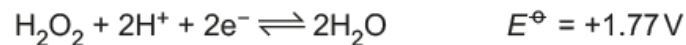


1. Nov/2023/Paper\_9701/41/No.3(e, f)

(e) The  $E^\ominus$  values for two electrode reactions are given.



(i) An electrochemical cell is constructed with the following half-cells (electrodes):

- an acidified solution of  $\text{H}_2\text{O}_2$ , a platinum wire
- $\text{Cr}^{2+}$  mixed with  $\text{Cr}^{3+}$ , a platinum wire.

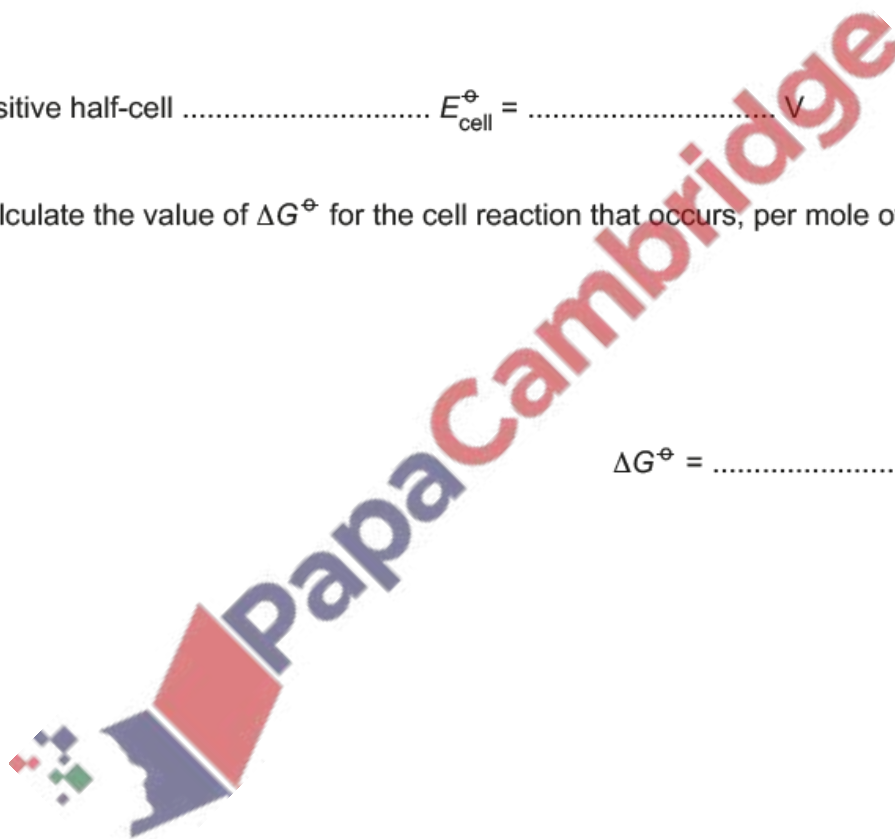
Identify the positive half-cell and calculate the standard cell potential,  $E_{\text{cell}}^\ominus$ .

positive half-cell .....  $E_{\text{cell}}^\ominus =$  ..... V

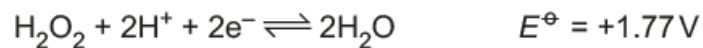
[1]

(ii) Calculate the value of  $\Delta G^\ominus$  for the cell reaction that occurs, per mole of  $\text{H}_2\text{O}_2$ .

$\Delta G^\ominus =$  .....  $\text{kJ mol}^{-1}$  [2]



(f) The  $E^\ominus$  values for two electrode reactions are given.



An electrochemical cell is constructed with the following half-cells.

half-cell 1 an acidified solution of  $\text{H}_2\text{O}_2$  under standard conditions, a platinum wire

