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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

$$
= 5 \times 10^{-23} \text{ atoms}
$$

6.

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 $\times 10^{1}$

Set 2: Errors

- 1. (a) first weighing = ± 2 mg ∴ 0.002/12.363 × 100 = 0.02%; final weighing ± 2 mg ∴ 0.002/0.834 × 100 = 0.2% (b) Total error ± 4 mg = ± 4 mg = 0.004/0.834 \times 100 = 0.5 %
- 2. $153/36671 \times 100 = 0.417 \%$
- 3. (a) 1 part in 3000 so $3000 \times 0.2 = 600$ mg (b) Min Mass = $(0.2 \times 100) / 0.01 = 2000$ 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 or 2.29 \times 10⁻² (the answers is expressed to the least number of significant figures, which is 3)
- 7. (a) 1.67×10^{2} °C or 167 °C (b) -2 $\times 10^{1}$ °C or -20 °C
- 8. (a) $13.6 \times 96.485 = 1.31 \times 10^3 \text{ kJ} \text{ mol}^{-1}$ (b) $24.6 \times 96.485 = 2.37 \times 10^3 \text{ kJ} \text{ mol}^{-1}$ (c) $9.32 \times 96.485 = 8.99 \times 10^2 \text{ kJ mol}^{-1}$
- 9. $350 / 1920 \times 100 = 18\%$
- 10. 0.24 or 2.4×10^{-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 $v = m/d$ ensity = 3.0 / 2.7 = 1.111 cm³ $v = 51 \times 3 \times$ 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 $c(NaHCO₃) = n/V = 0.0291 / 0.2500 = 0.1164$ mol L⁻¹ % error (assume no error for mass) = $(0.3 / 250.0) \times 100 = \pm 0.12$ % $c(NaHCO₃) = 0.1164$ mol $L^{-1} \pm 0.12$ %

absolute error / $0.11642 = 0.12 \%$ absolute error = $0.12/100 \times 0.11642 = 0.00014$ mol L⁻¹ $c(NaHCO₃) = 0.11642 \pm 0.00014$ mol L⁻¹

- 12. (a) Average titre = 22.53 mL (23.15 is outlier) Error due to glassware = $(2 \times 0.05) \times 3$ readings = 0.3 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 $0.06/20.00 \times 100 = 0.3\%$ $n(NaHCO3) = 0.0446 \times 0.02000$ = 8.92×10^{-4} mol (d) $n(H+) = n(NaHCO₃)$ $= 8.92 \times 10^{-4}$ 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 = $1.63/100 \times 3.96 \times 10^{-2}$ = 6.45×10^{-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 % 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 b

- 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 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 (+ΔH), 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 $T_1 < T_2$ 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

Set 6: Equilibrium systems

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Acids and bases

Set 8 Acids and bases

- 1. Conjugate acids: HF, HCO₃, HClO₄, HSO₃⁻
- 3. (a) $H_2SO_4(aq)$) + $H_2O(\ell)$ \rightarrow $H_3O^+(aq)$ + SO_4^{2-} (b) $H_2S(aq)$ + $H_2O(\ell)$ \Rightarrow $H_3O^+(aq)$ + HS
- , NH⁴⁺ 2. Conjugate bases: Br, SO_4^2 , HPO₄², S², NO₃, OH
	- (aq) (c) KOH(aq) \rightarrow K⁺(aq) + OH⁻(aq) (aq) (d) $N_2H4(aq) + H_2O(\ell) \rightleftharpoons H^+(aq) + HS^-(aq)$
- 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) (b) Concentrated solution of a weak acid (HY)

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

(c) Dilute solution of a strong acid (HX) (d) Dilute solution of a weak acid (HY)

Set 9: Acid and base strength

1. 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: $HCl(aq) \rightarrow H^+(aq) + Cl(aq)$ When the very soluble hydrogen fluoride dissolves in water there is little ionisation and it is classified as a weak acid: HF(aq) \Box H⁺(aq) + F⁻(aq)

- 2. (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) NH_3 , CO_3^2 ², CH_3NH_2 or any organic amine.
- 3. (a) Concentrated and weak (b) Dilute and strong (c) Concentrated and strong (d) Dilute and weak 4. (i) (a) $HBr(g) \rightarrow H^+(aq) + Br^ (b)$ ionisation (c) HBr, OH, Br, H_3O^+ , H_2O (ii) (a) $CH_3COOH(1) \rightleftharpoons H^+(aq) + CH_3COO^ (b)$ ionisation (c) OH, CH₃COO, H₃O⁺, CH₃COOH, H₂O (iii) (a) $H_2SO_4(\ell) \rightarrow H^+(aq) + HSO_4(aq)$ and $HSO_4(aq) \rightleftharpoons H^+(aq) + SO_4^{2}(aq)$ (b) ionisation (c) H_2SO_4 , OH , SO_4^2 , HSO_4 , H_3O^+ , H_2O (iv) (a) $NH_3(g) + H_2O(\ell) \rightleftharpoons NH_4^+(aq) + OH^ (b)$ ionisation (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 $HClO_4$ and the weaker base is ClO_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) H₃PO₄(ℓ) \rightleftharpoons H+(aq) + H₂PO₄ (aq) > H₂PO₄ (aq) \rightleftharpoons H⁺(aq) + HPO₄² (aq) > HPO₄² (aq) \rightleftharpoons H⁺(aq) + PO₄³ (aq) (b) H_2O , H_3PO_4 , H_3O^+ , $H_2PO_4^-$, $HPO_4^2^-$, $PO_4^3^-$, 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 ²(aq) + H₂O(ℓ) \rightleftharpoons HCO₃⁻(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). HCO₃ (aq) + H₂O(ℓ) \Rightarrow H₂CO₃(aq) + OH (aq) $(d_N) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$ $(e) CH₃COO(aq) + H₂O(l) \rightleftharpoons CH₃COOH(aq) + OH(aq)$

- 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)

 $^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$ calcium hydrogenphosphate $(\bigwedge_{p}H)$ 2 ⁻(aq) + H₂O(ℓ) \Rightarrow H2PO4⁻(aq) + OH⁻(aq) and H_2PO_4 ⁻(aq) + $H_2O(\ell) \rightleftharpoons H_3PO_4(aq) + OH$ ⁻(aq) potassium sulfate (Ψ pH) 2 (aq) + H₂O(ℓ) \Rightarrow HSO₄ (aq) + OH (aq) ammonium chloride $(\blacklozenge_{\text{pH}})$ $^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$ (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. $Ca(H_2PO_4)_2(s) \rightleftharpoons 2H_2PO_4(aq) + Ca^{2+}(aq)$ 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) → 2Na⁺(aq) + CO₃²(aq) and CO₃²(aq) + H₂O(ℓ) \rightleftharpoons HCO₃⁻(aq) + OH⁻(aq)
- 6. (a) $OH(aq) + CH_3COOH(aq) \rightleftharpoons H_2O(\ell) + CH_3COO$ (b) basic, $pH > 7$. $\text{(c) } CH_3COO \text{ (aq)} + H_2O \text{ (l)} \rightleftharpoons CH_3COOH \text{(aq)} + OH \text{ (aq)}$
- 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 1.8×10^{-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 × 10⁻¹⁴ / [H⁺] = 6.76 x 10⁻⁸ mol L⁻¹ (b) Acidic
- 3. $[H^+] = 1.00 \times 10^{-14} / [OH] = 6.33 \times 10^{-9} \text{ mol L}^{-1}$
- 4. (a) $[H^+] = 1.00 \times 10^{-14} / [OH] = 6.45 \times 10^{-11}$ mol L⁻¹ (b) $[OH] = 2 \times [Ba(OH)_2]$; $[H^+] = 1.00 \times 10^{-14} / [OH] = 1.00 \times 10^{-14} / 7.80 \times 10^{-2} = 1.28 \times 10^{-13}$ mol L⁻¹
- 5. (a) [OH⁻] = 1.00 × 10⁻¹⁴ / [H⁺] = 9.62 x 10⁻¹⁴ mol L⁻¹ (b) $H_2SO_4(\ell) \to H^+(aq) + HSO_4(aq)$ and $HSO_4(aq) \to 0.9HSO_4(aq) + 0.1H^+(aq) + 0.1SO_4^{2-}(aq)$ Overall $H_2SO_4(\ell) \to 0.9HSO_4(aq) + 1.H^+(aq) + 0.1SO_4^{2-}(aq)$ $[H^+] = 1.1$ $[H_2SO_4] = 1.1 \times 0.125 = 0.1375 \text{ mol } L^{-1}$ $[OH^+] = 7.27 \times 10^{-14} \text{ mol } L^{-1}$
- 6. (a) Ionisation of water is endothermic: heat + H₂O(ℓ) \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) $10^{-14}/[H^+] = [OH^-] = 10^{-14}/1.55 \times 10^{-5} = 6.45 \times 10^{-10}$ mol L⁻¹ OH⁻¹
	- (b) Some release of SO₂ from the smelter with elevated levels of $CO₂$ 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 $[H^+] = (1.6 \times 10^{-4} + 1.3 \times 10^{-6})/2 = 8.065 \times 10^{-5}$ mol L⁻¹ $[H^+] = 8.07 \times 10^{-5}$ mol L^{-1} As $10^{-14}/[H^+] = [OH^-] = 10^{-14}/8.07 \times 10^{-5} = 1.24 \times 10^{-10}$ mol L^{-1}

