# NSW INDEPENDENT TRIAL EXAMS – 2002 CHEMISTRY - SUGGESTED ANSWERS

#### **SECTION I - PART A**

1	A	2	В	3	Α	4	C	5	D	6	С	7	В	8	В
9	D	10	Α	11	С	12	D	13	C	14	С	15	В		

#### SECTION I - PART B

- 16. (a) The ethanol molecule is polar (1 mk) and has strong dipole-dipole attractions with other molecules, acting as a solvent for a range of polar substances (1 mk) (Hydrogen bonding as an alternative) (1 mk)
  - (b) Any two of: (2 mk)

temperature in range 30-40°C dilute aqueous sugar solution yeast to provide enzymes anaerobic environment

(c) As fermentation proceeds the reaction mixture decreases in mass due to the loss of carbon dioxide. (1

carbon dioxide. (1 mk)  $C_6H_{12}O_6 \rightarrow 2C_2H_6O + 2CO_{2(g)}$  (1 mk)

17. Advantage: any one of:

(1 mk)

water soluble (spills clean up and fire safety) easily ignited

renewable

Disadvantage: any one of:

(1 mk)

lower energy density

flash point below ambient temperature

- 18. (a)  $V^{2+} \rightarrow V^{3+} + e^{-}$  (1mk)
  - (b)  $VO_2^+ + 2H^+ + V^{2+} \rightarrow VO^{2+} + H_2O + V^{3+}$  (1 mk)
  - (c) 1.26 V (1 mk)
  - (d) The cell can be recharged. It has a long life. It is highly efficient. All reactions occur in solution, with inert electrodes. It is relatively inexpensive. It can be used for electric vehicles and thus reduce pollution from petrol driven cars. (2 marks for any one reason and relating it to society/environment) (2 mks)
- 19. Addition polymers are formed when unsaturated monomer units join without loss of atoms.

Chlomothers has the street CVI CVICI

Chloroethene has the structure CH<sub>2</sub>=CHCl (1 mk)

During polymerisation the double bonds open out, providing electrons for bonding to other monomer molecules. (1 mk)

- 20. (a)  ${}_{6}C^{14} \rightarrow {}_{7}N^{14} + \beta$  (1 mk)
  - (b) Geiger-Muller tube, scintillation counter, photo film or cloud/bubble chamber (1mk)
  - (c) Any valid example eg: Iodine-121 (1 mk) to treat hyperactive thyroid by destroying some cells, a less invasive treatment than surgery (1 mk) (2 mks)

```
21 Moles HCl (limiting reactant) = 1.5/24.47 = 0.0613 mol
                                                                                                     (1 \text{ mk})
    Mass NH<sub>4</sub>Cl formed = 0.0613 \times mol. mass = 0.0613 \times 53.5 = 3.28 g
                                                                                                     (1 \text{ mk})
22. eg: sodium hydrogen carbonate (or the hydrogen carbonate ion)
                                                                                                     (1 \text{ mk})
    This species acts as a proton donor and acceptor, shown by:-
         HCO_3 + H_3O^+ \rightarrow H_2CO_3 + H_2O \text{ (or } CO_2 + 2H_2O)
                                                                                                     (1 \text{ mk})
         HCO_3 + OH \rightarrow CO_3^2 + H_2O
                                                                                                      (1 \text{ mk})
23. (a) (i) methyl propanoate
                                                                                                      (1 \text{ mk})
          (ii) CH<sub>3</sub>CH<sub>2</sub>COOCH<sub>3</sub> or expanded formula
                                                                                                      (1 \text{ mk})
          (iii) any strong acid, phosphorus pentoxide etc
                                                                                                      (1 \text{ mk})
          (iv) Heating under reflux increases the reaction rate (higher temperature)
                                                                                                      (1 \text{ mk})
          while preventing loss of reactants or products by vaporisation to outside.
                                                                                                      (1 \text{ mk})
     (b) The ester has low polarity resulting in much weaker intermolecular forces
                                                                                                      (1 \text{ mk})
          than in pentanol and butanoic acid which both have polar OH groups.
                                                                                                      (1 \text{ mk})
          With an additional O atom butanoic acid is still more polar
                                                                                                          OR
          Pentanol and butanoic acid also form hydrogen bonds.
          The boiling points reflect the strengths of these intermolecular forces.
                                                                                                      (1 \text{ mk})
24. (a) Opening the bottle causes a decrease in pressure causing the equilibrium to shift to the
          side of greatest number of gaseous molecules, therefore formation of more gaseous
          CO<sub>2</sub>
                                                                                                      (2 \text{ mk})
     (b) As lowering the temperature favours the formation of aqueous CO<sub>2</sub>, this reaction is the
          reaction that produces heat, therefore the forward reaction as written is exothermic.
25. (a) H_2SO_4 + 2KOH \rightarrow K_2SO_4 + 2H_2O
                                                                                                      (1 \text{ mk})
     (b) no. of moles of KOH = 2 \times 10^{-5} x no. of moles of H_2SO_4
          25.0 \times [KOH] = 2 \times 0.200 \times 0.0200
                                                                                                      (1 \text{ mk})
          [KOH] = 0.320 \text{ mol } L^{-1}
     (c) Mass KOH = molarity x volume x mol. mass
                          = 0.32 \times 5 \times 56.1
                                                                                                    (1 \text{ mk})
                          = 89.8g
26. (a) N_{2(g)} + 3H_{2(g)} \rightarrow 2NH_{3(g)}
                                                                                                      (1 \text{ mk})
     (b) (i) As the forward reaction occurs with a decrease in the number of gas molecules, Le
          Chatelier's principle states that increasing the pressure causes the equilibrium to move
          to the right, increasing the yield of ammonia.
                                                                                                     (1 \text{ mk})
          (ii)As shown in the graph (or predicted by Le Chatelier) a higher temperature would
          reduce the equilibrium yield of ammonia, so that the reaction is carried out at the
          lowest temperature which affords an acceptably fast reaction.
                                                                                                     (1 \text{ mk})
     (c) Any one of:
                                                                                                    (1 \text{ mk})
          To maintain the 3:1 stoichiometric ratio of hydrogen:nitrogen, to prevent an excess of
          either in the chamber.
          To ensure no oxygen is present, to avoid explosion with hydrogen.
          To ensure no S compounds are present, to avoid inactivating the catalyst.
     (d) One of fertilisers, production of nitric acid, dyes, explosives, Solvay process etc (1 mk)
27. (a) moles of oxygen = 20 \times 0.2095/24.47 = 0.171 mol
                                                                                                     (1 \text{ mk})
     (b) moles of argon = 200 \times 0.00934/24.47 = 0.0763 mol
                                                                                                     (1 \text{ mk})
          mass of argon = moles x mol. mass = 0.0763 \times 39.95 = 3.05 \text{ g}
                                                                                                     (1 \text{ mk})
```

