$$
\mathrm{Cu}_{2} \mathrm{~S}(s)+\mathrm{O}_{2}(g) \rightarrow 2 \mathrm{Cu}(s)+\mathrm{SO}_{2}(g)
$$

What volume of sulfur dioxide gas will be released at $25^{\circ} \mathrm{C} \& 101.3 \mathrm{kPa}$ when 2.2 g of copper (I) sulfide is reacted?
(A) 338 mL
(B) 563 mL
(C) 10 mL
(D) 515 mL
2. Which acid below is not naturally occurring?
(A) 2-hydroxypropane-1,2,3-tricarboxylic acid
(B) HCl
(C) HBr
(D) $\mathrm{CH}_{3} \mathrm{COOH}$
3. $\mathrm{H}_{2} \mathrm{PO}_{4}^{-}$is an amphiprotic species.

Which of the following represents the conjugate acid and conjugate base respectively of $\mathrm{H}_{2} \mathrm{PO}_{4}{ }^{-}$?
(A) $\mathrm{H}_{3} \mathrm{PO}_{4}$ and $\mathrm{HPO}_{4}{ }^{2-}$
(B) $\mathrm{PO}_{4}{ }^{3-}$ and $\mathrm{H}_{3} \mathrm{PO}_{4}$
(C) $\mathrm{H}_{3} \mathrm{PO}_{4}$ and $\mathrm{HPO}_{4}^{-}$
(D) $\mathrm{HPO}_{4}{ }^{2-}$ and $\mathrm{H}_{3} \mathrm{PO}_{4}$
4. If equal volumes of the following aqueous solutions were mixed, which one would have the highest pH ?
(A) $1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{NaOH}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{CH}_{3} \mathrm{COOH}$
(B) $1 \mathrm{molLL}^{-1} \mathrm{NH}_{3}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{H}_{2} \mathrm{SO}_{4}$
(C) $1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{H}_{2} \mathrm{SO}_{4}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{Ba}(\mathrm{OH})_{2}$
(D) $1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{KOH}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{HCl}$
5. 25 mL of a solution of $\mathrm{H}_{2} \mathrm{SO}_{4}$ that has a pH of 3 is pipetted into a 250 mL volumetric flask and distilled water added up to 250 mL . What is the pH of the diluted solution?
(A) 0.5
(B) 4.5
(C) 4
(D) 5
6. In the Haber process, which of the following conditions would result in an industrially acceptable method of increasing the yield of ammonia?
(A) increasing the temperature of the reaction chamber
(B) channelling the ammonia to a cooling chamber
(C) increasing the amount of $\mathrm{N}_{2}(\mathrm{~g})$
(D) increasing the amount of $\mathrm{H}_{2}(\mathrm{~g})$
7. In the production of ammonia using the Haber process, which of the following statements is incorrect?
(A) At equilibrium, the yield is higher when the temperature is lower.
(B) Before reaching equilibrium, the rate is higher at a higher temperature.
(C) The rate of the reaction is lower at a higher temperature because the reaction is exothermic.
(D) At equilibrium, the yield is lower at a lower pressure.
8. In a water treatment plant, the monitoring system for the quantity of the flocculant added to the water system malfunctioned with less than the recommended amount being added to the water. What would be the effect of this on the water quality?
(A) an unusually high bacterial count
(B) a high value of TDS (Total Dissolved Solids)
(C) a water supply with a pH between 8 and 9
(D) a water supply of high turbidity
9. Which of the following pairs of compounds are isomers?
(A) 1,2-difluorobutane and 1-fluorobutane
(B) 3-chloro-2-methyl-2-pentene and 1-chloro-1-hexene
(C) 2-bromopropane and 2-bromo-2-propene
(D) 1,2-difluorobutane and 1,2-dichlorobutane
10. The concentration of ozone in the troposphere is $0.000002 \%(\mathrm{v} / \mathrm{v})$. What is this concentration in parts per million ( ppm ) ?
(A) 0.0002
(B) 0.002
(C) 0.02
(D) 20
11. Which of these displacement reactions can occur spontaneously?
(A) $2 \mathrm{Ag}(s)+\mathrm{Cu}^{2+} \rightarrow 2 \mathrm{Ag}^{+}+\mathrm{Cu}(s)$
(B) $\mathrm{Pb}^{2+}+\mathrm{Sn}(s) \rightarrow \mathrm{Sn}^{2+}+\mathrm{Pb}(s)$
(C) $\mathrm{Fe}(s)+\mathrm{Mg}^{2+} \rightarrow \mathrm{Fe}^{2+}+\mathrm{Mg}(s)$
(D) $2 \mathrm{Al}^{3+}+3 \mathrm{Ni}(s) \rightarrow 2 \mathrm{Al}(s)+3 \mathrm{Ni}^{2+}$
12. The vanadium redox cell currently under development at UNSW acts as a galvanic cell during the reaction,

$$
\mathrm{V}^{2+}+\mathrm{VO}_{2}^{+}+2 \mathrm{H}^{+} \rightarrow \mathrm{V}^{3+}+\mathrm{VO}^{2+}+\mathrm{H}_{2} \mathrm{O}
$$

Which of the species below is the reductant in this reaction?
(A) $\mathrm{V}^{3+}$
(B) $\mathrm{VO}_{2}{ }^{+}$
(C) $\mathrm{H}^{+}$
(D) $\mathrm{V}^{2+}$
13. The reaction shows the production of ethanol from ethene.

$$
\mathrm{C}_{2} \mathrm{H}_{4}+\mathrm{H}_{2} \mathrm{O} \underset{ }{\rightleftarrows} \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}
$$

