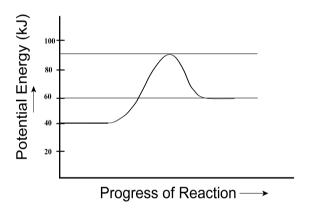
Part I Total Value: 50%

Instructions: Shade the letter of the correct answer on the computer scorable answer sheet provided.

- 1. Which observation can be explained using the kinetic molecular theory?
- \checkmark (A) A basketball is inflated with air.
 - (B) Butane C_4H_{10} burns faster than pentane C_5H_{12} .
 - (C) Powdered coal is more explosive than lumps of coal.
 - (D) Silver metal reacts slower than aqueous silver nitrate.
- 2. Which factor best explains why the combustion of candle wax, $C_{25}H_{52}$, is slower than the combustion of decane, $C_{10}H_{22}$?
 - (A) concentration
 - (B) nature of reactants
 - (C) pressure

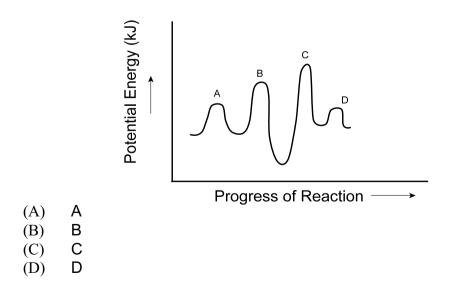
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- (D) temperature
- 3. For the diagram below, what is the activation energy for the reverse reaction?



	(A)	20 kJ
~	(B)	30 kJ
	(C)	40 kJ
	(D)	50 kJ

4. What is the rate determining step in the mechanism below?



5. Using the mechanism below, which species is the catalyst?

Step 1: $2 \operatorname{NO}(g) + \operatorname{O}_2(g) \rightarrow 2 \operatorname{NO}_2(g)$ Step 2: $2 \operatorname{NO}_2(g) \rightarrow 2 \operatorname{NO}(g) + 2 \operatorname{O}(g)$ Step 3: $2 \operatorname{O}(g) + 2 \operatorname{O}_2(g) \rightarrow 2 \operatorname{O}_3(g)$

 $\checkmark (A) NO(g)$ $(B) NO_2(g)$

$$(\mathbf{D}) \quad \mathbf{NO}_2(\mathbf{g})$$

(C)
$$O(g)$$

(D) $O_2(g)$

6. Given the table below, order the reaction trials from fastest to slowest.

Trial #	Concentration (mol/L)	Temperature (°C)
1	0.020	30
2	0.020	20
3	0.20	50
4	0.20	40

 $fastest \rightarrow slowest$

(A) $1 \rightarrow 2 \rightarrow 3 \rightarrow 4$ (B) $2 \rightarrow 1 \rightarrow 4 \rightarrow 3$ (C) $3 \rightarrow 4 \rightarrow 1 \rightarrow 2$ (D) $4 \rightarrow 3 \rightarrow 2 \rightarrow 1$

1

7. Which change will affect the value of K in the equilibrium system below?

 $2 \text{ NO}_2(g) \implies 2 \text{ NO}(g) + \text{ O}_2(g) \qquad \Delta H = 54 \text{ kJ}$

- \checkmark (A) decreasing temperature
 - (B) decreasing volume
 - (C) increasing [NO]
 - (D) increasing $[NO_2]$

8. Which change in the equilibrium below would result in the highest concentration of IF_3 ?

$$3 I_2(g) + 4 F_2(g) \implies 2 IF_3(g) + I_4F_2(g) + 34.2 kJ$$

- (A) decreasing $[I_2]$
 - (B) decreasing volume
 - (C) increasing $[I_4F_2]$
 - (D) increasing temperature

9. Which is true, for the equilibrium as written, if the system changes from blue to pink when placed in an ice bath?

 $\begin{array}{rcl} \operatorname{Co}(\operatorname{H}_2\operatorname{O})_6^{2+}(\operatorname{aq}) &+& 4\operatorname{Cl}^-(\operatorname{aq}) &\rightleftharpoons & \operatorname{Co}\operatorname{Cl}_4^{2-}(\operatorname{aq}) &+& 6\operatorname{H}_2\operatorname{O}(\ell) \\ \\ pink & & blue \end{array}$

	Reaction Type	ΔH
(A)	endothermic	negative
(B)	endothermic	positive
(C)	exothermic	negative
(D)	exothermic	positive

- 10. Which K value indicates the highest concentration of reactants?
- $(A) 1.9 \times 10^{-8}$ $(B) 1.3 \times 10^{-7}$ $(C) 2.6 \times 10^{7}$ $(D) 3.9 \times 10^{8}$

~

11. What is the equilibrium constant expression for the system below?

 $2 C(s) + 2 H_2O(g) \implies CH_4(g) + CO_2(g)$

$$\checkmark (A) \qquad \frac{[CH_{4}][CO_{2}]}{[H_{2}O]^{2}}$$

$$(B) \qquad \frac{[CH_{4}][CO_{2}]}{[C]^{2}[H_{2}O]^{2}}$$

$$(C) \qquad \frac{[H_{2}O]^{2}}{[CH_{4}][CO_{2}]}$$

$$(D) \qquad \frac{[C]^{2}[H_{2}O]^{2}}{[CH_{4}][CO_{2}]}$$

1

12. 0.100 mol of PCl₃, Cl₂, and PCl₅ are each introduced into the same 1.0 L evacuated flask. What is true of the concentration of each substance as the equilibrium below is established?

