- 1 Which of the following gives the largest number of moles of particles?
 - A ions in 0.5 dm³ of 0.1 mol dm⁻³ NaOH(aq)
 - **B** IO_3^- ions in 0.2 dm³ of 0.5 mol dm⁻³ KH(IO_3)₂
 - C molecules in 480 cm³ of ClO₂(g) at room temperature and pressure
 - D atoms in 1.92 g of O₃
 - 2 Use of the Data Booklet is relevant to this question.

Which of the following statements are correct?

- 1 The second ionisation of any element is always higher than its first ionisation energy.
- 2 The ionic size for Na⁺ is the same as Al³⁺ because they are isoelectronic.
- 3 The removal of the two electrons from Cr to form Cr²⁺ is from the 3d subshell.
- 4 An aluminium atom will require more energy to form a gaseous ion with the charge of +4 compared to a silicon atom.
- A 1 only
- B 1 and 3 only
- C 1 and 4 only
- D 1, 2, 3 and 4
- 3 The table gives the successive ionisation energies for an element E in Period 4.

	1st	2nd	3rd	4th	5th	6th
Ionisation energy / kJ mol ⁻¹	950	1800	2700	4800	6000	12300

What is the formula of the chloride of E?

A ECI

B ECl₂

C ECl₃

D ECla

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[Turn over

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4 Tetrathionic acid, H₂S₄O₅, has been used as an antidote in cyanide poisoning.

HO —
$$S^{1}$$
 — S^{2} — S^{3} — S^{4} — OH

In tetrathionic acid, the oxidation number of hydrogen is +1 and oxygen is -2.

What could be the oxidation number of the four sulfur atoms?

	S ¹	S ²	S³	S ⁴
A	+5	0	0	+5
В	+6	0	0	+6
С	+2	+2	+5	+5
D	+2	+2	+6	+6

Arsenic trioxide, As₂O₃, reacts with zinc in the presence of of a suitable acid. In this reaction, the oxidation number of zinc increases by 2 and the oxidation number of arsenic decreases by 6.

How many moles of zinc are needed to reduce 1 mole of As₂O₃ in this reaction?

- A $\frac{1}{3}$ mol
- $B = \frac{1}{6} \text{ mol}$
- C 3 mol
- D 6 mol

6 Consider the molecule,

What is the hybridisation of each of the carbon atoms, C-1 to C-4?

	C-1	C-2	C-3	C-4
A	sp ²	sp ³	sp²	sp
В	sp	sp³	sp²	sp³
С	sp²	sp³	sp	sp²
D	sp ³	sp ²	sp²	sp

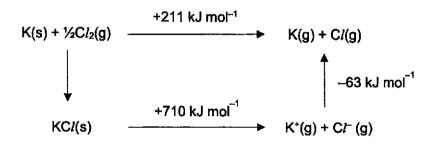
- 7 Which of the following cannot be explained by hydrogen bonding?
 - A The difference in the boiling points between water and ammonia.
 - B The difference in volatility between pentan-2-ol and octan-2-ol.
 - C The difference in boiling point between

- D The difference in the relative molecular mass of ethanoic acid molecules in benzene and in water.
- 8 Which of the following pairs consist of a planar molecule and a polar molecule?
 - A CH₂O and cis-CH₃CH=CHCH₃
 - B SO₃ and trans-CH₃CH=CHCH₃
 - C CH₃OCH₃ and CH₃CO₂H
 - D CH₃COCH₃ and CCI₄

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[Turn over

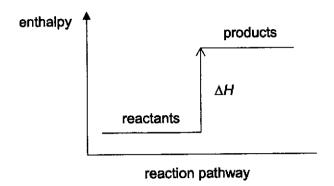
- 9 Which of the following reactions resulted in an increase in the bond angle of the underlined substance?
 - A $H_3O^+ + F^- \longrightarrow HF + H_2O$
 - B $NaNH_2 + H_2O \longrightarrow NaOH + NH_3$
 - C BH₃ + NaH → NaBH₄
 - D $CO_2 + H_2O \implies 2H^+ + CO_3^{2-}$
- 10 The enthalpy changes for some reactions are shown below.



Which of the following gives the correct signs of ΔH for the reactions?

	$K(s) + \frac{1}{2}C/_2(g) \rightarrow KC/(s)$	$K^+(g) + C\Gamma(g) \rightarrow K(s) + \frac{1}{2}CI_2(g)$	$KCI(s) \rightarrow K(g) + CI(g)$
A	-	+	+
В	–	_	· +
С	+	_	_
D	+	+	_

11 Which enthalpy change could be correctly represented by the enthalpy diagram shown?



- 1 neutralisation
- 2 combustion
- 3 formation
- A 1, 2 and 3
- **B** 1 and 2
- C 1 and 3
- D 3 only

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[Turn over

12 Use of the Data Booklet is relevant to this	is auestion
------------------------------------------------	-------------

What is the order of enthalpy changes of the following reactions from most negative to most positive?

$$\Delta H_1$$
 2NO₂(g) \rightarrow N₂O₄(g)

$$\Delta H_2$$
 CH₃(g) + C l (g) \rightarrow CH₃C l (g)

$$\Delta H_3$$
 CH₄(g) \rightarrow CH₃(g) + H(g)

- A $\Delta H_2 \rightarrow \Delta H_1 \rightarrow \Delta H_3$
- **B** $\Delta H_3 \rightarrow \Delta H_1 \rightarrow \Delta H_2$
- C $\Delta H_1 \rightarrow \Delta H_2 \rightarrow \Delta H_3$
- **D** $\Delta H_1 \rightarrow \Delta H_3 \rightarrow \Delta H_2$
- 13 Element X is in Period 3 of the Periodic Table. After mixing 0.5 mol of the chloride of X with limited cold water, 1.0 mol of HCl was formed.

What is element X?

- A Mg
- B Al
- C F

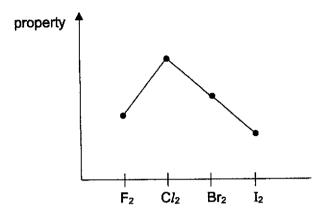
D Si

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14 Use of the Data Booklet is relevant to this question.

The following shows the variation of a property of the halogen molecules.



What is the property?

- A boiling point
- B bond energy
- C oxidising power
- D intensity of colour
- **X** is made up from two elements from Period 3 (excluding argon) of the Periodic Table. One of the elements has the highest melting point in the period. The other element has the smallest anion in the period.

What is the formula of X?

- A AICI3
- **B** Al₂S₃
- C SiCl4
- D SiS₂

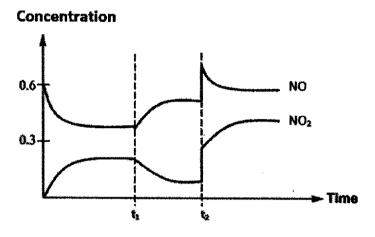
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[Turn over

16 At a temperature T K, 0.60 mol dm⁻³ of NO and 0.30 mol dm⁻³ of O₂ were introduced into a 2 dm³ vessel and allowed to reach equilibrium.

$$2 \text{ NO } (g) + O_2 (g) \iff 2 \text{NO}_2 (g) \qquad \Delta H < 0$$

The graph below shows the changes in the amounts of NO and NO_2 in the system with time. A change was made to the system at time, t_1 and t_2 .