9. (a) Stomach Acid : n = c x v = 1.5 x 2.5 x 10^{-4} = 3.75 x 10^{-4} mol H⁺ $\underline{Al_2O_3}$: n= m/M = 5 x 10⁻³/(26.98+ 3(17.00)) = 6.412 x 10⁻⁵ mol Al(OH)₃ ∴ there are 1.93 x 10⁻⁴ mol OH . available $Mg(OH)_2$: n= m/M = 5 x 10⁻³/(24.31+ 2(17.00)) = 8.575 x 10⁻⁵ mol Mg(OH)₂ ∴ there are 1.715 x 10⁻⁴ mol OH available Total OH^{$=$} = 3.645 x 10⁻⁴ mol OH⁻ available The acid is INXS by 3.75 x 10^{-4} - 3.645 x 10^{-4} = 1.05 x 10^{-5} mol $c = n/v = 1.05 \times 10^{-5} / 1.5 = 7.00 \times 10^{-6}$ [OH] = 1.43 x 10⁻⁹ mol 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 1.00×10^{-7} mol L⁻¹.

10. (a) $V(Mg(OH)_2) = 4.86$ mL (b) $[H^+] = 8.34 \times 10^{-3}$ mol L^{-1} [OH⁻ $[OH^-] = 1.20 \times 10^{-12}$ mol 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) \Rightarrow H⁺(aq) + (Litmus) (aq) Red Blue

- 2. (a) H(Indicator)(aq) \Rightarrow H⁺(aq) + (indicator)(aq)
	- (b) Addition of an acid increases $[H^+]$, causes more $H(Indication)$ 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.

4.

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.

8. (a) Phenolphthalein (b) More acid will be used (c) Concentration measured lower than it actually is

Set 13: The pH scale

- 1. (a) $[H^+] = [HC\ell] = 0.100 \text{ mol } L^{-1}$ $] = [HC\ell] = 0.100 \text{ mol L}^{-1}$ [OH '] = 1.00 x 10⁻¹³ mol L⁻¹; pH = 1.00 (b) $[H^+] = [HNO_3] = 0.00500 \text{ mol } L^{-1}$ $] = [HNO₃] = 0.00500 \text{ mol L}^{-1}$ [OH] = 2.00 x 10⁻¹² mol L⁻¹; pH = 2.30
	- (c) $[OH-] = [NaOH] = 0.0100$ mol L^{-1} (d) $[H^+] = [HC\ell] = 2.00 \text{ mol L}^{-1}$ [OH-
		- $] = 1.00 \times 10^{-12}$ mol L^{-1} ; pH = 12.0 $] = 5.00 \times 10^{-15}$ mol L⁻¹; pH = 0.300
		- (e) $[H^+] = [OH -] = 1.00 \times 10^{-14}$ mol L⁻¹ and the pH = 7.00. Neutral as solutions of Na⁺ ions and C ℓ ions do not hydrolyse in water.

- 3. Concentration changed by a factor of 1000
- 4. Depleted Solution: $pH = 2$: $[H^+] = 10^{-2}$ 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 $5 = 10^{-5}$ mol $L^{-1} = 0.00001$ mol L^{-1} pH $3.6 = 10^{-3.6}$ mol $L^{-1} = 2.52 \times 10^{-4}$ mol L^{-1} $c_1v_1 = c_2v_2 = 10^{-5} \times X = 0.025 \times 2.52 \times 10^{-4} = 630$ 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 ∴ 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

- 7. (a) Drain : $n = c \times v = 0.236 \times 0.200 = 0.0472$ mol Runoff : $n = c \times v = 0.156 \times 0.300 = 0.0468$ 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(HC ℓ) = n(H⁺) / c = 0.094 / 1 = 0.094 L = 94 mL per 500 mL ∴ 188 mL per litre 8. pH 5.5 \therefore [H⁺] = 10^{-5.5} = 3.162 × 10⁻⁶ mol L⁻¹
- $n = c$ x $v = 3.162$ x $10^{-6} \times 15000 = 4.743 \times 10^{-2}$ mol H⁺ NaOH : $10g \div n = m/M = 10/39.99 = 0.25$ mol of OH Result : 0.2026 mol NaOH INXS ∴ c=n/v = 0.2026/15000 = 1.351 × 10⁻⁵ mol L⁻¹ OH⁻ ∴ the pOH = 4.87 ∴ pH = $14 - 4.87 = 9.13$
- 9. pH 7.20 \therefore [H⁺] = 10^{-7.2} = 1.584 \times 10⁻⁸ mol L⁻¹ H⁺ $n = c \times v = 1.584 \times 10^{-8} \times 2.0 \times 10^{6} = 0.03168$ mol H⁺ pH 7.8 ∴ [H⁺] = 10^{-7.8} = 6.3095 × 10⁻⁸ mol L⁻¹ H⁺ $n = c$ x v = 6.3095 x 10⁻⁸ x 2.0 x 10^{6 =} 0.12619 mol H⁺ Needs to increase mol H⁺ by 0.12619 - 0.03168 = 0.09451 mol $V(HC\ell) = n(H^+_{required}) / c = 0.09451/12.0 = 7.87$ mL
- 10. (a) pH 6.75 ∴ [H⁺] = $10^{-6.75}$ = 1.778 × 10^{-7} mol L⁻¹ H⁺ pH 5.1∴ [H⁺] = 10^{-5.1} = 7.943 × 10⁻⁶ 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 × 10⁻⁷ mol L⁻¹ H⁺ pH 8.00 ∴ pOH = 6 ∴ [OH-] = 10^{-6} = 1 × 10^{-6} mol L⁻¹ OH Mix 1 L of each OH⁻ INXS by : 1 x 10⁻⁶ - 1.68 \times 10⁻⁷ = 8.22 \times 10⁻⁷ mol OH⁻ $c = n/v = 8.22 \times 10^{-7}/2 = 4.11 \times 10^{-7}$ 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.
- 4. (a) Hypochlorite ions added to pool water as sodium hypochlorite solution or calcium hypochlorite solid to kill micro organisms. Weak acid hypochlorous acid forms: OC $\ell^-(aq) + H_2O(\ell) \rightleftharpoons HOC\ell(aq) + OH(aq)$ The buffer uses up some of these OH⁻ ions stopping the pool water becoming alkaline too quickly.
	- (b) $OH(aq) + HCO₃(aq) \rightleftharpoons H₂O(l) + CO₃²(aq)$ and $HCO₃(aq) \rightleftharpoons CO₃²(aq) + H⁺(aq)$: H⁺ reacts with the OH. Both processes use up OH ions.
- 5. (a) $H₂O$ and $HCO₃$ ion ion (b) $\text{CO}_2(\text{aq}) + \text{H}_2\text{O}(\ell) \rightleftharpoons \text{HCO}_3(\text{aq}) + \text{H}^+(\text{aq})$ H^+ ion is used up by reacting with the HCO₃ and OH ions is used up by reacting with CO₂ directly, OH (aq) + CO₂(aq) \rightleftharpoons HCO₃ (aq) or with H⁺ ions, results in more H⁺ ions. H⁺(aq) + OH (aq) \rightleftharpoons H₂O(ℓ)
- 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 1½ times number of moles of hydrochloric acid. This reacts with half the citrate ion: $C_6H_5O_7^{3.7}(aq) + 3H^+(aq) \rightleftharpoons H_3C_6H_5O_7(aq)$
- 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

