

JANUARY 1997

PROVINCIAL EXAMINATION

MINISTRY OF EDUCATION, SKILLS AND TRAINING

CHEMISTRY 12

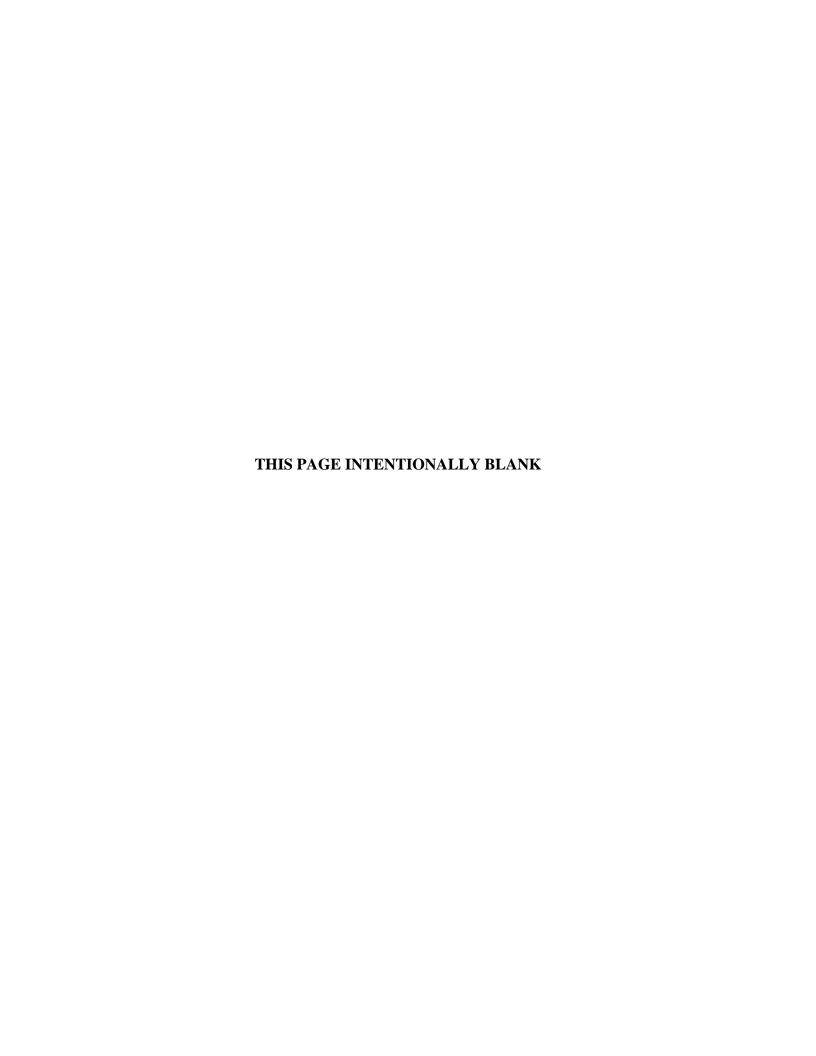
GENERAL INSTRUCTIONS

- 1. Insert the stickers with your Student I.D. Number (PEN) in the allotted spaces above. Under no circumstance is your name or identification, other than your Student I.D. Number, to appear on this paper.
- 2. Take the separate Answer Sheet and follow the directions on its front page.
- 3. Be sure you have an **HB pencil** and an eraser for completing your Answer Sheet. Follow the directions on the Answer Sheet when answering multiple-choice questions.
- 4. For each of the written-response questions, write your answer in the space provided.
- 5. When instructed to open this booklet, **check the numbering of the pages** to ensure that they are numbered in sequence from page one to the last page, which is identified by

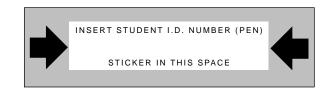
END OF EXAMINATION.

6. At the end of the examination, place your Answer Sheet inside the front cover of this booklet and return the booklet and your Answer Sheet to the supervisor.

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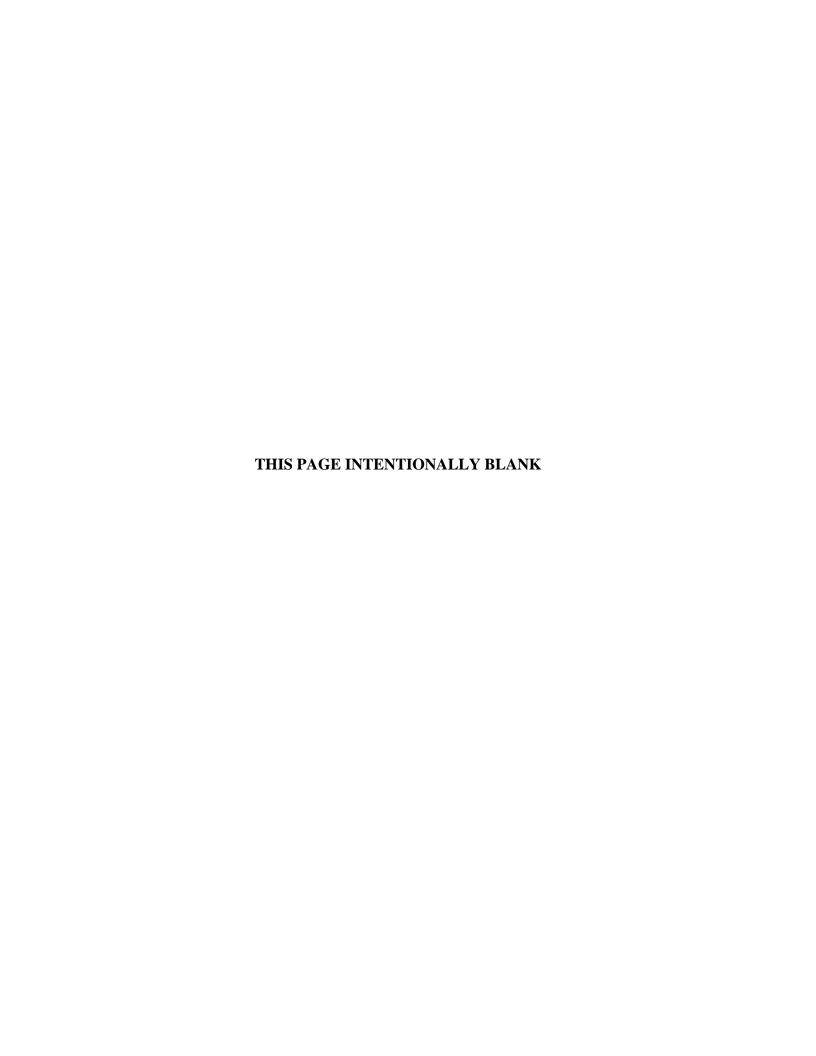
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CHEMISTRY 12 JANUARY 1997 PROVINCIAL