28.	(a)	Wash the precipitate with distilled water  Dry the precipitate	(1 mk)
	(b)	Moles of ppte. = $0.125/\text{mol.}$ mass = $0.125/222.56 = 5.62 \times 10^{-4}$	(1 mk)
	(-)	Moles of P present = 2 x moles ppte	(1 mk)
		$=1.123 \times 10^{-3} \text{ mol}$	(1 1111)
		% P in detergent = $1.123 \times 30.97 \times 100/2.85 = 1.22\%$	(1 mk)
	(c)	Excess nutrients such as nitrates and phosphates can cause excessive growth of a	lgae
		and other organisms in waterways. (1 mk) One type of alga that may grow excess	sively
		is blue/green algae which is toxic to livestock and can affect humans. The algal	
		blocks the entry of sunlight, and increases BOD (1 mk) creating an anoxic	
		environment leading to death of aquatic organisms.(1mk) (	3 mks)
20	7-1	FF 1 0 6 50/10 10	
29.		$[Fe] = 2.6 \times 50/10 = 13 \text{ ppm}$	(1 mk)
		Mass of Fe in 260 g = $13 \times 260 \times 10^{-6} = 3.38 \text{ mg}$	(1 mk)
		% daily allowance = 3.38 x 100/18 = 18.8%  The undiluted decide has a concentration beyond the calibration and a fauthority and the calibration	(1 mk)
	( <b>u</b> )	The undiluted drink has a concentration beyond the calibration scale for the	(11-)
		spectrometer used.	(1 mk)
SU	GGI	ESTED ANSWERS TO OPTIONS	
<b>^</b>		TOMAS TANDUSCHIA COURT PROPERTY.	
. —		TON 30 – INDUSTRIAL CHEMISTRY	
(a)		$k = [CH_3OH]/([H_2]^2 \times [CO])$	(1  mk)
	(ii	) Concentrations = $1.0/10$ , $0.9/10$ and $0.1/10$	
		$k = 0.01/(0.09^2 \times 0.1) = 12.3$	(1  mk)
	(ii	i) As the forward reaction is exothermic the equilibrium constant would increase	<b>,</b>
		with the equilibrium shifting to the right.	(1  mk)
(b)	(i)		(1  mk)
		Products: sodium carbonate & calcium chloride (solution)	(1  mk)
	(ii	· · · · · · · · · · · · · · · · · · ·	
		Proximity of limestone deposit	
		Means of disposal of calcium chloride waste	
		Energy supply	
(a)	c.	Proximity to markets and/or workforce  Any two	(2  mk)
(c)	20	pap is manufactured by the saponification of fats and oils with sodium hydroxide	,
		lution to form glycerol and the sodium salts of fatty acids (soaps).	(2 mk)
		ne anion of the soap is the cleaning agent, consisting of a long hydrophobic alkyl	
		ined to the hydrophilic carboxyl group. ~~~~~~~COO-	(1 mk)
	- W	reasy 'dirt' particles become dissolved into the alkyl chains of a collection of soap	
	in	olecules, the hydrophilic carboxyl groups allowing the assembly (a micelle) to dis	sperse
		It; a suitable diagram and short description)	(0 1)
(d)	(i)		(2 mk)
(u)		20/ 20/	(1 mk)
	(ii		
		Lower the temperature	
		Increase the pressure	,,
	7	(Use a catalyst to reach equilibrium more quickly)  Any two	(2  mk)
	(ii	i) Sulfuric acid is a very strong acid, oxidising agent and dehydrating agent.	(11-)
		The acid must be handled only with protective glasses and gloves.	(1  mk)
		Dilution must be carried out by slowly adding the acid to water, with stirring.	(1 mk) (1 mk)
			1.1 1116

30.(e) The mercury cell electrolyses sodium chloride solution with an inert anode, where chlorine is produced, and a mercury pool cathode into which sodium metal dissolves. The mercury amalgam flows into a tank of water into which the sodium dissolves forming sodium hydroxide and hydrogen. Thus the main products are sodium hydroxide, chlorine and hydrogen. (3 mk)

Technical considerations:

Proximity to electric power, transport facilities, workforce, raw materials.

