Intermolecular Forces

1. Arrange the following in order of increasing boiling point. Explain your reasoning.

D) CH₃CH₂CH₂Cl E) CH₃CH₂CH₂OH A) CH₄ B) CH₃CH₂CH₂CH₃

2. What is the change in enthalpy when 47.5 g of water freezes? ($\Delta H_{fus} = 6.02 \text{ kJ/mol for H}_2\text{O}$)

- 3. Platinum crystallizes with the face centered cubic unit cell. The radius of a platinum atom is 139 pm. Calculate the edge length of the unit cell and the density of platinum in g/cm³.
- 4. NaCl crystalizes in a face centerd cubic lattice of chloride ions, with the smaller sodium ions occupying holes between the chloride ions. How many Cl⁻ ions are in contact with any single Na⁺ ion?
 - a. 4

b. 6

c. 8

d. 12

Solutions

1. An aqueous NaCl solution is made using 133 g of NaCl diluted to a total solution volume of 1.00 L. Calculate the molarity, molality, and mass percent of the solution. (Assume the solution has a density of 1.08 g/mL.)

2. A solution contains 55.8 g of glucose ($C_6H_{12}O_6$) in 455 g of water. What is the freezing point and boiling point of the solution?

For water: f.p. = 0.000°C

 $K_f = 1.858^{\circ}\text{C/}m$

b.p. = 100.000°C $K_b = 0.512$ °C/m

Kinetics

1. For the reaction given below, express the rate of reaction with respect to each of the reactants and products.

$$2 N_2(g) + 5 O_2(g) \rightarrow 2 N_2O_5(g)$$

2. Given the following experimental data, find the rate law and the rate constant for the reaction:

$$NO(g) + NO_2(g) + O_2(g) \rightarrow N_2O_5(g)$$

Run	[NO] ₀	[NO ₂] ₀	$[O_2]_0$	Initial Rate (M·s ⁻¹)
1	0.10 M	0.10 M	0.10 M	2.1 x 10 ⁻²
2	0.20 M	0.10 M	0.10 M	4.2 x 10 ⁻²
3	0.20 M	0.30 M	0.10 M	1.26 x 10 ⁻¹
4	0.10 M	0.10 M	0.20 M	2.1 x 10 ⁻²

What is the overall reaction order?

- 3. Indicate the order of reaction for each of the following observations.
 - a. A plot of the concentration of the reactant versus time which yields a straight line.
 - b. Reactions having a half-life that is independent of initial concentration.
 - b. A plot of the reciprocal of the concentration versus time yields a straight line.
- 4. The rate law for the reaction $A + B \rightarrow C + D$ is first order in [A] and second order in [B]. If [A] is halved and [B] is doubled, wthe rate of the reaction will
 - a. Remain the same.

- b. Be increased by a factor of 2
- c. Be increased by a factor of 4
- d. Be increased by a factor of 8

5. Consider the following overall reaction which is experimentally observed to be second order in AB and zero order in C.

$$AB + C \rightarrow A + BC$$

Determine whether the mechanism below is valid for this reaction

$$AB + AB \xrightarrow{k_1} AB_2 + A$$
 (slow)

$$AB_2 + C \xrightarrow{k_2} AB + BC$$
 (fast)

Equilibrium

1. Consider the following exothermic reaction:

$$C_2H_4(g) + CI_2(g) \rightleftarrows C_2H_4CI_2(g)$$

If you were a chemist trying to maximize the amount of C₂H₄Cl₂ produced, choose all of the following that create the desired shift in equilibrium. Assume that the reaction mixture reaches equilibrium.

- A. Increasing the reaction volume
- b. Removing C₂H₄Cl₂ from the reaction mixture as it forms.
- c. Lowering the reaction temperature d. Adding Cl₂

2. The reversible reaction

$$CO(g) + 2 H_2(g) \rightleftharpoons CH_3OH(g) + heat$$

is carried out by mixing carbon monoxide and hydrogen gases in a closed vessel under high pressure with a suitable catalyst. After equilibrium established at high temperature and pressure, all three substances are present. If the pressure on the system is lowered, with the temperature held constant, what will be the result?