What were the changes made at time, t₁ and t₂?

	t ₁	t ₂
A	The temperature was increased	Volume of the system is increased
В	The temperature was decreased	More O₂ was added
С	An inert gas was added at constant volume	More NO₂ was added
D	The temperature was increased	Volume of the system is decreased

17 An equilibrium can be represented by the following equation:

$$CH_3COOH + C_2H_5OH \Rightarrow CH_3COOC_2H_5 + H_2O$$

The equilibrium concentration of $\mathrm{CH_3COOH}$ is found to be 10 mol dm⁻³. At equilibrium, 5 moles of $\mathrm{CH_3COOH}$ is added to a 1 dm³ mixture.

Which of the following statements are correct?

- 1 The value of equilibrium constant will increase.
- 2 The position of equilibrium will shift to the right.
- 3 The new equilibrium concentration of CH₃COOH will be less than 10 mol dm⁻³
- 4 The new equilibrium concentration of CH₃COOH will be between 10 mol dm⁻³ and 15 mol dm⁻³
- A 1 and 2 only
- B 2 and 4 only
- C 1, 2 and 3 only
- D 1, 2 and 4 only

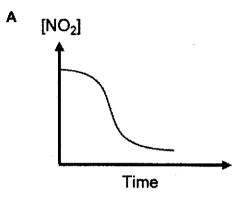
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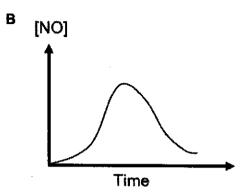
18 Nitrogen dioxide and methanoic acid reacted according to the equation below:

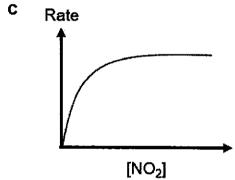
$$NO_2(g) + HCOOH(I) \rightarrow NO(g) + CO_2(g) + H_2O(I)$$

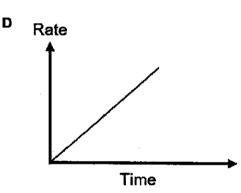
The progress of the reaction is monitored by the changes in concentration of NO₂. This reaction is unique as one of the products, NO, catalysed the reaction.

Which graph best describes the above reaction?









19 The amount of carbon dioxide dissolved in a fizzy drink is affected by the following reversible reaction.

$$CO_2(g) + H_2O(l) \rightleftharpoons H_2CO_3(aq)$$

The reaction of carbon dioxide with water to reach equilibrium with carbonic acid has been shown to have first order kinetics with a half-life of 0.030 s.

What is the time taken for the carbonic acid to reach 75% of its equilibrium concentration?

- A 0.060 s
- **B** 0.180 s
- C 0.250 s
- D 0.414 s
- 20 Na₂S₂O₃ reacts with dilute HCl to form a pale yellow precipitate.

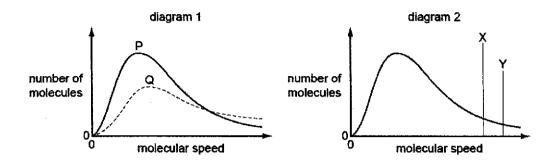
If 1 cm³ of 0.1 mol dm⁻³ HCl is added to 10 cm³ of 0.02 mol dm⁻³ Na₂S₂O₃, the precipitate forms slowly. If the experiment is repeated with 1 cm³ of 0.1 mol dm⁻³ HCl and 10 cm³ of 0.05 mol dm⁻³ Na₂S₂O₃, the precipitate forms faster.

What is a possible explanation for the observation above?

- A The activation energy of the reaction is lower when 0.05 mol dm⁻³ Na₂S₂O₃ is used.
- B The rate constant increases when 0.05 mol dm⁻³ Na₂S₂O₃ is used.
- C The reactant particles collide with more energy when 0.05 mol dm⁻³ Na₂S₂O₃ is used.
- ${f D}$ The reactant particles collide more frequently when 0.05 mol dm⁻³ Na₂S₂O₃ is used.

[Turn over

21 Different Maxwell-Boltzmann distribution curves are shown in the following diagrams.



In diagram 1, curve P or Q corresponds to a temperature higher than that of the other curve.

In diagram 2, line **X** or **Y** corresponds to the activation energy for a catalysed reaction and the other line corresponds to the activation energy of the same reaction when uncatalysed.

Which of the following statements are correct?

- 1 Using greater amount of catalyst does not further lower the activation energy.
- 2 The increase in temperature causes the curve in diagram 1 to change from P to Q.
- 3 Curve Q corresponds to the reaction occurring at a higher temperature while line X corresponds to a catalysed reaction.
- When the temperature is increased, the fraction of molecules that have energy more than the activation energy will be increased.
- A All of the above
- **B** 1, 2 and 3 only
- C 2 and 3 only
- D 2 and 4 only

22 Methanesulfonic acid is a non-volatile strong acid which is used to remove calcium carbonate from kettles.

Methanesulfonic acid

Which statement about methanesulfonic acid is incorrect?

- A The gas evolved when methanesulfonic acid reacts with calcium carbonate is CO₂.
- B 0.1 mol dm⁻³ methanesulfonic acid has a pH value of 1.
- C The Brønsted-Lowry conjugate base of methanesulfonic acid is the CH₃SO₃⁻ ion.
- D The K_a value of methanesulfonic acid is very small.
- 23 Which of the following will give an acidic buffer solution when dissolved in 1 dm3 of water?
 - 1 1 mole of CH₃COOH and 0.5 mole of NaOH
 - 2 1 mole of HCi and 1 mole of CH₃COONa
 - 3 1 mole of CH₃COOH and 1 mole of CH₃COONa
 - A 1 and 2 only
 - B 1 and 3 only
 - C 2 and 3 only
 - D 3 only

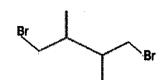
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24 A compound **Z**, upon mono-chlorination with chlorine in the presence of UV light, forms 3 possible structural isomers.

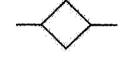
Which of the following compounds could be Z?

1

2



3



- A 1, 2 and 3
- **B** 1 and 3
- C 2 and 3
- D 1 only
- Different chlorinated pentanes were separately treated with hot ethanolic sodium hydroxide. Two of these gave the same hydroxarbon, C_5H_8 .

Which of the following pairs of chlorinated pentanes can form C₅H₈?

