$\qquad$
$\qquad$

- This is practice - Do NOT cheat yourself of finding out what you are capable of doing. Be sure you follow the testing conditions outlined below.
- DO NOT USE A CALCULATOR. You may use ONLY the green periodic table.
- Try to work at a pace of 1.3 min per question. Time yourself. It is important that you practice working for speed.
- Then when time is up, continue working and finish as necessary.

| $\exp$ | inital [A] <br> $\left(\mathrm{mol} \mathrm{L}^{-1}\right)$ | inital [B] <br> $\left(\mathrm{mol} \mathrm{L}^{-1}\right)$ | initial rate of formation of [C] <br> $\left(\mathrm{mol} \mathrm{L}^{-1} \mathrm{sec}^{-1}\right)$ |
| :---: | :---: | :---: | :---: |
| 1 | 0.10 | 0.10 | $2.5 \times 10^{-4}$ |
| 2 | 0.20 | 0.10 | $5.0 \times 10^{-4}$ |
| 3 | 0.20 | 0.50 | $1.25 \times 10^{-2}$ |

1. The initial-rate data in the table above were obtained for the reaction represented below. What is the experimental rate law for the reaction.

$$
\mathrm{A}+\mathrm{B} \rightleftarrows 2 \mathrm{C}
$$

a. $\quad$ rate $=k[A][B]$
b. $\quad$ rate $=k[A][B]^{2}$
c. $\quad$ rate $=k[A][B]^{3}$
d. rate $=k[A]^{2}[B]^{2}$
e. rate $=k \frac{[A]}{[B]}$
2.

$$
\text { rate }=k[\mathrm{~A}]^{2}
$$

For the reaction for which rate law is given above, a plot of which of the following is a straight line?
a. $\frac{1}{[A]}$ versus time
b. [A] versus $\frac{1}{\text { time }}$
c. [A] versus time
d. $\ln [\mathrm{A}]$ versus time
e. $\ln [\mathrm{A}]$ versus $\frac{1}{\text { time }}$
3.

$$
2 \mathrm{X}_{(\mathrm{g})}+\mathrm{Y}_{(\mathrm{g})} \rightleftarrows 2 \mathrm{Z}_{(\mathrm{g})}
$$

When the concentration of substance Y in the reaction above is doubled, all other factors being held constant, it is found that the rate of the reaction quadruples. The most probable explanation for this observation is that
a. substance X is not involved in any of the steps of the mechanism.
b. substance Y is involved in the rate-determining step of the mechanism but is not involved in subsequent steps.
c. substance Y is probably a catalyst and thus speeds up the rate of the reaction.
d. the reactant with the smallest coefficient in the balanced equation generally has the greatest effect on the rate of the reaction.
e. the order of the reaction with respect to Y is 2 .
4. The proposed steps for a catalyzed reaction between $\mathrm{X}^{4+}$ and $\mathrm{Z}^{+}$are represented below.

$$
\begin{array}{ll}
\text { step 1: } & \mathrm{X}^{4+}+\mathrm{Y}^{2+} \rightarrow \mathrm{X}^{3+}+\mathrm{Y}^{3+} \\
\text { step 2: } & \mathrm{X}^{4+}+\mathrm{Y}^{3+} \rightarrow \mathrm{X}^{3+}+\mathrm{Y}^{4+} \\
\text { step 3: } & \mathrm{Y}^{4+}+\mathrm{Z}^{+} \rightarrow \mathrm{Z}^{3+}+\mathrm{Y}^{2+}
\end{array}
$$

