

CHEMISTRY Stage 3 WACE Examination 2014 Marking Key

Marking keys are an explicit statement about what the examiner expects of candidates when they respond to a question. They are essential for fair assessment because their proper construction underpins reliability and validity.

CHEMISTRY STAGE 3

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Section One: Multiple-choice 25% (25 Marks)

| Question No. | Answer |
|-----------------------------------------------------------------------|-------------------------------------------------------------|
| 1 | В |
| 2 | D |
| 3 | D |
| 4 | В |
| 5 | D |
| 6 | D |
| 7 | С |
| 8 | Α |
| 9 | С |
| 10 | С |
| 11 | Α |
| 12 | В |
| 13 | D |
| 14 | С |
| 15 | D |
| 16 | D |
| 17 | Α |
| 18 | С |
| 19 | С |
| 20 | В |
| 1 2 3 4 5 6 6 7 8 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 | B D D B D C A C C A B D D D C A C C B A C C B A C C A A A A |
| 22 | С |
| 23 | В |
| 24 | A |
| 25 | A |

End of Section One

Section Two: Short answer

35% (72 Marks)

Question 26 (4 marks)

For the following reactions, describe fully the observed changes, including any

- colour changes
- odours
- precipitates (give the colour)
- gases evolved (give the colour or describe as colourless).
- (a) Lead(II) nitrate solution is added to potassium iodide solution.

(1 mark)

$$Pb^{2+}(aq) + 2\Gamma(aq) \rightarrow PbI_2(s)$$

(b) Solid copper(II) carbonate is added to dilute nitric acid solution.

(3 marks)

$$CuCO_3(s) \quad + \quad 2 \; H^{\scriptscriptstyle +}(aq) \quad \rightarrow \quad Cu^{2^{\scriptscriptstyle +}}(aq) \quad + \quad CO_2(g) \quad + \quad H_2O(\ell)$$

| Description | Marks |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| (a) Yellow precipitate formed | 1 |
| (b) Green solid dissolves to give blue solution and a colourless and (odourless) gas is evolved (1 for solid dissolving; 1 for colour change green to blue and 1 for colourless,(odourless) gas) | 1–3 |
| Incorrect | 0 |
| Total | 4 |

Question 27 (4 marks)

Write balanced ionic equations to represent the reactions described below.

(a) Chlorine gas is bubbled through an aqueous solution of sodium bromide. (2 marks)

$$C\ell_2(g)$$
 + 2 Br $\bar{}$ (aq) \rightarrow 2 C $\ell\bar{}$ (aq) + Br $_2$ (aq)

(b) Solid nickel oxide is added to dilute hydrochloric acid solution.

(2 marks)

NiO(s) + 2 H⁺(aq)
$$\rightarrow$$
 Ni²⁺(aq) + H₂O(ℓ)

| Description | Marks |
|---------------------------------------------|-------|
| Correct formulae for reactants and products | 1–2 |
| Balanced equations | 1–2 |
| Incorrect | 0 |
| Total | 4 |

NB: State symbols not required.

Award 1 mark for correctly balanced molecular equation.

Question 28 (8 marks)

Complete the table below by **either** drawing the Lewis structures **or** naming the shape of the molecules. For Lewis structures, any lone pairs must be shown.

All electron shell pairs should be represented as either: or as —.

| Molecule/Ion | Lewis structure | Name of shape |
|------------------------------|-------------------------------------|----------------------|
| NCℓ ₃ | :Ċŀ:N:Ċŀ: :Ċŀ: | (Trigonal) pyramidal |
| H₂S | H:S:H | Bent (V-shaped) |
| SiH₄ | H H—Si—H :: H :Si:H H or H | tetrahedral |
| O ₃ | ;ö: | Bent (V-shaped) |
| NO ₃ ⁻ | [::::] | Trigonal planar |

| Description | Marks |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Shapes correctly named | 1–2 |
| All Lewis structures correct including lone pairs | 6 |
| Octet around central atom is correctly shown (i.e. bonds are correct) for all structures but lone pairs are missing; or Lewis structures correct but square brackets and charge missing around NO₃⁻ | 5 |
| Octet around central atom is correctly shown (i.e. bonds are correct) for all structures but lone pairs are missing; and Lewis structures correct but square brackets and charge missing around NO₃⁻ | 4 |
| Award 3 marks if each structure has the correct number of valence electrons shown but not correctly distributed | 3 |
| Incorrect | 0 |
| Total | 8 |

NB: Lewis structure does not need to represent the shape.

GE 3

Question 29 (9 marks)

5

Ethanol C₂H₅OH, and hexane, C₆H₁₄, are two common industrial solvents.