Course Code = CH Examination Type = P

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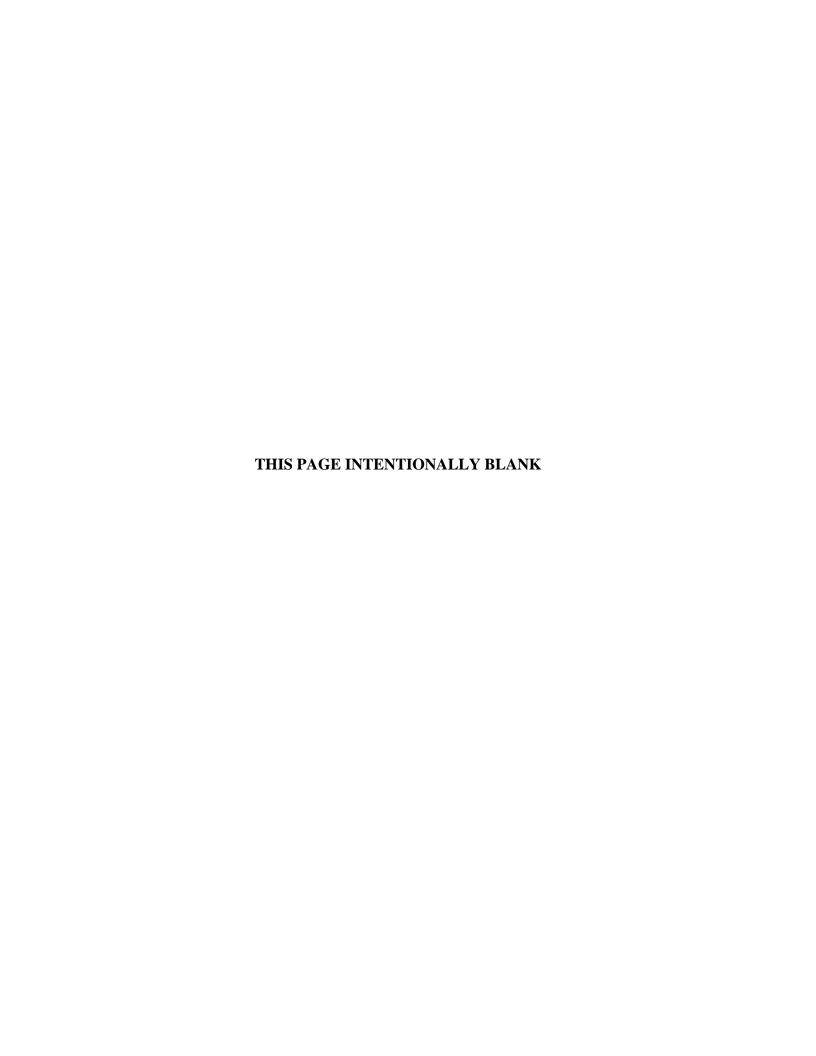
CHEMISTRY 12 PROVINCIAL EXAMINATION

			V	⁷ alue	Suggested Time
1.	This exam	ination consists of two parts:			
	PART A:	48 multiple-choice questions		48	70
	PART B:	11 written-response questions		32	50
			Total:	80 marks	120 minutes

- 2. The following tables can be found in the separate **Data Booklet:**
 - Periodic Table of the Elements
 - Atomic Masses of the Elements
 - Names, Formulae, and Charges of Some Common Ions
 - Solubility of Common Compounds in Water
 - Solubility Product Constants at 25°C
 - Relative Strengths of Brönsted-Lowry Acids and Bases
 - Acid-Base Indicators
 - Standard Reduction Potentials of Half-cells

No other reference materials or tables are allowed.

- 3. An approved scientific calculator is essential for the examination. The calculator must be a hand-held device designed **only** for mathematical computations such as logarithmic and trigonometric functions. It **can be** programmable, but **must not** contain any graphing capabilities. You **must not** bring into the examination room any devices to support calculators such as manuals, printed or electronic cards, printers, memory expansion chips or cards, or keyboards.
- 4. You have **two hours** to complete this examination.



PART A: MULTIPLE CHOICE

Value: 48 marks Suggested Time: 70 minutes

INSTRUCTIONS: For each question, select the **best** answer and record your choice on the Answer

Sheet provided. Using an HB pencil, completely fill in the circle that has the letter

corresponding to your answer.

- 1. Which of the following factors affects the rate of heterogeneous reactions only?
 - A. nature of reactants
 - B. temperature of system
 - C. surface area of reactants
 - D. concentration of reactants
- 2. A 25.0 mL sample of hydrogen peroxide decomposes producing 50.0 mL of oxygen gas in 137 s. The rate of formation of $\,{\rm O}_2$ in mL/min is
 - A. 0.182 mL/min
 - B. 0.365 mL/min
 - C. 10.9 mL/min
 - D. 21.9 mL/min
- 3. For collisions to be successful, reactants must have
 - A. favourable geometry only.
 - B. sufficient heat of reaction only.
 - C. sufficient potential energy only.
 - D. sufficient kinetic energy and favourable geometry.
- 4. Consider the following reaction:

$$\frac{1}{2}H_{2(g)} + \frac{1}{2}I_{2(g)} \to HI_{(g)}$$
 $\Delta H = +28 \text{ kJ}$

The activation energy for the formation of HI is 167 kJ. The activation energy for the decomposition of HI is

- A. 28 kJ
- B. 139 kJ
- C. 167 kJ
- D. 195 kJ

5. Consider the following reaction mechanism:

Step 1:
$$NO_{(g)} + O_{3(g)} \rightarrow NO_{2(g)} + O_{2(g)}$$

Step 2:
$$O_{(g)} + NO_{2(g)} \rightarrow NO_{(g)} + O_{2(g)}$$

The catalyst is

- A. O_2
- B. O_3
- C. NO
- D. NO₂

6. Consider the following:

$$2NH_{3(g)} \rightleftarrows N_{2(g)} + 3H_{2(g)}$$

A flask is initially filled with $\,\mathrm{NH}_3$. As the system approaches equilibrium, the rate of the forward reaction

- A. increases as the rate of the reverse reaction decreases.
- B. decreases as the rate of the reverse reaction increases.
- C. increases as the rate of the reverse reaction increases.
- D. decreases as the rate of the reverse reaction decreases.

7. Consider the following reaction:

$$Na_2CO_{3(s)} + 2HCl_{(aq)} \rightarrow 2NaCl_{(aq)} + CO_{2(g)} + H_2O_{(\ell)}$$
 $\Delta H = -27.7 \text{ kJ}$

In this reaction,

- A. minimum enthalpy and maximum entropy both favour products.
- B. minimum enthalpy and maximum entropy both favour reactants.
- C. minimum enthalpy favours products and maximum entropy favours reactants.
- D. minimum enthalpy favours reactants and maximum entropy favours products.