Which of the following conditions are necessary for this reaction?
(A) warmth, yeast
(B) heating, refluxing
(C) heating, fractional distillation
(D) heating, pressurising, sulfuric acid catalyst
14. Which of the following represent the empirical formulas of polyethene and polyvinyl chloride?
(A) $\mathrm{CH}_{2}$ and CHCl
(B) $\mathrm{CH}_{2}$ and $\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{Cl}$
(C) $\left(\mathrm{CH}_{2}-\mathrm{CH}_{2}\right) \mathrm{n}$ and $\left(\mathrm{CH}_{2}-\mathrm{CHCl}\right) \mathrm{n}$
(D) $\left(\mathrm{CH}_{2}-\mathrm{CH}_{2}\right) \mathrm{n}$ and $\left(\mathrm{CH}_{2}-\mathrm{CCl}_{2}\right) \mathrm{n}$
15. The equation shows the bombardment of U-235 with a neutron which initiates a fission reaction,

$$
{ }_{92}^{235} U+{ }_{0}^{1} n \rightarrow{ }_{56}^{147} B a+\mathrm{X}+3{ }_{0}^{1} n
$$

Which of the following correctly identifies species X ?
(A) $\mathrm{Kr}-36$
(B) $\mathrm{Kr}-86$
(C) $\mathrm{Pa}-91$
(D) $\mathrm{Np}-93$

## Answer Sheet

## Section I

## Part A

| 1. | $\mathrm{A} \AA$ | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |
| :--- | :--- | :--- | :--- | :--- |
| 2. | $\mathrm{A} \AA$ | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |
| 3. | $\mathrm{A} \AA$ | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |
| 5. | $\mathrm{A} \AA$ | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |
| 6. $\AA$ | $\mathrm{A} \AA$ | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |
|  | $\mathrm{B} \AA$ | $\mathrm{C} \AA$ | $\mathrm{D} \AA$ |  |

7. $\mathrm{A} \AA$

B Å
C A
D $\AA$
8. $\mathrm{A} \AA$

B Å
C Å
D $\AA$
9. $\mathrm{A} \AA$

B A
C A
D $\AA$
10. $\mathrm{A} \AA$

B Å
C Å
D Å
11. $\mathrm{A} \AA$

B Å
C Å
D Å
12. $\mathrm{A} \AA$

B Å
C A
D $\AA$
13. $\mathrm{A} \AA$

B Å
C Å
D $\AA$
14. $\mathrm{A} \AA \AA$

B Å
C Å
D $\AA \AA$
15. $\mathrm{A} \AA$

B Å
C Å
D Å

# JAMES RUSE AGRICULTURAL HIGH SCHOOL <br> 2002 CHEMISTRY TRIAL HSC EXAM <br> Section I (continued) 

## Part B-60 marks

Attempt Questions 16-28
Allow about 1 hour and 45 minutes for this part
Answer the questions in the spaces provided
Show all relevant working in questions involving calculations

Question 16 (3 marks)
A 5.00 mL volume of vinegar was found to weigh 4.50 g . The vinegar was placed into a conical flask and diluted with 20.0 mL of distilled water. The concentration of acetic acid (ethanoic acid) in the vinegar was determined by titration with $0.100 \mathrm{~mol} \mathrm{~L}^{-1}$ sodium hydroxide. At the endpoint, the titre was 23.3 mL .
(a) Calculate the percentage mass of acetic acid in the original undiluted vinegar.
$\qquad$
..
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(b) What is the concentration $\left(\mathrm{mol} \mathrm{L}^{-1}\right)$ of acetic acid in the undiluted vinegar?
$\qquad$
$\qquad$
$\qquad$

## Student Number.

$\qquad$

Question 17 (3 marks)
(a) (i) Explain why ammonium chloride can form an acidic solution in water and therefore classified as an acidic salt. Use equation(s) in your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) Explain why sodium hydrogen carbonate can form a basic solution in water and is therefore classified as a basic salt. Use equation(s) in your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(b) Name a salt which should form a neutral solution in water.

## Question 18 (8 marks)

During your course, you determined the heat of neutralisation of an acid. . Describe the procedure you followed and justify the appropriateness of the procedure you adopted in order to obtain valid and reliable results.
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## Student Number

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Question 19 (4 marks)
Naturally occurring citric acid has the molecular formula $\mathrm{C}_{6} \mathrm{H}_{8} \mathrm{O}_{7}$. Like all acids, it reacts with carbonate solutions to form carbon dioxide gas. When 1.537 g citric acid was added to a solution containing excess sodium carbonate, 295 mL carbon dioxide (measured at $25^{\circ} \mathrm{C}$ and 101.3 kPa ) was formed.
(a) Write an equation for the reaction of hydrogen ions with carbonate ions. How many moles of carbon dioxide were formed?
$\qquad$
$\qquad$
$\qquad$
(b) Determine the number of moles of hydrogen ions produced by 1.537 g citric acid.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) How many replaceable hydrogens (acidic hydrogens) are there in citric acid? Explain your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

## Question 20 (2 marks)

Using ethanoic acid and nitric acid in your answer. Draw diagrams to represent: (a) a concentrated, weak acid solution.
(b) a strong, dilute acid solution

Make sure you use correct formulas in your diagrams.

concentrated weak acid solution

strong, dilute acid solution

## Student Number

$\qquad$

Question 21 (6 marks)
(a) Describe the procedure you used to quantitatively analyse a manufactured product
$\qquad$
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(b) Identify one problem you encountered in the procedure.
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(c) Propose a solution to this problem.
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Question 22 (3 marks)
Evaluate the effectiveness of atomic absorption spectrophotometric (AAS) measurements in pollution control.
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## Student Number

$\qquad$

Question 23. ( 6 marks)
Describe the test for the biochemical oxygen demand (BOD) and evaluate its importance with respect to the monitoring of the possible eutrophication of waterways.
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## Question 24 (5 marks)

Discuss the problems associated with the use of CFCs and the steps taken to alleviate these problems. 5
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## Student Number

$\qquad$

Question 25 (5 marks)
(a) A student prepares 250 mL of a $5 \%(\mathrm{w} / \mathrm{v})$ glucose solution and adds 1 gram of yeast. Write a balanced chemical equation for the fermentation which occurs.
(b) Calculate the mass of ethanol produced.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) Describe conditions which promote fermentation.
$\qquad$
$\qquad$
$\qquad$
(d) Relate the structure of the ethanol molecule to its use as a solvent.
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## Question 26 (6 marks)

Using examples of named polymers, compare and contrast addition polymerisation with condensation polymerisation. Make reference to the sources of reactants and the processes used in the manufacture of the reactants.
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$\qquad$

## Student Number

$\qquad$

Question 27 (4 marks)
(a) Give a reason why some nuclei are unstable.
$\qquad$
(b) The reaction mechanism for esterification was studied and verified using a tracer. It was proved that water was formed from an H from the alcohol and an OH from the acid.