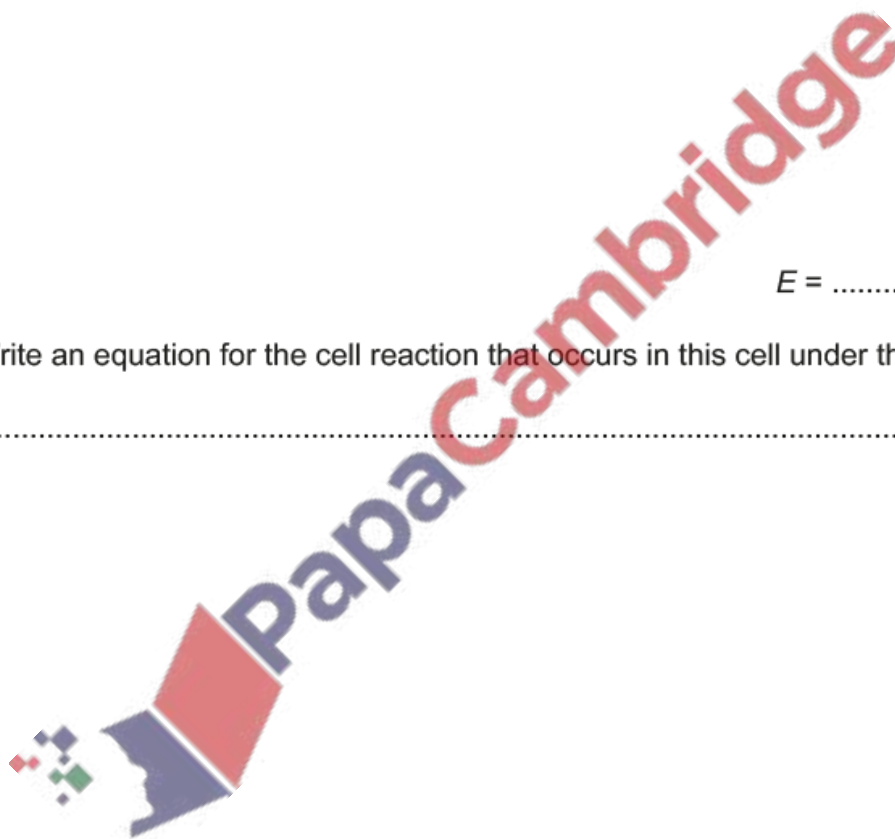
half-cell 2 a solution containing  $0.020\text{ mol dm}^{-3}\text{Co}^{3+}$  and  $2.0\text{ mol dm}^{-3}\text{Co}^{2+}$ , a platinum wire

(i) Use the Nernst equation to calculate the value of  $E$ , the electrode potential of half-cell 2 under these conditions.

$E = \dots\dots\dots\text{V}$  [2]

(ii) Write an equation for the cell reaction that occurs in this cell under these conditions.

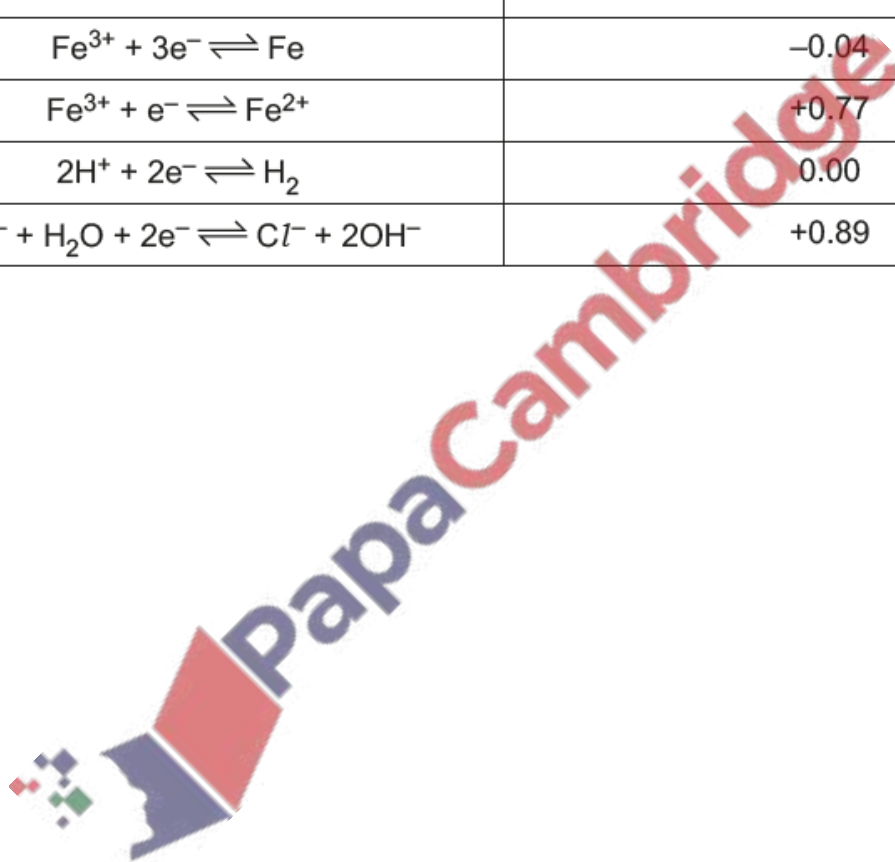
..... [1]