(a) (i) Identify the polarity of molecules that are soluble in each of these solvents. (2 marks)

| | Polarity of molecules/substances that are soluble in the solvent | |
|---------|------------------------------------------------------------------|-------------|
| | (circle | e one only) |
| Ethanol | Polar | Non-polar |
| | (circle | e one only) |
| Hexane | Polar | Non-polar |

| Description | Marks |
|-----------------------------------------------------------|-------|
| Recognition that ethanol will dissolve polar molecules | 1 |
| Recognition that hexane will dissolve non-polar molecules | 1 |
| Incorrect | 0 |
| Total | 2 |

(ii) Explain the interactions between solute and solvent particles in solutions of these two solvents. (3 marks)

| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------------|-------|
| Recognition that: | |
| solutions using ethanol is a result of dipole/dipole or hydrogen bond interactions between solute and solvent | 1 |
| solutions using hexane is a result of dispersion forces between solute and solvent | 1 |
| interactions between solute/solvent must be of similar strength to the solute/solute and solvent/solvent interactions | 1 |
| Incorrect | 0 |
| Total | 3 |

Question 29 (continued)

(b) The 'loss' of 4 mL on mixing water with ethanol implies strong interactions between the water and ethanol molecules. Describe these interactions and explain the origin of their strength. (4 marks)

| Description | Marks |
|--------------------------------------------------------------------------------|-------|
| Recognition that the interactions between water and ethanol are of | 1 |
| similar strength to those in the individual liquids | |
| Recognition that water and ethanol have hydrogen bonding | 1 |
| Explains hydrogen bonding in terms of: | |
| oxygen being extremely electronegative leading to the high | 1 |
| polarity of the O-H bond or | 1 |
| a big difference in electronegativity of O and H | |
| Explains the strength of the H-bond (any one of the following) | |
| highly polar | |
| high charge density of H | 1 |
| low electron density | |
| appropriate annotated diagram | |
| Incorrect | 0 |
| Total | 4 |

Question 30 (9 marks)

Hydrogen can be made by reacting methane (natural gas) with water (steam). The reaction can form the chemical equilibrium represented below.

$$CH_4(g) + H_2O(g) \longrightarrow 3 H_2(g) + CO(g) \Delta H = +206 \text{ kJ mol}^{-1}$$

State the conditions of temperature and pressure that would optimise the yield of hydrogen at a reasonable rate of reaction. Using collision theory and principles of chemical equilibrium, explain your choice of conditions.

| | Optimum conditions | Explanation |
|-------------|--------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Temperature | high temperature | High temperature increases the proportion of molecules colliding with energy above the E _a and so increasing the reaction rates for both the forward and reverse reactions but the (forward) endothermic direction will increase more so increasing yield of H ₂ . |
| | (1 mark) | (3 marks) |
| Pressure | moderate pressure | High pressure increases frequency of collisions between molecules and increases rates for both the forward and reverse reactions but increases reverse reaction rate more (because there are fewer gas molecules on reactant side). Low pressure will increase yield of H ₂ but the rate of the reaction will be too slow so a compromise moderate pressure is needed. |
| | (1 mark) | (4 marks) |

| Description | Marks |
|-----------------------------------------------------------------------------------------|-------|
| Recognition of high temperature and moderate pressure | 1–2 |
| Recognition that high temperature increases the proportion of molecules | 1 |
| colliding with energy above the E _a | ı |
| Recognition that high temperature increases rates of forward (and reverse) | 1 |
| reaction(s) | ı |
| Recognition that high temperature increases the rate of forward reaction | |
| more than rate of reverse reaction or | 1 |
| accept Le Chatelier's Principle explanation | |
| Recognition that high pressure increases frequency of collisions between | 1 |
| molecules | 1 |
| Recognition that high pressure increases rates of both forward and reverse | 1 |
| reactions | |
| Recognition that high pressure increases rate of reverse reaction more than | 1 |
| forward | 1 |
| Recognition that low pressure will increase yield of H ₂ but the rate of the | 1 |
| reaction will be too slow. Compromise between yield and reaction rate | 1 |
| Incorrect | 0 |
| Total | 9 |

NB: Where the student chooses the incorrect optimum conditions but their explanation matches the marking key, they may be awarded marks for the explanation.

Question 31 (5 marks)

(a) State the role of the standard hydrogen half-cell in determining the table of Standard Reduction Potentials. (2 marks)

| Description | Marks |
|----------------------------------------------------------------------------|-------|
| Recognition that the hydrogen half-cell: | |
| is the reference half-cell | 1 |
| against which the reduction potential of all other half-cells are measured | 1 |
| Incorrect | 0 |
| Total | 2 |

(b) State **three** limitations of Standard Reduction Potential tables.

