Name\_\_\_\_\_

#### MULTIPLE CHOICE. Choose the one alternative that best completes the statement or answers the question.

- 1) At equilibrium, \_\_\_\_\_.
  - A) the rates of the forward and reverse reactions are equal
  - B) the rate constants of the forward and reverse reactions are equal
  - C) all chemical reactions have ceased
  - D) the value of the equilibrium constant is 1
  - E) the limiting reagent has been consumed
- 2) Which one of the following will change the value of an equilibrium constant?
  - A) adding other substances that do not react with any of the species involved in the equilibrium
  - B) varying the initial concentrations of reactants
  - C) changing temperature
  - D) varying the initial concentrations of products
  - E) changing the volume of the reaction vessel
- 3) The value of  $K_{eq}$  for the following reaction is 0.25:

 $SO_2(g) + NO_2(g) \rightleftharpoons SO_3(g) + NO(g)$ 

The value of  $K_{eq}$  at the same temperature for the reaction below is \_\_\_\_\_.

$$2SO_2(g) + 2NO_2(g) \rightleftharpoons 2SO_3(g) + 2NO(g)$$
  
A) 0.062 B) 16 C) 0.25 D) 0.50 E) 0.12

4) The value of  $K_{eq}$  for the equilibrium

$$H_2(g) + I_2(g) \rightleftharpoons 2 HI(g)$$

is 794 at 25°C. At this temperature, what is the value of Keq for the equilibrium below?

HI (g) 
$$\rightleftharpoons 1/2$$
 H<sub>2</sub> (g) + 1/2 I<sub>2</sub> (g)  
A) 0.035 B) 0.0013 C) 28 D) 397 E) 1588

5) The K<sub>eq</sub> for the equilibrium below is  $7.52 \times 10^{-2}$  at  $480^{\circ}$ C.

$$2Cl_2(g) + 2H2O(g) \implies 4HCl(g) + O_2(g)$$

What is the value of  $K_{\mbox{eq}}$  at this temperature for the following reaction?

Cl<sub>2</sub> (g) + H<sub>2</sub>O (g) 
$$\rightleftharpoons$$
 2HCl (g) +  $\frac{1}{2}$ O<sub>2</sub> (g)  
A) 0.150 B) 0.274 C) 0.0376 D) 5.66 × 10<sup>-3</sup> E) 0.0752

6) At 1000 K, the equilibrium constant for the reaction

$$2NO(g) + Br_2(g) \implies 2NOBr(g)$$

is  $K_p = 0.013$ . Calculate  $K_p$  for the reverse reaction,

2NOBr (g) 
$$\rightleftharpoons$$
 2NO (g) + Br<sub>2</sub> (g).  
A) 0.99 B) 1.1 C) 0.013 D) 1.6 × 10<sup>-4</sup> E) 77

7) The expression for K<sub>eq</sub> for the reaction below is \_\_\_\_\_.

$$4$$
CuO (s) + CH<sub>4</sub> (g)  $\rightleftharpoons$  CO<sub>2</sub> (g) + 4Cu (s) + 2H<sub>2</sub>O (g)

A) 
$$\frac{P_{CO_2} P_{H_2O}^2}{P_{CH_4}}$$
  
B) 
$$\frac{P_{CH_4}}{P_{H_2O}^2 P_{CO_2}}$$
  
C) 
$$\frac{[Cu] P_{CO_2} P_{H_2O}^2}{[CuO]^4 P_{CH_4}}$$
  
D) 
$$\frac{P_{CH_4}}{P_{CO_2} P_{H_2}^2}$$
  
E) 
$$\frac{P_{CO_2} P_{H_2O}^2}{P_{CuO}}$$

8) Acetic acid is a weak acid that dissociates into the acetate ion and a proton in aqueous solution:

$$HC_{2}H_{3}O_{2}(aq) \rightleftharpoons C_{2}H_{3}O_{2}G(aq) + H^{+}(aq)$$

At equilibrium at 25°C a 0.100 M solution of acetic acid has the following concentrations: [HC2H3O2] = 0.0990 M, [C2H3O2G] =  $1.33 \times 10^{-3}$  M, and [H+] =  $1.33 \times 10^{-3}$  M. The equilibrium constant, K<sub>eq</sub>, for the ionization of acetic acid at 25°C is \_\_\_\_\_.

