Q1. Where appropriate, use the standard electrode potential data in the table below to answer the questions which follow.

							E°N
Zn²⁺(aq)	+	2e⁻	\rightarrow	Zn(s)			-0.76
V³⁺(aq)	+	e-	\rightarrow	V²⁺(aq)			-0.26
^{SO} 4 ^{2− (aq)} + 2H·(aq)	+	2e [.]	\rightarrow	$SO_3^{2-}(aq)$	+	$H_2O(I)$	+0.17
VO²⁺(aq) +2H⁺(aq)	+	e-	\rightarrow	V³⁺(aq)	+	$H_2O(I)$	+0.34
Fe³⁺(aq)	+	e⁻	\rightarrow	Fe²⁺(aq)			+0.77
^{VO} 2 ^(aq) + 2H·(aq)	+	e-	\rightarrow	VO²⁺(aq)	+	$H_2O(I)$	+1.00
Cl₂(aq)	+	2e-	\rightarrow	2Cl⁻(aq)			+1.36

(a)	From the table above select the species which is the most powerful reducing agent.	
		(1)

- (b) From the table above select
 - (i) a species which, in acidic solution, will reduce $VO_2^+(aq)$ to $VO_2^+(aq)$ but will **not** reduce $VO_2^+(aq)$ to $V^{3+}(aq)$,
 - (ii) a species which, in acidic solution, will oxidise VO²⁺(aq) to VO²⁺(aq).
- (c) The cell represented below was set up under standard conditions.

 $Pt|Fe^{2+}(aq), Fe^{3+}(aq)||TI^{3+}(aq),TI^{+}(aq)|Pt$

Cell e.m.f. = + 0.48 V

(2)

(i) Deduce the standard electrode potential for the following half-reaction.

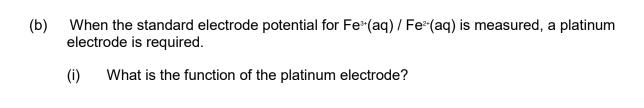
		TI³⁺(aq) + 2e⁻ → TI⁺(aq)	
	(ii)	Write an equation for the spontaneous cell reaction.	(3)
<i>(</i> 1)	A 51		
(d)	16.2	r acidification, 25.0 cm³ of a solution of hydrogen peroxide reacted exactly with cm³ of a 0.0200 mol dm⁻³ solution of potassium manganate(VII). The overall ation for the reaction is given below.	
	2Mn	$O_4^- + 6H^+ + 5H_2O_2 \rightarrow 2Mn^{2+} + 8H_2O + 5O_2$	
	(i)	Use the equation for this reaction to determine the concentration, in g dm $^{\!\scriptscriptstyle 3}\!$, of the hydrogen peroxide solution.	
	(ii)	Calculate the maximum volume of oxygen, measured at a pressure of 98 kPa and a temperature of 298 K, which would be evolved in this reaction.	
		(Total 14 n	(8) narks)

Q2.	Large blocks of magnesium are bolted onto the hulls of iron ships in an attempt to prevent the iron being converted into iron(II), one of the steps in the rusting process.							
	Use	the data below, where appropriate, to answer the questions which follow.						
		E^{Θ}/V $Mg^{2+}(aq) + 2e^{-} \longrightarrow Mg(s)$ -2.37 $Fe^{2+}(aq) + 2e^{-} \longrightarrow Fe(s)$ -0.44 $O_2(g) + 2H_2O(I) + 4e^{-} \longrightarrow 4OH^-(aq)$ $+0.40$						
	(a)	Calculate the e.m.f. of the cell represented by Mg(s) Mg²+(aq) Fe²+(aq) Fe(standard conditions. Write a half-equation for the reaction occurring at the electrode of this cell when a current is drawn.						
		Cell e.m.f.						
		Half-equation						
			(2)					
	(b)	Deduce how the e.m.f. of the cell Mg(s) Mg²+(aq) Fe²+(aq) Fe(s) changes w concentration of Mg²+ is decreased. Explain your answer.	hen the					
		Change in e.m.f.						
		Explanation						
			(3)					
	(c)	Calculate a value for the e.m.f. of the cell represented by $Pt(s) OH(qq) O_2(g) Fe^{2s}(qq) Fe(s)$ and use it to explain why iron corrodes we contact with water which contains dissolved oxygen.	hen in					
		Cell e.m.f.						
		Explanation						
			(2)					
			(Total 7 marks)					

Q3.Use the standard electrode potential data in the table below to answer the questions which follow.

