



MEGA LECTURE · A LEVEL CHEMISTRY

Chapter 4: Acids and Redox

OCR A Level Chemistry A (H432) — Structured Practice Worksheet

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iGCSE, O & A Level



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SPECIFICATION

2.1.4 Acids · 2.1.5 Redox · PAG
2 Titration

TOTAL MARKS

100

SUGGESTED TIME

2 hours

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Instructions to candidates

- Answer **all** questions in the spaces provided. Show all working in calculations.
- Include state symbols where equations are requested with them.
- Marks are shown in brackets [] at the end of each part.
- You may use a scientific calculator and the periodic table on the OCR Data Sheet.

Question 1 — Acids, bases and alkalis

2.1.4

- a. Give the formulae of sulfuric acid and nitric acid. [2]
-
- b. State what is released when an acid is dissolved in water. [1]
-
- c. Explain the difference between a *strong* acid and a *weak* acid, giving one example of each. [3]
-
-
-
- d. Define the terms *base* and *alkali*, making the difference between them clear. [2]
-
-
-

Question 1 total: 8 marks**Question 2 — Neutralisation**

2.1.4

- a. Write the full equation for the reaction of sulfuric acid with sodium hydroxide. [2]
-
-
- b. Write the full equation, with state symbols, for the reaction of calcium carbonate with hydrochloric acid. [2]
-
-
- c.

Write the full equation for the reaction of magnesium oxide with nitric acid.

[2]

d. Write the equation for the reaction of ammonia with hydrochloric acid, and name the product.

[2]

e. Write the ionic equation, with state symbols, for the reaction of any carbonate ion with any acid.

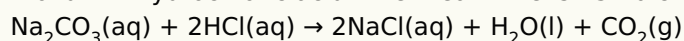
[2]

Question 2 total: 10 marks

Question 3 — Acid-base titration

PAG 2

A student titrates 25.0 cm³ portions of aqueous sodium carbonate of unknown concentration with 0.150 mol dm⁻³ hydrochloric acid. The mean titre is 23.40 cm³.



a. Describe the procedure for carrying out one accurate titration, from filling the burette to reading the endpoint.

[4]

b. Name a suitable indicator for this titration and state its colour change at the endpoint.

[2]

c. Explain why a conical flask, rather than a beaker, is used for the alkali.

[1]

d. Calculate the concentration of the sodium carbonate solution in mol dm⁻³.

[3]

e. Convert your answer to a concentration in g dm⁻³.

[2]

Question 4 — Oxidation numbers

2.1.5

a. Deduce the oxidation number of the stated element in each species.

[6]

Species	Element	Oxidation number
SO_4^{2-}	S	
NO_3^-	N	
$\text{Cr}_2\text{O}_7^{2-}$	Cr	
NaClO	Cl	
MnO_4^-	Mn	
Fe_2O_3	Fe	

b. Oxygen usually has oxidation number -2 and hydrogen $+1$. Give one example of a compound in which each of these rules is *not* followed, stating the oxidation number in your example.**[2]**

c. Use oxidation numbers to explain the name *potassium chlorate(V)* and write its formula.**[2]**

Question 4 total: 10 marks**Question 5 — Identifying redox reactions**

2.1.5

Zinc metal is added to aqueous copper(II) sulfate: $\text{Zn(s)} + \text{CuSO}_4\text{(aq)} \rightarrow \text{ZnSO}_4\text{(aq)} + \text{Cu(s)}$ a. Define *oxidation* and *reduction* in terms of electron transfer.**[2]**

b. Use oxidation numbers to identify which species is oxidised and which is reduced in this reaction. Justify your answer.

[3]

c. Write the two half-equations for this reaction.

[2]

d.

Magnesium reacts with dilute hydrochloric acid. Write the full equation and use oxidation numbers to show that this is a redox reaction.

[3]

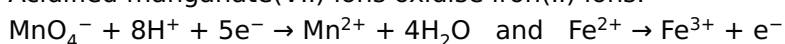
Question 5 total: 10 marks

Question 6 — Redox equations

2.1.5

a. Acidified manganate(VII) ions oxidise iron(II) ions:

[4]



Combine the two half-equations to give the overall ionic equation.

b. Chlorine reacts with water: $\text{Cl}_2 + \text{H}_2\text{O} \rightarrow \text{HCl} + \text{HClO}$. Assign oxidation numbers to chlorine in each species and explain why this reaction is described as *disproportionation*.

