## Jordan University of Science & Technology Department of Applied Chemistry CH102 Exam II 11/5/2004

Student's Name: Instructor's Name: Student's No.: Cuch The O Section: Serial No.: Question Answer X Question Answer

## Circle the best correct answer for all the following questions?

1. If the half-life, t <sub>1/2</sub> ,	for a first	order reaction	is 495	seconds,	then the rate	constant
for this reaction is:	•					

A) 3.20x10<sup>-4</sup> s<sup>-1</sup> B) 1.40x10<sup>-3</sup> s<sup>-1</sup> C) 2.4x10<sup>-3</sup> s<sup>-1</sup> D) 2.0x10<sup>-2</sup> s<sup>-1</sup>

 Consider the reaction A + B → Products. If the rate expression is given by rate = k[A][B], which of the following is not correct for this reaction?

A) The reaction is first order in A

B) The reaction is first order in B

The reaction is overall second order

Doubling the concentration of both A and B should double the rate.

3-5. Consider the following rate data for the reaction below at certain temperature:

 $2A + 3B \rightarrow D$ 

Experiment	Initial [A], M	Initial [B], M	Initial rate, M.s-1
1.	0.10	0.30	.7.22x10 <sup>-5</sup>
2.	0.10	0.60	1.44x10 <sup>-4</sup>
3.	0.20	0.60	5.67x10 <sup>-4</sup>

3. The reaction is ---- order in A and ---- order in B, respectively:

A) first, first B) second, second (C) second, first D) first, second

4. What is the rate constant for the reaction?

A) 0.024 M<sup>-1</sup>s<sup>-1</sup>

(B) 0.024 M<sup>-2</sup>s<sup>-1</sup>

C) 0.004 M<sup>-1</sup>s<sup>-1</sup>

D) 0.004 M<sup>-2</sup>s<sup>-1</sup>

5. If the rate of disappearance of A is 0.02 Ms<sup>-1</sup>. What is the rate of disappearance of

A) 0.06 Ms<sup>-1</sup>

(B) 0.03 Ms<sup>-1</sup> C) 0.02 Ms<sup>-1</sup>

D) 0.04 Ms<sup>-1</sup>

6. The value of the rate constant for a reaction can be increased if:

The temperature of the reaction is increased.

B) The concentration of the reactants is increased

C) The activation energy of the reaction is increased.

D) The concentration of the products is increased.

√7. The reaction 2A → Products is second order in A. If the initial concentration of A is 0.80 M, and it took 500 seconds for the concentration of A to drop to 0.45 M. The concentration of A after 1800 seconds will be:

A) 0.41 M

B) 0.21 M

C) 0.15 M

20.32 M

8. According to the collision theory:

A) All collisions will give products

B) only active collisions may give products

C) only collisions with proper orientation of the reacting atoms and/or molecules will give products

D) both B and C are important factors to give products.



9. The reaction A  $\rightarrow$  products is a first order with k = 0.12 s<sup>-1</sup>. How many seconds it will take for the initial concentration to drop to 40 % of its value?

