EXPLORING CHEMISTRY

YEAR 12 - EXPERIMENTS, INVESTIGATIONS & PROBLEMS

Worked Solutions

The STAWA *Worked Solutions* have been developed through the collaboration of teachers working in Department of Education, Catholic Education WA and Association of Independent Schools of WA. Funding assistance was provided by the Department of Education.

The *Worked Solutions* are intended to support the Chemical understanding and problem solving section of the STAWA ATAR Exploring Chemistry Year 12: experiments, investigations and problems.

In an endeavour to provide the highest quality publication, the STAWA *Worked Solutions* were written and checked by different teachers. This does not guarantee that all answers are correct. Teachers are advised to work through disputed solutions with their students. If they are sure there is an error then they are asked to forward corrections to STAWA by email: admin@stawa.net

The STAWA *Worked Solutions* are a great example of teachers helping teachers for the benefit of all students.



Measurement in Chemistry

Set	Set 1: Significant figures and unit conversion									
1.	(a) (f)	3 4	(b) (g)	3 6	(c) (h)	3 3	(d)	3	(e)	3
2.	(a) (d)	$\begin{array}{c} 6.409\times10^3\\ 5.38\times10^1\end{array}$	(b) (e)	$\begin{array}{c} 3.2 \times 10^{\text{-2}} \\ 6.1 \times 10^{\text{-6}} \end{array}$	(c)	$8.91 imes 10^5$				
3.	(a) (b) (c)	$8 \times 10^{-4} (\times 10)$ $4.5 \times 10^{3} (\div 10)$ $9.0 \times 10^{-2} (\times 10)$	$(1^{3}) = 0.$ $(1^{3})^{-3} = 4$ $(1^{3})^{-3} = 9$	8 mm 5 kg 00 g	(d) (e) (f)	$7.03 \times 10^{5} (-0.05 \times 10^{4})$ $2.59 (\div 10^{9})$	$(+10^3) =$ $(\times 10^3) =$ (-2.59)	= 703 L = 5 × 10 ⁵ n 9 × 10 ⁻⁹ m	ηL	
4.	In a are 3. 0. 16. <u>+0.</u> 20.	ddition or subt contained in the 104 72 2 002 026 g = 20.0	to the t	n calculations surement wit	the ar h the lo place	nswer can hav east number o Answe	ve no n of digit er: 2.00	nore digits to the right 0×10^{1}	s to t ght c	he right of the decimal point than of the decimal point.
5.	20 k	$g = 20 x 10^3 g$: 1	Number of ato	oms = = = =	total mass \div (20 x 10 ³) \div (20 / 3.819) 5 × 10 ⁻²³ ato	- mass - (3.81) $\times (10^{-1})$	per atom 9×10^{-23}) 3^{+23})		

6.

	Pressure - mmHg	Pressure - atm	Pressure - Pa
(a)	760	1.00	1.01×10^{5}
(b)	750	$9.87 imes 10^{-1}$	9.97×10^{-1}
(c)	1.56×10^{3}	2.05	2.07×10^5
(d)	100	1×10^{-1}	1×10^{5}
(e)	4.83×10^{3}	$6.36 imes 10^{0}$	7.31×10^{3}

7. In addition or subtraction calculations the answer can have no more digits to the right of the decimal point than are contained in the measurement with the least number of digits to the right of the decimal point (see qn 4) 0.103 + 11.45 + 0.01 + 0.001 + 68.53 = 80.094 = 80.09 g (to 2nd decimal place) Answer: 8.009×10^{1}

Set 2: Errors

- 1. (a) first weighing = $\pm 2 \text{ mg} \div 0.002/12.363 \times 100 = 0.02\%$; final weighing $\pm 2 \text{ mg} \div 0.002/0.834 \times 100 = 0.2\%$ (b) Total error $\pm 4 \text{ mg} = \pm 4 \text{ mg} = 0.004/0.834 \times 100 = 0.5\%$
- 2. $153/36671 \times 100 = 0.417 \%$
- 3. (a) 1 part in 3000 so 3000 × 0.2 = 600 mg
 (b) Min Mass = (0.2 × 100) / 0.01 = 2000 mg or 2 g
- 4. (a) Smallest mass: 80.1 40.2 = 39.9 g (b) $(80.3 40.0) \pm (0.2 + 0.2) = 40.3 \pm 0.4$ g
- 5. $6.8245 \div 1.13 = 6.03938$ (the answer is expressed to the least number of significant figures, which is 3) = 6.04
- 6. $0.0229 \text{ or } 2.29 \times 10^{-2}$ (the answers is expressed to the least number of significant figures, which is 3)
- 7. (a) $1.67 \times 10^2 \,^{\circ}$ C or $167 \,^{\circ}$ C (b) $-2 \times 10^1 \,^{\circ}$ C or $-20 \,^{\circ}$ C
- 8. (a) $13.6 \times 96.485 = 1.31 \times 10^3 \text{ kJ mol}^{-1}$ (b) $24.6 \times 96.485 = 2.37 \times 10^3 \text{ kJ mol}^{-1}$ (c) $9.32 \times 96.485 = 8.99 \times 10^2 \text{ kJ mol}^{-1}$
- 9. $350 / 1920 \times 100 = 18\%$
- 10. $0.24 \text{ or } 2.4 \times 10^{-1}$ (the answers is expressed to the least number of significant figures, which is 2)



Set 3: Random and systematic and errors

- 1. Random Error: b and d Systematic Error: a and c
- 2. No one was right. The instrument can only be read to half a graduation. The bottom of the meniscus should be read so 3.6 or 3.7 mL volume would be acceptable readings.
- 3. Add percentage errors: (0.12/17)100 + (0.16/273)100 = 0.7655 = 0.77 %
- 4. density = m/v $v = m/density = 3.0 / 2.7 = 1.111 \text{ cm}^3$ $v = 51 \times 3 \times \text{thickness}$ thickness = 1.111 / (51 x 3) = 0.007262 = 7.26 x 10⁻³ cm
- 5. (a) Jenny (b) No only random errors can be eliminated by repeating an experiment (c) Obtain new glassware and solutions
- 6. (a) actual error in each titre is half the smallest scale division 0.5 mL or 0.0005 L
 (b) % error: (0.5/20.9)100 = 2.39%.
- 7. parallax error note the view of the circles around the burette
- 8. (a) Lyndon has random errors, Jenny has a close set of inaccurate results systematic error
 (b) Jenny has high precision and low accuracy, while Lyndon has low accuracy and low precision
 (c) Jenny needs to adjust the sights or obtain new equipment. Lyndon needs coaching
- 9. (a) (0.6/600)100 = 0.1%
 - (b) (0.6/250)100 = 0.24%
 - (c) (0.6/30)100 = 2%
- 10. (a) (0.05/10.5)100 = 0.48%
 - (b) (0.05/25.3)100 = 0.20%
 - (c) (0.05/37.2)100 = 0.13%
- 11. $M(NaHC_3) = 22.99 + 1.008 + 12.01 + 48 = 84.008$ $n(NaHC_3) = m/M = 2.445 / 84.008 = 0.0291$ mol $c(NaHCO_3) = n/V = 0.0291 / 0.2500 = 0.1164$ mol L⁻¹ % error (assume no error for mass) = (0.3 / 250.0) ×100 = ± 0.12 % $c(NaHCO_3) = 0.1164$ mol L⁻¹ ± 0.12 %

absolute error / 0.11642 = 0.12 %absolute error = $0.12/100 \times 0.11642 = 0.00014 \text{ mol } \text{L}^{-1}$ c(NaHCO₃) = $0.11642 \pm 0.00014 \text{ mol } \text{L}^{-1}$

- 12. (a) Average titre = 22.53 mL (23.15 is outlier)Error due to glassware = $(2 \times 0.05) \times 3 \text{ readings} = 0.3 \text{ mL}$ % error = $0.3/22.53 \times 100 = 1.33\%$
 - (b) Error due to range of titres = 22.60 22.45 = 0.15 mL % error = $0.15/22.53 \times 100 = 0.67\%$
 - (c) % uncertainty in conical flask will be due to pipette $0.06/20.00 \times 100 = 0.3\%$ $n(NaHCO3) = 0.0446 \times 0.02000 = 8.92 \times 10^{-4} \text{ mol}$ (d) $n(H+) = n(NaHCO_3) = 8.92 \times 10^{-4} \text{ mol}$
 - (d) $n(H^+) n(NaHCO_3)$ 8.92×10^{-1} mole $C(HCl) = 8.92 \times 10^{-4}/0.02253$ $= 3.96 \times 10^{-2}$ mol L⁻¹ % uncertainty (using glassware tolerance) = 1.33 + 0.3 = 1.63 % Absolute uncertainty $= 1.63/100 \times 3.96 \times 10^{-2}$ $= 6.45 \times 10^{-4}$



13.	(a)	average titre = $(5.44 + 5.60 + 5.55)/3$	= 5.53
		Absolute Uncertainty (glassware) = $(2 \times 0.10) \times 3$	= 0.60 mL
		% uncertainty = $0.6/5.53 \times 100 = 11\%$	
	(b)	average titre = $(24.50 + 24.45 + 24.30)/3$	= 24.42 mL

- (b) average titre = (24.30 + 24.45 + 24.30)/3 = 24.42 % uncertainty = $0.6/24.42 \times 100$ = 2.5 %
- (c) The smaller the average titre, the greater the effect of the tolerance of the glassware on the uncertainty. From the data above, it can be seen that diluting reduces the error significantly.

Chemical equilibrium

Set 4: Reaction rates and energy

- 1. (a) Perform the experiment on a balance and observe the rate of loss of mass
 - (b) Capture the hydrogen by downward delivery of water and read the level of gas at regular intervals
- 2. Ions in solution need no activation energy, have an infinite surface area while the sugar requires large activation energy and has relatively small surface area
- 3. (a) increased concentration more collisions higher rate of forward reaction
 - (b) increasing the concentration of the acid more collisions and a higher rate of reaction
- 4. (a) increasing pressure decreases volume means concentration increases so the rate will increase
- (b) no effect as reactants are solutions

6.

5. The catalyst provides a new pathway of lower activation energy. Offering a new pathway with lower activation energy means more particles can achieve activation and the rate will increase.



- (b) The heat of reaction of an endothermic reaction is positive $(+\Delta H)$, heat is absorbed from the surroundings so the reaction vessel fells cold, while the exothermic reaction releases energy to the surroundings, fells hot, expressed as a negative heat of reaction (- ΔH).
- (c) I Reaction A has the lower activation energy so you would expect it (given all other factors equal) to be the faster reaction
 - II The reverse of Reaction B would be faster as it has the lower activation energy, 70 kJ.



7. Increasing the temperature of a reaction increases its rate. More molecules or reacting particles have energy equal or greater than the activation energy as shown by the shaded section of the graph below. The higher temperature increases the number of collisions and the probability of a collision resulting in a reaction.



E(T) = average energy at temperature T T₁ < T₂ The proportion of molecules with energy equal to or greater than the activation energy, E_A (shaded areas) increases with an increase in temperature

Set 5: Equilibrium constant expressions

1.	$K = \underline{[H_2O]^2}$	8. K	$= [CO_2]$
	$[H_2]^2[O_2]$		
2.	$\mathbf{K} = \underline{[\mathbf{N}_2]^2[\mathbf{O}_2]}$	9. K	$= [CO_3^2][NH_4^+]$
	$[N_2O]^2$		$[HCO_3^-][NH_3]$
3.	$K = \underline{[CH_3OH]}$	10. K	= 1
	$[H_2]^2[CO_2]$		$[SO_3]$
4.	$K = [Ag^{+}]^{2}[CrO_{4}^{2}]$	11. K	$= [H_2O][CO_2]$
5.	$K = \underline{[HCO_3^-][H^+]}$	12. K	= <u>1</u>
	$[CO_2]$		$[C\ell_2]$
6.	$K = [SO_4^{2-}][H_3O^+]$	13. K	$= [Ca^{2+}][HCO_3^{-}]^2$
	$[HSO_4]$		$[CO_2]$
7.	$K = [CrO_4^{2}]^2 [H^+]^2$	14. K	$= \underline{[HC\ell]^2}$
	$[Cr_2O_7^{2-}]$		$[H_2O][CO_2]$

Set 6: Equilibrium systems

1.	(a) K	$= \underline{[NH_3]^2}$				
	(b)	(i) decrease	(ii) increase	(iii) increase	(iv) no change	
2.	(a) K	$= \underline{[H^+]^4[OC\ell^-]^2}$ $[C\ell_2]$				
	(b)	(i) increase	(ii) decrease	(iii) increase	(iv) decrease	
3.	(a)	(i) decrease	(ii) decrease	(iii) increase		
	(b)	(i) decrease	(ii) no change	(iii) increase		
4.	(a)	(i) increase	(ii) increase	(iii) decrease		
	(b)	(i) increase	(ii) decrease	(iii) decrease		
5.	(a)	(i) increase	(ii) decrease	(iii) no change the	n decrease	
	(b)	(i) no effect	(ii) same concentra	ation (pressure) but	more CO_2 is present	(iii) increase

Acids and bases

Set 8 Acids and bases

- 1. Conjugate acids: HF, HCO₃⁻, HCℓO₄, HSO₃⁻, NH⁴⁺
- 3. (a) $H_2SO_4(aq)$) + $H_2O(\ell) \rightarrow H_3O^+(aq) + SO_4^{2-}(aq)$ (b) $H_2S(aq)$) + $H_2O(\ell) \rightleftharpoons H_3O^+(aq) + HS^-(aq)$
- 2. Conjugate bases: Br⁻, SO₄⁻²⁻, HPO₄⁻²⁻, S²⁻, NO₃⁻, OH⁻
 - (c) KOH(aq) \rightarrow K⁺(aq) + OH⁻(aq) (d) N₂H4(aq) + H₂O(ℓ) \rightleftharpoons H⁺(aq) + HS⁻(aq)
- 4. A strong acid completely ionises in solution, while a weak acid only partially ionises in solution. Acid concentration refers to the amount of acid dissolved. A concentrated solution contains a high proportion of acid, while a dilute acid solution is mainly water with a very small proportion of acid.

(a) Concentrated solution of a strong acid (HX)



(c) Dilute solution of a strong acid (HX)



(b) Concentrated solution of a weak acid (HY)



(d) Dilute solution of a weak acid (HY)



I. conjugate acid-base pairs	II. Reaction favours the:
a. HF (aq) + H ₂ O (l) \Box F (aq) + H ₃ O ⁺ (aq)	
acid base conjugate base conjugate acid	forward reaction
b. $HSO_4(aq) + NH_3(aq) \square SO_4^{2-}(aq) + NH_4(aq)$	
acid base conjugate base conjugate acid	forward reaction
c. $H_2CO_3(aq) + F(aq) \square HCO_3(aq) + HF(aq)$	
acid base conjugate base conjugate acid	reverse reaction
d. $H_2PO_4^-(aq) + H_2O(l) \square H_3O^+(aq) + HPO_4^{2-}(aq)$	
acid base conjugate acid conjugate base	forward reaction

Set 9: Acid and base strength

5.

