

JUNE 1998

PROVINCIAL EXAMINATION

MINISTRY OF EDUCATION

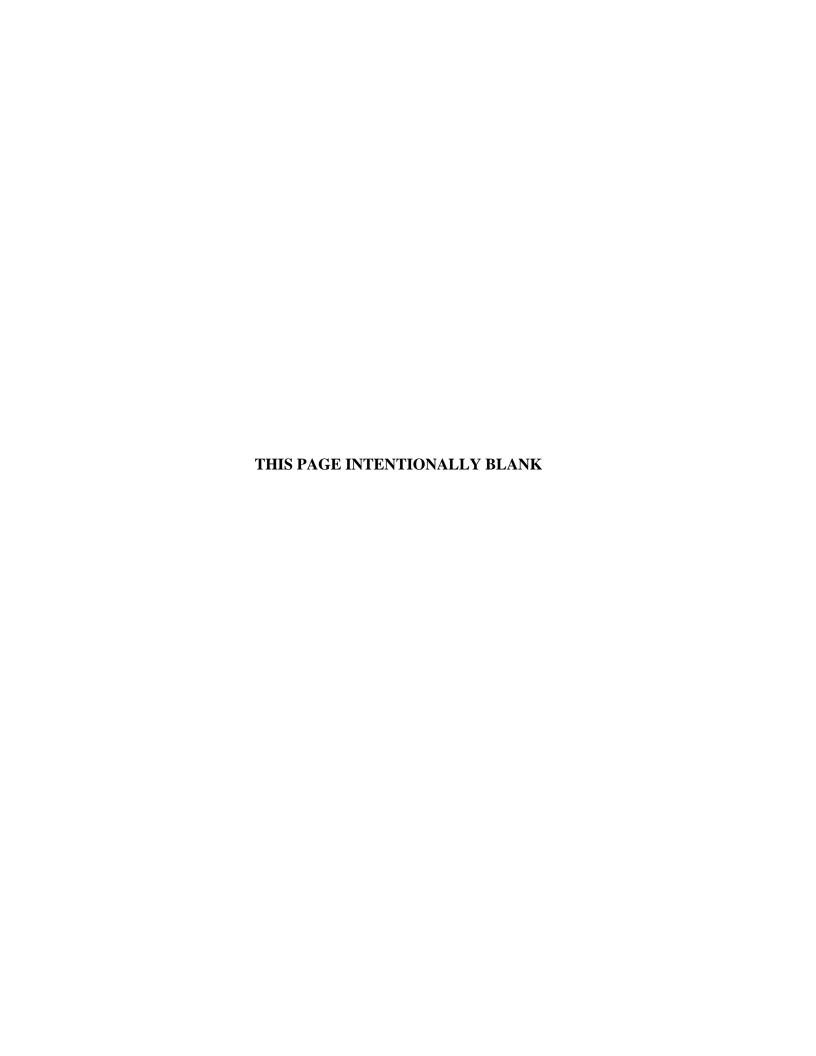
CHEMISTRY 12

GENERAL INSTRUCTIONS

- 1. Insert the stickers with your Student I.D. Number (PEN) in the allotted spaces above and on the back cover of this booklet. Under no circumstance is your name or identification, other than your Student I.D. Number, to appear on this booklet.
- 2. Ensure that in addition to this examination booklet, you have a **Data Booklet** and an **Examination Response Form**. Follow the directions on the front of the Response Form.
- 3. **Disqualification** from the examination will result if you bring books, paper, notes or unauthorized electronic devices into the examination room.
- 4. All multiple-choice answers must be entered on the Response Form using an **HB pencil**. Multiple-choice answers entered in this examination booklet will **not** be marked.
- 5. For each of the written-response questions, write your answer in the space provided in this booklet.
- 6. 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 .

7. At the end of the examination, place your Response Form inside the front cover of this booklet and return the booklet and your Response Form to the supervisor.