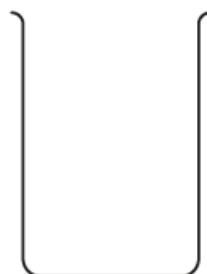
Some electrode potentials are shown in Table 3.1.

**Table 3.1**

electrode reaction	$E^\ominus/V$
$V^{2+} + 2e^- \rightleftharpoons V$	-1.20
$V^{3+} + e^- \rightleftharpoons V^{2+}$	-0.26
$VO^{2+} + 2H^+ + e^- \rightleftharpoons V^{3+} + H_2O$	+0.34
$VO_2^+ + 2H^+ + e^- \rightleftharpoons VO^{2+} + H_2O$	+1.00
$Fe^{2+} + 2e^- \rightleftharpoons Fe$	-0.44
$Fe^{3+} + 3e^- \rightleftharpoons Fe$	-0.04
$Fe^{3+} + e^- \rightleftharpoons Fe^{2+}$	+0.77
$2H^+ + 2e^- \rightleftharpoons H_2$	0.00
$ClO^- + H_2O + 2e^- \rightleftharpoons Cl^- + 2OH^-$	+0.89



- (a) (i) Complete the diagram to show a standard hydrogen electrode. Label your diagram. Identify all substances. You do **not** need to state standard conditions.



[1]

- (ii) An electrochemical cell is set up using an  $\text{Fe}^{3+}/\text{Fe}^{2+}$  electrode and a standard hydrogen electrode.

Identify the positive electrode in the electrochemical cell and the direction of electron flow in the external circuit.

positive electrode .....

Electrons flow from the ..... electrode to the ..... electrode.

[1]



(b) The vanadium-containing species in the electrode reactions given in Table 3.1 are V,  $V^{2+}$ ,  $V^{3+}$ ,  $VO^{2+}$  and  $VO_2^+$ .

(i) Identify **one** vanadium-containing species that does **not** react with  $Fe^{2+}$  ions under standard conditions.

Use data from Table 3.1 to explain your answer.

.....  
..... [1]

(ii) Identify **all** the vanadium-containing species that will react with  $Fe^{2+}$  ions under standard conditions.

..... [1]

(iii) Write an equation for **one** of the possible reactions identified in (ii).

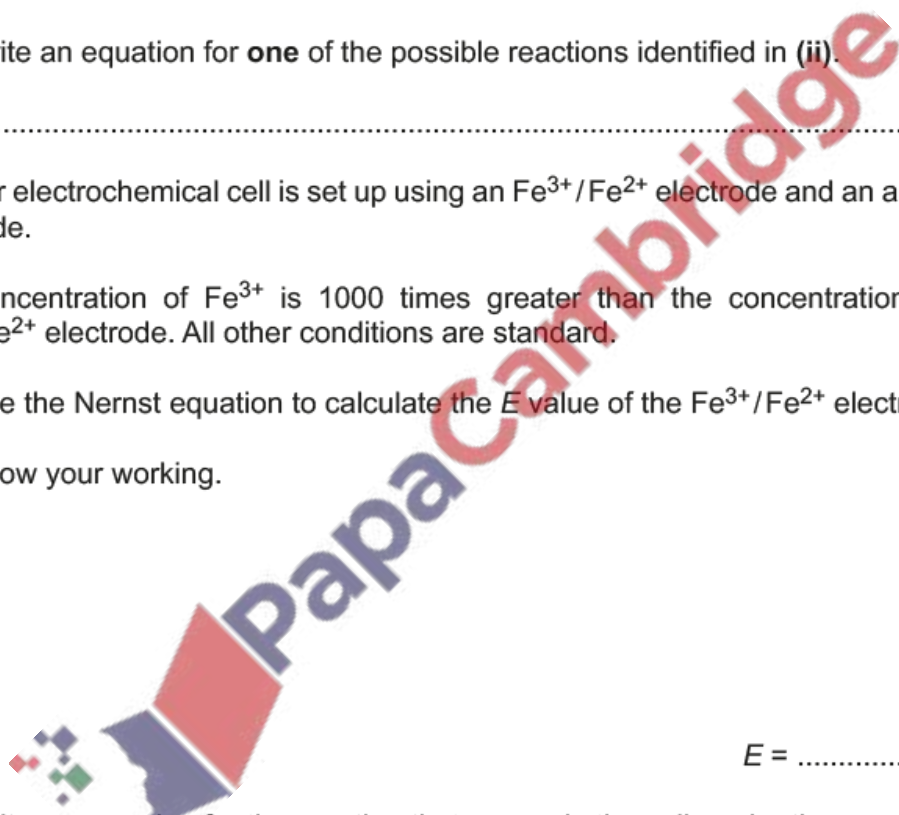
..... [1]