- a. The amount of CH₃OH will increase.
- b. The amount of CH₃OH will decrease.
- c. The amount of each substance will be unchanged d. The amount of each substance will increase.
- 3. Carbon monoxide gas reacts with hydrogen gas at elevated temperatures to form methanol according to this equation:

$$CO(g) + 2 H_2(g) \rightleftharpoons CH_3OH(g) + heat$$

When 0.40 mol of CO and 0.30 mol of H₂ are allowed to reach equilibrium in a 1.0 L container, 0.060 mol of CH₃OH are formed. What is the value of K_c ?

- a. 0.50
- b. 0.98
- c. 1.7
- d. 5.4

Acid and Base Chemistry

1. Classify each acid as strong or weak. If the acid is weak, write an expression for the acid ionization constant (K_a)

a. HF

b. HCl

c. H₂SO₄

d. H₂CO₃

2. Determine the percent of ionization of a 0.225 M solution of benzoic acid ($K_a = 6.5 \times 10^{-5}$)?

3. Determine the pH of the following solutions:

a. 0.0195 M KOH

b. 0.318 M KC₆H₅O (K_a phenol = 1.3×10^{-10})

Aqueous Ionic Equilibrium

- 1. Consider a titration of a 35.0 mL sample of 0.175 M CH_3COOH (K_a = 1.800×10^{-5}) is titrated with 0.200 M KOH. Determine:
 - a. Initial pH
 - b. equivalence volume
 - c. pH when 10.0 mL 0.200 M KOH has been added
 - d. pH at the equivalence point
 - E. pH at 5.0 mL past the equivalence point
 - F. Draw a rough graph of what the titration curve expected.

2. If $Q_{sp} > K_{sp}$ is the solution saturated, unsaturated, or supersaturated?

Free Energy and Thermodynamics

1. Calculate ΔS°_{surr} at the given temperature for a reaction with the following conditions:

 $\Delta H_{rxn}^{o} = -287 \text{ kJ}; T = 298 \text{ K}$

2. Determine the value of ΔH^o_{rxn} , ΔS^o_{rxn} and ΔG^o_{rxn} at 25°C and the spontaneity. If nonspontaneous, is it possible that a change in temperature will make it spontaneous? If so, should the temperature be raised or lowered to accomplish spontaneity?

 $3 H_2 (g) + Fe_2O_3 (s) \rightarrow 2 Fe (s) + 3 H_2O (g)$

	ΔH_f^o (kJ/mol):	ΔS ^o (J/mol·K):	
H ₂ (g)	0.0	130.7	
Fe ₂ O ₃ (s)	-824.2	87.4	
Fe(s)	0.0	27.3	
H ₂ O (g)	-241.8	188.8	

3. Find the equilibrium constant for the following reaction at 25°C:

$$2 H_2S(g) \leftrightarrows 2 H_2(g) + S_2(g)$$

A.
$$\Delta G_{rxn}^{o} = -33.4 \text{ kJ/mo}$$

$$h AG^0 = 0.0 \, kI/mo$$

A.
$$\Delta G_{rxn}^o = -33.4 \text{ kJ/mol}$$
 b. $\Delta G_{rxn}^o = 0.0 \text{ kJ/mol}$ c. $\Delta G_{rxn}^o = 79.7 \text{ kJ/mol}$

Electrochemistry

1. Balance the following redox reaction in acidic solution:

$$BrO_3^-(aq) + N_2H_4(g) \rightarrow Br^-(aq) + N_2(g)$$

2. Sketch the cell for the following reaction:

$$Cr(s) + Fe^{3+}(aq) \rightarrow Fe^{2+}(aq) + Cr^{3+}(aq)$$

3. Express the following reaction in cell notation:

$$2 \text{ Ag}^+(aq) + Pb (s) \rightarrow 2 \text{ Ag}(s) + Pb^{2+}(aq)$$

4. Balance the following reaction in basic solution:

$$Al(s) + MnO_4(aq) \rightarrow MnO_2(s) + Al(OH)_4(aq)$$

5. Calculate E^o_{cell} for the following redox reaction and determine spontaneity. 2 Cu (s) + Mn²⁺(aq) \rightarrow 2 Cu⁺ (aq) + Mn (s)

6. Which of the following metal cations is the best oxidizing agent? Why?

Nuclear Chemistry

1. One of the nuclides in spent nuclear fuel is U-235, an alpha emitter with a half-life of 703 million years. How long would it take for the amount of U-235 to reach one-eighth of its initial mass? What is the order of this process?