Identify a named radioisotope which could have been used in this research.
$\qquad$
(c) Cobalt-60 is a multi-purpose commercial radioisotope.

Describe how a non-transuranic isotope like cobalt-60 can be produced in a nuclear reactor and give an equation to illustrate your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

## Question 28 (5 marks)

The graph shows the boiling points of the alkane series from methane to octane.

(a) Identify which gaseous alkane (plotted on the graph) would be the easiest to liquefy and give a reason for your answer.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(b) Sketch a curve on the graph showing the relative boiling points of the alkanoic acids from ethanoic acid to hexanoic acid in relation to the corresponding alkanes.
(c) Explain the difference in the boiling points of the alkanes versus the alkanoic acids
$\qquad$
$\qquad$
$\qquad$
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$\qquad$

## Section II

## 25 marks

## Attempt Question 29

Allow about 45 minutes for this section.
Answer the question in a writing booklet provided
Show all relevant working in questions involving calculations
Question 29. (25 marks)
(a) Describe the work of Davy and Faraday in increasing the understanding of electron transfer reactions.
(b) Various methods can be used to protect the hulls of ships from corrosion. Explain four different methods that are used for protection.
(c) Describe a passivating metal.
(d) (i) Compare the concentrations of gases normally dissolved in the oceans to their concentrations in the atmosphere.
(ii) Explain how the solubilities of these 'soluble' gases vary with ocean depth
(e) Three experiments represented in diagrams $X, Y$, and $Z$ were set up.


In which of these experiments, all at the same temperature, will the iron corrode the most and which will corrode the least? Explain your answer.

## Please turn over

(f) Silver plated objects are obtained in an electrolytic cell in which the object is one electrode. The other electrode is a block of silver, and silver cyanide solution, AgCN , is the electrolyte. The cell is illustrated below:

(i) Write a balanced equation to represent the oxidation process. At which electrode does this occur?
(ii) Write a balanced equation to represent the reduction process. At which electrode does this occur?
(g) Consider the electrolysis set-up below:


Describe using half-equations what happens at the anode and at the cathode during electrolysis.

## End of Question 29

END OF TEST $\pi$

## Section I <br> Total Marks (75)

## Part A

Total marks (15)
Attempt Questions 1-15
Allow about 30 minutes for this part

Use the multiple choice answer sheet.
Select the alternative A, B, C or D that best answers the question. Fill in the response square completely.

Sample 2+4=(A) $2 \quad$ (B) $6 \quad$ (C) $8 \quad$ (D) 9
A Å
B $\zeta$
C Å
D $\AA$

If you think you have made a mistake, put a cross through the incorrect answer and fill in the new answer.
$A \zeta$
B $\zeta$
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D Å

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correct
$A \zeta$
B $\zeta$
C Å
D $\AA$

## SECTION I <br> Part A <br> Multiple Choice

1. The extraction of copper from copper(I) sulfide produces sulfur dioxide as a by-product according to the equation:

$$
\mathrm{Cu}_{2} \mathrm{~S}(s)+\mathrm{O}_{2}(g) \rightarrow 2 \mathrm{Cu}(s)+\mathrm{SO}_{2}(g)
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What volume of sulfur dioxide gas will be released at $25^{\circ} \mathrm{C} \& 101.3 \mathrm{kPa}$ when 2.2 g of copper (I) sulfide is reacted?
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3. $\mathrm{H}_{2} \mathrm{PO}_{4}^{-}$is an amphiprotic species.

Which of the following represents the conjugate acid and conjugate base respectively of $\mathrm{H}_{2} \mathrm{PO}_{4}{ }^{-}$?
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(B) $\mathrm{PO}_{4}{ }^{3-}$ and $\mathrm{H}_{3} \mathrm{PO}_{4}$
(C) $\mathrm{H}_{3} \mathrm{PO}_{4}$ and $\mathrm{HPO}_{4}^{-}$
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4. If equal volumes of the following aqueous solutions were mixed, which one would have the highest pH ?
(A) $1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{NaOH}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{CH}_{3} \mathrm{COOH}$
(B) $1 \mathrm{molLL}^{-1} \mathrm{NH}_{3}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{H}_{2} \mathrm{SO}_{4}$
(C) $1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{H}_{2} \mathrm{SO}_{4}+1 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{Ba}(\mathrm{OH})_{2}$
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5. 25 mL of a solution of $\mathrm{H}_{2} \mathrm{SO}_{4}$ that has a pH of 3 is pipetted into a 250 mL volumetric flask and distilled water added up to 250 mL . What is the pH of the diluted solution?
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7. In the production of ammonia using the Haber process, which of the following statements is incorrect?
(A) At equilibrium, the yield is higher when the temperature is lower.
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$$
\mathrm{V}^{2+}+\mathrm{VO}_{2}^{+}+2 \mathrm{H}^{+} \rightarrow \mathrm{V}^{3+}+\mathrm{VO}^{2+}+\mathrm{H}_{2} \mathrm{O}
$$

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(C) $\mathrm{H}^{+}$
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$$
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(C) $\left(\mathrm{CH}_{2}=\mathrm{CH}_{2}\right) \mathrm{n}$ and $\left(\mathrm{CH}_{2}=\mathrm{CHCl}\right) \mathrm{n}$
(D) $\left(\mathrm{CH}_{2}=\mathrm{CH}_{2}\right) \mathrm{n}$ and $\left(\mathrm{CH}_{2}=\mathrm{CCl}_{2}\right) \mathrm{n}$
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$$