8. Consider the following equilibrium:

$$CH_{4(g)} + H_2O_{(g)} + heat \rightleftharpoons CO_{(g)} + 3H_{2(g)}$$

In which of the following will both stresses shift the equilibrium right?

- A. a decrease in temperature and a decrease in volume
- B. an increase in temperature and a decrease in volume
- C. a decrease in temperature and an increase in volume
- D. an increase in temperature and an increase in volume
- 9. Consider the following equilibrium:

$$2SO_{2(g)} + O_{2(g)} \rightleftharpoons 2SO_{3(g)} \qquad \Delta H = -198 \text{ kJ}$$

There will be no shift in this equilibrium when

- A. more O_2 is added.
- B. a catalyst is added.
- C. the volume is increased.
- D. the temperature is increased.
- 10. Consider the following equilibrium:

$$2Fe_{(s)} + 3H_2O_{(g)} \rightleftharpoons Fe_2O_{3(s)} + 3H_{2(g)}$$

The equilibrium constant expression is

A.
$$K_{eq} = \frac{[Fe_2O_3][H_2]^3}{[Fe]^2[H_2O]^3}$$

B.
$$K_{eq} = \frac{[Fe_2O_3][3H_2]}{[2Fe][3H_2O]}$$

C.
$$K_{eq} = \frac{\left[H_2\right]^3}{\left[H_2O\right]^3}$$

D.
$$K_{eq} = [H_2]^3$$

11. Consider the following equilibrium:

$$N_2O_{4(g)} \rightleftharpoons 2NO_{2(g)}$$
 $K_{eq} = 0.133$

At equilibrium, the $\left[N_2O_4\right]$ is equal to

A.
$$\frac{0.133}{[NO_2]}$$

B.
$$\frac{[NO_2]}{0.133}$$

$$C. \quad \frac{0.133}{\left[NO_2\right]^2}$$

D.
$$\frac{[NO_2]^2}{0.133}$$

12. Consider the following equilibrium:

$$CaCO_{3(s)} + 556 \text{ kJ} \rightleftharpoons CaO_{(s)} + CO_{2(g)}$$

The value of the equilibrium constant will increase when

- A. CO_2 is added.
- B. CO₂ is removed.
- C. the temperature is increased.
- D. the temperature is decreased.
- 13. Consider the following equilibrium:

$$C_{(s)} + H_2O_{(g)} \rightleftharpoons CO_{(g)} + H_{2(g)}$$

The contents of a 1.00 L container at equilibrium were analyzed and found to contain 0.20 mol C, 0.20 mol H_2O , 0.60 mol CO and 0.60 mol H_2 . The equilibrium constant is

- A. 0.11
- B. 0.56
- C. 1.8
- D. 9.0

- 14. When a student mixes equal volumes of $0.20 \text{ M} \text{ Na}_2\text{S}$ and $0.20 \text{ M} \text{ Sr}(\text{OH})_2$,
 - A. no precipitate forms.
 - B. a precipitate of only SrS forms.
 - C. a precipitate of only NaOH forms.
 - D. precipitates of both NaOH and SrS form.
- 15. A student wishes to identify an unknown cation in a solution. A precipitate does not form with the addition of SO_4^{2-} , but does form with the addition of S^{2-} . Which of the following is the unknown cation?
 - A. Ag⁺
 - B. Mg^{2+}
 - C. Ca²⁺
 - D. Cu²⁺
- 16. When dissolved in water, which of the following produces an ionic solution?
 - A. O_2
 - B. CH₄
 - C. CaCl₂
 - D. $C_{12}H_{22}O_{11}$
- 17. The solubility of MnS is $4.8 \times 10^{-7} \, \mathrm{M}$, at $25^{\circ} \, \mathrm{C}$. The $\, \mathrm{K}_{sp} \,$ value is
 - A. 2.3×10^{-13}
 - B. 4.8×10^{-7}
 - C. 9.6×10^{-7}
 - D. 6.9×10^{-4}
- 18. Which of the following units is commonly used to describe solubility?
 - A. mL/s
 - B. g/°C
 - $C. \quad mol/L \\$
 - D. °C/mol

- 19. A 200.0 mL solution contains 0.050 mol of $Ba(NO_3)_2$. The $[NO_3^-]$ is
 - A. 0.050 M
 - B. 0.10 M
 - C. 0.25 M
 - D. 0.50 M
- 20. Consider the following solubility equilibrium:

$$MgCO_{3(s)} \rightleftharpoons Mg^{2+}_{(aq)} + CO_3^{2-}_{(aq)}$$

The addition of which of the following substances would decrease the solubility of MgCO₃?

- A. H₂O
- B. NaCl
- C. NaOH
- D. Na₂CO₃
- 21. A basic solution
 - A. tastes sour.
 - B. feels slippery.
 - C. does not conduct electricity.
 - D. reacts with metals to release oxygen gas.
- 22. The balanced formula equation for the neutralization of H_2SO_4 by KOH is
 - A. $H_2SO_4 + KOH \rightarrow KSO_4 + H_2O$
 - $\text{B.} \quad \text{H}_2\text{SO}_4 + \text{KOH} \quad \rightarrow \ \text{K}_2\text{SO}_4 + \text{H}_2\text{O}$
 - C. $H_2SO_4 + 2KOH \rightarrow K_2SO_4 + H_2O$
 - D. $H_2SO_4 + 2KOH \rightarrow K_2SO_4 + 2H_2O$

- 23. An Arrhenius base is defined as a substance which
 - A. donates protons.
 - B. donates electrons.
 - C. produces H⁺ in solution.
 - D. produces OH⁻ in solution.
- 24. Consider the following equilibrium:

$$HS^- + H_3PO_4 \rightleftharpoons H_2S + H_2PO_4^-$$

The order of Brönsted-Lowry acids and bases is

- A. acid, base, acid, base.
- B. acid, base, base, acid.
- C. base, acid, acid, base.
- D. base, acid, base, acid.
- 25. The equation representing the reaction of ethanoic acid with water is
 - A. $CH_3COO^- + H_2O \rightleftharpoons CH_3COOH + OH^-$
 - B. $CH_3COO^- + H_2O \rightleftharpoons CH_2COO^{2-} + H_3O^+$
 - C. $CH_3COOH + H_2O \rightleftharpoons CH_3COO^- + H_3O^+$
 - D. $CH_3COOH + H_2O \rightleftharpoons CH_3COOH_2^+ + OH^-$
- 26. Consider the following equilibrium:

$$2H_2O_{(\ell)} + 57 \text{ kJ} \ \rightleftharpoons \ H_3O^+_{(aq)} + OH^-_{(aq)}$$

When the temperature is decreased, the water

- A. stays neutral and $[H_3O^+]$ increases.
- B. stays neutral and $\left[H_3O^+\right]$ decreases.
- C. becomes basic and $[H_3O^+]$ decreases.
- D. becomes acidic and $\left[H_3O^+\right]$ increases.