A) 
$$5.71 \times 10^4$$
 B)  $1.79 \times 10^{-5}$  C)  $1.75 \times 10^{-7}$  D)  $5.71 \times 10^6$  E) 0.100

9) At 200°C, the equilibrium constant for the reaction below is  $2.40 \times 10^3$ .

 $2NO(g) \Longrightarrow N_2(g) + O_2(g)$ 

A closed vessel is charged with 36.1 atm of NO. At equilibrium, the partial pressure of O<sub>2</sub> is \_\_\_\_\_\_ atm. A) 35.7 B) 18.1 C)  $1.50 \times 10^{-2}$  D) 6.00 E) 294

10) How is the reaction quotient used to determine whether a system is at equilibrium?

A) The reaction is at equilibrium when  $Q < K_{eq}$ .

B) The reaction is at equilibrium when  $Q > K_{eq}$ .

C) At equilibrium, the reaction quotient is undefined.

D) The reaction quotient must be satisfied for equilibrium to be achieved.

E) The reaction is at equilibrium when  $Q = K_{eq}$ .

11) In the coal-gasification process, carbon monoxide is converted to carbon dioxide via the following reaction:

 $CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$ 

In an experiment, 0.35 mol of CO and 0.40 mol of H<sub>2</sub>O were placed in a 1.00–L reaction vessel. At equilibrium, there were 0.19 mol of CO remaining. K<sub>eq</sub> at the temperature of the experiment is \_\_\_\_\_\_.

A) 0.75 B) 1.0 C) 5.47 D) 1.78	E) 0.56
--------------------------------	---------

12) Nitrosyl bromide decomposes according to the following equation.

 $2NOBr(g) \rightleftharpoons 2NO(g) + Br_2(g)$ 

A sample of NOBr (0.64 mol) was placed in a 1.00–L flask containing no NO or Br<sub>2</sub>. At equilibrium the flask contained 0.46 mol of NOBr. How many moles of NO and Br<sub>2</sub>, respectively, are in the flask at equilibrium?

A) 0.46, 0.23 B) 0.18, 0.090 C) 0.46, 0.46 D) 0.18, 0.360 E) 0	).18, 0.18
--	------------

- 13) In which of the following reactions would increasing pressure at constant temperature <u>not</u> change the concentrations of reactants and products, based on Le Chatelier's principle?
  - A)  $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ B)  $2N_2(g) + O_2(g) \rightleftharpoons 2N_2O(g)$ C)  $N_2(g) + 2O_2(g) \rightleftharpoons 2NO_2(g)$ D)  $N_2O_4(g) \rightleftharpoons 2NO_2(g)$ E)  $N_2(g) + O_2(g) \rightleftharpoons 2NO(g)$

14) Consider the following reaction at equilibrium:

 $2CO_2(g) \Longrightarrow 2CO(g) + O_2(g) \qquad \Delta H^\circ = -514 \text{ kJ}$ 

Le Chatelier's principle predicts that adding  $O_2(g)$  to the reaction container will \_\_\_\_\_\_.

A) decrease the partial pressure of CO<sub>2</sub> (g) at equilibrium

B) decrease the value of the equilibrium constant

C) increase the partial pressure of  $CO_2$  (g) at equilibrium

D) increase the value of the equilibrium constant

E) increase the partial pressure of CO (g) at equilibrium

15) Consider the following reaction at equilibrium:

 $2CO_2(g) \implies 2CO(g) + O_2(g) \qquad \Delta H^\circ = -514 \text{ kJ}$ 

Le Chatelier's principle predicts that an increase in temperature will \_\_\_\_\_\_.

A) increase the partial pressure of O<sub>2</sub> (g)

- B) decrease the value of the equilibrium constant
- C) increase the partial pressure of CO
- D) decrease the partial pressure of  $CO_2(g)$
- E) increase the value of the equilibrium constant
- 16) Consider the following reaction at equilibrium.

 $2CO_2(g) \rightleftharpoons 2CO(g) + O_2(g) \qquad \Delta H^\circ = -514 \text{ kJ}$ 

Le Chatelier's principle predicts that the equilibrium partial pressure of CO (g) can be maximized by carrying out the reaction \_\_\_\_\_.

