MINISTRY USE ONLY



Place Personal Education Number (PEN) here.

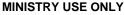












		_

Chemistry 12 AUGUST 2002

Course Code = CH

Student Instructions

- 1. Place the stickers with your Personal Education Number (PEN) in the allotted spaces above. Under no circumstance is your name or identification, other than your Personal Education 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. 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 .

5. 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.

Question 1:	Question 8:
1. (3)	8. [3]
Question 2: 2	Question 9: 9. (4)
Question 3:	Question 10:
3 (2)	10 (3)
Overtion 4:	Ougstion 11:
Question 4: 4 (4)	Question 11: 11. (4)
Question 5:	Question 12:
5.	
(3)	12 (2)
(3)	(2)
(3) Question 6: 6. (3)	(2) Question 13:
(3) Question 6: 6	(2) Question 13:

CHEMISTRY 12

AUGUST 2002

COURSE CODE = CH

GENERAL INSTRUCTIONS

- 1. Aside from an approved calculator, electronic devices, including dictionaries and pagers, are **not** permitted in the examination room.
- 2. 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.
- 3. For each of the written-response questions, write your answer in the space provided in this booklet.
- 4. Ensure that you use language and content appropriate to the purpose and audience of this examination. Failure to comply may result in your paper being awarded a zero.
- 5. This examination is designed to be completed in **two hours**. *Students may, however, take up to 30 minutes of additional time to finish.*

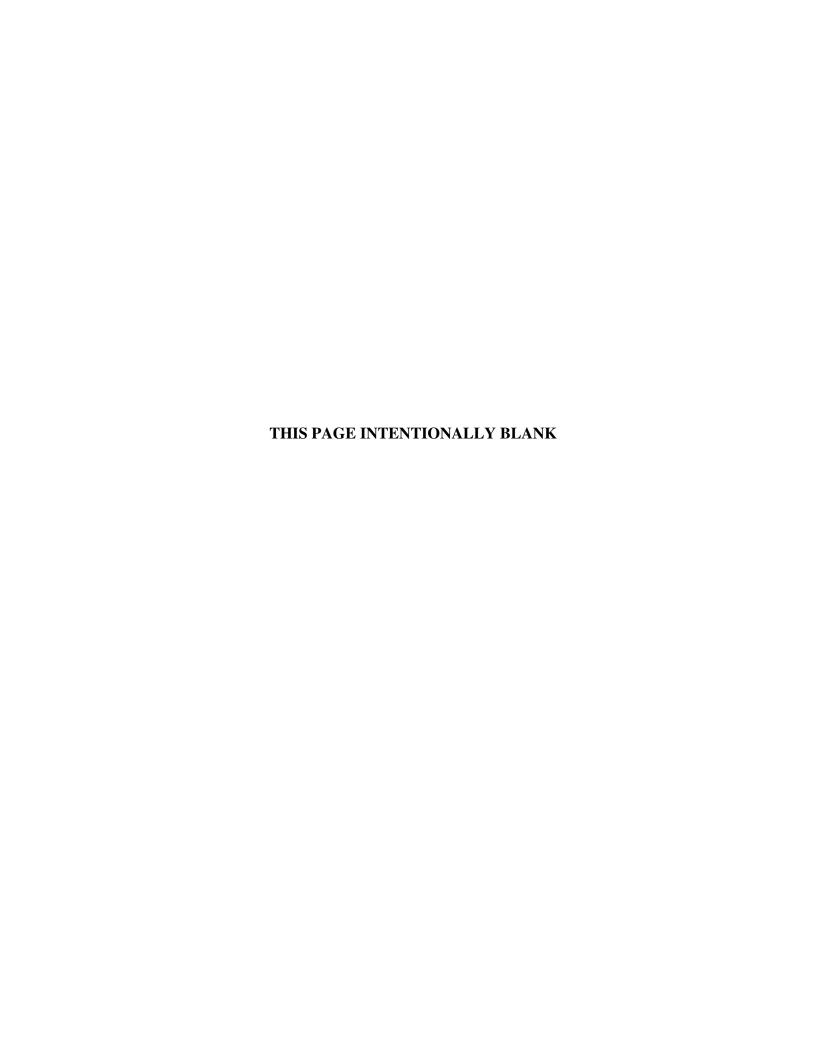
CHEMISTRY 12 PROVINCIAL EXAMINATION

1.	This exam	ination consists of two parts:	Value	Suggested Time
	PART A:	48 multiple-choice questions	60	70
	PART B:	13 written-response questions	40	50
		Tota	l: 100 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 Brønsted-Lowry Acids and Bases
 - Acid-Base Indicators
 - Standard Reduction Potentials of Half-cells

No other reference materials or tables are allowed.

3. 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 and may also include graphing 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 must not have the ability to either transmit or receive electronic signals. In addition to an approved calculator, students will be allowed to use rulers, compasses, and protractors during the examination.



PART A: MULTIPLE CHOICE

Value: 60 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.

Selected multiple-choice questions are worth 2 marks.

1. Which of the following could be used to describe the rate of a reaction?

- A. $\frac{\text{change in time}}{\text{change in concentration}}$
- B. change in mass change in concentration
- C. $\frac{\text{change in concentration}}{\text{change in time}}$
- $D. \quad \frac{\text{change in concentration}}{\text{change in mass}}$
- 2. Which factor explains why coal dust is explosive?

(1 mark)

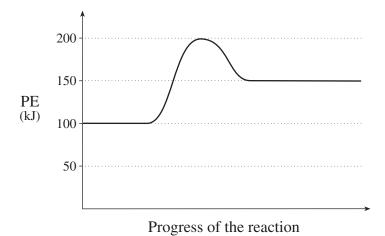
(1 mark)

- A. temperature
- B. surface area
- C. concentration
- D. addition of catalyst
- 3. As an activated complex changes into products, what changes occur in the chemical bonds of the activated complex and the PE of the system?