- 27. In a solution at 25°C, the $\left[H_3O^+\right]$ is $3.5 \times 10^{-6} M$. The $\left[OH^-\right]$ is
 - A. $3.5 \times 10^{-20} \,\mathrm{M}$
 - B. $2.9 \times 10^{-9} \text{ M}$
 - C. $1.0 \times 10^{-7} \text{ M}$
 - D. $3.5 \times 10^{-6} \,\mathrm{M}$
- 28. In a solution with a pOH of 4.22, the $\left[\text{OH}^{-} \right]$ is
 - A. $1.7 \times 10^{-10} \,\mathrm{M}$
 - B. $6.0 \times 10^{-5} \text{ M}$
 - C. 6.3×10^{-1} M
 - D. $1.7 \times 10^4 \,\mathrm{M}$
- 29. An aqueous solution of NH₄CN is
 - A. basic because $K_a < K_b$
 - B. basic because $K_a > K_b$
 - C. acidic because $K_a < K_b$
 - D. acidic because $K_a > K_b$
- 30. The net ionic equation for the predominant hydrolysis reaction of $KHSO_4$ is
 - A. $HSO_4^- + H_2O \rightleftharpoons SO_4^{2-} + H_3O^+$
 - B. $HSO_4^- + H_2O \rightleftharpoons H_2SO_4 + OH^-$
 - C. $KHSO_4 + H_2O \rightleftharpoons K^+ + SO_4^{2-} + H_3O^+$
 - D. $KHSO_4 + H_2O \rightleftharpoons K^+ + H_2SO_4 + OH^-$

31. Consider the following equilibrium for an indicator:

$$HInd + H_2O \rightleftharpoons Ind^- + H_3O^+$$

- At the transition point,
- A. $[HInd] > [Ind^-]$
- B. $[HInd] = [Ind^-]$
- C. $[HInd] < [Ind^-]$
- D. $[HInd] = [H_3O^+]$
- 32. The equivalence point in a titration is reached when 20.0 mL of H_2SO_4 is added to 20.0 mL of 0.420 M KOH. The $\left[H_2SO_4\right]$ in the original solution is
 - A. 0.00840 M
 - B. 0.210 M
 - C. 0.420 M
 - D. 0.840 M
- 33. In a titration between a weak acid and a strong base, the pH at the equivalence point is
 - A. 3
 - B. 5
 - C. 7
 - D. 9
- 34. The pH of 100.0 mL of 0.0050 M NaOH solution is
 - A. 2.30
 - B. 3.30
 - C. 10.70
 - D. 11.70
- 35. A buffer solution is prepared by adding 1.0 mol of $NaCH_3COO$ to 1.0 L of 1.0 M CH_3COOH . The molar concentration of CH_3COO^- is approximately
 - A. 0.0
 - B. 0.5
 - C. 1.0
 - D. 2.0

- 36. The equation for the reaction of Cl₂O with water is
 - A. $Cl_2O + H_2O \rightleftharpoons 2HClO$
 - B. $Cl_2O + H_2O \rightleftharpoons 2ClO + H_2$
 - C. $Cl_2O + H_2O \rightleftharpoons Cl_2 + H_2O_2$
 - D. $Cl_2O + H_2O \rightleftharpoons Cl_2 + O_2 + H_2$
- 37. Consider the following redox reaction:

$$2 \text{MnO}_4^{\;-} + 5 \text{CH}_3 \text{CHO} + 6 \text{H}^+ \rightarrow 5 \text{CH}_3 \text{COOH} + 2 \text{Mn}^{2+} + 3 \text{H}_2 \text{O}$$

The species that loses electrons is

- A. H₂O
- B. MnO_4^-
- C. CH₃CHO
- D. CH₃COOH
- 38. A spontaneous redox reaction occurs when a piece of iron is placed in 1.0 M CuSO₄. The reducing agent is
 - A. Fe
 - B. Cu²⁺
 - C. H_2O
 - D. SO₄²⁻
- 39. Consider the following redox reaction:

$$3SO_2 + 3H_2O + ClO_3^- \rightarrow 3SO_4^{2-} + 6H^+ + Cl^-$$

The reduction half-reaction is

- A. $ClO_3^- + 6H^+ \rightarrow Cl^- + 3H_2O + 6e^-$
- B. $ClO_3^- + 6H^+ + 6e^- \rightarrow Cl^- + 3H_2O$
- C. $SO_2 + 2H_2O \rightarrow SO_4^{2-} + 4H^+ + 2e^-$
- D. $SO_2 + 2H_2O + 2e^- \rightarrow SO_4^{2-} + 4H^+$

- 40. Bromine, Br₂, will react spontaneously with
 - A. I⁻
 - B. I_2
 - C. Cl
 - D. Cl₂
- 41. The substances H₂O₂, H₃PO₄ and H₂SO₃ in order of increasing strengths as oxidizing agents are:
 - A. H_2O_2 , H_3PO_4 , H_2SO_3
 - B. H_2SO_3 , H_3PO_4 , H_2O_2
 - C. H_3PO_4 , H_2SO_3 , H_2O_2
 - D. H_2O_2 , H_2SO_3 , H_3PO_4
- 42. The oxidation number of platinum in $Pt(H_2O)_4^{2+}$ is
 - A. +2
 - B. 0
 - C. +4
 - D. $+\frac{1}{2}$
- 43. Consider the following half-reaction:

$$BrO^- \rightarrow Br^-$$
 (basic)

The balanced equation for the half-reaction is

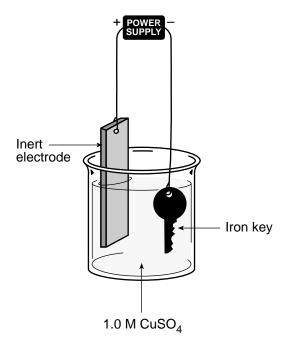
- A. $BrO^- + 2H^+ + 2e^- \rightarrow Br^- + H_2O$
- B. $BrO^- + 2H^+ \rightarrow Br^- + H_2O + 2e^-$
- C. $BrO^- + H_2O \rightarrow Br^- + 2OH^- + 2e^-$
- D. $BrO^- + H_2O + 2e^- \rightarrow Br^- + 2OH^-$