 $PCl_3(g) + Cl_2(g) \implies PCl_5(g) \qquad K_{eq} = 0.45$

l ₂] [PCl ₅]		
eases increases		
eases decreases		
eases increases		
eases decreases		

- 13. Which pH value would best describe a substance that tastes sour?
- ✓ (A) 6.8
 (B) 7.0
 (C) 7.2
 (D) 14.0

14. What is the conjugate base of $H_2PO_4^{-}(aq)$?

- ✓ (A) $HPO_4^{2-}(aq)$ (B) $H_3PO_4(aq)$ (C) $OH^{-}(aq)$ (D) $PO_4^{3-}(aq)$
- 15. Which substance is amphoteric?
 - (A) $CH_{3}COO^{-}(aq)$
 - (B) $CO_3^{2}(aq)$
- $\checkmark \quad (C) \quad HCO_3^{-}(aq)$

~

~

- (D) $SO_3^{2}(aq)$
- 16. Which is an acid-base conjugate pair?

$HCO_3^{-}(aq) + HNO_2(aq) \implies H_2CO_3(aq) + NO_2^{-}(aq)$

	Acid	Conjugate Base	
(A)	HCO_3^-	H_2CO_3	
(B)	HCO_3^{-}	NO_2^-	
(C)	HNO_2	H_2CO_3	
(D)	HNO_2	NO_2^-	

- 17. Which best describes a 10.0 mol/L solution of hydrofluoric acid?
 - (A) strong and concentrated
 - (B) strong and dilute
- \checkmark (C) weak and concentrated
 - (D) weak and dilute
- 18. The pH of a sample of rainwater has changed from 5.4 to 4.4. What is true of the rainwater?

	$[H_3O^+]$	Factor
(A)	decrease	1
(B)	decrease	10
(C)	increase	1
(D)	increase	10

- 19. What is the pH of a 0.00946 mol/L solution of barium hydroxide, Ba(OH)₂?
 - (A) 1.723
 - (B) 2.024
 - (C) 11.976
 - (D) 12.277

1

- 20. What is the pH of a solution made by diluting 10.0 mL of a 0.214 mol/L solution of nitric acid, HNO₃(aq), to a volume of 200.0 mL?
 - (A) 0.670
- **/** (B) 1.971
 - (C) 12.029
 - (D) 13.330
- 21. What is $[H_3O^+]$ of a solution with a pH of 2.00?
 - (A) 0.0010 mol/L
- ✔ (B) 0.010 mol/L
 - (C) 0.020 mol/L
 - (D) 2.0 mol/L
- 22. Which is representative of the weakest acid?
- ✓ (A) $K_a = 6.9 \times 10^{-8}$ (B) $K_a = 3.6 \times 10^{-6}$ (C) $K_a = 2.6 \times 10^{-4}$ (D) $K_a = 1.9 \times 10^{-2}$
- 23. Which acid would be expected to have the lowest K_a value?

	Volume (mL)	pН	Conductivity
(A)	50.2	3.25	poor
(B)	40.5	3.25	good
(C)	20.7	5.50	good
(D)	15.0	5.50	poor

- 24. What is the K_a of a weak acid, HA, if the pH at equilibrium for the acid is 2.20 and the equilibrium concentration of HA is 0.930 mol/L?
- $(A) \quad \begin{array}{l} 4.3 \times 10^{-5} \\ (B) \quad 6.8 \times 10^{-3} \\ (C) \quad 1.5 \times 10^{2} \\ (D) \quad 2.3 \times 10^{4} \end{array}$

~

- 25. The blood's natural buffering system $(H_2CO_3(aq)/HCO_3^-(aq))$ allows it to maintain a pH of 7.35. When a small amount of base enters the blood, with what species does it react?
 - (A) $H_3O^+(aq)$
 - (B) $HCO_3^{-}(aq)$
- $\checkmark \quad (C) \qquad H_2CO_3(aq)$
 - (D) $H_2O(\ell)$

26. A solution was tested with three indicators to give the results shown. What is the approximate pH of the solution?

Indicator	Final Colour
bromocresol green	blue
indigo carmine	blue
thymolphthalein	blue

- (A) 5.00
- 9.00 (B)
- 1 (C) 10.8 (D) 11.6
- 27. Which term describes the colour change when an acid-base indicator is used?
- ~ end point (A)
 - (B) equilibrium point
 - (C) equivalence point
 - (D) neutralization point

If the K_a expression for an acid is $K_a = \frac{\left[H_a O^*\right] \left[NO_a^-\right]}{\left[HNO_a\right]}$, what is the K_b expression for its 28.

conjugate base?