	Bonds	PE
A.	form	increases
B.	form	decreases
C.	break	increases
D.	break	decreases

4. Consider the following PE diagram:

(1 mark)

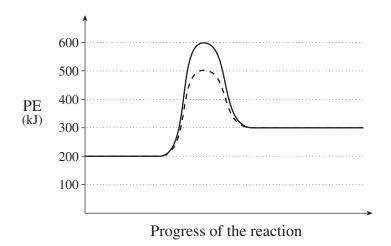


Which of the following describes ΔH and the type of reaction in the forward direction?

	ΔH (kJ)	Type of Reaction
A.	-50	exothermic
B.	-50	endothermic
C.	+50	exothermic
D.	+50	endothermic

5. Consider the following PE diagram for a catalyzed and uncatalyzed reaction:

(2 marks)



Which of the following describes the forward reaction?

	Reaction	Activation Energy (kJ)	ΔH (kJ)
A.	catalyzed	300	-100
B.	uncatalyzed	300	-100
C.	catalyzed	400	+100
D.	uncatalyzed	400	+100

6. Consider the following reaction mechanism:

(1 mark)

Step 1	$NOCl \rightarrow NO + Cl$
Step 2	$NOCl + Cl \rightarrow NO + Cl_2$

Identify the reaction intermediate.

- A. Cl
- $B.\quad Cl_2$
- C. NO
- D. NOCl

7. Consider the following:

(1 mark)

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

Initially, NH₃ is added to an empty flask. How do the rates of the forward and reverse reactions change as the system proceeds to equilibrium?

	Forward Rate	Reverse Rate
A.	increases	increases
B.	increases	decreases
C.	decreases	increases
D.	decreases	decreases

8. Consider the following:

(2 marks)

$$H_{2(g)} + Br_{2(\ell)} \stackrel{?}{\rightleftharpoons} 2HBr_{(g)} + energy$$

What positions do minimum enthalpy and maximum entropy tend toward?

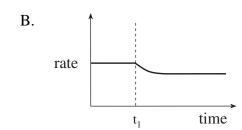
	Minimum Enthalpy	Maximum Entropy
A.	products	products
B.	products	reactants
C.	reactants	products
D.	reactants	reactants

Use the following equilibrium equation to answer questions 9 and 10.

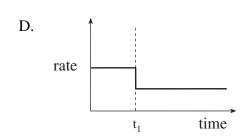
$$H_{2(g)} + I_{2(g)} \rightleftharpoons 2HI_{(g)} + energy$$

- 9. Which of the following stresses will **not** cause a shift in equilibrium? (1 mark)
 - A. decrease $[I_2]$
 - B. increase $[H_2]$
 - C. decrease volume
 - D. increase temperature
- 10. Which of the following shows the **reverse** rate of reaction when the volume is decreased at time = t_1 ? (1 mark)

A. rate t_1 time



 t_1 time



11. Consider the following:

(1 mark)

$$2SO_{3(g)} \quad \rightleftarrows \quad 2SO_{2(g)} + O_{2(g)}$$

Initially, some SO_3 is placed into a 3.0 L container. At equilibrium there is $0.030 \, \text{mol } SO_2$ present. What is the $\left[O_2\right]$ at equilibrium?

- A. 0.0050 mol/L
- B. 0.010 mol/L
- C. 0.015 mol/L
- D. 0.030 mol/L
- 12. Which reaction has the following equilibrium expression?

$$K_{eq} = \frac{[PCl_5]}{[PCl_3][Cl_2]}$$

- A. $PCl_{3(g)} + Cl_{2(g)} \rightleftharpoons PCl_{5(g)}$
- $\text{B.}\quad \text{PCl}_{3(g)} + \text{Cl}_{2(\ell)} \quad \rightleftarrows \quad \text{PCl}_{5(g)}$
- C. $PCl_{5(g)} \rightleftharpoons PCl_{3(g)} + Cl_{2(g)}$
- $\text{D.} \quad \operatorname{PCl}_{5(g)} \;\; \rightleftarrows \;\; \operatorname{PCl}_{3(g)} \, + \, \operatorname{Cl}_{2(\ell)}$
- 13. What will cause the value of K_{eq} for an exothermic reaction to decrease? (1 mark)
 - A. increasing the pressure
 - B. increasing the temperature
 - C. decreasing the temperature
 - D. decreasing the surface area

14. Consider the following equilibrium:

$$H_{2(g)} + I_{2(g)} \rightleftharpoons 2HI_{(g)}$$

An equilibrium mixture contains 1.0×10^{-3} mol H_2 , 2.0×10^{-3} mol I_2 and 1.0×10^{-2} mol HI in a $1.0\,L$ container. What is the value of K_{eq} ?

- A. 2.0×10^{-2}
- B. 5.0×10^{1}
- C. 5.0×10^3
- D. 1.0×10^4
- 15. Which of the following could be used to express solubility?

- A. mol
- B. M/s
- C. g/mL
- D. mL/min
- 16. When $100.0 \,\text{mL}$ of a saturated solution of BaF_2 is heated and all the water is evaporated, 3.6×10^{-4} mol of solute remains. The solubility of BaF_2 is (1 mark)
 - A. $1.9 \times 10^{-10} \,\mathrm{M}$
 - B. 1.3×10^{-5} M
 - C. $3.6 \times 10^{-4} \text{ M}$
 - D. $3.6 \times 10^{-3} \text{ M}$

17. A solution contains both $0.2\,\mathrm{M}~\mathrm{Mg}^{2+}_{(aq)}$ and $0.2\,\mathrm{M}~\mathrm{Sr}^{2+}_{(aq)}$. These ions can be removed separately through precipitation by adding equal volumes of $0.2\,\mathrm{M}$ solutions of

(1 mark)

- A. OH^- , and then S^{2-}
- B. Cl⁻, and then OH⁻
- C. CO_3^{2-} , and then SO_3^{2-}
- D. SO_4^{2-} , and then PO_4^{3-}
- 18. Consider the following equilibrium:

(1 mark)

$$CaSO_{4(s)} \rightleftharpoons Ca^{2+}_{(aq)} + SO_{4(aq)}^{2-}$$

Which of the following would shift the above equilibrium to the left?

