## Assessment Schedule - 2006

## Chemistry: Describe oxidation-reduction processes (90696)

## **Evidence Statement**

Q	Evidence	Achievement	Achievement with Merit	Achievement with Excellence
1(a)	In Cr <sub>2</sub> O <sub>7</sub> <sup>2-</sup> ox number of Cr is +6 but in Cr <sup>3+</sup> it is +3, so decrease of oxidation number or gain of electrons therefore a reduction.	Correct oxidation numbers with valid statement.		
1(b)	$Cu \rightarrow Cu^{2+} + 2e OR Cu - 2e \rightarrow Cu^{2+}$ $Cr_2O_7^{2-} + 14H^+ + 6e \rightarrow 2Cr^{3+} + 7H_2O$ $Cr_2O_7^{2-} + 14H^+ + 3Cu \rightarrow 2Cr^{3+} + 7H_2O + 3Cu^{2+}$	Two half equations correctly balanced.	All three equations correct.	
1(c)	Changes in half-cells are: In copper half-cell  • Copper electrode erodes or starts to disappear.  • Solution goes (darker) blue (Cu <sup>2+</sup> forming). In dichromate half-cell  • Solution goes from orange (Cr <sub>2</sub> O <sub>7</sub> <sup>2-</sup> ) to blue/green (Cr <sup>3+</sup> ).	Identifies one observed change in each half cell.		
2(a)	The $E_{\text{cell}}^{o}$ for the reaction between $\text{MnO}_4^-$ and $\text{Mn}^{2+}$ is -0.64V, and being negative means the reaction will not spontaneously occur. <b>OR</b> Recognizes reduction potential of $E^{o}$ ( $\text{MnO}_4^-/\text{Mn}^{3+}$ ) is less than $E^{o}$ ( $\text{Mn}^{3+}/\text{Mn}^{2+}$ ) so $\text{MnO}_4^-$ cannot be spontaneously reduced by $\text{Mn}^{2+}$ .	Identify a negative value of $E^o_{cell}$ means no reaction <b>OR</b> Recognises reduction potential of $E^o(MnO_4^-/Mn^{3+})$ is less than $E^o(Mn^{3+}/Mn^{2+})$ .	Shows calculation to determine $E^{o}_{cell}$ value of $-0.64$ V and links to fact that no reaction occurs.  OR  Explanation that correctly links reduction potentials to strength of oxidant and reductant so that no reaction occurs.	

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2(b)	<ul> <li>(i)</li> <li>• In the acidified solution, observation is purple to colourless or pale pink solution because the purple MnO<sub>4</sub><sup>-</sup> is reduced to colourless Mn<sup>2+</sup> (both the sulfite and sulfate ion produced by oxidation are colourless species).</li> <li>• In the weakly alkaline solution, the purple solution of MnO<sub>4</sub><sup>-</sup> forms a black or brown precipitate of MnO<sub>2</sub> solid.</li> <li>• In the strongly alkaline solution the MnO<sub>4</sub><sup>-</sup> solution is reduced to a green solution of MnO<sub>4</sub><sup>2-</sup>.</li> <li>(ii)</li> <li>5SO<sub>3</sub><sup>2-</sup> + 2MnO<sub>4</sub><sup>-</sup> + 6H<sup>+</sup> → 5SO<sub>4</sub><sup>2-</sup> + 2Mn<sup>2+</sup> + 3H<sub>2</sub>O</li> </ul>	At least two correct observations linked to species involved <b>OR</b> Correct observations for all three reactions of MnO <sub>4</sub> <sup>-</sup> . <b>OR</b> All four species correctly linked to their colours <b>OR</b> Redox reactants and products identified: SO <sub>3</sub> <sup>2-</sup> and MnO <sub>4</sub> <sup>-</sup> , Mn <sup>2+</sup> and SO <sub>4</sub> <sup>2-</sup>	All observations correct and linked to species involved.  OR  Fully balanced equation correct.	All correct observations with links to species and correctly balanced equation.
	In Cell One the beakers are connected by a bridge containing methanol, which is a non-electrolyte and therefore will not allow flow of ions. Needs to be replaced with a salt bridge or solution of an ionic substance such as KNO <sub>3</sub> .  In Cell Two a spontaneous reaction occurs in the right-hand beaker where the Zn is oxidised and Cu <sup>2+</sup> reduced. There would be no reaction in the left-hand beaker and no flow of charge. Reversing the electrodes or the solutions in the beakers would result in a flow of charge between the half-cells.	Correctly explains the problem in one cell and how the problem can be fixed.	Identifies and explains the problem in both cells.	
4(a)	(i) • e <sup>-</sup> flow from left to right • left electrode negative/right electrode positive.  (ii)  E <sup>o</sup> <sub>cell</sub> = +0.74 V = E <sup>o</sup> (H <sup>+</sup> /H <sub>2</sub> ) – E <sup>o</sup> (X <sup>3+</sup> /X)  E <sup>o</sup> (X <sup>3+</sup> /X) = -0.74 V  The H <sup>+</sup> /H <sub>2</sub> electrode is positive because electrons are flowing towards this electrode. This electrode has a standard reduction potential of 0.00 V, so the voltage on the voltmeter gives the difference between this electrode where reduction occurs and the other electrode where oxidation occurs. The other electrode must be more negative than 0.00 V by the reading on the voltmeter.	Direction of electron flow consistent with sign on electrodes, ie as written or both direction of electron flow and signs on electrodes reversed. <b>OR</b> $E^{o}(X^{3+}/X) = -0.74 \text{ V with no working shown.}$ <b>OR</b> Recognition that difference in reduction potential between $E^{o}(X^{3+}/X) \text{ and } E^{o}(H^{+}/H_{2}) = 0.00 \text{ V}$ gives value of 0.74 V.	$E^{o}(X^{3+}/X) = -0.74 \text{ V}$ is calculated or explained correctly by comparison with $E^{o}(H^{+}/H_{2}) = 0.00 \text{ V}$ and either direction of $e^{-}$ flow or labeling of electrodes is correct. <b>OR</b> Comparison with $E^{o}(H^{+}/H_{2}) = 0.00 \text{ V}$ gives $E^{o}(X^{3+}/X) = +0.74 \text{ V}$ , and direction of electron flow and signs of electrodes both reversed. <b>OR</b> Direction of electron flow and reduction potential correct, but no or limited explanation/calculation.	All correct with valid explanation / calculation for electrode potential.