(A)
$$K_{n} = \frac{\left[HNO_{n}\right]}{\left[H_{n}O^{+}\right]\left[NO_{n}\right]}$$

(B)
$$K_{b} = \frac{\left[H_{3}O^{+}\right]\left[HNO_{2}\right]}{\left[NO_{2}\right]}$$

$$\checkmark \qquad (C) \qquad K_{b} = \frac{\left[OH^{-}\right]\left[HNO_{2}\right]}{\left[NO_{2}^{-}\right]}$$

(D)
$$K_{1_{0}} = \frac{\left[OH^{-}\right]\left[NO_{3}^{-}\right]}{\left[HNO_{3}\right]}$$

29.

What is a measure of the average kinetic energy of particles?

- (A) heat capacity
- **(B)** molar enthalpy
- (C) specific heat
- temperature 1 (D)

30. What is the molar enthalpy change for the reaction below?

Chemical	$\Delta H_{f}^{\circ}(kJ/mol)$	
MgCO ₃ (s)	-1113	
MgO(s)	-602	
$CO_2(g)$	-394	

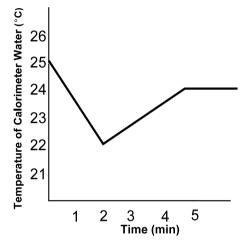
 $MgCO_3(s) \rightarrow MgO(s) + CO_2(g)$

-2109 kJ/mol (A)

(B)
$$-117 \text{ kJ/mol}$$

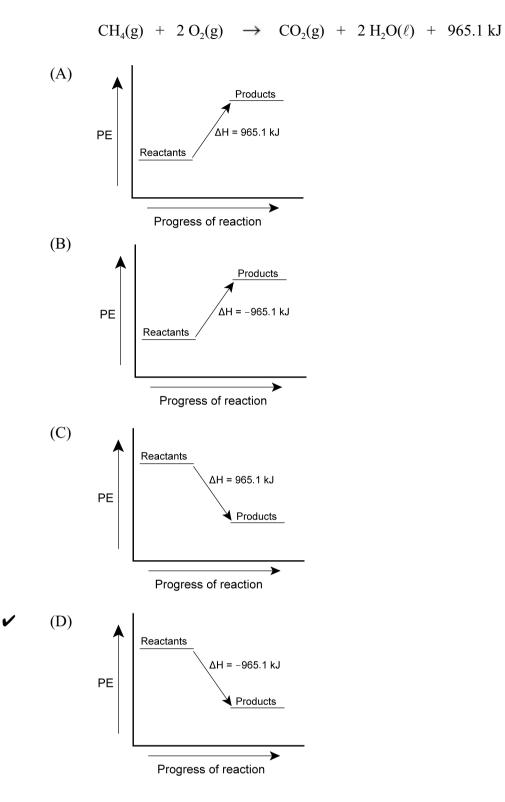
(C) 117 kJ/mol 1

- 2109 kJ/mol (D)
- A piece of ice was dropped into liquid water to determine the molar heat of fusion of 31. water. What is the best interpretation of the results as shown in the graph below?



- (A) The ice finishes melting at 2 minutes and heat enters the container.
 - The ice finishes melting at 2 minutes and heat leaves the container. (B)
 - The ice finishes melting at 5 minutes and heat enters the container. (C)
 - The ice finishes melting at 5 minutes and heat leaves the container. (D)
- 32. If it takes 588 J of energy to change the temperature of a 28.7 g strip of copper from 26.3 °C to 79.5 °C, what is the specific heat capacity of copper?
- 0.258 J/g·°C (A)
- 0.385 J/g·°C **(B)**
 - (C) 2.60 J/g·°C
 - $20.5 \text{ J/g} \cdot ^{\circ}\text{C}$ (D)
- 33. Which phase change is exothermic?
- 1 (A) gas to liquid
 - liquid to gas (B)
 - (C) solid to gas
 - solid to liquid (D)
- 34. Which energy change occurs when water is heated from 37.0 °C to 95.0 °C?
- (A) kinetic energy decreases 1
 - kinetic energy increases **(B)**
 - (C) potential energy decreases
 - potential energy increases (D)

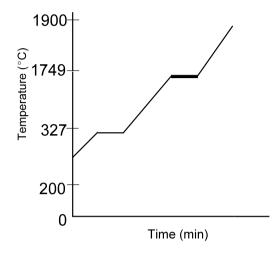
35. Which enthalpy diagram best represents the reaction below?