- A. adding $CaSO_{4(s)}$
- B. adding $MgSO_{4(s)}$
- C. removing some Ca²⁺_(aq)
- D. removing some $SO_{4(aq)}^{2-}$
- 19. Calculate the solubility of CaC_2O_4 .

- A. $2.3 \times 10^{-9} \,\mathrm{M}$
- B. $1.2 \times 10^{-5} \text{ M}$
- C. $4.8 \times 10^{-5} \,\mathrm{M}$
- D. $8.3 \times 10^{-4} \,\mathrm{M}$

20. How many moles of dissolved solute are present in 100.0 mL of a saturated SrCO₃ solution?

(2 marks)

- A. $5.6 \times 10^{-11} \text{ mol}$
- B. 2.4×10^{-6} mol
- C. 2.4×10^{-5} mol
- D. 2.4×10^{-4} mol
- 21. What happens when equal volumes of 0.2 M AgNO₃ and 0.2 M NaCl are combined?

(2 marks)

- A. A precipitate forms because the trial ion product $> K_{sn}$
- B. A precipitate forms because the trial ion product $< K_{sp}$
- C. No precipitate forms because the trial ion product $> K_{sp}$
- D. No precipitate forms because the trial ion product $< K_{sp}$
- 22. Determine the maximum $[Na_2CO_3]$ that can exist in 1.0 L of 0.0010 M Ba $(NO_3)_2$ without forming a precipitate.

(1 mark)

- A. $2.6 \times 10^{-12} \text{ M}$
- B. 2.6×10^{-9} M
- C. 2.6×10^{-6} M
- D. 5.1×10^{-5} M
- 23. A Brønsted-Lowry acid is defined as a substance that

- A. releases H⁺_(aq)
- B. releases $OH^-_{(aq)}$
- C. accepts a proton
- D. donates a proton

- 24. Which of the following represents the reaction of $H_2PO_4^-$ acting as an acid? (1 mark)
 - A. $H_2PO_4^- + H_2O \rightleftharpoons H_3PO_4 + OH^-$
 - B. $H_2PO_4^- + H_2O \rightleftharpoons H_3O^+ + H_3PO_4$
 - C. $H_2PO_4^- + H_2O \rightleftharpoons H_3O^+ + HPO_4^{2-}$
 - D. $H_2PO_4^- + 2H_2O \rightleftharpoons H_4PO_4^+ + 2OH^-$
- 25. Consider the following equilibrium:

(1 mark)

$$HS^- + H_3BO_3 \rightleftharpoons H_2BO_3^- + H_2S$$

The two species acting as Brønsted-Lowry bases in the above equilibrium are

- A. HS^- and H_2S
- B. H₃BO₃ and H₂S
- C. HS⁻ and H₂BO₃⁻
- D. H₃BO₃ and H₂BO₃
- 26. List the bases $C_2O_4^{\ 2^-}$, NH_3 , and $PO_4^{\ 3^-}$ in order from strongest to weakest. (1 mark)
 - A. $PO_4^{3-} > NH_3 > C_2O_4^{2-}$
 - B. $C_2O_4^{2-} > NH_3 > PO_4^{3-}$
 - C. $NH_3 > PO_4^{3-} > C_2O_4^{2-}$
 - D. $PO_4^{3-} > C_2O_4^{2-} > NH_3$

27. A basic solution can be defined as one in which

(1 mark)

- A. $\left[H_3 O^+ \right]$ is not present
- B. $\left[H_3 O^+ \right]$ is equal to $\left[OH^- \right]$
- C. $\left[H_3 O^+ \right]$ is less than $\left[OH^- \right]$
- D. $\left[H_3O^+\right]$ is greater than $\left[OH^-\right]$
- 28. What is the $\left[H_3O^+\right]$ in 0.025 M HNO₃ ?

(1 mark)

- A. $4.0 \times 10^{-13} \,\mathrm{M}$
- B. 0.025 M
- C. 1.60 M
- D. 12.40 M
- 29. Write the base ionization constant expression for

$$NH_{3(aq)} + H_2O_{(\ell)} \rightleftharpoons NH_{4(aq)}^+ + OH_{(aq)}^-$$

A.
$$K_b = \frac{\left[NH_3\right]}{\left[NH_4^+\right]\left[OH^-\right]}$$

B.
$$K_b = \frac{\left[NH_4^+\right]\left[OH^-\right]}{\left[NH_3\right]}$$

C.
$$K_b = \frac{[NH_3][H_2O]}{[NH_4^+][OH^-]}$$

D.
$$K_b = \frac{[NH_4^+][OH^-]}{[NH_3][H_2O]}$$

- 30. The equation for the predominant hydrolysis of NH_4NO_3 can be represented by (1 mark)
 - A. $NH_4NO_{3(s)} \rightleftharpoons NH_4^+_{(aq)} + NO_3^-_{(aq)}$
 - $B. \quad NH_{4 \; (aq)}^{\;\; +} + H_2O_{(\ell)} \;\; \rightleftarrows \;\; H_3O_{\; (aq)}^{+} + NH_{3(aq)}$
 - C. $NO_{3(aq)}^- + H_2O_{(\ell)} \rightleftharpoons HNO_{3(aq)} + OH_{(aq)}^-$
 - D. $NH_4NO_{3(aq)} + H_2O_{(\ell)} \rightleftharpoons H_3O^+_{(aq)} + NH_3NO_{3(aq)}^-$
- 31. A solution made from baking soda (NaHCO₃) has an amphiprotic anion which is (2 marks)
 - A. basic since $K_a < K_b$
 - B. basic since $K_a > K_b$
 - C. acidic since $K_a < K_b$
 - D. acidic since $K_a > K_b$
- 32. A chemical indicator in solution consists of