- 44. In an operating electrochemical cell, the anions migrate
 - A. towards the anode through the wire.
 - B. towards the cathode through the wire.
 - C. towards the anode through the salt bridge.
 - D. towards the cathode through the salt bridge.
- 45. A piece of iron can be prevented from corroding by
 - A. making it a cathode.
 - B. placing it in an acidic solution.
 - C. attaching a small piece of lead to it.
 - D. attaching a small piece of gold to it.
- 46. Consider the following overall equation for an electrochemical cell:

$$3Ag^+ + Cr \rightarrow Cr^{3+} + 3Ag$$

- At standard conditions, the initial cell voltage is
- A. +0.06 V
- B. +0.39 V
- C. +1.21 V
- D. +1.54 V

47. Consider the following diagram:



The half-reaction at the cathode is

A.
$$Cu^{2+} + 2e^{-} \rightarrow Cu_{(s)}$$

B.
$$2SO_4^{2-} \rightarrow S_2O_8^{2-} + 2e^-$$

C.
$$H_2O \rightarrow \frac{1}{2}O_{2(g)} + 2H^+ + 2e^-$$

D.
$$2H_2O + 2e^- \rightarrow H_{2(g)} + 2OH^-$$

48. An electrolytic process is used to purify impure lead. The electrodes are

	ANODE	CATHODE
A.	carbon	impure lead
B.	pure lead	carbon
C.	pure lead	impure lead
D.	impure lead	pure lead

This is the end of the multiple-choice section.

Answer the remaining questions directly in this examination booklet.

PART B: WRITTEN RESPONSE

Value: 32 marks	Suggested Time:	50 minutes					
INSTRUCTIONS:	You will be expected to communicate your knowledge and understanding of chemical principles in a clear and logical manner.						
	Your steps and assumptions leading to a solution must be written in the below the questions.	spaces					
	Answers must include units where appropriate and be given to the corresignificant figures.	ect number of					
	For questions involving calculation, full marks will NOT be given for only an answer.	or providing					
1. Define the term	m activated complex. (2 marks)						
		Score for Question 1:					
		1					
	n theory, explain why a mixture of natural gas and air does not react at ture but explodes when a piece of platinum is placed in the gas mixture.						
	(2 marks)						
-							
-							

Score for Question 2:

3. Identify four characteristics of a chemical equilibrium.

(2 marks)

- (2)

Score for Ouestion 3:

- 4. At high temperature, 0.500 mol HBr was placed in a 1.00 L container where it decomposed to give the equilibrium:

$$2HBr_{(g)} \rightleftharpoons H_{2(g)} + Br_{2(g)}$$

At equilibrium, the $\left[Br_{2} \right]$ is 0.0855 mol/L. What is the value of the equilibrium constant? (3 marks)

> Score for Question 4:

5. A saturated solution of BaF₂ has a $\left[Ba^{2+} \right]$ of $3.6 \times 10^{-3} M$. Calculate the K_{sp} value.

(2 marks)

Score for Question 5:

5. _____

6. Calculate the maximum mass of Na_2SO_4 which can be dissolved in 2.0 L of 1.5 M $Ca(NO_3)_2$ without a precipitate forming.

(3 marks)

Score for Question 6:

5. <u>(3)</u>

7. a) Write two equations showing the amphiprotic nature of water as it reacts with HCO_3^- . (2 marks)

b) Calculate the K_b for HCO_3^- .

(1 mark)

Score for Question 7:

8. Calculate the $\left[H_3O^+ \right]$ in 0.550 M $\, C_6H_5COOH.$

(3 marks)

Score for Question 8:

9. Calculate the pH of the solution formed by mixing 20.0 mL of 0.500 M HCl with 30.0 mL 0.300 M NaOH. (4 marks)

Score for Question 9:

9. ____

10. Consider the following reaction:

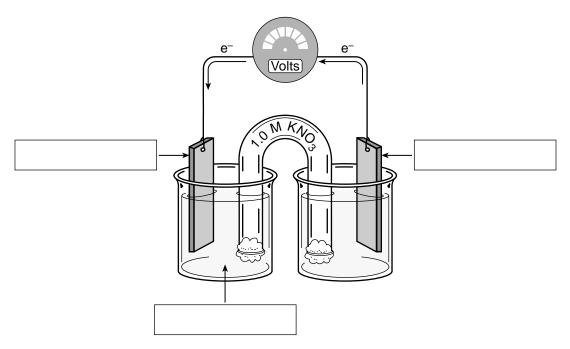
$${\rm Cr_2O_7}^{2-} + 6{\rm Br}^- + 14{\rm H}^+ \rightarrow 2{\rm Cr}^{3+} + 3{\rm Br}_2 + 7{\rm H}_2{\rm O}$$

In a redox titration, 15.58 mL of 0.125 M $\rm Cr_2O_7^{2-}$ was needed to completely oxidize the Br $^-$ in a 25.00 mL sample of NaBr. Calculate the [Br $^-$] in the original solution. (3 marks)

Score for Question 10:

0. ____

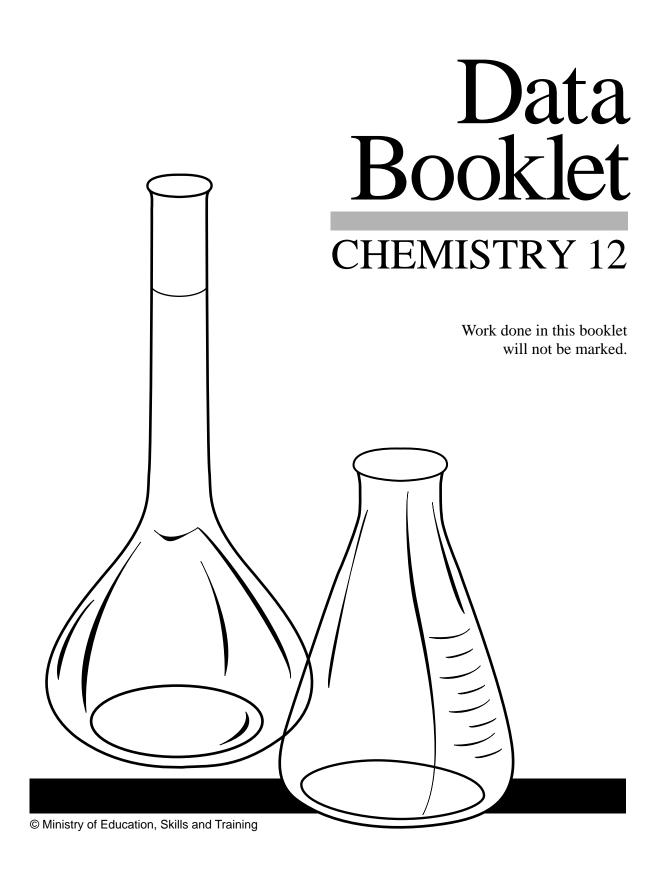
- 11. Consider the following materials and cell diagram:
 - silver, aluminum and nickel electrodes
 - 1.0 M solutions of $AgNO_3$, $Al(NO_3)_3$ and $Ni(NO_3)_2$
 - a) From the above list, select the materials that are capable of producing the greatest voltage, then label the diagram below. (3 marks)



- b) Calculate the initial voltage for the electrochemical cell in part a). (1 mark)
- c) Which two metals from the above list would produce an electrochemical cell with the smallest initial voltage? (1 mark)

Score for Question 11:

11. ____(5)



Revised November 1994

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5	Solubility Product Constants at 25°C
6	Relative Strengths of Brönsted-Lowry Acids and Bases
7	Acid-Base Indicators
8	Standard Reduction Potentials of Half-Cells

D.R. Lide, CRC Handbook of Chemistry and Physics, 74 edition, CRC Press, Boca Raton, 1993.