(3 marks)

| Description | Marks |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Any three of: The values of E⁰ depend upon concentration. (1 mol L⁻¹) Applies only to aqueous solutions The emf of a cell can depend on temperatures The values of E⁰ give no indication of reaction rate/high activation energy Predictive tool – reaction may not occur All gases at 100kPa (standard pressure) | 1–3 |
| Incorrect | 0 |
| Total | 3 |

Question 32 (6 marks)

Nitrogen gas from the atmosphere undergoes a series of redox reactions to transform it into nitrate ions that are absorbed by plants. The process can be simplified into the following three steps.

Step 1 – Nitrogen-fixing soil bacteria reduce nitrogen gas to ammonium ions.

Step 2 – Nitrifying bacteria then oxidise ammonium ions to nitrite ions.

Step 3 – Nitrifying bacteria then oxidise nitrite ions to nitrate ions.

Write the half-equations for each of these steps. Assume acidic conditions.

$$N_2 + 8 H^+ + 6 e^- \rightarrow 2 NH_4^+$$
 $NH_4^+ + 2 H_2 O \rightarrow NO_2^- + 8 H^+ + 6 e^-$

$$NO_2^- + H_2O \rightarrow NO_3^- + 2 H^+ + 2 e^-$$

| Description | Marks |
|-------------------------------------|-------|
| 2 marks for a correct half-equation | 0–6 |
| Incorrect | 0 |
| Total | 6 |

NB: For an incorrect half-equation, award 1 mark if reactants and products are correct but electrons or balancing incorrect; if atoms are consistently balanced for all the half-equations but electrons incorrect award 4 marks.

Question 33 (9 marks)

(a) Methanoic acid, HCOOH, may be produced by oxidation of an alcohol with acidified potassium permanganate, MnO₄⁻, solution.

Write the oxidation and reduction half-equations and the final redox equation for this reaction. (5 marks)

| Oxidation half-equation | $CH_3OH(\ell) + H_2O(\ell) \rightarrow HCOOH(\ell) + 4 H^+(aq) + 4 e^- (\times 5)$ |
|-------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Reduction half-equation | $MnO_4^-(aq) + 8 H^+(aq) + 5 e^- \rightarrow Mn^{2+}(aq) + 4 H_2O(\ell)$ (×4) |
| Final redox equation | $5 \text{ CH}_3\text{OH}(\ell) + 4 \text{ MnO}_4^-(\text{aq}) + 12 \text{ H}^+(\text{aq}) \rightarrow 5 \text{ HCOOH}(\ell) + 4 \text{ Mn}^{2+}(\text{aq}) + 11 \text{ H}_2\text{O}(\ell)$ |

| Description | Marks |
|----------------------------------------------------------------------|-------|
| Correct oxidation half-equation; for an incorrect oxidation half- | |
| equation, award 1 mark if reactants and products are correct but | 1–2 |
| electrons or balancing incorrect | |
| Correct reduction half-equation | 1 |
| Correct final redox equation; for an incorrect redox equation, award | 1–2 |
| 1 mark if reactants and products are correct but balancing incorrect | 1–2 |
| Incorrect | 0 |
| Total | 5 |

NB: State symbols not required.

If oxidation/reduction reversed award a maximum of 4 marks.

Balancing includes cancelling of H⁺ and H₂O

(b) Methanoic acid reacts with ethanol in the presence of sulfuric acid to produce a sweet smelling compound.

Write the balanced equation for the reaction of methanoic acid with ethanol. (2 marks)

| Description | | Marks |
|------------------------------------------|-------|-------|
| Correct equation | | 2 |
| Only 3 formulae correct or water missing | | 1 |
| Incorrect | | 0 |
| | Total | 2 |

(c) Draw the structural formula for the sweet smelling compound and give its IUPAC name. Show **all** H atoms in the structure. O (2 marks)

Name: ethyl methanoate

| Description | Marks |
|----------------------------------------------------|-------|
| Correct structure (condensed structure acceptable) | 1 |
| Correct name – ethyl methanoate | 1 |
| Incorrect | 0 |
| Total | 2 |

Question 34 (5 marks)

Alcohols can be classified as primary, secondary or tertiary.

Complete the table below by drawing the structure for a primary alcohol, a secondary alcohol and a tertiary alcohol, each with the molecular formula $C_5H_{12}O$. Show **all** H atoms in your structures.