(c) Another electrochemical cell is set up using an  $Fe^{3+}/Fe^{2+}$  electrode and an alkaline  $ClO^-/Cl^-$  electrode.

The concentration of  $Fe^{3+}$  is 1000 times greater than the concentration of  $Fe^{2+}$  in the  $Fe^{3+}/Fe^{2+}$  electrode. All other conditions are standard.

(i) Use the Nernst equation to calculate the  $E$  value of the  $Fe^{3+}/Fe^{2+}$  electrode.

Show your working.



$E = \dots\dots\dots$  V [2]

(ii) Write an equation for the reaction that occurs in the cell, under these conditions.

..... [1]

(d) Another electrochemical cell is set up using an  $Fe^{2+}/Fe$  electrode and an alkaline  $ClO^-/Cl^-$  electrode under standard conditions.

Calculate the value of  $\Delta G^\ominus$  for the cell.

$\Delta G^\ominus = \dots\dots\dots$   $\text{kJ mol}^{-1}$  [3]

- (e) A solution of iron(II) sulfate,  $\text{FeSO}_4(\text{aq})$  is electrolysed with iron electrodes. Under the conditions used, no gas is evolved at the cathode.

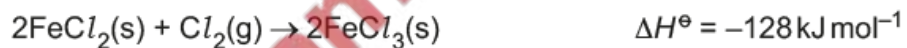
A current of 0.640A is passed for 17.0 minutes. The mass of the cathode increases by 0.185g.

Use these results to calculate an experimental value for the Avogadro constant,  $L$ .

Show your working.

$$L = \dots\dots\dots \text{mol}^{-1} \quad [3]$$

- (f) Iron(II) chloride,  $\text{FeCl}_2$ , is oxidised by chlorine to form iron(III) chloride,  $\text{FeCl}_3$ , under standard conditions.



**Table 3.2**

species	$S^\ominus / \text{JK}^{-1} \text{mol}^{-1}$
$\text{Cl}_2(\text{g})$	223
$\text{FeCl}_2(\text{s})$	120
$\text{FeCl}_3(\text{s})$	142

- (i) Use Table 3.2 and other data to calculate the Gibbs free energy change,  $\Delta G^\ominus$ , for this reaction.

Show your working.

$$\Delta G^\ominus = \dots\dots\dots \text{kJ mol}^{-1} \quad [3]$$

(ii) Predict whether this reaction becomes more or less feasible at a higher temperature.

Explain your answer.