36. The molar heat of solution for NaOH(s) is -44.6 kJ/mol. What is true if a sample of NaOH(s) is dissolved in water in a calorimeter?

	Temperature of Water	Reaction	
(A)	decreases	absorbs energy	
(B)	decreases	releases energy	
(C)	increases	absorbs energy	
(D)	increases	releases energy	

- 37. Samples of two compounds X and Y have identical masses and initial temperatures. They are placed in an insulated container and a quantity of heat is added. What can be concluded if the final temperature of X is lower than that of Y?
 - (A) The molar mass of X is greater than that of Y.
 - (B) The molar mass of X is less than that of Y.
 - (C) The specific heat capacity of X is greater than that of Y.
 - (D) The specific heat capacity of X is less than that of Y.
- 38. The graph below shows the heating curve of a substance. What is its melting point?



- (A) 250 °C ✓ (B) 327 °C
 - (B) 327 C (C) 1749 °C
 - (D) 1749 C (D) 1900 °C
- A student walks to school in 0.50 hours travelling at 6 km/h. Before she leaves her house she eats 60.0 g of granola. How much energy will remain when she arrives at school? (FV of granola is 20.3 kJ/g; walking at 6 km/h burns 1675 kJ/h of energy.)
- ✓ (A) 380 kJ
 - (B) 460 kJ
 - (C) 840 kJ
 - (D) 1200 kJ
- 40. Given the reaction below, what is the molar enthalpy of formation for $NH_3(g)$?

 $N_2(g) + 3 H_2(g) \rightarrow 2 NH_3(g) + 92 kJ$

- (A) −92 kJ/mol (B) −46 kJ/mol
 - (C) 46 kJ/mol
 - (D) 92 kJ/mol

41. Which is a balanced oxidation half-reaction?

- (A) $\operatorname{Br}_2(\ell) + 2 e^- \rightarrow 2 \operatorname{Br}^-(\operatorname{aq})$
- $(B) \qquad 2 H^+ + 2 e^- \rightarrow H_2(g)$
- (C) $\operatorname{Sn}^{2+}(\operatorname{aq}) \longrightarrow \operatorname{Sn}(\operatorname{s}) + 2 \operatorname{e}^{-}$
- $(D) \quad \operatorname{Sn}^{2+}(\operatorname{aq}) \rightarrow \operatorname{Sn}^{4+}(\operatorname{aq}) + 2 e^{-1}$

42. Which species is the reducing agent in the reaction below?

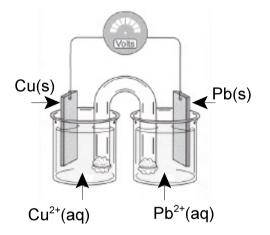
 $5 \text{ Fe}^{2+}(aq) + \text{MnO}_4^{-}(aq) + 8 \text{ H}^+(aq) \rightarrow 5 \text{ Fe}^{3+}(aq) + \text{Mn}^{2+}(aq) + 4 \text{ H}_2O(\ell)$

- ✓ (A) $Fe^{2+}(aq)$ (B) $Fe^{3+}(aq)$ (C) $Mn^{2+}(aq)$
 - (D) $MnO_4^{-}(aq)$

43. What is the balanced equation for the reaction below?

 $\mathrm{Co}^{^{3+}}\!(\mathrm{aq}) \ + \ \mathrm{Cd}(\mathrm{g}) \ \longrightarrow \ \mathrm{Co}^{^{2+}}\!(\mathrm{aq}) \ + \ \mathrm{Cd}^{^{2+}}\!(\mathrm{aq})$

- $(A) \quad Co^{3+}(aq) + Cd(g) \rightarrow Co^{2+}(aq) + Cd^{2+}(aq)$ $(B) \quad Co^{3+}(aq) + 2 Cd(g) \rightarrow Co^{2+}(aq) + 2 Cd^{2+}(aq)$ $(C) \quad 2 Co^{3+}(aq) + Cd(g) \rightarrow 2 Co^{2+}(aq) + Cd^{2+}(aq)$ $(D) \quad 2 Co^{3+}(aq) + 2 Cd(g) \rightarrow 2 Co^{2+}(aq) + 2 Cd^{2+}(aq)$
- 44. Which is the cathode in the diagram below?