- A) at high temperature and high pressure
- B) at high temperature and low pressure
- C) at low temperature and low pressure
- D) at low temperature and high pressure
- E) in the presence of solid carbon

17) The effect of a catalyst on an equilibrium is to \_\_\_\_\_

- A) increase the rate at which equilibrium is achieved without changing the composition of the equilibrium mixture
- B) increase the rate of the forward reaction only
- C) shift the equilibrium to the right
- D) increase the equilibrium constant so that products are favored
- E) slow the reverse reaction only

18) The following equilibrium is readily established:

 $SO_2Cl_2(g) \rightarrow SO_2(g) + Cl_2(g)$ 

At equilibrium at 373 K, a 1.00-L reaction vessel contains 0.0106 mol of  $SO_2Cl_2$  and 0.0287 mol each of  $SO_2$  and  $Cl_2$ . What is  $K_{eq}$  for the reaction at 373 K?

A) 12.8 B) 2.72 C) 0.0781 D) 2.39 E) 0.418

19) Dinitrogentetraoxide partially decomposes according to the following equilibrium:

 $N_2O_4(g) \rightarrow 2NO_2(g)$ 

A 1.000–L flask is charged with  $3.00 \times 10^{-2}$  mol of N<sub>2</sub>O<sub>4</sub>. At equilibrium,  $2.36 \times 10^{-2}$  mol of N<sub>2</sub>O<sub>4</sub> remains. K<sub>eq</sub> for this reaction is \_\_\_\_\_\_.

A) 0.723 B) 1.92 × 10<sup>-4</sup> C) 6.93 × 10<sup>-3</sup>

- D) 0.391
- E) 0.212

20) The K<sub>eq</sub> for the reation below is  $1.49 \times 108$  at  $100^{\circ}$ C:

$$CO(g) + Cl_2(g) \rightarrow COCl_2(g)$$

In an equilibrium mixture of the three gases,  $P_{CO} = P_{Cl_2} = 8.60 \times 10^{-4}$  atm. The partial pressure of the product, phosgene (COCl<sub>2</sub>), is \_\_\_\_\_\_ atm. A)  $2.01 \times 10^{14}$  B)  $1.72 \times 10^{11}$  C)  $1.28 \times 10^5$  D)  $4.96 \times 10^{-15}$  E)  $1.10 \times 10^2$ 

21) Phosphorous trichloride and phosphorous pentachloride equilibrate in the presence of molecular chlorine according to the reaction:

 $PCl_3(g) + Cl_2(g) \rightarrow PCl_5(g)$ 

 $K_{eq} = 2.01$  at 500 K. A 1.000-L reaction vessel is charged with 0.990 mol of PCl<sub>5</sub> and allowed to equilibrate at this temperature. The equilibrium partial pressure of PCl<sub>3</sub> is \_\_\_\_\_ atm.

A) 0.702 B) 4.25 C) 4.50 D) 36.4 E) 0.496

22) Phosphorous trichloride and phosphorous pentachloride equilibrate in the presence of molecular chlorine according to the reaction:

 $PCl_3(g) + Cl_2(g) \rightarrow PCl_5(g)$ 

 $K_{eq} = 2.01$  at 500 K. A 1.000-L reaction vessel is charged with 0.300 mol of PCl<sub>5</sub> and allowed to equilibrate at this temperature. The equilibrium partial pressure of PCl<sub>5</sub> is \_\_\_\_\_ atm.

A) 10.1 B) 0.386 C) 2.24 D) 2.48 E) 0.211

2004 Free Response - Form B

1.  $N_2(g) + 3 H_2(g) \rightleftharpoons 2 NH_3(g)$ 

For the reaction represented above, the value of the equilibrium constant,  $K_p$  is  $3.1 \times 10^{-4}$  at 700 K.

- a) Write the expression for the equilibrium constant, K<sub>p</sub>, for the reaction.
- b) Assume that the initial partial pressures of the gases are as follows:

 $P(N_2) = 0.411$  atm,  $P(H_2) = 0.903$  atm, and  $P(NH_3) = 0.224$  atm.

