## AP Chem Rate of Reaction Review!

1. For the Reaction: $\quad 2 \mathrm{~A}_{(\mathrm{g})}+\mathrm{BC}_{2(\mathrm{~g})} \quad------>\mathrm{A}_{2} \mathrm{~B}_{(\mathrm{g})}+2 \mathrm{C}_{(\mathrm{g})}$

| Trial \# | Initial concentration of $\mathrm{A}(\mathrm{M}):$ | Initial Concentration of $\mathrm{BC}_{2}(\mathrm{M})$ | Initial rate of formation of $\mathrm{A}_{2} \mathrm{~B}(\mathrm{M} / \mathrm{s})$ |
| :--- | :---: | :---: | :---: |
| 1 | 0.050 | 0.050 | 0.00016 |
| 2 | 0.050 | 0.100 | 0.00064 |
| 3 | 0.10 | 0.200 | 0.00512 |

a. Find the rate law and the value of k , including units.
b. In trial 1, what was the initial rate of formation of C ?
c. Predict the rate of disappearance of $\mathrm{A}_{(\mathrm{g})}$ if both reactant starting concentrations are 0.030 M .
d. Calculate the intial rate of reaction (in terms of $\mathrm{A}_{2} \mathrm{~B}$ ) if the initial concentrations of A and $\mathrm{BC}_{2}$ are 0.15 M and 0.25 M , respectively.


Suppose the above reaction is done by pouring 100.0 mL of 0.80 Molar nitric acid at $20.0^{\circ} \mathrm{C}$ onto 50 . grams of copper in the form $1.0 \mathrm{~cm}^{3}$ cubes. How would the rate of reaction change if you changed the following things? (Increase? Decrease? Or no change?)
$\qquad$ a. Change the concentration of nitric acid to 0.60 M .
b. Use $50 . \mathrm{g}$ of copper powder, instead of the cubes.
c. Add 50.0 mL of water to the 100.0 mL of 0.80 M nitric acid, and then add this solution to the $1.0 \mathrm{~cm}^{3}$ copper cubes, instead of the original solution. (still at $20 .{ }^{\circ} \mathrm{C}$ ) d. Change the temperature of the acid to $30.0^{\circ} \mathrm{C}$

| 3.$2 \mathrm{C}_{5} \mathrm{H}_{6}$ <br> (cyclopentadiene) | $-\cdots-->$ <br> (dicyclopentadiene) | $\mathrm{C}_{10} \mathrm{H}_{12}$ |
| :--- | :---: | :---: |
|  | Initial Concentration of $\mathrm{C}_{5} \mathrm{H}_{6}(\mathrm{M} / \mathrm{s})$ | Initital rate of formation of $\mathrm{C}_{10} \mathrm{H}_{12}(\mathrm{M} / \mathrm{s})$ |
| Experiment 1 | 0.040 | 0.000267 |
| Experiment 2 | 0.060 | 0.000601 |
| Experiment 3 | 0.120 | 0.00240 |

a. Determine the rate law and the value of the rate constant.
b. How will the reaction rate be affected if the temp increases?

Explain your answer mathematically
Explain your answer in terms of what the molecules are doing
c. How will the reaction rate be affected if a catalyst is added?

Explain your answer mathematically
Explain in terms of what the molecules are doing
d. How is the reaction rate affected when the concentration of cyclopentadiene is increased?

Explain in terms of what the molecules are doing
e. If you start with a $\mathrm{C}_{5} \mathrm{H}_{6}$ molarity of 0.120 Molar, what concentration of $\mathrm{C}_{5} \mathrm{H}_{6}$ will remain after 2.00 minutes?
4. Consider the following 2-step reaction mechanism involving ozone $\left(\mathrm{O}_{3}\right)$ depletion:
Elementary Step \#1
$\mathrm{O}_{3(\mathrm{~g})}+\mathrm{NO}_{(\mathrm{g})}--->\mathrm{NO}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}$
(slow)
Elementary Step \#2

$$
\mathrm{O}_{(\mathrm{g})}+\mathrm{NO}_{2(\mathrm{~g})}^{(\mathrm{g})}-\cdots-->\mathrm{NO}_{(\mathrm{g})}+\mathrm{O}_{2(\mathrm{~g})} \quad \text { (fast) }
$$

a. Determine the overall reaction.
b. Identify any catalyst(s) and intermediate(s)
c. Was the catalyst in this reaction a homogeneous or heterogeneous catalyst?
d. Determine the rate law for the reaction.
e. Determine the "molecularity" of each elementary step (classify each elementary step as bimolecular, unimolecular, or termolecular)
f. What is the difference between an elementary step and an overall reaction, in terms of collisions?
g. Suggest a heterogeneous catalyst for the reaction. $\qquad$
5. Reaction: $2 \mathrm{NO}+\mathrm{O}_{2}---->2 \mathrm{NO}_{2}$