The catalyst in this process is
a. $\mathrm{X}^{4+}$
b. $\mathrm{X}^{3+}$
c. $\mathrm{Y}^{4+}$
d. $\mathrm{Y}^{2+}$
e. $Z^{+}$
5. The balanced equation for the reaction of nitrogen dioxide and fluorine is

$$
2 \mathrm{NO}_{2}+\mathrm{F}_{2} \rightarrow 2 \mathrm{NO}_{2} \mathrm{~F}
$$

The proposed mechanism is
step 1: $\mathrm{NO}_{2}+\mathrm{F}_{2} \rightarrow \mathrm{NO}_{2} \mathrm{~F}+\mathrm{F}$
(slow)
step 2: $\mathrm{F}+\mathrm{NO}_{2} \rightarrow \mathrm{NO}_{2} \mathrm{~F}$
Which of the following are correct?
I. The mechanism supports an experimentally determined rate law of rate $=\mathrm{k}\left[\mathrm{NO}_{2}\right]^{2}\left[\mathrm{~F}_{2}\right]$
II. F is an intermediate
III. The reaction is first order in $\mathrm{F}_{2}$
a. I only
b. I and II only
c. I and III only
d. II and III only
e. I, II, and III
6. Each of the following is true about heterogenous catalyst EXCEPT
a. Its presence changes the rate of chemical change.
b. It does not undergo a permanent change.
c. It is in the same phase as the reacting particles.
d. It's presence lowers the activation energy of the overall reaction.
e. Its presence decreases the potential energy of the activated complex.

## The Reaction below should be used for the next 3 questions

$$
\mathrm{IO}_{3}^{-}+5 \mathrm{I}^{-}+6 \mathrm{H}^{+} \rightarrow 3 \mathrm{I}_{2}+3 \mathrm{H}_{2} \mathrm{O}
$$

7. The oxidation of iodide ions by iodate ions in acidic aqueous solution occurs according to the stoichiometry shown above. The experimentally determined rate law of the reaction is

$$
\text { rate }=k\left[\mathrm{IO}_{3}^{-}\right]\left[\mathrm{I}^{-}\right]
$$

What is the order with respect to $\mathrm{IO}_{3}{ }^{-}$?
a. 1
b. 2
c. 3
d. 5
e. 6
8. According to the rate law for the reaction, an increase in the concentration of hydrogen ion has what effect on this reaction?
a. The rate of reaction increases.
b. The rate of the reaction decreases.
c. The value of the rate law constant increases.
d. The value of the rate law constant decreases.
e. Neither the rate nor the value of the rate law constant is changed.
9. According to the rate law for the reaction, a decrease in the concentration of iodide ion has what effect on this reaction?
a. The rate of the reaction decreases and the value of the rate law constant decreases.
b. The rate of the reaction decreases and the value of the rate law constant remains the same.
c. The rate of the reaction remains the same and the value of the rate law constant increases.
d. The rate of the reaction decreases and the value of the rate law constant increases.
e. Neither the rate nor the value of the rate law constant is changed.
10.

$$
\mathrm{A}_{(\mathrm{g})}+\mathrm{B}_{(\mathrm{g})} \rightarrow \mathrm{C}_{(\mathrm{g})}
$$

All of the following apply to the reaction above as it is carried out at constant temperature in a sealed rigid container EXCEPT
a. The total pressure decreases.
b. The rate of the reaction decreases.
c. The entropy of the system decreases
d. The number of molecules of C decreases.
e. The frequency of collisions between molecules A and $B$ decreases.
11. Consider the hypothetical reaction.

$$
\begin{gathered}
\mathrm{X}_{(\mathrm{g})}+2 \mathrm{Y}_{(\mathrm{g})} \rightarrow \mathrm{XY}_{2 \mathrm{~g})} \\
\frac{\Delta[Y]}{\Delta t}=-5.0 \times 10^{-2} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{sec}^{-1}
\end{gathered}
$$