Skilled staff monitoring operation to prevent release of toxic substances (chlorine, mercury, sodium hydroxide) or explosion (hydrogen); a very high level of plant safety is needed.

Any 2 (2 mk)

#### Environmental:

Minimise impact of plant (noise, light, emissions) and transport operations on surrounding population.

Need to remove all mercury from waste discharge to waterways. Release of chlorine to be kept below maximum permitted level.

Any 2 (2 mk)

# **QUESTION 31 - SHIPWRECKS & SALVAGE**

(a) (i) manganese (1 mk)

(ii)  $Mn_{(s)} \rightarrow Mn^{2+} + 2e^{-} +1.18 \text{ V}$  $Sn^{2+} + 2e^{-} \rightarrow Sn_{(s)} -0.14 \text{ V}$ 

 $Mn_{(s)} + Sn^{2+} \rightarrow Mn^{2+} + Sn_{(s)}$  +1.04 V (2 mk)

Criteria	Marks
Correct net equation and correct voltage calculation	2
Either correct net equation OR Correct voltage calculation	1

(b) (i) Zinc metal electrode is the Anode. Reference to an E<sup>o</sup> Potential Table indicates that zinc is more readily oxidised than iron. (2 mk)

Zine is more readily exicused than from.	
Criteria	Marks
Zinc electrode identified as ANODE AND	2
a correct supporting piece of information eg. reference to E <sup>0</sup> Table	
Zinc electrode identified as Anode, no justification or an incorrect	1
justification	<sup></sup> .

(ii) Bubbles of a gas (most likely to be oxygen gas) forming around the location of the anode and at the cathode, copper metal is electroplating the electrode. (2 mk)

Criteria Criteria	Marks
Correct observation occurring at the Anode AND Correct observation occurring at the Cathode	2
A correct observation occurring at the Cathode  A correct observation occurring at one electrode	1

(c) Preparing

Removed of any colors and described to the second second

Removal of any calcareous deposits by physical means such as chipping with a small drill as other methods of chemical treatment such as acid baths would damage the wood. Long immersion in fresh water with repeated water changes to permit removal of soluble salts so as to prevent damage from rapid drying out of the wood

<u>Preserving:</u> Impregnation the timber with an inert chemical material to fill the empty spaces inside the wood that were filled previously with the sea water as the wood will shrink and distort without this preservative step. Maintain the timber in conditions of controlled humidity, temperature and light as this will minimize any further degradation of the wood.

31. (c)

Criteria	Marks
Detailed descriptions of all processes	4 - 5
At least TWO methods of preparing and TWO methods of preserving of wooden artifacts	4-3
At least ONE method of preparation AND ONE method of preserving wooden artifacts	2 - 3
One description to be detailed	ļ
A method of preparing wood OR preserving wooden artifacts after long immersion in the sea.	1

(d) (i) Any suitable passivating metal e.g. aluminium

(1 mk)

(ii) Organise and weigh multiple pieces of equal sized masses of magnesium, iron and lead. Place a single sample of each metal into a single test-tube.

Use similar shaped & sized test-tubes. Add equivalent amounts of water to each tube. Leave some tubes with only metal & water.

To some tubes add 5 droplets of and to others add 10 drops of 2M H<sub>2</sub>SO<sub>4</sub>.

Leave all test-tubes for a similar specified period of time e.g.7 days

Wash and dry all pieces of metal then reweigh to determine any metal loss through corrosion.

Determine if any relationship exists between the degree of corrosion and the acidity of the conditions in which they were placed.

Criteria	
Main features of a feasible method described AND the concept of controlling all features except acidity and possibly the metal type	Marks 2
described	
Main features of a feasible method described	1

(iii) The dilemma exists because recent research is conflicting with traditionally accepted chemistry thinking.

Traditional thinking was that at the bottom of the ocean, it is absolutely dark and extremely cold (almost 0 °C) and conditions of low oxygen levels, these conditions would lower reaction rates thereby reducing the rate of corrosion of iron is shipwrecks.

Recent discoveries are showing that the waste products of particular types of chemosynthetic bacteria are acidic in nature, these acidic conditions cause acceleration of the rate of corrosion of steel in the very deep ocean.

(3 mk)

Criteria Sociali.	Marks
Discusses why a dilemma exists	3
Mentions a reason for slow corrosion e.g. low temps slowing reaction rates OR	3
causing acidic conditions which speed corrosion	·
Mentions a reason for slow corrosion e.g. low temps slowing reaction rates AND	2
Mentions a reason for newly discovered corrosion rates i.e. bacterial wastes causing acidic conditions which speed corrosion	
Mentions a reason for slow corrosion e.g. low temps slowing reaction rates OR Mentions a reason for newly discovered corrosion rates i.e. bacterial wastes causing acidic conditions which speed corrosion	1

## 31.(e) Possible Corrosion Protection Alternatives

(7 mk)

Painting to prevent water & oxygen reacting with the iron in steel.

Painting with special resins to prevent water & oxygen reacting with the iron in steel. Impressed Electrical Currents run through the metal of the hull as one electrode from the ships generators and another electrode external to the ship sets up a circuit to reduce the oxidation of the hull.