- 7. CaO + 2HC $\ell \rightarrow$ CaC ℓ_2 + H₂O $n(HC\ell) = cV = (1.50 \times 0.250) = 0.375$ mol $n(CaO) = \frac{1}{2} n(HC\ell) = \frac{1}{2} (0.375) = 0.1875$ mol $m(CaO) = nM = 0.1875 \times (40.08 + 16.00) = 10.5 g$ $m(CaO) = 10.5 g$
- 8. 2NaHCO₃ + H₂SO₄ \rightarrow Na₂SO₄ + 2CO₂ + 2H₂O $m = 600 g$ m? V? $M (NaHCO₃) = 22.99 + 1.008 + 12.01 + 48 = 84.008$ $M(H_2SO_4) = 2(1.008)+32.07+4(16.00) = 98.086$
	- (a) $n(NaHCO₃) = m/M = 600 / 84.008 = 7.142$ mol $n(H_2SO_4) = \frac{1}{2} n(NaHCO_3) = 3.571$ 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)
	-

11. 25.2 mL of KOH

10. 7.34×10^{-2} mol L⁻¹

- 12. 57.5%
- 13. 192 g mol⁻¹
- 14. 0.954 g of Na₂CO₃

Set 16 : Acid-bases titrations 2

- 1. $[Acid] = 1.38 \times 10^{-5} \text{ mol L}^{-1}$ 2. $[Acid] = 2.53 \text{ mol L}^{-1}$
- 3. $[HCO₃]$
- 5. [lactic acid] = 9.97×10^{-6} mol L⁻¹ 6.
- 7. $[H_3C_6H_5O_7]_{ppm} = 31.4$ 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 $10. \, \text{V}$ ϕ Pb = 90.4%

- $]= 6.32 \times 10^{-6}$ mol L⁻¹ and L^{-1} 4. [OH-] = 1.59 x 10⁻⁶ mol L⁻¹
	- $pH = -log_{10}[H^+] = -log_{10}(11.2) = -1.05$

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

Oxidation and reduction

Set 18: Oxidation number

- (k) None (this is a precipitation reaction)
- (l) None (this is an acid/carbonate equation)
- **Set 19: Balancing half equations**

Set 20: Balancing redox equations

- 1. $Br_2 + 2I \rightarrow 2Br + I_2$
- 2. $Cu + 2Ag^{+} \rightarrow Cu^{2+} + 2Ag$
- 3. $Mg + Pb^{2+} \rightarrow Mg^{2+} + Pb$
- 4. $Mg + 2H^{+} \rightarrow Mg^{2+} + H_2$
- 5. $2A1 + 6H^{+} \rightarrow 2Al^{3+} + 3H_{2}$
6. $Mg + Cu^{2+} \rightarrow Mg^{2+} + Cu$
- $Mg + Cu^{2+} \rightarrow Mg^{2+} + Cu$
- 7. $2A1 + 3Zn^{2+} \rightarrow 2A1^{3+} + 3Zn$
- 8. $Cu + 2Fe^{3+} \rightarrow Cu^{2+} + 2Fe^{2+}$
- 9. $Zn + 2Ag^+ \rightarrow Zn^{2+} + 2Ag$
- 19. (a) $I_2 + 2e^- \rightarrow 2I^-$ (c) $S_2O_3^2 + 5H_2O + 4I_2 \rightarrow 2SO_4^2 + 10H^+ + 8I^-$
- 20. (a) $CH_3CH_2OH + H_2O \rightarrow CH_3COOH + 2H^+ + 2e^-$
- (c) $2CH_3CH_2OH+O_2 \rightarrow 2CH_3COOH$
-
- (c) O₂ is the oxidising agent, $C_6H_{12}O_6$ is the reducing agent
- 22. (a) Ox: $Mg \rightarrow Mg^{2+} + 2e^-$ Red: $Ti^{4+} + 4e^- \rightarrow Ti$ RedOx: $2Mg + TiCl_4 \rightarrow 2MgCl_2 + Ti^{4+}$
- (b) $TiC\ell_4$ is reduced, Mg metal is oxidised
- 23. (a) $3NO_2 + H_2O \rightarrow 2HNO_3 + NO$ (b) NO_2 is both (c) A disproportionation reaction

Set 21: Galvanic cells

1.

- 10. $C\ell_2 + 21 \rightarrow 2C\ell + I_2$.
- 11. $2Li + 2H_2O \rightarrow 2Li^+ + 2OH^- + H_2$
- 12. $Cu + 2NO_3 + 4H^+ \rightarrow Cu^{2+} + 2NO_2 + 2H_2O$
- 13. $Cu + SO_4^2 + 4H^+ \rightarrow Cu^{2+} + SO_2 + 2H_2O$
- 14. $2H_2O_2 \rightarrow 2H_2O + O_2$
15. $Cr_2O_7^{2-} + 8H^+ + 3C_2H_2$
- 15. $Cr_2O_7^{2+} + 8H^+ + 3C_2H_5OH \rightarrow 2Cr^{3+} + 3CH_3CHO + 7H_2O$
- 16. $Mg + 2H_2O \rightarrow Mg^{2+} + H_2 + 2OH^{-}$
- 17. $Cu_2O + 2H^+ \rightarrow Cu + Cu^{2+} + H_2O$
- 18. $4Au + 16CN^- + 3O_2 + 12H^+ \rightarrow 4[Au(CN)_4] + 6H_2O$

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

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

21. (a) $C_6H_{12}O_6 + 6O_2 \rightarrow 6CO_2 + 6H_2O$ (b) O_2 is reduced and C in the glucose is oxidised

 (i) (ii)

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

Set 22: Electrolytic Cells

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

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

 $\textcolor{red}{\Phi}\ \mathbb{G}$

Anode $\left|\begin{matrix} 1 \\ 2 \end{matrix}\right|$ Cathode

Set 23: Oxidation and Reduction: no answers provided

Set 25: Reactions of organic compounds

1. (a) $CH_3CH_2CH_3 + 5O_2 \rightarrow 3CO_2 + 4H_2O$

2. (a)
$$
CH_4 + C\ell_2 \rightarrow CH_3C\ell + HC\ell
$$

C H H H H + H Cl C H H H Cl + H Cl

chloromethane, dichloromethane, trichloromethane, tetrachloromethane (b) $CH_2=CH_2 + Br_2 \rightarrow BrCH_2CH_2Br$ 1,2-dibromomethane

Butan-2-ol

Propane

(e) $C_6H_{10} + HBr \rightarrow C_6H_{11}Br$

(d) $CH_3CHCH_2 + H_2 \rightarrow CH_3CH_2CH_3$

3. (a) ethene and water.

$$
\overset{\mathbf{H}}{\underset{\mathbf{H}}{\uparrow}}\overset{\mathbf{C}}{=}\overset{\mathbf{C}}{\underset{\mathbf{H}}{\uparrow}}\qquad \ \ \, +\qquad \ \ \, \mathbf{H}-\mathbf{O}-\mathbf{H}\qquad \ \ \, \longrightarrow \qquad \ \ \, \overset{\mathbf{H}}{\underset{\mathbf{H}}{\uparrow}}\overset{\mathbf{H}}{\unders
$$

(b) Step 1: methane, chlorine, ultraviolet light.