2. Write a nuclear equation for the fusion of two H-2 atoms to form He-3 and one neutron.

3. If 1.0 g of matter were converted to energy, how much energy would be formed?

Answer Key:

Intermolecular forces

1. A<C<B<D<E. Boiling point increases with increasing intermolecular forces. The molecules increase in their intermolecular forces as follows: A - dispersion forces; forces (broader electron cloud); C - dispersion forces over a condensed area. B - stronger dispersion D - dispersion forces and dipole-dipole interactions; E - dispersion forces, dipole-dipole interactions, and hydrogen bonding.

2. - 15.9 kJ

$$47.5 \text{ H}_2\text{O} \times \frac{1 \text{ mol H}_2\text{O}}{18.016 \text{ g H}_2\text{O}} \times \frac{6.02 \text{ kJ}}{1 \text{ mol H}_2\text{O}} = -15.9 \text{ kJ}$$

3. $l=393 \text{ pm}, d=21.3 \text{ g/cm}^3$

$$l = 2\sqrt{2} * r = 139 \times 10^{-12} * 2\sqrt{2} = 393 \text{ pm}$$

$$V = l^3 = \left(\frac{393*10^{-12}}{10^{-2}}\right)^3 = 6.0698 * 10^{-23} \text{cm}^3$$

$$\text{mass of unit cell} = \frac{195.08 \text{ g}}{6.022*10^{23} \text{ atoms}} = \frac{3.239*10^{-22} \text{ g}}{\text{atom}} \times \frac{4 \text{ atoms}}{1 \text{ cell}} = 1.296*10^{-21} \text{g}$$

$$\text{Density} = \frac{\text{mass}}{\text{Volume}} = \frac{1.296*10^{-21}}{6.0698*10^{-23}} = 21.3 \frac{\text{g}}{\text{cm}^3}$$

4. b.

Solutions

1. moles NaCl = 133g NaCl / 58.44 g/mol = 2.28 mol

$$M = mol/L 2.28 mol/ 1 L = 2.28M,$$

$$m = \text{mol solute / kg solvent} = 2.28 / (1.080 \text{ kg} - .133 \text{ kg}) = 2.41 m$$

% mass = g solute/grams solution * 100% = .133/1.08 * 100% = 12.3% by mass

2. molar mass = 55.8/180.156 g/mol = 0.3097

$$0.3097/0.455 = 0.6807 \ molal$$

$$\Delta T = m \times K_f = .6807 * 1.86$$

$$\Delta T = m \times K_b = .6807 * 0.512$$

Kinetics

1.
$$2 N_2(g) + 5 O_2(g) \rightarrow 2 N_2 O_5(g)$$

Rate =
$$-\frac{1}{2}\frac{\Delta[N_2]}{\Delta t}$$
 = $-\frac{1}{5}\frac{\Delta[O_2]}{\Delta t}$ = $+\frac{1}{2}\frac{\Delta[N_2O_5]}{\Delta t}$

2. Rate = $k[NO][NO_2]$

$$\frac{Rate}{[NO][NO_2]} = k = 2.1 \text{ M}^{-1} \cdot \text{s}^{-1}$$

2nd order reaction.

- 3. a. zero order
- b. first order c. second order

- 4. b.
- 5. Yes it's valid.

Equilibrium

- 1. B, C, D
- 2. b.
- 3. d. $0.060/(0.34*0.18^2) = 5.4$

Acid and Base Chemistry

- 1. Classify each acid as strong or weak. If the acid is weak, write an expression for the acid ionization constant (Ka). If Ka is available, it is a weak acid.
 - a. HF: weak acid

$$\label{eq:Ka} \begin{split} &HF\ (aq)+H_2O\ (I)\rightarrow H_3O^+\ (aq)+F^-\ (aq)\\ K_a=\frac{[H3O^+\][F^-]}{[HF]} \end{split}$$

- b. HCI: strong acid
- c. H₂SO₄: strong acid
- d. H₂CO₃:

$$H_2CO_3$$
 (aq) + H_2O (I) \rightarrow H_3O^+ (aq) + HCO_3^- (aq)
 $K_a = \frac{[H3O+][H2CO_3^-]}{[H2CO_3]}$

2. Find the % ionization of a 0.225 M benzoic acid solution.

H ₂ O	$C_6H_5CO_2H$	H₃O⁺	C ₆ H ₅ CO ₂ -
1	0.225M	0	0
С	-X	+χ	+χ
E	0.225-x	X	X

$$K_{\text{a}} = \frac{\text{[H3O+][HSO3-]}}{\text{[H2SO3]}} = \frac{\text{[x^2]}}{\text{[0.225-x]}} = 6.5 \times 10^{-5}$$

Assume x is small. x= .00382

Get percent ionization.