Sacrificial Anodes where pieces of more reactive metal e.g. zinc are attached to the ship so that they are corroded in preference to the metal of the ships hull.

# Comparisons

Painting is the cheapest and lowest technology approach.

Sacrificial anodes are more expensive as they require regular replacement of the more active metal which has been corroded away. Not a viable option for extremely large vessels.

Impressed current systems are very expensive to establish but are fairly cost effective over the lifetime of the vessel and have the advantage of weighing less than sacrificial anode systems.

#### Recommendation

The recommendation to the Silver Star Line is to paint the vessels with special resin based paints as these form hard abrasion resistant surfaces that protect the steel from the conditions which cause corrosion. This choice is done because it is cost effective (relatively cheap), it is simple to apply and provides a good initial level of protection to the metal.

Criteria	Marks
Chooses a preferred method of corrosion protection and gives one or more reasons to justify that choice	4 - 5
Makes a COMPARISON of the TWO or more methods of corrosion protection for steel	2 - 3
Outlines at least TWO methods of corrosion protection for steel	1

# **QUESTION 32 - BIOCHEMISTRY OF MOVEMENT**

(a) (i) for drawing the correct structure

COOH | (1 mk) H<sub>2</sub>N—C—H | R

R represents the side chain or an R group which is different in each amino acid

(ii) identifying and explaining the four kinds

identifying and explaining the four kinds of interacting forces. (2 mk) α is the hydrophobic forces between the hydrophobic R groups of some non-polar amino acids like phenylalanine, leucine, isoleucine, methionine, valine, tryptophan. β is the electrostatic forces between oppositely charged R groups, e.g. the negatively charged COOH— of a glutamic acid residue with the -+NH<sub>3</sub> of a lysine residue in an adjacent loop.

 $\chi$  is the disulfide bonds formed from a cysteine residue with another cysteine residue in an adjacent loop. This is the strongest of all the four factors.

 $\delta$  is the hydrogen bonding forces between the hydroxyl group of a serine, for example with a ring nitrogen atom of a histidine residue in an adjacent loop of the same chain.

These non-covalent interactions ( $\alpha$ ,  $\beta$  and  $\delta$ ) are individually weak but collectively strong.  $\gamma$  is a covalent cross-linkage which is strong in force.

(Although the native tertiary conformations of globular proteins are the stablest forms their polypeptide chains can assume, the tertiary conformation of globular proteins must not be regarded as absolutely rigid and fixed. Many globular proteins normally undergo changes in their conformation in the course of their biological function, e.g. enzyme molecules can change their conformation when they bind their substrate.)

## 32. (a) (ii) continued

Without the tertiary structure, the protein will not be able to function properly. Those four interactions are of vital importance to the function of the protein. Heat and various other treatments can denature or unfold native globular proteins by disturbing the various kinds of interacting forces.

(b) (i) 2 marks for assessing the validity of the statement. No mark will be given if a student says the statement is correct. (2 mk)

The statement is incorrect and misleading, since it wrongly suggests that the bond itself contains the energy. This is not the case. In fact the breaking of the chemical bonds requires input of energy.

 $ATP + H_2O \rightarrow ADP + P_1$ 

The free energy released by the hydrolysis of the phosphate esters thus does not come from the specific bond that is broken but results from the fact that the products of the reaction have a smaller free energy content than the reactants. It is however quite appropriate to use the term "high-energy phosphate compound" to referring to ATP having a large negative  $\Delta G^{\circ}$  of hydrolysis.

(ii) 2 marks for describing briefly how ATP can be replenished from oxidative phosphorylation or 1 mark only for just mentioning oxidative phosphorylation without further descriptions.

(2 mk) The oxidation of NADH and FADH is coupled with the phosphorylation of ADP to produce ATP. This oxidative phosphorylation is a multi-stage pathway, where a total of about ten redox reactions occur. It can be briefly represented by the following equation:

 $NADH + H^{+} + 3P_{1} + 3ADP + \frac{1}{2}O_{2} \rightarrow NAD^{+} + 3ATP + H_{2}O$ 

(c) I mark for stating the similarities

(5 mk)

4 marks for stating the differences. Out of these 4 marks, 2 marks for structural differences, 1 mark for functional differences and 1 mark for differences in the use of substrates.

# Similarities:

Both Type 1 and Type 2 muscle cells are for contraction which bring about movement of the body and part of the body.

Both types of muscle cells contain multi-nucleated cells made from a bundle of about one thousand fibrils. These fibrils consist of alternating sections of thick and thin filaments. Thick filaments consist of myosin. Thin filaments consist of actin. During muscle contraction, the thin actin filaments and the thick myosin filaments slide past each other. The area of overlap is extended as a muscle contracts.

Both types of muscle cells require the presence of both calcium ions and ATP.

**Differences** 

·	Type 1	Type 2
Structures	fewer contractile filaments, many mitochondria, richly supplied with blood vessels	many contractile filaments, fewer mitochondria, reduced supply of blood vessels
Functions	slow twitching, used for light, endurance exercises	fast twitching, used for heavy sprinting style exercises
Use of substrates	can use glucose, fatty acids, amino acids as fuels; adequate oxygen supply, ATP is derived from the process of oxidative phosphorylation	use muscle-stored glycogen as fuel.

