



A LEVEL

Examiners' report

CHEMISTRY B (SALTERS)

H433

For first teaching in 2015

H433/03 Summer 2022 series



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Introduction

Our examiners' reports are produced to offer constructive feedback on candidates' performance in the examinations. They provide useful guidance for future candidates.

The reports will include a general commentary on candidates' performance, identify technical aspects examined in the questions and highlight good performance and where performance could be improved. A selection of candidate answers are also provided. The reports will also explain aspects which caused difficulty and why the difficulties arose, whether through a lack of knowledge, poor examination technique, or any other identifiable and explainable reason.

Where overall performance on a question/question part was considered good, with no particular areas to highlight, these questions have not been included in the report.

A full copy of the question paper and the mark scheme can be downloaded from OCR.

Advance Information for Summer 2022 assessments

To support student revision, advance information was published about the focus of exams for Summer 2022 assessments. Advance information was available for most GCSE, AS and A Level subjects, Core Maths, FSMQ, and Cambridge Nationals Information Technologies. You can find more information on our <u>website</u>.

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Paper 3 series overview

H433/03 is one of the three examination units for GCE Chemistry B. This largely synoptic unit links together different areas of chemistry within different contexts, some practical, some familiar and some novel. To do well on this paper, candidates need to be comfortable applying their knowledge and understanding to unfamiliar contexts and be familiar with a range of practical techniques.

H433/03 is much more application based that the other two A level Chemistry Components, H433/01 and H433/02, which have a greater emphasis on knowledge and understanding of the assessment outcomes from the specification.

H433/03 also contains more questions set in a practical context, including an insert based on a practical procedure, than H433/01 and H433/02.

This particular exam series, coming in the wake of the COVID outbreak and the difficulty centres and centres experienced in being able to deliver hands on practical activities, resulted in many students providing less structured answers to those questions requiring a knowledge of practical procedure or analysis of practical observations.

Examiners were well aware of this and, while striving to maintain consistent standards of assessment, actively tried to give marks where possible.

Candidates who did well on this paper generally did the following:	Candidates who did less well on this paper generally did the following:
 Recognised that the detail required in longer text based answers was commensurate with the marks available. Carefully laid out the steps involved in calculations; thus minimising marks lost on any slip made in the calculation Linked the procedure or observations in questions involving practical investigations to the important underlying chemistry 	 Gave explanations for the longer answer questions which were not logically sequenced and missed fine detail Struggled to link essential chemistry to practical observations Gave a less successful or indeed no explanation for those questions traditionally regarded as more challenging, e.g. reaction mechanisms and/or electrochemistry

Question 1 (a) (i)

- 1 This question is about haloalkanes and their reactivity.
 - (a) Haloalkanes are useful intermediates for preparing a range of organic compounds.

A flowchart showing the synthesis of three organic compounds is shown below.



(i) Complete the flowchart showing the structures of compounds A and C.

On the dotted line show the reagent needed to form compound **B**.

[3]

There were many good responses. The structure of compound C, a ketone, proved the most difficult.

Question 1 (a) (ii)

(ii) Which homologous series does compound **B** belong to?

.....[1]

The majority of answers were correct. Wrong choices included amides and amino acids.

Question 1 (b)

(b) Compound **B** behaves as a base.

Complete the diagram of a mechanism that shows how a molecule of compound **B** reacts with water to give a basic solution.

Show 'curly arrows'.

[2]

This question produced a wide range of answers. The 'curly arrows', as ever proved difficult for many candidates. The lack of a positive charge on the alkyl ammonium cation product caught many students out. This question proved discriminating.

Question 1 (c)

(c) Some haloalkanes act as greenhouse gases that absorb infrared radiation in the troposphere.

Give the source of this infrared radiation.

......[1]

Answers often polarised between 'Earth' (correct) or 'Sun' (incorrect).

Question 1 (d)*

(d)* A student wishes to compare the relative reactivities of haloalkanes.

The student investigates the rates of hydrolysis of 1-chlorobutane, 1-bromobutane and 1-iodobutane by following the reaction in the presence of aqueous silver ions.

Design a suitable procedure that the student should follow.

Give the expected results and how they can be interpreted.