- i) Calculate the value of the reaction quotient, Q, at these initial conditions.
- ii) Predict the direction in which the reaction will proceed at 700. K if the initial partial pressures are those given above. Justify your answer.
- c) Calculate the value of the equilibrium constant,  $K_c$ , given that the value of  $K_p$  for the reaction at 700. K is  $3.1 \times 10^{-4}$ .
- d) The value of  $K_p$  for the reaction represented below is  $8.3 \times 10^{-3}$  at 700. K.

 $NH_3(g) + H_2S(g) \rightleftharpoons NH_4HS(g)$ 

Calculate the value of K<sub>p</sub> at 700. K for each of the reactions represented below.

i) 
$$NH_4HS(g) \rightleftharpoons NH_3(g) + H_2S(g)$$

ii)  $2 H_2S(g) + N_2(g) + 3 H_2(g) \rightleftharpoons 2 NH_4HS(g)$ 

### MULTIPLE CHOICE. Choose the one alternative that best completes the statement or answers the question.

1) A ID: chem9b 15.1-1 2) C ID: chem9b 15.1-7 3) A ID: chem9b 15.1-12 4) A ID: chem9b 15.1-15 5) B ID: chem9b 15.1-18 6) E ID: chem9b 15.1-24 7) A ID: chem9b 15.1-27 8) B ID: chem9b 15.1-31 9) B ID: chem9b 15.1-35 10) E ID: chem9b 15.1-38 11) E ID: chem9b 15.1-39 12) B ID: chem9b 15.1-42 13) E ID: chem9b 15.1-47 14) C ID: chem9b 15.1–50 15) B ID: chem9b 15.1-51 16) C ID: chem9b 15.1-52 17) A ID: chem9b 15.1-54 18) D ID: chem9b 15.2-2 19) C ID: chem9b 15.2-3 20) E ID: chem9b 15.2-4 21) B ID: chem9b 15.2-8

Answer Key Testname: CH\_13\_PRAC\_TEST\_EQUILIBRIUM.TST

22) A ID: chem9b 15.2-9

(PAGE ONE) AP Chem Practice Test Ch. 13 - Equilibrium (3) The second equation is ZX the stoichiometry of the 1stone, so the equilibrium consant for the 2nd equation is the square of the first one. (4) The 2<sup>nd</sup> equation was arrived at by reversing the 1st, and then cutting it in half stoichiometrically, Thus the K for the second equation is  $\sqrt{794} = 0.035$ 5 The 2nd equation is 1/2 of the first equation, so the 2nd equation's K = VIST equation's K leave put (6)  $K_p \# 1 = 0.013$ , so  $K_p \# 2 = 0.013 = 77$ Spiles For that problem  $\begin{cases} 8 \\ K = \frac{[H+][C_2H_3O_2^-]}{[H_2H_3O_2^-]} = \frac{[I,33 \times 10^{-3}][I,33 \times 10^{-3}]}{[0,0990]} \\ \hline \end{cases}$ = 1.79 × 10-5  $K = 2400 = \frac{x^2}{(36.1 - 2x)^2}$ (9)  $2N0 = N_2 + 0_2$ 0 0 +X +X 36.1  $2400 = \left(\frac{x}{36 \cdot 1 - 2x}\right)^2$ -2X ×Χ 36.1-2X V2400 = 36.1-2x  $49.0 = \frac{1}{36.1 - 2 \times 10^{-2}}$ S x=Poz= 17.9 (36.1-2×)(49.0) = × 1768.53 - 97,98×=× 1768.53 = 98,98 X -17.9 = x

PAGE TWO  $M = \frac{N}{V} = \frac{35meP}{1L}$  $= (0_2 + H_z)$  $CO + H_2O$ =0.35MI,35M 0,40M +0.16 +0.16 0, 40mal = 0,40 M C -0.16 -0.16 0.16 0.16 E 0,19M 0,24  $K = \frac{6.16)^2}{(0,19)(0.24)} = 0.56$ 0.19 - 0.35 = -0.16 MZNOBr = ZNO + BIG 12 M= 0.64 mor = 0,64M 0 +0,18 I 0.64 0 +0-09 -0.18 +Note: NOBr: Brz = 2:1 mol 0.09 0.18 0,46

A[NOB] = 0.46-0.64= -0.18

(3) E, DN=0 (same # of molif gab on each side gab on each side f eqn.)