2 marks for describing the experiment. Out of these 2 marks, 1 mark for briefly explaining the mathematical treatment

1 mark for describing how to improve the accuracy 1 mark for describing how to improve the reliability

- (i) The amount of heat released when 1 mole of a substance undergoes complete combustion in oxygen.
- (ii) An example of the named lipid can be glyceryl tristearate. A known mass of the lipid is burnt in a small spirit burner. A suitable wick is used. Heat produced by the combustion of the lipid is absorbed in 200 mL of water contained in a 250 mL conical flask, the bottom of which is about 4 cm above the burner. By finding the rise in temperature of the water, the amount of heat can be calculated using the formula H = m.c. T

  Heat change = mass of water x specific heat capacity x rise in temperature By dividing the mass of the lipid burnt with the molecular mass of the lipid, the number of mole of the lipid used can be calculated.

  The molar heat of combustion can be calculating by dividing the heat evolved with the number of mole of lipid.
- (iii) The accuracy can be improved by reducing the heat loss to the environment by insulating the flask and shielding the apparatus from all drafts. The heat loss to the flask can be taken into account by calculating the heat absorbed by the glass if the specific heat capacity of the glass is known.
  The reliability of the experiment can be improved if complete combustion of the lipid is insured. Oxygen gas can be supplied during the combustion. The experiment can be carried out with a bomb calorimeter.
- (e) 2 marks for describing how acetyl Co-A is formed via glycolysis (7 mk)
   1 mark for describing how the acetyl-CoA is fed into the TCA cycle.
   2 marks for describing how fatty acids are converted to fatty acyl-CoA
   1 mark for describing how acetyl-CoA from β oxidation of fatty acids is fed into TCA cycle.
   1 mark for mentioning the role of the β oxidation of fatty acids in the inhibition of the pyruvate conversion to acetyl CoA

The metabolism of glucose consists of two stages – glycolysis and tricarboxylic acid cycle. In glycolysis, each glucose molecule is broken down into 2 molecules of pyruvate via several intermediate compounds.

Under aerobic conditions the two pyruvate produced by glycolysis enter the mitochondria where they continue to undergo oxidation. The pyruvate reacts with co-enzyme A in the presence of pyruvate dehydrogenase to form acetyl-coenzyme A (acetyl-CoA) and CO<sub>2</sub>, accompanied by the reduction of NAD+ to NADH.

pyruvate + coenzyme A + NAD $^+$   $\rightarrow$  acetyl-coenzyme A + CO $_2$  + NADH Acetyl-CoA is the starting compound for the TCA cycle. Acetyl Co-A reacts with oxaloacetate (a 4 carbon compound) to form citrate (a 6 carbon compound). 6 carbon compounds are converted to 5 carbon compounds and to 4 carbon compounds with the release of carbon dioxide and hydrogen. Hydrogen passes through the cytochrome chain to generate ATP.

Fatty acids must be transported to the mitochondrial matrix and activated before they can be oxidised. The metabolism fatty acids requires the presence of coenzyme A, the carrier molecule. Fatty acids are converted to an activated fatty acyl—CoA by the action of ATP as shown in the equation:

fatty acid + CoA + ATP  $\rightarrow$  fatty acyl-CoA + AMP + PP<sup>1</sup>

The fatty acyl—CoA is now ready for  $\beta$  oxidation. This is a four-step process with each step catalysed by an enzyme. The process involves the reduction of one molecule of each of the coenzymes NAD<sup>+</sup> and FAD. The end result of the oxidation is the production of acetyl—CoA and another fatty acyl—CoA molecule, which is now shorter by two carbon atoms than the original fatty acid.

fatty acyl-CoA + FAD + NAD<sup>+</sup> + CoA +  $H_2O \rightarrow$ 

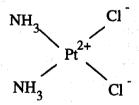
fatty acyl (containing 2 less C atoms)—CoA + acetyl—CoA + FADH<sub>2</sub> + NADH + H<sup>+</sup> This process can be repeated, removing two carbon atom sections each time, until the full acyl carbon chain of the fatty acid has been oxidised into acetyl—CoA. This oxidation of fatty acids produces large quantities of energy in the form of ATP.

Fatty acid oxidation is in direct competition with glucose metabolism, with both the fatty acids and pyruvate competing for CoA. There is a finite amount of CoA available in the cell's cytoplasm. Thus high levels of fatty acid entering a cell can inhibit the metabolism of pyruvate by consuming the CoA required.

When a person is starving, the β oxidation of fatty acids will dominate.

### **QUESTION 33 - CHEMISTRY OF ART**

(a)	(i)	$1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^5$	(1 mk)
	(ii)	Colour changes from purple to colourless	(1 mk)
		Oxidation state changes from +7 to +2	(1 mk)
(b)	(i)	At room temperature all atoms have their electron in the n=1 level,	(1 1111)
		so that no transitions involving the n=2 level are possible.	(1 mk)
	(ii)	Features of Bohr model for H atom:	()
		Electron in circular orbit around nucleus	
		Possible orbits restricted to certain energy values.	(1 mk)
		Light is emitted when electrons fall from higher to lower energy levels.	(1 mk)
		Line spectrum is generated by electron transitions between energy levels,	
		where the line frequency is proportional to the energy difference.	(1 mk)
(c)		eg., cisplatin Pt(Cl) <sub>2</sub> (NH <sub>3</sub> ) <sub>2</sub> which consists of a platinum ion Pt <sup>2+</sup> with two	(
	amn	nonia molecules and two chloride ions as ligands	