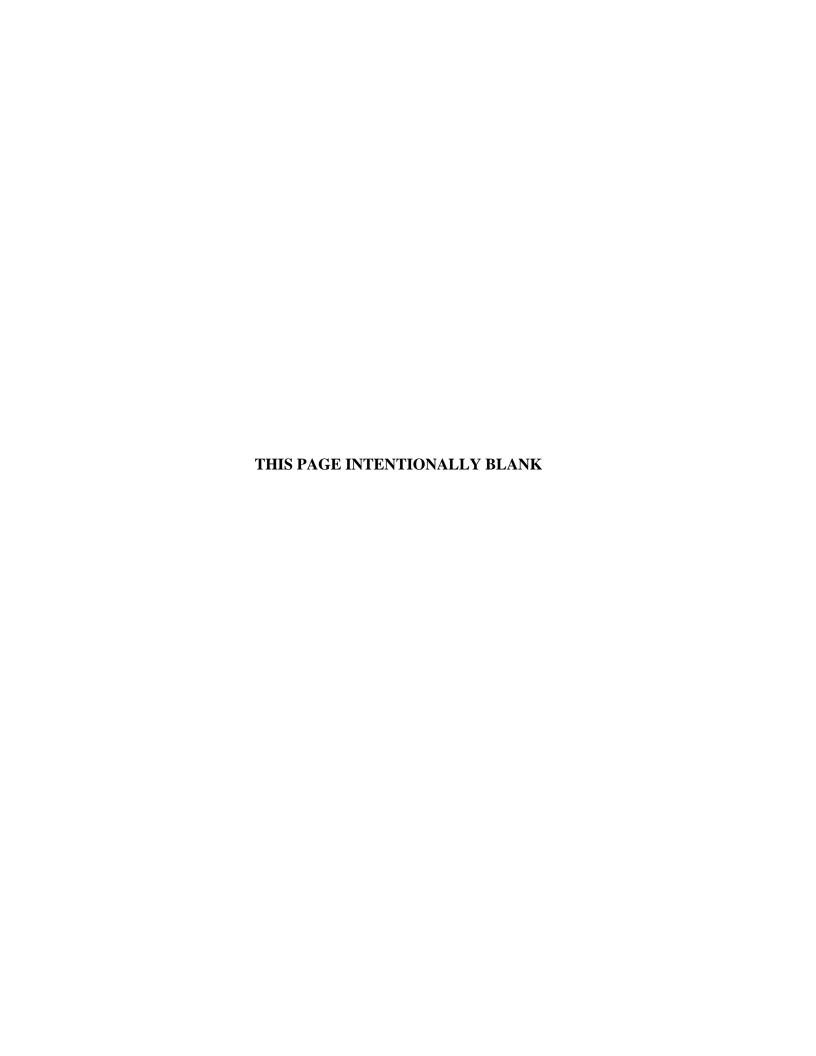
CHEMISTRY 12 PROVINCIAL EXAMINATION

1.	This exami	nation consists of two parts:		Value	Suggested Time
	PART A:	48 multiple-choice questions		48	70
	PART B:	11 written-response questions		32	50
			Total:	80 marks	120 minutes

- 2. Aside from an approved calculator, electronic devices, including dictionaries and pagers, are **not** permitted in the examination room.
- 3. 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.

- 4. A calculator is essential for the Chemistry 12 Provincial Examination. The calculator must be a hand-held device designed primarily for mathematical computations involving logarithmic and trigonometric functions. Computers, calculators with a QWERTY keyboard, and electronic writing pads will not be allowed. Students must not bring any external devices to support calculators such as manuals, printed or electronic cards, printers, memory expansion chips or cards, or external keyboards. Students may have more than one calculator available during the examination. Calculators may not be shared, and communication between calculators is prohibited during the examination. In addition to an approved calculator, students will be allowed to use rulers, compasses, and protractors during the examination.
- 5. The time allotted for this examination is **two hours**.



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 Response

Form provided. Using an HB pencil, completely fill in the circle that has the letter corresponding to your answer.

1. Which of the following reactions occurs most rapidly at room temperature?

A.
$$H_{2(g)} + I_{2(g)} \to 2HI_{(g)}$$

B.
$$2\operatorname{Fe}_{(s)} + \operatorname{O}_{2(g)} \to 2\operatorname{FeO}_{(s)}$$

C.
$$Cu^{2+}(aq) + S^{2-}(aq) \to CuS_{(s)}$$

D.
$$C_6H_{12}O_{6(aq)} + 6O_{2(g)} \rightarrow 6CO_{2(g)} + 6H_2O_{(g)}$$

2. Consider the following reaction:

$$CO + NO_2 \rightarrow CO_2 + NO$$
 $\Delta H = -234 \text{ kJ}$

The activation energy of the forward reaction is 134 kJ. What is the activation energy for the reverse reaction?

B.
$$-100 \text{ kJ}$$

- 3. Which of the following will decrease the number of effective collisions during a chemical reaction?
 - A. Adding a catalyst.
 - B. Increasing the surface area.
 - C. Decreasing the temperature.
 - D. Increasing reactant concentrations.

4. Consider the following reaction:

$$2\mathrm{N}_2\mathrm{O}_{5(g)} + 110~\mathrm{kJ} \rightarrow 4\mathrm{NO}_{2(g)} + \mathrm{O}_{2(g)}$$

The rate of reaction is increased by

- A. adding a catalyst.
- B. removing some O_2 .
- C. decreasing the temperature.
- D. increasing the volume of the container.

5. Which of the following apply to all equilibrium systems?

I Forward and reverse rates are equal	
II	Macroscopic properties are constant
III	Mass of reactants equals mass of products

- A. I and II only
- B. I and III only
- C. II and III only
- D. I, II and III

6. Consider the following equilibrium:

$$N_{2(g)} + 3H_{2(g)} \rightleftharpoons 2NH_{3(g)} + 92 \text{ kJ}$$

The forward reaction is

- A. exothermic and entropy is increasing.
- B. exothermic and entropy is decreasing.
- C. endothermic and entropy is increasing.
- D. endothermic and entropy is decreasing.

7. Consider the following equilibrium:

$$PCl_{5(g)} \rightleftharpoons PCl_{3(g)} + Cl_{2(g)}$$

The equilibrium concentration of PCl₅ will increase when

- A. PCl₃ is added.
- B. Cl₂ is removed.
- C. a catalyst is added.
- D. the volume of the container is increased.
- 8. Consider the following equilibrium:

$$N_2O_{4(g)} \rightleftharpoons 2NO_{2(g)}$$

If the volume of the container is decreased, the

- A. K_{eq} decreases.
- B. $[N_2O_4]$ increases.
- C. equilibrium does not shift.
- D. equilibrium shifts to the right.
- 9. Consider the following equilibrium:

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

For this reaction,

A.
$$K_{eq} = [CO_2]$$

B.
$$K_{eq} = \frac{1}{[CO_2]}$$

C.
$$K_{eq} = \frac{[CaCO_3]}{[CO_2][CaO]}$$

D.
$$K_{eq} = \frac{[CO_2][CaO]}{[CaCO_3]}$$

10. Which of the following reactions most favours products?

	REACTION	K_{eq}
I	$2SO_{2(g)} + O_{2(g)} \rightleftharpoons 2SO_{3(g)}$	2.6×10^2
II	$2 \operatorname{NO}_{(g)} + \operatorname{O}_{2(g)} \ \rightleftarrows \ 2 \operatorname{NO}_{2(g)}$	6.4×10^5
III	$2\operatorname{CO}_{(g)} + \operatorname{O}_{2(g)} \ \rightleftarrows \ 2\operatorname{CO}_{2(g)}$	2.5×10 ¹⁵
IV	$2H_{2(g)} + O_{2(g)} \rightleftharpoons 2H_2O_{(g)}$	1.7×10^{27}

- A. I
- B. II
- C. III
- D. IV

11. Consider the following:

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

At equilibrium in a 1.0 L container, there are 1.60×10^{-2} mol C, 1.50×10^{-2} mol H₂O, 3.00×10^{-1} mol CO, and 1.00×10^{-1} mol H₂. The value of K_{eq} is

