### Answer **all** the questions.

	Steel pipelines carrying natural gas occasionally corrode leading to the formation of 'green rust', Fe(OH) <sub>2</sub> .		
(a)	Write equations or half-equations to explain how green rust forms on steel.		
	In each case,	name the <b>type</b> of reaction involved.	
	equation 1	Give a half-equation for the formation of the Fe <sup>2+</sup> ions.	
	equation:		
	reaction type		
	equation 2	Give a half-equation for the formation of hydroxide ions.	
	equation:		
	reaction type		
	equation 3	Give an ionic equation for the formation of green rust.	
	equation:		
	reaction type	[6]	
(b)	Steel usually	corrodes to form 'red-brown' rust.	
	(i) Give the	oxidation state of Fe in red-brown rust.	
		[1]	
	(ii) Suggest steel pipe	why, in some conditions, green rust rather than red-brown rust forms on buried elines.	
		[1]	
(c)	Green rust of compound.	n a steel surface can be removed by converting the rust into a soluble iron	
	One simple m	nethod is to rub the surface with sulfuric acid.	
	Give the syste	ematic <b>name</b> of the soluble iron compound formed from green rust.	
		[1]	

(d) Another method used to remove green rust is to convert the rust into a soluble complex ion.

 ${\rm Fe}({\rm OH})_2$  will form a soluble complex ion with 'citrate' ions. Citrate ions are polydentate ligands. The structural formula of a citrate ion is shown below.

citrate ion ( $C_6H_5O_7^{3-}$ )

(i)	Explain the meaning of the terms complex, ligand and polydentate.	
	complex	
	ligand	
	polydentate	
		[4]
(ii)	The ratio of iron ion to citrate ion in the complex ion is 1:1.  The iron has a coordination number of 3.	
	Give the formula and charge of the complex ion formed.	

[2]

(e)	After some time, green rust can change into 'black rust'. The equation for this reaction is given
	below.

$$\label{eq:SFeOH} \mathrm{3Fe(OH)}_2 \longrightarrow \mathrm{(Fe^{2+})(Fe^{3+})}_2\mathrm{O_4} \ + \ \mathrm{H_2} \ + \ 2\mathrm{H_2O}$$
   
 
$$\mathbf{black} \ \mathbf{rust}$$

	(i)	Use <b>oxidation states</b> to determine what has been oxidised and reduced.
		is oxidised because
		and is reduced because
		[4]
	(ii)	Calculate the volume of hydrogen gas produced when 100g of green rust changes into black rust.
		Assume that 1.0 mole of gas occupies 24 dm <sup>3</sup> at room temperature and pressure.
		volume of hydrogen = dm <sup>3</sup> [2]
(f)	If se	eawater is present, the green rust is often oxidised to $[{\rm FeC} l_4]^-$ ions.
	Dra	w diagrams to show <b>two</b> possible shapes for $[FeCl_4]^-$ .

[2]

[Total: 23]

2 Oust is a common descaling treatment for kettles. Its main ingredient is the acid A, C<sub>3</sub>H<sub>6</sub>O<sub>3</sub>.

(a) Give the systematic name for acid A.

.....[2]

**(b)** A sachet of *Oust* contains 25.0 g of a liquid. The information on the packet indicates that the only acid present is acid **A** and that its content is 30–50% by mass.

Some students decide to investigate the percentage of acid  $\bf A$  in *Oust*. They dilute the liquid from one sachet to  $100\,{\rm cm}^3$  with water.  $25.0\,{\rm cm}^3$  of this solution reacts exactly with  $33.6\,{\rm cm}^3$  of  $1.00\,{\rm mol\,dm}^{-3}$  sodium hydroxide solution.

Calculate the actual percentage by mass of acid A in Oust.

Give your answer to an appropriate number of significant figures.

percentage of acid **A** in *Oust* = ...... % [5]

Turn over

(c) The students use indicator **B** to determine the end point of their titration in (b). They find that the indicator used for the titration is pink in very dilute alkali, but that the colour slowly fades in more concentrated alkali.

The slow reaction of indicator **B** in alkali can be represented as:

$$B(aq) + OH^-(aq) \rightarrow BOH^-(aq)$$

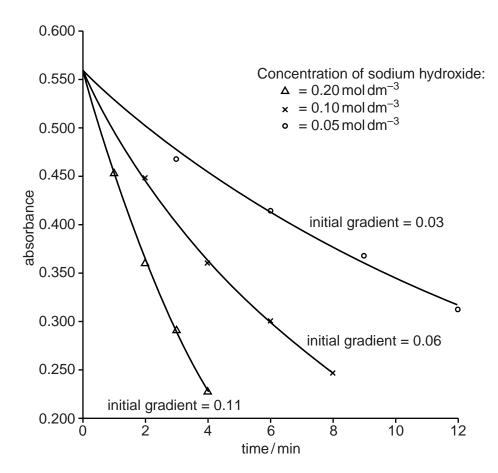
The students decide to follow this reaction by measuring the absorbance in a colorimeter.