 \checkmark (A) Cu(s)

(B)
$$Cu^{2+}(aq)$$

(D)
$$Pb^{2+}(aq)$$

45. What is the overall cell potential for the cell in which the reaction below occurs?

$$\operatorname{Au}^{3+}(\operatorname{aq}) + 3 \operatorname{NO}_2(g) + 3 \operatorname{H}_2O(\ell) \rightarrow \operatorname{Au}(s) + 3 \operatorname{NO}_3^{-}(\operatorname{aq}) + 6 \operatorname{H}^+(\operatorname{aq})$$

(A)	-0.90 V
(B)	-0.70 V
(C)	0.70 V

(D) 0.90 V

1

- 46. How many moles of electrons are transferred if a current of 1.80 A is allowed to run for 125 s?
- (A) $3.89 \times 10^{-5} \text{ mol}$ (B) $2.33 \times 10^{-3} \text{ mol}$ (C) $4.29 \times 10^{2} \text{ mol}$ (D) $1.39 \times 10^{3} \text{ mol}$

- 47. Which is the strongest reducing agent in the table below?
 - \checkmark indicates evidence of reaction
 - $\pmb{\mathsf{X}}$ indicates no evidence of reaction

	W(s)	X(s)	Q(s)	Z(s)
W -(s)	×	~	1	×
X ^{2–} (s)	×	×	1	×
Q ^{2–} (s)	×	×	×	×
$Z^{3-}(s)$	1	~	1	×

- (A) Q(s)
- (B) $Q^{-}(aq)$
- (C) Z(s)
- ✓ (D) $Z^{3-}(aq)$
- 48. Nickel (II) subsulfide, Ni_3S_2 , is mined at Voisey's Bay. What is the oxidation number of S in Ni_3S_2 ?
- ✓ (A) -3
 - (B) -2 (C) -1
 - (C) (D) 0
- 49. What type of cell is a rechargeable nickel-cadmium cell?
 - (A) fuel
 - (B) hydrogen
 - (C) primary
- ✓ (D) secondary
- 50. Which describes the energy conversion that occurs in an electrolytic cell?
 - (A) chemical to electrical
 - (B) chemical to mechanical
- \checkmark (C) electrical to chemical
 - (D) electrical to mechanical

Part II Total Value: 50%

Instructions: Complete all items in this section. Your responses should be clearly presented in a well-organized manner with proper use of units, formulae and significant digits where appropriate.

Value 2%

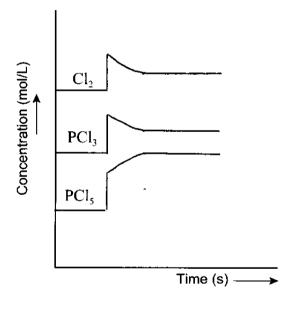
51.(a) For the reaction shown, explain how increasing the temperature would increase the rate of production of $O_2(g)$.

$$6 \operatorname{CO}_2(g) \ + \ 6 \operatorname{H}_2 O(\ell) \ \longrightarrow \ C_6 \operatorname{H}_{12} O_6(s) \ + \ 6 \operatorname{O}_2(g)$$

Increase in temperature causes reactant particles to move faster [1/2 mark]				
thus colliding more often $[\frac{1}{2}$ mark] and with more intensity $[\frac{1}{2}$ mark].				
More collisions will excess the E_a and have proper orientation $\ \left[\frac{1}{2} \mbox{ mark} \right]$,				
thus the rate increases.				

3% (b) For the equilibrium reaction shown, sketch the change in concentration for each reactant and product as the volume of the container is decreased and the system re-establishes equilibrium.

$$PCl_5(g) \rightleftharpoons Cl_2(g) + PCl_3(g)$$



[1.0 mark] for conc increase[1.0 mark] for the change[1.0 mark] for re-establish eqbm

3% 51.(c)

(c) Explain three changes that can be imposed on the system below that will allow the maximum amount of C_6H_6 to be produced.

$$3 C_2 H_2(g) \rightleftharpoons C_6 H_6(\ell) + 633 kJ$$

Any three of the following four:

1) increase $[C_2H_2]$ - system shifts right to use it up producing C_6H_6

- 2) remove C_6H_6 as it forms system shifts right to produce more of it
- 3) decrease temperature system shifts right to produce energy thus

producing C₆H₆

4) increase pressure/decrease volume - system shifts right to produce less gas molecules thus producing C₆H₆

[¹/₂ mark] x 3 for the change; [¹/₂ mark] x 3 for the explanation

(d) If 2.00 mol of HCl(g) is placed in an evacuated 3.00 L container and the equilibrium below is established, calculate the equilibrium concentration of all reactants and products.

2 HCl(aq) \implies H₂(g) + Cl₂(g) K = 3.4 × 10⁻²

 $c = \underline{n} = \frac{2.00 \text{ mol}}{3.00 \text{ L}} = 0.667 \text{ mol/L}$

I 0.667 n	ol/L 0	0	0
C - 2x	+x	+x	+ <i>x</i>
Е 0.667 -	2x x	x	x

$$\mathbf{K} = \underline{[\mathbf{H}_2] \ [\mathbf{Cl}_2]}_{[\mathbf{H}\mathbf{Cl}]^2}$$
[¹/₂ mark]

$$0.034 = \frac{x^2}{(0.667 - 2x)^2}$$
 [½ mark]

taking square root of both sides is much quicker than solving the quadratic since the assumption will not work

$0.18(4) = \frac{x}{0.667 - 2x}$	[½ mark]
0.12(3) - 0.36(8) x = x	
x = 0.089(9)	[½ mark]
$[H_2] = [Cl_2] = x = 0.090 \text{ mol/L}$	[½ mark]

$$[\text{HCl}] = 0.667 - 2x = 0.667 - (2 \times 0.089(9)) = 0.487 \text{ mol/L}$$
 [½ mark]

[1.0 mark] for science communication

5%

thus

4%

52.(a) Use an appropriate acid-base theory to explain why an aqueous solution of lithium hydrogen sulfate (LiHSO₄(aq)) turns blue litmus red.