[4]

c. Write the equation for the reaction of chlorine with cold dilute aqueous sodium hydroxide.

[2]

Question 6 total: 10 marks

Question 7 — Metals and acids

2.1.4 / 2.1.5

a. State two observations when calcium is added to dilute sulfuric acid.

[2]

b. Write the full equation, with state symbols, for this reaction.

[2]

c. Identify, with oxidation numbers, the element oxidised and the element reduced.

[3]

d.

Write the ionic equation for the reaction of calcium with dilute hydrochloric acid, with state symbols.

[3]

Question 7 total: 10 marks

Question 8 — Gas tests

2.1.4

a. Describe the test for carbon dioxide gas and its positive result.

[2]

b. Explain, with an equation, why limewater turns cloudy in a positive test.

[1]

c. Describe the test for hydrogen gas and its positive result.

[2]

d. Describe how to show that a solid contains ammonium ions, NH_4^+ , including the observation for a positive result.

[3]

Question 8 total: 8 marks

Question 9 — Percentage purity

2.1.3 / 2.1.4

A 1.00 g sample of impure calcium carbonate (the impurities do not react) is added to an excess of dilute hydrochloric acid. 220 cm³ of carbon dioxide is collected at RTP.

a. Write the full equation for the reaction of calcium carbonate with hydrochloric acid.

[2]

b. Calculate the percentage by mass of calcium carbonate in the sample.

[5]

c.

State one assumption made in this calculation.

[1]

Question 9 total: 8 marks

Question 10 — Planning a titration

PAG 2

Vinegar is a dilute solution of ethanoic acid, CH_3COOH . Plan an experiment to determine the concentration of ethanoic acid in a sample of vinegar by titration with a standard solution of sodium hydroxide. Include the apparatus, procedure, choice of indicator, and an outline of the calculation.

[6]

This question is marked using a level of response.

a. Define the term *standard solution*.

[2]

Question 10 total: 8 marks

Question 11 — Oxidation number drill

2.1.5

a. Deduce the oxidation number of the stated element in each species.

[6]

Species	Element	Oxidation number
H_2O_2	O	
NaH	H	
NH_4^+	N	
PO_4^{3-}	P	
SO_3^{2-}	S	
CO_3^{2-}	C	

Question 11 total: 6 marks

PAPER TOTAL: 100 MARKS

MARK SCHEME — Chapter 4: Acids and Redox

Award marks independently unless stated. ECF applies to calculations.

Q1 (a) [2]

- H_2SO_4 (1); HNO_3 (1)

Q1 (b) [1]

- H^+ ions (protons) are released. (1)

Q1 (c) [3]

- Strong acid: fully/completely dissociates in aqueous solution, e.g. HCl / H_2SO_4 / HNO_3 . (1 + 1 for example)
- Weak acid: only partially dissociates, e.g. CH_3COOH . (1 including example)

Q1 (d) [2]

- Base: a substance that neutralises an acid / accepts H^+ (e.g. metal oxide, hydroxide, carbonate). (1)
- Alkali: a base that dissolves in water, releasing OH^- ions. (1)

Q2 (a) [2]

- $\text{H}_2\text{SO}_4 + 2\text{NaOH} \rightarrow \text{Na}_2\text{SO}_4 + 2\text{H}_2\text{O}$ (1 species, 1 balancing)

Q2 (b) [2]

- $\text{CaCO}_3(\text{s}) + 2\text{HCl}(\text{aq}) \rightarrow \text{CaCl}_2(\text{aq}) + \text{H}_2\text{O}(\text{l}) + \text{CO}_2(\text{g})$ (1 balanced, 1 state symbols)

Q2 (c) [2]

- $\text{MgO} + 2\text{HNO}_3 \rightarrow \text{Mg}(\text{NO}_3)_2 + \text{H}_2\text{O}$ (1 species, 1 balancing)

Q2 (d) [2]