- A. a weak acid and its conjugate acid.
- B. a weak acid and its conjugate base.
- C. a strong acid and its conjugate acid.
- D. a strong acid and its conjugate base.
- 33. A chemical indicator has a transition point at a pOH = 8.0. Calculate its K_a value and identify the indicator. (2 marks)
 - A. $K_a = 1 \times 10^{-8}$, phenol red
 - B. $K_a = 1 \times 10^{-6}$, methyl red
 - C. $K_a = 1 \times 10^{-8}$, thymol blue
 - D. $K_a = 1 \times 10^{-6}$, chlorophenol red

- 34. In acid-base titrations, the solution of known concentration is called a(n) (1 mark)
 - A. basic solution.
 - B. acidic solution.
 - C. standard solution.
 - D. indicating solution.
- 35. During a titration, $25.0 \,\text{mL}$ of $H_3 PO_{4(aq)}$ is completely neutralized by $42.6 \,\text{mL}$ of $0.20 \,\text{M}$ NaOH . Calculate the concentration of the acid. (2 marks)
 - A. 0.11M
 - B. 0.17 M
 - C. 0.34 M
 - D. 1.0 M
- 36. Write the net ionic equation for the neutralization of $HF_{(aq)}$ with $Sr(OH)_{2(aq)}$. (2 marks)
 - A. $HF_{(aq)} + OH_{(aq)}^{-} \to H_2O_{(\ell)} + F_{(aq)}^{-}$
 - B. $HF_{(aq)} + H_2O_{(\ell)} \rightarrow H_3O^+_{(aq)} + F^-_{(aq)}$
 - C. $2HF_{(aq)} + Sr(OH)_{2(aq)} \rightarrow SrF_{2(aq)} + 2H_2O_{(\ell)}$
 - D. $2H^{+}_{(aq)} + 2F^{-}_{(aq)} + Sr^{2+}_{(aq)} + 2OH^{-}_{(aq)} \rightarrow SrF_{2(aq)} + 2H_{2}O_{(\ell)}$
- 37. Consider the buffer equilibrium:

(1 mark)

What happens when a small amount of $HCl_{(aa)}$ is added to the equilibrium system?

- A. The pH increases slightly.
- B. The pH decreases slightly.
- C. The equilibrium shifts to the right.
- D. The equilibrium does not shift due to the levelling effect.

20	A 11 1	
38.	An oxidizing agent is	,

(2 marks)

- A. reduced as it loses electrons.
- B. reduced as it gains electrons.
- C. oxidized as it loses electrons.
- D. oxidized as it gains electrons.

39. Consider the following spontaneous reaction:

$$Ga^{3+} + 3Rb \rightarrow 3Rb^{+} + Ga$$

What happens in this reaction?

(1 mark)

- A. Rb is reduced.
- B. Rb gains electrons.
- C. Ga³⁺ loses electrons.
- D. Ga³⁺ acts as an oxidizing agent.

40. What is the oxidation number of S in $S_2O_6^{2-}$?

(1 mark)

- A. +3
- B. +5
- C. +6
- D. +7

41. Which of the following is the weakest oxidizing agent?

(1 mark)

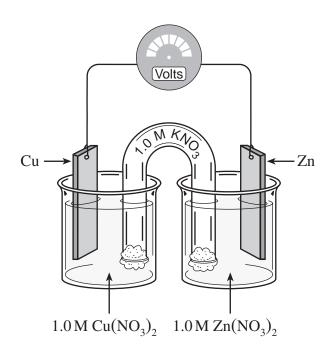
- A. Cl_2
- B. Al³⁺
- C. Sn²⁺
- D. acidified Cr₂O₇²⁻

42. Which of the following could react spontaneously with Ag metal?

- A. Cl
- B. Fe²⁺
- C. acidified SO_4^{2-}
- D. acidified NO₃

- 43. Which of the following could be titrated using acidified MnO_4^- ions? (1 mark)
 - A. Na⁺
 - B. IO_3^-
 - C. SO₄²⁻
 - $D. \quad H_2O_2$

Use the following diagram to answer questions 44 and 45.



44. What happens to the zinc electrode?

(2 marks)

- A. Mass increases as it is reduced.
- B. Mass decreases as it is reduced.
- C. Mass increases as it is oxidized.
- D. Mass decreases as it is oxidized.
- 45. Calculate the E° for the above cell.

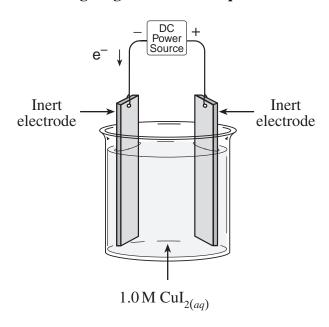
- A. -0.42 V
- B. 0.91 V
- C. 1.10 V
- D. 1.30 V

46. What happens to iron as it corrodes?

(2 marks)

- A. It loses electrons and is reduced.
- B. It gains electrons and is reduced.
- C. It loses electrons and is oxidized.
- D. It gains electrons and is oxidized.

Use the following diagram to answer questions 47 and 48.



47. What reaction occurs at the cathode?

(1 mark)

A.
$$2I^- \rightarrow I_2 + 2e^-$$

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

C.
$$H_2O \rightarrow \frac{1}{2}O_2 + 2H^+ + 2e^-$$

D.
$$2H_2O + 2e^- \rightarrow H_2 + 2OH^-$$

48. What happens to the $[I^-]$ in the operating cell?

(1 mark)

- A. $[I^-]$ increases overall.
- B. $[I^-]$ decreases overall.
- C. $[I^-]$ remains constant overall.
- D. $[I^-]$ decreases near the anode and increases near the cathode.

This is the end of the multiple-choice section.

Answer the remaining questions directly in this examination booklet.