A mechanism for the above reaction is shown below.
a. Show that the mechanism is consistent with the overall stoichiometry of the reaction.
b. Identify any catalysts or intermediates.
c. Which elementary step would you expect to have a higher activation energy?
d. Determine the rate law for the reaction.
e. What is the overall order of the reaction?
f. Based on the rate law, one might propose that the reaction occurred in a single elementary step. Why is this unlikely?
$\begin{array}{lllll}\text { Step } 1 \text { (fast eqm) } & \mathrm{NO}+\mathrm{NO} & \text { <-------> } & \mathrm{N}_{2} \mathrm{O}_{2} \\ \text { Step } 2 \text { (slow) } & \mathrm{N}_{2} \mathrm{O}_{2}+ & \mathrm{O}_{2} & ------> & 2 \mathrm{NO}_{2}\end{array}$
6. Y -----> Z

The concentration of Y vs time was determined at $25^{\circ} \mathrm{C}$.
At $25^{\circ} \mathrm{C}$, A plot of $\ln [\mathrm{Y}]$ vs time was not linear.
A plot of $1 /[\mathrm{Y}]$ vs time was linear, with a slope of $0.080 \mathrm{~L} / \mathrm{mol}-\mathrm{s}$.
a. Determine the rate law and the value of k . (Assume $25^{\circ} \mathrm{C}$ unless otherwise noted)
b. Will this reaction have a constant half-life? If so, calculate the half-life.
c. Make a graph showing the concentration of Y vs time.

Explain how the rate of reaction changes over time, why this happens, and how this relates to your graph.
d. If you start with a $[\mathrm{Y}]$ of 0.200 M , what will be the initial rate of reaction?
e. If you start with a $[\mathrm{Y}]$ of 0.200 M , what will be the concentration of Y after 40 . seconds?
f. If you start with a [Y] of 0.200 M , what will be the rate of reaction after 40 . seconds?
g. If the reaction has an activation energy of $42 \mathrm{~kJ} / \mathrm{mol}$, what is the value of the Arrhenius parameter A ?
h. Find the rate constant at $55^{\circ} \mathrm{C}$.
7. W ----> 2 Z at $25^{\circ} \mathrm{C}, \mathrm{k}=0.000059 \mathrm{~s}^{-1}$

Note: The rate constant for this reaction was calculated in terms of the rate of disappearance of $\mathbf{W}$.
a. Based on the units of k , what is the overall order of the reaction?
b. Which of these plots would be linear?
$\ln [\mathrm{W}]$ vs $(1 /$ time $) \quad \ln [\mathrm{W}]$ vs time $\quad[\mathrm{W}]$ vs time $\quad 1 /[\mathrm{W}]$ vs time
c. If you start with a [W] of 0.200 Molar, what will be the molarity after 4.00 hours, if the reaction occurs at $25^{\circ} \mathrm{C}$ ?
d. Does this reaction have a constant half life at $25^{\circ} \mathrm{C}$ ? If so, what is the value?
e. What is the initial rate of formation of Z , if W has a starting concentration of 0.200 M ? $\left(25^{\circ} \mathrm{C}\right)$
8. $\mathrm{N}_{2} \mathrm{O}_{4}+$ heat $<-\cdots 2 \mathrm{NO}_{2}$

Suppose that $\mathrm{N}_{2} \mathrm{O}_{4}$ and $\mathrm{NO}_{2}$ are at equilibrium in a sealed container with a fixed volume.
a. How will the following things be affected if the temperature is increased?

Keq $\qquad$
Forward rate of rxn $\qquad$
Reverse Rate of Rxn $\qquad$
Activation energy $\qquad$
b. How will the following things be affected if a catalyst is added?