			E [⊕] / \
Ce⁴(aq) + e⁻	-	Ce³∗(aq)	+1.70
MnO⁻(aq) + 8H⁺(aq)+ 5e⁻	\rightleftharpoons	Mn²⁺(aq) + 4H₂O(I)	+1.51
Cl ₂ (g) + 2e ⁻	$\overline{}$	2Cl-(aq)	+1.36
VO₂⁺(aq) +2H⁺(aq) + e⁻	$\overline{}$	$VO^{2+}(aq) + H_2O(I)$	+1.00
Fe³-(aq) + e⁻	\rightleftharpoons	Fe²⁺(aq)	+0.77
SO₄²-(aq) + 4H⁺(aq) + 2e⁻	$\overline{}$	H₂SO₃(aq) + H₂O(I)	+0.17

(a)	Name the standard reference electrode against which all other electrode potentials are measured.				
		(1)			



(11)	What are the standard conditions which apply to Fe³-(aq)/Fe²-(aq) whe measuring this potential?

(3)

(c)	The	cell represented below was set up under standard conditions.	
		$Pt H_{2}SO_{3}(aq),\ SO_{4}^{2-}(aq) MnO_{4}^{-}(aq),\ Mn^{2+}(aq) Pt$	
	Calci react	ulate the e.m.f. of this cell and write an equation for the spontaneous cell ion.	
	Cell	e.m.f	
	Equa	ation	
			(3)
(d)	(i)	Which one of the species given in the table is the strongest oxidising agent?	
	(ii)	Which of the species in the table could convert Fe²-(aq) into Fe³-(aq) but could not convert Mn²-(aq) into MnO₄-(aq)?	(3)
(e)	woul	data from the table of standard electrode potentials to deduce the cell which d have a standard e.m.f. of 0.93 V. Represent this cell using the convention <i>i</i> n in part (c).	
		(Total 12 ma	(2) arks)
Jse the	e data	in the table below to answer this question.	

Q4.U

E ∕ ∨ MnO_{4}^{-} (aq) + $8H^{+}$ (aq) + $5e^{-} \rightarrow Mn^{2+}$ (aq) + $4H_{2}O(I)$ + 1.52 $\text{Cr}_2\text{O}_7^{2-}(\text{aq}) + 14\text{H}^{\scriptscriptstyle +}(\text{aq}) + 6\text{e}^{\scriptscriptstyle -} \rightarrow 2\text{Cr}^{\scriptscriptstyle 3+}(\text{aq}) + 7\text{H}_2\text{O}(\text{I})$ + 1.33 + 0.77 $Fe^{3+}(aq) + e^- \rightarrow Fe^{2+}(aq)$

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$\operatorname{Cr}^{\scriptscriptstyle 3+}(\operatorname{aq}) + \operatorname{e}^{\scriptscriptstyle -} \to \operatorname{Cr}^{\scriptscriptstyle 2+}(\operatorname{aq})$	- 0.41
Zn²⁺(aq) + 2e⁻ → Zn(s)	- 0.76

Which one of the following statements is **not** correct?

- A Fe²⁺(aq) can reduce acidified MnO₄(aq) to Mn²⁺(aq)
- **B** $\operatorname{CrO}_7^{2-}(\operatorname{aq})$ can oxidise acidified $\operatorname{Fe}^{2+}(\operatorname{aq})$ to $\operatorname{Fe}^{3+}(\operatorname{aq})$
- **C** Zn(s) can reduce acidified $Cr_2O_7^{2-}$ (aq) to Cr^{2+} (aq)
- **D** Fe²⁺(aq) can reduce acidified Cr³⁺(aq) to Cr²⁺(aq)

(Total 1 mark)

Q5.Use the data in the table below to answer this question.

MnO_{4}^{-} (aq) + 8H ⁺ (aq) + 5e ⁻ \rightarrow Mn^{2+} (aq) + 4H ₂ O(I)	+ 1.52
$\text{Cr}_2\text{O}_7^{2-}(\text{aq}) + 14\text{H}^*(\text{aq}) + 6\text{e}^- \rightarrow 2\text{Cr}^{3*}(\text{aq}) + 7\text{H}_2\text{O}(\text{I})$	+ 1.33
Fe³+(aq) + e⁻ → Fe²+(aq)	+ 0.77
$\operatorname{Cr}^{\scriptscriptstyle 3+}(\operatorname{aq}) + \operatorname{e}^{\scriptscriptstyle -} \to \operatorname{Cr}^{\scriptscriptstyle 2+}(\operatorname{aq})$	- 0.41
$Zn^{2+}(aq) + 2e^- \rightarrow Zn(s)$	- 0.76

The most powerful oxidising agent in the table is

- A Mn²⁺(aq)
- **B** Zn(s)
- \mathbf{C} MnO $\frac{1}{4}$ (aq)
- **D** Zn²⁺(aq)

(Total 1 mark)

Q6.In this question consider the data below.