- B CH₃CH₂CH₂CH₂CH₂Cl and ClCH₂CH₂CH₂CH₂Cl
- C CH₃CH₂CHC*I*CH₂CH₃ and C*I*CH₂CH₂CH₂CH₂CHC*I*₂
- D CH₃CHC/CH₂CHC/CH₃ and C/CH₂CH₂CH₂CH₂CH₂CI

26 T can be converted to U as shown below.

Which of the following statements are correct?

- 1 Stage I may involve the use of sodium borohydride.
- 2 Stage I may involve the use of lithium aluminium hydride.
- 3 Stage II may involve the use of DCC and methylamine.
- A 1, 2 and 3
- **B** 1 and 3
- C 2 and 3
- **D** 1 only
- Which of the following products are formed when CH₃CH₂CO₂CH₂NHCOCH₂CH₃ is heated with excess aqueous sodium hydroxide?
 - A CH₃CH₂CO₂CH₂NH₂ and CH₃CH₂COOH
 - B CH₃CO₂CH₂NH₂ and CH₃CH₂COONa
 - C CH₃CH₂COOH and HOCH₂NH₂
 - D CH₃CH₂COONa and HOCH₂NH₂

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Turn over

28 Which of the following reactions forms an organic product that contains a deuterium atom? [Deuterium, D: ²H]

A

В

C

D

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[Turn over

BP~678

30 The repeat unit of a polymer is shown below.

Which deductions about this polymer can be made from the repeat unit?

- 1 It is a polyester.
- 2 It could be readily made from HOCH₂CH₂CH₂CO₂H.
- 3 There are hydrogen bonds formed between two polymer chains.
- A 1 only
- B 1 and 3 only
- C 2 and 3 only
- **D** 1, 2 and 3

SECTION A

Answer all the questions in the spaces provided.

Elements can be synthetically produced via nuclear reactions. 1 (a)

> During the process, protons or neutrons are injected 'like bullets' at the nucleus of an atom in making the synthetic element.

> A scientist attempts to produce 32P using a neutron and an isotope E of another element.

> > one neutron + isotope E $\rightarrow {}^{32}_{15}P + {}^{3}_{1}H$

(i)	Identify isotope E.	
		r

Draw the energy level diagram, showing the electronic configuration of $^{32}_{15}\mathrm{P}$ in (ii) the ground state.



[Turn over

(iii) Draw a clearly-labelled diagram to show the shape of one of the *singly-occupied* orbital in ³²P.

[1]

(iv) Describe one similarity and one difference between a 3s and a 3p orbital.

	Similarity	Difference
3s and 3p		
orbital		

[1]

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[1]

(b) Elements with an unstable atomic nucleus undergo radioactive decay.

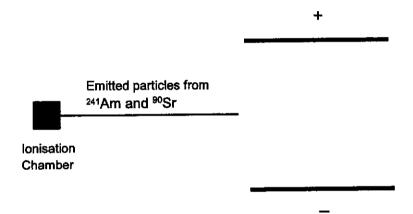
Americium–241 is used commonly in smoke detectors. It works by emitting a constant stream of α -particles, which are similar to the nucleus of Helium–4.

Strontium–90 is another radioactive substance. On decaying, it emits β -particles, which can be considered as electrons.

A small amount of Americium-241 and Strontium-90 are placed separately in an ionisation chamber to emit a constant stream of radiation and the emitted particles are passed through an electric field.

(i) On the diagram below, sketch the deflection path for the emitted particles in an electric field.

You may label the emitted particles from Americium–241 and Strontium–90 as α –particles and β –particles respectively.



(ii) Explain the difference in deflection path and angle respectively for the emitted particles in c(i).

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(iii) A deflection of -8° was observed when a beam of α -particles was passed through an electric field.

In another experiment, a beam of doubly charged particles \mathbf{X} was passed through the electric field under the same experimental conditions. The angle of deflection was found to be $+2^{\circ}$.

Identify the ion X. Show your working.

[2]

[Total: 9]

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[Turn Over

2	(a)	Boron and its compounds have many uses. Boron trichloride, BCl3 can be used in
	• •	refining process of aluminium. Boric acid, H ₃ BO ₃ sometimes known as B(OH) ₃ , is one
		of the most commonly produced borates and is widely used in the pharmaceutical and
		cosmetic industries.

Boric acid and borate salts such as sodium tetraborate, NaB(OH)₄, exist naturally in rocks, soil, plants and water.

(i)	With reference to the Valence Shell Electron Pair Repulsion Theory, explain the shape and bond angle with respect to the B atom in B(OH) ₃ .		
	,		
	[2]		

(ii) Draw the dot-and-cross diagrams for BCl₃ and B(OH)₄⁻. Clearly indicate any co-ordinate (dative covalent) bonds present.

[3]

Table 2.1 shows the melting points for boron compounds.

Table 2.1

Compound	Melting point / °C
Boron trichloride	-107.3
Boric acid	170.9
Sodium tetraborate	741

	(iii)	With reference to structure and bonding, explain the difference in melting points of the three componds in Table 2.1.
		[2]
(b)	(i)	BCl_3 can undergo hydrolysis to form boric acid and a gas that turns Universal Indicator red. Write an equation for this reaction forming $B(OH)_3$ from BCl_3 .
		[1]
	(ii)	Using appropriate bond energy values from the <i>Data Booklet</i> , and taking the bond energies for the B–C <i>l</i> bond and B–O bond to be 456 kJ mol ⁻¹ and 536 kJ mol ⁻¹ respectively, calculate the enthalpy change for the reaction in (b)(i) .
		(2)
	(iii)	[2]
	(111)	Suggest a reason why the enthalpy change calculated in (b)(ii) is only an approximate value.
		[1]

8873/TJC Prelim/2020

[Turn Over

(c) Boronic acid is an alkyl or aryl substituted boric acid.

Ethylboronic acid, CH₃CH₂B(OH)₂, has shown potency to be used as a drug in chemotherapy where certain proteolytic enzymes are blocked which would otherwise degrade proteins.

(i) The equation below represents the enthalpy change of combustion of ethylboronic acid.

 $aCH_3CH_2B(OH)_2(s) + bO_2(g) \longrightarrow cB_2O_3(s) + dCO_2(g) + eH_2O(l)$

Define the standard enthalpy change of combustion of ethylboronic acid and determine the values of a to e in the equation.

a :	 , b :	_, c :	_, d :	_, e:	
	 		**********************		**************
	 				[2]

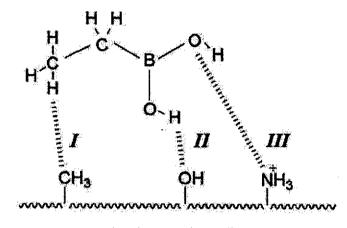
(ii) In an experiment, a scientist burns 1.5 g of ethylboronic acid with 2.5 dm³ of pure oxygen gas.

The reaction mixture is then allowed to cool to room temperature and pressure. Calculate the change in volume of gas, assuming ethylboronic acid is completely burned in the experiment.

(iii) For a drug to achieve the therapeutic effect, it must form interactions with the target active sites.