Give the IUPAC names for the primary and secondary alcohols you have drawn.

| | Structure | Name |
|----------------------|-------------------------|--------------------------------------------------------------|
| Primary alcohol | H H H H H | Pentan-1-ol 2-methylbutan-1-ol 2,2-dimethylpropan-1-ol |
| Secondary alcohol | H H H OH H | Pentan-2-ol Pentan-3-ol 3-methylbutan-2-ol |
| Tertiary alcohol | H H CH ₃ | Name not required. |

| Description | Marks |
|--------------------------------------------------------------------------------------|-------|
| A correct primary alcohol structure with MF C ₅ H ₁₂ O drawn | 1 |
| Name matches alcohol structure drawn | 1 |
| A correct secondary alcohol structure with MF C ₅ H ₁₂ O drawn | 1 |
| Name matches alcohol structure drawn | 1 |
| Correct tertiary alcohol structure with MF C ₅ H ₁₂ O drawn | 1 |
| Incorrect | 0 |
| Total | 5 |

NB: Award 2 of the possible 3 marks for structures if all H atoms not shown; condensed structures acceptable.

Question 35 (4 marks)

(a) The structure below represents a segment of polyacrylic acid.

Draw the structure for the monomer of this addition polymer.

(1 mark)

| Description | Marks |
|-------------------|-------|
| Correct structure | 1 |
| Incorrect | 0 |
| Total | 1 |

(b) (i) What is the name of the interactions occurring between water molecules and sodium ions to enable the removal of the latter from the polymer? (1 mark)

| Description | Marks |
|-----------------------------------------------------------------------------|-------|
| The water molecules and sodium ions interact through ion-dipole attractions | 1 |
| Incorrect | 0 |
| Total | 1 |

(ii) Explain how the polymer sodium polyacrylate can absorb large quantities of water. (2 marks)

| Description | Marks |
|-----------------------------------------------------------------|-------|
| Recognition that water molecules form hydrogen bonds with the | 1 |
| carboxylate groups (accept ion-dipole) | ı |
| Recognition that there are a large number of carboxylate groups | |
| in the polymer so a large number of water molecules can | 1 |
| hydrogen bond to the polymer | |
| Incorrect | 0 |
| Total | 2 |

Question 36 (9 marks)

(a) What is the independent variable in this investigation?

(1 mark)

| Description | Marks |
|---------------------------------------------------------------------------------------------------|-------|
| Concentrations of solution (accept concentration of one of Cu ²⁺ or Zn ²⁺) | 1 |
| Incorrect | 0 |
| Total | 1 |

(b) What is the dependent variable in this investigation?

(1 mark)

| Description | Marks |
|------------------------------------|-------|
| Electrical potential/voltage/volts | 1 |
| Incorrect | 0 |
| Total | 1 |

(c) Why did the volumes and temperatures of solutions and surface areas of the electrodes need to be the same in each trial? (1 mark)

| Description | Marks |
|------------------------------------------------------------------------------------------------|-------|
| To be confident that any changes in electrical potential are due to concentration changes only | 1 |
| Incorrect | 0 |
| Total | 1 |

(d) Explain the increase in electrical potential as the concentration of Cu²⁺ ions increased and the decrease in electrical potential as the concentration of Zn²⁺ ions increased.

(2 marks)

| Description | Marks |
|-------------------------------------------------------------------------------------------------------------------------------|-------|
| The rate of forward reaction increases as concentration of Cu ²⁺ ions increased (Accept forward reaction favoured) | 1 |
| The rate of reverse reaction increases as concentration of Zn ²⁺ ions increased (Accept reverse reaction favoured) | 1 |
| Incorrect | 0 |
| Total | 2 |

(e) The student also observed that as the cells were allowed to run for a while their electrical potential slowly decreased from its maximum value. Why did this happen? (2 marks)

| Description | Marks |
|------------------------------------------------------------------------------------------|-------|
| Recognition that as the cell operates the concentration of reactants decreases | 1 |
| Recognition that forward reaction rate decreases or system approaches equilibrium | 1 |
| Incorrect | 0 |
| Total | 2 |

The student concluded:

'As the concentration of the oxidant increases, so does the cell voltage (electrical potential).'

(f) List **two** ways to improve the investigation.

(2 marks)

| Description | Marks |
|------------------|-------|
| Test other cells | 1 |
| Repeat trials | 1 |
| Incorrect | 0 |
| Total | 2 |

STAGE 3

Section Three: Extended answer 40% (82 Marks)

Question 37 (9 marks)

Aqua regia is a mixture of concentrated hydrochloric acid and nitric acid that is able to dissolve gold. One of its uses is in the analysis of gold content in gold ore.