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(A) $\mathrm{Kr}-36$
(B) $\mathrm{Kr}-91$
(C) $\mathrm{Pa}-91$
(D) $\mathrm{Np}-93$

## Answer Sheet

## Section I

## Part A

1. $\mathrm{A} \zeta$
B Å
C Å
D $\AA$
2. $\mathrm{A} \AA$
B Å
$C \zeta$
D Å
3. $\mathrm{A} \zeta$

B Å
C Å
D Å
4. $\mathrm{A} \zeta$

B Å
C Å
D Å
5. $\mathrm{A} \AA$

B Å
$C \zeta$
D $\AA$
6. $\mathrm{A} \AA$

B $\zeta$
C Å
D $\AA$
7. $\mathrm{A} \AA$

B Å
$C \zeta$
D $\AA$
8. $\mathrm{A} \AA$

B Å
C Å
D $\zeta$
9. $\mathrm{A} \AA$

B $\zeta$
C Å
D Å
10. $\mathrm{A} \AA$

B Å
$C \zeta$
D Å
11. $\mathrm{A} \AA$

B $\zeta$
C Å
D Å
12. $\mathrm{A} \AA$

B Å
C A
D $\zeta$
13. $\mathrm{A} \AA$

B Å
C Å
D $\zeta$
14. $\mathrm{A} \AA$

B $\zeta$
C A
D $\AA$
15. $\mathrm{A} \zeta$

B Å
C A
D $\AA$

JAMES RUSE AGRICULTURAL HIGH SCHOOL
2002 CHEMISTRY TRIAL HSC EXAM
Section I (continued)

## Part B - 60 marks

Attempt Questions 16-28
Allow about 1 hour and 45 minutes for this part
Answer the questions in the spaces provided
Show all relevant working in questions involving calculations

Question 16 (3 marks)
A 5.00 mL volume of vinegar was found to weigh 4.50 g . The vinegar was placed into a conical flask and diluted with 20.0 mL of distilled water. The concentration of acetic acid (ethanoic acid) in the vinegar was determined by titration with $0.100 \mathrm{~mol} \mathrm{~L}^{-1}$ sodium hydroxide. At the endpoint, the titre was 23.3 mL .
(a) Calculate the percentage mass of acetic acid in the original undiluted vinegar.

| Criteria | Mark |
| :--- | :--- | :--- |
| mole $\mathrm{CH}_{3} \mathrm{COOH}$ $=$ mole $\mathrm{NaOH}=\left(0.100 \mathrm{molL}^{-1}\right)(0.0233 \mathrm{~L})=2.33 \times 10^{-3} \mathrm{~mol}$ <br> mass $\mathrm{CH}_{3} \mathrm{COOH}$ $=$ mole $\times$ molar mass <br>  $=2.33 \times 10^{-3}$ mol $\times[2(12.01)+2(16.00)+4(1.008)]$ <br>  $=0.13992 \sim 0.140 \mathrm{~g}$ |  |
| $\% \mathrm{CH}_{3} \mathrm{COOH}$ | $=($ mass $/$ volume $) 100 \%=\frac{0.140}{5.00} \times 100=\mathbf{2 . 8 \%}$ |$\quad 2$|  |
| :--- |

(b) What is the concentration $\left(\mathrm{mol} \mathrm{L}^{-1}\right)$ of acetic acid in the undiluted vinegar?

|  | Criteria |
| :--- | :--- |
| $\left.\mathrm{CH}_{3} \mathrm{COOH}\right]$ $=$ mole $\mathrm{CH}_{3} \mathrm{COOH} / \mathrm{vol} \mathrm{in} \mathrm{L}$ <br>  $2.33 \times 10^{-3} \mathrm{~mole} / 0.005 \mathrm{~L}$ <br>  $=\mathbf{0 . 4 6 6}$ mole $/ \mathrm{L}$ |  |
| Correct calculation | 1 |

Question 17 (3 marks)
(a) (i) Explain why ammonium chloride can form an acidic solution in water and therefore classified as an acidic salt. Use equation(s) in your answer.

| Criteria | Mark(s) |
| :--- | :--- |
| The $\mathrm{NH}_{4}{ }^{+}$ion is a better proton donor, ie a stronger acid than water and hence, is able to donate a <br> proton to water to increase $\mathrm{H}_{3} \mathrm{O}^{+}$concentration in water. <br> $\mathrm{NH}_{4}{ }^{+}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{+}$ | 1 |
| any answer with a similar reasoning | 1 |

(ii) Explain why sodium hydrogen carbonate can form a basic solution in water and is therefore classified as a basic salt. Use equation(s) in your answer.

| Criteria | Mark(s) |
| :--- | :--- |
| The $\mathrm{HCO}_{3}^{-}$ion is amphiprotic substance but tends to accept rather than donate protons to water. | 1 |
| Abstraction of a proton from water results in a basic solution |  |
| $\mathrm{HCO}_{3}{ }^{-}+\mathrm{H}_{2} \mathrm{O} \Leftrightarrow \mathrm{H}_{2} \mathrm{CO}_{3}+\mathrm{OH}^{-}$. Free $\mathrm{OH}^{-}$ions in solution make the solution basic | 1 |
| any similar reasoning with an equation |  |

(b) Name a salt which should form a neutral solution in water.

| Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :--- |
| Sodium chloride | 1 |
| any other neutral salt | 1 |

Question 18 (8 marks)
During your course, you determined the heat of neutralisation of an acid. . Describe the procedure you followed and justify the appropriateness of the procedure you adopted in order to obtain valid and reliable results.