What is the rate of formation of $\mathrm{XY}_{2(\mathrm{~g})}$
a. $-5.0 \times 10^{2-} \mathrm{mol} \mathrm{L}^{-1} \mathrm{sec}^{-1}$
b. $-2.5 \times 10^{2-} \mathrm{mol} \mathrm{L}^{-1} \mathrm{sec}^{-1}$
c. $1.0 \times 10^{-2} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{sec}^{-1}$
d. $2.5 \times 10^{-2} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{sec}^{-1}$
e. $\quad 5.0 \times 10^{-2} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{sec}^{-1}$
12. Consider the hypothetical reaction below taking place in a syringe at constant temperature. The reaction is started by placing $\mathrm{A}_{(\mathrm{g})}$ and $\mathrm{B}_{(\mathrm{g})}$ in the syringe.
No chemical equilibrium is established.

$$
\mathrm{A}_{(\mathrm{g})}+2 \mathrm{~B}_{(\mathrm{g})} \rightarrow \mathrm{AB}_{2(\mathrm{~g})}
$$

Which change causes an immediate increase in the partial pressure $\mathrm{B}_{(\mathrm{g})}$ ?
I. addition of $\mathrm{B}_{(\mathrm{g})}$ at constant volume
II. decrease in the volume of the reaction vessel
III. addition of $\mathrm{AB}_{2(\mathrm{~g})}$ at constant volume
a. I only
b. II only
c. I and II only
d. II and III only
e. I, II, and III
13. For two first order reactions of different substances $A$ and X

$$
\begin{array}{ll}
\mathrm{A} \rightarrow \mathrm{~B} & \mathrm{t} 1 / 2=30.0 \mathrm{~min} \\
\mathrm{X} \rightarrow \mathrm{Y} & \mathrm{t} 1 / 2=60.0 \mathrm{~min}
\end{array}
$$

This means that
a. doubling the concentration of A will have $1 / 2$ the effect on half-life that doubling the concentration of B will have on its half-life
b. a certain number of grams of A will react twice as fast as the same number of grams of $X$
c. a certain number of grams of $X$ will react twice as fast as the same number of grams of A
d. the rate constant of $\mathrm{A} \rightarrow \mathrm{B}$ is lower than the rate constant of $\mathrm{X} \rightarrow \mathrm{Y}$.
e. 3 moles of A will react more rapidly than 3 moles of X.

Consider the reaction and its rate law given below for the next six questions.

$$
\begin{gathered}
2 \mathrm{~A}_{(\mathrm{g})}+\mathrm{B}_{(\mathrm{g})} \rightarrow \mathrm{C}_{(\mathrm{g})} \\
\text { rate }=k[\mathrm{~A}]^{2}[\mathrm{~B}]
\end{gathered}
$$

At the beginning of one trial of this reaction
$[\mathrm{A}]=4.0$ and $[\mathrm{B}]=1.0$.
The rate of formation of C was $0.048 \mathrm{~mole}^{-1} \mathrm{sec}^{-1}$
14. The numerical value of k , the rate constant, for this reaction is closest to
a. $3 \times 10^{2}$
b. $4 \times 10^{0}$
c. $8 \times 10^{-1}$
d. $1 \times 10^{-2}$
e. $3 \times 10^{-3}$
15. Which is the label for $k$, the rate constant?
a. $\mathrm{mol}^{2} \mathrm{~L}^{-2} \mathrm{sec}^{-1}$
b. $\mathrm{L} \mathrm{mol}^{-1} \mathrm{sec}^{-1}$
c. $\quad \mathrm{L}^{2} \mathrm{~mol}^{-2} \mathrm{sec}^{-1}$
d. $\mathrm{L}^{2} \mathrm{sec} \mathrm{mol}^{-1}$
e. $\mathrm{L}^{2} \mathrm{sec} \mathrm{mol}^{-2}$
16. When [B] decreases 0.4 M , what will be the value of [A]?
a. $\quad 0.8 \mathrm{M}$
b. $\quad 1.6 \mathrm{M}$
c. $\quad 2.8 \mathrm{M}$
d. $\quad 3.4 \mathrm{M}$
e. $\quad 3.6 \mathrm{M}$
17. Which of the following describes how the rate for this trial reaction, at constant temperature, changes as [B] approaches $0.4 \mathrm{~mol} \mathrm{~L}^{-1}$ ?
a. The rate decreases because the concentration of the products increases.
b. The rate remains the same because the rate constant remains the same.
c. The rate remains the same because the temperature remains the same.
d. The rate decreases because the concentration of the reactants decreases.
e. The rate remains the same because the energy of activation remains the same.
18. Which applies to this reaction as it proceeds at constant temperature?
I. The rate of the reaction decreases.
II. The effectiveness of collisions between reactant molecules remains the same.
III. The frequency of collisions between reactant molecules remains the same.
a. I only
b. II only
c. I and II only
d. II and III only
e. I, II, and III
19. Which applies to this system when its temperature increases at constant volume?
I. [A] decreases at a greater rate.
II. The value for $k$, the rate constant, remains the same.
III. The rate of the reaction increases.
a. I only
b. II only
c. I and III only
d. II and III only
e. I, II, and III
20. Which applies to any reaction mechanism?
I. It is a list of steps that produce the overall chemical reaction.
II. It includes only unimolecular steps.
III. It cannot include reaction intermediates.
a. I only
b. II only
c. III only
d. I and II only
e. I and III only
21. The first order decomposition of some radioactive isotope is 3 days. Approximately what percentage of the original substance will have decayed after 12 days have passed?
a. $94 \%$
b. $88 \%$
c. $50 \%$
d. $25 \%$
e. $6 \%$