A) 14.3 s

B) 8.52 s

C) 4.26 s

D) 7.64 s

10. Which of	the followin	ig is correct ab	out the activation ener	rgy for a certain
reaction?			The same of the sa	
the first of the second			e of the reaction	
		s of temperatu		
			n of the reactants.	
D) It depends	on the initia	al concentratio	n of the products.	
2A →	2C + I (sl	ow)	2C + D, if the mechan	nism of this reaction is:
	D (fa:	St)		
11.This reacti		t makes in D	D) second order in	A and zero order in B
A) First order		rst order in B		A and zero order in B
C) Second ord	ier in B		second order in	A and first order in I.
10 771	/ 1: - t t	iion io		
		is reaction is	D) D	
A) C	B) B	X)1	D) D	
_13. Which of	the followin	g is correct ab	out a catalyst for a rea	ction at equilibrium?
A) Increases t	he amount o	of products pre	sent at equilibrium.	
B) Increases the	he rate at wh	hich the equilil	brium is reached but d	ecreases the equilibrium
constant.				
(A) Increases the	he rate at wl	nich the equilil	brium is reached withou	out changing the
equilibrium co	onstant.			
D) The seasting	J. Lee	and the same of th		
D) The reaction	on needs less	s amounts of re	eactants to reach equil	ibrium.
14. Which of	the followin	g is considered	eactants to reach equil	
14. Which of Bronsted-Low	the followin	g is considered	d a conjugate acid-base	e pair according to
14. Which of	the followin	g is considered		e pair according to
14. Which of Bronsted-Low A) H <sub>2</sub> S/HS <sup>-</sup>	the following very concept B) H <sub>3</sub> PC the equilibri	ig is considered? $O_4/PO_4^{3-}$ C) tum $N_2O_{4(g)} \leftrightarrow$	d a conjugate acid-base	e pair according to
14. Which of Bronsted-Low A) H <sub>2</sub> S/HS <sup>-</sup> 15-17. Given concentration	the following very concept B) H <sub>3</sub> PC the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> 8	ig is considered $^{?}$ $^{0_4/PO_4}^{3-}$ C) $^{0_4/PO_4}^{3-}$ C) $^{0_4/PO_4}^{0_4}$ in equilibrium	d a conjugate acid-base $H_2CO_3/CO_3^2$ No $N_2(g)$ , $K = 0.21$ equal 0.0045 M:	e pair according to
14. Which of a Bronsted-Low A) H <sub>2</sub> S/HS 15-17. Given concentration 15. Calculate	the following very concept B) H <sub>3</sub> PC the equilibrit of N <sub>2</sub> O <sub>4 (g)</sub> at the concentr	ig is considered $^{2}$ $^{2}$ $^{2}$ $^{3}$ $^{4}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{2}$ $^{4}$ $^{4}$ $^{5}$ $^{6}$ $^{7}$	d a conjugate acid-base $H_2CO_3/CO_3^2$ No $N_2(g)$ , $K = 0.21$ equal 0.0045 M:	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the
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14. Which of Bronsted-Low A) H <sub>2</sub> S/HS 15-17. Given concentration 15. Calculate A) 0.0166 M	the following very concept B) H <sub>3</sub> PC the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> at the concentr	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  tum N <sub>2</sub> O <sub>4 (g)</sub> ↔ at equilibrium  ration of NO <sub>2</sub> at 0.0310 M	d a conjugate acid-base $H_2CO_3/CO_3^2$ No $N_2(g)$ , $K = 0.21$ equal 0.0045 M: at equilibrium?	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the
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14. Which of Bronsted-Low A) H <sub>2</sub> S/HS 15-17. Given concentration 15. Calculate A) 0.0166 M	the following very concept B) H <sub>3</sub> PC the equilibris of N <sub>2</sub> O <sub>4 (g)</sub> at the concentrate the initial contentrate the initial con	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  tum N <sub>2</sub> O <sub>4 (g)</sub> ↔ at equilibrium  ration of NO <sub>2</sub> at 0.0310 M	d a conjugate acid-base $H_2CO_3/CO_3^2$ No $N_2(g)$ , $K = 0.21$ equal 0.0045 M: at equilibrium?	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M
14. Which of Bronsted-Low A) H <sub>2</sub> S/HS 15-17. Given concentration 15. Calculate A) 0.0166 M 16. Calculate is zero?	the following very concept B) H <sub>3</sub> PC the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> at the concentrathe initial control B) the equilibrium and the equilibrium the equilibrium at the equilibrium the equilib	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  tum N <sub>2</sub> O <sub>4 (g)</sub> ↔ at equilibrium  ration of NO <sub>2</sub> at 0.0310 M  oncentration of 0.0155 M	d a conjugate acid-base  H <sub>2</sub> CO <sub>3</sub> /CO <sub>3</sub> <sup>2</sup> N  2NO <sub>2 (g)</sub> , K = 0.21  equal 0.0045 M:  at equilibrium?  C) 0.0022 M  N <sub>2</sub> O <sub>4</sub> ; assuming initial	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M  al concentration of NO <sub>2</sub> D) 0.0096 M