C H H $H - C - H$ **+** $Cl - Cl$ \longrightarrow $H - C$ **H H** $H - C$ -Cl + **H Cl**

butan-2-one

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- 7. (a) butan-2-ol, and acidified MnO₄ or acidified Cr₂O₇².
	- $5CH_3CHOHCH_2CH_3 + 2MnO_4 + 6H + \rightarrow 5 CH_3COCH_2CH_3 + 2Mn^{2+} + 8H_2O$ (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 ℓ finally $CH_2C\ell_2 + C\ell_2 \rightarrow CHC\ell_3$ + HC ℓ
	- (c) pentan-l-ol, $CH_3CH_2CH_2CH_2OH$, and acidified MnO₄ or acidified $Cr_2O_7^2$. $5CH_3CH_2CH_2CH_2OH + 4MnO_4 + 12H^+ \rightarrow 5 CH_3CH_2CH_2CH_2COOH + 4Mn^{2+} +11H_2O$
	- (d) propanoic acid, propan-l-ol, $CH_3CH_2CH_2OH$ and conc sulfuric acid. $CH_3CH_2COOH + CH_3CH_2CH_2OH \rightarrow CH_3CH_2COOCH_2CH_3 + H_2O$
	- (e) Cyclopentene and water.

- (f) hex-2-ene, $CH_3CH=CHCH_2CH_2CH_3$ and bromine. $CH_3CH=CHCH_2CH_2CH_3 + Br_2 \rightarrow CH_3CHBrCHBrCH_2CH_2CH_3$
- (g) propene, CH₃CH=CH₂ and hydrogen chloride: CH₃CH=CH₂ + HC $\ell \rightarrow$ CH₃CHC ℓ CH₃
- (h) pentan-l-ol, $CH_3CH_2CH_2CH_2OH$, 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, CH₃CH₂OH, methanoic acid and sulfuric acid: CH₃CH₂OH + HCOOH \rightarrow HCOOCH₂CH₃ + H₂O
- (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(CH₃CH=CH₂) = nM = 16.64 x 42.078 = 700.1 g = 0.700 kg
- 2. 2 CH₃OH + 2 Na \rightarrow 2 CH₃ONa + H₂
	- (a) $n(CH_3OH) = m/M = 250 / 32.04 = 7.803$ mol $m(CH_3ONa) = nM = 7.803 \times 54.024 = 42155 = 422 g$
	- (b) $n(H_2) = \frac{1}{2} n(CH_3OH)$ $V(H₂)$ 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 $(273+23.0) / 102.4 = V_2 = 93.8$ L
- 3. $C_3H_8 + 5O_2 \rightarrow 3CO_2 + 4H_2O$ $V(CO₂) = 3V(C₃H₈) = 3 \times 1.00 = 3.00$ L
- 4. CH₃(CH₂)₁₁C₆H₄SO₃H + NaOH \rightarrow 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 $m(CH_3(CH_2)_{11}C_6H_4SO_3Na) = 348.5 \times 196 = 68.3 \text{ kg}$ (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) $n[C_{10}H_8] = m/M = 50.0 / 128.2 = 0.3900$ mol $= n/v = 0.3900 / 0.250 = 1.56$ mol L⁻¹ (b) $C_1V_1 = C_2V_2 = 1.56 \times 0.25 = 2.80 \times V_2$ ∴ $V_2 = 0.1393$ L $V(H₂O)_{to remove} = V₁ - V₂ = 0.250 -0.1393 = 0.1106 = 0.111 L = 111 mL$
- 6. $2 (CH_3(CH_2)_16COONa(aq) + CaCl_2(aq) \rightarrow ((CH_3(CH_2)_16COO)_2)Ca(s) + 2 NaCl(aq)$ $n((CH₃(CH₂)₁₆COO)₂)Ca = m/M = 1.25 / 607.02 = 2.059 x 10⁻³ mol$ ∴ there was 2 x 2.059 x 10^{-3} mol of soap $M = n \times M = 2 \times 2.059 \times 10^{-3} \times 306.45 = 1.262$ g soap % = soap / mixture x $100 = 1.262 / 10.0$ x $100 = 12.6%$

- 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 2.462×10^{-7} mol HTa in 20 mL must [titratable acid] = $n/V = 2.462 \times 10^{-7} / 0.02 = 1.23 \times 10^{-5}$ mol L⁻¹
	- (b) Ionised to 2% means there is 1.23 x 10⁻⁵ x 0.02 = 2.46 x 10⁻³ mol H⁺ $pH = 2.609$
- 8. $\text{MnO}_4^- + 8\text{H}^+ + 5\text{e}^- \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O}$ $CH_3CH_2OH + H_2O \rightarrow CH_3COOH + 4H^+ + 4e^ 5 \text{ CH}_{3}CH_{2}OH + 4 \text{ MnO}_{4} + 12H^{+} \rightarrow 4 \text{ Mn}^{2+} + 5 \text{ CH}_{3}COOH + 11H_{2}O$ 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 $n(MnO₄)$ = c x V = 0.0216 x 0.023085 = 0.000498636 mol ∴ as MR = 4:5 there was $5/4$ x 0.000498636 mol alcohol in the 20 mL sample = 0.000623295 mol So in the whole 500 mL there was $25 \times 0.000623295 = 0.015582375$ mol in 5 mL wine So in 1L there was 200 x 0.015582375 = 3.116 mol L^{-1} = 3.12 mol L^{-1}
	- (b) $[CH_3CH_2OH]_{\text{wire in g per L}} = n \times M = 3.12 \times 46.06 = 144 \text{ g L}^{-1}$
- 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 $n(HCl) = c \times V = 0.0507 \times 0.024915 = 0.001263$ mol HCl ∴ there were 0.001263 mol of methanamine in the 20 mL sample ∴ in the diluted sample there were $12\frac{1}{2}$ times as much = 12.5 x 0.001263 = 0.0157899 mol methanamine in 25 mL original sample ∴ in 1L there was 40 x as much = 40 x 0.0157899 = 0.632 mol L⁻¹

- (b) $m = n x M = 0.632 x 31.0571 = 19.628 g/L$ $ρ = m/V$ so m = $ρ$ x V = 1.07 x 1000 = 1070 g/L % CH₃NH₂ = 19.628 / 1070 x 100 = 1.83 %
- 10. $CH_3CHOHCH_3 \rightarrow CH_3COCH_3 + 2H^+ + 2e^ MnO_4$ ⁻ + 8H⁺ + 5e⁻ → Mn²⁺ + 4H₂O $5CH_3CHOHCH_3 + 2MnO_4 + 6H^+ + \rightarrow 5CH_3COCH_3 + 2Mn^{2+} + 8H_2O$ 5.00 mL sample Diluted to 250 mL 20 mL sample taken $n(MnO₄)$ = c x V = 0.0203 x 0.02873 = 0.0005832 mol As MR is 2: 5 there were $5/2 \times 0.0005832$ mol propan-2-ol in the 20 mL sample $= 0.001458$ mol \therefore in the 250 mL solution there was $12\frac{1}{2}$ 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

1. (a) Let there be 100 g of substance

(b) PV = nRT so n = PV / RT = (105 x 1.18) / (8.314 x 298) = 0.0500 mol If 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 C_4H_8

- (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)

(c)

 $C_{0.150}H_{0.4498}O_{0.0747}$ divide though by $0.0747 = C_{2}H_{6}O_{1}$

- (b) PV = nRT so n = PV / RT = (98.0×0.950) / (8.314×373) = 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: CH_3CH_2OH and CH_3-O-CH_3
- (d) CH_3CH_2OH
- 3. (a)

CHO + O_2 → CO_2 + H_2O 0.682 g - 0.2642 g C -0.06649 g H ∴ 0.35131 g of O $n = m/M = 0.35131 / 16 =$ 0.021956 mol 0.968 g $n = m/M$ $=0.968/44 = 0.022$ mol ∴ 0.022 mol of C $m = n x M = 0.022 x$ $12.01 = 0.2642$ g of C 0.594 g $n = m/M$ =0.594/18.01 $= 0.03298$ mol H₂O ∴ 0.06596 mol of H $= 0.06649$ g of H Formula : $C_{0.022}H_{0.06596}O_{0.021956} = CH_3O$

(b) For 0.744 g sample $PV = nRT$ so $n = PV / RT = (95.0 \times 0.497) / (8.314 \times 473) = 0.0120062$ mol If 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.