Each ligand bonds to the Pt<sup>2+</sup> ion using an electron pair to form a coordinate covalent bond. (Lewis bases) - diagram (3 mks) Cisplatin is used as an anticancer agent in treating tumours. It binds to DNA chains in cancer cells, thus preventing their replication. (2 mks)

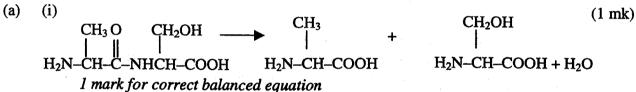
(d) (i)Transition metals have incomplete d-subshells of the n=3, n=4 and n=5 shells.(1 mk)
 (ii) The electrons of the d and s subshells have similar energy values. (1 mk)These metals can form ions by losing varying numbers of electrons from the d subshell and the preceding s subshell, leading to a variety of oxidation states (1 mk) (2 mks)
 (iii)The melting points increase from Sc to V then decrease, with Zn having the lowest value. The trend reflects the number of d electrons available for bonding and the number of empty orbitals available. (1 mk)
 The density increases from Sc to Cu due to increasing atomic mass and decreasing

atomic radius. (Zinc, with filled d subshell has a lower density) (1 mk)
Magnetism is shown only by Fe, Co and Ni which have unpaired d electrons in their structure. (1 mk)

(e) eg: only (select pigment, origin, composition (3mks), extraction (1mk), application (3mks).

Ochres are found in various colours, these reflecting varying degrees of hydration of iron oxides. They are formed by chemical weathering near the earth's surface. Red ochre contains anhydrous iron(III) oxide Fe<sub>2</sub>O<sub>3</sub>, yellow ochre approximates Fe<sub>2</sub>O<sub>3</sub>.H<sub>2</sub>O, while brown ochre is nearly pure limonite FeO(OH). All ochres are highly insoluble, their extraction consisting of excavation and grinding to a fine powder. Aborigines sometimes applied ochre directly to cave walls or mixed the ochre with a spreading agent such as orchid juice, wax, honey, egg yolk or plant gums. Modern paints use spreading agents, such as oil or aqueous solutions in a similar way. The agent must be sufficiently viscous to prevent running and bind the pigment particles together as the agent evaporates or dries. The agent itself has little colour so that the paint deposit retains the colour of the original pigment.

# **QUESTION 34 - FORENSIC CHEMISTRY**



(ii) During electrophoresis charged particles migrate through an electrolyte when a voltage is applied. The speed at which the particles move depends on their size and charge. When separating amino acids, the pH of the electrolyte will influence the charge on each amino acid. At low pH, the amino acid molecules will be positively charged, at high pH, they will be positively charged and at an intermediate pH they will be neutral. The actual pH at which these charges form is different for each amino acid.

Criteria	Marks
an outline of the electrophoresis process a description of the charges on amino acids at high, low and intermediate pH's an understanding that different amino acids become charged at different pH's	2
only 1 of the above criteria	1

(b) (i) Atoms need to absorb energy in order to promote one or more of its electrons to energy levels than those fill when in ground state. This energy may be given to the atom by heating to a high temperature, by applying a high electrical voltage or by subjecting the atom to certain frequencies of electromagnetic radiation. The light is emitted when the electrons in the excited states fall back to their "normal" ground state energy level.

Criteria	Marks
demonstrate an awareness of at least two types of energy that can be ab	
to excirte an atom	
reference to absorbing energy	1

(b) (ii) Each element has its own unique emission spectrum, as seen by the Cu, Li and Cr spectra shown above. This spectrum is not affected by the presence of atoms of other elements. A spectrum produced by a mixture of elements will contain spectral lines of all the elements present, although closely spaced lines may not be completely resolved and weak lines may not show up.

The sample spectrum shown shows all the spectral lines in the Cu and Cr spectra, indicating their presence in the sample. Some of the lines in the Li spectrum are not present indicating that Li is not present in the sample. The presence of additional lines not in the Cu, Li or Cr samples suggests the presence of another element.

In soil analysis, detection of an unusual element in a sample can sometimes pinpoint the location from which the sample originated. While this technique identifies the elements present in the sample, it does not give any indication of the nature of the compounds present

Criteria	Marks
discussion of how the presence of an element can be identified in a mixed sample spectrum	2
discusison of how the absence of spectral lines can indicate the absence of an element in any significant quantity	
identifying that another unknown element is present in the sample	
discussion of how the presence of an element can be identified in a mixed sample spectrum with reference to the sample shown OR	1
identifying that CU and Cr are both present in sample the spectrum shown	

(c) KMnO4 is a strong oxidising agent changing from purple to brown in colour when it reacts. A cold, dilute, aqueous solution of KMnO<sub>4</sub> will react differently with the three compounds named.

KMnO<sub>4(aq)</sub> does not react with cyclohexane and so retains its original purple colour when added to a sample of cyclohexane.

KMnO<sub>4(aq)</sub> will oxidise both glycerol and 1-propanol, changing from purple to brown. It is the OH functional group in these compounds that reacts. Glycerol is a triol, so three times as much KMnO<sub>4</sub> will be required to react with an amount of glycerol compared with an equal amount of 1-propanol.

Therefore by measuring the amount of KMnO<sub>4</sub> required for complete reaction samples of glycerol and 1-propanol can be distinguished.