Concentration is a measure of the proportion of solute in a solution. Strength refers to the degree of ion formation that occurs when an acid or base is dissolved in water. For example, hydrogen chloride completely ionises in water and is classified as a strong acid: HCℓ(aq) → H⁺(aq) + Cl(aq) When the very soluble hydrogen fluoride dissolves in water there is little ionisation and it is classified as a weak acid: HF(aq) □ H⁺(aq) + F⁻(aq)



- (a) HCℓ, HNO₃, H₂SO₄, HBr, HI, HCℓO₄,
 (b) CH₃COOH, H₂S, HF, H₃PO₄, H₂C₂O₄, HSO₄⁻, or any organic acid.
 (c) NaOH, KOH, Ba(OH)₂, Ca(OH)₂, or any metal hydroxide or oxide.
 (d) NH₃, CO₃²⁻, CH₃NH₂ or any organic amine.
- (a) Concentrated and weak (b) Dilute and strong 3. (c) Concentrated and strong (d) Dilute and weak 4. (i) (a) HBr(g) \rightarrow H⁺(aq) + Br⁻(aq) (b) ionisation (c) HBr, OH^- , Br^- , H_3O^+ , H_2O (a) CH₃COOH(1) \rightleftharpoons H⁺(aq) + CH₃COO⁻(aq) (ii) (b) ionisation (c) OH^- , CH_3COO^- , H_3O^+ , CH_3COOH , H_2O (a) $H_2SO_4(\ell) \rightarrow H^+(aq) + HSO_4(aq)$ and $HSO_4(aq) \rightleftharpoons H^+(aq) + SO_4^{2-}(aq)$ (iii) (c) H_2SO_4 , OH^- , SO_4^{2-} , HSO_4^{-} , H_3O^+ , H_2O (b) ionisation (a) $NH_3(g) + H_2O(\ell) \rightleftharpoons NH_4^+(aq) + OH^-(aq)$ (b) ionisation (iv) (c) H_3O^+ or H^+ , $NH_4^{+,-}$ OH⁻, NH_3 , H_2O^- (v) (a) $Ba(OH)_2(aq) \rightarrow Ba^{2+}(aq) + 2OH^{-}(aq)$ (b) dissociation (c) $Ba(OH)_2$, H_3O^+ , Ba^{2+} , OH^- , H_2O
- 5. Stronger acid is $HC\ell O_4$ and the weaker base is $C\ell O_4$
- 6. (a) $H_2S(g) + H_2O(\ell) \rightleftharpoons H_3O^+(aq) + HS^-(aq)$ and $HS^-(g) + H_2O(\ell) \rightleftharpoons H_3O^+(aq) + S^{2-}(aq)$ (b) Strongest acid is H_3O^+ and the strongest base is S^{2-}
- 7. (a) $H_3PO_4(\ell) \rightleftharpoons H^+(aq) + H_2PO_4(aq) > H_2PO_4(aq) \rightleftharpoons H^+(aq) + HPO_4^{2-}(aq) > HPO_4^{2-}(aq) \rightleftharpoons H^+(aq) + PO_4^{3-}(aq)$ (b) $H_2O_1H_3PO_4, H_3O^+, H_2PO_4, HOO_4^{2-}, PO_4^{3-}, OH^{-}$
- 8. 99% sulfuric acid consists mostly of H_2SO_4 molecules, as there is little water to allow ionisation, so there are very few hydrogen ions to react with the iron. 2 mol L⁻¹ sulfuric acid is a strong acid; there is full ionisation for the release of one hydrogen ion represented by the equation: $H_2SO_4(\ell) \rightleftharpoons H^+(aq) + HSO_4^-(aq)$ There is therefore a high concentration of hydrogen ions which is the species that reacts with the iron in a reaction represented by the equation $2H^+(aq) + Fe(s) \rightarrow Fe^{2+}(aq) + H_2(g)$ Steel wool has a very high surface area and so there are more collisions and a higher rate of reaction.
- 9. For effective washing, large numbers of OH⁻ ions are required in the washing powder. To supply these substantial amounts of sodium hydroxide or sodium carbonate need to be present. The sodium hydroxide, being a strong base will produce a high concentration of OH⁻ ions which can be detrimental to fabrics. It is also dangerous to use, as concentrated OH⁻ solutions are corrosive to skin. Sodium carbonate, however, is a weak base that produces relatively low concentrations of OH⁻ ions but as these are consumed, more are produced until all of the carbonate ions are used up. The production of the OH⁻ ions from the carbonate ions is illustrated by the equation: $CO_3^{2-}(aq) + H_2O(\ell) \rightleftharpoons HCO_3^{--}(aq) + OH^{-}(aq)$
- 10. For this use the hydrogen ion concentration produced in the solution must be very low but there must be a process that replaces the hydrogen ions that are used up. It must therefore be a weak acid. As hydrochloric acid is a strong acid the concentration of hydrogen ions, even in dilute solutions, is relatively high and would damage human skin particularly in sensitive areas such as eyes and areas where fungal infections are likely to occur.

Set 10 : Hydrolysis

- 1. (a) neutral (b) basic (c) basic (d) acidic (e) basic
- 2. (b) $PO_4^{3-}(aq) + H_2O(\ell) \rightleftharpoons HPO_4^{2-}(aq) + OH^{-}(aq)$ (c). $HCO_3^{-}(aq) + H_2O(\ell) \rightleftharpoons H_2CO_3(aq) + OH^{-}(aq)$ (d_ $NH_4^{+}(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^{+}(aq)$ (e) $CH_3COO^{-}(aq) + H_2O(\ell) \rightleftharpoons CH_3COOH(aq) + OH^{-}(aq)$



- 3. (a) Ammonium nitrate, calcium hydrogenphosphate, potassium sulfate, and ammonium chloride.
 - (b) The salts contain either an anion of a weak acid or the cation of a weak base. (c) Ammonium nitrate (Ψ pH) $NH_4^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$
 - (c) Animomum intrate (\checkmark pH) $H_{4}^{(aq) + H_{2}O(\ell)} \rightleftharpoons H_{3}(aq) + H_{3}O(aq)$ calcium hydrogenphosphate (\Uparrow pH) $HPO_{4}^{2-}(aq) + H_{2}O(\ell) \rightleftharpoons H2PO4^{-}(aq) + OH^{-}(aq)$ potassium sulfate (\checkmark pH) $SO_{4}^{-2-}(aq) + H_{2}O(\ell) \rightleftharpoons HSO_{4}^{-}(aq) + OH^{-}(aq)$ ammonium chloride (\checkmark pH) $NH_{4}^{+}(aq) + H_{2}O(\ell) \rightleftharpoons NH_{3}(aq) + H_{3}O^{+}(aq)$ (d) These salts contain only ions derived from strong acids and strong bases so they do not react with water to produce the strong acid or strong base.

4.
$$Ca(H_2PO_4)_2(s) \rightleftharpoons 2H_2PO_4(aq) + Ca^{2+}(aq) \text{ and } H_2PO_4(aq) + H_2O(\ell) \rightleftharpoons HPO_4^{2-}(aq) + H_3O^{+}(aq)$$

- 5. (a) Yes, it becomes basic. (b) Na₂CO₃(aq) \rightarrow 2Na⁺(aq) + CO₃²⁻(aq) and CO₃²⁻(aq) + H₂O(ℓ) \rightleftharpoons HCO₃⁻ (aq) + OH⁻(aq)
- 6. (a) $OH^{-}(aq) + CH_{3}COOH(aq) \rightleftharpoons H_{2}O(\ell) + CH_{3}COO^{-}(aq)$ (b) basic, pH > 7. (c) $CH_{3}COO^{-}(aq) + H_{2}O(\ell) \rightleftharpoons CH_{3}COOH(aq) + OH^{-}(aq)$
- 7. Yes it becomes basic as F⁻ is the anion of a weak acid: $F^{-}(aq) + H_2O(\ell) \rightleftharpoons HF(aq) + OH^{-}(aq)$
- 8. The pH of a solution of ammonium ethanoate depends on the relative strength of the weak base ammonia and the weak acid ethanoic acid. The hydrolysis process is represented by the equations: $NH_4^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$ and $CH_3COO^-(aq) + H_2O(\ell) \rightleftharpoons CH_3COOH(aq) + OH^-(aq)$ Their ionisation constants are both close to 1.8 x 10⁻⁵ so the solution will be neutral.

Set 11: Water equilibrium

- 1. $H_2O(\ell) + H_2O(\ell) \rightleftharpoons OH-(aq) + H_3O^+(aq)$
- 2. (a) $[OH^{-}] = 1.00 \times 10^{-14} / [H^{+}] = 6.76 \text{ x } 10^{-8} \text{ mol } L^{-1}$ (b) Acidic
- 3. $[H^+] = 1.00 \times 10^{-14} / [OH^-] = 6.33 \times 10^{-9} \text{ mol } L^{-1}$
- 4. (a) $[H^+] = 1.00 \times 10^{-14} / [OH^-] = 6.45 \text{ x } 10^{-11} \text{ mol } \text{L}^{-1}$ (b) $[OH^-] = 2 \times [Ba(OH)_2]; [H^+] = 1.00 \times 10^{-14} / [OH^-] = 1.00 \times 10^{-14} / 7.80 \text{ x } 10^{-2} = 1.28 \text{ x } 10^{-13} \text{ mol } \text{L}^{-1}$
- 5. (a) $[OH^{-}] = 1.00 \times 10^{-14} / [H^{+}] = 9.62 \times 10^{-14} \text{ mol } L^{-1}$ (b) $H_2SO_4(\ell) \rightarrow H^+(aq) + HSO_4(aq) \text{ and } HSO_4(aq) \rightarrow 0.9HSO_4(aq) + 0.1H^+(aq) + 0.1SO_4^{-2}(aq)$ Overall $H_2SO_4(\ell) \rightarrow 0.9HSO_4(aq) + 1.1H^+(aq) + 0.1SO_4^{-2}(aq)$ $[H^{+}] = 1.1 \quad [H_2SO_4] = 1.1 \times 0.125 = 0.1375 \text{ mol } L^{-1} \quad [OH^{-}] = 7.27 \times 10^{-14} \text{ mol } L^{-1}$
- 6. (a) Ionisation of water is endothermic: heat + $H_2O(\ell) \rightleftharpoons H^+(aq) + OH(aq)$ a change in temperature will change the proportion of reactants and products and so change the equilibrium constant.
 - (b) K value decreases. A decreased in temperature causes the equilibrium position to move to restore some of the heat removed. H^+ and OH^- combine to produce water and heat, concentration become smaller.
 - (c) If temperature is increased a greater concentration of hydrogen ions will result.
 - (d) As the temperature is increased the rate of both the forward and reverse reactions will increase, but the endothermic forward reaction increases more than reverse reaction so hydrogen ion concentration increases.
- 7. (a) $10^{-14}/[H^+] = [OH^-] = 10^{-14}/1.55 \text{ x } 10^{-5} = 6.45 \text{ x } 10^{-10} \text{ mol } L^{-1} \text{ OH}^-$
 - (b) Some release of SO_2 from the smelter with elevated levels of CO_2 dissolving in the rainwater.
 - (c) $SO_2(aq) + H_2O(\ell) \rightleftharpoons HSO_3^-(aq) + H^+(aq)$ and $CO_2(aq) + H_2O(\ell) + HCO_3^-(aq) + H^+(aq)$
- 8. As volumes were equal, the average $[H^+] = (1.6 \times 10^{-4} + 1.3 \times 10^{-6})/2 = 8.065 \times 10^{-5} \text{ mol } L^{-1}$ $[H^+] = 8.07 \times 10^{-5} \text{ mol } L^{-1} \text{ As } 10^{-14}/[H^+] = [OH^-] = 10^{-14}/8.07 \times 10^{-5} = 1.24 \times 10^{-10} \text{ mol } L^{-1}$



9. (a) Stomach Acid : $n = c \ge v = 1.5 \ge 2.5 \ge 10^{-4} = 3.75 \ge 10^{-4} \mod H^+$ <u>Al_2O_3</u> : $n = m/M = 5 \ge 10^{-3}/(26.98 + 3(17.00)) = 6.412 \ge 10^{-5} \mod Al(OH)_3 \therefore$ there are 1.93 $\ge 10^{-4} \mod OH^$ available <u>Mg(OH)_2</u> : $n = m/M = 5 \ge 10^{-3}/(24.31 + 2(17.00)) = 8.575 \ge 10^{-5} \mod Mg(OH)_2 \therefore$ there are 1.715 $\ge 10^{-4} \mod OH^$ available Total OH⁻ = 3.645 $\ge 10^{-4} \mod OH^-$ available The acid is INXS by 3.75 $\ge 10^{-4} - 3.645 \ge 10^{-4} = 1.05 \ge 10^{-5} \mod c = n/v = 1.05 \ge 10^{-5}/1.5 = 7.00 \ge 10^{-6}$ [OH⁻] = 1.43 $\ge 10^{-9} \mod L^{-1}$

(b) Excess $A\ell(OH)_3$ and $Mg(OH)_2$ would remain undissolved (both insoluble in water) and the concentration of both hydrogen and hydroxide ions would therefore be 1.00 x 10⁻⁷ mol L⁻¹.

10. (a) $V(Mg(OH)_2) = 4.86 \text{ mL}$ (b) $[H^+] = 8.34 \text{ x } 10^{-3} \text{ mol } \text{L}^{-1}$ [OH⁻] = 1.20 x $10^{-12} \text{ mol } \text{L}^{-1}$

Set 12: Indicators and their use

- 1. (a) An organic acid or base
 - (b) The acid must be distinctly different in colour from its conjugate base:

e.g.: H(Litmus) (aq) \rightleftharpoons H⁺(aq) + (Litmus) (aq) Red Blue

- 2. (a) H(Indicator)(aq) \rightleftharpoons H⁺(aq) + (indicator)(aq)
 - (b) Addition of an acid increases [H⁺], causes more H(Indicator) to be produced and a reduction in the concentration of (Indicator)⁻. The colour changes from the (Indicator)⁻ colour to the H(Indicator) colour.
 - (c) Addition of a base reduces [H⁺] as H⁺ reacts with OH- to produce water. This causes the production of more (Indicator)⁻ and less H(Indicator). The colour changes from H(Indicator) colour to (Indicator)⁻ colour.
 - (d) Addition of Acid: increase in the rate of the reaction producing H(Indicator) with no immediate change to the reaction producing (Indicator)⁻ hence a colour change from (Indicator)" colour to H(Indicator) colour. Addition of Base: a reduction in [H⁺] causes a reduction in the rate of the reaction producing H(Indicator) with no immediate change in the rate of the reaction producing (Indicator)⁻ hence a colour change from H(Indicator)⁻ colour to (Indicator)⁻ colour.
- 3.

(b) $< pH=8.3$ Colourless) $< pH = 10.1$ (Yellow) $< pH = 11.4$ (Blue	
	2)
>pH = 10 (Pink) $>pH = 12.0 (Red)$ $>pH = 13.0 (Yell)$	ow)
(c) 8.3 to 10 10.1 to 12.0 11.4 to 13.0	

4.

(a)	Methyl red	Methyl orange	Bromophenol blue	Methyl violet
(b)	< pH = 4.4 (Red)	< pH = 3.1 (Red)	< pH = 3.0 (Yellow)	< pH = 0 (Yellow)
	>pH = 6.2 (Yellow)	>pH = 4.4 (Yellow)	> pH = 4.6 (blue)	>pH = 1.6 (Violet)
(c)	4.4 to 6.2	3.1 to 4.4	3.0 to 4.6	0 to 1.6

5. (a) H_2O , Na^+ , CH_3COO^- , OH^- , CH_3COOH , H^+



- (c) The solution is basic: $CH_3COO^{-}(aq) + H_2O(\ell) \rightleftharpoons CH_3COOH(aq) + OH^{-}(aq)$
- (d) Phenolphthalein. The end point (colour changes) must occur at the equivalence point. Equivalence point is basic use indicator that changes colour within pH 7 and 11. Phenolphthalein changes in pH range 8.3 to 10.



- (d) End point pH = 7: Indicator that changes colour in the range of pH = about 3 to 11.
- 8. (a) Phenolphthalein (b) More acid will be used

(c) Concentration measured lower than it actually is

Set 13: The pH scale

- 1. (a) $[H^+] = [HC\ell] = 0.100 \text{ mol } L^{-1}$ (b) $[H^+] = [HNO_3] = 0.00500 \text{ mol } L^{-1}$ [OH '] = 1.00 x 10⁻¹³ mol L⁻¹; pH = 1.00 [OH] = 2.00 x 10⁻¹² mol L⁻¹; pH = 2.30
 - (c) $[OH-] = [NaOH] = 0.0100 \text{ mol } L^{-1}$ $[H^+]$
 - (d) $[H^+] = [HC\ell] = 2.00 \text{ mol } L^{-1}$
- $\begin{bmatrix} OH ' \end{bmatrix} = 1.00 \text{ x } 10^{-13} \text{ mol } L^{-1}; \text{ pH} = 1.00 \\ \begin{bmatrix} OH \end{bmatrix} = 2.00 \text{ x } 10^{-12} \text{ mol } L^{-1}; \text{ pH} = 2.30 \\ \begin{bmatrix} H^+ \end{bmatrix} = 1.00 \text{ x } 10^{-12} \text{ mol } L^{-1}; \text{ pH} = 12.0 \\ \begin{bmatrix} OH^- \end{bmatrix} = 5.00 \text{ x } 10^{-15} \text{ mol } L^{-1}; \text{ pH} = 0.300 \\ \end{bmatrix}$
- (e) $[H^+] = [OH -] = 1.00 \times 10^{-14} \text{ mol } \text{L}^{-1}$ and the pH = 7.00. Neutral as solutions of Na⁺ ions and C ℓ^- ions do not hydrolyse in water.