As part of quality control processes, a chemist in a gold analysis laboratory analysed aqua regia to ensure the required 3:1 mole ratio of hydrochloric acid to nitric acid. The chemist found that 20.0 mL of agua regia needed 28.6 mL of 8.00 mol L⁻¹ sodium hydroxide for complete neutralisation. The reaction for the neutralisation reaction between the sodium hydroxide and acid is represented by the equation below:

$$H^{+}(aq) + OH^{-}(aq) \rightarrow H_2O(\ell).$$

(a) Calculate the moles of hydrogen ions present in the 20.0 mL sample of agua regia. (2 marks)

| Description | Marks |
|--------------------------------------------|-------|
| $n(NaOH) = 0.0286 \times 8.00$ | 4 |
| = 0.2288 mol | ı |
| $n(H^{+}) = n(OH^{-}) = 0.229 \text{ mol}$ | 1 |
| Incorrect | 0 |
| Total | 2 |

The chemist analysed the chloride content of the agua regia by adding excess silver nitrate solution to a separate 20.0 mL sample of aqua regia. This resulted in the precipitation of 24.6 g of solid.

Write the balanced ionic equation for precipitation of silver chloride from agua regia. (b) (1 mark)

$$Ag^{+}(aq) + C\ell^{-}(aq) \rightarrow AgC\ell(s)$$

| Description | Marks |
|-----------------------------|-------|
| Correctly balanced equation | 1 |
| Incorrect | 0 |
| Total | 1 |

NB: State symbols not required.

Question 37 (continued)

(c) Calculate the moles of hydrochloric acid in the 20.0 mL of aqua regia. (3 marks)

| Description | Marks |
|-----------------------------------------------------------|-------|
| $M(AgC\ell) = 143.35 \text{ g mol}^{-1}$ | 1 |
| $n(AgC\ell) = \frac{24.6}{143.35} = 0.1716 \text{ mol}$ | 1 |
| $n(HC\ell) = n(C\ell^-) = n(AgC\ell) = 0.172 \text{ mol}$ | 1 |
| Incorrect | 0 |
| Total | 3 |

(d) Determine whether the aqua regia had the required ratio of hydrochloric acid to nitric acid. State clearly whether the ratio was as required and support your answer with clear workings. (3 marks)

| Description | Marks |
|---------------------------------------------------------------------|-------|
| $n(HNO_3) = n(H^+)total - n(HC\ell) = 0.2288 - 0.1716 = 0.0572 mol$ | 1 |
| Ratio = $\frac{n(HC\ell)}{n(HNO_3)} = \frac{0.1716}{0.0572} = 3.00$ | 1 |
| Yes the ratio of HCℓ to HNO₃ is 3:1 | 1 |
| Incorrect | 0 |
| Total | 3 |

Question 38 (12 marks)

(a) State and explain the trend in atomic radius across the Second Period. (3 marks)

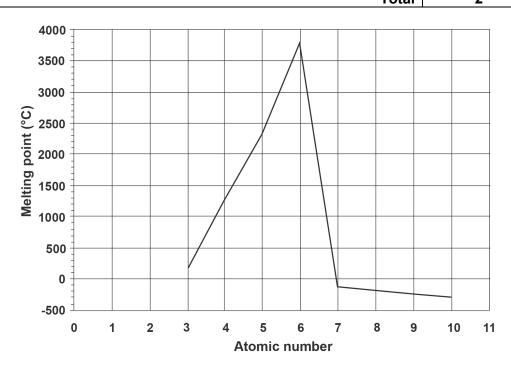
| Description | Marks |
|------------------------------------------------------------------------------------------------------|-------|
| Recognition that the atomic radius decreases across the period | 1 |
| Recognition that each successive electron is in the same valence shell | 1 |
| Recognition that increasing positive charge of the nucleus pulls the electrons closer to the nucleus | 1 |
| Incorrect | 0 |
| Total | 3 |

(b) State and explain the trend in first ionisation energy across the Second Period. (4 marks)

| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Recognition that the first ionisation energy increases across the period | 1 |
| Recognition that the valence shell is closer to nucleus (radius reduced) across the period | 1 |
| Recognition that the positive charge increases in nucleus. | 1 |
| Recognition that with increasing positive charge of the nucleus there is an increase in the force of attraction and more energy is needed to remove an electron | 1 |
| Incorrect | 0 |
| Total | 4 |

(c) Plot a graph of atomic number against melting point for the Second Period elements. (2 marks)

| Description | Marks |
|---------------------------------|--------|
| Accurate plotting of points | 1 |
| Straight line connecting points | 1 |
| Incorrect | 0 |
| Tr | otal 2 |



Question 38 (continued)

(d) Based on their bonding when solid, explain the difference in melting points of lithium, carbon and neon. (3 marks)

| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Recognition that as a metal lithium has a moderately high melting point due to moderate forces of attraction between cations and delocalised electrons | 1 |
| Recognition that carbon's very high melting point can be explained by its strong covalent bonding between all carbon atoms in the network structure | 1 |
| Recognition that neon's low melting point can be explained by the weak dispersion forces between neighbouring neon atoms (so only small amounts of energy are needed to overcome these attractive forces) | 1 |
| Incorrect | 0 |
| Total | 3 |

Question 39 (14 marks)

An organic compound that contains only carbon, hydrogen, oxygen and bromine, was analysed to determine its empirical formula. A combustion analysis of 1.50 g of the compound produced 1.58 g of carbon dioxide and 0.563 g water.