PART B: WRITTEN RESPONSE

Valu	e: 40 marks	Suggested Tir	me: 50 minutes
INST	RUCTIONS:	You will be expected to communicate your knowledge and understa chemical principles in a clear and logical manner.	nding of
		Your steps and assumptions leading to a solution must be written in below the questions.	the spaces
		Answers must include units where appropriate and be given to the consignificant figures.	orrect number of
		For questions involving calculations, full marks will NOT be given providing only an answer.	en for
1.	Consider the r	eaction:	(3 marks)
		$2\mathrm{H}_2\mathrm{O}_{(\ell)} \to 2\mathrm{H}_{2(\mathrm{g})} + \mathrm{O}_{2(\mathrm{g})}$	
		oduction of $\rm O_2$ is $\rm 1.2\times10^{-2}~mol/s$. How many seconds will it take 100.0 g $\rm H_2O$?	
2.	Define the term	m catalyst.	(2 marks)
3.	Describe the n	ature of dynamic equilibrium.	(2 marks)

4. Consider the following:

$$N_2O_{4(g)} \quad \rightleftarrows \quad 2NO_{2(g)} \qquad \qquad K_{eq} = 9.5 \times 10^{-3}$$

Initially, 0.060 mol $\rm N_2O_4$ and 0.020 mol $\rm NO_2$ are placed into a 2.00 L container. Use calculations to determine the direction in which the reaction proceeds in order to reach equilibrium.

5. A 100.0 mL saturated solution of FeF_2 contains 0.0787 g of solute. Determine $\left[Fe^{2+}\right]$ and $\left[F^{-}\right]$ in the solution.

(3 marks)

6. Consider the following information and accompanying diagram:

In a titration experiment, ${\rm AgNO}_{3(aq)}$ was used to determine the $\left[{\rm Cl}^{-}\right]$ in a water sample and the following data were recorded:

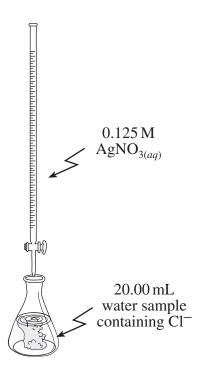
$$\left[\text{AgNO}_3 \right] = 0.125 \,\text{M}$$

Volume of water sample containing $Cl^- = 20.00 \, mL$ Initial buret reading of $AgNO_3 = 5.15 \, mL$ Final buret reading of $AgNO_3 = 37.15 \, mL$

The equation for this reaction is

$$Ag^{+}_{(aq)} + Cl^{-}_{(aq)} \rightarrow AgCl_{(s)}$$

Using the above data, determine the $\left[\text{Cl}^{-} \right]$ in the water sample.



(3 marks)

7. Consider the following equilibria:

I.	CH ₃ COOH + OCN [−] → HOCN + CH ₃ COO [−]
II.	CH ₃ COOH + ClO [−]

a) In equation \mathbf{I} above, the reactants are favoured. Identify the stronger acid.

(1 mark)

- b) In equation **II** above, the products are favoured. Identify the stronger acid. (1 mark)
- c) Consider the following reaction:

$$HOCN + CIO^- \rightleftharpoons OCN^- + HCIO$$

Does this reaction favour reactants or products? Explain.

(2 marks)

8. At 60°C, the pH = 6.51 for pure water. Determine the value of K_w at this temperature.

(3 marks)

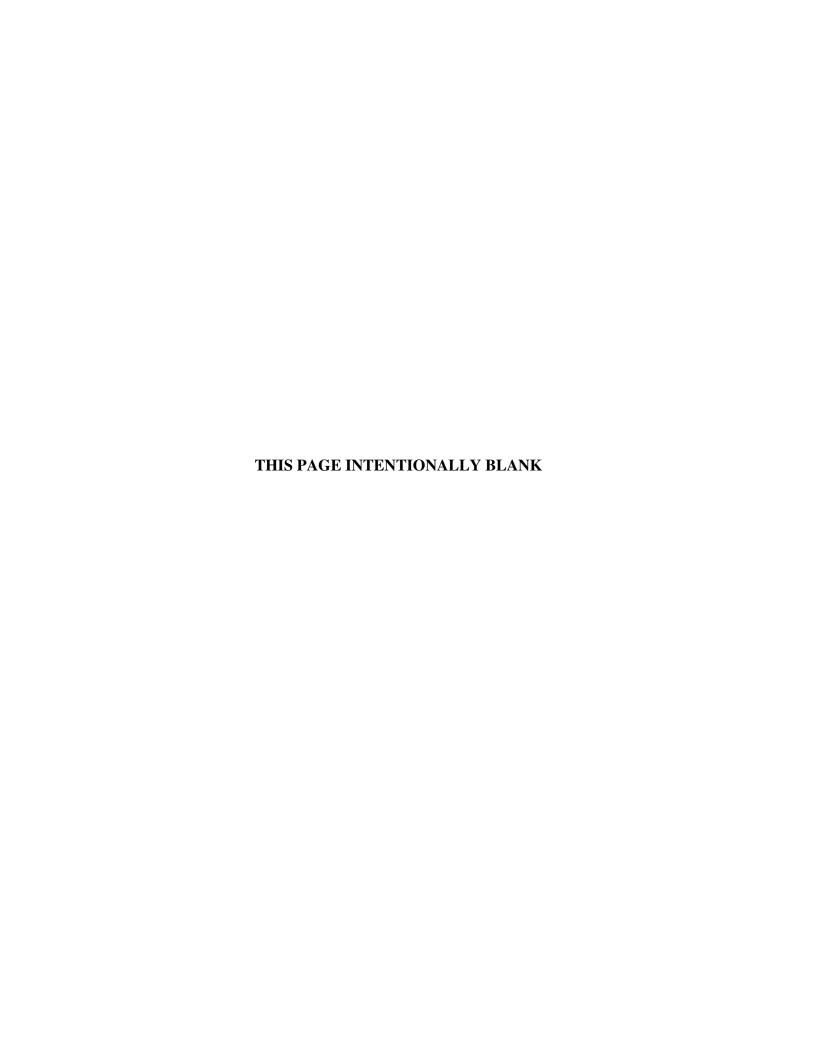
9. Calculate the pH of $0.35\,M\ H_2CO_3$.