H O C

H H C H H O H C H H C H O H O H

4.

CHNO + $O_2 \rightarrow CO_2$ + H₂O 1.180 g - 0.4804 g C -0.1007 g H -0.560268 g of N₂ ∴ 0.038632 g of O $n = m/M = 0.038632 / 16$ $= 0.0024145$ mol 1.758 g $n = m/M$ $=1.758/44 = 0.03995$ mol ∴ 0.0400 mol of C $m = n \times M = 0.0400$ $x 12.01 = 0.4804 g C$ 0.900 g $n = m/M$ =0.900/18.01 $= 0.04997$ mol H₂O ∴ 0.100 mol of H $= 0.1007$ g H

H

A second sample: 1.180 g $n = PV / RT = (105.0 \times 0.471) / (8.314 \times 298) = 0.0200$ mol nitrogen gas N₂ $m = n \times M = 0.0200 \times 28.013 = 0.560268$ g of N₂ $n(N_2) = m(N_2)/M(N_2) = 0.560268 / 28.02 = 0.02$ mol $n(N) = 2 n(N_2) = 0.04$ 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 C₂H₅NO 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, C_2H_5NO include. NH₂CH₂CHO CH₃CONH₂
- 5. (a)

CHN $+$ O₂ \rightarrow CO₂ $+$ H₂O 0.620 g -0.4804 g C -0.093 g of N ∴ 0.0466 g of H $n(H) = m/M = 0.0466 / 1.008$ $= 0.0462$ mol $n(N) = m/M = 0.093/14$ $= 0.00664$ mol 1.76 g $n = m/M$ $=1.76/44 = 0.04$ mol ∴ 0.0400 mol of C $m = n \times M = 0.0400 \times 12.01$ $= 0.4804$ g of C *2nd sample* 0.232 g n = PV / RT = (101.3 x 0.0295) / (8.314 x 288) = 0.001248 mol gas N₂ $m(N_2) = n \times M = 0.001248 \times 28.01 = 00.349 g$ ∴ % N_2 = 0.0349 / 0.232 x 100 = 15.04% ∴ in the original experiment there was $0.1504 \times 0.620 = 0.093$ g of nitrogen Empirical Formula $C_{0.0400}H_{0.0462}N_{0.00664} = C_6H_7N$ (b) 3^{rd} *sample – let* $V = IL$ $n = PV / RT = (101.3 \times 1) / (8.314 \times 373) = 0.03266$ $p = m/v$ so $m = p x V = 3.04 g$ ∴ 3.04 g = 0.03266 mol So 1 mol = 93.080 g $M_r(C_6H_7N) = 93.126$, So Molecular formula = Empirical Formula = C_6H_7N

-0.1612 g H ∴ 0.638 g of O $n = m/M = 0.638 / 16 =$ 0.03987 mol O $=3.52/44 = 0.08$ mol ∴ 0.08 mol of C $m = n x M = 0.08 x$ $12.01 = 0.9608$ g of C $=1.44/18.01$ $= 0.080$ mol H₂O ∴ 0.160 mol of H $= 0.1612$ g of H

Empirical formula = $C_{0.08}H_{0.160}O_{0.03987}$ = C_2H_4O

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

(c) (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.

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.

- 3. (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)
$$
O\underset{H \to N- C}{\underset{H \to N- C}{\underset{C}{\underset{C}{\underset{C}{\bigcirc}}}} O}
$$

6. (a) θ (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. (a) CH3COOH(") + H2O(") → CH3COO- (aq) + H3O+ (aq) (b) NH3(g) + H2O(") → NH⁴ (c) NaHCO3 (s) → HCO3 - (aq) + Na⁺ (aq) HCO3 (d) NaHSO4(s) → Na+ (aq) + HSO4 - (aq) HSO4 - (e) K2CO3 (s) -→ 2K⁺ (aq) + CO3 2- (aq) CO3 2- (f) NH4CH3COO(s) → NH4 ⁺ (aq) + CH3COO- (aq) NH4 CH3COO- (aq) + H2O(")⇌ CH3COOH(aq) + OH- (aq) 2. (a) i) H+ (aq) + OH- (aq) → H2O(") ii) Two colourless solutions are mixed together. No visible reaction; some heat evolved. (b) i) Ba(OH)2(s) + 2H⁺ (aq) → Ba2+ (aq) + 2H2O(") ii) A white solid dissolves in a colourless solution. (c) i) MgO(s) + 2H+ (aq) → Mg2+ (aq) + H2O(") ii) A white solid dissolves in a colourless solution. (d) i) CH3COOH(aq) + NH3 (aq) → NH4 ⁺ (aq) + CH3COO- (aq) (e) i) Zn(s) + 2H ⁺ (aq) → Zn2+ (aq) + H2 (g) ii) A silver solid dissolves in a colourless solution; colourless, odourless gas evolved (f) i) 2CH3COOH(aq) + Mg(s) → Mg2+ (aq) + 2CH3COO- (aq) + H2 (g) (g) i) Cu(s) + 4H⁺ (aq) + 2NO3 - (aq) → Cu2+ (aq) + 2NO2 (g) + 2H2O(") (h) i) Ni(s) + 2H⁺ (aq) → Ni2+ (aq) + H2 (g) (1) i) Fe(s) + 2H⁺ (aq) → Fe2+ (aq) + H2 (g) (aq) +I2 (s) b) i) Mg(s) + Fe2+ (aq) → Mg2+ (aq) + Fe(s) c) i) Cu(s) + 2Ag+ (aq) → Cu2+ (aq) + 2Ag(s) d) i) Zn(s) + Ni2+ (aq) → Zn2+ (aq) + Ni(s) e) i) 2Na(s) + 2H2O(") → 2Na+ (aq) + 2OH- (aq) + H2 (g) f) i) 2K(s) + 2HzO(")→ 2K⁺ (aq) + 2OH- (aq) + H2 (g) g) i) Cℓ² (g) + 2Br- (aq) → 2Cℓ- (aq) + Br2 (aq) ii) Two colourless solutions are mixed to form a white precipitate. b) i) Ag+ (aq) + Br- (aq) → AgBr(s) ii) Two colourless solutions are mixed to form a cream/white solid. c) i) Pb2+(aq) + 2I - (aq) → PbI2(s) ii) Two colourless solutions are mixed to form a yellow precipitate. d) i) Ca2+(aq) + SO4 2- (aq) → CaSO4(s) ii) Two colourless solutions are mixed to form a white precipitate e) i) Ba2+(aq) + OH- (aq) + SO4 2- (aq) + H+ (aq) → BaSO4(s) + H2O(") ii) Two colourless solutions are mixed to form a white precipitate. f) i) Fe2+(aq) + CO3 2- (aq) → FeCO3(s) g) i) 3 Zn2+(aq) + 2 PO4 3- (aq) → Zn3(PO4)2(s)

 $+$ HCO^{$-$} (aq)

 $-$ (aq) + H₂O(ℓ) \rightarrow CO₃² (aq) + H₃O⁺(aq) $(aq) + H_2O(\ell) \rightarrow SO_4^{2-}(aq) + H_3O^+(aq)$

- (aq) + H₂O(ℓ) \rightleftharpoons HCO₃ (aq) + OH (aq)
- $_+^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$

-
- n in the vinegar smell.
- ii) A silver solid dissolves in a colourless solution; colourless, odourless gas evolved.
-
- ras and a blue solution. as and green solution.
- ilved and a pale green soln.
- 3. (a) i) $Br_2 (\ell) + 2l^2 (aq) \rightarrow 2Br^2$
	- d dark brown solid forms
	- green soln to colourless.
	- ind brown solution turns blue.
	- tion fades to colourless.
	- $blourless, odourless gas.$
		- a colourless, odourless gas.
		- range solutions.
- 4. a) i) $\text{Ag}+(aq) + \text{C}\ell^-(aq) \rightarrow \text{AgC}\ell(s)$

ipitate. Green solution fades.