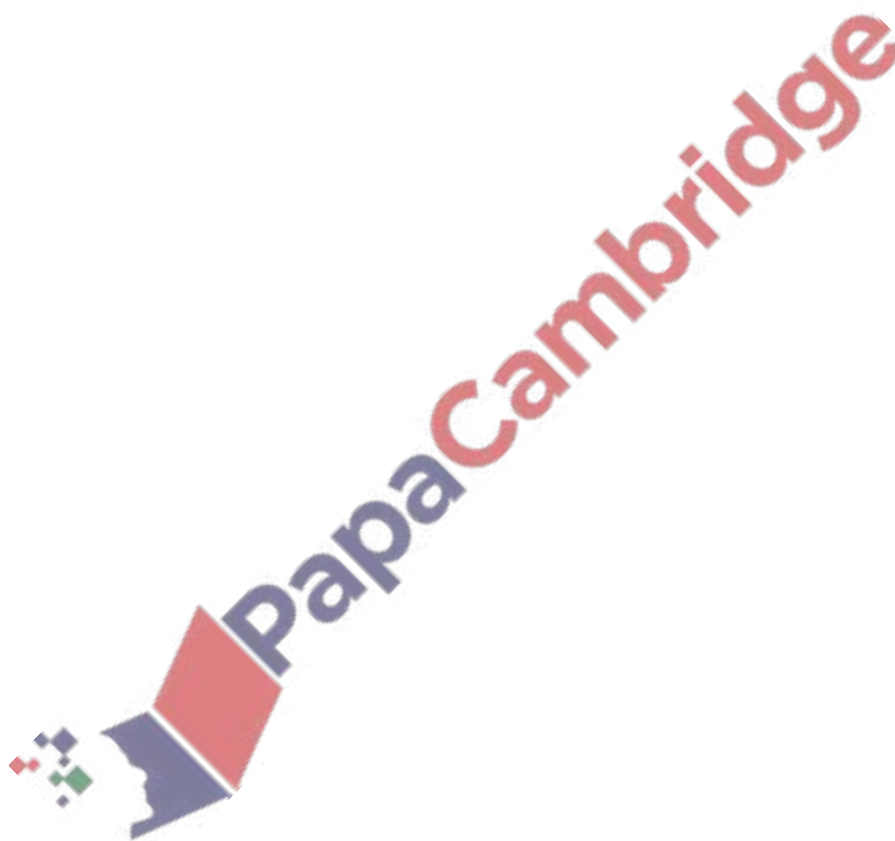
The reaction becomes ..... feasible.

explanation .....

.....

[1]

[Total: 18]



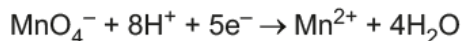
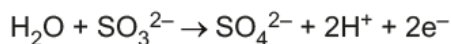
3. June/2023/Paper\_9701/42/No.1(c\_d)

(c) Potassium sulfite,  $K_2SO_3$ , is used as a food additive.

The concentration of sulfite ions,  $SO_3^{2-}$ , can be determined by titration using aqueous acidified manganate(VII) ions,  $MnO_4^-$ .

- A  $250\text{ cm}^3$  solution contains  $3.40\text{ g}$  of impure  $K_2SO_3$ .
- $25.0\text{ cm}^3$  of this solution requires  $22.40\text{ cm}^3$  of  $0.0250\text{ mol dm}^{-3}$  acidified  $MnO_4^-$  to reach the end-point. All the  $SO_3^{2-}$  ions are oxidised. None of the other species in the impure  $K_2SO_3$  are oxidised.

The reaction occurs as shown by the two half-equations.



(i) Give the ionic equation for the reaction between  $SO_3^{2-}$  and acidified  $MnO_4^-$ .

..... [1]

(ii) Calculate the percentage purity of the sample of  $K_2SO_3$ .  
Show your working.

percentage purity of  $K_2SO_3$  = ..... [3]

(d) Potassium disulfite,  $K_2S_2O_5$ , is another food additive. The disulfite ion,  $S_2O_5^{2-}$ , has the displayed formula shown in Fig. 1.2.

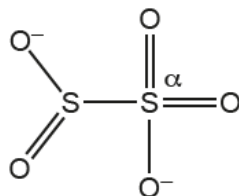


Fig. 1.2

Deduce the geometry (shape) around the  $S(\alpha)$  atom in  $S_2O_5^{2-}$ .

geometry around  $S(\alpha)$  ..... [1]

4. June/2023/Paper\_9701/42/No.9



(a) Define standard cell potential,  $E_{\text{cell}}^{\ominus}$ .

.....  
..... [1]

(b) An electrochemical cell is set up to measure  $E_{\text{cell}}^{\ominus}$  of a cell consisting of an  $\text{Fe}^{3+}/\text{Fe}^{2+}$  half-cell and a  $\text{Cl}_2/\text{Cl}^-$  half-cell.

Draw a labelled diagram of this electrochemical cell.

Include all necessary substances. It is **not** necessary to state conditions used.

PapaCambridge

[3]

(c) The cell reaction for the electrochemical cell in (b) is shown.



Calculate  $\Delta G^{\ominus}$ , in  $\text{kJ mol}^{-1}$ , for this cell reaction.

$\Delta G^{\ominus} = \dots\dots\dots \text{kJ mol}^{-1}$  [2]

(d) Another experiment is set up using the same electrochemical cell.

In this experiment the  $\text{Fe}^{2+}$  concentration is  $0.15 \text{ mol dm}^{-3}$ . All other concentrations remain at their standard values.