On treatment of 1.75 g of the compound to convert the bromine in the compound to bromide ions and further reaction with silver nitrate, 1.97 g of silver bromide was precipitated.

(a) Determine the empirical formula of the compound.

(10 marks)

| | Desc | ription | | | Marks |
|----------------------------------------------------------------------------------------|---------------|---------------------|-----------------------|---------------|-------|
| m(C) = $1.58 \times \frac{12.01}{44.01} =$ %(C) = $\frac{0.431}{1.50} \times 100 =$ | | | | | 1–2 |
| $m(H) = 0.563 \times \frac{2.016}{18.016}$ | - = 0.0630 g | | | | |
| $\%(H) = \frac{0.0630}{1.50} \times 100$ | | | | | 1–2 |
| $m(Br) = 1.97 \times \frac{79.7}{79.9 + 1}$ $\%(Br) = \frac{0.838}{1.75} \times 100\%$ | 07.0 | 3 g | | | 1–2 |
| % O = 100 - 28.7 - 4 | .2 – 47.9 = 1 | 9.2 | | | 1 |
| | С | Н | Br | 0 | |
| Ratio by mass | 28.7 | 4.20 | 47.9 | 19.2 | |
| Ratio by mol | 28.7 12.01 | 4.20 1.008 | 47.9 79.9 | 19.2 16 | |
| | 2.39 | 4.17 | 0.599 | 1.20 | 1–3 |
| Divide by smallest | 2.39 0.599 | 4.17 0.599 | $\frac{0.599}{0.599}$ | 1.20 0.599 | . • |
| | 3.98 | 6.96 | 1 | 2.00 | |
| Therefore empirical for | ormula is C₄⊦ | H ₇ BrO₂ | | | |
| | | | | Total | 10 |

Question 39 (continued)

(b) 1.95 g of the compound was vaporised and was found to occupy 0.387 L at 95.0 kPa and 105 °C. Determine the molecular formula of the compound. (2 marks)

| Description | Marks |
|-------------------------------------------------------------------------------------------|-------|
| $n = \frac{95 \times 0.387}{8.314 \times 378.15} = 0.0117 \text{ mol}$ | |
| $M = \frac{1.95}{0.0117} = 167$ | 1 |
| $EFM(C_4H_7BrO_2) = (4 \times 12.01) + (7 \times 1.008) + 79.9 + (2 \times 16) = 166.996$ | |
| Since empirical mass = molecular mass, the molecular formula is $C_4H_7BrO_2$ | 1 |
| Total | 2 |

(c) Further analysis of the organic compound revealed that it had a carboxylic acid functional group. Draw a possible structural formula of the organic compound. (2 marks)

| Description | Marks |
|--------------------------------------------------------------------------------------------------------------------------------|-------|
| CH ₂ BrCH ₂ COOH or CH ₃ CHBrCH ₂ COOH or CH ₃ CH ₂ CHBrCOOH | 1–2 |
| Total | 2 |

NB: If molecule drawn with carboxylic acid shown with one minor error, 1 mark. A correct structure is drawn for an incorrect molecular formula, if it represents a carboxylic acid, 2 marks.

Accept a correct bond-line structure, 2 marks

Question 40 (15 marks)

(a) Explain why sodium hydroxide is not suitable as a primary standard (2 marks)

| Description | Marks |
|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Any two of the following: | |
| does not have high molar mass absorbs moisture/is deliquescent/hygroscopic reacts with CO₂ from the atmosphere mass varies over time cannot be obtained pure | 1-2 |
| Incorrect | 0 |
| Total | 2 |

(b) Show that the concentration of the sodium hydroxide solution is $0.0916 \text{ mol } L^{-1}$. Show sufficient workings to justify your answer. (3 marks)

| Description | Marks |
|------------------------------------------------------------------------------------------------------|-------|
| $n(HC\ell) = 0.01745 \times 0.105 = 1.832 \times 10^{-3} \text{ mol}$ | 1 |
| $n(NaOH) = n(HC\ell) = 1.832 \times 10^{-3} \text{ mol}$ | 1 |
| c(NaOH) = $\frac{n}{v} = \frac{1.832 \times 10^{-3}}{0.02} = 9.16 \times 10^{-2} \text{ mol L}^{-1}$ | 1 |
| Incorrect | 0 |
| Total | 3 |