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		13	5	В	Boron 10.8	13	Al	Aluminum 27.0	; ;	31	Сa	Gallium 69.7	49	In	Indium 114.8	81	П	Thallium 204.4				
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								V	0 2	77	Ċ	Chromium 52.0	42	Mo	Molybdenum 95.9	74	≽	Tungsten 183.8	106	Sg	Seaborgium (1) (263)	
								u	C 8	73	>	Vanadium 50.9	41	Sp	-	73	Та	Tantalum 180.9	105	Ha	Hahnium (262)	
								-				Titanium 47.9		Zr	Zirconium 91.2		Hť	Hafnium 178.5	104	Rf	Rutherfordium (261)	
		-						r	S 2	717	Sc	Scandium 45.0	39	Y	Yttrium 88.9	57	La	Lanthanum F	68	Ac	Actinium Rui (227)	
		2	4	Be	Beryllium 9.0	12	Μα	Magnesium	24.3	70		Calcium 40.1		Sr	$\bar{\mathbf{x}}$	56			88		Radium (226)	
1 1	H Hydrogen 1.0		3	Ľ	Lithium 6.9	1		Sodium	- 1	19	K	Potassium 39.1	37	Rb	Rubidium 85.5	55	Cs	Cesium 132.9	87	Ή	Francium (223)	

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Values in parentheses are the masses of the most stable or best known isotopes for elements which do not occur naturally.

71	Lu	Lutetium 175.0	103	Lr	Lawrencium (262)
70	Yb	Ytterbium 173.0	102	$_{\rm O}^{ m N}$	Nobelium (259)
69	Tm	Thulium 168.9	101	Md	Mendelevium (258)
89	Er	Erbium 167.3	100	Fm	Fermium (257)
29	Но	Holmium 164.9	66	Es	Einsteinium (252)
99	Dy	Dysprosium 162.5	86	Cţ	Californium (251)
65	Tb	Terbium 158.9	76	Bk	Berkelium (247)
64	В	Gadolinium 157.3	96	Cm	Curium (247)
63	Eu	Europium 152.0	95	Am	Americium (243)
62	Sm	Samarium 150.4	94	Pu	Plutonium (244)
61	Pm	Promethium (145)	93	Np	Neptunium (237)
09	pN	Neodymium 144.2	92	n	Uranium 238.0
59	Pr	Praseodymium Neodymiu 140.9	91	Pa	Protactinium 231.0
28	Ce	Cerium 140.1	06	Th	Thorium 232.0

ATOMIC MASSES OF THE ELEMENTS

Based on mass of C^{12} at 12.00. Values in parentheses are the mass of the most stable or best known isotopes for elements which do not occur naturally.

Element	Symbol	Atomic Number	Atomic Mass	Element	Symbol	Atomic Number	Atomic Mass
Actinium	Ac	89	(227)	Mercury	Hg	80	200.6
Aluminum	Al	13	27.0	Molybdenum	Mo	42	95.9
Americium	Am	95	(243)	Neodymium	Nd	60	144.2
Antimony	Sb	51	121.8	Neon	Ne	10	20.2
Argon	Ar	18	39.9	Neptunium	Np	93	(237)
Arsenic	As	33	74.9	Nickel	Ni	28	58.7
Astatine	As At	85	(210)	Niobium	Nb	41	92.9
Barium	Ba	56	137.3	Nitrogen	N	7	14.0
Berkelium	Bk	97	(247)	Nobelium	No	102	(259)
Beryllium	Be	4	9.0	Osmium	Os	76	190.2
	Бе Bi	83	209.0		Os	8	190.2
Bismuth				Oxygen			
Boron	В	5	10.8	Palladium	Pd	46	106.4
Bromine	Br	35	79.9	Phosphorus	P	15	31.0
Cadmium	Cd	48	112.4	Platinum	Pt	78	195.1
Calcium	Ca	20	40.1	Plutonium	Pu	94	(244)
Californium	Cf	98	(251)	Polonium	Po	84	(209)
Carbon	C	6	12.0	Potassium	K	19	39.1
Cerium	Ce	58	140.1	Praseodymium	Pr	59	140.9
Cesium	Cs	55	132.9	Promethium	Pm	61	(145)
Chlorine	Cl	17	35.5	Protactinium	Pa	91	231.0
Chromium	Cr	24	52.0	Radium	Ra	88	(226)
Cobalt	Co	27	58.9	Radon	Rn	86	(222)
Copper	Cu	29	63.5	Rhenium	Re	75	186.2
Curium	Cm	96	(247)	Rhodium	Rh	45	102.9
Dysprosium	Dy	66	162.5	Rubidium	Rb	37	85.5
Einsteinium	Es	99	(252)	Ruthenium	Ru	44	101.1
Erbium	Er	68	167.3	Rutherfordium	Rf	104	(261)
Europium	Eu	63	152.0	Samarium	Sm	62	150.4
Fermium	Fm	100	(257)	Scandium	Sc	21	45.0
Fluorine	F	9	19.0	Selenium	Se	34	79.0
Francium	Fr	87	(223)	Silicon	Si	14	28.1
Gadolinium	Gd	64	157.3	Silver	Ag	47	107.9
Gallium	Ga	31	69.7	Sodium	Na	11	23.0
Germanium	Ge	32	72.6	Strontium	Sr	38	87.6
Gold	Au	79	197.0	Sulphur	S	16	32.1
Hafnium	Hf	72	178.5	Tantalum	Ta	73	180.9
Hahnium	Ha	105	(262)	Technetium	Tc	43	(98)
Helium	He	2	4.0	Tellurium	Te	52	127.6
Holmium	Но	67	164.9	Terbium	Tb	65	158.9
Hydrogen	Н	1	1.0	Thallium	Tl	81	204.4
Indium	In	49	114.8	Thorium	Th	90	232.0
Iodine	I	53	126.9	Thulium	Tm	69	168.9
Iridium	Îr	77	192.2	Tin	Sn	50	118.7
Iron	Fe	26	55.8	Titanium	Ti	22	47.9
Krypton	Kr	36	83.8	Tungsten	W	74	183.8
Lanthanum	La	57	138.9	Uranium	Ü	92	238.0
Lawrencium	Lr	103	(262)	Vanadium	V	23	50.9
Lead	Pb	82	207.2	Xenon	Xe	54	131.3
Lithium	Li	3	6.9	Ytterbium	Yb	70	173.0
Lutetium	Lu	71	175.0	Yttrium	Y	70 39	88.9
Magnesium	Mg	12	24.3	Zinc	Zn	30	65.4
Manganese	Mn	25	54.9	Zirconium	Zr	40	91.2
Mendelevium	Md	101	(258)				

