Which of the following reaction mechanisms is consistent with the rate law established in (a)? Explain your choice.

I.
$$NO + NO \rightleftharpoons N_2O_2$$
 (fast)

$$N_2O_2 + Br_2 \rightarrow 2 \text{ NOBr}$$
 (slow)

II.
$$Br_2 \rightarrow Br + Br$$
 (slow)

$$2(NO + Br \rightarrow NOBr)$$
 (fast)



$$N_2O_5(g) \to 4NO_2(g) + O_2(g)$$

Dinitrogen pentoxide gas decomposes according to the equation above. The first-order reaction was allowed to proceed at 40°C and the data below were collected.

$[N_2O_5]$ (M)	Time (min)	
0.400		
0.289	20.0	
0.209	40.0	
0.151	60.0	
0.109	80.0	

a= In[a]+ In[A] == KT = In[a89] In [0.40] = -k20 K= 0.0163 min = 1 t = 8.19 man

- Calculate the rate constant for the reaction using the values for concentration and time given C.) INE AJ. INE. 4] -,063 in the table. Include units with your answer. $N_2O_5 = e^{-2.55}$
- After how many minutes will [N₂O₅] be equal to 0.350 M? √(b)
 - What will be the concentration of N₂O₂ after 100 minutes have elapsed? (c)
 - Calculate the initial rate of the reaction. Include units with your answer. (d)

What is the half-life of the reaction?

Figaph or use formula =
$$\frac{603}{k} = \frac{603}{6003} = 42.5$$

= 0.0163(-4)

M = 0.078M

The following results were obtained in experiments designed to study the rate of the reaction above:

Experi- ment		oncentra- noles/L) [B]	Initial Rate of Formation of D (M/min)
1	0.10	0.10	1.5×10^{-3}
2	0.20	0.20	3.0×10^{-3}
3	0.20	0.40	6.0×10^{-3}

1-2 double a double B Rxn double: no help

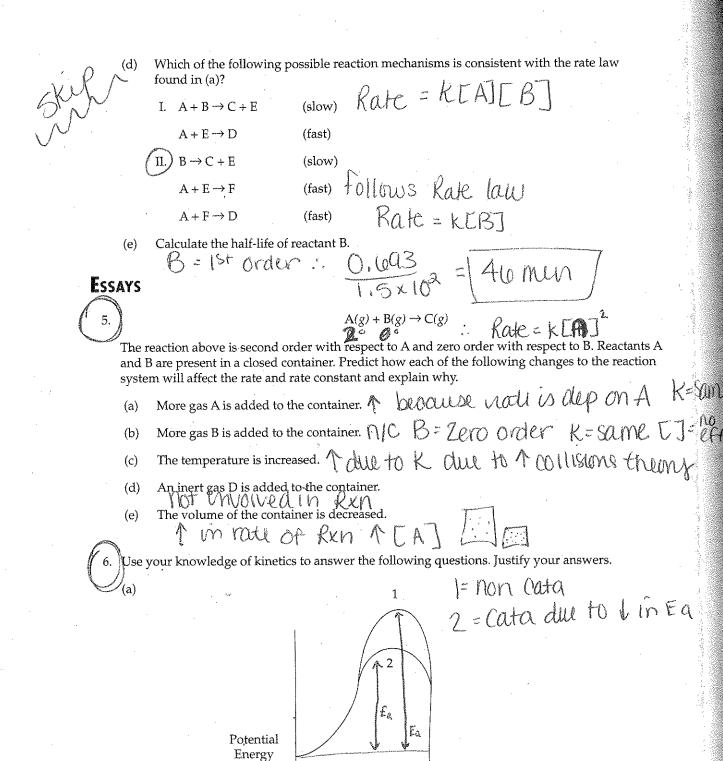
2-3 Achs Bdauble Rak dowbu.

Rak dowbu.

B = 2°

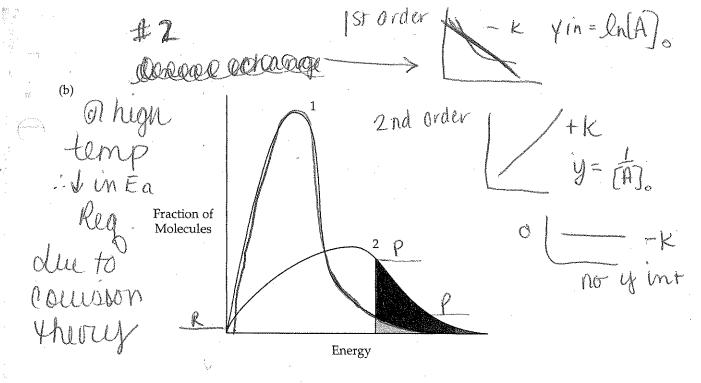
chie to no charge

- Write the rate law for the reaction. KAR = KER (a)
- Calculate the value of the rate constant, k, for the reaction. Include the units. 3rd order over all: NO eq (b)
- If experiment 2 goes to completion, what will be the final concentration of D? Assume that ... SOLVE (c) the volume is unchanged over the course of the reaction and that no D was present at the start of the experiment.