Ethylboronic acid forms interaction with certain proteolytic enzymes at the enzyme active site and blocks the enzymatic actions, which in turn achieves the chemotherapeutic effect.

The diagram below shows three interactions between ethylboronic acid and the enzyme active site.



Enzyme active site

Name all the interactions, I to III, shown in the diagram.

<i>I</i> :	••••••••		************		• • • • • • • • • • • • • • • • • • • •	• • • • • •
<i>II</i> :	***************************************			•••••	****************	
<i>III</i> :						

[3]

[Total: 18]

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[Turn Over

3 (a) A flameless ration heater (FRH) is a water activated chemical heater included with meals, ready-to-eat (MREs) used by the United States military.

The heater contains finely powdered magnesium metal alloyed with small amount of iron and sodium chloride. Water is added to activate the reaction.

$$Mg + 2H_2O \rightarrow Mg(OH)_2 + H_2$$

(i) The reaction between magnesium and excess water produces 6900 cm³ of hydrogen at s.t.p.

Calculate the mass of magnesium that has reacted.

(ii)	Suggest why sodium cannot be used in place of magnesium in the FRH.
	[1]

- (b) Exothermic reactions that do not produce hydrogen gas are being explored as alternatives in FRH. Some examples include the reaction of phosphorus(V) oxide and sodium oxide with water.
 - (i) Write an equation and give the approximate pH of the resulting solution when
 - phosphorus(V) oxide reacts with water
 - sodium oxide reacts with water

[2]

[1]

	(ii)	Sodium oxide reacts with phosphorus(V) oxide to produce sodium phosphate as the only product. Write a balanced equation for this reaction.
		[1]
	(iii)	Phosphorus(V) oxide differs in its bonding from that of sodium oxide, as reflected by the different ways in which they react with water.
		Explain why the difference in bonding arises between sodium oxide and $\operatorname{phosphorus}(V)$ oxide.
		······································
		•••••••••••••••••••••••••••••••••••••••
	-	
		[2]
	(iv)	The acid-base behaviour of beryllium oxide shows similarities to that of sodium oxide and phosphorus(V) oxide.
		Explain this behaviour of beryllium oxide.
		[2]
(c)	Mag	nesium oxide reacts reversibly with chlorine according to the following equation.
		$2MgO(s) + 2Cl_2(g) \Rightarrow 2MgCl_2(s) + O_2(g) \Delta H_r = -80 \text{ kJ mol}^{-1}$
	Und	er certain conditions, a dynamic equilibrium is established.
	(i)	Explain why magnesium oxide has a higher melting point than magnesium chloride.
		[1]
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(d)

(ii)		y change of formation of MgO is - 602 kJ mol ⁻¹ , cale formation of MgC l_2 (s).	culate
			[2]
		•	
(iii)	Predict and explain the at equilibrium.	effect of increasing temperature on the amount of o	xygen
			······
	•••••		
	*************************		[2]
iodio	ee test tubes were filled volume test tubes were filled volume. The test to th	with hydrogen chloride, hydrogen bromide and hydr A red-hot wire was placed into each test tube. in Table 3.1.	ogen The
		Table 3.1	
	Gas	Observation	
	hydrogen chloride	no observable change	
	hydrogen bromide	reddish brown vapour formed after some time	
	hydrogen iodide	purple fumes formed immediately	
	ervations.	nal stability of the hydrogen halldes, explain the a	
			•••••
		.,,	[2]
****	,		tal: 161

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4 Hydrazine, N₂H₄, is a popular choice of rocket fuels during outer space explorations. The stored hydrazine is passed over iridium catalyst and it decomposes to its elements. This rapid production of hot gaseous elements provides the thrust required. During the decomposition process, ammonia can also be formed as an intermediate.

The use of hydrazine as a fuel has been extended for use in aircraft as well. The first ever rocket-powered fighter plane was the Messerschmitt Me 163 Komet, The Komet was powered by the reaction between a hydrazine-methanol fuel mixture and oxidizer hydrogen peroxide. These reactants ignites spontaneously upon contact. Table 4.1 shows some properties about the fuels.

Table 4.1

	ΔH _c e / kJ mol ⁻¹	Density/ g cm ⁻³
N ₂ H ₄ (<i>I</i>)	-622.2	1.02
CH₃OH(<i>i</i>)	-726.0	0.792

(a) In the Komet, the hydrazine-methanol mixture reacts to propel the plane.
 A fully filled fighter plane carries 225 litres of hydrazine and 268 litres of methanol.

Using the data in Table 4.1, calculate the total heat evolved under standard conditions for combustion of this quantity of hydrazine-methanol mixture.

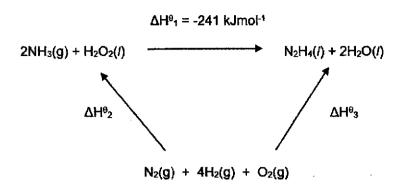
Assume that all the hydrazine and methanol are fully combusted.

[2]

(b)	Write a balanced equation for decomposition of hydrazine to form ammonia and nitrogen gas.
	[1]
(c)	Using the concept of activation energy and an appropriate sketch of the Boltzmann distribution, explain why the use of iridium catalyst speeds up the chemical reaction in (b) .

[2]

(d) An energy cycle involving hydrazine at 298K and 1bar is shown below.



Use the following data and the above energy cycle, calculate the standard enthalpy change of formation of hydrazine.

 $\Delta H^{\theta_{\rm f}}(NH_3) = -46.1~kJ~mol^{-1}, \\ \Delta H^{\theta_{\rm f}}(H_2O_2) = -187.8~kJ~mol^{-1}, \\ \Delta H^{\theta_{\rm c}}(H_2) = -285.8~kJ~mol^{-1}$

[2]

[Total: 7]

5

(a)	Ethanoyl chloride, CH ₃ COC <i>I</i> , reacts with ethanol to form ethyl ethanoate and hydrogen chloride.		
		CIOC COCI	
		arly, for , it will react with CH₂=CHCH₂OH to form compound P and ogen chloride.	
	Draw	the structure of P.	
(b)	P und	P [1] dergoes polymerization reaction to form polymer Q. Draw the structure of the polymer Q.	
		Q	
		[1]	
	(ii)	State the type of polymerisation for the formation of Q .	
		[1]	
	(iil)	Based on the structure drawn in (b)(i) , predict whether polymer Q is a thermosetting or a thermoplastic polymer. Explain your answer.	

		[2]	

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	(iv)	Suggest a use for polymer Q in consideration of its physical properties.
		[2]
(c)	CH ₂ =	=CHCH₂OH reacts with hot acidified K₂Cr₂O ₇ to form compound T .
	(i)	Draw the structure of compound T.
		T
	(ii)	Draw the structure of two repeat units of the polymer formed from compound T when mixed with potassium hydroxide in the presence of an initiator.
	(iii)	[1] The polymer formed in (c)(ii) has the capability to absorb <i>large</i> quantities of water. Explain why this is so.
		[1]

[Total:10]

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Turn Over

Section B

Answer one question from this section in the spaces provided.