Either:	
Modern Arrhenius: the HSO ₄ ⁻ reacts with wat	ter to form H ₃ O ⁺ ions
$HSO_4^- + H_2O \rightarrow H_3O^+ + SO_4^{-2-}$	
Or:	
Bronsted-Lowry: the HSO ₄ ⁻ donates a proton	
$HSO_4^- + H_2O \rightleftharpoons H_3O^+ + SO_4^{2-}$	
What is the pH of the resulting solution prepared by mixir 0.220 mol/L KOH(aq) solution with 30.00 mL of a 0.150	6
$HCl(aq) + KOH(aq) → H_2O(l) + KCl(aq)$ $H_3O^+(aq) + OH^-(aq) → 2 H_2O(l)$	(½ mark)
n _(HCl) = c v = (0.150 M) (0.03000 L) = 4.50 x 10^{-3} mol n _(KOH) = c v = (0.220 M) (0.02500 L) = 5.50 x 10^{-3} mol	(1.0 mark)
Ratio: HCl : KOH is 1 : 1	
Excess OH ⁻ : 4.50 x 10 ⁻³ mol of HCl reacts with 4.50 x leaving 0.00100 mol KOH	10 ⁻³ mol of KOH (½ mark)
$c_{(KOH)} = 0.00100 \text{ mol} = 0.0181(8) \text{ mol/L} = [OH^{-}]$ 0.0550 L	(1.0 mark)
$pOH = -\log [OH^{-}] = -\log 0.0181(8) = 1.74$ pH = 14.000 - pOH = 14.000 - 1.74 = 12.26	(½ mark) (½ mark)

4%

4% 52.(c) A 0.10 mol/L acid HA(aq) has a K_a value of 5.9×10^{-6} .

(i) What is the pH of the solution?

	HA	$+ H_2 O \rightleftharpoons$	H_3O^+	\mathbf{A}^{-}
Ι	0.10		0	0
С	- <i>x</i>		+x	+x
Ε	0.10 - <i>x</i>		+x	+x
check		0.10 > 500	assume good	(½ mark)
	K _a 5	5.9 x 10 ⁻⁶		

thus assume $0.10 - x \sim 0.10$ (½ mark)

$$K_{a} = [H_{3}O^{+}][A^{-}]$$
[HA] (½ mark)

$$5.9 \times 10^{-6} = \frac{x^2}{0.10}$$
 (½ mark)

$$x = 7.6(8) \times 10^{-4} = [H_3O^+]$$
 (½ mark)

$$pH = -\log [H_3O^+] = -\log 7.6(8) \times 10^{-4} = 3.11$$
 (½ mark)

(ii) Calculate the percent ionization of the acid.

% rxn =
$$[]_{change} x 100\% = \frac{7.6(8) x 10^{-4}}{0.10} x 100\% = 0.77\%$$

[1.0 mark]

(d) A pipette is used to transfer four 25.00 mL samples of hydrochloric acid, HCl(aq), to flasks. Each sample is then titrated to the equivalence point using a solution prepared by dissolving 0.200 g of sodium carbonate, Na₂CO₃, to form 100.0 mL of solution. The results below were obtained. What is the concentration of HCl(aq)?

Trial	1	2	3	4
Final burette reading (mL)	20.98	33.26	33.12	45.43
Initial burette reading (mL)	8.08	20.98	20.83	33.12
Volume of Na ₂ CO ₃ added (mL)	12.90	12.28	12.29	12.31

 $2 \text{ HCl}(aq) + \text{Na}_2\text{CO}_3(s) \rightarrow \text{H}_2\text{O}(\ell) + \text{CO}_2(g) + 2 \text{ NaCl}(aq)$ Avg Volume used = (12.28 + 12.29 + 12.31) mL / 3 = 12.29(3) mL (½ mark)

$$n (Na_{2}CO_{3}) = \frac{m}{M} = \frac{0.200 \text{ g}}{105.99 \text{ g/mol}} = 0.00188(7) \text{ mol} \qquad (1.0 \text{ mark})$$

$$c (Na_{2}CO_{3}) = \frac{n}{v} = \frac{0.00188(7) \text{ mol}}{0.1000 \text{ L}} = 0.0188(7) \text{ mol/L} \qquad (\frac{1}{2} \text{ mark})$$

