UNIT REVIEW: REDOX – ANSWERS

- 1. a) +6
 - b) +5
 - c) +3, +4
 - **d**) -1

| 4 | | 4 | 1 | |
|---|----|---|----|-----|
| | | A | | |
| | 12 | | V | d |
| w | | | ₩. | -11 |

| Question | Oxidized | Reduced | Oxidizing Ag. | Reducing Ag. |
|----------|----------|-----------------|-----------------|--------------|
| A | I | IO ₃ | IO ₃ | Ī |
| В | Zn | NO ₃ | NO ₃ | Zn |

- 3. a) $Cr_2O_7^{2-} + 3Bi^{+3} + 4OH^{-1} \rightarrow 2Cr^{+3} + 3BiO_3^{-1} + 2H_2O$
 - b) $MnO_4 + 3SO_3^{-1} + 2OH^{-1} \rightarrow MnO_2 + 3SO_4^{-2} + H_2O$ c) $6Br_2 + 12OH^{-1} \rightarrow 10Br^{-1} + 2BrO_3^{-1} + 6H_2O$

 - d) $2P + 3PO_4^{-3} + 2H_2O + OH^{-} \rightarrow 5HPO_3^{-2}$
 - e) $7H_2O + Cr_2O_7^{-2} + 3C_2O_4^{-2} \rightarrow 2Cr^{+3} + 6CO_2 + 14OH^{-1}$
 - f) $5H_2O + 2CrO_4^{2-} + Cl^{-} \rightarrow ClO_3^{-} + 2Cr^{+3} + 10OH^{-}$
 - g) $2MnO_4^- + 3CN^- + H_2O \rightarrow 2MnO_2 + 3CNO^- + 2OH^{-1}$
 - h) $2OH^{-} + SeO_{3}^{-2} + Cl_{2} \rightarrow SeO_{4}^{-2} + 2Cl^{-1} + H_{2}O$
- a) Yes. Cl_2 SRP = +1.36V, Br_2 SRP = +1.09 V, E^0 = +0.27 V, spontaneous
 - b) No. Al $^{3+}$ SRP = -1.66V, Hg SRP = +0.85, E 0 = -2.51V, non-spontaneous
- 5. Potassium (K) is the stronger reducing agent. Oxidation potential of K (+2.92 V) > Oxidation potential of Cu (-0.15 V). Potassium is an alkali metal, i.e. one valence electron, highly reactive and hence easily oxidized
- Sulfur has multiple oxidation states ranging from S⁻² (isoelectronic to Ar) to S⁺⁶ (isoelectronic to Ne). Hence SO₂ (S oxidation number of +4) can be reduced or oxidized towards an oxidation number of -2 or +6
- 7. a) $2Cr_2O_7^{2-} + C_2H_5OH + 16H^+ \rightarrow 4Cr^{3+} + 2CO_2 + 11H_2O$
 - b) 0.16% alcohol in blood
- 8. Fe (Iron) is both the oxidant and the reducing agent; Iron is a better reducing agent than chlorine gas and is most commonly found with oxidation numbers of +2 or +3, i.e. oxidized
- 9. $2Al_{(s)} + 6H_{(aq)}^+ + 3O_{2(aq)} \rightarrow 2Al_{(aq)}^{+3} + 3H_2O_{2(aq)}$

- 11. $E_{cell}^{o} = -0.36 \text{ V}$, negative voltage, non-spontaneous ($dG = -nFE^{o}$)
- 12. Electrolytes maintain a balance of charge between anode and cathode such that the cell produces a voltage consistently. Anode has excess positive charge in solution, must be balanced with anions; Cathode has excess negative charge in solution, must be balanced with cations.
- 13. The presence of salts or acids will increase water's conductivity and hence improve it's ability to carry electrons during the redox reaction.
- 14. Scandium (Sc, 44.95 g/mol)
- 16. a) 124 minutes
 - b) Water is more easily reduced than Calcium ions in an electrolytic aqueous cell (-0.83V > -2.87 V); Water will be discharged in cell instead, calcium ions do not undergo reduction
- 17. 41.1 minutes

18. i)
$$C_6H_8O_6 \rightarrow C_6H_6O_6 + 2H^+ + 2e^-$$

ii) $2Fe^{+3}C_6H_8O_6 \rightarrow 2Fe^{+2} + C_6H_6O_6 + 2H^+$

- 19. a) $Na^+ + e^- \rightarrow Na$; $2Cl^- \rightarrow Cl_2 + 2e^-$; Electron flow from battery to electrodes; Electron flow at the cathode (reduction of cations); Electron flow at the anode (oxidation of anions)
 - b) 0.2 mol of sodium; net equation is $2Na^+ + 2Cl^- \rightarrow 2Na + Cl_2$: For every mol of chlorine gas, 2 mols of solid sodium is formed; $(0.1 \times 2 = 0.2 \text{ mol})$
 - c) 0.25 mol Cl₂
 - d) Hydrogen gas and hydroxide formed at the cathode, Oxygen gas and H^+ at the anode; $H_2O_{(l)} \rightarrow H_{2(g)} + \frac{1}{2}O_{2(g)}$
 - e) Potential difference (voltage) obtained when a half-cell of an element is connected with a hydrogen electrode under standard conditions (298K, 1.0 moldm^{-3} , 1 atm)

20. a)
$$Cr_2O_7^{-2} + 3SO_3^{-2} + 8H^+ \rightarrow 2Cr^{+3} + 3SO4^{-2} + 4H_2O$$

b) $H_2O + MnO_4^- + 3VO^{+2} \rightarrow MnO_2 + 3VO_2^{+1} + 2H^+$

21. Water has a greater reduction potential than Na^+ (-0.83V > -2.71V); Water is more easily oxidized than F^- (-1.23V > -2.87 V), hence it is preferred in an electrolytic oxidization reaction at both the cathode and anode

22. a) Na⁺, H⁺, Ag⁺ b) Ag, H, Na