- A. 0.500
- B. 2.00
- C. 80.0
- D. 125

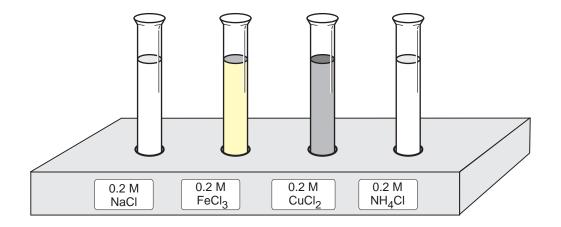
12. Consider the following equilibrium:

$$N_2O_{4(g)} \rightleftharpoons 2NO_{2(g)}$$
 $K_{eq} = 1.0 \times 10^{-2}$

At equilibrium, the $[NO_2] = 2.0 \times 10^{-2} \text{ mol/L}$ and the $[N_2O_4]$ is

- A. $4.0 \times 10^{-6} \text{ mol/L}$
- B. $4.0 \times 10^{-2} \text{ mol/L}$
- C. 2.0 mol/L
- D. 25 mol/L

- 13. Which of the following dissolves in water to form an ionic solution?
 - A. O₂
 - B. SiO₂
 - C. KMnO₄
 - D. $C_{12}H_{22}O_{11}$
- 14. A dilute solution of AgNO₃ is added dropwise to each of the following test tubes until a precipitate forms in each tube.



Which solution requires the lowest $\left[Ag^{+}\right]$ to just begin precipitation?

- A. NaCl
- B. FeCl₃
- $C. \quad CuCl_2$
- D. NH₄Cl
- 15. Which of the following is most soluble?
 - A. Na₂S
 - B. CaSO₄
 - C. PbCO₃
 - D. $Zn(OH)_2$

16.	The	solubility of PbI ₂ will increase with the addition of
	A.	PbI_2
	B.	heat.

D. $Pb(NO_3)_2$

C. water.

- 17. Which of the following saturated solutions has the lowest $\left[SO_4^{2-} \right]$ at 25°C?
 - A. SrSO₄
 - B. PbSO₄
 - C. CaSO₄
 - D. BaSO₄
- 18. The solubility of $MgCO_3$ is
 - A. $4.6 \times 10^{-11} \text{ M}$
 - B. 6.8×10^{-6} M
 - C. 1.4×10^{-5} M
 - D. 2.6×10^{-3} M
- 19. When equal volumes of 0.20 M Pb $(NO_3)_2$ and 0.20 M KI are mixed together,
 - A. a precipitate forms since Trial Ion Product $> K_{sp}$
 - B. a precipitate forms since Trial Ion Product $< K_{SD}$
 - C. no precipitate forms since Trial Ion Product $> K_{sp}$
 - D. no precipitate forms since Trial Ion Product $< K_{sp}$
- 20. Bubbles form when $Mg_{(s)}$ is added to $HCl_{(aq)}$. The gas produced is
 - A. O₂
 - B. H₂
 - C. Cl₂
 - D. H₂O

- 21. A substance which produces hydrogen ions in solution is the definition of
 - A. an Arrhenius acid.
 - B. an Arrhenius base.
 - C. a Brönsted-Lowry acid.
 - D. a Brönsted-Lowry base.
- 22. The number of conjugate pairs in a Brönsted-Lowry acid-base equation is
 - A. 1
 - B. 2
 - C. 3
 - D. 4
- 23. Which of the following is a stronger base than HPO_4^{2-} ?
 - A. H_2O
 - B. NO_2^-
 - C. CO_3^{2-}
 - D. $H_2PO_4^-$
- 24. Consider the following equilibria:

$$HTe^- + HSe^- \rightleftharpoons H_2Te + Se^{2-}$$

$$HTe^- + Se^{2-} \rightleftharpoons HSe^- + Te^{2-}$$

Reactants are favoured in both equilibria. The order of acids from strongest to weakest is

- A. HTe⁻, HSe⁻, H₂Te
- B. HSe⁻, H₂Te, HTe⁻
- C. H₂Te, HTe⁻, HSe⁻
- D. H₂Te, HSe⁻, HTe⁻