The Nernst equation is shown.

$$E = E^\ominus + (0.059/z) \log \frac{[\text{oxidised species}]}{[\text{reduced species}]}$$

(i) Use the Nernst equation to calculate the electrode potential,  $E$ , for the  $\text{Fe}^{3+}/\text{Fe}^{2+}$  half-cell in this experiment.

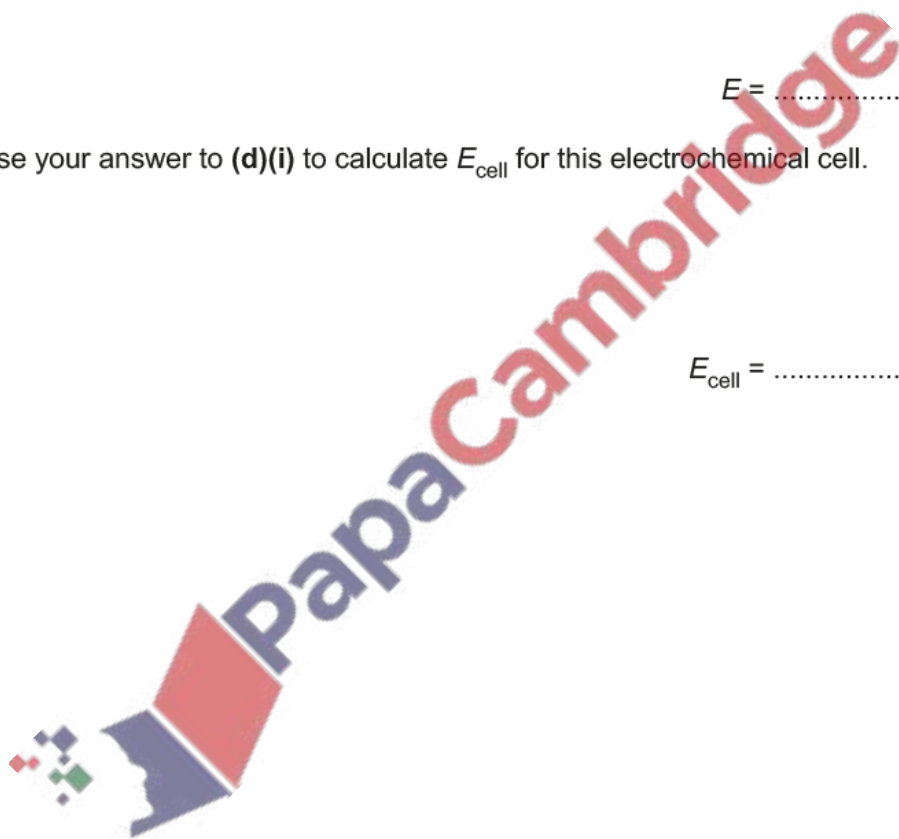
$[E^\ominus: \text{Fe}^{3+}/\text{Fe}^{2+} = +0.77 \text{ V}]$

$E = \dots\dots\dots \text{ V [1]}$

(ii) Use your answer to (d)(i) to calculate  $E_{\text{cell}}$  for this electrochemical cell.

$E_{\text{cell}} = \dots\dots\dots \text{ V [1]}$

[Total: 8]



5. March/2023/Paper\_9701/42/No.2(a, b)

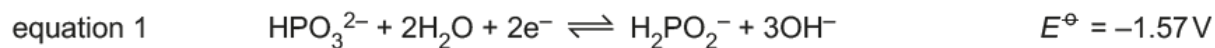
Hypophosphorous acid is an inorganic acid.

The conjugate base of hypophosphorous acid is  $\text{H}_2\text{PO}_2^-$ .

(a) Give the formula of hypophosphorous acid.

..... [1]

(b)  $\text{H}_2\text{PO}_2^-$  is a strong reducing agent. It can be used to reduce metal cations without the need for electrolysis.



(i) In an experiment, an alkaline  $\text{HPO}_3^{2-}/\text{H}_2\text{PO}_2^-$  half-cell is constructed with  $[\text{H}_2\text{PO}_2^-] = 0.050\text{ mol dm}^{-3}$ .

All other ions are at their standard concentration.

Predict how the value of  $E$  of this half-cell differs from its  $E^\ominus$  value.

Explain your answer.