2.	(a) $[H^+] = 1.00 \times 10^{-3} \text{ mol } L^{-1}$	$[OH^{-}] = 1.00 \times 10^{-11} \text{ mol } L^{-1}$
	(b) $[H^+] = 1.00 \times 10^{-11} \text{ mol } \text{L}^{-1}$	$[OH^{-}] = 1.00 \times 10^{-3} \text{ mol } \text{L}^{-1}$
	(c) $[H^+] = 10.0 \text{ mol } L^{-1}$	$[OH^{-}] = 1.00 \times 10^{-15} \text{ mol } \text{L}^{-1}$
	(d) $[H^+] = 2.75 \times 10^{-5} \text{ mol } L^{-1}$	$[OH^{-}] = 3.63 \times 10^{-10} \text{ mol } L^{-1}$
	(e) $[H^+] = 2.51 \times 10^{-8} \text{ mol } \text{L}^{-1}$	$[OH^{-}] = 3.98 \times 10^{-7} \text{ mol L}^{-1}$

- 3. Concentration changed by a factor of 1000
- 4. Depleted Solution: $pH = 2 \therefore [H^+] = 10^{-2}$ so $n = c \times v = 0.01 \times 2 = 0.02$ mol H^+ Added acid : $n = c \times v = 3.00 \times 3.00 = 9.00$ mol H^+ Total acid = 9.02 mol New $c = n/v = 9.02 / 5 = [H^+] = 1.80$ mol L^{-1}
- 5. $pH 5 = 10^{-5} \text{ mol } L^{-1} = 0.00001 \text{ mol } L^{-1} pH 3.6 = 10^{-3.6} \text{ mol } L^{-1} = 2.52 \times 10^{-4} \text{ mol } L^{-1}$ $c_1v_1 = c_2v_2 = 10^{-5} \times X = 0.025 \times 2.52 \times 10^{-4} = 630 \text{ mL}$ Water required = 630 - 25.0 = 605 mL
- 6. pH 12 \therefore pOH =2 so [OH⁻] = 0.01 mol L⁻¹ pH 11.7 \therefore pOH =2.3 so [OH⁻] = 10^{-2.3} = 0.0050 mol L⁻¹ \therefore he needs to add 0.01 - 0.0050 = 0.005 mol/L He has only 100mL so needs to add 0.0005 mol NaOH m(NaOH) to be added = nM = 0.0005 \times 39.998 = 0.00200 g = 20.0 mg



- (a) Drain : n = c × v = 0.236 × 0.200 = 0.0472 mol Runoff : n = c × v = 0.156 × 0.300 = 0.0468 mol New c = n/v = (0.0472 + 0.0468) / (0.2 + 0.3) = 0.188 mol L⁻¹ OH⁻ ∴ pOH = 0.726 ∴ pH = 14 - 0.726 = 13.3
 (b) V(HCℓ) = n(H⁺) / c = 0.094 / 1 = 0.094 L = 94 mL per 500 mL ∴ 188 mL per litre
 PH 5.5 ∴ [H⁺] = 10^{-5.5} = 3.162 × 10⁻⁶ mol L⁻¹
- n = c x v = $3.162 \times 10^{-6} \times 15000 = 4.743 \times 10^{-2} \text{ mol H}^+$ NaOH : $10g \therefore n=m/M = 10/39.99 = 0.25 \text{ mol of OH}^-$ Result : $0.2026 \text{ mol NaOH INXS} \therefore c=n/v = 0.2026/15000 = 1.351 \times 10^{-5} \text{ mol L}^{-1} \text{ OH}^ \therefore$ the pOH = $4.87 \therefore$ pH = 14 - 4.87 = 9.13
- 9. $pH 7.20 \therefore [H^+] = 10^{-7.2} = 1.584 \times 10^{-8} \text{ mol } L^{-1} \text{ H}^+$ $n = c \times v = 1.584 \times 10^{-8} \times 2.0 \times 10^{6} = 0.03168 \text{ mol } H^+$ $pH 7.8 \therefore [H^+] = 10^{-7.8} = 6.3095 \times 10^{-8} \text{ mol } L^{-1} \text{ H}^+$ $n = c \times v = 6.3095 \times 10^{-8} \times 2.0 \times 10^{6} = 0.12619 \text{ mol } H^+$ Needs to increase mol H^+ by 0.12619 - 0.03168 = 0.09451 mol $V(HC\ell) = n(H^+_{required}) / c = 0.09451/12.0 = 7.87 \text{ mL}$
- 10. (a) pH 6.75 : $[H^+] = 10^{-6.75} = 1.778 \times 10^{-7} \text{ mol } L^{-1} H^+$ pH 5.1: $[H^+] = 10^{-5.1} = 7.943 \times 10^{-6} \text{ mol } L^{-1} H^+$ Average = 4.0605 × 10⁻⁶ mol $L^{-1} H^+$ pH = $-\log_{10}[H^+] = -\log_{10}(4.059 \times 10^{-6}) = 5.39$

(b) pH 6.75 : $[H^+] = 10^{-6.75} = 1.778 \times 10^{-7} \text{ mol } L^{-1} H^+$ pH 8.00 : pOH = 6 : $[OH_-] = 10^{-6} = 1 \times 10^{-6} \text{ mol } L^{-1} OH^-$ Mix 1 L of each OH⁻ INXS by : 1 x 10⁻⁶ - 1.68 × 10⁻⁷ = 8.22 × 10⁻⁷ mol OH⁻ c =n/v = 8.22 × 10⁻⁷/2 = 4.11 × 10⁻⁷ mol L⁻¹ OH⁻ pOH = $-\log_{10}[OH^-] = -\log_{10}(4.11 \times 10^{-7}) = 6.39$: pH = 14-6.39 = 7.61

Set 14: Buffers

- 1. Buffer solutions resist a change in pH even with the addition of substantial amounts of hydrogen or hydroxide ions. Many specific reactions that occur in biological systems occur only at specific pH values. Some reactions produce or use up hydrogen ions in these solutions. Buffers prevent large changes in the pH of solutions such as blood, cell contents and lymph system allowing vital reactions to continue.
- 2. $NH_4^+(aq) + OH^-(aq) \rightarrow NH_3(aq) + H_2O(\ell)$ $NH_3(aq) + H^+(aq) \rightarrow NH_4^+(aq)$ Thus a buffer can accept or donate a proton to approximately maintain pH
- 3. Buffer solutions can be produced that have specific and known pH value. The fixed pH values are use as standards to calibrate the meters.
- (a) Hypochlorite ions added to pool water as sodium hypochlorite solution or calcium hypochlorite solid to kill micro organisms. Weak acid hypochlorous acid forms: OCℓ (aq) + H₂O(ℓ) ⇒HOCℓ(aq) + OH (aq) The buffer uses up some of these OH ions stopping the pool water becoming alkaline too quickly.
 - (b) $OH(aq) + HCO_3(aq) \rightleftharpoons H_2O(1) + CO_3^{2-}(aq)$ and $HCO_3(aq) \rightleftharpoons CO_3^{2-}(aq) + H^+(aq)$: H⁺ reacts with the OH⁻. Both processes use up OH⁻ ions.
- 5. (a) H_2O and HCO_3^- ion (b) $CO_2(aq) + H_2O(\ell) \rightleftharpoons HCO_3^-(aq) + H^+(aq)$ H^+ ion is used up by reacting with the HCO_3^- and OH^- ions is used up by reacting with CO_2 directly, $OH^-(aq) + CO_2(aq) \rightleftharpoons HCO_3^-(aq)$ or with H^+ ions, results in more H^+ ions. $H^+(aq) + OH^-(aq) \rightleftharpoons H_2O(\ell)$
- 6. Method 1: Add an equal volume of 1 mol L⁻¹ sodium ethanoate solution Method 2: Add an equal volume of a 0.5 mol L⁻¹ solution of sodium hydroxide. This would react with half the acetic acid to produce acetate ion as in the equation: $CH_3COOH(aq) + OH(aq) \rightarrow CH_3COO(aq) + H_2O(\ell)$

- 7. Method 1 Make a solution in water of the sodium citrate mixed with an equal number of moles of citric acid. Method 2 Make a solution in water of the sodium citrate add 1¹/₂ times number of moles of hydrochloric acid. This reacts with half the citrate ion: $C_6H_5O_7^{3-}(aq) + 3H^+(aq) \rightleftharpoons H_3C_6H_5O_7(aq)$
- 8. A change in temperature changes the equilibrium concentration of the species in equilibrium and as one of the species is always either H⁺ or OH⁻ ions, changing their concentrations changes the pH.

Set 15: Acid-base titrations 1

1.	(a)	$H^+(aq) + OH^-(aq) \rightarrow H_2O(\ell)$ $n(H^+) = cV = (2.00 \times 0.100) = 0.200 \text{ mol}$ $n(OH^-) = n(N_2OH) = n(HCl) = n(H^+)$					
	(b)	V(NaOH) = n/(CH) - n(HC) - n	V = 0.400 L				
	(c)	V(NaOH) = n/c = 0.225 / 0.500 = 0.450 L or 450 mL $H^{+}(aq) + OH^{-}(aq) \rightarrow H_2O(\ell)$ $n(H^{+}) = 2n(H_2SO_4) = cV = 2(0.250 \times 0.020) = 0.0100 \text{ mol}$ $n(OH^{-}) = n(NaOH) = n(H^{+})$	V = 0.450 L				
	(d)	V(NaOH) = n/c = 0.0100 / 0.500 = 0.0200 L or 20.0 mL $H_3PO_4(aq) + 3OH^-(aq) \rightarrow PO_4^{-3}(aq) + 3H_2O(1)$ $n(H^+) = 3n(H_3PO_4) = cV = 3(0.800 \times 0.0750) = 0.180 \text{ mol}$ $n(OH^-) = n(NaOH) = n(H^+)$	V = 0.0200 L				
		V(NaOH) = n/c = 0.180 / 0.500 = 0.360 L or 360 mL	V = 0.360 L				
2.	(a)	H ⁺ (aq) + OH ⁻ (aq) → H ₂ O(ℓ) n(OH ⁻) = n(NaOH) = cV = (0.600 × 0.200) = 0.120 mol n(OH ⁻) = n(H ⁺) n(H SO) = 1/ n(H ⁺) = 0.0600 mol					
		$V(H_2SO_4) = n/c = 0.0600 / 0.200 = 0.300 L \text{ or } 300 \text{ mL}$	V = 0.300 L				
	(b)	H ⁺ (aq) + OH ⁺ (aq) → H ₂ O(ℓ) n(H ₂ SO ₄) = n (Ba(OH) ₂) = cV = (0.100 × 0.0500) = 0.00500 mol V(H ₂ SO ₄) = n/c = 0.00500 / 0.200 = 0.0250 L or 25.0 mL	V = 0.0250 L				
3.	H ⁺ (ac n(H ⁺) n(OH c(KO 8.86 2	$\begin{aligned} q) + OH^{-}(aq) &\to H_2O(\ell) \\ p = cV = (0.105 \times 0.0211) = 0.00221 \text{ mol} \\ \Gamma) = n(KOH) = n(H^{+}) \\ PH) = n/V = 0.0221 / 0.0250 = 0.0886 \text{ mol} \text{ L}^{-1} \\ x \ 10^{-2} \text{ mol} \text{ L}^{-1} \end{aligned}$					
4.	$\begin{array}{c} H_2SO\\ n(H_2S)\\ c(H_2S)\end{array}$	D_4 (aq) + 2NaOH (aq) → Na ₂ SO ₄ (aq) + 2H ₂ O(ℓ) SO ₄) = ¹ / ₂ n(NaOH) = ¹ / ₂ (cV) = ¹ / ₂ (2.00 × 0.0222) = 0.0222 mol SO ₄) = n/V = 0.0222 / 0.00500 = 4.44 mol L ⁻¹					
5.	$\begin{split} &Na_2CO_3 \ (aq) + 2HNO_3 \ (aq) \to 2NaNO_3 \ (aq) + CO_2 \ (g) + 2H_2O(\ell) \\ &n(HNO_3) = cV = (1.00 \times 0.00350) = 0.00350 \ mol \\ &n(Na_2CO_3) = \frac{1}{2} \ n(HNO_3) = 0.00175 \ mol \\ &c(Na_2CO_3) = n/V = 0.00175 \ / \ 0.0250 = 0.0700 \ mol \ L^{-1} \\ &7.00 \ x \ 10^{-2} \ mol \ L^{-1} \end{split}$						
6.	Mg(C M(M) n(Mg n(HC V(HC V(HC	$\begin{array}{l} DH_{2} + 2HC\ell \rightarrow MgC\ell_{2} + 2H_{2}O \\ g(OH)_{2}) &= 24.31 + 2(16 + 1.008) = 58.326 \\ g(OH)_{2}) &= m/M = 0.450 / 58.326 = 0.00771 \text{ mol} \\ \ell \ \text{reacting} &= 2n(Mg(OH)_{2}) = 0.0153 \text{ mol} \\ \ell \ \text{reacting} &= n/c = 0.0154 / 0.150 = 0.103 \text{ L} \\ \ell \ \text{reacting} &= 0.103 \text{ L} \end{array}$					



- 7. $CaO + 2HC\ell \rightarrow CaC\ell_2 + H_2O$ $n(HC\ell) = cV = (1.50 \times 0.250) = 0.375 \text{ mol}$ $n(CaO) = \frac{1}{2} n(HC\ell) = \frac{1}{2} (0.375) = 0.1875 \text{ mol}$ $m(CaO) = nM = 0.1875 \times (40.08 + 16.00) = 10.5 \text{ g}$ m(CaO) = 10.5 g
- 8. $2NaHCO_3 + H_2SO_4 \rightarrow Na_2SO_4 + 2CO_2 + 2H_2O$ m = 600 g m? V? $M (NaHCO_3) = 22.99 + 1.008 + 12.01 + 48 = 84.008$ $M (H_2SO_4) = 2(1.008) + 32.07 + 4(16.00) = 98.086$ (a) $n(NaHCO_3) = m/M = 600 / 84.008 = 7.142 \text{ mol}$
 - (a) $n(NaFCO_3) = m/M = 6007 84.008 = 7.142 mor$ $n(H_2SO_4) = \frac{1}{2} n(NaHCO_3) = 3.571 mol$ $m(H_2SO_4) = nM = 3.571 \times 98.086 = 350 g$ $(b) V(H_2SO_4) = n/c = 3.571 / 12.0 = 0.297 L (297 mL)$
- 9. (a) i) 0.107 mol L^{-1} (ii) 3.89 g L^{-1}
- (b) 0.935 L (935 mL)

(c) 0.140 L (140 mL)

- 10. 7.34 x 10⁻² mol L⁻¹ 11. 25.2 mL of KOH
- 11. 25.2 mL of KO
- 12. 57.5%
- 13. 192 g mol⁻¹
- 14. 0.954 g of Na₂CO₃

Set 16 : Acid-bases titrations 2

- 1. $[Acid] = 1.38 \times 10^{-5} \text{ mol } \text{L}^{-1}$
- 3. $[HCO_3^-] = 6.32 \times 10^{-6} \text{ mol } \text{L}^{-1}$
- 5. [lactic acid] = $9.97 \times 10^{-6} \mod L^{-1}$
- 7. $[H_3C_6H_5O_7]_{ppm} = 31.4 \text{ ppm}$
- 8. (a) $NH_3(aq) \rightleftharpoons H^+(aq) + NH_4^+(aq)$
 - (b) pH = $-\log_{10}[H^+] = -\log_{10}(8.737 \times 10^{-3}) = 2.06$
 - (ii) methyl orange, methyl red or bromothymol blue.
 - (iii) As the equivalence point is acidic an indicator that changes colour at a pH between about 3 and 7 is required. Methyl orange (3.1 4.4), methyl red (4.4 6.2) and bromothymol blue (3.0 4.6) all change colour within this range.
- 9. [Fe]ppm = 20.1 ppm



4. $[OH-] = 1.59 \times 10^{-6} \text{ mol } \text{L}^{-1}$

6.
$$pH = -log_{10}[H^+] = -log_{10}(l \ 1.2) = -1.05$$



12

10.
$$\%$$
Pb = 90.4%

Set 17: Acids and bases in action: no answers provided

Oxidation and reduction

Set 18: Oxidation number

1.(a) (g) (m) (s) (y)	+4 -3 -2 +5 +2	(b) (h (n) (t) (z)	-2 +2 0 -2 +3	(c) (i) (o) (u)	+6 +5 +2 +4	(d) (j) (p) (v)	+2 +5 +4 +2	(e) (k) (q) (w)	+6 -4 +1 +1	(f) (l) (r) (x)	+5 +4 -3 +2
2.(a)	Mg(0+2)) O (0	-2)				(f)	Sn (+2 +4) O (0 -2)		
(b)	Na $(0 + 1)$) H (+1	1 0)				(g)	Br (-1 0)	S (+6 +4)		
(c)	I (-1 0) C	(0 - 1)					(h)	Fc (+2+3)) Cr (+6 +3)		
(d)	S (-2 0) N	(+5 +	2)				(i)	None (this	is an acid/b	ase read	ction)
(e)	S (-2 0) M	In (+7	+2)				(j)	N (-3 0) N	(+3 0)		
STAWA	12T001 Explo	oring Ch	emistry Year	12 worke	d solutions				രെ		
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- None (this is a precipitation reaction) (k)
- (1)None (this is an acid/carbonate equation)
- Set 19: Balancing half equations