% ionization =
$$\frac{concentration\ of\ ionized\ acid}{initial\ concentration\ of\ acid} \times 100\% = \frac{0.00383}{0.225} \times 100\% = 1.70\%$$

3. a.
$$pH = -log(H_3O^+) = -log(0.0195) = 12.29$$

b. Get
$$K_b = \frac{K_w}{K_a} = \frac{1 \times 10^{-14}}{1.3 \times 10^{-10}} = 7.7 \times 10^{-5}$$
. Ice table to get $x = 4.95 \times 10^{-3}$. pOH = -log (x) = 2.31, pH = 14- pOH = 14- 2.31 = 11.69

Aqueous Ionic Equilibrium

1. a.
$$1.800 \times 10^{-5} = \frac{x^2}{0.175}$$
 $x = 0.010141$

b.
$$mol CH_3COOH = 0.0350L \times 0.175M = 6.125 \times 10^{-3} mol CH_3COOH = mol NaOH$$

$$6.125 \times 10^{-3} \times \frac{1 L}{0.200 \ mol} \times \frac{1000 \ mL}{1 \ L} = 30.625 \ mL = 30.6 \ mL$$

c. initial moles CH₃COOH - moles NaOH = remaining moles CH₃COOH

 $0.0350L \times 0.175M - 0.0100L \times 0.200M = 0.004125 \ mol \ CH_3COOH$

$$[CH_3COOH] = \frac{0.004125 \, mol}{0.0350L + 0.0100L} = 0.916667M$$

$$0.0100L \times 0.200M \ NaOH = 0.000200 \ mol \ NaOH \times \frac{1 \ mol \ CH_3COO^-}{1 \ mol \ NaOH} \times \frac{1}{0.0450L} = 0.0444M$$

$$pH = -\log(1.800 \times 10^{-5}) - \log\left(\frac{0.0444M}{0.091667M}\right) = 4.685$$

d. $6.125 \times 10^{-3} \text{ mol CH}_3\text{COO}^-\text{ produced at equivalence point; } [\text{CH}_3\text{COO}^-] = \frac{6.125 \times 10^{-3} mol}{0.0656L} = 0.09337 \text{M}$

$$K_b = \frac{1.00 \times 10^{-14}}{1.800 \times 10^{-5}} = \frac{x^2}{0.09337\text{M}}$$
 $x = [OH^-] = 7.02 \times 10^{-6}$

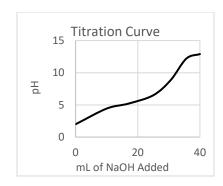
$$pH = 14 - (-log[OH^{-}]) = 14 - (-log(1.02x10^{-6}) = 8.857)$$

e. 0.200 M * 0.0356 L = 0.000712 mol NaOH

 $0.000712 \text{ mol NaOH} - 6.125 \text{ x } 10^{-3} \text{ mol CH}_3\text{COOH} = 9.95\text{x} 10^{-4} \text{ mol NaOH remaining}$

$$[OH^-] = \frac{9.95 \times 10^{-4} mol}{0.0656} = 0.015213$$
 $pH = 14 - (-\log(0.15213)) = 12.182$

f.



2. The solution is supersaturated. Solid will begin to precipitate out of solution when Q > K_{sp} or a seed crystal could cause a precipitate to form if the supersaturated solution was heated to allow increased solubility.

Free Energy and Thermodynamics

Spontaneity:

ΔΗ	ΔS	High Temp	Low Temp
-	+	spontaneous	Spontaneous
+	-	nonspontaneous	nonspontaneous
-	-	nonspontaneous	Spontaneous
+	+	spontaneous	nonspontaneous

Mathematically:

$$\Delta G_f^o = \Delta H_f^o - T\Delta S_f^o$$

If ΔH is negative and you are subtracting a positive number, the result must be negative (spontaneous).

If ΔH is positive and you are subtracting a negative number (+# - -#), the result must be positive (nonspontaneous).