\begin{tabular}{|c|c|}
\hline Criteria \& Mark(s) \\
\hline \begin{tabular}{l}
The heat of neutralisation is the heat released when 1 mole of an acid is neutralised with 1 mole of a base \(e g \mathrm{HCl}+\mathrm{NaOH} \rightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{O}(l)\)
\[
\text { or } \mathrm{H}^{+}+\mathrm{OH}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}(l) \quad \Delta \mathrm{H}_{\text {neut }}=(-) \text { exothermic }
\] \\
The experiment was done in the following manner: \\
ò \(\quad 50.0 \mathrm{~mL}\) of \(1.00 \mathrm{molL}^{-1} \mathrm{HCl}\) was pipetted into a polystyrene cup and its temperature measured and recorded. \\
ò 50.0 mL of \(1.00 \mathrm{molL}^{-1} \mathrm{HCl}\) was placed into a 50.0 mL burette and its temperature measured and recorded \\
ò The NaOH solution was then added to the HCl solution; stirred (carefully) with the thermometer and the maximum temperature reached noted. The mixture was stirred before recording the temperature to avoid recording localised temperature changes as those of the bulk of the solution. \\
A polystyrene cup (good insulator) is used to prevent heat transfer through the walls of the container. Accurate measurements of volumes, use of pipette and burette is needed for an accurate calculation of mass of solution and number of moles of acid or base. Repeating the experiment several times could add to the reliability of the results. The acids and the bases were both accurately known by standardisation so when \(\Delta \mathrm{H}\) is calculated, \(\Delta \mathrm{H}=(\mathrm{mCg} \Delta \mathrm{t}) / \mathrm{n}\). the value of n is an accurate one. The initial temperatures of the NaOH and the HCl may differ hence, the initial temperatures of the HCl and the NaOH were determined and averaged.. Assumption of a density of \(1 \mathrm{~g} / \mathrm{mL}\) for water incurred very little error or a databook. Alternatively, a data book could have been consulted for the density of water at specific temperatures.
\end{tabular} \& 1

3

4 <br>
\hline Heat of neutralisation is defined, appropriate equations given or heat of neutralisation explained without equations. Detailed procedure is given and with an analysis of the procedure \& 8 <br>
\hline
\end{tabular}

Question 19 (4 marks)
Naturally occurring citric acid has the molecular formula $\mathrm{C}_{6} \mathrm{H}_{8} \mathrm{O}_{7}$. Like all acids, it reacts with carbonate solutions to form carbon dioxide gas. When 1.537 g citric acid was added to a solution containing excess sodium carbonate, 295 mL carbon dioxide (measured at $25^{\circ} \mathrm{C}$ and 101.3 kPa ) was formed.
(a) Write an equation for the reaction of hydrogen ions with carbonate ions. How many moles of carbon dioxide were formed?

| Criteria | Mark(s) |
| :--- | :--- |
| $2 \mathrm{H}^{+}+\mathrm{CO}_{3}{ }^{2-} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{CO}_{2}(g)$ 1 <br> ${\mathrm{Moles} \mathrm{CO}_{2}}=$ volume $\mathrm{CO}_{2} /$ molar volume at SLC  <br> $=0.295 \mathrm{~L} / 24.47 \mathrm{~L} /$ mole $=\mathbf{0 . 0 1 2 1}$ moles  | 1 |

(b) Determine the number of moles of hydrogen ions produced by 1.537 g citric acid.

| Criteria | Mark(s) |
| :--- | :--- |
| Moles $\mathrm{H}^{+}=2 \times$ moles $\mathrm{CO}_{2}=2 \times 0.0121=0.0242$ moles |  |
| Moles of citric acid $=1.537 \mathrm{~g} /[6(12.01)+8(1.008)+7(16.00)] \mathrm{g} / \mathrm{mole}=\mathbf{8 . 0 1} \times \mathbf{1 0}^{-3}$ | 1 |
| Moles $\mathrm{H}^{+}=$moles $\mathrm{H}^{+}$from citric acid? |  |
|  |  |

(c) How many replaceable hydrogens (acidic hydrogens) are there in citric acid? Explain your answer.

| Criteria | Mark(s) |
| :--- | :--- |
| Moles of $\mathrm{H}^{+}$by stoichiometry $=0.0242$ while | 1 |
| Moles of citric acid $=8.01 \times 10^{-3}$ |  |
| There is 3 times as much $\mathrm{H}^{+}$reacting with $\mathrm{CO}_{2}$ as there are moles of citric acid, therefore, there are |  |
| 3 replaceable hydrogens in citric acid. |  |

## Question 20 (2 marks)

Using ethanoic acid and nitric acid in your answer. Draw diagrams to represent: (a) a concentrated, weak acid solution.
(b) a strong, dilute acid solution

Make sure you use correct formulas in your diagrams.

concentrated weak acid solution

strong, dilute acid solution

Question 21 (6 marks)
(a) Describe the procedure you used to quantitatively analyse a manufactured product

| Criteria | Mark(s) |
| :--- | :--- |
| Analysis of citric acid in orange juice. Exactly 25.00 mL of orange juice was pipetted into a <br> conical flask. The pipette was previously rinsed with orange juice and the conical flask, rinsed with <br> demineralised water. The orange juice was titrated with standard NaOH solution with <br> phenolphthalein as the indicator. The NaOH solution was added until the yellow solution turned <br> orange. The number of moles of citric acid $=3$ times the number of moles of $\mathrm{NaOH} ;$ <br> $\mathrm{H}_{3} \mathrm{Cit}+3 \mathrm{NaOH} \rightarrow \mathrm{Na}_{3} \mathrm{Cit}+3 \mathrm{H}_{2} \mathrm{O}$ | 4 |
| Complete procedure including equation(s), the principle behind the analysis and the stoichiometry <br> of the reaction | 4 |

(b) Identify one problem you encountered in the procedure.

| Criteria | Mark(s) |
| :--- | :--- |
| One problem is the difficulty in judging the endpoint of the titration since the orange juice is itself <br> highly coloured | 1 |
| any valid problem | 1 |

(c) Propose a solution to this problem.

| Criteria | Mark(s) |
| :--- | :--- |
| A solution to this problem is the use of a pH glass electrode to monitor the titration. The electrode <br> should be rinsed well after the titration to prevent any orange juice component from fouling the <br> glass membrane | 1 |
| any reasonable solution to the analysis problem given in (b) | 1 |