- ii) Two colourless solutions are mixed together to form a white precipitate.
- h) i) $Cu^{2+}(aq) + 2 \text{OH}(aq) \rightarrow Cu(OH)₂(s)$
	- ii) Blue solution mixed with a colourless solution to form a blue precipitate. Blue solution colour fades.
- i) 2 Cr³⁺(aq) + 3 CO₃²-(aq) \rightarrow Cr₂(CO₃)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) $M(Fe₂O₃) = 159.7 g mol⁻¹$ %Fe≔ 111 7 / 159.7 x100 = 69.9% 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: $2.08/2.71 \times 100 = 76.8\%$
- 3 a) $n(Br_2) = 125 / (79.9 \text{ x } 2) = 0.782 \text{ mol}$ $n(C_6H_6) = 60.0 / 78.108 = 0.768$ mol 1 mol of Br_2 requires 1 mol of C_6H_6 0.782 mol of Br₂ requires 0.782 mol of C_6H_6 $n(C_6H_6 \text{ required}) > n(C_6H_6 \text{ available})$ C_6H_6 is LR $n(C_6H_3Br) = n(C_6H_6) = 0.768$ mol $m(C_6H_5Br) = 0.768 \times 157 = 121 g$ b) % % yield= $93\frac{2}{121} \times 100 = 77.3$ %

4. a)
$$
Ca(OH)_2 \rightarrow CaO + H_2O
$$

b) $%CaO: 4\ 33/5.67 \times 100 = 76.4\%$

- 5. a) $Ba^{2+}(aq) + SO_4^{2-}(aq) \rightarrow BaSO_4(s)$ $n(Ba^{2+}) = n(CuSO_4.5H_2O)$ $n(CuSO_4.5H_2O) = 1.11/249.69 = 4.45 \times 10^{-3}$ mol = $n(BaC\ell_2)$ $m(BaC\ell_2) = (4.45 \times 10^{-3})\times (137.3 + 70.9) = 0.926$ g
	- c) $n(CuC\ell_2.2H_2O) = n(BaC\ell_2 \text{ used}) = 4.45 \times 10^{-3} \text{ mol}$ $m(CuC\ell_2.2H_2O) = (4.45 \times 10^{-3}) \times 170.482 = 0.758$ g %yield:- 0 345/0.758 x $100 = 45.5$ %
- 6. $n(Na_2S_2O_7) = \frac{1}{2} n(S)$ $n(S) = 17500/32.06 = 5.46 \times 10^2$ mol $n(Na_2S_2O_7) = \frac{1}{2} (5.46 \times 10^2) = 2.73 \times 10^2$ mol $m(Na_2S_2O_7) = (2.73 \times 10^2) \times (45.98 + 64.12 + 112) = 6.06 \times 10^4 g$ %yield:50000/60617 x 100 = 82.5 %
- 7. let $X = m(NaHCO₃)$ $Y = m(Na_2CO_3)$ $X + Y = 100$ $Y = 100 - X$

 $n(Na_2CO_3) = 90.7 / 106 = 0.856$ mol $n(Na_2CO_3 \text{ total}) = \frac{1}{2} n(NaHCO_3 \text{ initial}) + n(Na_2CO_3 \text{ initial})$ $=$ ($\frac{1}{2}$ x X/84.01) + Y/100 $0.856 = X/168 + (100-x)/106$ $62X = 1556.352$ $X=25.1 g$ $Y = 74.9 g$ $\%Na_2CO_3 = 74.9 / 100 \times 100 = 74.9 \%$

8.

Step 5

Reduction: Cr_2O_7^2 + 14H⁺ + 6e⁻ \longrightarrow 2Cr³⁺ + 7H₂O Oxidation: $Fe^{2+} \longrightarrow Fe^{3+} + e^{-}$ Combined : Cr₂O₇² + 14H⁺ + 6e⁻ + 6Fe²⁺ - 2Cr³⁺ + 7H₂O + 6Fe³⁺ + 6e⁻ Mol Ratio $Cr_2O_7^{2}$: Fe = 1:6 In 20mL sample $n (Cr_2O_7^2)$ = c xv = 0.0181 x 0.0209 = 0.0003783 mol ∴ n(Fe²⁺) = 6 x 0.00037829 = 0.002269 mol Fe²⁺ $m(Fe) = 0.002269 \text{ x } 55.85 = 0.1267 \text{ g}$ ∴ In 250 mL there was 12.5 x 0.1267 = 1.584 g Fe

∴ percentage purity of the wire = $1.5844/1.63 \times 100 = 97.2\%$ pure

9.

1 $\vert 2 \vert$ 3 1.27 g mineral Converted to $K_2Cr_2O_7$ Titrated with 37.5 mL of 0.400 M FeSO₄ Step 3 Reduction: Cr_2O_7^2 + 14H⁺ + 6e⁻ \longrightarrow 2Cr³⁺ + 7H₂O Oxidation: $\mathsf{Fe}^{2+} \longrightarrow \mathsf{Fe}^{3+} + \mathsf{e}^{-}$ Combined : Cr₂O₇² + 14H⁺ + 6e⁻ + 6Fe \longrightarrow 2Cr³⁺ + 7H₂O + 6Fe²⁺ + 6e⁻ Mol Ratio $Cr_2O_7^{2}$: Fe = 1:6 $n (Fe^{2+})$ - = c xv = 0.0375 x 0.400 = 0.015 mol Fe²⁺ ∴ n(Cr₂O₇²) = 1/6 x 0.015 = 0.0025 mol ∴ n(Cr) = 2 x 0.0025 = 0.005 mol ∴ m(Cr) = 52.00 x 0.005 = 0.26 g

∴ percentage purity of mineral = $0.26/1.27 \times 100 = 20.5 \%$ pure

10.

1 2 $\sqrt{0.752 \text{ g}}$ impure Na₂SO₃ $\sqrt{\frac{\text{T}}{\text{I}} \cdot \text{mL}}$ of 0.0993 M K₂Cr₂O₇ Reduction: Cr_2O_7^2 + 14H⁺ + 6e⁻ \longrightarrow 2Cr³⁺ + 7H₂O Oxidation: $SO_3^2 \longrightarrow SO_4^2 + 2e^-$ Combined : Cr_2O_7^2 + 14H⁺ + 6e⁻ + 3SO₃² \longrightarrow 2Cr³⁺ + 7H₂O + 3SO₄²⁺ + 6e⁻ Mol Ratio Cr₂O₇²⁻: SO₃²⁻ = 1:3 $n (Cr_2O_7^2)$ - = c xv = 0.0172 x 0.0993 = 0.0017079 mol ∴ n(SO₃²) = 3 x 0.0017079 = 0.005123 mol ∴ m(Na₂SO₃) = 126.043 x 0.005123 = 0.6458 g

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

11.

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

 $n \left(\text{MnO}_4 \right)$ = 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

∴ m(Fe₂O₃) = 0.0018655 x 159.688 = 0.297898 g

In whole sample

∴ m(Fe₂O₃) = 10 x 0.297898 = 2.97898 g

∴ percentage purity = 2.97898 / 3.08 x 100 = 96.7 % pure

12.