25. The equilibrium expression for the ion product constant of water is

A.
$$K_w = [H_3O^+][OH^-]$$

B.
$$K_w = \frac{1}{\left[H_3O^+\right]\left[OH^-\right]}$$

C.
$$K_w = \frac{[H_2O]^2}{[H_3O^+][OH^-]}$$

D.
$$K_w = \frac{[H_3O^+][OH^-]}{[H_2O]^2}$$

26. Calculate the pH of 0.01 M NaOH.

A.
$$1.0 \times 10^{-12}$$

B.
$$1.0 \times 10^{-2}$$

27. Consider the following:

TEMPERATURE	K_w
25°C	1.0×10^{-14}
50°C	5.5×10^{-14}

When water is heated,

A.
$$\left[H_3 O^+ \right]$$
 and $\left[OH^- \right]$ both increase.

B.
$$\left[H_3O^+\right]$$
 and $\left[OH^-\right]$ both decrease.

C.
$$\left[H_3O^+\right]$$
 increases and $\left[OH^-\right]$ decreases.

D.
$$\left[H_3 O^+ \right]$$
 decreases and $\left[OH^- \right]$ increases.

28. The $[H_3O^+]$ of a solution with a pOH of 4.60 is

A.
$$4.0 \times 10^{-10}$$
 M

B.
$$2.5 \times 10^{-5}$$
 M

C.
$$6.6 \times 10^{-1}$$
 M

D.
$$9.7 \times 10^{-1}$$
 M

29. The K_b expression for aniline, $C_6H_5NH_2$, is

A.
$$K_b = \left[C_6 H_5 N H_3^+\right] \left[O H^-\right]$$

B.
$$K_b = \left[C_6 H_5 N H^- \right] \left[H_3 O^+ \right]$$

C.
$$K_b = \frac{\left[C_6 H_5 N H_3^+\right] \left[O H^-\right]}{\left[C_6 H_5 N H_2\right]}$$

D.
$$K_b = \frac{\left[C_6 H_5 N H^- \right] \left[H_3 O^+ \right]}{\left[C_6 H_5 N H_2 \right]}$$

30. A solution of $0.10 \text{ M HPO}_4^{2-}$ is

A. acidic because
$$K_a < K_b$$

B. acidic because
$$K_a > K_b$$

C. basic because
$$K_b < K_a$$

D. basic because
$$K_b > K_a$$

31. Consider the following equilibrium:

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

At the transition point for an acid-base indicator,

A.
$$K_a = [Ind^-]$$

B.
$$K_a = [HInd]$$

C.
$$K_a = [H_3O^+]$$

D.
$$K_a = [H_3O^+][OH^-]$$

- 32. A solution is amber with neutral red and colourless with phenolphthalein. The approximate pH of the solution is
 - A. 4
 - B. 6
 - C. 8
 - D. 10
- 33. When a 1.0 M acidic solution is titrated with a 1.0 M basic solution, the pH at the equivalence point is 8.5. The reactants could be
 - A. HBr and KOH
 - B. HNO₃ and NH₃
 - C. HCl and NaOH
 - D. HNO₂ and NaOH
- 34. The net ionic equation for the reaction between $KOH_{(aq)}$ and $HBr_{(aq)}$ is
 - A. $K^+_{(aq)} + Br^-_{(aq)} \rightarrow KBr_{(s)}$
 - B. $H^+_{(aq)} + OH^-_{(aq)} \rightarrow H_2O_{(\ell)}$
 - C. $KOH_{(aq)} + HBr_{(aq)} \rightarrow KBr_{(aq)} + H_2O_{(\ell)}$
 - D. $K^{+}_{(aq)} + OH^{-}_{(aq)} + H^{+}_{(aq)} + Br^{-}_{(aq)} \to K^{+}_{(aq)} + Br^{-}_{(aq)} + H_{2}O_{(\ell)}$
- 35. Consider the following equilibrium for a buffer solution:

$$CH_3COOH + H_2O \rightleftharpoons CH_3COO^- + H_3O^+$$

When a small amount of acid is added to this system, and equilibrium is reestablished,

- A. $\left[\text{CH}_3 \text{COO}^- \right]$ and pH have both increased.
- B. [CH₃COOH] and pH have both decreased.
- C. $\left[\text{CH}_3 \text{COO}^- \right]$ has decreased and pH remains relatively constant.
- D. [CH₃COOH] has decreased and pH remains relatively constant.

- 36. Which of the following dissolves in water to form a basic solution?
 - A. NO
 - B. SO₃
 - C. CO₂
 - D. K₂O
- 37. Which of the following is an equation representing a redox reaction?
 - A. $2NO_{2(g)} \rightarrow N_2O_{4(g)}$
 - B. $\operatorname{Mg}_{(s)} + \operatorname{Cl}_{2(g)} \to \operatorname{MgCl}_{2(s)}$
 - C. $\operatorname{Ag}^+(aq) + \operatorname{Cl}^-(aq) \to \operatorname{AgCl}_{(s)}$
 - D. $NH_{3(aq)} + H^{+}_{(aq)} \rightarrow NH_{4(aq)}^{+}$
- 38. Consider the following:

$$O_3 + H_2O + SO_2 \rightarrow SO_4^{2-} + O_2 + 2H^+$$

- In the redox reaction above, the chemical species oxidized is
- A. H⁺
- B. O₃
- $C. SO_2$
- D. SO_4^{2-}
- 39. A product of the oxidation of NO_2 is
 - A. NO
 - B. N₂O
 - C. NO₂
 - D. NO₃

- 40. Which of the following is the strongest reducing agent?
 - A. Sn²⁺
 - B. Fe²⁺
 - C. Au³⁺
 - D. Mn²⁺
- 41. At standard conditions, a spontaneous reaction will occur between
 - A. Pb and Fe^{2+}
 - B. Cu and Sn²⁺
 - C. Cu^{2+} and Sn^{2+}
 - D. Pb^{2+} and Fe^{2+}
- 42. Consider the following:

$$2MnO_4^- + 5H_2O_2 + 6H^+ \rightarrow 5O_2 + 2Mn^{2+} + 8H_2O$$

In an experiment, 2.00×10^{-4} mol of MnO₄⁻ is required to titrate 10.0 mL of H₂O₂ to the equivalence point. The concentration of H₂O₂ is

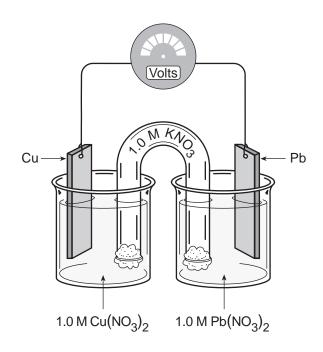
- A. 5.00×10^{-3} M
- B. 8.00×10^{-3} M
- C. $2.00 \times 10^{-2} \text{ M}$
- D. 5.00×10^{-2} M
- 43. A lead storage battery contains electrochemical cells which involve the following half-reactions:

$$PbO_2 + 4H^+ + SO_4^{2-} + 2e^- \rightleftharpoons PbSO_4 + 2H_2O$$
 $E^\circ = +1.69 \text{ V}$
 $PbSO_4 + 2e^- \rightleftharpoons Pb + SO_4^{2-}$ $E^\circ = -0.36 \text{ V}$

Use the data above to calculate the E° for one of these cells.

- A. $E^{\circ} = +1.33 \text{ V}$
- B. $E^{\circ} = -1.33 \text{ V}$
- C. $E^{\circ} = -2.05 \text{ V}$
- D. $E^{\circ} = +2.05 \text{ V}$

Use the following diagram to answer questions 44, 45 and 46.