Names, Formulae, and Charges of Some Common Ions

Positive ions (ca	tions)	Negative ions (anions)		
Aluminum	Al ³⁺	Bromide	Br ⁻	
Ammonium	$\mathrm{NH_4}^+$	Carbonate	CO_3^{2-}	
Barium	Ba^{2+}	Chlorate	ClO ₃	
Calcium	Ca^{2+}	Chloride	Cl ⁻	
Chromium(II), chromous	Cr ²⁺			
Chromium(III), chromic	Cr ³⁺	Chlorite	ClO_2^-	
Copper(I)*, cuprous	Cu^+	Chromate	CrO ₄ ²⁻	
Copper(II), cupric	Cu^{2+}	Cyanide	CN ⁻	
Hydrogen	$\mathrm{H}^{\scriptscriptstyle +}$	Dichromate	$\operatorname{Cr_2O_7}^{2-}$	
Hydronium	H_3O^+	Dihydrogen phosphate	$\mathrm{H_2PO_4}^-$	
Iron(II)*, ferrous	Fe^{2+}	Ethanoate, Acetate	CH ₃ COO ⁻	
Iron(III), ferric	Fe^{3+}	Fluoride	F^{-}	
Lead(II), plumbous	Pb^{2+}	Hydrogen carbonate, bicarbonate	HCO ₃	
Lead(IV), plumbic	Pb ⁴⁺	Hydrogen oxalate, binoxalate	$HC_2O_4^-$	
Lithium	$\mathrm{Li}^{\scriptscriptstyle +}$	Hydrogen sulphate, bisulphate	HSO_4^-	
Magnesium	$\mathrm{Mg}^{2^{+}}$		•	
Manganese(II), manganous	Mn^{2+}	Hydrogen sulphide, bisulphide	HS ⁻	
Manganese(IV)	Mn^{4+}	Hydrogen sulphite, bisulphite	HSO_3^-	
Mercury(I)*, mercurous	${\rm Hg_2}^{2+}$	Hydroxide	OH^-	
Mercury(II), mercuric	Hg^{2+}	Hypochlorite	ClO ⁻	
Potassium	K^{+}	Iodide	I ⁻	
Silver	Ag^+	Monohydrogen phosphate	$\mathrm{HPO_4}^{2-}$	
Sodium	Na^+	Nitrate	NO_3^-	
Tin(II)*, stannous	Sn^{2+}	Nitrite	NO_2^-	
Tin(IV), stannic	Sn^{4+}	Oxalate	$C_2O_4^{2-}$	
Zinc	$\mathbf{Z}\mathbf{n}^{^{2+}}$	Oxide**	0^{2-}	

* Aqueous solutions are readily oxidized by air.

** Not stable in aqueous solutions.

Perchlorate	ClO ₄
Permanganate	$\mathrm{MnO_4}^-$
Phosphate	PO ₄ ³⁻
Sulphate	SO_4^{2-}
Sulphide	S^{2-}
Sulphite	SO ₃ ²⁻
Thiocyanate	SCN ⁻

SOLUBILITY OF COMMON COMPOUNDS IN WATER

The term soluble here means $> 0.1 \ mol/L \ at \ 25^{o}C$.

NEGATIVE IONS (Anions)	POSITIVE IONS (Cations)	SOLUBILITY OF COMPOUNDS		
All	Alkali ions: Li ⁺ , Na ⁺ , K ⁺ , Rb ⁺ , Cs ⁺ , Fr ⁺	Soluble		
All	Hydrogen ion, H ⁺	Soluble		
All	Ammonium ion, NH ₄ ⁺	Soluble		
Nitrate, NO ₃ ⁻	All	Soluble		
Chloride, Cl or Bromide, Br	All others	Soluble		
or Iodide, I	Ag ⁺ , Pb ²⁺ , Cu ⁺	Low Solubility		
Sulphate, $SO_4^{\ 2-}$	All others	Soluble		
Sulphate, 50 ₄	Ag ⁺ , Ca ²⁺ , Sr ²⁺ , Ba ²⁺ , Pb ²⁺	Low Solubility		
Sulphide, S ²⁻	Alkali ions, H^+ , NH_4^+ , Be^{2+} Mg^{2+} , Ca^{2+} , Sr^{2+} , Ba^{2+}	Soluble		
	All others	Low Solubility		
Hudravida OH-	Alkali ions, H^+ , NH_4^+ , Sr^{2+}	Soluble		
Hydroxide, OH ⁻	All others	Low Solubility		
Phosphate, PO ₄ ³⁻ or Carbonate, CO ₃ ²⁻	Alkali ions, H ⁺ , NH ₄ ⁺	Soluble		
or Sulphite, SO_3^{2-}	All others	Low Solubility		

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	\mathbf{K}_{sp}
barium carbonate	BaCO ₃	2.6×10^{-9}
barium chromate	BaCrO ₄	1.2×10^{-10}
barium sulphate	${ m BaSO}_4$	1.1×10^{-10}
calcium carbonate	CaCO ₃	5.0×10^{-9}
calcium oxalate	CaC ₂ O ₄	2.3×10^{-9}
calcium sulphate	CaSO ₄	7.1×10^{-5}
copper(I) iodide	CuI	1.3×10^{-12}
copper(II) iodate	Cu(IO ₃) ₂	6.9×10^{-8}
copper(II) sulphide	CuS	6.0×10^{-37}
iron(II) hydroxide	Fe(OH) ₂	4.9×10^{-17}
iron(II) sulphide	FeS	6.0×10^{-19}
iron(III) hydroxide	Fe(OH) ₃	2.6×10^{-39}
lead(II) bromide	PbBr ₂	6.6×10^{-6}
lead(II) chloride	PbCl ₂	1.2×10^{-5}
lead(II) iodate	Pb(IO ₃) ₂	3.7×10^{-13}
lead(II) iodide	PbI ₂	8.5×10^{-9}
lead(II) sulphate	PbSO ₄	1.8×10^{-8}
magnesium carbonate	MgCO ₃	6.8×10^{-6}
magnesium hydroxide	Mg(OH) ₂	5.6×10^{-12}
silver bromate	$AgBrO_3$	5.3×10^{-5}
silver bromide	AgBr	5.4×10^{-13}
silver carbonate	Ag_2CO_3	8.5×10^{-12}
silver chloride	AgCl	1.8×10^{-10}
silver chromate	Ag ₂ CrO ₄	1.1×10^{-12}
silver iodate	${\rm AgIO_3}$	3.2×10^{-8}
silver iodide	AgI	8.5×10^{-17}
strontium carbonate	SrCO ₃	5.6×10^{-10}
strontium fluoride	SrF ₂	4.3×10^{-9}
strontium sulphate	SrSO ₄	3.4×10^{-7}
zinc sulphide	ZnS	2.0×10^{-25}