Criteria	Marks
describes the use of dilute aqueous KMnO <sub>4</sub> as a oxidising agent describes observations made when KMnO <sub>4</sub> is mixed with all three compounds named	4-5
explains why glycerol reacts with more KMnO <sub>4</sub> than 1-propanol	<b>.</b>
describes observations made when KMnO <sub>4</sub> is mixed with some of the compounds named OR	2-3
explains why glycerol reacts withmore KMnO <sub>4</sub> than 1-propanol	
describes observations made when KMnO <sub>4</sub> reacts	1

(d) (i) Glucose (1 mk)

(ii) Drops of iodine solution, when added to an aqueous solution or suspension of starch, will form a deep blue colour. Drops of a solution of calcofluor, when added to a finely divided sample of cellulose dispersed in water will produce a mixture that will emit a blue glow when placed under a UV light source.

Criteria	Marks
describes tests (including observations) for both cellulose or starch	2
identifies tests for both cellulose and starch OR	1
describes a test for cellulose or starch	

(iii) Cellulose, starch and glycogen are all biopolymers made glucose monomer units. Glucose may exist in a number of isomeric forms, two of which are called α-glucose and β-glucose. Cellulose consists of long chains of β-glucose units (typically between 200 to 400 units) joined together. Because of the β- arrangement of the OH group on the carbonyl carbon in glucose, every second glucose unit in the cellulose chain is inverted producing a very linear, unbranched chain. These linear chains intertwine, forming hydrogen bonds with one another, to form fibres. Starch and glycogen consist of α-glucose units joined together, each unit the same way up, producing chains that curl around into globular shapes. Starch may consist of between 60 to 6000 glucose units, the shorter chains being unbranched while the longer chains are branched. Glycogen is a highly branched chain typically only a few hundred glucose units long.

Criteria Marks identifies that cellulose is compossed of different isomers of glucose than is 3 starch or glycogen description of the arrangement of the glucose units resulting in different shaped chains in cellulose compared to starch or glycogen identifying the branching of the polymer chains in stgarch and glycogen and the lack of branching in cellulose outlining some differences between stach and glycogen identifies glucose as the common monomer unit in at least two of cellulose, 2 starch or glycogen description of the inverted arrangement of alternate glucose uints in cellulose OR identifying the n\branching of the polymer chains in starch &/or glycogen identifies that cellulose, starch or glycogen are polymers made of glucose 1 monomer units.

(e) Scanning tunnelling microscopy is a valuable tool to the forensic chemist. Even though it cannot be used on all objects, it does given detailed information about the surface of an object that cannot be obtained by most other analytical techniques. It can be performed on very small samples of material and it is a non-destructive analytical technique, although the small electric current used may damage some sensitive materials. Although it can only be used for analysing the surface of conducting materials, when used in conjunction with other techniques that analyse the composition of the material, it can allow the identification of an individual source of the material, allowing the matching of unique surface details. These details will be a result of the unique conditions to which an individual object will be exposed over time such as weathering conditions, stains and surface deposits. Similar detailed surface analysis of non-conducting materials requires a slightly different technique.

Scanning tunnelling microscopy uses a very fine needle which is moved at a specific height above the surface features. When a voltage is applied between the needle and the surface, electrons in the surface move to the tip of the needle producing a small current. Vertical movement of the needle maintains a constant current and hence a constant nheight above the surface. This vertical movement can be used to produce a detailed image of the surface. Surface features of less than one nanometre in height or width can be detected.

Criteria	Marks
clear outline of the how the scanning tunnelling microsocpe works	7
an explanation of the types of images produced by a scanning tunnelling	
microscope	
demonstrates an understanding of some of the advantages and some of the	
limitations of using scanning tunnelling microscopy with reference to the value	
or the infoermation gained to the forensic chemist	
outlien of how the scanning tunnelling microscope works	5-6
description of the types of images produced by a scanning tunnelling	
micrscope	
description of some of the advantages and liitaions of using scanning	
tunneuing microscopy	
description of the types of images produced by a scanning tunnelling	3-4
microscope ND	3.4
outline of how the c\scanning tunnelling microscope works OR	
a descrition of some of the advantages and limitations of using scanning	
tunnelling microscopy	
description of the types of images produced by a scanning tunnelling	1-2
microscope OR	1-4
description of some of the advantages and/or limitations of using scanning	
tunnelling microscopy	

The Trial HSC examination, marking guidelines/suggested answers and 'mapping grid' have been produced to help prepare students for the HSC to the best of our ability.

Individual teachers/schools may alter parts of this product to suit their own requirements.