44. In the electrochemical cell above, the half-reaction at the cathode is

A.
$$Pb \rightarrow Pb^{2+} + 2e^{-}$$

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

C.
$$Pb^{2+} + 2e^- \rightarrow Pb$$

D.
$$Cu^{2+} + 2e^{-} \rightarrow Cu$$

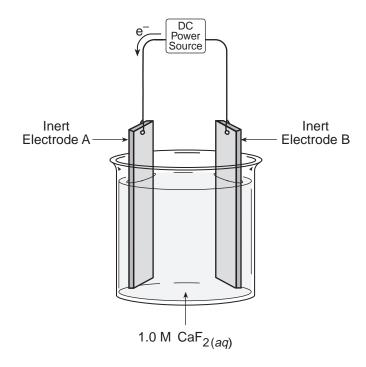
45. In the electrochemical cell above, the anode

- A. loses mass as Pb is oxidized.
- B. gains mass as Pb is oxidized.
- C. loses mass as Cu is oxidized.
- D. gains mass as Cu is oxidized.

46. In the salt bridge in the electrochemical cell above,

- A. electrons flow into the Pb half-cell.
- B. electrons flow into the Cu half-cell.
- C. anions migrate into the Pb half-cell.
- D. anions migrate into the Cu half-cell.

Use the following diagram to answer questions 47 and 48.



- 47. The predominant half-reaction occurring at Electrode B is
 - A. $2F^- \rightarrow F_2 + 2e^-$
 - B. $Ca^{2+} + 2e^{-} \rightarrow Ca$
 - C. $2H_2O \rightarrow O_2 + 4H^+ + 4e^-$
 - D. $2H_2O + 2e^- \rightarrow H_2 + 2OH^-$
- 48. In the electrolytic cell above, a suitable substance for Electrode B is
 - A. carbon.
 - B. lithium.
 - C. sodium.
 - D. calcium.

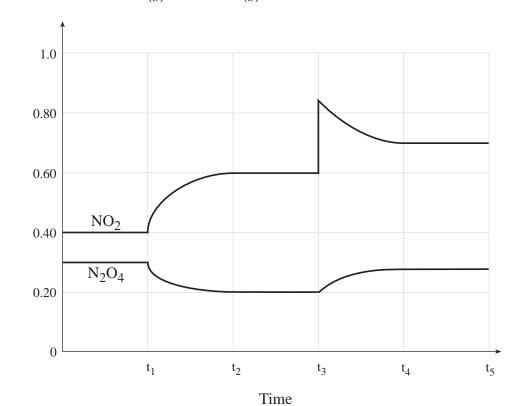
This is the end of the multiple-choice section. Answer the remaining questions directly in this examination booklet.

PART B: WRITTEN RESPONSE

Value: 32 marks	Sug	ggested Time: 50 minutes
INSTRUCTIONS:	You will be expected to communicate your knowledge an chemical principles in a clear and logical manner.	nd understanding of
	Your steps and assumptions leading to a solution must be below the questions.	written in the spaces
	Answers must include units where appropriate and be giv significant figures.	en to the correct number of
	For questions involving calculation, full marks will NO only an answer.	OT be given for providing
	es not always occur when two reactant particles collide. Gi	(2 marks)
ii)		
2. Consider the f	following reaction mechanism:	
	Step 1: $NO + NO \rightarrow N_2O_2$	
	Step 2: $N_2O_2 + H_2 \rightarrow N_2O + H_2O$	
	Step 3: $N_2O + H_2 \rightarrow N_2 + H_2O$	
a) Write the e	equation for the overall reaction.	(1 mark)
b) Identify the	e reaction intermediate(s).	(1 mark)

3. Consider the following graph for the reaction:

energy +
$$N_2O_{4(g)} \rightleftharpoons 2NO_{2(g)}$$



a) What is the stress imposed at time t_1 ?

Concentration (mol/L)

(1 mark)

b) What is the stress imposed at time t_3 ?

(1 mark)

c) Calculate $K_{\it eq}$ for the equilibrium between t_2 and t_3 .

(2 marks)

4. Consider the following net ionic equation:

$$Ag^{+}_{(aq)} + SCN^{-}_{(aq)} \rightarrow AgSCN_{(s)}$$

A 20.00 mL sample of 0.200 M $\,\mathrm{NH_4SCN}$ is used to titrate a 30.00 mL sample containing $\,\mathrm{Ag}^+$. Calculate the $\left[\mathrm{Ag}^+\right]$ in the original sample. (3 marks)

5. A solution contains $0.020 \, M \, Ba^{2+}$ and an unknown concentration of Sr^{2+} . When dilute Na_2CO_3 is slowly added to the mixture, both Ba^{2+} and Sr^{2+} start to precipitate at the same time.

(3 marks)

a) Calculate the $\left[{{{\rm{CO}}_3}^{2^ - }} \right]$ when ${{\rm{BaCO}}_3}$ starts to precipitate.

b) Calculate the initial $\left[Sr^{2+} \right]$.

6.	a) Define the term <i>amphiprotic</i> .	(1 mark)

b) Give an example of an amphiprotic anion.

(1 mark)

7. The salt Na_2CO_3 undergoes hydrolysis to produce a basic solution. Calculate the $\left[OH^-\right]$ in 0.100 M Na_2CO_3 . (4 marks)

8. A student titrated a 25.00 mL sample of a 0.20 M HX (unknown) acid with 0.20 M NaOH. The following data were collected.

Volume of base added (mL)	рН
0.00	2.72
10.00	4.57
24.90	7.14
24.99	8.14
25.00	8.88
25.01	9.60
26.00	11.59
35.00	12.52

a)	Describe the acid HX as strong or weak. Support your answer with two observations from the data table. (3 marks)
b)	Select an appropriate indicator for this titration and identify the colour at the equivalence point. (1 mark)

9. Balance the following redox reaction:

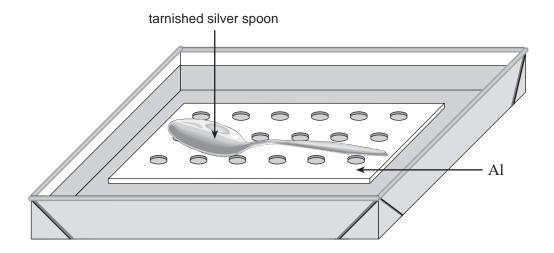
(3 marks)

$$Ni + ClO_4^- \rightarrow Ni^{2+} + Cl^-$$
 (acid)

10. Define the term *electrolysis*.

(2 marks)

11. Consider the following diagram:



On a silver spoon, the black tarnish, Ag_2S , can be removed spontaneously by placing the spoon in contact with aluminum in a conducting solution.

a) Write the equations for the two half-reactions.