Q	Evidence	Achievement	Achievement with Merit	Achievement with Excellence
		Method correct but mathematical error.  This includes correct balanced equation but use inverted ratio to get n(X) =5.46 mol.  OR incorrectly balanced eqn but application of ratio in that eqn correct.	Answer correct.	
5(a) (b)	Zn > Ga > Fe  Answer discusses fact that $E^{\circ}(\text{Fe}^{2^+}/\text{Fe}) > (E^{\circ}(\text{Ga}^{3^+}/\text{Ga}) > E^{\circ}(\text{Zn}^{2^+}/\text{Zn})$ . Ie as zinc can reduce both $\text{Ga}^{3^+}$ and $\text{Fe}^{2^+}$ it is the strongest reductant ( $E^{\circ}(\text{Ga}^{3^+}/\text{Ga})$ and $E^{\circ}(\text{Fe}^{2^+}/\text{Fe})$ must be greater than $E^{\circ}(\text{Zn}^{2^+}/\text{Zn})$ ). Ga is a stronger reductant than Fe as Ga can reduce $\text{Fe}^{2^+}$ , but Fe cannot reduce $\text{Ga}^{3^+}$ and $E^{\circ}(\text{Fe}^{2^+}/\text{Fe})$ must be greater than $E^{\circ}(\text{Ga}^{3^+}/\text{Ga})$ .	Order of reductants (in part a) correct without valid justification. <b>OR</b> In part (b) identifies order of $E^{\circ}(Fe^{2^{+}}/Fe) > (E^{\circ}(Ga^{3^{+}}/Ga) > E^{\circ}(Zn^{2^{+}}/Zn)$ .	$>(E^{\circ}(Ga^{3+}/Ga) > E^{\circ}(Zn^{2+}/Zn)$ but lists	Answer shows correct order of metals as reductants and discussion clearly relates the order to the relevant $E^{\circ}$ values. eg best reductant will have most negative $E^{\circ}$ value and will be metal reacting with both of the other metal ions.

## **Judgement Statement**

Chemistry: Describe oxidation-reduction processes (90696)

Achievement	Achievement with Merit	Achievement with Excellence
FIVE questions answered correctly.	SIX questions answered correctly, including at least FOUR at Merit	SEVEN questions answered correctly, including at least THREE
Minimum of 5 × A	level.	at Merit level and at least TWO at Excellence level.
	Minimum of $4 \times M + 2 \times A$	
		Minimum of $2 \times E + 3 \times M + 2 \times A$