(a) 0.10 mole of an organic compound C_xH_yO_z, gives 0.40 mole of carbon dioxide and 0.40 mole of water vapour upon complete combustion.
 8.8 g of C_xH_yO_z, when vaporised, was found to occupy 2.4 dm³ at room temperature and pressure.

Determine the molecular formula of this organic compound.

[3]

(b) Ethanoic acid, also commonly known as acetic acid, is an important chemical due to its varied uses in the industry.

The preferred industrial method for its manufacture is by carbonylation of methanol, CH₃OH accounting for 65% of the total world ethanoic acid, CH₃CO₂H manufacturing capacity.

In an experiment, the above process is carried out in a 2 dm³ reaction vessel. 3.40 moles of methanol and 2.75 moles of carbon monoxide are reacted at 200 °C and allowed to reach dynamic equilibrium. 1.20 mole of ethanoic acid is found to be present in the equilibrium mixture. Calculate the value for the equilibrium constant, K_{c_i} for the reaction. State its units.

c)		cm ³ of 3.00 mol dm ⁻³ ethanoic acid is added to 50.0 cm ³ of 1.40 mol dm ⁻³ lm hydroxide in a polystyrene cup. The maximum temperature rise recorded was °C.
	Give	n that the specific heat capacity of the solution is 4.18 J g ⁻¹ K^{-1} .
	(i)	Define what is meant by the standard enthalpy change of neutralisation.
		[1]
	(ii)	Write a balanced chemical equation for the neutralisation of ethanoic acid with barium hydroxide.
		[1]
	(iii)	Given that the neutralisation process was 70% efficient, calculate the standard enthalpy change of neutralisation for the reaction in (ii).
		· ·
		[3]
(d)		ered ethanoic acid mixes have been used as preservatives. To ensure its ctiveness, the ethanoic acid/sodium ethanoate buffer solution is maintained 16.
	(i)	What is meant by the term buffer solution?
		[1]

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(ii)	Explain, with the aid of equations, how the mixture can act as a buffer upon addition of a small amount of acid and alkali.
	[2]

(e) Explain, with the aid of a diagram, why the relative molecular mass of ethanoic acid is found to be 120 when dissolved in a suitable solvent.

[2]

(f) V is a fragment of polymer.

(i) Draw the structural formulae of the monomers found in V.

[1]

State the reagent and conditions that could be used to completely hydrolyse	V.
	[1]
ii) Name the two compounds that can be polymerised together to produce polyamide with four carbon atoms per repeat unit.	
v) State the condition required for the formation of the polyamide found in (f)(iii)	
	[1]
(Total:	201

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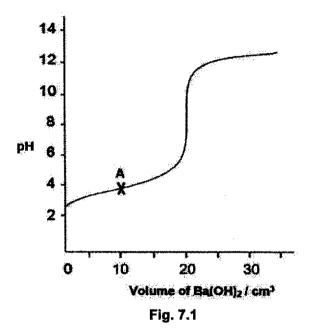
	21		
(a)	20 cm ³ of a hydride of nitrogen, N _x H _y is combut 10 cm ³ of nitrogen and 30 cm ³ of steam are hydride.	usted in an excess of oxygen at e formed. Deduce the formula	110 °C. for this
			19
(b)	Draw the two molecules, NH_3 and H_2O , and in each of the bonds it contains, and the overall	ndicate for each one the polarit polarity of the molecule.	
(b)	Draw the two molecules, NH ₃ and H ₂ O, and in each of the bonds it contains, and the overall NH ₃	ndicate for each one the polarit polarity of the molecule. H ₂ O	
(b)	each of the bonds it contains, and the overall	polarity of the molecule.	
(b)	each of the bonds it contains, and the overall	polarity of the molecule.	
(b)	each of the bonds it contains, and the overall	polarity of the molecule.	
(b)	each of the bonds it contains, and the overall	polarity of the molecule.	[2
(b)	each of the bonds it contains, and the overall	polarity of the molecule.	ty of
(b)	each of the bonds it contains, and the overall	polarity of the molecule. H₂O	

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.....[1]

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Fig. 7.1 shows the pH change that occurs when 15.0 cm 3 of 0.30 mol dm $^{-3}$ of lactic acid, CH₃CH(OH)COOH is titrated with barium hydroxide.



(ii) Write a balanced equation for the reaction between lactic acid and barium hydroxide.

[1]

(iii) Calculate the concentration of barium hydroxide used in this titration and its pH

value.

(iv) The solution at point A can act as a *buffer*.

Write a chemical equation to show how the solution at point A can act as a buffer when a small amount of H⁺ is added to it.

[1]

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Turn Over

[2]

(v) From the following list, suggest with reasoning, a suitable indicator to be used for the titration.

Indicators	pH range
Methyl violet	0 – 1
Methyl orange	3 – 4
Phenolphthalein	9 –10

••••	
	[2]

(d) The standard enthalpy change of neutralisation for the reaction in (c)(ii) can be determined experimentally by mixing known volumes of 1.0 mol dm⁻³ lactic acid and 1.0 mol dm⁻³ barium hydroxide. The following results are obtained.

Volume of lactic acid used = 40.0 cm³

Volume of barium hydroxide used = 20.0 cm³

Initial temperature of mixture = 26 °C

Final temperature of mixture = 34 °C

Specific heat capacity of the solution is 4.18 J g^{-1} K⁻¹.

(i) Use the data given to calculate the standard enthalpy change of neutralisation.

[2]

(ii) Using the value calculated in **d(i)**, draw a labelled energy profile diagram for the reaction, given that the activation energy is +5 kJ mol⁻¹.

[2]

(e) Two synthetic routes are shown below for converting C₁CH₂CH₂CHO into two different polymers.

Synthesis 1

CICH₂CH₂CHO

reaction 1

$$C_3H_4O$$

reaction 2

 C_3H_6O

reaction 3

 $C_3H_6O_3$

reaction 4

polymerisation

 $C_3H_6O_3$

polymerisation

one repeat unit

(i)	Name the reagents and conditions used in reactions 1 to 4.
	reaction 1:
	reaction 2:
	reaction 3:
	reaction 4:
	[4]

(ii) Draw the structure of compound X.

[1]

[Total:20]

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2020TJC JC2 H1 Chemistry Prelim MCQ Worked Solutions

_		_	-	_	
>	21	0	11	B	_
0	23	٨	12	0	2
80	23	C	13	0	3
A	24	8	14	A	4
D	25	c	15	-	O1
8	26	٥	16	>	6
0	27	B	17		7
>	28	>	18	>	~
C	28	>	19		40
	8	0	29		1

Question 1 Answer: B

no. of moles of Na⁺ and OH⁻ ions $= (0.5 \times 0.1) \times 2 = 0.10$ mol

no, of moles of molecules = 480 / 24000 = 0.02 mol no. of moles of IO_3^- lons = $(0.2 \times 0.5) \times 2 = 0.20$ mol

no. of moles of O atoms = (1.92 + 48.0) x 3 = 0.12 mol

Question 2 Answer: C

Option 1 is correct as the 2rd electron removed is experienced greater attraction.