(4 marks)

10.	A		
	a)	What causes the difference in these pH values?	(2 marks)
	b)	Select one indicator which could be used for both titrations.	(1 mark)
11.	Ва	alance the following redox equation:	(4 marks)
		$ClO_3^- + S_2O_3^{2-} \rightarrow S_4O_6^{2-} + Cl^-$ (acidic)	

12.	State two characteristics of the overall reaction in an electrochemical cell.	(2 marks)
	i)	
	ii)	
13.	Describe two chemically different methods of preventing the corrosion of iron. Explain how each method works.	(3 marks)
	Method 1:	
	Explanation:	
	Method 2:	
	Explanation:	

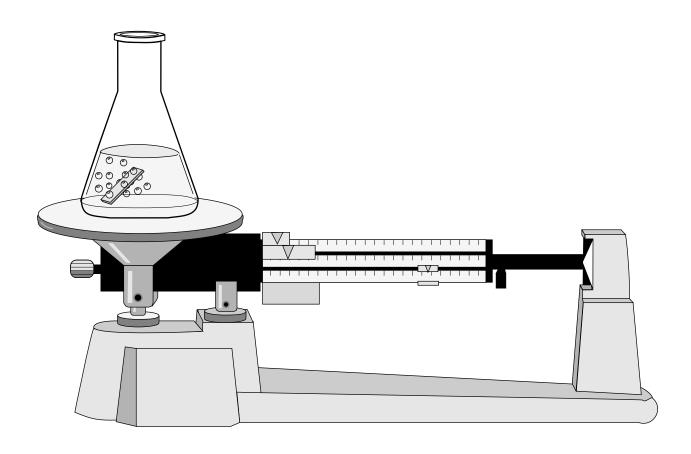
END OF EXAMINATION



Data Booklet

CHEMISTRY 12

Work done in this booklet will not be marked.



© Ministry of Education Revised January 2000

CONTENTS

Page	Table
1	Periodic Table of the Elements
2	Atomic Masses of the Elements
3	Names, Formulae, and Charges of Some Common Ions
4	Solubility of Common Compounds in Water
5	Solubility Product Constants at 25°C
6	Relative Strengths of Brønsted-Lowry Acids and Bases
7	Acid-base Indicators
8	Standard Reduction Potentials of Half-cells

7	,
č	_
-	_
•	_
Ĺ	T`
¥	$\overline{}$
5	-
L	T
-	
Ė	T.
Ę	T
۲	Т
ē	
Ļ	T
(
	_
Ę	T
Ĺ	_
7	~
-	
_	Ϥ
H	-
7	
١	_
7	$\overline{}$
۲	=
(
۲	_
¢	Y
	I,
ř	٦

	18		2	Ę	Helium	4.0	10	Se	Neon	20.2	18	Ā	Argon	39.9	36	궃	Krypton	83.8	54	Xe	Xenon	131.3	98	R	Radon	(222)					
	17						6	щ	Fluorine	19.0	17	ច	Chlorine	35.5	35	ğ	Bromine	79.9	53	_	lodine	126.9	85	Αŧ	Astatine	(210)					
	16						8	0	Oxygen	16.0	16	ဟ	Sulphur	32.1	34	Se	Selenium	79.0	52	<u>e</u>	Tellurium	127.6	84	S	Polonium	(508)					
	15				:		7	z	Nitrogen	14.0	15	_	Phosphorus	31.0	33	As	Arsenic	74.9	51	Sb	Antimony	121.8	83	區	Bismuth	209.0					
	14						9	ပ	Carbon	12.0	14	<u>ت</u>	Silicon	28.1	32	ge	Germanium	72.6	20	Sn	트	118.7	82	Ъ	Lead	207.2					
	13						2	Ф	Boron	10.8	13	₹	Aluminum	27.0	31	Ga	Gallium	2.69	49	드	Indium	114.8	81	F	Thallium	204.4					
CINI	12					L									30	Zn	Zinc	65.4	48	ဦ	Cadmium	112.4	80	Η̈́	Mercury	200.6					
JODIC LABLE OF THE ELEMENTS	11														29	చె	Copper	63.5	47	Ag	Silver	107.9	62	Αu	Gold	197.0					
I I I I I I	10														28	Z	Nickel	28.7	46	Pd	Palladium	106.4	78	芷	Platinum	195.1					
ADLE	6					Atomic Number	10		Atomic Mass						27	ပိ	Cobalt	58.9	45	Rh	Rhodium	102.9	77	<u>-</u>	Iridium	192.2	109	Ĭ	Meitnerium	(592)	
	8					— Atomi	—— Symbol	— Name	— Atomi						26	Ъ	lron	55.8	44	Ru	Ruthenium	101.1	92	Os	Osmium	190.2	108	Ŧ	Hassium	(592)	•
LEKI	7					41	i <u>s</u>	Silicon	28.1						25	M	Manganese	54.9	43	ဦ	Technetium	(86)	75	Re	Rhenium	186.2	107	뮵	Bohrium	(262)	
	9														24	ပ်	Chromium	52.0	42	Ø	Molybdenum	92.9	74	>	Tungsten	183.8	106	Sg	Seaborgium	(263)	
	2														23	>	Vanadium	6.03	41	S N	Niobium	92.9	73	Б	Tantalum	180.9	105	Ср	Dubnium	(262)	•
	4														22	F	Titanium	47.9	40	Zr	Zirconium	91.2	72	¥	Hafnium	178.5	104	¥	Rutherfordium	(261)	
	3														21	လွ	Scandium	45.0	39	>	Yttrium	88.9	22	ľ	Lanthanum	138.9	89	Ac	Actinium	(227)	
	2						4	Be	Beryllium	9.0	12	Mg	Magnesium	24.3	20	Sa	Calcinm	40.1	38	Š	Strontium	97.8	56	Ba	Barium	137.3	88	Ra	Radium	(226)	
	_	7	- I	Hydrogen	1.0		က	=	Lithium	6.9	11	Na	Sodium	23.0	19	¥	Potassium	39.1	37	Rb	Rubidium	85.5	55	ဌ	Cesium	132.9	87	上	Francium	(223)	
																															-

Based on mass of C^{12} at 12.00.