1.	$Mg \rightarrow Mg^{2+} + 2e^{-}$	Oxidation	8.	$Au^+ + e^- \rightarrow Au$	Reduction
2.	$S + 2e^{-} \rightarrow S^{2-}$	Reduction	9.	$2H^+ + 2e^- \rightarrow H_2$	Reduction
3.	$2C1 \rightarrow Cl_2 + 2e^{-1}$	Oxidation	10.	$Cu^{2+} + 2e^{-} \rightarrow Cu$	Reduction
4.	$Ca \rightarrow Ca^{2+} + 2e^{-}$	Oxidation	11.	$AsO_3^{3-} + H_2O \rightarrow AsO_4^{3-} + 2H^+ + 2e^-$	Oxidation
5.	$I_2 + 2e^- \rightarrow 2I^-$	Reduction	12.	$S_2O_3^{2-} + 5H_2O \rightarrow 2SO_4^{2-} + 10H^+ + 4e^-$	Oxidation
6.	$Zn \rightarrow Zn^{2+} + 2e^{-}$	Oxidation	13.	$NO_3^- + 10H^+ + 8e^- \rightarrow NH_4^+ + 3H_2O$	Reduction
7.	$Cu^+ \rightarrow Cu^{2+} + e^-$	Oxidation	14.	$MnO_4 + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O$	Reduction

Set 20: Balancing redox equations

- $Br_2 + 2I \rightarrow 2Br + I_2$ 1.
- $Cu^{+} 2Ag^{+} \rightarrow Cu^{2+} + 2Ag$ $Mg + Pb^{2+} \rightarrow Mg^{2+} + Pb$ $Mg + 2H^{+} \rightarrow Mg^{2+} + H_{2}$ 2.
- 3.
- 4.
- $2\overline{A1} + 6\overline{H^+} \rightarrow 2\overline{A1^{3+}} + 3\overline{H_2}$ 5.
- $Mg + Cu^{2+} \rightarrow Mg^{2+} + Cu$ 6. 7.
- $2A1 + 3Zn^{2+} \rightarrow 2A1^{3+} + 3Zn$ $Cu + 2Fe^{3+} \rightarrow Cu^{2+} + 2Fe^{2+}$ 8.
- $Zn + 2Ag^+ \rightarrow Zn^{2+} + 2Ag$ 9.
- $I_2 + 2e^- \rightarrow 2I^-$ 19. (a) $S_2O_3^2 + 5H_2O + 4I_2 \rightarrow 2SO_4^2 + 10H^+ + 8I^-$ (c)
- $CH_3CH_2OH + H_2O \rightarrow CH_3COOH + 2H^+ + 2e^-$ 20. (a)
- $2CH_3CH_2OH+O_2 \rightarrow 2CH_3COOH$ (c)
- 21. (a) $C_6H_{12}O_6 + 6O_2 \rightarrow 6CO_2 + 6H_2O$

(b) $S_2O_3^{2-} + 5H_2O \rightarrow 2SO_4^{2-} + 8e^- + 10H^+$

(b) O_2 is reduced and C in the glucose is oxidised

(b) $O_2+2H_2O+8e^- \rightarrow 4OH^-$

(b) NO_2 is both

- O_2 is the oxidising agent, $C_6H_{12}O_6$ is the reducing agent (c) Ox: Mg \rightarrow Mg²⁺+2e⁻ Red: Ti⁴⁺+4e⁻ \rightarrow Ti RedOx: 2Mg + TiC $\ell_4 \rightarrow$ 2MgC ℓ_2 + Ti⁴⁺ 22. (a)
- TiC ℓ_4 is reduced. Mg metal is oxidised (b)
- 23. (a) $3NO_2 + H_2O \rightarrow 2HNO_3 + NO$

(c) A disproportionation reaction

Set 21: Galvanic cells



@••\$=

- (m) None (acid/base neutralisation equation)
- (n) None (this is a precipitation reaction)

 $2Li + 2H_2O \rightarrow 2Li^+ + 2OH^- + H_2$ 11. 12.

10. $C\ell_2 + 21^- \rightarrow 2C\ell^- + I_2$.

- $\begin{array}{c} Cu + 2NO_{3}^{-} + 4H^{+} \rightarrow Cu^{2+} + 2NO_{2} + 2H_{2}O \\ Cu + SO_{4}^{-2-} + 4H^{+} \rightarrow Cu^{2+} + SO_{2} + 2H_{2}O \end{array}$ 13.
- $2\mathrm{H}_2\mathrm{O}_2 \rightarrow 2\mathrm{H}_2\mathrm{O} + \mathrm{O}_2$ 14.
- $Cr_2O_7^{2-} + 8H^+ + 3C_2H_5OH \rightarrow 2Cr^{3+} + 3CH_3CHO + 7H_2O$ 15.
- $Mg + 2H_2O \rightarrow Mg^{2+} + H_2 + 2OH^2$ 16.
- $Cu_2O + 2H^+ \rightarrow Cu + Cu^{2+} + H_2O$ 17.
- $4Au + 16CN^{-} + 3O_2 + 12H^{+} \rightarrow 4[Au(CN)_4]^{-} + 6H_2O$ 18.

2.	(a) (b) (c) (d) (e)	$Cr + 3Ag^{+} \rightarrow Cr^{3+} + 3Ag$ $Mg + Cu^{2+} \rightarrow Mg^{2+} + Cu$ $Mg + 2Ag^{+} \rightarrow Mg^{2+} + 2Ag$ $Cr_{2}O_{7}^{2^{-}} + 14H^{+} + 6Fe^{2^{+}} \rightarrow 2Cr^{3^{+}} + 7H_{2}O + 6Fe^{3^{-}}$ $C\ell_{2} + 2I^{-} \rightarrow 2C\ell^{-} + I_{2}$	$E^{o} = +1.54 V$ $E^{o} = +2.70 V$ $E^{o} = +3.61 V$ $E^{o} = +0.59 V$ $E^{o} = +0.82 V$
3.	(a)	yes $E^{\circ} = +0.97$ (b) no $E^{\circ} = -0.94$	(c) no $E^{\circ} = -1.53$ (d) no $E^{\circ} = -0.34$
4.	(a) (b)	$\begin{array}{lll} Mg + 2H^{+} \rightarrow Mg^{2+} + H_{2} & E^{\circ} = + 2.36 \text{ V} \\ Sn + 2H^{+} \rightarrow Sn^{2+} + H_{2} & E^{\circ} = + 0.14 \text{ V} \\ Zn + 2H^{+} \rightarrow Zn^{2+} + H_{2} & E^{\circ} = + 0.76 \text{ V} \\ Sr + 2H^{+} \rightarrow Sr^{2+} + H_{2} & E^{\circ} = + 2.90 \text{ V} \\ Cu + 2H^{+} \rightarrow Cu^{2+} + H_{2} & E^{\circ} = - 0.34 \text{ V} \\ Ag + H^{+} \rightarrow Ag^{+} + H_{2} & E^{\circ} = - 0.80 \text{ V} \\ Only those with a positive E^{\circ} will be spontaneoon \\ The total E^{\circ} must be positive for the first but nega \\ (i) Sn or Ni & (ii) acidified H_{2}O_{2} \text{ on } \\ (iv) O_{2}/4H^{+}, Cr_{2}O_{7}^{-2}, C\ell_{2} & (v) \text{ Au, } C\ell^{-1} \end{array}$	us. Mg, Sn, Zn, Sr tive for the second. $f MnO_4^-$ (iii) Pb, Sn, Ni, Co
5.	(a) (b) (c)	$Fe^{2+} + Fe^{2+} \rightarrow Fe^{3+} + Fe$ $H_2O_2 + H_2O_2 \rightarrow 2 H_2O + O_2$ $C\ell_2 + C\ell_2 \rightarrow HOC\ell + HC\ell$	$E^{o} = -1.21V$ not spontaneous $E^{o} = +1.06 V$ spontaneous $E^{o} = -0.27 V$ not spontaneous
6.	(a) (c) (e) (g)	$C\ell_2 + 2Br^- \rightarrow 2C\ell^- + Br_2 + 0.28 V$ 2Aℓ+ 6H ⁺ → 2Aℓ ³⁺ + 3H ₂ +1.68 V No Reaction No Reaction	(b) No reaction (d) Fe + Sn ²⁺ \rightarrow Fe ²⁺ + Sn +0.30 V (f) Cr ₂ O ₇ ²⁻ +8H ⁺ + 3H ₂ S \rightarrow 2Cr ³⁺ + 7H ₂ O +3S +1.19 V (h) Both Br ⁻ and Fe ²⁺ are oxidized

Set 22: Electrolytic Cells

- 1. (a) See diagram
 - Cathode : $Cu^{2+}(aq) + 2e^{-} \rightarrow Cu(s)$ (b) Anode: $2I^{-}(aq) \rightarrow 2e^{-} + I_{2}(aq)$
 - (c) -0.54 + 0.34 = -0.20 V The cell would require at least 0.20 V to be supplied
 - Anode (d) The blue solution would fade, salmon pink deposit on electrode and brown solution formed.

	HBr	NiI ₂	KCℓ
Products	H ₂ and Br ₂	Ni and I ₂	$C\ell_2$ and K
Anode	$2Br(aq) \rightarrow 2e^{-} + Br_2(aq)$	$2\Gamma(aq) \rightarrow I_2(aq) + 2e^{-1}$	$2\text{Cl}^- \rightarrow 2\text{e}^- + \text{Cl}_2$
Cathode	$2\text{H}^+(\text{aq}) + 2e^- \rightarrow \text{H}_2(g)$	$Ni^{2+}(aq) + 2e^{-} \rightarrow Ni(s)$	$K^+ + e^- \rightarrow K$
Overall	$2H^+(aq) + 2Br^-(aq) \rightarrow H_2(g) +$	$2I^{-}(aq) + Ni^{2+}(aq) \rightarrow Ni(s) +$	$2K^+ + 2Cl^- \rightarrow K + Cl_2$
	$Br_2(aq)$	$I_2(aq)$	
Min Voltage	1.08 V	0.78 V	Cannot be determined from data sheet as not standard conditions
Observe	Colourless gas and orange solution	Colourless gas and brown solution	Green pungent gas and silver solid

	PbBr ₂	CoCl ₂
Products	Pb and Br ₂	Co and Cl ₂
Anode	$2Br(aq) \rightarrow 2e + Br_2(aq)$	$2\text{Cl}^{-} \rightarrow 2\text{e}^{-} + \text{Cl}_2$
Cathode	$Pb^{2+}(aq) + 2e^{-} \rightarrow Pb(s)$	$\operatorname{Co}^{2+}(\operatorname{aq}) + 2e^{-} \rightarrow \operatorname{Co}(s)$
Overall	$Pb^{2+} + 2Br^{-} \rightarrow Pb + B_{r2}$	$\operatorname{Co}^{2^+} + 2\operatorname{Cl}^- \to \operatorname{Co}(s) + \operatorname{Cl}_2(g)$
Min Voltage	Cannot be determined from data	Cannot be determined from data
_	sheet as not standard conditions	sheet as not standard conditions
Observe	Red liquid and silver/grey solid	Green pungent gas and silver
		solid

(a) Aluminium metal and chlorine gas (b) Hydrogen and oxygen gas. E^o favours less reactive substances 3.

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Cathode

Set 23: Oxidation and Reduction: no answers provided

Set 25: Reactions of organic compounds

 $\begin{array}{c} H \\ H \\ - C \\ H \end{array} + C I \\ H \\ - C \\ - H \end{array} + C I \\ - C \\ + \\ H \\ - C \\ - C \\ - C \\ + \\ - C \\ - C \\ - C \\ + \\ - C \\ - C \\ - C \\ - C \\ + \\ - C \\$

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- 7. (a) butan-2-ol, and acidified MnO_4^- or acidified $Cr_2O_7^{2-}$.
 - 5CH₃CHOHCH₂CH₃ + 2MnO₄⁻ + 6H+ \rightarrow 5 CH₃COCH₂CH₃ + 2Mn²⁺ + 8H₂O (b) methane, CH₄, and chlorine. CH₄ + C $\ell_2 \rightarrow$ CH₃C ℓ + HC ℓ then
 - $CH_3C\ell + C\ell_2 \rightarrow CH_2C\ell_2 + HC\ell$ finally $CH_2C\ell_2 + C\ell_2 \rightarrow CHC\ell_3 + HC\ell$ (c) pentan-l-ol, $CH_3CH_2CH_2CH_2CH_2OH$, and acidified MnO_4^- or acidified $Cr_2O_7^{-2-}$.
 - 5CH₃CH₂CH₂CH₂OH + 4MnO₄⁻ + 12H⁺ \rightarrow 5 CH₃CH₂CH₂CH₂COOH + 4Mn²⁺ +11H₂O
 - (d) propanoic acid, propan-l-ol, CH₃CH₂CH₂OH and conc sulfuric acid. CH₃CH₂COOH + CH₃CH₂CH₂OH → CH₃CH₂COOCH₂CH₃ + H₂O
 - (e) Cyclopentene and water.

- (f) hex-2-ene, CH₃CH=CHCH₂CH₂CH₃ and bromine. CH₃CH=CHCH₂CH₂CH₃ + Br₂ \rightarrow CH₃CHBrCHBrCH₂CH₂CH₃
- (g) propene, CH₃CH=CH₂ and hydrogen chloride: CH₃CH=CH₂ + HC ℓ → CH₃CHC ℓ CH₃
- (h) pentan-l-ol, $CH_3CH_2CH_2CH_2CH_2OH$, and acidified $Cr_2O_7^{2-}$ in limited quantities. $3CH_3CH_2CH_2CH_2CH_2OH + Cr_2O_7^{2-} + 8H^+ \rightarrow 3 CH_3CH_2CH_2CH_2CHO + 2Cr^{3+} + 7H_2O$
- (i) ethanol, CH_3CH_2OH , methanoic acid and sulfuric acid: $CH_3CH_2OH + HCOOH \rightarrow HCOOCH_2CH_3 + H_2O$
- (j) but-1-ene, $CH_2=CHCH_2CH_3$, and water: $CH_2=CHCH_2CH_3 + H2O \rightarrow CH_3CHOHCH_2CH_3$

Set 26: Calculations involving carbon compounds

- 1. $CH_3CH=CH_2 + H_2O \rightarrow CH_3CH_2CH_2OH$ n(propanol) = m/M = 1000 / 60.09 = 16.64 mol m(CH_3CH=CH_2) = nM = 16.64 x 42.078 = 700.1 g = 0.700 kg
- 2. 2 CH₃OH + 2 Na \rightarrow 2 CH₃ONa + H₂
 - (a) $n(CH_3OH) = m/M = 250 / 32.04 = 7.803 \text{ mol}$ $m(CH_3ONa) = nM = 7.803 \text{ x} 54.024 = 42155 = 422 \text{ g}$
 - (b) n(H₂) = ½ n(CH₃OH) V(H₂)stp = n x 22.71 = 3.901 x 22.71 = 88.6 L
 (c) Using PV = nRT, V = nRT/P = 7.803 x 8.314 x (273+23.0) / 102.4 = V₂ = 93.8 L
 - (c) Using $1^{\circ} = 11(1^{\circ}, \sqrt{-11}(1^{\circ}) = 1.005 \times 0.514 \times (275 + 25.0) + 102.4 = \sqrt{-100}$
- 3. $C_3H_8 + 5 O_2 \rightarrow 3 CO_2 + 4 H_2O$ $V(CO_2) = 3V(C_3H_8) = 3 \times 1.00 = 3.00 L$
- 4. CH₃(CH₂)₁₁C₆H₄SO₃H + NaOH → CH₃(CH₂)₁₁C₆H₄SO₃Na + H₂O
 (a) Sulfonic acid : n = m/M = 196 mol Sodium hydroxide : n = m/M = 8500 / 39.99 = 212.6 mol ∴ NaOH is INXS and sulfonic acid is the limiting reagent m(CH₃(CH₂)₁₁C₆H₄SO₃Na) = 348.5 x 196 = 68.3 kg
 (b) m(NaOH)_{left} = 212.6 - 196 = 16.6 mol M = n x M = 16.6 x 39.99 = 664 g = 0.664 kg
- 5. (a) $n[C_{10}H_8] = m/M = 50.0 / 128.2 = 0.3900 \text{ mol}$ = $n/v = 0.3900 / 0.250 = 1.56 \text{ mol } L^{-1}$ (b) $C_1V_1 = C_2V_2 = 1.56 \text{ x } 0.25 = 2.80 \text{ x } V_2$ $\therefore V_2 = 0.1393 \text{ L}$ $V(H_2O)_{\text{to remove}} = V_1 - V_2 = 0.250 \text{ -} 0.1393 = 0.1106 = 0.111 \text{ L} = 111 \text{ mL}$
- 6. 2 $(CH_3(CH_2)_{16}COONa(aq) + CaCl_2(aq) \rightarrow ((CH_3(CH_2)_{16}COO)_2)Ca(s) + 2 NaCl(aq)$ $n((CH_3(CH_2)_{16}COO)_2)Ca = m/M = 1.25 / 607.02 = 2.059 \times 10^{-3} mol$ \therefore there was 2 x 2.059 x 10⁻³ mol of soap $M = n \times M = 2 \times 2.059 \times 10^{-3} \times 306.45 = 1.262 \text{ g soap}$ $\% = \text{soap} / \text{mixture x } 100 = 1.262 / 10.0 \times 100 = 12.6\%$

