## CHEM1612 (Chemistry Pharmacy 1B) - November 2007

## 2007-N-2

- -78.2 kJ mol<sup>-1</sup>
- $-350 \text{ kJ mol}^{-1}$

# 2007-N-3

•  $\Delta G^{\circ} = \Delta H^{\circ} - T\Delta S^{\circ}$ = -198.4 - (298 × -187.9) × 10<sup>-3</sup> = -142.4 kJ mol<sup>-1</sup> As  $\Delta G^{\circ}$  is negative, the reaction is spontaneous

To the left

 $K = 9.15 \times 10^{24}$ 

 $T > 1056 {
m K}$ 

# 2007-N-4

• 
$$K = 6.52 \times 10^2$$

86.2%

(i) add more glucose (ii) decrease the temperature

(removing either (or both) of the products isn't really simple)

## 2007-N-5

- $1.40 \times 10^4 \text{ g mol}^{-1}$
- 334 g

## 2007-N-6

0.557 g

decrease

Increased pressure in lungs during deep sea diving causes an increased solubility of all gases in the blood. On ascending too quickly, these gases can bubble out of the blood. This is a serious problem with nitrogen as the bubbles can rupture blood vessels causing "the bends". The He atom is much smaller than the  $N_2$  molecule, has a smaller electron cloud and is less polarisable. It therefore is less soluble in blood than nitrogen and is preferred as the above dangers are reduced.

#### 2007-N-7

• +1.67 kJ mol<sup>-1</sup>

 $+7.37 \text{ kJ mol}^{-1}$ 

To the left. As written, the reaction is non-spontaneous (as  $\Delta G^{\circ} > 0$ ). It must therefore run spontaneously in the opposite direction.



extent of reaction



- $1.0 \times 10^{-7} \text{ M}$
- •



dichlorobisethylenediaminecobalt(III) ion

 sodium hexafluoroaluminate(III) pentaamminebromocobalt(III) sulfate  ${}^{186}_{74}W + 2{}^{1}_{0}n \rightarrow {}^{188}_{74}W \rightarrow {}^{188}_{75}\text{Re} + {}^{0}_{-1}\beta$   ${}^{186}_{74}W + 2{}^{1}_{1}p \rightarrow {}^{188}_{76}\text{Os} \rightarrow {}^{188}_{75}\text{Re} + {}^{0}_{+1}\beta$ 

 $^{188}_{76}$ Os is a stable nuclide so proton bombardment doesn't work.

Neutron production is much cheaper than proton production.

Neutron bombardment is much easier reaction as the positively charged nucleus repels the positively charged protons.

 $1.36 \times 10^{-6}$  g

## 2007-N-10

- $1.1 \times 10^3$  minutes
- 0.86 mol

### 2007-N-11

• 0.51 V

0.24 V

(b)  $Ag^+(aq)$  ions are produced. They will react with the  $Cl^-(aq)$  ions in solution and a white precipitate of AgCl will be seen to form around the electrode.

## 2007-N-12

- $Zn^{2+}(aq) + 4CN^{-}(aq) \implies [Zn(CN)_4]^{2-}(aq)$  $7.8 \times 10^{-23} M$
- Colloids can be stabilised electrostatically like charges on the surface of the particles cause the particles to repel each other. The addition of electrolytes neutralises these surface charges and the particles can now come together.

#### 2007-N-13

• Rate =  $k[NO_2]^2[H_2]$ 2.9 × 10<sup>2</sup> M<sup>-2</sup> s<sup>-1</sup> 4.1 × 10<sup>-5</sup> mol L<sup>-1</sup> s<sup>-1</sup> Step 1: 2NO(g)  $\iff N_2O_2(g)$  fast equilibrium with  $K = [N_2O_2] / [NO]^2$   $K = [N_2O_2] / [NO]^2 \implies [N_2O_2] = K[NO]^2$ Step 2:  $N_2O_2(g) + H_2(g) \rightarrow N_2O(g) + H_2O(g)$  slow (rate determining) Rate =  $k[N_2O_2][H_2]$   $= kK[NO]^2[H_2]$ which is consistent with rate law of Rate =  $k[NO]^2[H_2]$ .