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

70 Yb Ytterbium 173.0	102 No Nobelium (259)
69 Tm Thulium 168.9	101 Md Mendelevium (258)
68	100
Er	Fm
Erbium	Fermium
167.3	(257)
67	99
Ho	Es
Holmium	Einsteinium
164.9	(252)
66	98
Dy	Cf
Dysprosium	Californium
162.5	(251)
65	97
Tb	BK
Terbium	Berkelium
158.9	(247)
64	96
Gd	Cm
Gadolinium	Curium
157.3	(247)
63 Eu Europium 152.0	95 Am Americium (243)
62 Sm Samarium 150.4	94 Pu Plutonium (244)
61	93
Pm	Np
Promethium	Neptunium
(145)	(237)
60	92
Nd	U
Neodymium	Uranium
144.2	238.0
59 Pr Praseodymium 140.9	91 Pa Protactinium 231.0
58	90
Ce	Th
Cerium	Thorium
140.1	232.0

103 **Lr** Lawrencium (262)

71 **Lu** Lutetium 175.0

ATOMIC MASSES OF THE ELEMENTS

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

Element	Symbol	Atomic Number	Atomic Mass
Actinium	Ac	89	(227)
Aluminum	Al	13	27.0
Americium	Am	95	(243)
Antimony	Sb	51	121.8
Argon	Ar	18	39.9
Arsenic	As	33	74.9
Astatine	At	85	(210)
Barium	Ba	56	137.3
Berkelium	Bk	97	(247)
Beryllium	Be	4	9.0
Bismuth	Bi	83	209.0
Boron	В	5	10.8
Bromine	Br	35	79.9
Cadmium	Cd	48	112.4
Calcium	Ca	20	40.1
Californium	Cf	98	(251)
Carbon	C	6	12.0
Cerium	Ce	58	140.1
Cesium	Cs	55	132.9
Chlorine	Cl	17	35.5
Chromium	Cr	24	
			52.0
Cobalt	Co	27	58.9
Copper	Cu	29	63.5
Curium	Cm	96	(247)
Dubnium	Db	105	(262)
Dysprosium	Dy	66	162.5
Einsteinium	Es	99	(252)
Erbium	Er	68	167.3
Europium	Eu	63	152.0
Fermium	Fm	100	(257)
Fluorine	F	9	19.0
Francium	Fr	87	(223)
Gadolinium	Gd	64	157.3
Gallium	Ga	31	69.7
Germanium	Ge	32	72.6
Gold	Au	79	197.0
Hafnium	Hf	72	178.5
Helium	He	2	4.0
Holmium	Но	67	164.9
Hydrogen	Н	1	1.0
Indium	In	49	114.8
Iodine	I	53	126.9
Iridium	Ir	77	192.2
Iron	Fe	26	55.8
Krypton	Kr	36	83.8
Lanthanum	La	57	138.9
Lawrencium	Lr	103	(262)
Lead	Pb	82	207.2
Lithium	Li	3	6.9
Lutetium	Lu	71	175.0
	Mg	12	24.3
MIABLICATION			
Magnesium Manganese	Mn	25	54.9

Element	Symbol	Atomic Number	Atomic Mass
Mercury	Hg	80	200.6
Molybdenum	Mo	42	95.9
Neodymium	Nd	60	144.2
Neon	Ne	10	20.2
Neptunium	Np	93	(237)
Nickel	Ni	28	58.7
Niobium	Nb	41	92.9
Nitrogen	N	7	14.0
Nobelium	No	102	(259)
Osmium	Os	76	190.2
Oxygen	O	8	16.0
Palladium	Pd	46	106.4
Phosphorus	P	15	31.0
Platinum	Pt	78	195.1
Plutonium	Pu	94	(244)
Polonium	Po	84	(209)
Potassium	K	19	39.1
Praseodymium	Pr	59	140.9
Promethium	Pm	61	(145)
Protactinium	Pa	91	231.0
Radium	Ra	88	(226)
Radon	Rn	86	(222)
Rhenium	Re	75	186.2
Rhodium	Rh	45	102.9
Rubidium	Rb	37	85.5
Ruthenium	Ru	44	101.1
Rutherfordium	Rf	104	(261)
Samarium	Sm	62	150.4
Scandium	Sc	21	45.0
Selenium	Se	34	79.0
Silicon	Si	14	28.1
Silver	Ag	47	107.9
Sodium	Na	11	23.0
Strontium	Sr	38	87.6
Sulphur	S	16	32.1
Tantalum	Ta	73	180.9
Technetium	Tc	43	(98)
Tellurium	Te	52	127.6
Terbium	Tb	65	158.9
Thallium	Tl	81	204.4
Thorium	Th	90	232.0
Thulium	Tm	69	168.9
Tin	Sn	50	118.7
Titanium	Ti	22	47.9
Tungsten	W	74	183.8
Uranium	U	92	238.0
Vanadium	V	23	50.9
Xenon	Xe	54	131.3
Ytterbium	Yb	70	173.0
Yttrium	Y	39	88.9
Zinc	Zn	30	65.4
Zirconium	Zr	40	91.2

NAMES, FORMULAE, AND CHARGES OF SOME COMMON IONS

- * Aqueous solutions are readily oxidized by air. ** Not stable in aqueous solutions.

Positive Ions

	(Cations)	
Al ³⁺	Aluminum	Pb ⁴⁺	Lead(IV), plumbic
NH_4^{+}	Ammonium	Li ⁺	Lithium
Ba^{2+}	Barium	Mg^{2+}	Magnesium
Ca ²⁺	Calcium	Mn^{2+}	Manganese(II), manganous
Cr^{2+}	Chromium(II), chromous	Mn^{4+}	Manganese(IV)
Cr ³⁺	Chromium(III), chromic	${\rm Hg_2}^{2+}$	Mercury(I)*, mercurous
Cu^+	Copper(I)*, cuprous	Hg^{2+}	Mercury(II), mercuric
Cu^{2+}	Copper(II), cupric	K^+	Potassium
H^+	Hydrogen	Ag^+	Silver
H_3O^+	Hydronium	Na^+	Sodium
$\mathrm{Fe^{2+}}$	Iron(II)*, ferrous	Sn^{2+}	Tin(II)*, stannous
Fe ³⁺	Iron(III), ferric	Sn^{4+}	Tin(IV), stannic
Pb^{2+}	Lead(II), plumbous	Zn^{2+}	Zinc