RELATIVE STRENGTHS OF BRÖNSTED-LOWRY ACIDS AND BASES

in aqueous solution at room temperature

Strength of Acid	Name of Acid	Acid	Base K _a	Strength of Base
Strong	Perchloric	$\text{HClO}_{4} \rightarrow$	$H^+ + ClO_4^-$ very large	Weak
	Hydriodic		$H^+ + I^-$ very large	
	Hydrobromic		$H^+ + Br^-$ very large	
	Hydrochloric		$H^+ + Cl^-$ very large	
	Nitric		$H^+ + NO_3^-$ very large	
	Sulphuric	$_{\text{H}_2\text{SO}_4} \rightarrow$	$H^+ + HSO_4^-$ very large	
	Hydronium Ion	2 .	$H^+ + H_2O$ 1.0	
	Iodic	3	$H^+ + IO_3^- \dots 1.7 \times 10^{-1}$	
	Oxalic		$H^+ + HC_2O_4^-$ 5.9×10 ⁻²	
	Sulphurous $(SO_2 + H_2O)$		$H^+ + HSO_3^- \dots 1.5 \times 10^{-2}$	
	Hydrogen sulphate ion	$HSO_4^- \iff$	$H^+ + SO_4^{2-}$ 1.2×10^{-2}	
	Phosphoric	•	$H^+ + H_2 PO_4^- \dots 7.5 \times 10^{-3}$	
	Hexaaquoiron ion, iron(III) ion	3 7	$H^+ + Fe(H_2O)_5(OH)^{2+}$ 6.0×10 ⁻³	
	Citric		$H^+ + H_2C_6H_5O_7^- \dots 7.1 \times 10^{-4}$	
	Nitrous	3 0 3 7	$H^+ + NO_2^- \dots 4.6 \times 10^{-4}$	
	Hydrofluoric	_	$H^+ + F^-$	
	Methanoic, formic	нсоон ↔	$H^+ + HCOO^- \dots 1.8 \times 10^{-4}$	
	Hexaaquochromium ion, chromium(III) ion	$Cr(H_2O)_6^{3+} \iff$	$H^+ + Cr(H_2O)_5(OH)^{2+}$ 1.5×10 ⁻⁴	
	Benzoic	` , -	$H^+ + C_6 H_5 COO^-$ 6.5×10 ⁻⁵	
	Hydrogen oxalate ion	0 5	$H^+ + C_2O_4^{2-}$ 6.4×10 ⁻⁵	
	Ethanoic, acetic	CH₃COOH ←	H ⁺ + CH ₃ COO ⁻ 1.8×10 ⁻⁵	
	Dihydrogen citrate ion	$H_2C_6H_5O_7^- \iff$	$H^+ + HC_6H_5O_7^{2-}$	
	Hexaaquoaluminum ion, aluminum ion	$Al(H_2O)^{3+} \iff$	$H^+ + Al(H_2O)_5(OH)^{2+}$ 1.4×10 ⁻⁵	
	Carbonic $(CO_2 + H_2O)$		$H^+ + HCO_3^- \dots 4.3 \times 10^{-7}$	
	Monohydrogen citrate ion	$HC_6H_5O_7^{2-} \iff$	$H^+ + C_6 H_5 O_7^{3-}$ 4.1×10^{-7}	
	Hydrogen sulphite ion	$HSO_3^- \iff$	$H^+ + SO_3^{2-}$ 1.0×10^{-7}	
	Hydrogen sulphide	$H_2S \iff$	$H^+ + HS^- \dots 9.1 \times 10^{-8}$	
	Dihydrogen phosphate ion	$H_2PO_4^- \iff$	$H^+ + HPO_4^{2-}$ 6.2×10 ⁻⁸	
	Boric	$H_3BO_3 \iff$	$H^+ + H_2BO_3^- \dots 7.3 \times 10^{-1}$	0
	Ammonium ion		$H^+ + NH_3 \dots 5.6 \times 10^{-1}$	
	Hydrocyanic	HCN ←	$H^+ + CN^- \dots 4.9 \times 10^{-1}$	0
	Phenol	$C_6H_5OH \iff$	$H^+ + C_6 H_5 O^- \dots 1.3 \times 10^{-10}$)
	Hydrogen carbonate ion		$H^+ + CO_3^{2-}$ 5.6×10 ⁻¹	
	Hydrogen peroxide	$H_2O_2 \iff$	$H^+ + HO_2^-$ 2.4×10 ⁻¹	2
	Monohydrogen phosphate ion	$HPO_4^{2-} \iff$	$H^+ + PO_4^{3-}$ 2.2×10 ⁻¹	3
	Water	•	$H^+ + OH^ 1.0 \times 10^{-14}$	
	Hydroxide ion	OH⁻ ←	$H^+ + O^{2-}$ very small	
	Ammonia	$NH_3 \leftarrow$	$H^+ + NH_2^-$ very small	\downarrow
Weak		<u> </u>		Strong

ACID-BASE INDICATORS

INDICATOR	pH RANGE IN WHICH COLOUR CHANGE OCCURS	COLOUR CHANGE AS pH INCREASES
Methyl violet	0.0 – 1.6	yellow to blue
Thymol blue	1.2 – 2.8	red to yellow
Orange IV	1.4 – 2.8	red to yellow
Methyl orange	3.2 – 4.4	red to yellow
Bromcresol green	3.8 - 5.4	yellow to blue
Methyl red	4.8 – 6.0	red to yellow
Chlorophenol red	5.2 - 6.8	yellow to red
Bromthymol blue	6.0 – 7.6	yellow to blue
Phenol red	6.6 - 8.0	yellow to red
Neutral red	6.8 - 8.0	red to amber
Thymol blue	8.0 – 9.6	yellow to blue
Phenolphthalein	8.2 - 10.0	colourless to pink
Thymolphthalein	9.4 – 10.6	colourless to blue
Alizarin yellow	10.1 – 12.0	yellow to red
Indigo carmine	11.4 – 13.0	blue to yellow