- 7. HTa + NaOH \rightarrow NaT + H₂O (a) n(NaOH) = c x V = 1.506 x 10⁻⁵ x 0.01635 = 2.462 x 10⁻⁷ mol NaOH used As mol ratio is 1:1 there were 2.462 x 10⁻⁷ mol HTa in 20 mL must [titratable acid] = n/V = 2.462 x 10⁻⁷ / 0.02 = 1.23 x 10⁻⁵ mol L⁻¹
 - (b) Ionised to 2% means there is $1.23 \times 10^{-5} \times 0.02 = 2.46 \times 10^{-3} \text{ mol H}^+$ pH = 2.609
- 8. $MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O$ $CH_3CH_2OH + H_2O \rightarrow CH_3COOH + 4H^+ + 4e^ 5 CH_3CH_2OH + 4 MnO_4^- + 12H^+ \rightarrow 4 Mn^{2+} + 5 CH_3COOH + 11H_2O$ 5.00 mL sample Diluted to 500 mL 20 mL sample taken Titres: 23.56, 23.38, 23.12, 23.05 So mean of last two = 23.085 mL $n(MnO_4^-) = c \ x \ V = 0.0216 \ x \ 0.023085 = 0.000498636 \text{ mol}$ $\therefore \text{ as } MR = 4:5 \text{ there was } 5/4 \ x \ 0.000498636 \text{ mol alcohol in the } 20 \ \text{mL sample} = 0.000623295 \text{ mol}$ So in the whole 500 mL there was 25 x $0.000623295 = 0.015582375 \text{ mol in } 5 \ \text{mL wine}$ So in 1L there was 200 x $0.015582375 = 3.116 \ \text{mol } L^{-1} = 3.12 \ \text{mol } L^{-1}$
 - (b) $[CH_3CH_2OH]_{\text{wine in g per L}} = n \times M = 3.12 \times 46.06 = 144 \text{ g L}^{-1}$
- 9 $CH_3NH_2 + HCl \rightarrow CH_3NH_3Cl$

 25.00 mL sample
 Diluted to 250 mL
 20 mL sample taken

 Titres : 25.08, 24.86, 25.13, 24.79, 24.88 Mean of last 4 = 24.915 mL

 n(HCl) = c x V = 0.0507 x 0.024915 = 0.001263 mol HCl

 ∴ there were 0.001263 mol of methanamine in the 20 mL sample

 ∴ in the diluted sample there were 12½ times as much = 12.5 x 0.001263 = 0.0157899 mol methanamine in 25 mL original sample

 ↓ in 1L there were 40 m 0.0157800 = 0.622 mol L⁻¹

: in 1L there was 40 x as much = 40 x $0.0157899 = 0.632 \text{ mol } \text{L}^{-1}$

- (b) $m = n \ge M = 0.632 \ge 31.0571 = 19.628 \ g / L$ $\rho = m/V \ so \ m = \rho \ge V = 1.07 \ge 1000 = 1070 \ g/L$ % (CH₃NH₂) = 19.628 / 1070 \u2202 1000 = 1.83 %
- 10. CH₃CHOHCH₃ → CH₃COCH₃ + 2H⁺ + 2e⁻ MnO₄⁻ + 8H⁺ + 5e⁻ → Mn²⁺ + 4H₂O 5CH₃CHOHCH₃ + 2MnO₄⁻ + 6H⁺ + → 5CH₃COCH₃ + 2Mn²⁺ + 8H₂O <u>5.00 mL sample</u> Diluted to 250 mL <u>20 mL sample taken</u> n(MnO₄⁻) = c x V = 0.0203 x 0.02873 = 0.0005832 mol As MR is 2: 5 there were 5/2 x 0.0005832 mol propan-2-ol in the 20 mL sample = 0.001458 mol ∴ in the 250 mL solution there was 12¹/₂ times as much = 12.5 x 0.001458 = 0.018225 mol This was originally from a 5.00 mL C = n/V = 0.018225 / 0.005 = 3.645 mol L-1 in the original substance m = n x M = 3.645 x 60.0950 = 219 g L⁻¹

Set 27: Empirical, molecular and structural formula

(a) Let there be 100 g of substance

1.

(c)

Let there be 100	5 of substance	
Symbol	Mass	Mol
Ċ	85.7g	85.7/12.01 = 7.1357
Н	14.3 g	14.3 / 1.008 = 14.186
Ratio = $1:2 \div \text{the}$	e Empirical Formu	ıla is CH ₂

(b) PV = nRT so n = PV / RT = (105 x 1.18) / (8.314 x 298) = 0.0500 molIf 2.80 g is 0.005 mol then 1 mol is 2.80/0.05 = 56 g Mass of the EF is 14 – the MF mass is 56. The ratio is 4:1 so the MF is C₄H₈

- (d) Bromine adds to atoms either end of the double bond so the double bond is between atoms 2 and 3 therefore formula is but-2-ene
- 2. (a) CHO O_2 CO_2 H₂O 3.45 g 6.6 g 4.05 g n = m/M- 1.802 C n = m/M-0.45337 H =6.6/44 = 0.1500=4.05/18.01 $= 0.22487 \text{ mol H}_2\text{O}$ mol ∴ 1.19463 g of O : 0.4498 mol of H ∴ 0.150 mol of C n = m/M = 1.19463 / 16 = $m = n \ge M = 0.150 \ge 0.150 \ge 0.150 \ge 0.150 \ge 0.0150 = 0.$ = 0.45337 g of H 0.0747 mol 12.01 = 1.802 g of C

 $C_{0.150}H_{0.4498}O_{0.0747}$ divide though by $0.0747 = C_2H_6O_1$

- (b) PV = nRT so $n = PV / RT = (98.0 \times 0.950) / (8.314 \times 373) = 0.030021$ mol If 1.38 g is 0.0300 mol then 1 mol is 1.38/0.03 = 46 g $M_r(C_2H_6O) = 46.068$ Relative molecular mass is the same as the relative empirical formula mass so the molecular formula is C_2H_6O
- (c) Possible structures include: CH₃CH₂OH and CH₃-O-CH₃
- (d) CH₃CH₂OH
- 3. (a)

 O_2 CHO CO_2 H_2O 0.682 g 0.968 g 0.594 g - 0.2642 g C n = m/Mn = m/M-0.06649 g H =0.968/44 = 0.022=0.594/18.01 mol $= 0.03298 \text{ mol H}_2\text{O}$: 0.35131 g of O ∴ 0.06596 mol of H $\therefore 0.022 \text{ mol of C}$ n = m/M = 0.35131 / 16 = $m = n \ge M = 0.022 \ge 1000$ = 0.06649 g of H 0.021956 mol 12.01 = 0.2642 g of С Formula : $C_{0.022}H_{0.06596}O_{0.021956} = CH_3O$

(b) For 0.744 g sample

PV = nRT so n = PV / RT = (95.0 x 0.497) / (8.314 x 473) = 0.0120062 molIf 0.744 g is 0.0120062 mol then 1 mol is 0.744 / 0.0120062 = 61.97 g $M_r = 61.975$, C_2H_6O

(c) Possible structures are.

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4.

CHNO H_2O O_2 CO_2 1.758 g 0.900 g 1.180 g - 0.4804 g C n = m/Mn = m/M-0.1007 g H =1.758/44 = 0.03995=0.900/18.01-0.560268 g of N₂ mol $= 0.04997 \text{ mol H}_2\text{O}$ ∴ 0.038632 g of O ∴ 0.0400 mol of C ∴ 0.100 mol of H n = m/M = 0.038632 / 16 $m = n \ge M = 0.0400$ = 0.1007 g H= 0.0024145 molx 12.01 = 0.4804 g C

A second sample: 1.180 g $n = PV / RT = (105.0 \times 0.471) / (8.314 \times 298) = 0.0200 \text{ mol nitrogen gas } N_2$ $m = n \times M = 0.0200 \times 28.013 = 0.560268 \text{ g of } N_2$ $n(N_2) = m(N_2)/M(N_2) = 0.560268 / 28.02 = 0.02 \text{ mol}$ $n(N) = 2 n(N_2) = 0.04 \text{ mol}$

(a) Empirical Formula : $C_{0.0400}H_{0.100}N_{0.0400}O_{0.0024145} = C_2H_5N_2O$

A third sample: 0.5896 g

 $n = PV / RT = 95.5 \times 0.281 / 8.314 \times 323.15 = 9.988 \times 10^{-3} mol$

 $M = m/n = 0.5896 / 9.988 \times 10^{-3} = 59.03$

- (b) E.F. mass = 73 yet M = 59 as per sample 3 the Molecular formula C_2H_5NO has M = 59 but this is not the EF based on sample 2 so there is an error in the question.
- (c) Possible structures for a compound with this molecular formula, C₂H₅NO include. NH₂CH₂CHO CH₃CONH₂
- 5. (a)

CHN O_2 CO_2 H_2O 1.76 g 0.620 g -0.4804 g C n = m/M=1.76/44 = 0.04 mol -0.093 g of N : 0.0400 mol of C ∴ 0.0466 g of H n(H) = m/M = 0.0466 / 1.008 $m = n \ge M = 0.0400 \ge 12.01$ = 0.0462 mol= 0.4804 g of C n(N) = m/M = 0.093/14= 0.00664 mol2nd sample 0.232 g n = PV / RT = (101.3 x 0.0295) / (8.314 x 288) = 0.001248 mol gas N₂ $m(N_2) = n \times M = 0.001248 \times 28.01 = 00.349 g$ \therefore % N₂ = 0.0349 / 0.232 x 100 = 15.04% \therefore in the original experiment there was 0.1504 x 0.620 = 0.093 g of nitrogen Empirical Formula $C_{0.0400}H_{0.0462}N_{0.00664} = C_6H_7N$ (b) 3^{rd} sample – let V = 1Ln = PV / RT = (101.3 x 1) / (8.314 x 373) = 0.03266 $\rho = m / v$ so $m = \rho x V = 3.04 g$ $\therefore 3.04 \text{ g} = 0.03266 \text{ mol}$ So 1 mol = 93.080 g $M_r(C_6H_7N) = 93.126$, So Molecular formula = Empirical Formula = C_6H_7N

Let there be 1L of gas n = PV / RT = (105.0 x 1) / (8.314 x 423) = 0.02985 mol $\rho = m /v \text{ so } m = \rho \text{ x } V = 2.62 \text{ g}$ $\therefore 2.62 \text{ g} = 0.02985 \text{ mol so } 1 \text{ mol} = 2.62 / 0.02985 = 87.7721$ $Mr(C_2H_4O) = 44.0526 \therefore MF = EF \text{ x } 2 = C_4H_8O_2$

(c) Three solutions all esters e.g.

Set 28: Amino Acids

- 2. Enzymes are biological catalysts that speed up chemical reactions. In biological systems, other ways of speeding up reactions, such as increasing temperature, concentration or pressure are often not achievable without killing the organism. Without enzyme catalysts, many cellular reactions would not take place under cellular conditions.
- 3. 3 you would need a water molecule to break each peptide bond (as one water was released when each peptide bond was formed)
- 4. 99 one water molecule is created with each peptide bond formed, so 1 water is released when the first 2 amino acids join then one water is released for each peptide bond formed when adding an addition amino acid.

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Name	Symbol	Name	Symbol	Name	Symbol
aspartic acid	Asp	glycine	Gly	lysine	Lys
				Structur	e
Structure O H N-C- H' C C C C C C C C	D [⊖] H 2	Structur O H N-C- H H H	e ∠O [⊖] - H	O H N-C- H CH CH CH CH CH CH CH CH	.0 ⁹ - Н I ₂ I ₂ I ₂ Н

Set 29: Proteins

- 1. The primary structure is the linear sequence of amino acid residues in a protein.
 - The secondary structure is the structure that arises from the arrangement of hydrogen bonds that occurs between the oxygen and the hydrogen atoms of the amide bonds that occur in the proteins backbone. Oxygen's and hydrogen's on the side chains also form hydrogen bonds, but these ones do not participate in secondary structure. Some secondary structures are alpha helix (shown below) and beta sheet.

Hydrogen bonds between amino acids at different locations in polypeptde chain

α-helix

• The tertiary structure is the overall shape that a polypeptide chain forms. This shape arises largely through the interactions of the side chains, and includes hydrogen bonding, hydrophobic interactions, ionic bonding (NH₃⁺ COO⁻), and disulphide bonding between cysteine side chains (cys-S-S-cys). See figure opposite.

2.

6.

Protein structure	Bonding forces
Primary	covalent bonds
Secondary	hydrogen bonds
Tertiary	dispersion forces, hydrogen bonds, ionic bonds, disulfide bonds

- (a) The α-helix secondary structure, every N-H group donates a hydrogen bond to the C=O group of an amino acid four residues earlier in the chain. Beta sheets consist of strands connected by at least two backbone hydrogen bonds, forming a pleated sheet
 - (b) A quaternary structure because there are multiple peptide chains (protein molecules) in each complex.
 - (c) Glutamic acid is hydrophilic and would aggregate around water molecules, valine is hydrophobic. It could also disrupt any hydrogen bonding or ionic bonding (salt bridges) the glutamic acid residue was involved in.
- 4. The different amino acid sequences and consequently their 3d structure, makes different proteins unique.
- 5. A protein's shape is determined by its sequence of amino acids. The different side chains on these determine the mix of hydrogen bonds, ionic bonds, disulfide bridges or dispersion forces that give it its shape.

(a)
$$O_{C}O$$

 $H_{\oplus}I_{H}-N-C-H$
 $H_{H}B$

(b) The length and nature of the amino acids and their affinity for other species.

- 7. (a) α -helices are held together by hydrogen bonds.
 - (b) N-H and C=O
 - (c) β -sheets are held together by multiple hydrogen bonds
 - (d) N-H and C=O
 - (e) Hydrogen bonds are secondary bond and will rupture at high temperatures eg: cooking meat.
- 8. Disulfide bridges are the strongest and most thermo stable. They are covalent bonds requiring approximately 60kJ/mole to break, compared to 20kJ/mol for hydrogen bonds.
- 9. The presence of concentrated sodium hydroxide will cause the amide groups to hydrolyse. This means that proteins will split into their component amino acids. At lower concentrations, hydrogen bonding and tertiary structure will be disrupted. The flesh will dissolve.

Properties, reactions and chemical synthesis

Set 30: Reaction types

1

2.