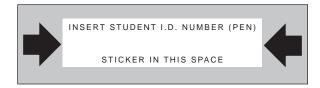
(2 marks)

b) Write the equation for the redox reaction.

(1 mark)

END OF EXAMINATION

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CHEMISTRY 12

June 1998

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CHEMISTRY 12

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Score fo	r
Question	1

1. _____

Score for Question 8:

8. ____

Score for Question 2:

2. _____(2)

Score for Question 9:

9. ____

Score for Question 3:

3. _____

Score for Question 10:

10. ____

Score for Question 4:

4. ____(3)

Score for Question 11:

11. _____(3)

Score for Question 5:

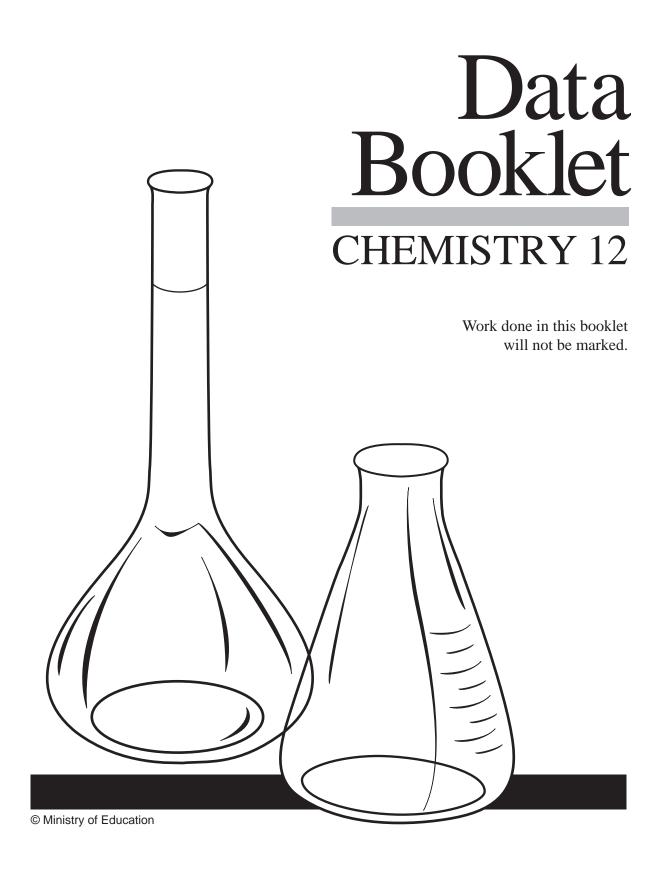
5. _____

Score for Question 6:

6. _____

Score for Question 7:

7. _____



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6	Relative Strengths of Brönsted-Lowry Acids and Bases
7	Acid-Base Indicators
8	Standard Reduction Potentials of Half-Cells

REFERENCE

1	PERIODIC TABLE OF THE ELEMENTS										18						
H Hydrogen 1.0																	2 He Helium
	2	ı										13	14	15	16	17	4.0
3	4					14 -	Aton	nic number				5	6	7	8	9	10
Li Lithium	Be Beryllium					Si -	Sym					В	C	N	О	F	Ne
6.9	9.0					Silicon - 28.1 -	Nam Aton	nic mass				Boron 10.8	Carbon 12.0	Nitrogen 14.0	Oxygen 16.0	Fluorine 19.0	Neon 20.2
11	12											13	14	15	16	17	18
Na	Mg											Al	Si	P	S	Cl	Ar
Sodium	Magnesium			_		_						Aluminum 27.0	Silicon 28.1	Phosphorus 31.0	Sulphur 32.1	Chlorine 35.5	Argon 39.9
23.0	24.3	3	4	5	6	7	8	9	10	11	12			-			
19 K	Ca	21 Sc	22 T i	23 V	24 Cr	25 Mn	26 Fe	27 Co	28 Ni	Cu	30 Z n	Ga 31	Ge 32	33 As	34 Se	35 Br	36 Kr
Potassium	Calcium	Scandium	I I Titanium	V Vanadium	Chromium	Manganese		Cobalt	Nickel	Cu	ZII	Gallium	Germanium	AS Arsenic	Selenium	Bromine	Krypton
39.1	40.1	45.0	47.9	50.9	52.0	54.9	55.8	58.9	58.7	63.5	65.4	69.7	72.6	74.9	79.0	79.9	83.8
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr	Y	Zr	Nb	Mo	Тс	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe
Rubidium 85.5	Strontium 87.6	Yttrium 88.9	Zirconium 91.2	Niobium 92.9	Molybdenum 95.9	Technetium (98)	Ruthenium 101.1	Rhodium 102.9	Palladium 106.4	Silver 107.9	Cadmium 112.4	Indium 114.8	Tin 118.7	Antimony 121.8	Tellurium 127.6	Iodine 126.9	Xenon 131.3
55	56	57	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba	La	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn
Cesium	Barium	Lanthanum	Hafnium	Tantalum	Tungsten	Rhenium	Osmium	Iridium	Platinum	Gold	Mercury	Thallium	Lead	Bismuth	Polonium	Astatine	Radon
132.9	137.3	138.9	178.5	180.9	183.8	186.2	190.2	192.2	195.1	197.0	200.6	204.4	207.2	209.0	(209)	(210)	(222)
87	88	89	104	105	106	107	108	109									
Fr Francium	Ra Radium	Ac Actinium	Rf Rutherfordium	Ha Hahnium	Sg Seaborgium	Uns	Uno Unniloctium	Une Unnilennium									
(223)	(226)	(227)	(261)	(262)	(263)	Unnilseptium (262)	(265)	(266)									
Based on	Based on mass of C^{12} at 12.00.																
			\ 1	58	59	60	61	62	63	64	65	66	67	68	69	70	71
	Values in parentheses are the masses of the most stable or best			Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Но	Er	Tm	Yb	Lu
known isc	otopes for	elements	(31)	Cerium 140.1	Praseodymium 140.9	Neodymium 144.2	Promethium (145)	Samarium 150.4	Europium 152.0	Gadolinium 157.3	Terbium 158.9	Dysprosium 162.5	Holmium 164.9	Erbium 167.3	Thulium 168.9	Ytterbium 173.0	Lutetium 175.0
which do	not occur	naturally.	\	90	91	92	93	94	95	96	97	98	99	100	101	102	103
			\	Th	Pa	Ü	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr
					Protactinium	Uranium	Neptunium	Plutonium	Americium	Curium	Berkelium	Californium	Einsteinium	Fermium	Mendelevium	Nobelium	Lawrencium
	232.0 231.0 238.0 (237) (244) (243) (247) (247) (251) (252) (257) (258) (259) (262)																