[6]

A majority of students knew that key to this investigation was the formation of a silver halide precipitate, but struggled to put together a logical, structured procedure. There were also polarised ideas about the order of precipitation and so interpretation. A significant minority of candidates wrongly linked the reactivity of the haloalkane to that of the halogen. The best answers also included an equation the hydrolysis reaction as well as for the ionic precipitation reaction.

Question 2 (a) (i)

- 2 This question is about the enthalpy change of the reaction of zinc with copper(II) sulfate solution.
 - (a) A student investigates the temperature change when Reaction 2.1 occurs.

 $CuSO_4(aq) + Zn(s) \rightarrow Cu(s) + ZnSO_4(aq)$ Reaction 2.1

(i) The student adds 1.20 g of powdered zinc to 50.0 cm³ of 0.200 mol dm⁻³ copper(II) sulfate solution in a glass beaker.

Use calculations to show which is the limiting reagent.

limiting reagent is[2]

Well answered by the majority of candidates.

Question 2 (a) (ii)

(ii) The student measures the temperature of the contents of the beaker over a period of 20 minutes and plots the data on the graph below.



The student uses the graph to determine the highest temperature as shown.

The student uses this temperature to calculate the heat energy given to the solution per mole of limiting reagent.

Show the student's calculation, giving the answer in $kJ mol^{-1}$.

Give your answer to an **appropriate** number of significant figures.

heat energy given to the solution = \dots kJmol⁻¹ [3]

This question discriminated well. Common mistakes included the use of the wrong mass for the solution and a failure to calculate the heat energy for one mole of 'reaction'. Examiners did allow ecf marks for individual errors correctly carried forward.

Question 2 (b)

(b) Calculate $\Delta_r H^{\oplus}$ for **Reaction 2.1** using the data shown below.

	Δ _f H [◆] / kJ mol ^{−1}
Cu ²⁺ (aq)	+ 64.4
Zn ²⁺ (aq)	-152.4

 $\Delta_r H^{\oplus} = \dots kJ \operatorname{mol}^{-1} [1]$

A pleasing majority of candidates calculated this value correctly; however some lost the mark by not including the negative sign.

Question 2 (c)*

(c)* The value obtained from the student's experiment is considerably less exothermic than the value in part (b).

Evaluate the student's experiment, identifying limitations in both the experimental procedure and the measurements taken. You should also comment on how any of the limitations you have identified will affect the final value.

Suggest possible improvements to the procedure, apparatus and measurements.

The vast majority of candidate's knew that heat loss to the environment was the most significant factor causing the final value to be lower than the theoretical. However the stem of the question requires candidate answers to not only identify limitations in both the procedure and the measurements but also suggest possible improvements to the procedure, apparatus and measurements in order to be given the higher marks for this question. The majority of candidates just mentioned heat loss to the environment and the use of polystyrene cups, without considering any other, albeit possibly less significant, factors.

Assessment for learning

In questions such as 2(c) students should be encouraged to consider/interrogate in detail the key formula at the heart of the practical, $q = m x c x \Delta T$.

It might be helpful to ask students what each of the symbols actually mean. e.g. 'q' is the heat energy in Joules transferred in the reaction – is this per mole of reaction?

m is mass/g – of what? Water, solution, zinc? **c** is specific heat capacity – water or solution or even container (glass/polystyrene)? Δ **T** is temperature <u>change</u> (conversion to Kelvin scale makes no difference) – has the graph been used in the best way [6]

Question 3 (a) (i)

3 This question is about redox and electrochemical cells.

Table 3.1 shows standard electrode potentials, some of which will be needed for the rest of this question.

Half reaction	Standard electrode potential, <i>E</i> ⁺ /V
Zn ²⁺ (aq) + 2e [−] ← Zn(s)	-0.76
$Fe^{2+}(aq) + 2e^{-} \rightleftharpoons Fe(s)$	-0.44
$2H^+(aq) + 2e^- \rightleftharpoons H_2(g)$	0.00
$Cu^{2+}(aq) + 2e^{-} \rightleftharpoons Cu(s)$	+0.34
$\frac{1}{2}O_2(aq) + H_2O(I) + 2e^- \rightleftharpoons 2OH^-(aq)$	+0.40
$\frac{1}{2}I_2(aq) + e^- \rightleftharpoons I^-(aq)$	+0.54
$Fe^{3+}(aq) + e^{-} \rightleftharpoons Fe^{2+}(aq)$	+0.77
$Ag^+(aq) + e^- \rightleftharpoons Ag(s)$	+0.80
$Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-(aq)$	+1.36
$MnO_4^{-}(aq) + 8H^+(aq) + 5e^- \Longrightarrow Mn^{2+}(aq) + 4H_2O(I)$	+1.51

Table 3.1

(a) The reaction between zinc metal and aqueous Cu²⁺ ions can be arranged in a cell to produce electrical energy.