- $CH_3COOH(\ell) + H_2O(\ell) \rightarrow CH_3COO^-(aq) + H_3O^+(aq)$ (a) $NH_3(g) + H_2O(\ell) \rightarrow NH_4^+(aq) + H_3O^+(aq)$ (b) $NaHCO_3 (s) \rightarrow HCO_3(aq) + Na^+ (aq)$ (c) $NaHSO_4(s) \rightarrow Na^+(aq) + HSO_4^-(aq)$ (d) K_2CO_3 (s) $\rightarrow 2K^+$ (aq) $+ CO_3^{2-}$ (aq) (e) $NH_4CH_3COO(s) \rightarrow NH_4^+ (aq) + CH_3COO^- (aq)$ (f) $CH_3COO-(aq) + H_2O(\ell) \rightleftharpoons CH_3COOH(aq) + OH^-(aq)$ i) $H^+(aq) + OH^-(aq) \rightarrow H_2O(\ell)$ (a) ii) Two colourless solutions are mixed together. No visible reaction; some heat evolved. i) $Ba(OH)_2(s) + 2H^+(aq) \rightarrow Ba^{2+}(aq) + 2H_2O(\ell)$ (b) ii) A white solid dissolves in a colourless solution. i) MgO(s) + 2H⁺ (aq) \rightarrow Mg²⁺ (aq) + H₂O(ℓ) (c) ii) A white solid dissolves in a colourless solution. i) $CH_3COOH(aq) + NH_3(aq) \rightarrow NH_4^+(aq) + CH_3COO^-(aq)$ (d) i) $Zn(s) + 2H^+(aq) \rightarrow Zn^{2+}(aq) + H_2(g)$ (e) (f) (g) i) Ni(s) + 2H⁺ (aq) \rightarrow Ni²⁺ (aq) + H₂ (g) (h) i) $Fe(s) + 2H^+(aq) \rightarrow Fe^{2+}(aq) + H_2(q)$ (1)i) $Br_2(\ell) + 2l^-(aq) \rightarrow 2Br^-(aq) + I_2(s)$ (a) i) $Mg(s) + Fe^{2+}(aq) \rightarrow Mg^{2+}(aq) + Fe(s)$ b) i) $Cu(s) + 2Ag^+(aq) \rightarrow Cu^{2+}(aq) + 2Ag(s)$ c) i) $Zn(s) + Ni^{2+}(aq) \rightarrow Zn^{2+}(aq) + Ni(s)$ d) i) $2Na(s) + 2H2O(\ell) \rightarrow 2Na^+(aq) + 2OH^-(aq) + H_2(g)$ e) i) $2K(s) + 2HzO(\ell) \rightarrow 2K^+(aq) + 2OH^-(aq) + H_2(q)$ f) i) $C\ell_2(g) + 2Br(aq) \rightarrow 2C\ell(aq) + Br_2(aq)$ g) i) $Ag+(aq) + C\ell^{-}(aq) \rightarrow AgC\ell(s)$ a) ii) Two colourless solutions are mixed to form a white precipitate. i) $Ag^{+}(aq) + Br^{-}(aq) \rightarrow AgBr(s)$ b) ii) Two colourless solutions are mixed to form a cream/white solid. i) $Pb^{2+}(aq) + 2I^{-}(aq) \rightarrow PbI_{2}(s)$ c) i) $Ca^{2+}(aq) + SO_4^{2-}(aq) \rightarrow CaSO4(s)$ d) ii) Two colourless solutions are mixed to form a white precipitate i) $Ba^{2+}(aq) + OH^{-}(aq) + SO_4^{2-}(aq) + H^{+}(aq) \rightarrow BaSO_4(s) + H_2O(\ell)$ e) ii) Two colourless solutions are mixed to form a white precipitate. i) $Fe^{2+}(aq) + CO_3^{2-}(aq) \rightarrow FeCO_3(s)$ f) i) $3 \operatorname{Zn}^{2+}(aq) + 2 \operatorname{PO}_4^{3-}(aq) \rightarrow \operatorname{Zn}_3(\operatorname{PO}_4)_2(s)$ g) STAWA12T001 | Exploring Chemistry | Year 12 worked solutions
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- $NaHCO_3 (s) \rightarrow Na^+ (aq) + HCO^- (aq)$
- $HCO_3^-(aq) + H_2O(\ell) \rightarrow CO_3^{2-}(aq) + H_3O^+(aq)$
- $HSO_4(aq) + H_2O(\ell) \rightarrow SO_4^{2}(aq) + H_3O^{+}(aq)$
- $\text{CO}_3^2(\text{aq}) + \text{H}_2\text{O}(\ell) \rightleftharpoons \text{HCO}_3^2(\text{aq}) + \text{OH}^2(\text{aq})$
- $NH_4^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$
- ii) Two colourless solutions mixed. No visible reactions. There is a reduction in the vinegar smell.
- ii) A silver solid dissolves in a colourless solution; colourless, odourless gas evolved
- i) $2CH_3COOH(aq) + Mg(s) \rightarrow Mg^{2+}(aq) + 2CH_3COO^{-}(aq) + H_2(g)$ ii) A silver solid dissolves in a colourless solution; colourless, odourless gas evolved.
- i) $Cu(s) + 4H^+(aq) + 2NO_3(aq) \rightarrow Cu^{2+}(aq) + 2NO_2(g) + 2H_2O(\ell)$
- ii) Brown solid dissolves in colourless solution to produce brown, pungent gas and a blue solution.
- ii) Silver solid dissolves in colourless solution forms colourless, odourless gas and green solution.
 - ii) Silver solid dissolves in colourless solution, colourless, odourless gas evolved and a pale green soln.
- 3.
 - ii) Brown/orange liquid added to colourless solution. Brown/orange fades and dark brown solid forms
 - ii) Silver solid to pale green solution. Black precipitate on silver solid; pale-green soln to colourless.
 - ii) Brown solid to colourless solution. Black precipitate on brown solid; colourless solution turns blue.
 - ii) Silver solid to green solution. Black precipitate on silver solid; green solution fades to colourless.
 - ii) A silver/white solid reacts vigorously with a colourless liquid to form a colourless, odourless gas.
 - ii) A silver/white solid reacts vigorously with a colourless liquid to produce a colourless, odourless gas.
 - ii) A green pungent gas dissolves in a colourless solution to form a brown/orange solutions.
- 4.
 - ii) Two colourless solutions are mixed to form a yellow precipitate.
 - ii) Pale green solution mixed with colourless solution forms pale green precipitate. Green solution fades.

- ii) Two colourless solutions are mixed together to form a white precipitate.
- i) $\operatorname{Cu}^{2+}(\operatorname{aq}) + 2 \operatorname{OH}^{-}(\operatorname{aq}) \to \operatorname{Cu}(\operatorname{OH})_2(\operatorname{s})$
- ii) Blue solution mixed with a colourless solution to form a blue precipitate. Blue solution colour fades. i) $2 \operatorname{Cr}^{3+}(aq) + 3 \operatorname{CO}_{3}^{2-}(aq) \rightarrow \operatorname{Cr}_{2}(\operatorname{CO}_{3})_{2}(s)$
- i) i) $2 \operatorname{Cr}^{3+}(\operatorname{aq}) + 3 \operatorname{CO}_{3}^{2-}(\operatorname{aq}) \rightarrow \operatorname{Cr}_{2}(\operatorname{CO}_{3})_{2}(s)$ ii) Green solution mixed with colourless solution to form a green precipitate. Green solution colour fades.

Set 31: Percentage composition and yield

1. a) $M(Fe_2O_3) = 159.7 \text{ g mol}^{-1}$ %Fe:= 111 7 / 159.7 x100 = 69.9%

h)

- b) %Heamatite: $65/69.9 \times 100 = 92.9\%$
- 2. $Zn + 2H^+ \rightarrow Zn^{2+} + H_2$ m(remaining) = m(Cu) = 0.630 g m(Zn) = 2.71 - 0.630 = 2.08 g %Zn: 2.08/2.71 x 100 = 76.8%
- 3 a) $n(Br_2) = 125 / (79.9 \text{ x } 2) = 0.782 \text{ mol}$ $n(C_6H_6) = 60.0 / 78.108 = 0.768 \text{ mol}$ 1 mol of Br₂ requires 1 mol of C_6H_6 $0.782 \text{ mol of } Br_2 \text{ requires } 0.782 \text{ mol of } C_6H_6$ $n(C_6H_6 \text{ required}) > n(C_6H_6 \text{ available})$ $C_6H_6 \text{ is } LR$ $n(C_6H_3Br) = n(C_6H_6) = 0.768 \text{ mol}$ $m(C_6H_5Br) = 0.768 \text{ x } 157 = 121 \text{ g}$ b) %yield= 93 2/121 x 100 = 77.3 %
- 4. a) $Ca(OH)_2 \rightarrow CaO + H_2O$ b) %CaO: 4 33/5.67 x100 = 76.4 %
- 5. a) $Ba^{2+}(aq) + SO_4^{2-}(aq) \rightarrow BaSO_4(s)$ $n(Ba^{2+}) = n(CuSO_4.5H_2O)$ $n(CuSO_4.5H_2O) = 1.11/249.69 = 4.45 \times 10^{-3} \text{ mol} = n(BaC\ell_2)$ $m(BaC\ell_2) = (4.45 \times 10^{-3}) \times (137.3 + 70.9) = 0.926 \text{ g}$
 - c) $n(CuC\ell_2.2H_2O) = n(BaC\ell_2 used) = 4.45 \times 10^{-3} \text{ mol}$ $m(CuC\ell_2.2H_2O) = (4.45 \times 10^{-3}) \times 170.482 = 0.758 \text{ g}$ %yield:- 0 345/0.758 x 100 = 45.5 %
- 6. $n(Na_2S_2O_7) = \frac{1}{2} n(S)$ $n(S) = 17500/32.06 = 5.46 \times 10^2 \text{ mol}$ $n(Na_2S_2O_7) = \frac{1}{2} (5.46 \times 10^2) = 2.73 \times 10^2 \text{ mol}$ $m(Na_2S_2O_7) = (2.73 \times 10^2) \times (45.98+64.12+112) = 6.06 \times 10^4 \text{ g}$ $\% \text{ yield:} 50000/60617 \times 100 = 82.5 \%$
- 7. let $X = m(NaHCO_3)$ $Y = m(Na_2CO_3)$ X + Y = 100Y = 100 - X

$$\begin{split} n(\text{Na}_2\text{CO}_3) &= 90.7 \ / \ 106 = 0.856 \ \text{mol} \\ n(\text{Na}_2\text{CO}_3 \ \text{total}) &= \frac{1}{2} \ n(\text{Na}\text{HCO}_3 \ \text{initial}) + n(\text{Na}_2\text{CO}_3 \ \text{initial}) \\ &= (\frac{1}{2} \ x \ X/84.01) + Y/100 \\ 0.856 &= X/168 + (100\text{-}x)/106 \\ 62X &= 1556.352 \\ X &= 25.1 \ \text{g} \\ Y &= 74.9 \ \text{g} \\ \%\text{Na}_2\text{CO}_3 &= 74.9 \ / \ 100 \ x \ 100 = 74.9 \ \% \end{split}$$

1	2	3	4	5
1.63g wire	Dissolved in acid	Made up to 250 mL	20 mL sample	Titrated with 18.1 mL of 0.0209 M K ₂ Cr ₂ O ₇

Step 5

Reduction: $Cr_2O_7^{2^-} + 14H^{+} + 6e^- \longrightarrow 2Cr^{3^+} + 7H_2O$ Oxidation: $Fe^{2^+} \longrightarrow Fe^{3^+} + e^-$ Combined : $Cr_2O_7^{2^-} + 14H^{+} + 6e^- + 6Fe^{2^+} \longrightarrow 2Cr^{3^+} + 7H_2O + 6Fe^{3^+} + 6e^-$ Mol Ratio $Cr_2O_7^{2^-}$: Fe = 1:6In 20mL sample n ($Cr_2O_7^{2^-}$) - = c xv = 0.0181 x 0.0209 = 0.0003783 mol ∴ n(Fe^{2^+}) = 6 x 0.00037829 = 0.002269 mol Fe^{2^+} m(Fe) = 0.002269 x 55.85 = 0.1267 g ∴ In 250 mL there was 12.5 x 0.1267 = 1.584 g Fe

 \therefore percentage purity of the wire = 1.5844/1.63 x 100 = 97.2% pure

9.

8.

 $\begin{array}{|c|c|c|c|c|} \hline 1 & 2 & 3 \\ \hline 1.27 \text{ g mineral} & Converted to K_2Cr_2O_7 & Titrated with 37.5 mL of 0.400 M FeSO_4 \\ \hline \text{Step 3} \\ \hline \text{Reduction: } Cr_2O_7^{-2} + 14H^{\dagger} + 6e^{-} \longrightarrow 2Cr^{3+} + 7H_2O \\ \hline \text{Oxidation: } Fe^{2+} & \longrightarrow Fe^{3+} + e^{-} \\ \hline \text{Combined : } Cr_2O_7^{-2} + 14H^{\dagger} + 6e^{-} + 6Fe & \longrightarrow 2Cr^{3+} + 7H_2O + 6Fe^{2+} + 6e^{-} \\ \hline \text{Mol Ratio } Cr_2O_7^{-2} : Fe = 1:6 \\ n (Fe^{2+}) - = c xv = 0.0375 x 0.400 = 0.015 \text{ mol Fe}^{2+} \\ \therefore n(Cr_2O_7^{-2-}) = 1/6 x 0.015 = 0.0025 \text{ mol} \\ \therefore n(Cr) = 2 x 0.0025 = 0.005 \text{ mol} \\ \therefore m(Cr) = 52.00 x 0.005 = 0.26 \text{ g} \\ \end{array}$

 \therefore percentage purity of mineral = 0.26/1.27 x 100 = 20.5 % pure

10.

 $\frac{1}{0.752 \text{ g impure Na}_2\text{SO}_3} = \frac{2}{10.752 \text{ g impure Na}_2\text{SO}_4} = \frac{2}{10.752 \text{ g impure Na}_2\text{SO}_3} = \frac{2}{10.752 \text{ g impure Na}_2\text{SO}_4} = \frac{2}$

: percentage purity = $0.6458/0.752 \times 100 = 85.9$ % pure

11.

1		2	3	4	5
3	.08 g sample	Dissolved in	Made up to 250 mL	25 mL sample	Titrated with 28.7 mL of
		sulfuric acid			0.0260 M KMnO ₄
Red	luction: MnO_4^{-1}	' + 8H ⁺ + 5e ⁻ —	\rightarrow Mn ²⁺ + 4H ₂ O		
Oxidation: $Fe^{2^+} \longrightarrow Fe^{3^+} + e^-$					
Combined : $MnO_4^- + 8H^+ + 5e^- + 5Fe^{2+} \longrightarrow 5Fe^{3+} + 5e^- + Mn^{2+} + 4H_2O$					

Mol Ratio MnO_4^- : $Fe^{2+} = 1:5$

n (MnO₄⁻)- = c xv = 0.0287 x 0.0260 = 0.0007462 mol In 25 mL sample ∴ n(Fe) = 5 x 0.0007462 = 0.003731 mol ∴ n(Fe₂O₃) = 0.003731 / 2 = 0.0018655 mol

 \therefore m(Fe₂O₃) = 0.0018655 x 159.688 = 0.297898 g

In whole sample

 \therefore m(Fe₂O₃) = 10 x 0.297898 = 2.97898 g

: percentage purity = $2.97898 / 3.08 \times 100 = 96.7 \%$ pure

12.

1	2	3	4
FeO and	Dissolved in	Two equal	1. Titrated with 15.0 mL of 6.30 g L^{-1} KMnO ₄
Fe ₂ O ₃	sulfuric acid	aliquots	
		*	2. Reduced with zinc and Titrated with 25.1 mL of
			$6.30 \text{ g L}^{-1} \text{ KMnO}_4$

Zinc has the effect of reducing all the Fe^{3+} to Fe^{2} Step 4 6.30 g L-1 = $6.3 / 158.0339 = 0.03986 \text{ mol} / \text{L}^{-1} \text{ MnO}_4^{-1}$ Reduction: $MnO_{4}^{-} + 8H^{+} + 5e^{-} \longrightarrow Mn^{2+} + 4H_{2}O$ Oxidation: $Fe^{2+} \longrightarrow Fe^{3+} + e^{-}$ Combined : $MnO_4^- + 8H^+ + 5e^- + 5Fe^{2+} \longrightarrow 5Fe^{3+} + 5e^- + Mn^{2+} + 4H_2O$ Mol Ratio MnO_4^- : $Fe^{2+} = 1:5$ Titration 1 $n (MnO_4) = c xv = 0.03986 x 0.0150 = 0.0005979 mol$ \therefore n(Fe²⁺) = 5 x 0.0005979 = 0.0029895 mol Titration 2 $n (MnO_4) = c xv = 0.03986 x 0.0251 = 0.001000 mol$ \therefore n(Fe²⁺) = 5 x 0.001000 = 0.005002 mol \therefore n(Fe²⁺) =n(FeO)= 0.002990 mol $n(Fe^{3+}) = 2 \times n(Fe_2O_3) = 0.005002 - 0.002990 = 0.002013$ \therefore n(Fe₂O₃) = $\frac{1}{2}$ x 0.002013 = 0.001006 For the half sample \therefore m(FeO) = 0.002990 x 71.844 = 0.2148 g \therefore m(Fe₂O₃) = 0.001006 x 159.688 = 0.1607 g For the whole sample \therefore m(FeO) = 0.2148 x 2 = 00.4296 g \therefore m(Fe₂O₃) = 0.1607 x 2 = 0.3214 g

Set 32: Limiting reagents

 a) 1 mole of Pb(NO)₃)₂ requires 2 moles of KI 0300 mole of Pb(NO)₃)₂: requires 2 x 0.300 = 0.600 mol of KI n(KI required) > n(KI available) ∴ KI is LR
 b) n(PbI₂) = ½ n(KI)

b)
$$n(PbI_2) = \frac{1}{2} n(KI)$$

= $\frac{1}{2} \times 0.400 = 0.200 \text{ mol}$
 $n(PbI_2) = 0.200 \times 461.0 = 92.2 \text{ g}$