1.
$$\Delta S_{surr}^o = \frac{-\Delta H_{sys}^o}{T} \Rightarrow \Delta S_{surr}^o = -\frac{-287000}{298} = 963 \frac{J}{K}$$

2.
$$\Delta H_{rxn}^o = \sum n_p \times \Delta H_f^o(products) - \sum n_r \times \Delta H_f^o(reactants)$$

$$\Delta H_{ryn}^{o} = +98.8 \text{ kJ}$$

$$\Delta S_{rxn}^o = \sum n_p \times \Delta S_f^o(products) - \sum n_r \times \Delta S_f^o(reactants)$$

$$\Delta S_{rxn}^o$$
 = +141.5 J/K T= 25°C + 273.15 = 298 K

 $\Delta G^o = \Delta H^o_{rxn}$ -T ΔS^o_{rxn} $\Delta G^o = 98800 \, \text{J}$ - (298 K) (141.5 $\frac{J}{K}$) = +56.6 kJ. Reaction is non-spontaneous. The reaction can be made spontaneous by increasing the temperature.

$$\Delta G_{rxn}^o = \sum n_p \times \Delta G_f^o(products) - \sum n_r \times \Delta G_f^o(reactants)$$

$$\Delta G_{rxn}^o = [2(0.0) + 1(79.7)] - [2(-33.4)] = +146.5 \text{ kJ}$$

3.
$$\Delta G_{rxn}^o = -RT \ln K$$
 a. $K_{eq} = e^{\frac{-\Delta G_{rxn}^o}{RT}} = e^{\frac{-33400 \text{J}}{\left(8.314 \frac{J}{K \cdot mol}\right)^{(298K)}}} = 7.16 \times 10^5$

$$\text{b. } K_{eq} = e^{\frac{-\Delta G_{TXN}^0}{RT}} = \ e^{\frac{0J}{\left(8.314\frac{J}{K\cdot mol}\right)(298K)}} = \ 1 \qquad \text{c. } K_{eq} = e^{\frac{-\Delta G_{TXN}^0}{RT}} = \ e^{\frac{79700J}{\left(8.314\frac{J}{K\cdot mol}\right)(298K)}} = \ 1.07\times 10^{-14}$$

Electrochemistry

1.
$$BrO_3^-(aq) + N_2H_4(g) \rightarrow Br^-(aq) + N_2(g)$$

1/2 RXNS:

$$BrO_3^-(aq) \rightarrow Br^-(aq)$$

$$N_2H_4(g) \rightarrow N_2(g)$$

Balance O's, Add H₂O:

$$BrO_3^-(aq) \rightarrow Br^-(aq) + 3 H_2O(I)$$

Balance H w/H+:

$$6H^{+}(aq) + BrO_{3}^{-}(aq) \rightarrow Br^{-}(aq) + 3 H_{2}O(I)$$

$$N_2H_4(g) \rightarrow N_2(g) + 4 H^+(aq)$$

Add electrons:

$$6H^{+}(aq) + BrO_{3}^{-}(aq) + 6 e^{-} \rightarrow Br^{-}(aq) + 3 H_{2}O(I)$$

$$N_2H_4(g) \rightarrow N_2(g) + 4 H^+(aq) + 4 e^{-1}$$

Equalize electrons:

$$12H^{+}(aq) + 2BrO_{3}^{-}(aq) + 12e^{-} \rightarrow 2Br^{-}(aq) + 6H_{2}O(l)$$

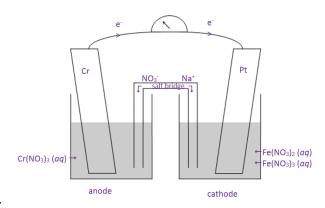
$$3N_2H_4(g) \rightarrow 3N_2(g) + 12 H^+(aq) + 12 e^{-1}$$

Add ½ rxns:

$$3N_2H_4(g) + 12H^+(aq) + 2BrO_3^-(aq) + 12e^- \rightarrow 2Br^-(aq) + 6H_2O(I) + 3N_2(g) + 12H^+(aq) + 12e^-$$

Cancel stuff (12 H^+ , 12 e^- leaves):

$$3N_2H_4(g) + 2BrO_3^-(aq) \rightarrow 2Br^-(aq) + 6H_2O(l) + 3N_2(g)$$



2.