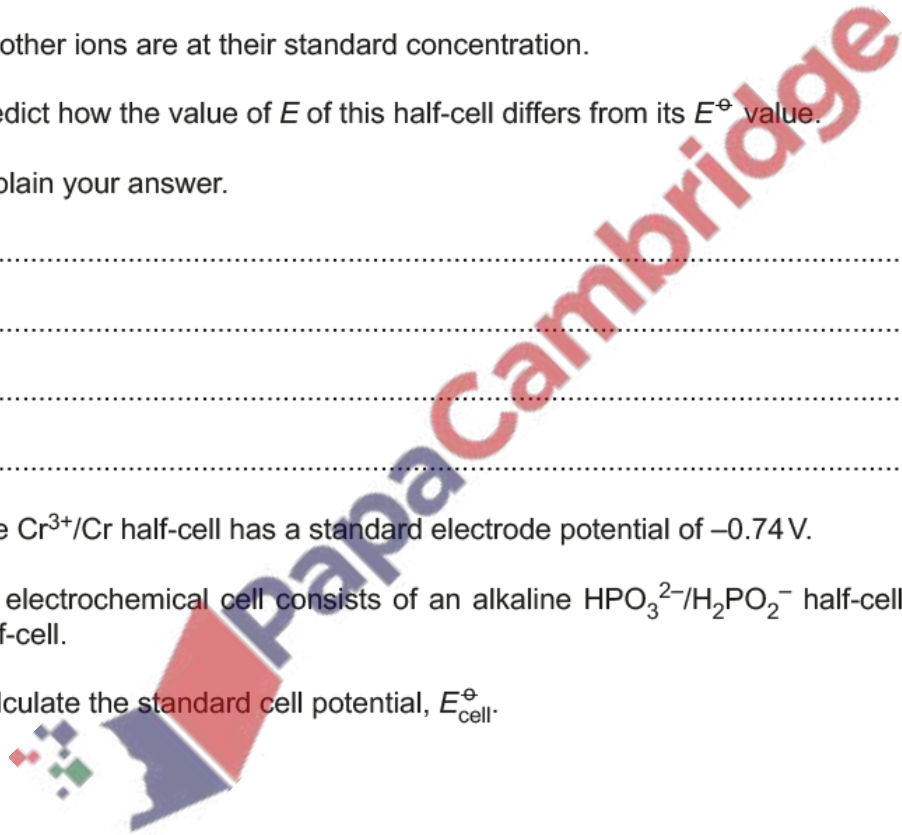
.....  
.....  
.....  
..... [2]

(ii) The  $\text{Cr}^{3+}/\text{Cr}$  half-cell has a standard electrode potential of  $-0.74\text{V}$ .

An electrochemical cell consists of an alkaline  $\text{HPO}_3^{2-}/\text{H}_2\text{PO}_2^-$  half-cell and a  $\text{Cr}^{3+}/\text{Cr}$  half-cell.

Calculate the standard cell potential,  $E_{\text{cell}}^\ominus$ .

$E_{\text{cell}}^\ominus = \dots\dots\dots\text{V}$  [1]



- (iii) Complete the diagram in Fig. 2.1 to show how the standard electrode potential of the  $\text{Cr}^{3+}/\text{Cr}$  half-cell can be measured relative to that of the standard hydrogen electrode.

Identify the chemicals, conditions and relevant pieces of apparatus.

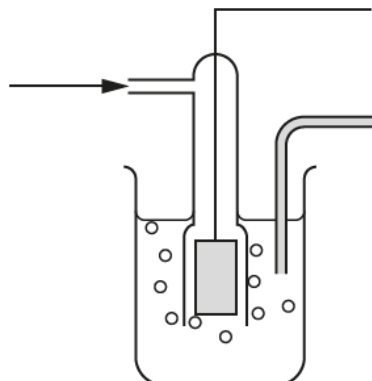


Fig. 2.1

[3]

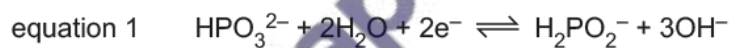
- (iv) Label Fig. 2.1 to show:

- which is the positive electrode
- the direction of electron flow in the external circuit.

[1]

- (v)  $\text{H}_2\text{PO}_2^-$  reduces  $\text{Ni}^{2+}$  to Ni in alkaline conditions.

Use equation 1 to construct the ionic equation for this reaction.



[1]