They are provided with:

- three aqueous solutions of sodium hydroxide with different concentrations
- a suitable pink solution of indicator B.

They carry out experiments with the three sodium hydroxide solutions.

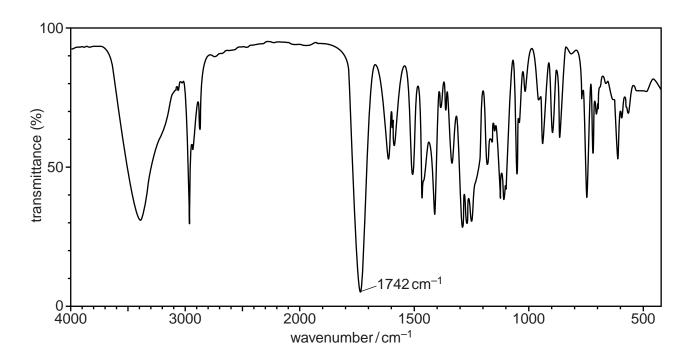
The results are shown on the graph and the magnitude of each initial gradient is given.



	Des	scribe how the students could have carried out the experiments to obtain these results.
		[4]
(d)	The	absorbance is proportional to the concentration of <b>B</b> .
	(i)	How would the students confirm that the absorbance was proportional to the concentration of <b>B</b> ?
		[3]
(	(ii)	Use the data on the graph to determine the order of reaction with respect to hydroxide ion.
		Order is because
		[2]
(	iii)	The students then set out to determine the order with respect to <b>B</b> .
		How would the students do this?
		[2]

**(e)** Indicator **B** is an organic compound which gives a purple colour with neutral iron(III) chloride solution.

The infrared spectrum of indicator **B** is shown below.



(i)	Use the spectrum to identify <b>two</b> functional groups present in indicator <b>B</b> .
	Give your reasoning.
	[3]
(ii)	Describe how the reaction of <b>B</b> with neutral iron(III) chloride solution gives more information about one of these groups.
	[1]

/£\	۸ م: ۵ ۸			4:440 40 4	010 400 010 0 400
(T)	ACIG A	can exist	as two	amerent	stereoisomers.

	Exp ther	lain why $f A$ can exist as two different stereoisomers and describe the relationship between m.
	In y	our answer, you should use technical terms, spelled correctly.
		[2]
(g)		d <b>A</b> is made from glucose and can be polymerised to form a thermoplastic material which be used in food packaging.
	Enz	rymes in bacteria present in soil will hydrolyse the polymer.
	(i)	Explain why acid <b>A</b> can be polymerised and name the functional group which joins the monomer units together.
		explanation
		functional group
		[2]
	(ii)	Give <b>two</b> advantages of using this polymer in the production of food packaging.
		[2]

(h) Enzymes present in the soil can also be hydrolysed. One such product of hydrolysis is compound **C**.

Give structural formulae for the ions that compound  ${\bf C}$  will form in highly acidic soil and in highly alkaline soil.

acidic soil:

alkaline soil:

[3]

[Total: 31]

3 Some engineering plastics contain heat-resistant polymers and are used to make parts for machinery. The table below shows some details for two heat-resistant polymers.

Name of polymer	Structure of repeating unit	T <sub>g</sub> /°C	Arrangement of chains
PEEK		143	ordered
PPO	<del></del>	215	tangled

(a)	Name <b>two</b> different functional groups, other than a benzene ring, present in PEEK.
	and

(b) PPO can be made by reacting compound  ${\bf D}$  with  ${\bf O}_2$  in a 2:1 mole ratio.

compound D

(i)	Name the other product of the reaction that forms PPO.	
		[1]
(ii)	State the <b>type</b> of polymerisation reaction in which PPO is formed.	
		[1]

(iii)	Suggest why the value of $T_{\rm g}$ differs for PPO and PEEK.	
	In your a	answer:
	•	state what is meant by $T_{\rm g}$
	•	describe ${f and}$ explain what happens when a polymer is heated above its ${\cal T}_{{f g}}$
	•	suggest why, in terms of the arrangement of chains, $\textit{T}_{\rm g}$ for PPO is higher than that for PEEK.

(c) Polyaramids are also heat-resistant polymers.