# Chemistry 2002 Trial Examination Mapping Grid

Question	Marks	Content 9.	Syllabus outcomes H	Targeted performance bands
1	1	2.1	3,9	2-3
2	1	2.2	3,9	2-3
3	1	2.4	3,6,7	3-4
4	1	2.3	4,11	3-4
5	1	2.5	4,6,14	4-5
6	1	3.1,3.3	14	3-4
7	1	3.1,3.4	12,14	3-4
8	1	1,3.4	11,12	2-3
9	1	3.2	8,14	4-5
10	1 .	3.3	10	4-5
11	1	4.4	6,14	3-4
12	1	4.2	6,8	2-3
13	1	4.5	4,14	2-3
14	1	4.3	1,11	2-3
15	1	4.5	8,14	3-4
16(a)	2	2.3	2,6	3-5
16(b)	2	1,2.3	7,12,14	2-6
16(c)	2	1,2,3	11,9,13	2-6
17(a)	2	2.3	4,14,13	2-5
18(a)	1	2.4	8,13,14	3-4
18(b)	1	2.4	8,13	3-4
18(c)	1	2.4	8,14	3-4
18(d)	2	2.4	3,4	2-5
19	3	2.1	6,9,13	3-5
20(a)	1	2.5	2,6,13	3-4
20(b)	1	1,2.5	11,13	2-3
20(c)	2	2.5	1,3,4,11	3-6
21	2	3.2	8,10,13	4-5
22	3	3.4	6,8,13	3-6

Question	Marks	Content 9.	Syllabus outcomes H	Targeted performance bands
23(a)(i)	1	3.5	9,13	4-6
23(a)(ii)	1	3.5	6,13	3-4
23(a)(iii)	1	3.5	9	4-5
23(a)(iv)	2	1,3.5	11.13	4-6
23(b)	3	3.5	2,6	3-4
24(a)	2	3.2	8,13	3-4
24(b)	2	3.2	8,13,14	3-5
25(a)	1	3.4	6,8,13	3-4
25(b)	2	3.4	10,13	3-4
25(c)	1	3.4	10,13	2-4
26(a)	1	4.1	8,13	2-3
26(b)(i)	1	4.1	8,11,13	3-4
26(b)(ii)	1	4.1	8,11,13	2-3
26(c)	2	4.1	3,11,13	2-3
26(d)	1	4.1	4	3-5
27(a)	1	4.4	10,13	4-6
27(b)	2	2.1,4.4	10,13	3-4
28(a)	2	1,4.5	11,13	3-4
28(b)	2	4.5	10,13	3-4
28(c)	3	4.5	4,8,13	4-6
29(a)	1	1,4.3	10,14	
29(b)	1	1,4.3	10,13	
29(c)	1	1,4.3	10,13	
29(d)	1	1	11,12,13	
TOTAL	=75	†	<del>                                     </del>	

Question	Marks	Content 9.	Syllabus outcomes H.,	Targeted performance
Option 1: i	ndustrial		П.,	bands
30(a)(i)	1	5.2	8,10	2-4
30(a)(ii)	1	5.2	10.13	2-4
30(a)(iii)	1	5.2	12,8,13	4-5
30(b)(i)	2	5.6	3,7	2-3
30(b)(ii)	2	5.6	4,11,13	4-5
30(c)	5	5.5	2,3,6	4-6
30(d)(l)	1	5.3	8,10	2-4
30(d)(ii)	2	5.3	3,8,13	2-4
30(d)(iii)	3	5.3	1,11,13	3-5
30(e)	7	5.4	34813	3-6
Option 2: S	hlowrec	ks and Salv	/80e	1 3-0
31(a)(i)	1	6.3,6.4	7,10	3-4
31(a)(ii)	2	6.3	7,8,10	3-4
31 (b) (i)	2	6.3	8,11,13	3-4
31(b)(ii)	2	1,6.3	6,8,13	2-4
31(c)	5	6.7	3,8,16	3-6
31(d)(i)	1	6.6	6.13	2-3
31(d)(ii)	2	6.6	11,12	3-5
31(d)(lii)	3	6.5,6.6	3,14,15	3-6
31(e)	7	6.4	3,8,13,14,16	2-6
Option 3: E	lochemi	try of May	ement	
32(a)(i)	1	7.4	2,13	2-3
32(a)(ii)	2	7.4	2,13	3-5
32(b)(l)	2	7.1	2,7,9,10,13	3-4
32(b)(ii)	2	7.1	2,7,9,10,13	3-4
32(c)	5	7.5,7.6	2,13	3-6
32(d)(i)	1	1.7.3	2,5,7,8,9,10,11,	2-6
32(d)(ii)	2	1,7.3	12,13,14,15,16	2-6
32(d)(lii)	3	1,7.3		2-6
32(e)	7	7.8,7.9, 7.10	2,7,9,10,13,16	3-6

Question	Marks	Content 9.	Syllabus outcomes H	Targeted performance bands
Option 4: 0	hemistr	of Art		
33(a)(i)	1	8.4	2,6,13	3-4
33(a)(ii)	2	8.5	6,13	2-4
33(b)(l)	1	8.3,8.4	6,14	4-5
33(b)(N)	3	8.3	2,6	3-6
33(c)	5	8.7	3,4,13	3-6
33(d)(i)	1	8.4	6	2-3
33(d)(ii)	2	8.5	6,7,13	3-4
33(d)(iii)	3	8.6	6,8,13	4-6
33(e)	7	8.1,8.2	1,3,4,8,13	2-6
Option 5: F	orensic	Chemistry	· · · · · · · · · · · · · · · · · · ·	<del></del>
34(a)(i)	1	9.4	9	3-5
34(a)(ii)	2	9.4	4,8,9	5-6
34(b)(l)	2	9.7	6,7	3-5
34(b)(ii)	2	9.7	3,4	2-5
34(c)	5	9.1,9.3	8,9,10,14	3-6
34(d)(i)	1	9.2	9	2-3
- 34(d)(ii)	2	9.2	8,8,11	2-4
34(d)(iii)	3	9.2	9	3-5
34(e)	7	9.6	1,3,4	3-6