0, 18M= × mol xmel=0,18mplNO and 0.09melBiz

HAdd Oz = causes Q>K = Caused rantoshift left in response, when ran shifts left: d) K dog not change a) Proz increases e) Pco dermases b) K does not change C) Pco2 increases

 $200_2 = 200 + 0_2 + 514 k^3$ PAGE 15 THREE " Attern = -514 KJ" means that heat is produced in this exothermic reaction. You can taink of heat as a product. The reaction shifts to the left, since increasing the Traquires that heat (a product) be added to the system, Also, the equilibrium [costant will decrease? [co] and [U2] decrease and [cos] increases. (16) c, low temp (romones heat, drives rxn to right because this produces head) and low P (Fener moles of gas on rightside). (17) A, nore Freeton K SO2Ch2 = SU2+Ch2 (8) $K = \frac{[so_2][cl_2]}{[so_2cl_2]} = \frac{(0.0287)(0.0287)}{0.0106} = 0.0778$ I amost sure I amnot sure why the answer ter is wrong.

(19) N2U4 = ZNO2 I 0,0300M O C -x +2x E 0.0236M 2X

 $\Delta [N_204] = 0.0236 - 0.0390 = 0.0064M = X$   $2x = 2 \times 0.0064 = 0.0128M$  $K = \frac{[.0128]^2}{.0236} = 6.94 \times 10^{-3}$ 

 $K = \frac{P_{coro_2}}{P_{co}} = 1.49 \times 10^{8}$  $\frac{20}{\text{equilibrium } 8:60 \times 10^4} \quad \begin{array}{c} CO(l_2) \\ \hline CO(l_2) \\$  $P_{core_z} = (1,49\times10^{\circ})(8,60\times10^{-4})^{\circ}$ Plucez = 110 atm  $= 1.10 \times 10^{2} atm$ 21) That Keg is a Kp," As Far as I know, this fact shald have been stated in the problem. PV=NRT p= (0,990 mal) (0,08206 Latm) (500K) 40.6 atm 1.000 L  $K_p = 2.01 = \frac{40.6 - X}{x^2}$ PCR3 + CR2 = PCR5 40.6 atm  $\mathcal{O}$ 0 I 2.01 X2= 40.6-X -X+XC + X $\boldsymbol{\boldsymbol{\kappa}}$ 701+2+x=40.6 X 40.6-X F 2.01×2+×-40.6=0  $x = -b \pm \sqrt{b^2 - 0ac}$ (1=2,0) b=1 20 C=-40.6  $= -1 \pm \sqrt{1^{2} - (4)(2.01)(-40.6)}$ (2)(2.01) =-1 ± 1+ 326.4 2 (2.01)  $\frac{17.09}{4.02} \neq 4.25 = 1000$  $= -1 + \sqrt{3274} = -1 + 18.09 = -18.09 = -18.0$ 

$$\begin{array}{l} (22) \\ P(l_{3} + (l_{2} \implies P(l_{5} = \frac{P(l_{5})}{Y} = \frac{(0.30 \text{ me})(0.00206 \frac{l_{0} + 1}{K \text{ me}})(500 \text{ K})}{1.00 \text{ L}} \\ K_{P} = 2.01 \\ T = 500 \text{ K} = \frac{P(l_{5})}{Y} = \frac{(0.30 \text{ me})(0.00206 \frac{l_{0} + 1}{K \text{ me}})(500 \text{ K})}{1.00 \text{ L}} \\ = 12.3 \text{ atm} \\ P(l_{3} + (l_{2} \implies P(l_{5} = 12.3 \text{ atm})) \\ C = +X + X - X \\ E \times (12.3 \text{ K}) \\ C = +X + X - X \\ E \times (12.3 \text{ K}) \\ C = -12.3 \\ X = -\frac{b \pm \sqrt{b^{2} - 4ac}}{12} \\ C = -12.3 \\ X = -\frac{b \pm \sqrt{b^{2} - 4ac}}{12} \\ = -\frac{1 \pm \sqrt{1 - (4)(2e0)(-123)}}{(2)(2e0))} \\ = -1 \pm \sqrt{1 + 98.892} \\ -\frac{4.02}{4.02} \\ Y = -\frac{1 \pm \sqrt{99.9892}}{4.02} \\ Y = -\frac{1 \pm \sqrt{99.9892}}{4.02} \\ Y = -\frac{1 \pm \sqrt{99.9892}}{4.02} \\ Y = -\frac{1 \pm \sqrt{99.9892}}{12.3 - 2.24} = 10.1 \text{ latm} \end{array}$$

## AP<sup>®</sup> CHEMISTRY 2004 SCORING GUIDELINES (Form B)

### **Question 1**

 $N_2(g) + 3 H_2(g) \rightleftharpoons 2 NH_3(g)$ 

- 1. For the reaction represented above, the value of the equilibrium constant,  $K_p$ , is  $3.1 \times 10^{-4}$  at 700. K.
  - (a) Write the expression for the equilibrium constant,  $K_p$ , for the reaction.