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	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
Bismuth	Bi	83	209.0	Oxygen	O	8	16.0
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	На	105	(262)	Technetium	Tc	43	(98)
Helium	He	2	4.0	Tellurium	Te	52	127.6
Holmium	Но	6 7	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	39	88.9
Magnesium	Mg	12	24.3	Zinc	Zn	39	65.4
Manganese	Mn	25	24.3 54.9	Zirconium	Zn	40	91.2
	Md	101	(258)	Ziicomuni	Δl	40	91.2
Mendelevium	IVIU	101	(238)				

NAMES, FORMULAE, AND CHARGES OF SOME COMMON IONS

Positive ions (ca	tions)	Negative ions (anions)			
Aluminum	Al^{3+}	Bromide	Br ⁻		
Ammonium	$NH_4^{}$	Carbonate	CO_3^{2-}		
Barium	Ba^{2+}	Chlorate	ClO ₃		
Calcium	Ca ²⁺	Chloride	Cl ⁻		
Chromium(II), chromous	Cr ²⁺	Chlorite	ClO ₂		
Chromium(III), chromic	Cr ³⁺	Chromate	CrO_2 CrO_4^{2-}		
Copper(I)*, cuprous	Cu^+		•		
Copper(II), cupric	Cu^{2+}	Cyanide	CN ⁻		
Hydrogen	H^+	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	Mg^{2+}	Hydrogen sulphide, bisulphide	HS ⁻		
Manganese(II), manganous	Mn ²⁺	Hydrogen sulphite, bisulphite	HSO ₃		
Manganese(IV)	Mn ⁴⁺				
Mercury(I)*, mercurous	${\rm Hg_2}^{2+}$	Hydroxide	OH ⁻		
Mercury(II), mercuric	Hg^{2+}	Hypochlorite	ClO-		
Potassium	\mathbf{K}^{+}	Iodide	Ι-		
Silver	Ag^+	Monohydrogen phosphate	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	Zn^{2+}	Oxide**	O^{2-}		
* Aqueous solutions are readily or	kidized by air.	Perchlorate	ClO ₄		
** Not stable in aqueous solutions.		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
H 1 11 0H	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	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	$HClO_4 \rightarrow$	$H^+ + ClO_4^-$ very large	Weak
1	Hydriodic	•	$H^+ + I^-$ very large	
	Hydrobromic		H ⁺ + Br ⁻ very large	
	Hydrochloric		$H^+ + Cl^-$ very large	
	Nitric		$H^+ + NO_3^-$ very large	
	Sulphuric	,	$H^+ + HSO_4^-$ very large	
	Hydronium Ion	2 .	$H^+ + H_2O$	
	Iodic	5	$H^+ + IO_3^-$ 1.7×10 ⁻¹	
	Oxalic		$H^+ + HC_2O_4^-$ 5.9×10 ⁻²	
	Sulphurous $(SO_2 + H_2O)$		$H^+ + HSO_3^ 1.5 \times 10^{-2}$	
	Hydrogen sulphate ion		$H^+ + SO_4^{2-}$ 1.2×10^{-2}	
	Phosphoric	•	$H^+ + H_2PO_4^-$ 7.5×10 ⁻³	
	Hexaaquoiron ion, iron(III) ion	$Fe(H_2O)^{3+} \iff$	$H^+ + Fe(H_2O)_5(OH)^{2+}$ 6.0×10 ⁻³	;
	Citric	` '0	$H^+ + H_2C_6H_5O_7^-$ 7.1×10 ⁻⁴	
	Nitrous	5 0 5 7	$H^+ + NO_2^ 4.6 \times 10^{-4}$	
	Hydrofluoric	$_{\mathrm{HF}}\overset{_{_{_{_{_{_{_{_{_{_{_{_{_{_{_{_{}}}}}}}$	$H^+ + F^- = 3.5 \times 10^{-4}$,
	Methanoic, formic	нсоон ↔	$H^+ + HCOO^- \dots 1.8 \times 10^{-4}$	
	Hexaaquochromium ion, chromium(III) ion	$Cr(H_2O)_6^{3+} \iff$	$H^+ + Cr(H_2O)_5(OH)^{2+} \dots 1.5 \times 10^{-4}$	
	Benzoic	, , ,	$H^+ + C_6 H_5 COO^-$ 6.5×10 ⁻⁵	
	Hydrogen oxalate ion		$H^+ + C_2O_4^{2-}$ 6.4×10 ⁻⁵	
	Ethanoic, acetic	CH₃COOH ←	$H^+ + CH_3COO^-$ 1.8×10 ⁻⁵	
	Dihydrogen citrate ion	$H_2C_6H_5O_7^- \iff$	$H^+ + HC_6H_5O_7^{2-}$ 1.7×10 ⁻⁵	
	Hexaaquoaluminum ion, aluminum ion	$Al(H_2O)_6^{3+} \iff$	$H^+ + Al(H_2O)_5(OH)^{2+} \dots 1.4 \times 10^{-5}$	
	Carbonic $(CO_2 + H_2O)$		$H^+ + HCO_3^- \dots 4.3 \times 10^{-7}$	
	Monohydrogen citrate ion	$HC_6H_5O_7^{2-} \iff$	$H^+ + C_6 H_5 O_7^{3-} \dots 4.1 \times 10^{-7}$	
	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	3
	Boric	$H_3BO_3 \iff$	$H^+ + H_2BO_3^ 7.3 \times 10^{-1}$.0
	Ammonium ion	$NH_4^+ \iff$	$H^+ + NH_3$ 5.6×10 ⁻¹	.0
	Hydrocyanic	$HCN \iff$	$H^+ + CN^- \dots 4.9 \times 10^{-1}$	10
	Phenol	$C_6H_5OH \iff$	$H^+ + C_6 H_5 O^- \dots 1.3 \times 10^{-10}$	0
	Hydrogen carbonate ion	$HCO_3^- \iff$	$H^+ + CO_3^{2-}$ 5.6×10 ⁻¹	.1
	Hydrogen peroxide	$H_2O_2 \iff$	$H^+ + HO_2^-$ 2.4×10 ⁻¹	12
	Monohydrogen phosphate ion	$HPO_4^{2-} \iff$	$H^+ + PO_4^{3-} \dots 2.2 \times 10^{-1}$	13
	Water	$H_2O \iff$	$H^+ + OH^- \dots 1.0 \times 10^{-1}$	4
	Hydroxide ion	OH⁻ ←	$H^+ + O^{2-}$ very small	I
	Ammonia	$NH_3 \leftarrow$	$H^+ + NH_2^-$ very small	ı
Weak				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