3. $2 \text{ Ag}^+(aq) + \text{Pb (s)} \rightarrow 2 \text{ Ag(s)} + \text{Pb}^{2+}(aq)$ in cell notation:

Pb (s) | Pb²⁺ (aq) | Ag⁺ (aq) | Ag(s)

4. Al (s) +MnO₄ (aq) \rightarrow MnO₂(s) + Al(OH)₄ (aq)

Two ½ RXNS: MnO_4^- (aq) $\rightarrow MnO_2$ (s) \underline{AND} Al(s) \rightarrow Al(OH)₄ (aq)

Balance Everything but O and H:

already done

Balance O by adding H₂O:

$$MnO_4^-(aq) \to MnO_2(s) + 2 H_2O(l)$$

Al(s) + 4 H₂O(l) \to Al(OH)₄-(aq)

Balance H w/H+:

$$4 \text{ H}^+ (aq) + \text{MnO}_4^- (aq) \rightarrow \text{MnO}_2 (s) + 2 \text{ H}_2\text{O (l)}$$

 $Al(s) + 4 \text{ H}_2\text{O (l)} \rightarrow Al(OH)_4^- (aq) + 4 \text{ H}^+ (aq)$

Neutralize H+ with OH-, adding the same number of OH- to each side of the equations:

$$4 \text{ OH}^{-}(aq) + 4 \text{ H}^{+}(aq) + \text{MnO}_{4}^{-}(aq) \rightarrow \text{MnO}_{2}(s) + 2 \text{ H}_{2}\text{O}(l) + 4 \text{ OH}^{-}(aq)$$

 $4 \text{ OH}^{-}(aq) + \text{Al}(s) + 4 \text{ H}_{2}\text{O}(l) \rightarrow \text{Al}(OH)_{4}^{-}(aq) + 4 \text{ H}^{+}(aq) + 4 \text{ OH}^{-}(aq)$

Cancel waters:

2
$$H_2O(I) + MnO_4^-(aq) \rightarrow MnO_2(s) + 4 OH^-(aq)$$

4 $OH^-(aq) + AI(s) \rightarrow AI(OH)_4^-(aq) + 2 H_2O(I)$

Equalize charge by adding electrons:

$$3 e^{-} + 2 H_2O (I) + MnO_4^{-} (aq) \rightarrow MnO_2 (s) + 4 OH^{-} (aq)$$

 $4 OH^{-} (aq) + AI(s) \rightarrow AI(OH)_4^{-} (aq) + 3 e^{-}$

Add ½ rxns:

$$3 e^{-} + 2 H_2O(I) + MnO_4^{-}(aq) + 4 OH^{-}(aq) + Al(s) \rightarrow MnO_2(s) + 4 OH^{-}(aq) + Al(OH)_4^{-}(aq) + 3 e^{-}$$

Cancel stuff (4 OH and 3 e leaves):

$$2 H_2O(I) + MnO_4(aq) + Al(s) \rightarrow MnO_2(s) + Al(OH)_4(aq)$$

5. E^ocell and spontaneity for:

$$2 \text{ Cu } (s) + \text{Mn}^{2+}(aq) \rightarrow 2 \text{ Cu}^{+}(aq) + \text{Mn } (s)$$

2 ½ rxns w/electrons:

2 Cu (s) + 2
$$e^{-} \rightarrow 2 Cu^{+} (aq)$$

$$Mn^{2+}(aq) \to Mn(s) + 2e^{-}$$

Look up potentials:

Mn reduced,
$$E^{\circ}_{red}$$
 = -1.18 V

Cu oxidized,
$$E^{\circ}_{ox} = -E^{\circ}_{red} = -0.52V$$

$$E^{\circ}_{cell} = E^{\circ}_{ox} + E^{\circ}_{red} = -1.70V$$

Negative E°_{cell}, nonspontaneous.

6. Pb²⁺ is the strongest oxidizing agent because it has the highest E^o (most positive in the table).

Relevant equation:

$$\Delta G = -nFE_{cell}^o$$
F = 96,485 C/mol e⁻

Nuclear Chemistry

- 1. 2.11 x 10⁹ yr, 1st order.
- 2. ${}_{1}^{2}H + {}_{1}^{2}H \rightarrow {}_{2}^{3}He + {}_{0}^{1}n$
- 3. $E = mc^2$, $c = 2.998*10^8$

$$1/1000 * (2.998*10^8)^2 = 8.988*10^{13} J = 9.0 \times 10^{13} J$$