Keq $\qquad$
Forward rate of rxn $\qquad$
Reverse Rate of Rxn $\qquad$
Activation energy $\qquad$
9. $2 \mathrm{NO}_{2}<---->\mathrm{N}_{2} \mathrm{O}_{4}+$ heat

Suppose that $\mathrm{N}_{2} \mathrm{O}_{4}$ and $\mathrm{NO}_{2}$ are at equilibrium in a sealed container with a fixed volume. a. How will the following things be affected if the temperature is increased? Keq $\qquad$
Forward rate of rxn $\qquad$
Reverse Rate of Rxn $\qquad$
Activation energy $\qquad$
b. How will the following things be affected if a catalyst is added? Keq $\qquad$
Forward rate of rxn $\qquad$
Reverse Rate of Rxn $\qquad$
Activation energy $\qquad$
10. Reaction for the decomposition of $\mathrm{N}_{2} \mathrm{O}_{5}$ :

$$
2 \mathrm{~N}_{2} \mathrm{O}_{5(\mathrm{~g})}--\cdots--->4 \mathrm{NO}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}
$$

The energy diagram for this reaction is shown to the right.
a. Is this reaction exothermic or endothermic? $\qquad$
b. What is the value of $\Delta H r x n ?$ $\qquad$
c. Write the heat term into the equation on the correct side.
d. What is the value for Ea (Activation energy?)
e. Sketch the energy diagram (on the same graph) if a catalyst is added.
(for 10 and 11 , use a scale of 1 square $=10 \mathrm{~kJ}$ )
11. Rxn: $\mathrm{CO}_{2(\mathrm{~g})}+\mathrm{NO}_{(\mathrm{g})} \cdots--->\mathrm{CO}_{(\mathrm{g})}+\mathrm{NO}_{2(\mathrm{~g})}$

This reaction has an $\mathrm{Ea}=360 \mathrm{~kJ}$ and a $\Delta \mathrm{Hrxn}$ of 230 kJ .
a. Is this reaction exothermic or endothermic?
b. Write the heat term into the equation on the correct side.
c. Sketch the energy diagram for this reaction.
d. What is the $\Delta \mathrm{H}$ of the reverse reaction?
e. What is the Ea of the reverse reaction? $\qquad$

For \#12, 13, and 14, determine the rate law, the value and units of $k$, and the overall order of the reaction.
12. $3 \mathrm{~A}+\mathrm{B}-\cdots \mathrm{C}+2 \mathrm{D}$

| Initial [A](M) | Initial [B] (M) | Initial rate of formation of D (M/s) |
| :--- | :---: | :---: |
| 0.300 | 0.100 | 0.001157 |
| 0.200 | 0.100 | 0.000771 |
| 0.450 | 0.200 | 0.00694 |

13. $\mathrm{C} 3 \mathrm{PO}+\mathrm{R} 2 \mathrm{D} 2$-----> $\mathrm{BB} 8+\mathrm{K} 2 \mathrm{SO}$

| Initial [C3PO](M) | Initial [R2D2] (M) | Initial rate of formation of BB8(M/min) |
| :--- | :---: | :---: |
| 0.100 | 0.100 | 0.013 |
| 0.900 | 0.200 | 0.078 |
| 0.400 | 0.100 | 0.026 |

14. $2 \mathrm{X}+\mathrm{Y}--->$ Bees?

Initial [X](M) Initial [Y] (M) Initial rate of formation of Bees? (M/s)

| 0.150 | 0.100 | 0.114 |
| :--- | :--- | :--- |
| 0.300 | 0.100 | 0.456 |
| 0.450 | 0.200 | 1.026 |

Answers!!!!
1a. $\mathrm{r}=\mathrm{k}[\mathrm{A}]^{1}\left[\mathrm{BC}_{2}\right]^{2}, \mathrm{k}=1.2(8) \mathrm{M}^{-2} \mathrm{~s}^{-1}$ or $\mathrm{L}^{2} / \mathrm{mol}^{2}-\mathrm{s} \quad$ b. $0.00032 \mathrm{M} / \mathrm{s} \quad$ c. $0.000069 \mathrm{M} / \mathrm{s} \quad \mathrm{d} .0 .012 \mathrm{M} / \mathrm{s}$.
2. Decrease, increase, decrease, increase
3. a. $\mathrm{r}=\mathrm{k}\left[\mathrm{C}_{5} \mathrm{H}_{10}\right]^{2}, \mathrm{k}=0.17 \mathrm{M}^{-1} \mathrm{~s}^{-1} \quad$ 3e. 0.035 M
b. As Temp increases, rate increases (see key for explanation)
c. Adding a catalyst will cause the rate to increase (see key for explan.)
d. Increasing concentration of the reactant will cause the rate to increase (see key for explan.)