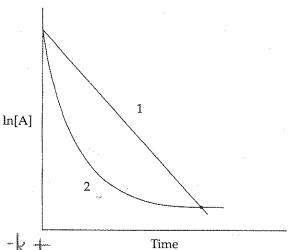
Reaction Coordinate

The two lines in the diagram above show different reaction pathways for the same reaction. Which of the two lines shows the reaction when a catalyst has been added?



Which of the two lines in the energy distribution diagram shows the conditions at a higher temperature?

(c)



n(A). In [A] = - k+

Which of the two lines in the diagram above shows the relationship of ln[A] to time for a first order

reaction with the following rate law:

reaction with the reaction with the reaction with the reaction with the reaction Rate = k[A] $n[A]_{+} = -k + ln[A]_{o} = Rate = k[A]$ $n[A]_{+} = -k + ln[A]_{o} = ln[A]$ Rate = k[A] $ln[A]_{+} = -k + ln[A]_{o} = ln[A]$ $ln[A]_{+} = -k + ln[A]_{o} = ln[A]$

5.

 $A(g) + B(g) \rightarrow C(g)$ $A(g) + B(g) \rightarrow C(g)$

The reaction above is second order with respect to A and zero order with respect to B. Reactants A and B are present in a closed container. Predict how each of the following changes to the reaction system will affect the rate and rate constant and explain why.

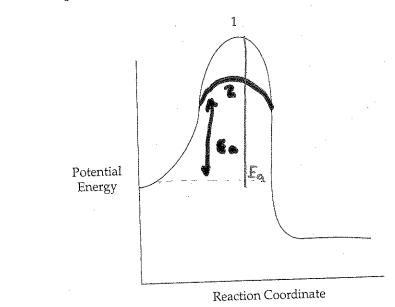
- (a) More gas A is added to the container.
- (b) More gas B is added to the container.
- (c) The temperature is increased.
- (d) An inert gas D is added to the container.
- (e) The volume of the container is decreased.
- a) A CAJ Arate because rate depends on CAJ due to rate lawy (CAJE INTE
- 6) 1 CB] no rate change since 5 ms
- C) I temp I rate + K

 It I KE ! more gas molecules conide w/eraugh
 evergy to overcome Eq.
- D) Dio not involved in the RXN or the rate law so rate + K are not effected
- E.) Volume 1 concentration 1 [A] trate
 rate constant K is independent of [A] so it
 wont change.

6. Use your knowledge of kinetics to answer the following questions. Justify your answers.

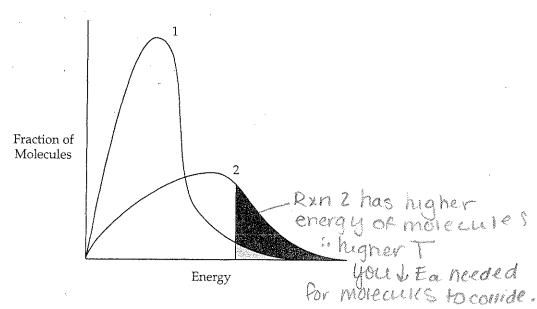
(a)

(b)



The two lines in the diagram above show different reaction pathways for the same reaction. Which of the two lines shows the reaction when a catalyst has been added?

-RXN 2 since it has a lower activation energy.



Which of the two lines in the energy distribution diagram shows the conditions at a higher temperature?

Which of the two lines in the diagram above shows the relationship of ln[A] to time for a first order reaction with the following rate law:

Rate = k[A] $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ - |M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$ $|M \subseteq AJ \cap M \subseteq AJ \circ : -Kt|$

Concentration (moles/L)

Concentration (moles/L)

Time

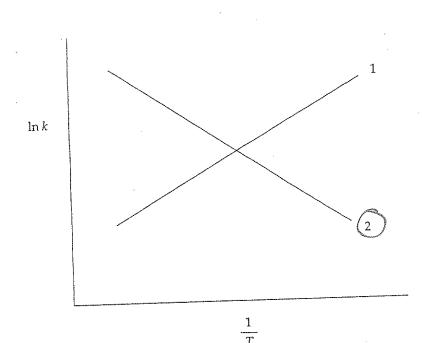
Concentration (moles/L)

A

Time

Which of the two graphs above shows the changes in concentration over time for the following reaction?

(e)

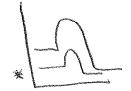


- Which of the two lines in the diagram above shows the relationship of lnk to $\frac{1}{T}$ for a reaction?
- b . How is the slope of the line related to the activation energy for the reaction?

a. Krastt : \- K or slope since x = +

Use your knowledge of kinetics to explain each of the following statements:

- (a) An increase in the temperature at which a reaction takes place causes an increase in reaction
- (b) The addition of a catalyst increases the rate at which a reaction will take place.
- (c) A catalyst that has been ground into powder will be more effective than a solid block of the same catalyst.
- (d) Increasing the concentration of reactants increases the rate of a reaction.
- A) AT = 1 Energy = 1 collisions = 1 reaction rate
- B.) alternative path at a lower Ea.: collisions that make contact at a k energy will be able to go to completion. Where prior to catalyst that energy would not have been sufficient.



- C.) Surface area has an effect on Run rate.

 A Surface area Aran rate. : pwd = 1 Surface area.
- D.) N'E reactants] = 1 probability of collisions