

TOPIC 15 ANSWERS TO EXERCISES

Topic 15 Exercise 1

- $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}4s^{2}3d^{3}$ $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}4s^{1}3d^{5}$ $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}4s^{1}3d^{10}$ 1. b) a) 1s²2s²2p⁶3s²3p⁶4s²3d⁷ 1s²2s²2p⁶3s²3p⁶4s²3d¹⁰ d) c) d) $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}3d^{7}$ $\frac{1s^22s^22p^63s^23p^63d^{10}}{1s^22s^22p^63s^23p^63d^3}$ 2. b) a) $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}3d^{2}$ c) d) $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}3d^{5}$ d)
- 3. a) They do not form any stable ions with partially filled d-orbitals
 - b) They have empty d-orbitals of low energy which can accept electron pairs
 - c) The presence of ligands causes the d-orbitals to split. In the presence of visible light, electrons are excited from low energy d-orbitals to high energy d-orbitals, absorbing the light in the process. The resultant light is coloured.
 - d) Cu^+ has a full 3d sublevel, so there are no available d-orbitals into which electrons can be excited.
- Prepare samples of the transition metal ion over a range of known concentrations. Add a small quantity of a suitable ligand to each in order to identify the colour. Choose the filter which gives the largest absorbance, and measure the absorbance of each sample using this filter.

Plot a graph of absorbance against concentration.

Take the sample of unknown concentration, add the ligand to intensify the colour, and measure its absorbance using the same filter.

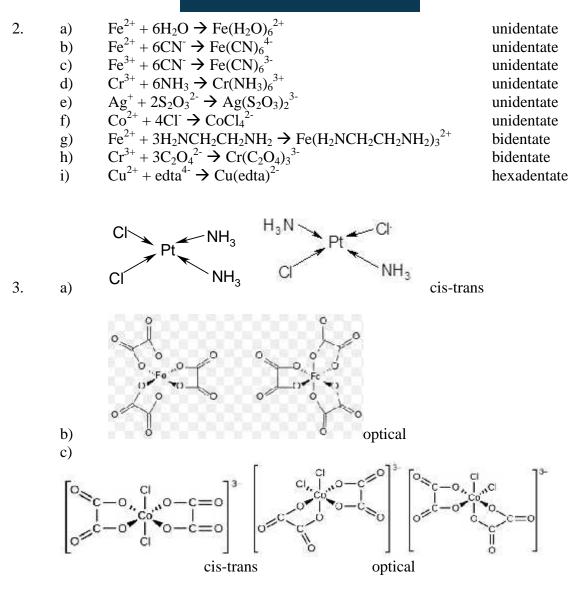
Use the graph to deduce the concentration of the solution.

Topic 15 Exercise 2

- 1. a) ion containing a central metal ion attached to one or more ligands by means of co-ordinate bonds
 - b) species containing a lone pair of electrons which can form a co-ordinate bond with a metal ion
 - c) the total number of co-ordinate bonds formed between the metal ion and the ligands

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Topic 15 Exercise 3

1. a) acid-base $\operatorname{Fe}(\operatorname{H_2O}_6^{2+}(\operatorname{aq}) + 2\operatorname{OH}^{-}(\operatorname{aq}) \rightarrow \operatorname{Fe}(\operatorname{H_2O}_4(\operatorname{OH}_2(\operatorname{s}) + 2\operatorname{H_2O}(\operatorname{l}))$ acid-base b) $Al(H_2O)_6^{3+}(aq) + 3OH^{-}(aq) \rightarrow Al(H_2O)_3(OH)_3(s) + 3H_2O(l)$ $Al(H_2O)_3(OH)_3(s) + 3OH^{-}(aq) \rightarrow Al(OH)_6^{3-}(aq) + 3H_2O(1)$ acid-base c) $Cu(H_2O)_6^{2+}(aq) + 2NH_3(aq) \rightarrow Cu(H_2O)_4(OH)_2(s) + 2NH_4^+(aq)$ Ligand exchange $[Cu(H_2O)_4(OH)_2](s) + 4NH_3(aq) == [Cu(NH_3)_4(H_2O)_2]^{2+}(aq) + 2H_2O(1) + 2OH^{-}(aq)$ d) ligand exchange $[Co(H_2O)_6]^{2+}(aq) + 4Cl^{-}(aq) = [CoCl_4]^{2-}(aq) + 6H_2O(l)$ ligand exchange $[CoCl_4]^{2-}(aq) + 6H_2O(l) == [Co(H_2O)_6]^{2+}(aq) + 4Cl^{-}(aq)$