4a. $\mathrm{O} 3+\mathrm{O}--->2 \mathrm{O} 2$ b. NO is a catalyst, NO 2 is an intermediate. c. Homogeneous
d. $\mathrm{r}=\mathrm{k}[\mathrm{O} 3]^{1}[\mathrm{NO}]^{1} \quad$ e. Step 1 and Step 2 are each bimolecular $\mathbf{f} .$. . see key
g. Pt!

5a. The mechanism adds up to the original reaction, so, yes, it is consistent.
b. N 2 O 2 is an intermediate. c. Step 2 , since it has the slower rate.
d. $\mathrm{r}=\mathrm{k}[\mathrm{NO} 2]^{2}[\mathrm{O} 2]$ (see key for explanation/work)
e. $3^{\text {rd }}$ order overall f. See key...termolecular collisions are rare!
6. a. $\mathrm{r}=\mathrm{k}[\mathrm{Y}]^{2}, \mathrm{k}=0.080 \mathrm{~L} / \mathrm{mol}-\mathrm{s}$
b. No! Shut up! Only $1^{\text {st }}$ order reactions have constant half life. This rxn is $2^{\text {nd }}$ order.
c. See key d. initial rate $=0.0032 \mathrm{M} / \mathrm{s} \quad$ e. $[\mathrm{Y}]=0.12$ Molar $\quad$ f. $0.0012 \mathrm{M} / \mathrm{s}$
g. $\mathrm{A}=1.8 \times 10^{6} \mathrm{M}^{-1} \mathrm{~s}^{-1} \quad$ h. $\mathrm{k}=0.38 \mathrm{M}^{-1} \mathrm{~s}^{-1}$ at $55^{\circ} \mathrm{C}$
7. a. First order (based on units of k)
b. Since it is first order, $\ln [W]$ vs time would be linear (with a slope of -k )
c. $[\mathrm{W}]=0.086 \mathrm{M}$
d. Yes! $\mathrm{t}_{1 / 2}=12000$ seconds, or 3.3 hours
e. $0.000024 \mathrm{M} / \mathrm{s}$
8. a. increase, increase, increase (the fwd rate increases by a larger factor than reverse rate), no change.
b. no change, increase, increase (fwd and reverse rates increase by same factor), decrease.
9. decrease, increase, increase (the reverse rate increases by a larger factor), no change.
b. no change, increase, increase (fwd and reverse rates increase by same factor), decrease.
10. a. exo b. $-23 \mathrm{~kJ} /$ mole (anywhere from -20 to -25 kJ is acceptable)
c. $100 \mathrm{~kJ}(95-100$ is ok) d. write " +23 kJ " on the right side e. see key!
11. a. endo b. write " +230 kJ " on left side.
c. See key. d. $\Delta$ Hreverse $\mathrm{rxn}=-230 \mathrm{~kJ} /$ mole
e. Ea reverse $r x n=130 \mathrm{~kJ} /$ mole .
12. $\mathrm{r}=\mathrm{k}[\mathrm{A}]^{1}[\mathrm{~B}]^{2}, \mathrm{k}=0.386 \mathrm{M}^{-2} \mathrm{~s}^{-1}, 3 \mathrm{rd}$ order overall.
13. $\mathrm{r}=\mathrm{k}[\mathrm{C} 3 \mathrm{PO}]^{0.5}[\mathrm{R} 2 \mathrm{D} 2]^{1}, \mathrm{k}=0.41 \mathrm{M}^{-0.5} \mathrm{~min}^{-1}$, the reaction is $11 / 2$ order overall.
14. $\mathrm{r}=\mathrm{k}[\mathrm{X}]^{2}[\mathrm{Y}]^{0} \quad$ or $\mathrm{r}=\mathrm{k}[\mathrm{X}]^{2} . \quad \mathrm{k}=5.07 \mathrm{M}^{-1} \mathrm{~s}^{-1}, 2^{\text {nd }}$ order overall.