STANDARD REDUCTION POTENTIALS OF HALF-CELLS

Ionic Concentrations are at 1M in Water at 25° C

STRENGTH OF OXIDIZING AGENT	OXIDIZING AGENTS		REDUCING AGENTS	E*(VOLTS)	STRENGTH OF REDUCING AGENT
strong			2F ⁻		weak
↑			2SO ₄ ²⁻		
			2H ₂ O		
			$Mn^{2+} + 4H_2O$		
	$Au^{3+} + 3e^{-}$	\rightleftharpoons	Au _(s)	+1.50	
			$\frac{1}{2}Br_{2(1)} + 3H_2O$		
			$C1^- + 4H_2O$		_
	$Cl_{2(g)} + 2e^{-}$	\rightleftharpoons	2C1 ⁻	+1.36	
	$Cr_2O_7^{2-} + 14H^+ + 6e^-$				gc
			H ₂ O		Overpotential Effect
			$Mn^{2+} + 2H_2O$		i le
			$\frac{1}{2} I_{2(s)} + 3H_2O \dots$		inti
	= (-)		2Br ⁻		ote
			$Au_{(s)} + 4C1^{-}$: dr
	$NO_3^- + 4H^+ + 3e^-$	\rightleftharpoons	$NO_{(g)} + 2H_2O$	+0.96	
	$Hg^{2+} + 2e^{-}$	\rightleftharpoons	Hg _(l)	+0.85	
	$\frac{1}{2}$ O _{2(g)} + 2H ⁺ (10 ⁻⁷ M)+ 2e ⁻	\rightleftharpoons	H ₂ O	+0.82	"i
			N ₂ O ₄ + 2H ₂ O		
			Ag _(s)		
			Hg ₍₁₎		
	$Fe^{3+} + e^{-}$	\rightleftharpoons	Fe ²⁺	+0.77	
			H ₂ O ₂		
			$MnO_{2(s)} + 4OH^{-}$		
			2I ⁻		
			Cu _(s)		
			$S_{(s)} + 3H_2O$		
			Cu _(s)		
	$SO_4^{2-} + 4H^+ + 2e^-$	\rightarrow	$H_2SO_3 + H_2O$	+0.17	
			Cu ⁺		
			Sn ²⁺		
			H ₂ S _(g)		
			H _{2(g)}		
	$Pb^{2+} + 2e^{-}$	→	Pb _(s)	0.13	
	$Sn^{2+} + 2e^{-}$	\rightarrow	Sn _(s)	=0.14	
	$Ni^{2+} + 2e^{-}$	\rightarrow	Ni _(s)	-0.26	
	$H_{2}PO_{4} + 2H^{+} + 2e^{-}$	$\stackrel{\longleftarrow}{\leftarrow}$	$H_3PO_3 + H_2O$	-0.28	
	$Co^{2+} + 2e^{-}$		Co ₍₈₎		
		`	H ₂ Se		
	2.1		Cr ²⁺		
	2H ₂ O+ 2e ⁻	•	$H_2 + 2OH^-(10^{-7}M)$		
	\Box $Fe^{2+} + 2e^{-}$		Fe _(s)		
	$\begin{array}{cccccccccccccccccccccccccccccccccccc$		$2Ag_{(s)} + S^{2-}$		
	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		$\operatorname{Cr}_{(s)}$		
	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		$\operatorname{Zn}_{(s)}$		
	$Te_{(s)} + 2H^{+} + 2e^{-}$	-	H ₂ Te		
	O 2H_O_ 2e-		$H_{2(g)} + 2OH^{-}$		
			Mn _(s)		
			Al _(s)		
	$M_{\alpha}^{2+} + 2e^{-}$	$\stackrel{\leftarrow}{\rightarrow}$	$Mg_{(s)}$	-2 37	
	$N_2^+ \perp a^-$	$\leftarrow \rightarrow$	Na _(s)	-2 71	
	C_2^{2+} C_2^{2-}	$\leftarrow \rightarrow$	Ca _(s)	-2 87	
	Sr2+ 20-	$\stackrel{\leftarrow}{\leftarrow}$	$Sr_{(s)}$	-2.80	
	R 2 ²⁺ + 22-	\leftarrow	Ba _(s)	-2 Q1	
	V+	\leftarrow	K _(s)	-2 93	
	D b +	\leftarrow	Rb _(s)	-2.08	
	$\mathbf{K}\mathbf{D} + \mathbf{e}$	✓	Cs _(s)	-2 02	\downarrow
weak			Cs _(s)		strong
Weak	LI + e		L1(8)	3.04	