## Consider the information below for the next five questions.

$$
5 \mathrm{Br}^{-}{ }_{(\mathrm{aq})}+\mathrm{BrO}_{3^{-(a q)}}^{-}+6 \mathrm{H}_{(\mathrm{aq})}^{+} \rightarrow 3 \mathrm{Br}_{2(\mathrm{~L})}+3 \mathrm{H}_{2} \mathrm{O}_{(\mathrm{L})}
$$

The reaction between bromide ions and bromate ions in acidic water solution occurs according to the equation above. The rate law for this reaction is known to be

$$
\text { rate }=k\left[\mathrm{Br}^{-}\right]\left[\mathrm{BrO}_{3}^{-}\right]\left[\mathrm{H}^{+}\right]^{2}
$$

22. The overall order for this reaction is
a. 2
b. 3
c. 4
d. 6
e. 12
23. One proposed reaction mechanism has three steps up to and including the slow step. The first step in the mechanism is

$$
\mathrm{Br}^{-}+\mathrm{H}^{+} \rightarrow \text { intermediate-A }
$$

Which statement(s) must be true about this first step in the reaction mechanism?
I. The coefficients of the reactants must correspond to the exponents on those same terms in the overall rate law.
II. No further step can include intermediate-A as a reactant.
III. A subsequent step must include $\mathrm{BrO}_{3}{ }^{-}$as a reactant.
a. I only
b. II only
c. III only
d. I and II only
e. I and III only
24. What is the effect of increasing $\left[\mathrm{H}^{+}\right]$in this reaction at a constant temperature?
a. The value of the rate constant increases.
b. The potential energy of the products decreases.
c. The potential energy of the activated complex decreases.
d. The number of collisions between $\mathrm{H}^{+}$and $\mathrm{Br}^{-}$ions increases.
e. The number of collisions between $\mathrm{H}^{+}$and $\mathrm{Br}^{-}$ions decreases.
25. Which change will cause a decrease in the rate of the reaction?
I. addition of $\mathrm{OH}^{-}$ions
II. removal of $\mathrm{H}^{+}$ions
III. addition of $\mathrm{H}_{2} \mathrm{O}$ molecules
a. I only
b. II only
c. I and II only
d. II and III only
e. I, II, and III
26. What is the effect of adding $\mathrm{Br}_{2(\mathrm{~L})}$ to the system? (Assume negligible change in volume.)
I. The mass of the system increases.
II. The rate of reduction of $\mathrm{BrO}_{3}{ }^{-}$increases.
III. The rate of oxidation of $\mathrm{Br}^{-}$decreases.
a. I only
b. II only
c. I and II only
d. II and III only
e. I, II, and III
27. The values for the change in enthalpy, $\Delta \mathrm{H}$, and the activation energy, $\mathrm{E}_{\mathrm{a}}$, for a given reaction are known. The value of $E_{a}$ for the reverse reaction equals
a. Ea for the forward reaction.
b. $-\left(E_{a}\right)$ for the forward reaction.
c. the sum of $-(\Delta H)$ and $E_{a}$
d. the sum of $\mathrm{E}_{\mathrm{a}}$ and $\Delta \mathrm{H}$
e. the difference between $\Delta \mathrm{H}$ and $\mathrm{E}_{\mathrm{a}}$
28. Which accounts for the increase in the rate of reaction when a catalyst is added to a reaction system?
a. decrease in $\Delta \mathrm{H}$ for the reaction
b. increase in $\Delta \mathrm{S}$ for the reaction