Option 2 is incorrect because the lonic size is also dependent on the nuclear charge, i.e. the no. of protons. Isoelectronic only implies they have the same number of electrons but not the same no. of

Option 3 is incorrect. The two electrons are from 3d and 4s.

Option 4 is correct as the further electron removed from AI is from inner quantum shell

Question 3 Answer: C

Using the iE values, the greatest jump is between $5^{\rm th}$ and $6^{\rm th}$ IE. Hence, E belongs to Group 15. E is likely to be P.

Question 4 Answer: A

Let average O.N of S be x.

2(+1) + 4x + 6(-2) = 0

×# +2.5

Option A is correct as the average O.N of S = (0 + 0 + 5 + 5) / 4 = +2.5

Alternative - calculate O.N of each S (refer to Redox notes page 6)

O.N of S1 = O.N of S4

(± + (†

ţ

(S-O bond) (S-S bond) (S=O bond) (S=O bond)

 $O.N \text{ of } S^2 = O.N \text{ of } S^3 = 0 \text{ (S-S bond)}$

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Question 5 Answer: D

Amount of electrons gained by 1 mole of As₂O₃

= 6 x 2 = 12 mol

amount of electrons lost by Zn

Since each mole of Zn loses 2 moles of electrons, amount of Zn needed to react with 1 mole of As₂O₃

m 12/2

= 6 mo!

Question 6

Answer: A

Answer: B

C-1 and C-3 are unsaturated with sp² hybridisation. C-2 is sp³ hybridised, C-4 is -C≡N so it is a sp C atom

BH₃ + NaH NaBH₄ (120° to 109°)

CO2 + H2O === 2H" + CO32" (180° to 120°)

Question 10 Answer: B

$KCl(s) \rightarrow K(g) + Cl(g)$	$K^*(g) + CF(g) \rightarrow K(s) + \frac{1}{2}CI_2(g)$	K(s) + ½C/₂(g) → KC/(s)
ΔH = +710 + (-63) = +647 kJ mol ⁻¹	$K^*(g) + CF(g) \Rightarrow K(s) + \frac{1}{2}CI_2(g)$ $\triangle H = -63 - (+211) = -274 \text{ kJ mo}^{-1}$	$\Delta H = +211 - (-63) - (+710) = -436 \text{ kJ mol}^{-1}$

Question 11 Answer: D

Enthalpy change of neutralisation and combustion are always negative. Enthalpy change of formation could be negative or positive.

= +410 kJ mol-1 --160 kJ mol-1 = -340 kJ mot1 ∆H when C-Cl bond is formed From data booklet, AH when N-N bond is formed AH when C-H bond is broken Question 12 Answer: A

Question 13

(hydrolyses only to a small extent in water) $[Mg(H_2O)_6]^{2*} \rightleftharpoons [Mg(H_2O)_5(OH)]^{2*} + H^*$ $MgC_{I_2} + 6H_2O \rightarrow [Mg(H_2O)_6]^{2+} + 2CF$

A/CI₃ + 3H₂O → A/(OH)₃ + 3HCI

PCIs + 4H₂O → POCIs + 2HCI

1.0 mol

SIC/4 + 2H2O → SIO2 + 4HC/

0.5 mol

Answer: B Question 14 Bo∰ng point increases down the group due to increasing electron cloud size.

From the data booklet, BE (F-F) = 158 kJ moi⁻¹, BE (CI-CI) = 244 kJ moi⁻¹, BE (Br-Br) = 193 kJ moi⁻¹ and BE第-1) = 151 kJ mol-1.

Oxigising power of halogens decreases down the group due to increasing atomic radius.

intensity of colour increases down the Group (see Data Booklet for colour).

Question 15 Answer: C

SI has the highest meiting point due to its giant covalent structure with numerous strong covalent bordes between atoms.

Cr≦nas the smallest lonic radius among anions P⁵- to Ci⁻ due to its highest nuclear charge.

Question 16

As the reaction is exothermic, an increased in the temperature at t₁ will favour the endothermic reaction, position of equilibrium shifts to the left. At t₂, the pressure of the vessel is increased which shift the position of equilibrium to the right to reduce the number of gaseous molecules.

Auestion 17

Option 3 is incorrect as the equilibrium concentration will increase until it reaches the new equilibrium is Option 1 is incorrect as the equilibrium constant will change when temperature change.

Question 18 Answer: A

Reaction rate of an autocatalysis reaction will be fow initially, high in between, and low eventually because, the product, NO formed in the reaction acts as a catalyst Tangent of a concentration-time graph gives reaction rate. Tangent is low initially, high in between, and low eventually.

Faster rate because auto-catalyst is now present Rate slow again as NO2 is depleted /ery slow rate because auto-catalyst not yet formed ime / s \[\bar{1}^2\] mol dm

For this shape of graph to correctly represent an autocatalysis reaction - the correct axis abe! should be rate vs time. 00

not an This rate-concentration graph is typical of an enzyme-substrate reaction, autocatalysis reaction. ပ

This rate-time graph is increasing throughout, not an autocatalysis reaction. ۵

Question 19

Analysis: When H2CO3 reaches 75% equilibrium conc, only 25% CO2 is left

0.25= 1/2

Let n = no of half-lifes for CO₂ to drop to 1%

n=2

Time taken = $2 \times 0.03 = 0.060 \text{ s}$

Question 20 Answer: D

When higher [Na₂S₂O₃] is used, there is greater no of reactant molecules having energy > or α Ea hence frequency of collisions increases leading to faster rate.

Question 21 Answer: A

Option1: is correct as the use of catalyst does lower the activation energy, but increasing the amount of the catalyst does not make it even lower.

Option 2: is correct as the shape of the graph will shift towards the right (showing more molecules

having more energy) when temperature rises

reaction occurring at a higher variance of number or recommendates. When temperature is increased, number or results in the peak of curve P shifting rightwards to give curve Q.

Option 4: Is correct. When temperature increases, the rate of reaction will increase as the fraction of molecules to that have energy more than the activation energy will increase. Option 3: is correct as the use of catalyst lower the activation energy from Y to X. Curve Q corresponds to the

Ġ

For strong acid, because it is fully dissociated in solution, its Ka value will be a large value

Question 22 Answer: D

Question 23 Answer: B

Option 1: excess weak acid CH₂COOH and salt CH₂COONa present Acidic buffer includes a weak acid and its sait

Option 2: no weak acid present

Option 3: weak acid - CH₃COOH and salt - CH₃COONa present

Question 24 Answer: A

Question 25 Answer: D

Hydrocarbon, C₂H₈ hence it is a diene (with 2 C=C)

Hence there must be 2 elimination to form 2 C≖C

CH₂CHC/CH₂CHC/CH₃ → CH₂=CHCH₂CH=CH₂

CICH2CH2CH2CH2CH2CH2CH → CH2=CHCH2CH=CH2

Question 26 Answer:B

Option 2 is incorrect as lithium aluminium hydride will reduce the carboxylic acid to primary alcohol

Question 27 Answer: D

Ester and amide undergo basic hydrolysis simultaneously to form CH₂CH₂COONa and HOCH₂NH₂.