(b) Assume that the initial partial pressures of the gases are as follows:

 $p_{\rm N2} = 0.411$  atm,  $p_{\rm H2} = 0.903$  atm, and  $p_{\rm NH3} = 0.224$  atm.

(i) Calculate the value of the reaction quotient, Q, at these initial conditions.

$Q = \frac{p_{\rm NH_3}^2}{p_{\rm N_2} \times p_{\rm H_2}^3} = \frac{(0.224)^2}{(0.411)(0.903)^3}$	1 point for calculation of $Q$ with correct mass action expression
Q = 0.166	Note: must be consistent with part (a)

(ii) Predict the direction in which the reaction will proceed at 700. K if the initial partial pressures are those given above. Justify your answer.

Since $Q > K_p$ , the numerator must decrease and the	1 point for direction or for stating that $Q > K_p$
denominator must increase, so the reaction must	1 maint fan annlanation
proceed from right to left to establish equilibrium.	I point for explanation

(c) Calculate the value of the equilibrium constant,  $K_c$ , given that the value of  $K_p$  for the reaction at 700. K is  $3.1 \times 10^{-4}$ .

$K_p = K_c(RT)\Delta n$ $\Delta n = 2 - 4 = -2$ $K_p = K_c(RT)^{-2}$	1 point for calculating $\Delta n$
$3.1 \times 10^{-4} = K_c (0.0821 \frac{\text{L atm}}{\text{mol K}} \times 700 \text{ K})^{-2}$	1 point for correct substitution and value of $K$
$3.1 \times 10^{-4} = K_c(3.0 \times 10^{-4})$ 1.0 = K	T point for correct substitution and value of $K_c$

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# AP<sup>®</sup> CHEMISTRY 2004 SCORING GUIDELINES (Form B)

#### Question 1 (cont'd.)

(d) The value of  $K_p$  for the reaction represented below is  $8.3 \times 10^{-3}$  at 700. K.

$$\mathrm{NH}_3(g) + \mathrm{H}_2\mathrm{S}(g) \rightleftharpoons \mathrm{NH}_4\mathrm{HS}(g)$$

Calculate the value of  $K_p$  at 700. K for each of the reactions represented below.

(i)  $NH_4HS(g) \rightleftharpoons NH_3(g) + H_2S(g)$ 

$K_p = \frac{1}{8.3 \times 10^{-3}} = 1.2 \times 10^2$	1 point for the calculation of $K_p$
--	--------------------------------------

(ii)  $2 \operatorname{H}_2 S(g) + \operatorname{N}_2(g) + 3 \operatorname{H}_2(g) \rightleftharpoons 2 \operatorname{NH}_4 \operatorname{HS}(g)$ 

$2 \times [\mathrm{NH}_3(g) + \mathrm{H}_2\mathrm{S}(g) \rightleftharpoons \mathrm{NH}_4\mathrm{HS}(g)]$ $\mathrm{N}_2(g) + 3 \mathrm{H}_2(g) \rightleftharpoons 2 \mathrm{NH}_3(g)$	$\begin{split} K_p &= (8.3 \times 10^{-3})^2 \\ K_p &= 3.1 \times 10^{-4} \end{split}$	1 point for squaring $K_p$ for NH <sub>4</sub> HS or for multiplying $K_p$ 's
$2 \text{ H}_2\text{S}(g) + \text{N}_2(g) + 3 \text{ H}_2(g) \rightleftharpoons 2 \text{ NH}_4\text{HS}(g)$ $K_p = (8.3 \times 10^{-3})^2 (3.1 \times 10^{-4}) = 2.1 \times 10^{-8}$		1 point for correct $K_p$