2. a) n(NaOH) = 5.55/39.998 = 0.139 mol n(HCℓ) = 4.88/36.458 = 0.134 mol 1 mol of NaOH requires 1 mol of HCℓ 0.139 mol of NaOH requires 0.139 mol of HCℓ $n(HC\ell \text{ required}) > n(HC\ell \text{ available})$ HC ℓ is LR

- b) $N(NaC\ell) = n(HC\ell) = 0.134 \text{ mol}$ $m(NaC\ell) = 0.134 \text{ x } 58.44 = 7.82 \text{ g}$
- 3. a) $n(CH_3COOH) = 4.78/60.052 = 7.96 \times 10^{-2} \text{ mol}$ $N(CaCO_3) = 2.22/100.08 = 2.22 \times 10^{-2} \text{ mol}$ 1 mole of CaCO₃ requires 2 moles of CH₃COOH 2.22 x 10⁻² mol of CaCO₃ requires 2 x (2.22 x 10⁻²) = 4.44 x 10⁻² mol CH₃COOH $n(CH_3COOH \text{ requ}) < n(CH_3COOH \text{ avail})$ $\therefore CaCO_3 \text{ is LR}$
 - b) $n(CO_2) = n(CaCO_3) = 2.22 \text{ x } 10^{-2} \text{mol}$ $m(CO_2) = (2.22 \text{ x } 10^{-2}) \text{ x } 44.01 = 0.976 \text{ g}$
 - c) $n(Ca(CH_3COO)_2) = n(CaCO_3) = 2,22 \times 10^{-2} \text{ mol}$ $m(Ca(CH_3COO)_2) = (2.22 \times 10^{-2}) \times 158.148 = 3.51 \text{ g}$
- 4. a) n(Mg)=6.08/24.3 = 0.250 mol $n(H_2SO_4) = 20.0/98.076 = 0.2039 \text{ mol}$ 1 mol of Mg requires 1 mol of H₂SO₄ 0.250 mol of Mg requires 0.250 mol of H₂SO₄ $n(H_2SO_4 \text{ req}) > n(H_2SO_4 \text{ avail})$ $\therefore H_2SO_4 \text{ is LR}$
 - b) $n(H_2) = n(H_2SO_4) = 0.2039 \text{ mol}$ $m(H_2) = 0.2039 \text{ x } 2.016 = 0.411 \text{ g}$
 - c) $n(MgSO_4.7H_2O) = n(H_2SO_4) = 0.2039 \text{ mol}$ $m(MgSO_4.7H_2O) = 0.2039 \text{ x} 246.472 = 50.3 \text{ g}$
- 5. a) $n(NaOH) = 1.600/3998 = 4.00 \times 10^{-2} \text{ mol}$ $n(H_2SO_4) = 1.472/98.076 = 1.5 \times 10^{-2} \text{ mol}$ 1 mol of H_2SO_4 requires 2 mol of NaOH 1.5 x 10⁻² mol of H_2SO_4 requires 2 x (1.50 x 10⁻²) = 3.00 x 10⁻² mol n(NaOH req) < n(NaOH avail) $\therefore H_2SO_4$ is LR
 - b) $n(Na_2SO_4) = n(H_2SO_4) = 1.50 \times 10^{-2} \text{ mol}$ $m(Na_2SO_4) = (1.50 \times 10^{-2}) \times 142.04 = 2.13 \text{ g}$ c) $n(NaOH \text{ rem}) = 4.00 \times 10^{-2} - 3.00 \times 10^{-2} = 1.00 \times 10^{-2} \text{ mol}$
 - m(NaOH) = $(1.00 \times 10^{-2}) \times 39.998 = 0.400 \text{ g}$
- a) n(Ag) = 16.25/107.9 = 0.151 mol n(HNO3) = 18.4/63.018 = 0.292 mol 1 mol of Ag requires 4/3 mol of HNO3 0.151 mol of Ag requires 4/3 x 0.151 = 0.201 mol n(HNO3 req) < n(HNO3) avail)
 ∴ Ag is LR
 - b) n(NO) = 1/3 n(Ag) = 1/3x0.151 = 0.0503 mol m(NO) = 0.0503 x 30.01 = 1.51 g
 - c) $n(HNO_3rem) = 0.292 0.201 = 0.091 mol m(HNCh) = 0.091 x 63.018 = 5.75 g$
- 7. a) $n(KO_2) = 5.00/71.1 = 7.03 \times 10^{-2} \text{ mol}$ $n(CO_2) = 9.00/44.01 = 0.204 \text{ mol}$ 1 mol of CO₂ requires 2 mol of KO₂ 0.204 mol requires 2 x 0.204 = 0.408 mol of KO₂ $n(KO_2 \text{ required}) > n(KO_2 \text{ avail})$ $\therefore KO_2 > \text{ is LR}$

 $n(K_2CO_3) = \frac{1}{2} n(KO_2) = \frac{1}{2} (7.03 \times 10^{-2}) = 3.52 \times 10^{-2} \text{ mol}$ $m(K_2CO_3) = (3.52 \times 10^{-2}) \times 138.21 = 4.86 \text{ g}$

- b) $n(O_2) = 3/2 n(KO_2) = 3/2 x (7.03 x 10^{-2}) = 0.105 mol m(O_2) = 0.105 x 32.00 = 3.36 g n(CO_2 rem) = 0.204 \frac{1}{2} (7.02 x 10^{-2}) = 0.269 mol m(CO_2) = 0.269 x 44.01 = 7.43 g$
- 8. $n(Ca_3(PO_4)_2) = 25.0 \times 10^6 / 310.18 = 8.06 \times 10^4 \text{ mol}$ $N(H_3PO_4) = 30.0 \times 10^6 / 97.94 = 3.06 \times 10^5 \text{ mol}$ 1 mol of $Ca_3(PO_4)_2$ requires 4 mol of H_3PO_4 8.06 x 10⁴ mol of $Ca_3(PO_4)_2$ requires 4 x (8.06 x 10⁴) = 3.224 x 10⁵ mol of H_3PO_4 $n(H_3PO_4 \text{ req}) > n(H_3PO_4 \text{ avail})$ $\therefore H_3PO_4 \text{ is LR}$ $n(Ca(H_2PO_4)_2) = \frac{3}{4} (n(H_3PO_4 \text{ avail}) = \frac{3}{4} (3.06 \times 10^5) = 2.296 \times 10^5 \text{ mol}$ $m(Ca(H_2PO_4)_2) = (2.296 \times 10^5) \times 234.052 = 5.37 \times 10^7 \text{ g} (53.7 \text{ tonne})$
- 9. $n(CO2) = 2.94/44.01 = 6.68 \times 10^{-2} \text{ mol}$ $n(Na_2CO_3) = n(CO_2) = 6.68 \times 10^{-2} \text{ mol}$ $m(Na_2CO_3) = (6.68 \times 10^{-2}) \times 105.99 = 7.08 \text{ g}$ %Na₂CO₃:= 7.08/7.20 x 00 = 98.3%
- 10. $n(C\ell_2) = 2.84/70.9 = 4.01 \times 10^{-2} \text{ mol}$ mol $n(MnO_2) = n(C\ell_2) = 4.01 \times 10^{-2} \text{ mol}$ $m(MnO2) = (4.01 \times 10^{-2}) \times 86.94 = 2.49 \text{ g}$ %MnO₂ = 3.49/3.52 X 100 = 99.0%

Set 33: Calculations involving gases

1.	a n=0.0	P 120.1 217 mol	V 0.45	n	R 8.314	T 299
	b n=0.0	P 99.3 938 mol	V 0.889	n	R 8.314	T 314.15
	c n=1.4	P 145 62 mol	V 27.5	n	R 8.314	T 328.15
2.(a)	P 105 n= 0.0 If Then	V 0.59 02367861 0.023678 1 mol = 9	n 3 mol 3613 mol = 93.8 g	R 8.314 = 2.22 g	T 298.15	Mass 2.22
(b)	P 98.5 n= 0.0 If 0.0 Then	V 1.22 04151639 41516396 1 mol = 1	n 6 mol 6 mol = $0.$ 11.0 g	R 8.314 456 g	T 348.15	Mass 0.456
(c)	P 68.4 n= 0.0 If Then	V 3.33 06968383 0.069683 1 mol = 9	n 4 mol 3834 92.7 g	R 8.314 mol =	T 393.15 6.46 g	Mass 6.46

- 4. P V n R T $105 \ 0.0255 \ 8.314 \ 298.15$ n= 0.001080151 mol of CO₂ \therefore there were 0.001080151 mol of CaCO₃ = 100 x 0.001080151 = 0.108 g CaCO₃
- 5. $2NH_3(g) + H_2SO_4(aq) \longrightarrow (NH_4)_2SO_4$
 - $nNH_3 = 1.50 / 22.71 = 0.06605 mol$
 - $nH_2SO_4 = c \times V = 2.54 \times 0.050 = 0.127 \text{ mol}$
 - (a) To use all the NH₃ we need 1/2 that number of moles of $H_2SO_4 = 1/2 \ge 0.03303$ mol acid needed and have 0.127 mol so acid is in excess and the LR is NH₃
 - (b) $n(NH_4)_2SO_4 \text{ produced} = \frac{1}{2} n(NH_3 \text{ used}) = \frac{1}{2} (0.06605) = 0.03303 \text{ moles}$:
 - mass ammonium sulfate = M x n = $132.094 \times 0.03303 = 4.36 \text{ g}$
 - (c) nH_2SO_4 left over = 0.127 available 0.03303 used = 0.09397 mol in excess
- 6. NaHCO₃ + HCl \longrightarrow CO₂ + H₂O + NaCl nNaHCO₃ = 0.273 / 84.00 = 0.0032497 mol
 - $nHC\ell = c \times V = 2.50 \times 0.050 = 0.125 mol$
 - (a) \therefore NaHCO₃ is the LR
 - (b) $n(CO_2) = n(NaHCO_3)$: $V(CO_2) = nRT/P = 0.085 L$
- 7. $\operatorname{Cu} + 4\operatorname{HNO}_3 \longrightarrow \operatorname{Cu}(\operatorname{NO}_3)_2 + 2\operatorname{NO}_2 + 2\operatorname{H}_2\operatorname{O}_3$
 - nCu = 1.33 / 63.55 = 0.0208 mol
 - $nHNO_3 = c x V = 6.00 x 0.025 = 0.15 mol$
 - (a) To use all the Cu we need 0.0208 x 4 mol HNO₃= 0.0832 mol We have more so the Cu is the LR
 (b) trNO = 2 x 0.0208 = 0.0416 mol
 - (b) \therefore nNO₂ = 2 x 0.0208 = 0.0416 mol V = 1.02 L
- 8. $MnO_2 + 4H^{\dagger} + 2Cl^{-} \longrightarrow Mn^{2^{+}} + Cl_2 + 2H_2O$ $Or: MnO_2 + 4HC\ell \rightarrow MnC\ell_2 + C\ell_2 + 2H_2O$ $nMnO_2 = 3.44 / 86.93 = 0.0396 \text{ mol}$ $nHC\ell = 6.20 \text{ x } 0.150 = 0.093 \text{ mol}$ $nHC\ell \text{ needed} = nH^{+} \text{ used} = 4 \text{ x } nMnO_2 = 0.158 \text{ mol}$ You need more HC ℓ than you have so HC ℓ is the limiting reagent $\therefore nC\ell_2 = \frac{1}{4} \text{ HC}\ell = \frac{1}{4}(0.093) = 0.02325 \text{ mol}$ P = nRT/V = 238.26 kPa = 238 kPa
- 9. $(NH_4)_2SO_4 + 2KNO_3 \longrightarrow K_2SO_4 + 4H_2O + 2N_2$ $n(NH_4)_2SO_4 = 30.0/132.140 = 0.227031 \text{ mol}$ $nKNO_3 = 34.0/101.1032 = 0.33629 \text{ mol}$ To use all the $(NH_4)_2SO_4$ we would need 2 x 0.22703 mol $KNO_3 = 0.45406$ mol (a) KNO_3 is the LR and is used up.
 - (b) n(NH4)2SO4 used = 1/2 nKNO3 = 0.16814 mol: n(NH4)2SO4left over = 0.22701 - 0.16814 = 0.05887 mol is INXS $m = n \times M = 0.058886 \times 132.140 = 7.78 \text{ g}$
 - (c) $nN_2 = nKNO_3 : P = nRT/V = 1814 \text{ kPa}$

Set 34: Empirical formulas 1

1.	(a) m n	Fe 2.76 2.76 55.85	Cl 3.51 3.51 35.45	
	Ratio (÷- by smallest)	0.0494 0.0494 0.0494 I	0.0990 0.990 0.494 2	
	$\mathbf{E.F.} = \mathbf{FeC}\ell_2$	1	2	
	(b)	C	ц	
	m n	2.06 2.06 12.01 0.172	0.430 0.430 1.008 0.427	
	Ratio (÷ by smallest)	0.172 0.172 0.172 2	0.427 0.172 5	
	EF C ₂ H ₅	-	C .	
	(c)	K	Mn	0
	m	3.71 3.71 39.1 9.49 x 10 ⁻²	5.21 5.21 54.94 9.48 x 10^{-2}	6.07 6.07 16.00 0.379
	Ratio (÷ by smallest)	9.49 x 10 ⁻² 9.48 x 10 ⁻²	9.48 x 10^{-2} 9.48 x 10^{-2}	0.379 9.48 x 10 ⁻²
	$\mathbf{E.F.} = \mathbf{KMnO_4}$			
	(d)	Na	S	0
	m n	3.86 3.86 22.99	5.38 5.38 32.06	10.7 10.7 16.00
	Ratio (÷ by smallest)	0.168 1.68 0.168 1	0.168 0.168 0.168 1	0.6688 0.6688 0.168 4
	$E.F = NaSO_4$			
2.	(a)	Ca 54.1 54.1 40.08	O 43.2 43.2 16.00	H 2.70 2.70 1.008
	Ratio (÷ by smallest)	1.35 1.35 1	2.70 2.70 1.35 2	2.68 1.35 2
	\mathbf{L} . \mathbf{r} . = $\mathbf{C}\mathbf{a}\mathbf{U}_{2}\mathbf{H}_{2}$ ($\mathbf{C}\mathbf{a}(\mathbf{O})$	[] [2]		

			Answers	3
	(b)			
		Pb	Ν	0
	m in 100 g	62.5	8.50	29.0
	n	62.5	8.50	29.0
		207.2	14.01	16.00
		0.302	0.607	1.812
	Ratio (÷ by smallest)	0.302	0.607	1.812
		0.302	0.302	0.302
		1	2	6
	$E.F. = PbN_2O_6, (Pb(NC))$	(₃) ₂)		
	(c)			
		С	0	Н
	m in 100 g	60.0	26.7	13.3
	n	60.0	26.7	13.3
		12.01	16.00	1.008
		5.00	1.669	13.2
	Ratio (÷ by smallest)	5.00	1.669	13.2
	× • /	1.669	1.669	1.669
		3	1	8
	$\mathbf{E.F.} = \mathbf{C_3H_8O}$			
3		S	0	
5.	m in 100 σ	40.0	60.0	
	n n n n n n n n n n n n n n n n n n n	40.0	60.0	
	11	32.06	16.00	
		1 25	3 75	
	Ratio (÷ by smallest)	1.25	3.75	
		1.25	1.25	
	$\mathbf{E.F.} = \mathbf{SO}_3$	1	3	
1	$0/C = 1.25/2.12 \times 100 =$	- 59 040/		
4.	$70C = 1.23/2.12 \times 100 =$ $9/U = 0.161/1.64 \times 100 =$	- 0.820%		
	$\frac{70\Pi - 0.101}{1.04} \times 100^{-1}$	-9.8270 -9.10270		
	/00 = 100-(38.90 + 9.8.	C = 51.2270	0	н
	m in 100 a	58.06	31.22	0.82
	n n n n n n n n n n	58.96	31.22	9.82
	11	12 01	16.00	1.008
		4 91	1 95	9 74
	Ratio (÷ by smallest)	4 91	1.95	9 74
	itatio (by sinanost)	1.95	1.95	1.95
		2.5	1	5
	x2	5	2	10
	$E.F. = C_5 H_{10} O_2$			
5.		Те	0	
	m	4.00	1.00	
	n	4.00	1.00	
		127.6	16.00	
	_ • / • · · ·	3.13 x 10 ⁻²	6.25×10^{-2}	
	Ratio (÷ by smallest)	3.13×10^{-2}	6.25×10^{-2}	
		3.13 x 10 ⁻²	3.13×10^{-2}	
		1	2	
	$E.F. = TeO_2$			