The polyaramid X-Fiper™ is formed from the monomer shown below.

$$\mathsf{H_2N} - \bigcirc \mathsf{COC} \mathit{l}$$

(i) Draw the repeating unit for X-Fiper™.

[1]

(ii) Circle **one** of the following terms which best describes the reaction that forms X-Fiper<sup>™</sup> from its monomer.

acid-base acylation addition substitution [1]

(d) Twaron™ is another polyaramid with the repeating unit shown below.

(i) Name the functional group linking arene units together in Twaron™.		
	[1]	
/ii\	Garments made with Twaron™ are hullet-proof, unlike those made from X-Finer™. This	

(ii) Garments made with Twaron<sup>™</sup> are bullet-proof, unlike those made from X-Fiper<sup>™</sup>. This is because the chains of Twaron<sup>™</sup> are straighter.

the polymer chains.

Explain this difference in garment strength in terms of intermolecular bonding between

[Total: 14]

4	A group of researchers has discovered a type of bacterium on the shores of a lake in California.
	The lake contains one of the highest natural concentrations of arsenic in the world.

This led them to consider that bacteria may be able to incorporate arsenic into biomolecules in place of phosphorus. They claimed that they had isolated DNA containing arsenic instead of phosphorus in the 'phosphate' sugar backbone.

(a)	(i)	Give a chemical reason why arsenic might be able to replace phosphorus.
		[1]

(ii) The structural formula for the dihydrogenar senate ion,  $\rm H_2AsO_4^-$  is shown below.

Draw a 'dot-and-cross' diagram for  ${\rm H_2AsO_4^-}.$ 

Show outer electrons only.

### As

(iii)	Use the electron pair repulsion theory to name the shape of $\rm H_2AsO_4^-$ and give an approximate bond angle for an O–As–O bond.
	Explain your answer.
	In your answer, you should indicate how the points that you make link together.

	9	,	,	, ,	,
[4					

[2]

	(iv)	Name the <b>type</b> of reaction which occurs if H <sub>2</sub> AsO <sub>4</sub> <sup>-</sup> reacts with deoxyribose to for backbone for DNA.	m a
			. [1]
	(v)	In this question you should refer to the information provided in the Data sheet.	
		Draw the structure of the ion formed when a $H_2AsO_4^-$ ion reacts with a <b>primary</b> hydrogroup of a deoxyribose molecule.	oxyl
			[3]
(b)	H <sub>2</sub> A	AsO <sub>4</sub> exists in weakly acidic solutions.	
	(i)	Suggest a formula for the arsenic compound formed from ${\rm H_2AsO_4^-}$ in strongly adsolutions.	cidic
			. [1]
	(ii)	In a strongly alkaline solution, OH <sup>-</sup> ions remove two protons from H <sub>2</sub> AsO <sub>4</sub> <sup>-</sup> .	
		Write an ionic equation for the reaction.	
	(iii)	Circle <b>two</b> proton acceptors in your equation in (ii).	[2] [1]

**(c)** Many other scientists have concerns about the validity of the evidence provided by the researchers.

One concern is that arsenates are more reactive than phosphates. The arsenate-sugar backbone in DNA would break down very easily in the presence of sulfur compounds in the environment.

Half-reaction	E <sup>⊕</sup> /V
$H_3PO_4(aq) + 2H^+(aq) + 2e^- \rightarrow H_3PO_3(aq) + H_2O(l)$	-0.28
$SO_4^{2-}(aq) + 4H^+(aq) + 2e^- \rightarrow SO_2(aq) + 2H_2O(l)$	+0.17
$H_3AsO_4(aq) + 2H^+(aq) + 2e^- \rightarrow H_3AsO_3(aq) + H_2O(I)$	+0.56

	(i)	Use the data in the table to show why $\rm H_3AsO_4$ will react with sulfur dioxide but $\rm H_3PO_4$ will not.	
		[2]	
	(ii)	Write the equation for the reaction between ${\rm H_3AsO_4}$ and sulfur dioxide in acid solution.	
		[2]	
(d) Another concern about the existence of arsenic-containing DNA is that the arsenate backbone would be quickly hydrolysed in the body. The arguments are based on the re of As–O bonds in <b>small</b> organic molecules. Scientists have estimated a constant hal 10 minutes for the hydrolysis of As–O bonds.			
	(i)	State the order, with respect to the arsenic compound, for the hydrolysis reaction.	
		Explain your answer.	
		[2]	
	(ii)	Suggest one reason why some scientists might disagree that the arsenate-sugar backbone in DNA will hydrolyse easily.	
		[1]	

**END OF QUESTION PAPER** 

[Total: 22]