 $CuSO_4(aq) + Zn(s) \rightarrow Cu(s) + ZnSO_4(aq)$

A diagram of a copper/zinc cell is shown in Fig. 3.1.



Fig. 3.1

When the cell delivers a current, oxidation takes place in one half-cell and reduction in the other.

(i) Write half-equations for the reactions that take place in each half-cell.

Show state symbols.

oxidation reaction:

reduction reaction:

This question was generally well answered.

Question 3 (a) (ii)

(ii) Explain the purpose of the salt bridge.

.....[2]

The movement of ions from the salt bridge into or from the half cells was the key here, but the easier mark of 'completing the circuit' was often missed; again, worth students checking the mark allocation.

Question 3 (a) (iii)

(iii) Calculate E_{cell}° for the copper/zinc cell in **Fig. 3.1**.

E[•]_{cell} = V [1]

This question was generally well answered.

Question 3 (b) (i)

(b) Standard electrode potentials can be used to decide whether a reaction is feasible.

Predict, with reasons, if any reaction could take place in each of the following situations.

Use the data in Table 3.1.

(i) Metallic silver is added to aqueous iron(II) sulfate.

.....[1]

This question proved difficult for many candidates, with vague suggested answers. The question relied on candidates realising that for silver metal to react silver atoms must lose electrons, and this can only happen to a half reaction with a greater positive (less negative) standard electrode potential.

Question 3 (b) (ii)

(ii) Aqueous potassium chloride is added to acidified potassium manganate(VII) solution.

Better answered than Question 3 (b) (i).

Question 3 (c) (i)

(c) The relationship between the ion concentration and the electrode potential for a metal/ion electrode is given (at 298 K) by **Equation 3.1**, where *n* is the number of electrons transferred in the half reaction.

 $E = E^{\circ} + \frac{0.059 \times \log_{10}[\text{ion}]}{n}$ Equation 3.1 (i) Explain how Equation 3.1 shows that $E = E^{\circ}$ under standard conditions. (i) Explain how Equation 3.1 shows that $E = E^{\circ}$ under standard conditions.

Only the better candidates explained this logically i.e. standard concentration is 1 mol dm⁻³ and log_{10} [1] is zero.

Question 3 (c) (ii)

(ii) Calculate the electrode potential of a $Zn^{2+}(aq)/Zn(s)$ half-cell when $[Zn^{2+}(aq)] = 0.20 \text{ mol dm}^{-3}$.

Half reaction	Standard electrode potential, <i>E</i> ^e / V	
$Zn^{2+}(aq) + 2e^{-} \rightleftharpoons Zn(s)$	-0.76	
Fe ²⁺ (aq) + 2e [−] ➡ Fe(s)	-0.44	
$2H^+(aq) + 2e^- \rightleftharpoons H_2(g)$	0.00	
$Cu^{2+}(aq) + 2e^{-} \rightleftharpoons Cu(s)$	+0.34	
$\frac{1}{2}O_2(aq) + H_2O(I) + 2e^- \Longrightarrow 2OH^-(aq)$	+0.40	
$\frac{1}{2}I_2(aq) + e^- \rightleftharpoons I^-(aq)$	+0.54	
$Fe^{3+}(aq) + e^{-} \rightleftharpoons Fe^{2+}(aq)$	+0.77	
$Ag^+(aq) + e^- \rightleftharpoons Ag(s)$	+0.80	
$Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-(aq)$	+1.36	
$MnO_4^{-}(aq) + 8H^+(aq) + 5e^- \Longrightarrow Mn^{2+}(aq) + 4H_2O(I)$	+1.51	

Table 3.1

This question was well answered.

Question 3 (d)

(d) The rusting of iron is an electrochemical process.

A solution of 'ferroxyl' indicator can be used to investigate the reactions taking place in rusting. This indicator turns blue in the presence of Fe^{2+} ions and pink in alkaline solution.