$$n_{used} (Na_{2}CO_{3}) = c \text{ x } v = (0.0188(7) \text{ mol/L}) (0.01229(3) \text{ L}) = 2.32 \text{ x } 10^{-4} \text{ mol} \qquad (\frac{1}{2} \text{ mark})$$

$$n (HCl) = 2.32 \text{ x } 10^{-4} \text{ mol } Na_{2}CO_{3} \text{ x } \frac{2 \text{ mol } HCl}{1 \text{ mol } Na_{2}CO_{3}} = 4.63(9) \text{ x } 10^{-4} \text{ mol} \qquad (1.0 \text{ mark})$$

$$c (HCl) = \frac{n}{v} = \frac{4.63(9) \text{ x } 10^{-4} \text{ mol}}{0.02500 \text{ L}} = 0.0186 \text{ mol/L} \qquad (\frac{1}{2} \text{ mark})$$

4%	53.(a) A 2.71 g sample of mercury at 72.1 °C is cooled to -38.8 °C and solidified.
	(i) Draw a cooling curve for this process.

(i)	Draw a cooling cur	ve for this process.	
	melting point Hg	-38.8 °C	
	specific heat	0.140 J/g·°C	
	ΔH_{fusion}	8.9 kJ/mol	
Tempera °C	38.8 Heat Energy	released during this pro	1.0 mark]
$\mathbf{q}_1 = \mathbf{mc} \Delta \mathbf{T}$	$f = (2.71 \text{ g}) (0.140 \text{ J/g} \circ \text{C}) (-3)$	$38.8^{\circ}\text{C} - 72.1^{\circ}\text{C}) = -4$	2.0(8) J [1.0 mark]
$\mathbf{q}_2 = \mathbf{n} \Delta \mathbf{H}_{sol}$	$= \frac{2.71 \text{ g}}{200.59 \text{ g/mol}} \text{ x} - 8.9 \text{ kJ/}$	mol = - 0.12 kJ	[1.0 mark]
$\Delta E_{\rm T} = -0.0$	420(8) kJ + -0.12 kJ = -0).16 kJ or 1.6 x 10 ² J	[1.0 mark]
NaO	mL of 0.500 mol/L HCl(aq) i H(aq) in a coffee cup calorim °C and the highest recorded t HCl(aq) + NaOl	eter. The initial temper	rature of each solution is
(i)	Calculate the entha	lpy of neutralization of	THC1.
$q_{rxn} = -q_{aq}$ $= -(100)$	^{soln} .0 g) (4.184 J/g•°C) (23.2°C	nark] [½ mark]	0 _{water} = 1.00 g/mL
= -20(9) = -2.0(9)	/	[1.0	mark]
$\mathbf{n}_{\mathrm{HCl}} = \mathbf{n}_{\mathrm{Nac}}$	on ratio 1:1 thus all H	ICl completely reacts	
$\mathbf{n}_{\mathrm{HCl}} = \mathbf{c} \mathbf{x} \mathbf{v}$	= (0.500 mol/L) (0.0500L) =	= 0.0250 mol [½ n	nark]
$\Delta {f H}$ _{neutralized H}	$_{\rm IC1} = {\bf q} = {\bf -2.0(9) kJ} = {\bf 0.0250 mol}$	= - 84 kJ [½ n	nark]
(ii)	*	[1.0 mark] for scie ons that must be made halpy of neutralization	
Any	y 2 of the following 4:		[½ mark @]
	1) $q_{calorimeter} = 0$		
	$D_{aq soln} = D_{water}$		
	3) $c_{aq soln} = c_{water}$		
	4) q outside surroundings =	0 (system is isolated)	

4% 53.(c). Using the data below, calculate the enthalpy of formation for $Bi_2O_3(s)$.