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- acid-base e) $2[Al(H_2O)_6]^{3+}(aq) + 3CO_3^{2-}(aq) \rightarrow 2[Al(H_2O)_3(OH)_3](s) + 3CO_2(g) + 3H_2O(l)$ f) precipitation $[Co(H_2O)_6]^{2+}(aq) + CO_3^{2-}(aq) \rightarrow CoCO_3(s) + 6H_2O(l)$
- Fe^{3+} has a greater charge density than Fe^{2+} 2.
 - So the electrons in the O-H bonds in the complex ion $Fe(H_2O)_6^{3+}$ are pulled more strongly towards the central metal ion than in $Fe(H_2O)_6^2$
 - which means the O-H bonds are more likely to dissociate to form H⁺ in $Fe(H_2O)_6^{3+}$

So $Fe(H_2O)_6^{3+}$ is more acidic and has a lower pH

3. The reaction involves breaking six O-H bonds and making six O-H bonds so H is close to 0

the reaction involves an increase in the number of species in aqueous solution from 2 to 7

- so there is a large increase in entropy
- so G is negative and the reaction is thermodynamically favoured

Topic 15 Exercise 4

- Solution turns yellow when acidified 1. Due to formation of VO_2^+ Yellow solution turns blue then green then violet as the vanadium is reduced from oxidation state +5 to +4 to +3 to +2
- 2. $= -2SO_{3}$ a) i) $2SO_2 + O_2$ a step in the manufacture of sulphuric acid
 - $N_2 + 3H_2 = 2NH_3$ ii) the manufacture of ammonia
 - $2CO + 2NO \rightarrow N_2 + 2CO_2$ iii) removal of pollutants in car engines
 - heterogeneous catalysts provide active sites at their surface where the b) reaction can take place. Catalyst poisons block these sites.
 - c) The catalyst is expensive and must be used in small quantities It needs to have a large surface area to maximize its effectiveness So it is spread thinly over an inert support.
- using Fe²⁺: 3. $S_{2}O_{8}^{2^{-}} + 2Fe^{2^{+}} \rightarrow 2SO_{4}^{2^{-}} + 2Fe^{3^{+}}$ $2Fe^{3^{+}} + 2I^{-} \rightarrow 2Fe^{2^{+}} + I_{2}$ using Fe³⁺: $2Fe^{3+} + 2I^{-} \rightarrow 2Fe^{2+} + I_2$ $S_2O_8^{2-} + 2Fe^{2+} \rightarrow 2SO_4^{2-} + 2Fe^{3+}$ these steps are all fast because they involve a collision between oppositely charged ions, which attract each other

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The uncatalysed reaction involves a collision between two anions, which repel, making the reaction slow

- 4. a) $2MnO_4^- + 16H^+ + 5C_2O_4^{-2-} \rightarrow 2Mn^{2+} + 8H_2O + 10CO_2$
 - b) Initially the reaction is slow because it requires a collision between two repelling anions
 But after the reaction has started the Mn²⁺ produced can catalyse the reaction, so it proceeds faster towards the end-point. It is an example of autocatalysis
- 5. $2Ag^+ + CH_3CHO + H_2O \rightarrow CH_3COOH + 2H^+ + 2Ag$ or $2Ag(NH_3)_2^+ + CH_3CHO + H_2O \rightarrow CH_3COOH + 2H^+ + 2Ag + 2NH_3$

Topic 15 Exercise 5

1. n = 12 2. 1.37 moldm⁻³ 3. x = 2 4. +5