In an experiment to investigate rusting, two iron nails are placed in a solution of 'ferroxyl' indicator (containing sodium chloride), in two separate petri dishes. One of the nails has copper wire wrapped around it, the other zinc wire.

The results are shown in Fig. 3.2 below.





A student suggests that the results in **Fig. 3.2** show that the attached zinc wire prevents the iron nail from rusting, but the attached copper wire does not.

Comment on the student's suggestion and use appropriate half-equations from **Table 3.1** to explain the colours in **Fig. 3.2**.

This question required candidates to link the results/observations from the investigation (Fig. 3.2) to the underlying chemistry. Many candidates gave other correct, but superfluous, chemistry involved in rusting which the practical was not designed to illustrate.

Exemplar 1 below shows a succinct, accurate answer gaining all 4 available marks.

Exemplar 1

The student is correct as the Zinc wire he a greate electrode potential than the Iron nail. Heldy reaction at is Zn (s) -7 Zn + 2e, 202 (ap) +H2 O(c) +2e -> 204 (ap) Hence no blie per for Fet and is pink for OH with is allalm. The copper wire does not prentert rusting as it has a lower electrode polerhial. The half equations are ! Fest-> Fertal) + 2e", 202 (aq) + H20 (1) +2e -> 20 H aq). Hence blue for Fett and pint again for OH- Wich is alkaline. [4]

This exemplar links the observations to the underlying chemistry.

Question 4 (a)

- 4 This question refers to the Practical Insert that is provided as an insert to this paper.
 - (a) Suggest how the students accurately made up a solution of 0.300 mol dm⁻³ copper(II) nitrate from their standard 0.400 mol dm⁻³ solution.

Name the apparatus involved.

A significant minority of students ignored the instruction to name the apparatus or used apparatus that would not meet the required precision.

Question 4 (b) (i)

(b) (i) Plot a graph of absorbance against concentration on the graph paper below and draw an appropriate line of best fit. Label the axes. [3]



A common error was for the x axis (concentration) scale not to be linear, and not using 0.0 as a perfectly valid point (helping out with the construction of a sensible line of best fit).

Question 4 (b) (ii)

(ii) Use your graph on page 12 and data from the insert to calculate the percentage by mass of copper in the brass sample.

percentage by mass of copper in the brass sample =	%	[3	31
percentage by made of copper in the brade dample	/0	ີ້	' 1

More successful responses had little trouble with this calculation.

Question 4 (b) (iii)

(iii) Explain why an orange filter is placed into the colorimeter before taking the readings.

Most candidates recognised it was the complimentary colour of the copper sulfate solution but missed the second mark. (Gives maximum absorbance.)

Question 4 (c) (i)

- (c) A copper(II) ion is said to form an octahedral complex with water ligands.
- (i) Explain the term ligand.[1]

A significant minority of candidates did not score this mark, giving rather vague answers.

Question 4 (c) (ii)

(ii) Complete the diagram below to show the **shape** of the octahedral copper(II) complex ion with water ligands.



[3]

Examiners were looking for use of 'wedges and dashes' to indicate the 3D nature of complex in order to gain the third mark.

Exemplar 2



The exemplar above shows best use of wedges and dashes to illustrate the 3D nature of the complex ion.

Question 4 (d)

(d) d-block elements form a variety of differently coloured complex ions.

Explain, in terms of electronic structure, why these complex ions are coloured and why different complexes of the same cation have different colours.

[4]

Many candidates' answers became quite jumbled in this question. The question did however prove very discriminating.

Misconception

There were at least two common misconceptions that occurred in some candidate answers. The most common being the idea that the colour of a transition complex is due to electrons dropping down electronic energy levels and releasing light energy (i.e. emission), rather than the colour of the complex being the reflected or transmitted frequencies of light not absorbed. Another misconception also seen, is that the d orbitals form a delocalised upper pi orbital and a lower ground state delocalised pi orbital.

Exemplar 3

uhen a ligand binds, the a black elements of substral is split sothat an energy gap conseponding to visible light is present, exectants are excited by attenting energy (E=hf) and male to the higher energy level, the complementary colour to that observed is seen of the rest of visible light is transmitted. The some attion can have different colours because different ugands kind which cause the d subshew to split by different amounts so energy gap is conseponding to onether wavelonghy of visible light so transmitted complementary colour is different

This candidate response offers a relatively succinct response gaining all 4 marks.

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