$BiCl_3(s) + Bi_2O_3(s)$	\rightarrow	3 BiOCl(s)	$\Delta H = -147.7 \text{ kJ}$	
Bi(s) + $\frac{3}{2}$ Cl ₂ (g)	\rightarrow	BiCl ₃ (s)	$\Delta H_f = -379.1 \text{ kJ}$] x -1
$2 Bi(s) + \frac{3}{2} O_2(g)$	\rightarrow	$Bi_2O_3(s)$	$\Delta H_{\rm f} = ?$] x -1
$Bi(s) + \frac{1}{2} O_2(g) + \frac{1}{2} Cl_2(g)$	\rightarrow	BiOCl(s)	$\Delta H_f = -366.9 \text{ kJ}$] x 3
BiCl ₃ (s)	\rightarrow	Bi(s) + $\frac{3}{2}$ C	$\Delta H = +379.1 \text{ kJ}$ [1]	mark]
			•	-
$Bi_2O_3(s)$	\rightarrow	2 Bi(s) + $\frac{3}{2}$	$\Omega_{2}(g) \Delta H = ? x - 1$	1 mark]
$Bi_2O_3(s)$ 3 Bi(s) + 3/2 $O_2(g)$ + 3/2 $Cl_2(g)$		2 Bi(s) + $\frac{3}{2}$ 3 BiOCl(s)	$O_2(g) \Delta H = ? x - 1$ [$\Delta H = -1100.7 k$	-
	\rightarrow	3 BiOCl(s)	$O_2(g)$ $\Delta H = ? x - 1$ [$\Delta H = -1100.7 k$	J mark]
3 Bi(s) + $3/2$ O ₂ (g) + $3/2$ Cl ₂ (g)	\rightarrow	3 BiOCl(s) 3 BiOCl(s)	$O_2(g)$ $\Delta H = ? x -1$ [$\Delta H = -1100.7 k$ [1 $\Delta H_{rxn} = -147.7 H$	J mark]
$3 \operatorname{Bi}(s) + 3/2 \operatorname{O}_2(g) + 3/2 \operatorname{Cl}_2(g)$ $BiCl_3(s) + Bi_2\operatorname{O}_3(s)$ Thus	→ → 100.7 k	3 BiOCl(s) 3 BiOCl(s) xJ) = - 147.7 kJ	$O_2(g)$ $\Delta H = ? x -1$ [$\Delta H = -1100.7 k$ [1] $\Delta H_{rxn} = -147.7 l$	J [mark]

$$\Delta H_{f} = -573.9 \text{ kJ} \qquad [1 \text{ mark}]$$

3% 54.(a) Under acidic conditions, balance the half-reaction below.

$$Ag^{+}(aq) + S_{2}O_{3}^{2^{-}}(aq) \rightarrow SO_{4}^{2^{-}}(aq) + Ag(s)$$

$$5 H_{2}O + S_{2}O_{3}^{2^{-}} \rightarrow 2 SO_{4}^{2^{-}} + 10 H^{+} + 8 e^{-}$$

$$Ag^{+} + e^{-} \rightarrow Ag$$

$$[1 mark]$$

$$I'_{2} mark]$$

$$[1'_{2} mark]$$

$$5 H_2O + S_2O_3^{2-} \rightarrow 2 SO_4^{2-} + 10 H^+ + 8 e^- 8 Ag^+ + 8 e^- \rightarrow 8 Ag$$
[½ mark]

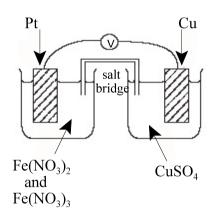
$$5 H_2O + S_2O_3^{2-} + 8 Ag^+ \rightarrow 2 SO_4^{2-} + 10 H^+ + 8 Ag$$
 [1 mark]

4% 54.(b) In an electrolytic cell, 0.061 g of Zn(s) was plated in 10.0 minutes from a solution of $ZnCl_2(aq)$. What current was used?

Plating at the cathode: $Zn^{2+} + 2e^- \rightarrow Zn$	[½ mark]
n (Zn) = \underline{m}_{M} = $\frac{0.061 \text{ g}}{65.38 \text{ g/mol}}$ = 9.3(3) x 10 ⁻⁴ mol	[½ mark]
n (e ⁻) = 9.3(3) x 10 ⁻⁴ mol Zn x $2 \mod e^{-}$ = 0.0018(7) mol 1 mol Zn	[½ mark]
$Q = n_{e} x F = (0.0018(7) mol) (96500 C/mol e^{-}) = 180 C$	[½ mark]
$I = Q_{t} = \frac{180 \text{ C}}{10.0 \text{ min } \text{x} \underline{60 \text{ s}}} = 0.30 \text{ A}$	[1.0 mark]

[1.0 mark] for science communication

3% (c) A student constructed an electrochemical cell as shown. The aqueous cell solutions had a concentration of 1 mol/L with respect to the metal ions present. The solution in one half-cell is initially an orange colour due to the mixture of the pale green Fe²⁺ ions and the orange Fe³⁺ ions. The other half is coloured blue due to the Cu²⁺ ions. Describe the colour changes the student would see in each half of the cell if the reaction proceeded until no further change took place.



 Fe^{3+} + $\mathrm{e}^{-} \rightarrow \mathrm{Fe}^{2+}$ $\mathrm{E}^{\circ} = +0.77 \mathrm{V}$

 $Cu^{2+} + 2e^- \rightarrow Cu E^\circ = +0.34 V$ this will become the oxidation at the anode

[1.0 mark]

Negative ions flow from cathode solution towards the anode solution.

Positive ions (blue Cu²⁺) flow from anode solution towards cathode solution.

[1.0 mark]

Orange solution will become orange-blue color as Cu²⁺ ions migrate in.

Blue solution will become a paler blue as Cu²⁺ ions leave

[1.0 mark]