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 mol / L⁻¹ MnO₄ Reduction: $MnO_4^- + 8H^+ + 5e^- \longrightarrow Mn^{2+} + 4H_2O$ Oxidation: $\mathsf{Fe}^{2+} \longrightarrow \mathsf{Fe}^{3+} + \mathsf{e}^{-}$ 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 $n \left(\text{MnO}_4 \right)$ = c xv = 0.03986 x 0.0150 = 0.0005979 mol ∴ n(Fe²⁺) = 5 x 0.0005979 = 0.0029895 mol Titration 2 $n \left(\text{MnO}_4 \right)$ = c xv = 0.03986 x 0.0251 = 0.001000 mol ∴ n(Fe²⁺) = 5 x 0.001000 = 0.005002 mol ∴ n(Fe²⁺) =n(FeO)= 0.002990 mol $n(Fe^{3+}) = 2 \times n(Fe_2O_3) = 0.005002 - 0.002990 = 0.002013$ ∴ n(Fe₂O₃) = ½ x 0.002013 = 0.001006 For the half sample ∴ m(FeO) = 0.002990 x 71.844 = 0.2148 g ∴ m(Fe₂O₃) = 0.001006 x 159.688 = 0.1607 g For the whole sample ∴ m(FeO) = 0.2148 x 2 = 00.4296 g ∴ m(Fe₂O₃) = 0.1607 x 2 = 0.3214 g

Set 32: Limiting reagents

- 1. 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) $n(PbI_2) = \frac{1}{2} n(Kl)$ $=$ $\frac{1}{2}$ x 0.400 = 0.200 mol $n(PbI₂) = 0.200 \times 461.0 = 92.2 g$

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

 $n(HC\ell \text{ required}) > n(HC\ell \text{ available})$ HCℓ is LR

- b) $N(NaC\ell) = n(HC\ell) = 0.134$ mol m(NaC ℓ) = 0.134 x 58.44 = 7.82 g
- 3. a) n(CH₃COOH)= $4.78/60.052 = 7.96 \text{ X } 10^{-2} \text{ mol}$ $N(CaCO₃) = 2.22/100.08 = 2.22 \times 10^{-2}$ 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(CH3COOH requ) < n(CH3COOH avail) ∴ CaCO₃ is LR
	- b) $n(CO_2) = n(CaCO_3) = 2.22 \times 10^{-2}$ mol $m(CO₂) = (2.22 \times 10^{-2}) \times 44.01 = 0.976$ g
	- c) $n(Ca(CH_3COO_2)) = n(CaCO_3) = 2.22 \times 10^{-2}$ mol $m(Ca(CH_3COO)_2) = (2.22 \times 10^{-2}) \times 158.148 = 3.51 g$
- 4. a) $n(Mg)=6.08/24.3=0.250 \text{ mol}$ $n(H₂SO₄) = 20.0/98.076 = 0.2039$ mol 1 mol of Mg requires 1 mol of $H₂SO₄$ 0.250 mol of Mg requires 0.250 mol of H_2SO_4 $n(H_2SO_4 \text{ req}) > n(H_2SO_4 \text{ avail})$ ∴ H_2SO_4 is LR
	- b) $n(H_2) = n(H_2SO_4) = 0.2039$ mol $m(H_2) = 0.2039 \times 2.016 = 0.411 g$
	- c) $n(MgSO_4.7H_2O) = n(H_2SO_4) = 0.2039$ mol $m(MgSO₄.7H₂O) = 0.2039$ x 246.472 = 50.3 g
- 5. a) $n(NaOH) = 1.600/3998 = 4.00 \times 10^{-2}$ mol $n(H_2SO_4) = 1.472/98.076 = 1.5 \times 10^{-2}$ mol 1 mol of H2SO4 requires 2 mol of NaOH 1.5 x 10⁻² mol of H₂SO₄ requires 2 x (1.50 x 10⁻²) = 3.00 x 10⁻² mol n(NaOH req) < n(NaOH avail) ∴ H_2SO_4 is LR
	- b) $n(Na_2SO_4) = n(H_2SO_4) = 1.50 \times 10^{-2}$ mol $m(Na_2SO_4) = (1.50 \times 10^{-2}) \times 142.04 = 2.13$ g c) n(NaOH rem) = 4.00 x 10^{-2} - 3.00 x 10^{-2} = 1.00 x 10^{-2} mol
	- $m(NaOH) = (1.00 \times 10^{-2}) \times 39.998 = 0.400 g$
- 6. 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 $4/3 \times 0.151 = 0.201$ mol $n(HNO₃ \text{req})$ < $n(HNO₃)$ avail) ∴ Ag is LR
	- b) $n(NO) = 1/3 n(Ag) = 1/3x0.151 = 0.0503$ mol $m(NO) = 0.0503 \times 30.01 = 1.51 g$
	- c) $n(HNO_3rem) = 0.292 0.201 = 0.091$ mol m(HNCh) = $0.091 \times 63.018 = 5.75 \text{ g}$
- 7. a) $n(KO_2) = 5.00/71.1 = 7.03 \text{ X } 10^{-2} \text{ mol}$ $n(CO₂) = 9.00/44.01 = 0.204$ mol 1 mol of $CO₂$ requires 2 mol of $KO₂$ 0.204 mol requires 2 x 0.204 = 0.408 mol of $KO₂$ $n(KO₂ required) > n(KO₂ avail)$ ∴ KO₂> is LR

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

- b) $n(O_2) = 3/2 n(KO_2) = 3/2 x (7.03 x 10^{-2}) = 0.105$ mol $m(O_2) = 0.105 \times 32.00 = 3.36 g$ $n(CO_2$ rem) = 0.204 – $\frac{1}{2}$ (7.02 x 10⁻²) = 0.269 mol $m(CO₂) = 0.269$ x 44.01 = 7.43 g
- 8. $n(Ca_3(PO_4)_2) = 25.0 \times 10^6 / 310.18 = 8.06 \times 10^4$ mol $N(H_3PO_4) = 30.0 \times 10^6 / 97.94 = 3.06 \times 10^5$ mol 1 mol of $Ca_3(PO_4)_2$ requires 4 mol of H_3PO_4 8.06 x 10⁴ mol of Ca₃(PO₄)₂ requires 4 x (8.06 x 10⁴) = 3.224 x 10⁵ mol of H₃PO₄ $n(H_3PO_4 \text{ req}) > n(H_3PO_4 \text{ avail})$ ∴ H_3PO_4 is LR $n(Ca(H_2PO_4)_2) = \frac{3}{4} (n(H_3PO_4 \text{ avail}) = \frac{3}{4} (3.06 \times 10^5) = 2.296 \times 10^5 \text{ mol}$ $m(Ca(H_2PO_4)_2) = (2.296 \times 10^5) \times 234.052 = 5.37 \times 10^7 g (53.7 \text{ tonne})$
- 9. $n(CO2) = 2.94/44.01 = 6.68 \times 10^{-2}$ mol $n(Na_2CO_3) = n(CO_2) = 6.68 \times 10^{-2}$ mol $m(Na_2CO_3) = (6.68 \times 10^{-2}) \times 105.99 = 7.08 \text{ g}$ $\%Na_2CO_3$ = 7.08/7.20 x 00 = 98.3%
- 10. $n(C\ell_2) = 2.84/70.9 = 4.01 \times 10^{-2}$ mol mol n(MnO₂) = n(C ℓ_2) = 4.01 x 10⁻² mol m(MnO2) = $(4.01 \times 10^{-2}) \times 86.94 = 2.49$ g $\%MnO_2 = 3.49/3.52 \times 100 = 99.0\%$