Answers %O = (52.0 + 13.0) = 35.0% 6. 0 Η С m in 100 g 52.0 35.0 13.0 52.0 35.0 13.0 n 12.01 16.00 1.008 4.33 2.19 12.9 Ratio (÷ by smallest) 4.33 2.19 12.9 2.19 2.19 2.19 2 1 6 $E.F. = C_2H_6O$ 7. Fe 0 7.83 11.2-7.83 = 3.37m 47.83 3.37 n 55.85 16.00 0.140 0.211 Ratio (÷ by smallest) 0.140 0.211 0.140 0.140 x 2 2 3 $E.F. = Fe_2O_3$ 8. Fe Cl Fe $C\ell$ m 44.0 100-44.0 = 5634.4 100-34.4 = 65.644.0 56.5 34.4 n 65.6 55.85 35.45 55.85 35.45 1.58 0.788 0.616 1.85 Ratio (÷ by smallest) 0.788 1.58 0.616 1.85 0.616 0.788 1.58 0.616 2 1 1 3 **E.F.** = $FeC\ell_2$ E.F.= $FeC\ell_3$ 9. Ο Ν m in 100 g 63.6 36.4 63.6 36.4 n 14.01 16.00 4.54 2.275 Ratio (÷ by smallest) 4.54 2.275 2.275 2.275 2 1 **E.F.** = N_2O 0 Ν 53.3 m in 100 g 46.7 53.3 46.7 n 14.01 16.00 3.33 3.33 Ratio (÷ by smallest) 3.33 3.33 3.33 3.33 1 3.33 E.F. = NO0 Ν 69.6 m in 100 g 30.4 69.6 30.4 n 14.01 16.00 2.17 4.35 Ratio (÷ by smallest) 2.17 4.35 2.17 2.17

2

			Answers	
10.	$n(Cl^{-}) = n(AgCl) = 9.47$ $m(Cl^{-}) = (6.61 \times 10^{-2}) \times 10^{-2}$	/ 143.35 = 6.61 x 1 35.45 = 2.34 g	0 ⁻² mol	
	m n	Ti 3.40 - 2.34 =1.06 1.06 47.88 2.21 - 10 ⁻²	Cl 2.34 2.34 35.45	
	Ratio (÷ by smallest)	$\begin{array}{c} 2.21 \times 10 \\ 2.21 \times 10^{-2} \\ 2.21 \times 10^{-2} \\ 1 \end{array}$	$\begin{array}{c} 6.61 \times 10 \\ 6.61 \times 10^{-2} \\ 2.21 \times 10^{-2} \end{array}$	
	E.F. = TIC ℓ_3	1	5	
11.	$\begin{split} n(CO_2) &= 0.660/12.01 = \\ n(H_2O) &= 0.270/18.016 = \\ n(C) &= n(CO_2) \\ m(CO_2) &= 12.01 \times 1.50 \times \\ n(H) &= 2(n(H_2O) \\ m(H_2) &= 2 \times 1.008 \times 1.50 \\ m(O) &= 0.290 - (0.180 + \\ n(O) &= 0.0798/16.00 = 4 \end{split}$	$1.50 \times 10^{-2} \text{ mol}$ = 1.50 x 10 ⁻² mol x 10 ⁻²) = 0.180 g 0 x 10 ⁻²) = 0.0302 g - 0.0302 = 0.0798 g .99 x 10 ⁻³ mol		
	n Ratio	C 1.50 x 10 ⁻² 1.50 x 10 ⁻² 4.99 x 10 ⁻³ 3	H 3.00 x 10 ⁻² 3.00 x 10 ⁻² 4.99 x 10 ⁻³ 6	O 4.99 x 10 ⁻³ 4.99 x 10 ⁻³ 4.99 x 10 ⁻³ 1
	$\mathbf{E.F.} = \mathbf{C_3H_6O}$	2	Ŭ	1
12.	$n(CO_2) = 0.403 / 12.01 =$ n(C) = n(CO2) $m(C) = 12.01 \times 9.16 \times 10^{-10}$	$= 9.16 \times 10^{-3} \text{ mol}$ $0^{-3}) = 0.110 \text{ g}$		

 $n(H_2O) = 0.165/18.016 = 9.16 \times 10^{-3} \text{ mol}$ $n(H) = 2 n(H_2O)$ $m(H) = 2 x 1.008 x (9.16 x 10^{-3}) = 0.0185 \text{ g}$ $C \qquad H$

	С	Н	Ο
n	9.16 x 10 ⁻³	1.83 x 10 ⁻²	9.16 x 10 ⁻³
Ratio	9.16 x 10 ⁻³	1.83 x 10 ⁻²	9.16 x 10 ⁻³
	9.16 x 10 ⁻³	4.99 x 10 ⁻³	4.99 x 10 ⁻³
	1	2	1

$\mathbf{E.F.}=\mathbf{CH_2O}$

13 $n(CO2) = 1.60$	$0 / 12.01 = 3.64 \times 10^{-2} \text{ mo}$	1		
$n(C) = n(CO_2)$				
m(C) = 12.01	$x (3.64 \times 10^{-2}) = 0.437 \text{ g}$			
$n(H_2O) = 0.770$	$0/18.016 = 4.274 \text{ x } 10^{-2} \text{ m}$	nol		
$n(H) = 2 n(H_2)$	D) =			
$m(H) = 2 \times 1.0$	$008 \text{ x} (4.274 \text{ x} 10^{-2}) = 0.08$	62 g		
n(N) = 0.1697/	$(14.01 = 1.21 \times 10^{-2})$	C		
m(O) = 1.279	-(0.437 + 0.0862 + 0.169)	(7) = 0.586		
n(O) = 0.586/1	$6.00 = 3.66 \text{ x } 10^{-2} \text{ mol}$,		
	С	Н	0	Ν
Ν	3.64 x 10 ⁻²	8.62 x 10 ⁻²	3.66 x 10 ⁻²	1.21 x 10 ⁻²
Ratio	3.64×10^{-2}	8.62 x 10 ⁻²	3.66×10^{-2}	1.21×10^{-2}
	1.21×10^{-2}	1.21×10^{-2}	1.21×10^{-2}	1.21×10^{-2}
	3	7	3	1

$$\mathbf{E.F.} = \mathbf{C}_3\mathbf{H}_7\mathbf{O}_3\mathbf{N}$$

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Set 35: Empirical formulas 2

 $n(CO_2) = 0.531/44.01 = 1.21 \times 10^{-2} \text{ mol} = n(CO_3^{2-})$ 1. $m(CO_3^{2}) = 60.01 \text{ x} (1.21 \text{ x} 10^{-2}) = 0.724 \text{ g}$ $n(H_2O) = 0.219 / 18.016 = 1.21 \times 10^{-2} \text{ mol} = n(OH^-)$ $m(OH^{-})=(1.21 \times 10^{-2}) \times 17.008 = 0.2067 \text{ g}$ m(Cu) = 2.088 - (0.724 + 0.2067) = 1.157 g $n(Cu) = 1.157 / 63.55 = 1.82 \times 10^{-2} \text{ mol}$

	Cu	CO_{3}^{2}	OH
n	1.82×10^{-2}	1.21 x 10 ⁻²	1.21 x 10 ⁻²
Ratio	1.82×10^{-2}	1.21 x 10 ⁻²	1.21 x 10 ⁻²
	1.21 x 10 ⁻²	1.21 x 10 ⁻²	1.21 x 10 ⁻²
	1.5	1	1
x 2	3 -	2	2
$E.F. = Cu_3(CO)$	$_{3})_{2}(OH)_{2}$		

$$\mathbf{E.F.} = \mathbf{Cu}_3(\mathbf{CO}_3)_2(\mathbf{OH})$$

2. (a)	С	Н
m in 100 g	85.7	14.3
n	85.7	143
	12.01	1.008
	= 7.14	= 14.18
Ratio	7.14	14.18
	7.14	7.14
	1	2

(b)
$$E.F. = CH_2$$

 $n = 1.18 \times 10^5 / 298 \times 8.315 = 5.00 \times 10^2 \text{ mol}$ $M = 2.80 / 5.00 \text{ x } 10^{-2} = 56.00 \text{ g mol}^{-1}$ MF = 56 / 14.00 = 4 $MF = C_4H_8$ (c)

%Pb = 2 93 / 3.41 x100 = 85.9 % 3. $n(AgCl) = 1.16 / (107.9 + 35.45) = 8.09 \times 10^{-3} \text{ mol} = n(C\ell^{-})$ $M(C\ell^{-}) = (8.09 \times 10^{-3}) \times 35.45 = 0.287 \text{ g}$ %Cl = 0287 / 2.93 x100 = 9.79 % %O = 100 - (85.9 + 9.79) = 4.31 %

	Pb	Cl	0
M in 100 g	859	9-79	431
Ν	85.9	9.79	4.31
	207.2	35.45	16.00
	=0.415 mol	= 0.276 mol	= 0.269
ratio	0.415	0.276	1
	0.269	0.269	
	=1.5	=1	1
$E.F. = Pb_3C\ell_2O_2$			

@**!**\$=

4.	(a) $n(CO_2) = 6.60 / 44.01 =$ $m(C) = 0.150 \times 12.01 =$ $n(H_2O) = 4.05 / 18.016$ $n(H) = 2 \times 0.224 = 0.45$ $m(H) = 0.450 \times 1.008 =$ m(O) = 3.45 - (1.80 + 0.000) n(O) = 1.197 / 16.0 = 0.000	= 0.150 mol= n(C) 1.80 g = 0.224 mol 50 mol = 0.453 g 0.453) = 1.197 g 0.0750 mol		
	n ratio	C 0.150 0.150 0.0750 2	H 0.450 0.450 0.0750 6	O 0.0750 0.0750 0.0750 1
	E.F. = C_2H_6O (b) $n = 0.950 \ge 98 / 373 \ge M = 1.38 / 3.00 \ge 10^{-2} \le EFM = MFM$ MF = C_2H_6O	$x = 8.315 = 3.00 \text{ x } 10^{-1}$ = 46.0 g mol ⁻¹	² mol	
	(c) CH ₃ CH ₂ OH			
5.	(a) $n(CoCO_3) = 0.849 / (5)$ $M(Co) = (7.14 \times 10^{-3}) \times (5)$ $M(Co) = (7.14 \times 10^{-3}) \times (7)$ M(C) = 3.43 / 44.01 = 7 $M(C) = (7.79 \times 10^{-3}) \times (7)$ $M(C) = (7.79 \times 10^{-3}) \times (7)$ M(C) = 100 - (34.5 + 28)	8.93 + 12.01 + 48.0 8.93 = 0.421 g 00 = 34.5 % $7.79 \times 10^{-3} \text{ mol}$ 12.01 = 0.936 g $00 \ 28.1 \%$ 1) = 37.4 %	$(0) = 7.14 \text{ x } 10^{-3} \text{ mol}$	l
	M in 100 g N	Co 34.5 34.5 58.93 = 0.585 mol	C 28.1 28.1 12.01 =2.34 mol	O 37.4 3.74 16.00 =2.34 mol
	Ratio	1	234 0.585	234 0.585
	$\mathbf{E.F.} = \mathbf{Co}\mathbf{C}_{4}\mathbf{O}_{4}$	1	4	4
	(b) EFM = $58.93 + (4 \times 12)$ Ratio = $341.9 / 120.93$ MF = $2 \times EF = Co_2C_8C$	$(.01) + (4 \times 16.00) =$ = 2 0_8	170.93	
6.	(a)	C	C	N
	n	2.64 44.01	1.62 x 2 18 016	0.236 x 10 ⁵ x 2 298 x 8 315
	n	$= 6.00 \times 10^{-2}$	=0.1798	$= 2.00 \times 10^{-2}$
	ratio	6.00×10^{-2} 2.00 x 10 ⁻²	0 1/98 2.00 x 10 ⁻²	I
	$E.F. = C_3H_9N$	3	9	

(b)	
n =	$0.254 \ge 95.5 / (19 + 273) \ge 8.315 = 9.99 \ge 10^{-3} \mod 10^{-3}$
M =	$0.5896 / 9.99 \ge 10^3 = 59.0 \text{ g}$
$EFM = (3 \times 12.01) + (9 \times 12.01)$	$(x \ 1.008) + 14.01 = 59.1 \text{ g}$
EFM = MFM	
$MF = C_3H_9N$	

7. (a)
C
C
C
$$\ell$$
H
1.189 x12.01
1.292 x35.45
0.662-(0.3245+0.3195)
44.01
143.35
= 0.3245 g
0.3195 g
0.0180
12.01
35.45
1.008
= 0.0271 mol
= 9.01 x 10⁻³ mol
= 0.0179 mol
0.0271
0.0179
9.01 x 10⁻³
= 3
= 1
= 2

 $\mathbf{E.F.} = \mathbf{C}_{3}\mathbf{H}_{2}\mathbf{C}\boldsymbol{\ell}$

8.

 $EFM = (3 \times 12.01) + (2 \times 1.008) + 35.45 = 73.5$ (b) Ratio = 147 / 73.5 = 2 $MF = 2 x EF = C_6H_4C\ell_2$ (c) Cl Cl Cl Cl С Η Ν 1.76 x12.01 0.420x2x1.008 0.0295x101.3x2x14.1 44.01 18.016 $(15 + 273) \times 8.315$ = 0.480g= 0.0470 g= 0.0350g% 0.480 x100 0.0470 0.0350

(m in 100 g) 0.232 0.620 0.620 = 77.4% = 7.58% = 15.1% 77.29 7.52 15.1 n 12.01 1.008 14.01 = 7.52= 6.44=1.08ratio 6.44 7.52 1 1.08 1.08 =6 =7 =1

$\mathbf{E.F.} = \mathbf{C_6H_7N}$

(b)
$$n = 100 \times 101.3 / (100 + 273) \times 8.315 = 3.27 \times 10^{-2} \text{mol}$$

 $\begin{array}{ll} M = & 3.04 \ / \ 3.27 x 10^{-2} = 93.1 g \ mol^{-1} \\ EFM = & (6 \ x \ 12.01) + (7 \ x \ 1.008) + 14.01 = 93 \\ MF = & EF \ = & C_6 H_7 N \end{array}$

Cl

9.	% $H_2O = (5 43 - 4 88) / 5.43 \times 100 = 10.4\%$ m(Ni) =0.578 / (58.69 + 32.06) x 58.69 = 0.374 g %Ni = 0.374 /((2.00/4.88) x 5.43) x 100 = 16.8% % $C_2O_4 = 50.5\%$ % $K = 100 - (10.4 + 16.8 + 50.5) = 22.3\%$					
		K	Ni	C.O.	H.O	
	0/_	N 22.3	16.8	C ₂ O ₄ 50 5	11_{2}	
	(m in 100g)	22.5	10.8	50.5	10.4	
	n	22.3	16.8	50.5	10.4	
		39.1	58.69	88.02	18.016	
		= 0.570mol	= 0.286mol	0.574mol	0.577mol	
	ratio	0.570	1	0.574	0.577	
		0.286		0.286	0.286	
		=2	=1	=2	=2	
	E.F. = $K_2Ni(C_2O_4)_2.2.J$	H ₂ O				
10.	$n(BaSO_4) = 0.4671 / (13)$ $m(S) = (2.00 \times 10^{-3}) \times 3$ $n(OH^{-}) = 0.250 \times 0.024$ $n(HC\ell) = n(NaOH) - 2n$ $n(C\ell^{-}) = n(OH^{-}) - 2n(S)$ $m(C\ell^{-}) = 35.45 \times (2.00)$	37.3 + 32.06 + 64.00 $32.06 = 6.42 \times 10^{-2} \text{ g}$ $x = 6.00 \times 10^{-3} \text{ mol}$ (H_2SO_4) $= 6.00 \times 10^{-3} - 4.00$ $\times 10^{-3} = 0.0709 \text{ g}$	$\begin{array}{l} \text{(b)} = 2.00 \text{ x } 10^{-3} \text{ mol} \\ \text{(c)} \\ $	= n(S)		
	$m(CC) = 55.45 \times (2.00)$	S S	Cl	0		
	%	$6.42 \times 10^{-2} \times 100$	0.0709 x100	100 -(23.7+26.2)		
	(m in 100 g)	0.2702	0.2702			
		= 23.7%	= 26.2 %	=50.1 %		
	n	23.7	26.2	50.1		
		32.06	35.45	16.00		

=0.739 mol

1

1

1

4

= 3.13/0.739 mol

3.13/0.739

4.25

4.25

17

= 0 739 mol

1

1

1

4

Set 36: Chemical Synthesis: no answers provided

ratio

x4

E.F. = $S_4C\ell_4O_{17}$