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Question 28

0

A) substitution gives CH₃CH₂OD.

B) oxidation to give CH₃CO₂:

C) Elimination to give CH2=CH2

D) Condensation to give CH₃CO₂CH:

Question 29

and.

Carboxylic acid functional group will condense with the alcohol group of the monomers to give the polymer.

Question 30 Answer: B

1: An ester linkage is formed when you join the repeat units together.

2: It is made from HOCH(CH₃)CH(OH)CO₂H.

3: This polyester has OH groups hence hydrogen bonding can be formed between polymers chains. where the page of the pa

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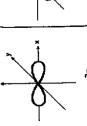
SECTION A

Answer all the questions in the spaces provided.

- 統 €
- <u>a</u>
- P: 1s² 2s² 2p⁸ 3s² 3p³ Energy / kJ €
- **≓**|& **=|**#
 - =|=

 $\widehat{\mathbf{E}}$

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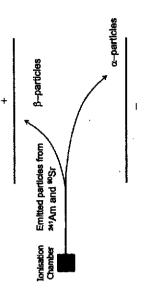


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Both 3s and 3p orbitals belong to the same principal quantum shell. Similarity 3s and 3p orbital

3s orbital has spherical shape white 3p orbital has dumb-bell shape.



R

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E

Analysis : a-particles = 4He2* ; β-particles = 0echarge for 4He2" = 2 €

 $\frac{\text{charge}}{\text{mass}} \text{ for e} = \frac{1}{1/1836}$

magnitude due to greater mass/charge ratio (or show numerical $\frac{q}{m}$) than a-particles will be deflected towards the negative plate and by a smaller

β-particles, which is towards positive plate and by a larger magnitude.

Since X is doubly charged and the deflection is towards the positive plate E

"ratio for α -particles (⁴He^{2*}) nuclei $= \frac{2}{4} = \frac{1}{2}$ g ratio for X2 = - 2

 $+2^{\circ} = (-\frac{2}{m} + \frac{1}{2}) \times -8^{\circ} \Rightarrow m = 16 \Rightarrow X \text{ is } O^{2^{\circ}}$

There are 3 bond pairs and no lone pairs around B atom. <u>To minimise</u> repuision and <u>maximize stability,</u> the 3 electron pairs are directed in a trigonal planar manner. Hence, shape is <u>trigonal planar</u> with <u>bond angle</u> 120°. €

N

[Total: 9]



 Sodium tetraborate has a glant lonic lattice structure, while boron trichtoride and boric acid are simple molecular structures.

€

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More energy is required to overcome the stronger ionic bonds between oppositely charge ions in sodium tetraborate than the weaker hydrogen higher melting point bonds between boric acid molecules, hence sodium tetraborate has a

3 BC/₃ + 3H₂O → B(OH)₃ + 3HC/

9

€

宁 Bonds broken: B-Cl : 3 (456) 6 (460) Bonds formed: B-O 3 (536)

: 3 (460) : 3 (431)

Total energy required: 4128 kJ mg/-1 Total energy released: 4281 kJ moi-1

 $\Delta H = \Sigma$ Bond energies of reactants – Σ Bond energies of products = +4128 - 4281

= - 153 kJ mol-1

3 It is an approximation as the bond energies from the Data Booklet are only average values and would not apply to the exact molecules in the reaction.

Bond energies are for breaking of bonds in the gaseous phase phase under standard conditions. However the substances in the reaction are not all in the gaseous

3 the energy released when 1 mole of ethylboronic acid is completel burnt in excess oxygen under standard conditions of 298K and 1 bar. The standard enthalpy change of combustion of ethylboronic acid is

0

a=1, b=7/2, c=1/2, d=2 and e=7/2

 \equiv Change in volume = Total vol of O₂ - [Vol of O₂ unreacted+ Vol CO₂] •Volume of O_2 reacted at rtp = 7/2 x 0.0203 x 24 = 1.71 dm³ Volume of CO₂ produced $= 2 \times 0.0203 \times 24 = 0.976 \text{ dm}^3$ Volume of unreacted $O_2 = 2.5 - 1.71 = 0.793 \text{ dm}$ •Decrease in volume of gas = $2.5 - [0.793 + 0.976] = 0.731 \text{ dm}^3$ Number of moles of ethylboronic acid = 1.5 / 73.8 = 0.0203 mol

3 I: instantaneous dipole-induced dipole II: hydrogen bond III: ion-dipole interactions

[Total: 18]

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3

number of moles of $H_2 = 6 900/22700 = 0.304$ mol

mass of Mg = $0.304 \times 24.3 = 7.39$ g

€ The reaction will be unsafe as sodium reacts explosively/ vigorously

be stored easily as sodium metal in the FRH. Sodium reacts readily with oxygen to form sodium oxide, so it cannot

3 P4O10(s) + 6H2O(l) -> 4H2PO4(aq)

3

- Na₂O(s) + H₂O(l) → 2NaOH(aq) PH=2
- 6Na₂O + P₄O₁₀ → 4Na₃PO,

3 3

- There is a great electronegativity difference between Na and O, hence Na₂O is an ionic compound.
- There is small electronegativity difference between P and O, hence P₄O₁₉ is a covalent compound.
- ₹ O2- lons to be mainly lonic with some covalent characteristics. BeO is an amphoteric oxide due to the high charge density of the Be2+ ion polarising the O2 anion causing the bonds between the Be2
- 3 Lattice energy of MgO is higher due to the higher charge of Q2 compared magnesium oxide. to CI. More energy is needed to overcome the stronger lonic bonds

 $\Delta H_r = \Sigma \Delta H_r$ products $-\Sigma \Delta H_r$ reactants ΔH₁ (MgC½) = - 642 kJ mol⁻¹ $80 = 2 \times \Delta H_1 (MgCh_2) - 2 \times (-802)$

€

- \equiv By Le Chateller's Principle, the system will counteract the increase Position of equilibrium will shift to the left and amount of oxygen will be temperature by favouring the endothermic reaction to remove heat. lower at equilibrium. 3
- ٥ and was able to decompose HI readily to form purple I2 furnes immediately. able to decompose some HBr to form reddish brown Br2 vapour after some time, Covalent band strength and hence bond energy of H-X decreases from H-C/ to H–Br to H⊣i. Hence thermal stability of HX decreases from HC/ to HBr to HI. The energy supplied by the red-hot wire was insufficient to decompose HC/, was [Total: 16]
- 1 Amount of hydrazine = 225000 x 1.02/32.0 = 7172 mol Total heat evolved = $7172 \times 622.2 + 6633 \times 726.0 = 9.28 \times 10^6$ Amount of methanol = $268000 \times 0.792/32.0 = 6633 \text{ mol}$
- 3N₂H₄ → N₂ + 4NH₅

3

9 In the presence of a catalyst, the number of reactant molecules having energy greater Hence the frequency of effective collisions increases and the rate increases han or equal to the lower activation energy. E. increases significantly.