Set 33: Calculations involving gases

- 4. P V n R T 105 0.0255 8.314 298.15 $n= 0.001080151$ mol of $CO₂$ ∴ there were 0.001080151 mol of CaCO₃ $= 100 \times 0.001080151 = 0.108$ g CaCO₃
- 5. $2NH_3(9) + H_2SO_4$ (aq) $\longrightarrow (NH_4)_2SO_4$
	- $nNH_3 = 1.50 / 22.71 = 0.06605$ mol
	- nH_2SO_4 = c x V = 2.54 x 0.050 = 0.127 mol
	- (a) To use all the NH₃ we need 1/2 that number of moles of $H_2SO_4 = 1/2 \times 0.06605 = 0.03303$ mol acid needed and have $0.127 \text{ mol} - \text{so } 20 \text{ gal}$ is in excess and the LR is NH₃
	- (b) $n(NH_4)_2SO_4$ produced = $\frac{1}{2}$ n(NH₃ used) = $\frac{1}{2}$ (0.06605) = 0.03303 moles:
	- mass ammonium sulfate = M x n = 132.094 x 0.03303 = 4.36 g
	- (c) nH₂SO₄ 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
	- $nH C \ell = c \times V = 2.50 \times 0.050 = 0.125$ mol
	- (a) ∴ NaHCO₃ is the LR
	- (b) $n(CO_2) = n(NaHCO_3): V(CO_2) = nRT/P = 0.085$ L
- 7. $Cu + 4HNO₃ \longrightarrow Cu(NO₃)₂ + 2NO₂ + 2H₂O$
	- $nCu = 1.33 / 63.55 = 0.0208$ mol
	- $nHNO₃ = c x V = 6.00 x 0.025 = 0.15$ mol
	- (a) To use all the Cu we need 0.0208×4 mol HNO₃= 0.0832 mol We have more so the Cu is the LR
	- (b) \therefore nNO₂ = 2 x 0.0208 = 0.0416 mol $V = 1.02$ L
- 8. $\text{MnO}_2 + 4H^+ + 2Cl^- \longrightarrow \text{Mn}^{2+} + Cl_2 + 2H_2O$

Or: MnO_2 + 4HC ℓ \rightarrow $MnC\ell_2$ + $C\ell_2$ + 2H₂O $nMnO₂ = 3.44 / 86.93 = 0.0396$ mol $nHCl = 6.20 \times 0.150 = 0.093$ mol nHCℓ needed = nH^+ used = 4 x nMnO₂ = 0.158 mol You need more HCℓ than you have so HCℓ is the limiting reagent ∴ nC ℓ_2 = ¼ HC ℓ = ¼(0.093) = 0.02325 mol $P = nRT/V = 238.26 kPa = 238 kPa$

- 9. $(\text{NH}_4)_2\text{SO}_4 + 2\text{KNO}_3 \longrightarrow \text{K}_2\text{SO}_4 + 4\text{H}_2\text{O} + 2\text{N}_2$ $n(NH_4)_2SO_4 = 30.0/132.140 = 0.227031$ mol $nKNO_3 = 34.0/101.1032 = 0.33629$ mol To use all the $(NH_4)_2SO_4$ we would need 2 x 0.22703 mol KNO₃ = 0.45406 mol
	- (a) $KNO₃$ is the LR and is used up.
	- (b) $n(NH4)2SO4$ used = $1/2$ $nKNO3 = 0.16814$ mol: $n(NH4)2SO4$ left over = $0.22701 - 0.16814 = 0.05887$ mol is INXS $m = n \times M = 0.058886 \times 132.140 = 7.78 g$
	- (c) $nN_2 = nKNO_3$: $P = nRT/V = 1814 kPa$

Set 34: Empirical formulas 1

Answers 6. $\%$ O = (52.0 + 13.0) = 35.0% C O H m in 100 g 52.0 35.0 13.0 n 52.0 35.0 13.0 12.01 16.00 1.008
4.33 2.19 12.9 4.33 2.19 12.9 Ratio (÷ by smallest) 4.33 2.19 12.9
2.19 2.19 2.19 2.19 2.19 2.19 2 1 6 $E.F. = C₂H₆O$ 7. Fe $\qquad \qquad$ \qquad O m 7.83 $11.2-7.83 = 3.37$ n 47.83 3.37 55.85 16.00 0.140 0.211 Ratio (\div by smallest) 0.140 0.211 0.140 0.140 $\begin{array}{ccc} \text{x 2} & \text{2} & \text{3} \end{array}$ $E.F. = Fe₂O₃$ 8. Fe Cl Fe Cl Fe Cl m 44.0 100-44.0 = 56 34.4 100-34.4 = 65.6 n 44.0 56.5 34.4 65.6 55.85 35.45 55.85 35.45 0.788 1.58 0.616 1.85 Ratio (\div by smallest) 0.788 1.58 0.616 1.85 0.788 1.58 0.616 0.616 1 2 1 3 $E.F. = FeCl₂$ **E.F.=** $FeCl₃$ 9. N O m in 100 g 63.6 36.4 n 63.6 36.4 14.01 16.00 4.54 2.275 Ratio (\div by smallest) 4.54 2.275 2.275 2.275 2 1 $E.F. = N₂O$ N O m in 100 g 46.7 53.3 n 46.7 53.3 14.01 16.00 3.33 3.33 Ratio (\div by smallest) 3.33 3.33 3.33 3.33 1 3.33 **E.F. = NO** N O m in 100 g 30.4 69.6 n 30.4 69.6 14.01 16.00 2.17 4.35 Ratio (\div by smallest) 2.17 4.35

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2.17 2.17 1 2

 $E.F. = C_3H_7O_3N$

 $n(H) = 2 n(H₂O) =$

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 $m(H) = 2 \times 1.008 \times (4.274 \times 10^{-2}) = 0.0862$ g

 $m(O) = 1.279 - (0.437 + 0.0862 + 0.1697) = 0.586$

C H O **N**

3 7 3 1

N

Ratio 3.64×10^{-2} 8.62×10^{-2} 3.66×10^{-2} 1.21×10^{-2}
 3.66×10^{-2} 1.21×10^{-2} Ratio 3.64 x 10^{-2} 8.62 x 10^{-2} 3.66 x 10^{-2} 1.21 x 10^{-2} 1.21 x 10^{-2} 1.21 x 10^{-2} 1.21 x 10^{-2} 1.21×10^{-2}

 $n(N) = 0.1697/14.01 = 1.21 \times 10^{-2}$

 $n(O) = 0.586/16.00 = 3.66 \times 10^{-2}$ mol

Set 35: Empirical formulas 2

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

$$
\mathbf{E}.\mathbf{F} = \mathbf{C} \mathbf{u}_3 (\mathbf{C} \mathbf{O}_3)_2 (\mathbf{O} \mathbf{H})_2
$$

 $E.F. = CH₂$ (b)

H

H H

H

 $n = 1.18 \times 10^5 / 298 \times 8.315 = 5.00 \times 10^2$ 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) **C H C H C H C H C H C H C H** \mathbf{H} \mathbf{C} **H H**

3. $\%Pb = 293 / 3.41 \times 100 = 85.9 \%$ $n(AgCl) = 1.16 / (107.9 + 35.45) = 8.09 \times 10^{-3}$ mol = $n(C\ell)$ $M(C\ell) = (8.09 \times 10^{-3}) \times 35.45 = 0.287 g$ $\%CI = 0287 / 2.93 \times 100 = 9.79 \%$ $\%$ O = 100 - (85.9 + 9.79) = 4.31 %

H

H

7. (a) C Cℓ 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 g n 0.3245 0.3195 0.0180 12.01 35.45 1.008 = 0.0271 mol = 9.01 x 10-3 mol =0.0179 mol Ratio 0.0271 1 0.0179 9.01 x 10-3 9.01 x 10-3 = 3 =1 = 2

 $E.F. = C₃H₂Cl$

(b) EFM = $(3 \times 12.01) + (2 \times 1.008) + 35.45 = 73.5$ Ratio = $147 / 73.5 = 2$ $MF = 2$ x $EF = C_6H_4C\ell_2$ (c) Cl Cl Cl Cl

$E.F. = C_6H_7N$

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

 $M = 3.04 / 3.27x10^{-2} = 93.1g \text{ mol}^{-1}$ EFM = $(6 \times 12.01) + (7 \times 1.008) + 14.01 = 93$ $MF = EF = C_6H_7N$

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

Set 36: Chemical Synthesis: no answers provided