- $\text{NH}_3 + \text{HCl} \rightarrow \text{NH}_4\text{Cl}$ (1)
- Ammonium chloride. (1)

Q2 (e) [2]

- $\text{CO}_3^{2-}(\text{aq}) + 2\text{H}^+(\text{aq}) \rightarrow \text{H}_2\text{O}(\text{l}) + \text{CO}_2(\text{g})$ (1 species, 1 state symbols)

Q3 (a) [4]

- Rinse and fill the burette with acid, removing the air bubble from the jet; record the initial reading at eye level. (1)
- Pipette 25.0 cm^3 of the alkali into a conical flask, add a few drops of indicator. (1)
- Add acid with swirling; near the endpoint add dropwise. (1)
- Stop at the first permanent colour change; record the final reading; repeat until concordant. (1)

Q3 (b) [2]

- Methyl orange. (1)
- Yellow $\rightarrow$ red (accept orange at endpoint). (1)
- *Phenolphthalein is unsuitable: it changes before the second equivalence point of carbonate.*

Q3 (c) [1]

- A conical flask can be swirled without splashing/loss of solution. (1)

Q3 (d) [3]

- $n(\text{HCl}) = 0.02340 \times 0.150 = 3.51 \times 10^{-3} \text{ mol}$ (1)
- $n(\text{Na}_2\text{CO}_3) = 3.51 \times 10^{-3} \div 2 = 1.755 \times 10^{-3} \text{ mol}$ (1)
- $c = 1.755 \times 10^{-3} \div 0.0250 = 0.0702 \text{ mol dm}^{-3}$. (1)

Q3 (e) [2]

- $M(\text{Na}_2\text{CO}_3) = 106 \text{ g mol}^{-1}$; 0.0702×106 (1) = 7.44 g dm^{-3} . (1)

Q4 (a) [6] — 1 mark each

- S in $\text{SO}_4^{2-} = +6$ · N in $\text{NO}_3^- = +5$ · Cr in $\text{Cr}_2\text{O}_7^{2-} = +6$
- Cl in $\text{NaClO} = +1$ · Mn in $\text{MnO}_4^- = +7$ · Fe in $\text{Fe}_2\text{O}_3 = +3$

Q4 (b) [2]

- O = -1 in peroxides, e.g. H_2O_2 . (1)
- H = -1 in metal hydrides, e.g. NaH. (1)

Q4 (c) [2]

- The (V) shows chlorine has oxidation number +5 in the chlorate ion. (1)
- KClO_3 . (1)

Q5 (a) [2]

- Oxidation: loss of electrons. (1)
- Reduction: gain of electrons. (1)

Q5 (b) [3]

- Zn: $0 \rightarrow +2$, so zinc is oxidised. (1)
- Cu: $+2 \rightarrow 0$, so copper(II) is reduced. (1)
- Correct justification linking increase/decrease of oxidation number to oxidation/reduction. (1)

Q5 (c) [2]

- $\text{Zn} \rightarrow \text{Zn}^{2+} + 2\text{e}^-$ (1)
- $\text{Cu}^{2+} + 2\text{e}^- \rightarrow \text{Cu}$ (1)

Q5 (d) [3]

- $\text{Mg} + 2\text{HCl} \rightarrow \text{MgCl}_2 + \text{H}_2$ (1)
- Mg: $0 \rightarrow +2$ (oxidised); H: $+1 \rightarrow 0$ (reduced). (1)
- Both oxidation and reduction occur together, so the reaction is redox. (Cl stays -1, a spectator.) (1)

Q6 (a) [4]

- Multiply the iron half-equation by 5 to balance the electrons. (1)
- $\text{MnO}_4^- + 8\text{H}^+ + 5\text{Fe}^{2+} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O} + 5\text{Fe}^{3+}$ (2 for fully correct; 1 if one error)
- No electrons remaining in the overall equation; charges balance (+17 each side). (1)

Q6 (b) [4]

- Cl in $\text{Cl}_2 = 0$. (1)
- Cl in HCl = -1; Cl in HClO = +1. (1)
- The same element (chlorine) is simultaneously oxidised ($0 \rightarrow +1$)... (1)
- ...and reduced ($0 \rightarrow -1$), which is disproportionation. (1)