Es.; activation energy for No. of molecules having energy ≥ Es No. of molecules having energy 2 Es.

No of molecules

Unicatialysed reaction

between the polymer chain and H₂O molecules enable polymer to absorb

large quantities of water.

Presence of hydrogen bonding and lon-dipole interactions [1]

Ê

€

E ... activation energy for catalysed reaction

By Hess' law, €

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 $-241 = \Delta H^{9}_{1}(N_{2}H_{4}) + 2(-285.8) - [2(-46.1) + (-187.8)]$ $\Delta H^{0}_{t}(N_{2}H_{4}) = -241 + 2(285.8) + [2(-46.1) + (-187.8)]$ = $+50.6 \text{ kJ mol}^{-1}$. AH⁶, = AH⁹3 - AH⁶2

Total:10]

Hydrogen bonding

[Total: 7]

Ξ

<u>s</u>

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Addition polymerisation

€

Ξ

- Q is a thermosetting polymer because there are extensive strong covalent bonds and cross-links between polymer chains. €
- components. It can retain its insulating properties, even when subjected Q is used for high-performance military and commercial electrical to high heat and high humidity over long time periods. Ξ

0-0-x r-0-r €

3

Section B

Answer one question from this section in the spaces provided

(a)
$$C_xH_yO_x(1) + \left(x + \frac{y}{4} - \frac{z}{2}\right)O_2(g) \Rightarrow$$

0.10 mol

次O2(g) $\frac{y}{2}$ H₂O(g)

3

=

A buffer solution is one which is capable of maintaining a fairly constant pH (by

= 53300 J mol⁻¹ or -53.3 kJ mol⁻¹

=-53316 =-7464 / 0.14

resisting pH change) when small amounts of acid or base are added to it.

3

When a small amount of H+ is added

CH3COOT + H* → CH3COOH

0,40 mol

0.40 mol

Amount of **A** at r.t.p = $\frac{2.4}{24.0}$ = 0.100 mol Molecular formula of $A = C_4H_8O_2$ Comparing mole ratio, x = 4, y = 8

 M_r of A = 88.0Molar mass of $A = \frac{8.8}{0.100} = 88.0 \text{ g mor}^4$

To solve for the value of z:

Ē

molecular formula of A Is C₄H₈O₂

4(12) + 8(1) + z(16) = 88

◉

Dimerisation has occurred between 2 molecules of ethanoic acid

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[OH] is slightly changed and pH of buffer solution remains fairly constant

The OH' is removed by CH3COOH

CH3COOH + OH → CH3COO + H2O When a small amount of OH- is added [H*] is slightly changed and pH of buffer solution remains fairly constant.

The H* is removed by CH₃COOT

	СН ₃ ОН(аq)	+	CO(g)	1	CH ₃ CO ₂ H(aq)
Initial no. of mol	3.40		2.75	0.	0
Change in no. of mol	-1.20		-1.20	0	+1.20
Eqm no. of	2.20		1.55		1.20

 $K_{\rm o} = (1.20/2) / [(2.20/2)(1.55/2)] = 0.704 \text{ mol}^{-1} \text{dm}^{-3}$

- O 3 Standard enthalpy change of neutralisation (ΔH,*) is the heat evolved when under standard conditions of 298K and 1 bar one mole of water is formed from the neutralisation between acid and alkal
- \equiv $2CH_3COOH(aq) + Ba(OH)_2(aq) \longrightarrow (CH_3COO-)_2Ba^{2+}(aq) + 2H_2O(l)$
- 2CH₃COOH (aq) + Ba(OH)₂ (aq) -----> (CH₃COOT)₂Ba²⁺(aq) + 2H₂O (f)

 \equiv

⇒ Ba(OH)₂ is the limiting reagent. Heat evolved = mc\DT

ΔH,0 = -Q

No. of moles of $CH_3COOH = 50/1000 \times 3.00 = 0.150$ mol

No. of moles of Ba(OH)₂ = 50/1000 x 1.4 = 0.0700 mol

= 7464J

Actual heat evolved = 5225/70 x 100

= 100 x 4.18 x 12.5

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3

3

Heat with aqueous NaOH/KOH/OH OR Heat with Aqueous HCI/H2SO4/H*

3

3

Ethanedioic acid & ethane-1,2-diamine OR Ethanedioyl chloride & ethane-1,2-diamine

3 DCC

[Total:20]

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[Total:20]

AH, 8 = -2006/0.0400 = -50.2 kJ mol-1 $= (4.18 \times 60.0) \times (34 - 26) = 2006 \text{ J}$ Heat evolved Q = mcAT €

Energy to mo! €

reaction 3: K₂Cr₂O₇, dilute H₂SO₄, heat or KMnO₄, dilute H₂SO₄, heat reaction 2: LIAM, in dry ether, rtp or NaBH, in ethanol, rtp reaction 1: ethanolic NaOH, heat €

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Comparing the coefficient of NxHy and H2O

X =]

 $\frac{1}{y} = \frac{20}{30}$

reaction 4: aqueous NaOH; heat

€

\$

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 $\frac{x}{2}N_2 + \frac{y}{2}H_2O$

 $N_xH_y + \frac{y}{4}O_2 \rightarrow$

Comparing the coefficient of N_xH_y and N₂

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Change in vol/cm3

<u>e</u>

The formula is NH3.

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Shape: Bent O-H bond is polar Shape: Trigonal pyramidal

A Bronsted Lowry base is a proton acceptor. \in

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There is an overall dipole moment and NH3 is polar.

There is an overall dipole moment and H₂O is polar.

€

 $n(lactic acid) = 0.0150 \times 0.30 = 0.00450 \text{ mol}$ $n[(Ba(OH)_2] = 0.00450/2 \text{ mol}$ €

= 0.00225 mol

Ba(OH)₂→ Ba²+ + 2OH-[Ba(OH)₂] ≡ 2[OH-] since barlum hydroxide is a strong base [Ba(OH)₂] = 0.00225/0.020 = 0.1125 mol dm⁻³

[OH*] = 0.225 mol dm⁻³

pOH = -log₁₀ [OH⁻] = 0.648 PH = 14 - 0.648 = 13.4

(w) CH3CH(OH)COO- + H+ → CH3CH(OH)COOH

Phenolphthalein Σ

Its working range of 9-10 lies within the long vertical part of the pH curve.

Partition List Finding 12020

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