	E /V
$Ag^{\scriptscriptstyle{+}}(aq) + e^{\scriptscriptstyle{-}} \to Ag(s)$	+0.80
$2H^{\scriptscriptstyle +}(aq) + 2e^{\scriptscriptstyle -} \to H_2(g)$	0.00
$Pb^{2+}(aq) + 2e^{-} \rightarrow Pb(s)$	-0.13

The e.m.f. of the cell $Pt(s) \mid H_2(g) \mid H^+(aq) \mid Ag^+(aq) \mid Ag(s)$ would be increased by

- A increasing the concentration of H⁺(aq).
- **B** increasing the surface area of the Pt electrode.
- **C** increasing the concentration of Ag⁺(aq).
- **D** decreasing the pressure of $H_2(g)$.

(Total 1 mark)

Q7. Use the data below, where appropriate, to answer the questions which follow.

Standard electrode potentials	E [⊕] /V
$2H^{\cdot}(aq) + 2e^{-} \rightarrow H_{2}(g)$	0.00
$Br_2(aq) + 2e^- \rightarrow 2Br^-(aq)$	+1.09
$^{2\text{BrO}_{3}^{-}}$ (aq) + 12H $^{+}$ (aq) + 10e $^{-}$ \rightarrow Br $_{2}$ (aq) + 6H $_{2}$ O(I)	+1.52

Each of the above can be reversed under suitable conditions.

(a) State the hydrogen ion concentration and the hydrogen gas pressure when, at 298 K, the potential of the hydrogen electrode is 0.00 V.

Hydrogen ion concentration

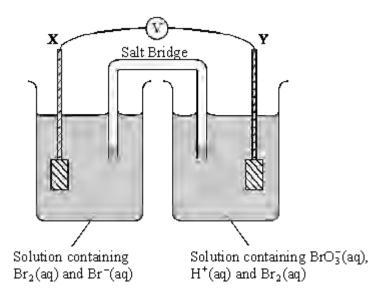
Hydrogen gas pressure

(2)

(b) The electrode potential of a hydrogen electrode changes when the hydrogen ion concentration is reduced. Explain, using Le Chatelier's principle, why this change occurs and state how the electrode potential of the hydrogen electrode changes.

Explanation of change

(c) A diagram of a cell using platinum electrodes **X** and **Y** is shown below.



(i) Use the data above to calculate the e.m.f. of the above cell under standard conditions.

(ii) Write a half-equation for the reaction occurring at electrode **X** and an overall equation for the cell reaction which occurs when electrodes **X** and **Y** are connected.

Half-equation	
Overall equation	

(Total 9 marks)

Q8.

$$Cr_{2}O_{7}^{2-}(aq) + 14H^{+}(aq) + 6e^{-} \rightarrow 2Cr^{3+}(aq) + 7H_{2}O(I) \qquad E^{\bullet} = +1.33 \text{ V}$$

$$Br_{2}(aq) + 2e^{-} \rightarrow 2Br^{-}(aq) \qquad E^{\bullet} = +1.09 \text{ V}$$

$$Fe^{3+}(aq) + e^{-} \rightarrow Fe^{2+}(aq) \qquad E^{\bullet} = +0.77 \text{ V}$$

$$VO^{2+}(aq) + 2H^{+}(aq) + e^{-} \rightarrow V^{3+}(aq) + H_{2}O(I) \qquad E^{\bullet} = +0.34 \text{ V}$$

$$SO_{4}^{2-}(aq) + 4H^{+}(aq) + 2e^{-} \rightarrow H_{2}SO_{3}(aq) + H_{2}O(I) \qquad E^{\bullet} = +0.17 \text{ V}$$

Based on the above data, which one of the following could reduce 0.012 mol of bromine to bromide ions?

- A 40 cm³ of a 0.10 mol dm⁻³ solution of $Cr_2O_7^{2-}$ (aq)
- **B** 80 cm³ of a 0.30 mol dm⁻³ solution of Fe³⁺(aq)
- C 50 cm³ of a 0.24 mol dm⁻³ solution of V³⁺(aq)
- C 50 cm³ of a 0.24 mol dm⁻³ solution of H₂SO₃(aq)

(Total 1 mark)

Q9.A disproportionation reaction occurs when a species **M**⁺ spontaneously undergoes simultaneous oxidation and reduction.

$$2M^+(aq) \rightarrow M^{2+}(aq) + M(s)$$

The table below contains E+ data for copper and mercury species.

	E⊕/∨
Cu²⁺(aq) + e⁻ → Cu⁺(aq)	+ 0.15
Cu⁺(aq) + e⁻ → Cu(s)	+ 0.52
$Hg^{2+}(aq) + e^- \rightarrow Hg^+(aq)$	+ 0.91
Hg⁺(aq) + e⁻ → Hg(l)	+ 0.80

Using these data, which one of the following can be predicted?

- A Both Cu(I) and Hg(I) undergo disproportionation.
- **B** Only Cu(I) undergoes disproportionation.

С	Only Hq(I)	undergoes	disprop	ortionation.
_	- / ()			

 $\label{eq:Delta} \textbf{D} \qquad \text{Neither Cu(I) nor Hg(I) undergoes disproportionation}.$

(Total 1 mark)