Question 22 (3 marks)
Evaluate the effectiveness of atomic absorption spectrophotometric (AAS) measurements in pollution control.

| Criteria | Mark(s) |
| :--- | :--- |
| Metal ions can be dangerous pollutants even at low levels. Therefore, studies utilising AAS are very <br> effective in pollution control because of the ability of AAS to detect metal ions at either very low or <br> very high levels. For very low levels, pre-concentration methods such as solvent extraction or <br> solvent evaporation may be done. Normally, the sample is simply directly introduced into the <br> instrument without pre-separation. Interference problems are minimal. This is due to the inherent <br> selectivity of the AAS technique which uses a specific hollow cathode lamp for each target ion. <br> Samples containing high levels of metal ions can be diluted to suit the requirements of the <br> instrument and the techniques. |  |
| The selectivity, low detection limit and ease of introducing the sample into the instrument should be <br> included. | 3 |

Question 23. (6 marks)
Describe the test for the biochemical oxygen demand (BOD) and evaluate its importance with respect to the monitoring of the possible eutrophication of waterways.

| Criteria | Mark(s) |
| :--- | :--- |
| Two identical water samples are collected and analysed for dissolved oxygen by the same technique <br> 5 days apart. After collection both samples are protected from light and oxygen exposure. Dissolved <br> oxygen is analysed with the oxygen electrode or the Winkler titration technique. The difference in <br> the dissolved oxygen level (DO) between the two samples is the biochemical oxygen demand <br> (BOD) of the water sample. BOD is a measure of the concentration of dissolved oxygen needed for <br> the complete breakdown of the organic matter in the water by aerobic bacteria which in turn uses up <br> oxygen and at the same time raises the nutrient level of the waterway. These result in the depletion <br> of oxygen, development of unpleasant odour and colonies of undesirable organisms such as algae. <br> When these organisms are broken down, further degradation and eutrophication of the waterway <br> result. Along with other tests, monitoring the BOD indicates the level of organic nutrients in the <br> water. The level of organic nutrients can indicate possible contamination with sewage making it <br> unfit for consumption and also can indicate the possibility of an enhaced rate of eutrophication in <br> that waterway. |  |
| Test for BOD described |  |
| Role of BOD in eutrophication and importance of monitoring BOD | 3 |

Question 24 (5 marks)
Discuss the problems associated with the use of CFCs and the steps taken to alleviate these problems.

| Criteria | Mark(s) |
| :--- | :--- |
| The main problem associated with the use of CFCs is their ability to destroy ozone in the <br> stratosphere. CFCs are broken down in the stratosphere under the influence of UV radiation to <br> produce chlorine free-radicals which can destroy ozone. The destruction of the Earth's radiation <br> "shield", the ozone layer, exposes the Earth's surface to dangerous levels of ozone which can cause <br> cell damage in both plants and animals, in humans leading to skin cancers and cataracts. The <br> CFCs are also greenhouse gases which contribute to global warming leading to global erratic <br> weather patterns. The Montreal Protocol in its various versions aimed to alleviate these problems <br> by halving and later on totally banning the manufacture of CFCs, the encouragement of the use of | 5 |
| CFC replacementss which are planned to be phased out as well and the setting up of funds to help <br> third world countries phase out CFCs. |  |
| Problems associated with CFCs | 3 |
| Montreal protocol regulations | 2 |

Question 25 (5 marks)
(a) A student prepares 250 mL of a $5 \%(\mathrm{w} / \mathrm{v})$ glucose solution and adds 1 gram of yeast. Write a balanced chemical equation for the fermentation which occurs.

| Criteria | Mark(s) |
| :--- | :--- |
| $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$-------> $2 \mathrm{CO}_{2}(g)+2 \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ | 1 |
| Yeast enzymes must be included to obtain the mark | 1 |

(b) Calculate the mass of ethanol produced.

| Criteria | Mark(s) |
| :--- | :--- |
| mass of glucose $=(0.05 \mathrm{~g} / \mathrm{mL}) \times 250 \mathrm{~mL}=12.5 \mathrm{~g}$  <br> mole $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ $=2 \times$ moles $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$  <br>  $=2 \times \frac{12.5}{6(12.01)+12(1.008)+6(16.00)}=0.1388 \sim \mathbf{0 . 1 3 9}$ mole  <br> mass of $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ $=$ mole of $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \times$ molar mass $=0.139 \times[2(12.01)+6(1.008)+16.00]$  <br>  $=\mathbf{6 . 3 6} \mathbf{g}$ 1 | 1 |

(c) Describe conditions which promote fermentation.

| Criteria | Mark(s) |
| :--- | :--- |
| Warm temperature, $\sim 37^{\circ} \mathrm{C}$ as well as neutral pH and the presence of yeast enzymes promote <br> fermentation | 1 |
| At least two conditions should be given to obtain the mark. | 1 |

(d) Relate the structure of the ethanol molecule to its use as a solvent.

| Criteria | Mark(s) |
| :--- | :--- |
| The ethanol molecule is a good solvent for polar substances because it is a polar molecule. The <br> highly electronegative oxygen depletes the carbon and the hydrogen of electrons making them <br> partially positive and itself, partially negative. It is able, therefore to dissolve polar substances by <br> dipole-dipole interaction or hydroxylated substances by hydrogen bonding | 1 |
| any reasonable explanation | 1 |

Question 26 (6 marks)
Using examples of named polymers, compare and contrast addition polymerisation with condensation polymerisation. Make reference to the sources of reactants and the and the processes used in their manufacture.