(c) Calculate the average titre volume to be used in the calculation of the citric acid content. (2 marks)

| Description | Marks |
|------------------------------------------------------------------------|-------|
| Differences in initial and final readings = 21.80, 20.85, 20.90, 20.95 | 1 |
| Titre volume = $\frac{20.85 + 20.90 + 20.95}{3}$ = 20.90 mL | 1 |
| Incorrect | 0 |
| Total | 2 |

Question 40 (continued)

(d) Given that citric acid ($C_6H_8O_7$) is a weak triprotic acid, determine the percentage composition by mass of citric acid in the cleaner. The molar mass of citric acid is 192.124 g mol⁻¹. (6 marks)

| Description | Marks |
|-------------------------------------------------------------------------------------------------------------|-------|
| $n(NaOH) = 0.02090 \times 0.0916 = 1.914 \times 10^{-3} \text{ mol}$ | 1 |
| In 20 mL of dilute citric acid, n(citric) = $\frac{1.914 \times 10^{-3}}{3}$ = 6.381 × 10 ⁻⁴ mol | 1 |
| n(citric) in 100 mL = $6.381 \times 10^{-4} \times 5 = 0.003191$ mol | 1 |
| hence in 10 mL original = 0.003191 mol | 1 |
| $m(citric) = n \times M = 0.003191 \times 192.124 = 0.613 g$ | 1 |
| Therefore % composition = $\frac{0.613}{10.4} \times 100 = 5.89\%$ | 1 |
| Incorrect | 0 |
| Total | 6 |

(e) Select a suitable indicator for this titration from the table below. Explain your choice. (2 marks)

| Indicator | Colour change (low pH – high pH) | pH range |
|------------------|-------------------------------------|-----------|
| Methyl yellow | red-yellow | 2.4 - 4.0 |
| Litmus | red-blue | 5.0 - 8.0 |
| Bromothymol blue | yellow-blue | 6.0 – 7.6 |
| Thymol blue | Yellow-blue | 8.0 – 9.6 |

| Description | Marks |
|---------------------------------------------------------------------------------------------------------------------------|-------|
| Thymol blue | 1 |
| The citrate ion hydrolyses to give hydroxide ions and so an equivalence point in the basic region or appropriate equation | 1 |
| Incorrect | 0 |
| Total | 2 |

Question 41 (24 marks)

(a) Write the balanced equation for the hydrolysis of urea.

(2 marks)

$$CO(NH_2)_2(aq) \quad + \quad H_2O(\ell) \quad \rightarrow \quad CO_2(aq) \quad + \quad 2 \; NH_3(aq)$$

| Description | Marks |
|---------------------------------------------|-------|
| Correct formulae for reactants and products | 1 |
| Balanced equation | 1 |
| Incorrect | 0 |
| Total | 2 |

NB: States for reactants and products not required.

(b) Explain briefly why hydrolysis of urea causes an **increase** in pH. Include an appropriate balanced equation in your answer. (2 marks)

| Description | Marks |
|---------------------------------------------------------------------------------------|-------|
| Statement showing recognition that NH ₃ hydrolyses to give OH ⁻ | 1 |
| Balanced hydrolysis equation | 1 |
| Incorrect | 0 |
| Total | 2 |

NB: States for reactants and products not required.

The pH increases because the ammonia produced by hydrolysis of urea in turn hydrolyses to produce hydroxide ions.

$$NH_3(aq) + H_2O(\ell)$$
 \longrightarrow $NH_4^+(aq) + OH^-(aq)$

(c) Determine the concentration, in grams per litre, of the phosphorus in the urine. Express your answer to **three** significant figures. (Assume all phosphorus has been precipitated.) (9 marks)

Molar masses (in g mol⁻¹): struvite 245.418; calcium hydroxyapatite 1004.636.

| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| $m(MgNH_4PO_4.6H_2O)$ in the precipitate = 0.823 × 25.3 = 20.823 g | 1 |
| $m(Ca_{10}(PO_4)_6(OH)_2)$ in the precipitate = 25.3 - 20.823 = 4.478 g | 1 |
| n(P) in struvite = n(MgNH ₄ PO ₄ .6H ₂ O) = $\frac{20.823}{245.418}$ = 8.484×10 ⁻² mol | 1 |
| $n(Ca_{10}(PO_4)_6(OH)_2) = \frac{4.478}{1004.636} = 4.457 \times 10^{-3} \text{ mol}$ | 1 |
| n(P) in calcium hydroxyapatite = $4.457 \times 10^{-3} \times 6 = 2.674 \times 10^{-2}$ mol Total n(P) = $8.484 \times 10^{-2} + 2.674 \times 10^{-2} = 1.116 \times 10^{-1}$ mol | 1 |
| | 1 |
| $m(P) = 1.116 \times 10^{-1} \times 30.97 = 3.456 g$ | 1 |
| Concentration = $\frac{3.456}{5.00}$ = 0.691 g L ⁻¹ | 1 |
| Answer expressed in three significant figures | 1 |
| Incorrect | 0 |
| Total | 9 |

