CHEMISTRY 1B (CHEM1102) - June 2005

2005-J-2

- All three are based on a cubic unit cell. Simple cubic has atoms stacked directly one on top of the other, least dense 52%, 1 atom/unit cell. BCC has one atom at the centre and 8 x 1/8 atoms at the corners of the unit cell, density 68%, 2 atoms/unit cell. FCC is most efficient packing with 74% density and 4 atoms/unit cell.
- $Ca_5(PO_4)_3OH(s) = 5Ca^{2+}(aq) + 3PO_4^{3-}(aq) + OH^{-}(aq)$

In acidic media, H^+ reacts with both PO_4^{3-} and OH^- to form the conjugate acids and hence shifting the equilibrium to the right.

Fluoridation can replace OH^- forming $Ca_5(PO_4)_3F(s)$. This is less soluble than hydroxyapatite and does not react with H to the same extent as OH^- .

2005-J-3

• Na₂O - basic; Cl_2O - acidic

Na₂O - pH increases; Cl₂O - pH decreases

Sodium oxide is ionic, releasing the very basic oxide ion on dissolution which reacts to form hydroxide. Dichlorine oxide is a covalent molecule and reacts with water to give an oxyacid.

• Alumina is amphoteric and will react with hydroxide to form a soluble complex:

 $Al_2O_3(s) + 2OH^{-}(aq) + 3H_2O(l) \rightarrow 2[Al(OH)_4]^{-}(aq)$

Iron oxide will not react with hydroxide and may now be removed by filtration.

 $2[\operatorname{Al}(\operatorname{OH})_4]^{-}(\operatorname{aq}) + 2\operatorname{CO}_2(g) \rightarrow \operatorname{Al}_2\operatorname{O}_3(s) + 2\operatorname{HCO}_3^{-}(\operatorname{aq}) + 3\operatorname{H}_2\operatorname{O}(l)$

Amphorteric

2005-J-4

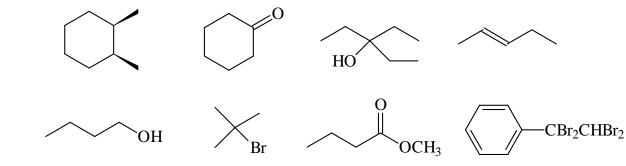
• $[Ni(en)_2(H_2O)_2]^{2+}$ N, O $3d^8$

• Rate = $k[NO]^2[Cl_2]$ $k = 180 \text{ mol}^{-2} \text{ L}^2 \text{ s}^{-1}$ 2005-J-5

• 2.73 4.94 0.024 mol or 1.9 g

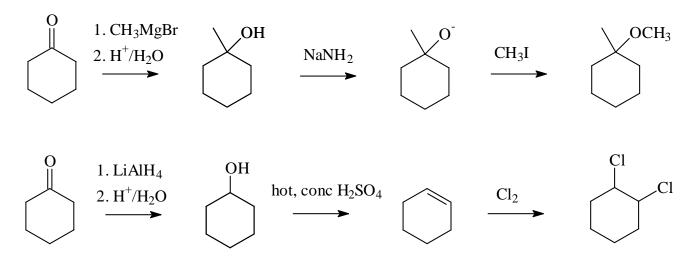
2005-J-6

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2005-J-7

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2005-J-8

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OH⁻ Hot, concentrated H₂SO₄ NaNH₂ CH₃COCl $Cr_2O_7^{2-}/H^+$ • Phenols are less acidic than carboxylic acids. Phenols react with aqueous hydroxide solution to form the phenoxide ion; carboxylic acids react with either aqueous hydroxide or aqueous hydrogen carbonate to form the carboxylate ion. The difference in stability arises from the relative stability of the conjugate base: there is more delocalisation of the charge in the carboxylate ion which is resonance stabilised.