Negative Ions (Anions)

Br^-	Bromide	OH^-	Hydroxide
CO ₃ ²⁻	Carbonate	ClO ⁻	Hypochlorite
ClO ₃	Chlorate	I ⁻	Iodide
Cl	Chloride	$\mathrm{HPO_4}^{2-}$	Monohydrogen phosphate
${ m ClO_2}^-$	Chlorite	NO_3^-	Nitrate
$\text{CrO}_4^{\ 2-}$	Chromate	NO_2^-	Nitrite
CN-	Cyanide	$C_2O_4^{\ 2-}$	Oxalate
$\operatorname{Cr_2O_7}^{2-}$	Dichromate	O^{2-}	Oxide**
$\mathrm{H_2PO_4}^-$	Dihydrogen phosphate	ClO ₄	Perchlorate
CH ₃ COO	Ethanoate, acetate	$\mathrm{MnO_4}^-$	Permanganate
F^{-}	Fluoride	PO_4^{3-}	Phosphate
HCO_3^-	Hydrogen carbonate, bicarbonate	SO_4^{2-}	Sulphate
$HC_2O_4^{-}$	Hydrogen oxalate, binoxalate	S^{2-}	Sulphide
${ m HSO_4}^-$	Hydrogen sulphate, bisulphate	SO_3^{2-}	Sulphite
HS^-	Hydrogen sulphide, bisulphide	SCN^-	Thiocyanate
HSO ₃	Hydrogen sulphite, bisulphite		

SOLUBILITY OF COMMON COMPOUNDS IN WATER

The term soluble here means > 0.1 mol/L at 25°C.

	Negative Ions Anions)	Positive Ions (Cations)	Solubility Compour	
	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	
or	Chloride, Cl	All others	Soluble	
or	Bromide, Br Iodide, I	Ag^+ , Pb^{2+} , Cu^+		Low Solubility
	g 1 1 + go 2-	All others	Soluble	
	Sulphate, SO_4^{2-}	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⁺		
_	Sulplide, S	All others		Low Solubility
	Hudronido OH	Alkali ions, H ⁺ , NH ₄ ⁺ , Sr ²⁺	Soluble	
	Hydroxide, OH - }	All others		Low Solubility
or	Phosphate, PO ₄ ³⁻	Alkali ions, H ⁺ , NH ₄ ⁺	Soluble	
or	Carbonate, CO_3^{2-} 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	BaSO_4	1.1×10^{-10}
Calcium carbonate	CaCO ₃	5.0×10^{-9}
Calcium oxalate	CaC_2O_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	$\mathrm{Ag_2CrO_4}$	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 BRØNSTED-LOWRY ACIDS AND BASES

in aqueous solution at room temperature.

STRONG

STRENGTH OF ACID

Name of Acid	Acid		Base	K_a
Perchloric	HClO ₄	\rightarrow	H ⁺ + ClO ₄ ⁻	very large
Hydriodic	НІ	\rightarrow	H ⁺ + I ⁻	very large
Hydrobromic	HBr	\rightarrow	H ⁺ + Br ⁻	very large
Hydrochloric	HCl	\rightarrow	H ⁺ + Cl ⁻	very large
Nitric	HNO ₃	\rightarrow	H ⁺ + NO ₃	very large
Sulphuric	H ₂ SO ₄	\rightarrow	H ⁺ + HSO ₄ ⁻	very large
Hydronium Ion	H ₃ O ⁺	\rightleftharpoons	H ⁺ + H ₂ O	1.0
lodic	HIO ₃	$\stackrel{\textstyle \rightarrow}{\leftarrow}$	H ⁺ + IO ₃	1.7×10^{-1}
Oxalic	$H_2C_2O_4$	\rightleftharpoons	$H^+ + HC_2O_4^-$	5.9×10^{-2}
Sulphurous (SO ₂ + H ₂ O)				
Hydrogen sulphate ion	HSO ₄ -	\rightleftharpoons	H ⁺ + SO ₄ ²⁻	1.2×10^{-2}
Phosphoric	H ₃ PO ₄	\rightleftharpoons	$H^+ + H_2PO_4^-$	7.5×10^{-3}
Hexaaquoiron ion, iron(III) ion				
Citric	$H_3C_6H_5O_7$	\rightleftharpoons	$H^+ + H_2C_6H_5O_7^-$	7.1×10^{-4}
Nitrous				
Hydrofluoric				
Methanoic, formic	НСООН	\rightleftharpoons	H ⁺ + HCOO ⁻	1.8×10^{-4}
Hexaaquochromium ion, chromium(III) ion	$Cr(H_2O)_6^{3+}$	\rightleftharpoons	$H^+ + Cr(H_2O)_5(OH)^{2+}$	1.5×10^{-4}
Benzoic				
Hydrogen oxalate ion	HC ₂ O ₄ -	\rightleftharpoons	$H^+ + C_2O_4^{2-}$	6.4×10^{-5}
Ethanoic, acetic	CH ₃ COOH	\rightleftharpoons	H ⁺ + CH ₃ COO ⁻	1.8×10^{-5}
Dihydrogen citrate ion	$H_2C_6H_5O_7$	\rightleftharpoons	$H^+ + HC_6H_5O_7^{2-}$	1.7×10^{-5}
Hexaaquoaluminum ion, aluminum ion				
Carbonic (CO ₂ + H ₂ O)				
Monohydrogen citrate ion				
Hydrogen sulphite ion				
Hydrogen sulphide				
Dihydrogen phosphate ion				
Boric				
Ammonium ion				
Hydrocyanic				
Phenol				
Hydrogen carbonate ion	HCO ₃ -	\rightleftharpoons	$H^+ + CO_3^{2-}$	5.6×10^{-11}
Hydrogen peroxide				
Monohydrogen phosphate ion				
Water				
Hydroxide ion				
Ammonia				-

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.