(d) Explain, using collision theory, why increasing the pH of the phosphorus-depleted urine converts the ammonium ions to ammonia. Support your answer with a balanced equation. (3 marks)

| Description | Marks |
|---------------------------------------------------------------------------------------|-------|
| Recognition that concentration of OH ⁻ ions increases so the frequency | 1 |
| of collisions between NH ₄ ⁺ and OH ⁻ ions increases | 1 |
| Recognition that rate of the forward reaction in the equilibrium | 1 |
| increases relative to the reverse reaction | |
| Balanced equation | 1 |
| Incorrect | 0 |
| Total | 3 |

NB: Increased pH increases concentration of OH⁻ so the frequency of collisions between NH₄⁺ and OH⁻ ions increases and thus the rate of the forward reaction in the equilibrium below increases relative to the reverse reaction and so increases production of ammonia. (The increase in hydroxide ion concentration shifts the equilibrium below to the production of ammonia.)

$$NH_4^+(aq) + OH^-(aq)$$
 \longrightarrow $NH_3(aq) + H_2O(\ell)$

(e) State the purpose of heating the solution to 40 °C.

(1 mark)

The solubility of NH₃ in water is reduced at high temperature.

| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| Statement showing recognition that solubility of NH_3 in water is reduced at high temperature; Accept $NH_3(aq) \rightarrow NH_3(g)$ is endothermic so increasing temperature favours the endothermic process and equilibrium shifts right producing more $NH_3(g)$ | 1 |
| Incorrect | 0 |
| Total | 1 |

(f) Write a balanced equation for the reaction in Step 3.

(2 marks)

Accept any of the following:

| Description | | Marks |
|---------------------------------------------|-------|-------|
| Correct formulae for reactants and products | | 1 |
| Balanced equation | | 1 |
| Incorrect | | 0 |
| | Total | 2 |

Question 41 (continued)

STAGE 3

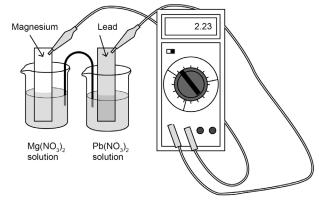
(g) The amount of ammonium sulfate recovered from the 5.00 L of urine in this experiment is 72.65 g. If the process is 78% efficient, what is the concentration of nitrogen, in grams per litre, in the phosphorus-depleted urine? (5 marks)

| Description | Marks |
|--------------------------------------------------------------------------|-------|
| $n((NH_4)_2SO_4) = \frac{72.65}{132.144} = 0.5498 \text{ mol}$ | 1 |
| $n(N) = 2 \times n((NH_4)_2SO_4) = 2 \times 0.5498 = 1.0996 \text{ mol}$ | 1 |
| For 78% efficient, m(N) = 1.0996 × 14.01 = 15.405 g | 1 |
| For 100%, $m(N) = \frac{15.405}{0.78} = 19.75 g$ | 1 |
| Concentration = $\frac{19.75}{5.00}$ = 3.95 g L ⁻¹ | 1 |
| Incorrect | 0 |
| Total | 5 |

Question 42 (8 marks)

Using the diagram below, explain the role of the following in the operation of an electrochemical (galvanic) cell:

- anode process
- cathode process
- lead(II) nitrate electrolyte
- salt bridge and ion migration
- electron flow in external circuit.



| Description | Marks |
|-----------------------------------------------------------------------------------------------------------------|-------|
| Anode oxidation occurs | 1 |
| Mg loses electrons to form Mg ions or equation for reaction at the anode Mg \rightarrow Mg ²⁺ + 2e | 1 |
| Cathode reduction occurs | 1 |
| Pb gains electrons to form Pb or equation for reaction at the cathode Pb ²⁺ + 2e ⁻ → Pb | 1 |
| Lead(II) nitrate electrolyte is the source of Pb ²⁺ ions | 1 |
| Salt bridge allows for the movement of ions between the two half-cells | 1 |
| Salt bridge allows electrical neutrality to be maintained or to complete the circuit | 1 |
| Electron follow provides energy to do work e.g. produces a reading on the meter | 1 |
| Total | 8 |

ACKNOWLEDGEMENTS

Section Three

Question 42

Adapted from: Megna, R. *Electrochemical cell* [Image]. Retrieved June 23, 2014, from http://fphoto.photoshelter.com/search?I_DSC= 45859901-2RM&_ACT=search

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