Q6 (c) [2]

- $\text{Cl}_2 + 2\text{NaOH} \rightarrow \text{NaCl} + \text{NaClO} + \text{H}_2\text{O}$ (1 species, 1 balancing)

Q7 (a) [2]

- Effervescence / bubbles of gas. (1)
- The metal dissolves / solution warms up. (Accept: white solid CaSO_4 may form.) (1)

Q7 (b) [2]

- $\text{Ca(s)} + \text{H}_2\text{SO}_4\text{(aq)} \rightarrow \text{CaSO}_4\text{(aq)} + \text{H}_2\text{(g)}$ (1 balanced, 1 state symbols)

Q7 (c) [3]

- Ca: $0 \rightarrow +2$, oxidised. (1)
- H: $+1 \rightarrow 0$, reduced. (1)
- S and O unchanged (+6, -2) — spectators. (1)

Q7 (d) [3]

- $\text{Ca(s)} + 2\text{H}^+\text{(aq)} \rightarrow \text{Ca}^{2+}\text{(aq)} + \text{H}_2\text{(g)}$ (1 species, 1 balancing/charges, 1 state symbols)

Q8 (a) [2]

- Bubble the gas through limewater. (1)
- Positive: limewater turns cloudy/milky. (1)

Q8 (b) [1]

- $\text{Ca(OH)}_2\text{(aq)} + \text{CO}_2\text{(g)} \rightarrow \text{CaCO}_3\text{(s)} + \text{H}_2\text{O(l)}$ — insoluble calcium carbonate forms. (1)

Q8 (c) [2]

- Hold a lighted splint at the mouth of the tube. (1)
- Positive: the gas ignites with a squeaky pop. (1)

Q8 (d) [3]

- Warm the solid with aqueous sodium hydroxide. (1)
- Test the gas released with damp red litmus paper. (1)
- Positive: litmus turns blue (ammonia gas is released). (1)

Q9 (a) [2]

- $\text{CaCO}_3 + 2\text{HCl} \rightarrow \text{CaCl}_2 + \text{H}_2\text{O} + \text{CO}_2$ (1 species, 1 balancing)

Q9 (b) [5]

- $n(\text{CO}_2) = 220 \div 24\,000 = 9.17 \times 10^{-3} \text{ mol}$ (1)
- 1 : 1 ratio $\rightarrow n(\text{CaCO}_3) = 9.17 \times 10^{-3} \text{ mol}$ (1)
- $m(\text{CaCO}_3) = 9.17 \times 10^{-3} \times 100.1 \text{ (1)} = 0.918 \text{ g}$ (1)
- % purity = 91.8% (accept 91.7–91.8). (1)

Q9 (c) [1]

- The impurities do not react with the acid / do not release gas (or: all CO_2 is collected). (1)

Q10 (a) [6] — Level of response

- **Level 3 (5–6):** Complete, workable plan with correct apparatus, procedure, indicator and calculation outline.
- **Level 2 (3–4):** Workable plan with minor omissions (e.g. no dilution consideration or vague endpoint description).
- **Level 1 (1–2):** Fragmentary plan; major steps missing.

Indicative content: pipette 25.0 cm³ of vinegar (diluted accurately if concentrated) into a conical flask; add phenolphthalein — suitable for a weak acid–strong base titration (colourless $\rightarrow$ pink); titrate with standard NaOH from a burette to the first permanent pale pink; repeat to concordance. Calculation: $n(\text{NaOH}) = c \times V$; 1:1 ratio $\rightarrow n(\text{CH}_3\text{COOH})$; $c = n \div 0.0250$ ($\times$ dilution factor if diluted).

Q10 (b) [2]

- A solution of accurately known concentration... (1)
- ...prepared by dissolving a known mass in a known volume (volumetric flask). (1)

Q11 (a) [6] — 1 mark each

- O in $\text{H}_2\text{O}_2 = -1$ · H in NaH = -1 · N in $\text{NH}_4^+ = -3$
- P in $\text{PO}_4^{3-} = +5$ · S in $\text{SO}_3^{2-} = +4$ · C in $\text{CO}_3^{2-} = +4$

END OF MARK SCHEME · 100 marks