| Criteria | Mark(s) |
| :---: | :---: |
| Addition polymerisation and condensation polymerisation both involve the combining together of small molecules (monomers) to form one large molecule (polymer). They differ in many respects: ü Addition polymerisation may either utilise an initiator or a catalyst for polymerisation whereas no general catalyst is utilised in condensation polymerisation. <br> ü The monomers in addition polymerisation are normally identical, for example for polyethene, ethene is the only monomer used in the manufacture of polyethene: $n \mathrm{CH}_{2}=\mathrm{CH}_{2}+\mathrm{CH}_{2}=\mathrm{CH}_{2} \rightarrow-\mathrm{CH}_{2}\left(\mathrm{CH}_{2}-\mathrm{CH}_{2}\right)_{\mathrm{n}}-\mathrm{CH}_{2}-$ <br> where as in condensation polymerisation, the monomers may be identical as in nylon, (6-aminohexanoic acid) or different as in the manufacture of polyester from ethylene glycol and terephthalic acid. <br> ü No other product is formed in addition polymerisation whereas a small molecule, for example, $\mathrm{H}_{2} \mathrm{O}$ or $\mathrm{NH}_{3}$ is produced in condensation polymerisation. <br> ü Reactants (ethene, mainly) in addition polymerisation are obtained from the cracking of petroleum while those of condensation polymerisation may come from the degradation of polymers of biological origins such cellulose or are industrially synthesised. | 6 |
| Discussion of similarity | 1 |
| Discussion of differences | 3 |
| Sources of monomers | 2 |

## MARKS

Question 27 (4 marks)
(a) Give a reason why some nuclei are unstable.

| Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :---: |
| proton to neutron ratio is too high or too low or the atom is too heavy $(\mathrm{Z}>83)$ | 1 |

(b) The reaction mechanism for esterification was studied and verified using a tracer. It was proved that water was formed from an H from the alcohol and an OH from the acid.


Identify a named radioisotope which could have been used in this research.

|  | Criteria |
| :--- | :---: | Mark(s) $\quad 1$

(c) Cobalt-60 is a multi-purpose commercial radioisotope.

Describe how a non-transuranic isotope like cobalt-60 can be produced in a nuclear reactor and give an equation to illustrate your answer.

| Criteria | Mark(s) |
| :--- | :---: |
| A non-transuranic isotope such as cobalt-60 can be produced by irradiating a stable |  |
| isotope such as cobalt-59 with neutrons in a nuclear reactor. | 2 |
| ${ }^{59} \mathrm{Co}+n \quad \rightarrow \quad{ }_{27} \mathrm{Co}$ | Co |
| Cobalt-60 decays to produce $\beta$ and $\gamma$ radiations. | 1 |
| Description of process | 1 |
| equation |  |

Question 28 (5 marks)


The graph shows the boiling points of the alkane series from methane to octane.
(a) Identify which gaseous alkane (plotted on the graph) would be the easiest to liquefy and give a reason for your answer.

| Criteria | Mark(s) |
| :--- | :---: |
| Butane, the gas with the highest boiling point and hence the easiest to liquify. If all <br> gases were cooled it will be the first to reach its "condensation" temperature which is <br> the same as its boiling point. | 2 |
| Butane | 1 |
| explanation of choice | 1 |

(b) Sketch a curve on the graph showing the relative boiling points of the alkanoic acids from ethanoic acid to hexanoic acid in relation to the corresponding alkanes.

| Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :---: |
| as indicated in the graph | 1 |

(c) Explain the difference in the boiling points of the alkanes versus the alkanoic acids

| Criteria | Mark(s) |
| :--- | :---: |
| The alkanes have only weak dispersion forces between molecules and hence require <br> only a small amount of energy to separate them. However, alkanoic acids are <br> extensively hydrogen-bonded to each other, hydrogen bonding is a stronger <br> intermolecular force than dispersion forces and hence require more energy to separate <br> the molecules from each other, i.e., higher boiling point. | 2 |
| discussion of two forces | 2 |

## Section II 25 marks

Question 29. (25 marks)
(a) Describe the work of Davy and Faraday in increasing the understanding of electron transfer reactions.

| Criteria | Mark(s) |
| :--- | :---: |
| Davy discovered electrolysis, the use of electricity to break down substances <br> chemically. His contribution includes the electrolysis of sodium and potassium, a <br> technique using electron-transfer processes. He demontrated these procedures without <br> specifically explaining them. | 2 |
| Faraday quantified electrolysis with his laws. He related the mass of substances <br> produced and the amount of electricity passing through. | 2 |

(b) Various methods can be used to protect the hulls of ships from corrosion. Explain four different methods that are used for protection.

|  | Criteria | Mark(s) |
| :--- | :---: | :---: |
| Four different methods used to protect the hulls of ships from corrosion are: <br> ü | Painting the hull of a ship with new polymer paints such as the Rustmaster Pro. This paint forms a <br> smooth impermeable (to oxygen and water) layer on the surface of the hull. It also incorporates <br> additives that react with the surface to form a very insoluble ionic substance pyroaurite. This layer | 2 |
| prevents ion migration on the surface and in conjunction with the polymer surface shields the surface <br> from oxygen and water, thus, inhibiting corrosion. |  |  |
| ü | Hulls of ships are bombarded with metal ions such as Ni and Cr in the plasma state. These metal ions <br> form surface alloys on the surface of the hulls to form a "stainless steel " - like finish. The protection <br> is similar to the surface passivation afforded by alloying metals such as Cr and Ni in stainless steel. | 2 |

ü Cathodic protection may be employed to protect the hulls of ships. This is done by attaching an active metal i.e. one which has a more negative reduction potential than iron (such as Zn and Mg ) to the hull of the ship. These metals, because of their greater reactivity act as sacrificial anodes. The hull becomes the cathode and the active metal becomes the sacrificial anode. The reactions are:
$\mathrm{O}_{2}+2 \mathrm{H}_{2} \mathrm{O}+4 e^{-} \rightarrow 4 \mathrm{OH}^{-}$ cathode reaction $\mathrm{Mg}(s) \quad \rightarrow \mathrm{Mg}^{2+} \quad+2 e^{-} \quad$ anode reaction
ü An impressed current may be applied to the hull of the ship with an inert anode and the ship's hull as the cathode. Any $\mathrm{Fe}^{2+}$ ions are reduced as well as $\mathrm{O}_{2}$ (The reduction of oxxygen is the main reduction process. The reactions are:
$\mathrm{O}_{2}+2 \mathrm{H}_{2} \mathrm{O}+4 \mathrm{e}^{-} \rightarrow 4 \mathrm{OH}^{-} \quad$ cathode reactions
$\mathrm{Fe}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Fe}(s)$
$2 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{O}_{2}+4 \mathrm{H}^{+}+4 \mathrm{e}^{-} \quad$ anode reaction at the inert electrode
(c) Describe a passivating metal.

| Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :---: |
| A passivating metal is a reactive metal that readily forms an unreactive surface coating <br> with substances such as $\mathrm{O}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ | 1 |

(d) Compare the concentrations of gases normally dissolved in the oceans to their concentrations in the atmosphere. How do the solubilities of these 'soluble' gases vary with ocean depth? Explain.

| Criteria | Mark(s) |
| :--- | :---: |
| The concentration of $\mathrm{O}_{2}$ in the atmosphere is about 30 times more than its <br> concentration in the surface waters of the ocean. $\mathrm{CO}_{2}$ on the other hand is so much <br> more soluble in ocean water than oxygen is with about $90 \%$ as much concentration as | 2 |
| atmospheric levels. |  |$\quad$| With ocean depth, the solubility of these gases should increase because of the dual |
| :--- |
| positive effect of temperature and pressure on solubility. As depth increases, |
| temperature decreases and hence, in general, gas solubility increases. Similarly an |
| increase in solubility should result from an increase in pressure which results from |
| increased depth. However, actual concentrations of the gases on the surface waters of |
| the ocean are still higher than the bulk of the ocean waters because of immdiate |
| contact with the atmosphere, the main source of these gases and because of the |
| inefficient mixing of surface waters with the bulk of the water of the ocean. |$\quad$.

(e) Three experiments represented in diagrams $X, Y$, and $Z$ were set up.


In which of these experiments, all at the same temperature, will the iron corrode the most and which will corrode the least? Explain your answer.

| Criteria | Mark(s) |
| :---: | :---: |
| Set-up $Z$ will have the iron corroding most. This is because in contact with tin, iron act as the anode and the tin as the cathode. This is due to the greater tendency of iron $\left(\mathrm{E}_{\text {red }}^{0}=-0.44 \mathrm{v}\right.$ ) i.e. more negative reduction potential, to be oxidised compared with Sn , ( $\mathrm{E}_{\text {red }}^{0}=-0.14 \mathrm{v}$ ). <br> The set-up which will corrode least is $Y$. Zinc in contact with iron protects the iron from corrosion by acting as the sacrificial anode in the "corrosion" galvanic cell. This is due to the greater reactivity of zinc ( $E^{0}$ red $=-0.76 \mathrm{v}$ ) compared with iron $\left(\mathrm{E}_{\text {red }}^{0}=-0.44 \mathrm{v}\right) .$ <br> The reaction in set-up $Y$ is <br> anode: $\mathrm{Zn}(s) \xrightarrow{\rightarrow} \mathrm{Zn}^{2+}+2 \mathrm{e}^{-}$ <br> cathode: $\mathrm{O}_{2}+\mathrm{H}_{2} \mathrm{O}+4 \mathrm{e}^{-} \rightarrow 4 \mathrm{OH}^{-}$ | 2 2 |

(f) Silver plated objects are obtained in an electrolytic cell in which the object is one electrode. The other electrode is a block of silver, and silver cyanide solution, AgCN , (actually, $\operatorname{Ag}(\mathrm{CN})_{2}{ }^{-}$complex) is the electrolyte. The cell is illustrated below:

(i) Write a balanced equation to represent the oxidation process. At which electrode does this occur?

|  | Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :--- | :--- |
| $\mathrm{Ag}(\mathrm{s})$ <br> or <br> $\mathrm{Ag}(\mathrm{s})+2 \mathrm{Ag}^{+}+\mathrm{e}^{-} \rightarrow \mathrm{Ag}(\mathrm{CN})_{2}{ }^{-}+\mathrm{e}^{-}$ | 1 |  |

(ii) Write a balanced equation to represent the reduction process. At which electrode does this occur?

|  | Criteria | $\operatorname{Mark}(\mathrm{s})$ |
| :--- | :--- | :---: |
| $\mathrm{Ag}^{+}+e^{-} \rightarrow \mathrm{Ag}(\mathrm{s})$ | 1 |  |
| or |  |  |
| $\mathrm{Ag}(\mathrm{CN})_{2}{ }^{-}+e^{-} \rightarrow \mathrm{Ag}(\mathrm{s})+2 \mathrm{CN}^{-}$ |  |  |

(g) Consider the electrolysis set-up below:


Describe using half-equations what happens at the anode and the cathode during electrolysis given the condition outlined in the table below:

| Cell | Electrodes | Electrolyte |
| :--- | :--- | :--- |
| 1 | copper | $\mathrm{CuSO}_{4}$ |
| 2 | carbon | $\mathrm{K}_{2} \mathrm{SO}_{4}$ |


|  | Criteria |
| :--- | :---: |
| Cell1 | Mark(s) |
| Anode (positive electrode): | 1 |
| $\mathrm{Cu}(\mathrm{s}) \rightarrow \mathrm{Cu}^{2+}+2 \mathrm{e}^{-}$ |  |
| Cathode (negative electrode) |  |
| $\mathrm{Cu}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Cu}(\mathrm{s})$ |  |
| Cell 2 |  |
| Anode:(positive electrode) |  |
| $\mathrm{H}_{2} \mathrm{O} \rightarrow 1 / 2 \mathrm{O}_{2}(\mathrm{~g})+2 \mathrm{H}^{+}+2 \mathrm{e}^{-}$ |  |
| Cathode: $\mathrm{H}_{2} \mathrm{O}+\mathrm{e}^{-} \rightarrow 1 / 2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{OH}^{-}$ |  |