14. Which of Bronsted-Low A) H <sub>2</sub> S/HS 15-17. Given concentration 15. Calculate A) 0.0166 M 16. Calculate is zero? 0.0200 M 17. Calculate 12N <sub>2</sub> O <sub>4(g)</sub> at 1	the following very concept B) H <sub>3</sub> PC the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> at the concentrate the initial concentrate B) the equilibrium of C?	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  cum N <sub>2</sub> O <sub>4 (g)</sub> ↔  at equilibrium  ration of NO <sub>2</sub> at 0.0310 M  oncentration of 0.0155 M  cum constant, I	d a conjugate acid-base $H_2CO_3/CO_3^2$ No $N_2O_2$ (g) , $K = 0.21$ equal 0.0045 M:  at equilibrium?  C) 0.0022 M $N_2O_4$ ; assuming initial C) 0.0400 M  K, for the following real	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M  al concentration of NO <sub>2</sub> D) 0.0096 M  action: 4NO <sub>2</sub> (g) ↔
14. Which of Bronsted-Low A) H <sub>2</sub> S/HS  15-17. Given concentration  15. Calculate A) 0.0166 M  16. Calculate is zero?  17. Calculate is 2N <sub>2</sub> O <sub>4(g)</sub> at 1	the following very concept B) H <sub>3</sub> PC the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> at the concentrate the initial concentrate B) the equilibrium of C?  B) 2.2	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  (um N <sub>2</sub> O <sub>4 (g)</sub> $\leftrightarrow$ at equilibrium  ration of NO <sub>2</sub> at 0.0310 M  oncentration of 0.0155 M  um constant, I	d a conjugate acid-base  H <sub>2</sub> CO <sub>3</sub> /CO <sub>3</sub> <sup>2</sup> N  2NO <sub>2 (g)</sub> , K = 0.21  equal 0.0045 M:  at equilibrium?  C) 0.0022 M  N <sub>2</sub> O <sub>4</sub> ; assuming initia  C) 0.0400 M  K, for the following rea  C) 9.4  D)	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M  al concentration of NO <sub>2</sub> D) 0.0096 M  action: 4NO <sub>2</sub> (g) ↔
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14. Which of Bronsted-Low A) H <sub>2</sub> S/HS  15-17. Given concentration  15. Calculate A) 0.0166 M  16. Calculate is zero?  0.0200 M  17. Calculate is 2N <sub>2</sub> O <sub>4(g)</sub> at 1  2N <sub>2</sub> O <sub>4(g)</sub> at 1  22  18. For the real 0.50 M, when is	the following the equilibrium of N <sub>2</sub> O <sub>4 (g)</sub> at the concentrate the initial concentrate the initial concentrate the initial concentrate the equilibrium of C?  B) 2.2 action A > 1 time of the rate o	g is considered?  0 <sub>4</sub> /PO <sub>4</sub> <sup>3</sup> C)  cum N <sub>2</sub> O <sub>4 (g)</sub> ↔  at equilibrium  ration of NO <sub>2</sub> at 0.0310 M  oncentration of 0.0155 M  rum constant, If the constant is the constant in the constant in the constant, If the constant is the constant in the consta	d a conjugate acid-base $H_2CO_3/CO_3^2$ No. $O_2(g)$ , $K = 0.21$ equal 0.0045 M: at equilibrium?  C) 0.0022 M  N <sub>2</sub> O <sub>4</sub> ; assuming initial C) 0.0400 M  C, for the following reaction of A droped from 1.00 minute to the concentration of A droped fr	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M  al concentration of NO <sub>2</sub> D) 0.0096 M  action: 4NO <sub>2</sub> (g) ↔  88  ped from 0.80 M to
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14. Which of Bronsted-Low A) H <sub>2</sub> S/HS  15-17. Given concentration  15. Calculate A) 0.0166 M  16. Calculate is zero?  17. Calculate is zero?  17. Calculate is 2N <sub>2</sub> O <sub>4 (g )</sub> at 1 A) 22  18. For the real 0.50 M, when is average rate of A) 0.005 M/s  19. A Bronester	the following the equilibrium of N <sub>2</sub> O <sub>4</sub> (g) at the concentrate the initial contraction A in the equilibrium of the equilibriu	g is considered?  04/PO4 <sup>3</sup> C)  1 um N <sub>2</sub> O <sub>4 (g)</sub> contained at equilibrium at equ	d a conjugate acid-base $H_2CO_3/CO_3^2$ No. $O_2(g)$ , $K = 0.21$ equal 0.0045 M: at equilibrium? C) 0.0022 M $O_2CO_4$ assuming initial C) 0.0400 M $O_3CO_4$ assuming initial C) 0.0400 M	e pair according to  H <sub>4</sub> /NH <sub>2</sub> 3 at 100 ° C and the  D) 0.0775 M  al concentration of NO <sub>2</sub> D) 0.0096 M  action: 4NO <sub>2</sub> (g) ↔  88  ped from 0.80 M to 0.1.50 minutes. The
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