c. increase in potential energy of the reactants
d. increase in potential energy of the products
e. decrease in potential energy of the activated complex
29. The rate constant for a gas phase reaction can be increased by doing which of the following?
I. Increasing the concentration of reactants
II. Increasing the temperature
III. Decreasing the size of the container
a. I only
b. II only
c. I and II only
d. I and III only
e. I, II, and III only
30. How does a catalyst increase the rate of a chemical reaction?
a. By changing the magnitude of the enthalpy change.
b. By providing a pathway for the reaction that has a lower activation energy.
c. By decreasing the potential energy of the products
d. By increasing the potential energy of the reactants
e. By giving energy to the reactants so more of them have energies greater than the activation energy.
31. Collision theory predicts all of the following EXCEPT that
a. a reaction will only occur if the collision geometry is correct
b. a reaction will not occur if the collision occurs with energy that is less than the activation energy.
c. a reaction will not occur if the reactants do not collide
d. frequency of collisions will increase with increasing temperature.
e. more successful collisions will occur for a reaction with a larger activation energy.
32. For the chemical reaction $\mathrm{H}_{2}+\mathrm{Br}_{2} \rightarrow 2 \mathrm{HBr}$, what can be said of the relative rates of consumption of the reactants when compared to the formation of product?
a. HBr will be produced at half the rate that bromine is consumed.
b. HBr will be produced at the same rate that bromine is consumed.
c. $\quad \mathrm{Br}_{2}$ and $\mathrm{H}_{2}$ will be consumed at different rates.
d. HBr will be produced at twice the rate that bromine is consumed
e. It is not possible to tell anything about relative rates without the rate law.
33. Which energy distribution diagram represents an increase in temperature?
(A)

(B)


(C)

(D)

(E)

34. Which pair of potential energy/reaction coordinate diagrams represents a comparison between the same reactants where one of the pair of diagrams represents a situation that includes a catalyst?
(A)

(B)

(B)

(C)

(C)

(D)

(D)

(E)

(E)


The following 4 questions refer to a reaction involving three reactants: $A, B, C$. Select from choices below:
(A) Rate $=k[\mathrm{~A}]^{0}$
(B) Rate $=k[\mathrm{~A}]^{2}$
(C) Rate $=k[\mathrm{~A}][\mathrm{B}]^{3}$
(D) Rate $=k[\mathrm{~A}][\mathrm{B}]^{2}[\mathrm{C}]^{2}$
(E) Rate $=k[\mathrm{~A}]^{2}[\mathrm{~B}]$
36. A rate law for which the units of rate are equal to the units of the rate constant
37. A rate law for a reaction that is 3 rd order overall
38. A rate law that represents a reaction in which doubling the concentration of A and doubling the concentration of B (keeping all other variables constant) will lead to a four fold increase in the rate
39. A rate law that represent a reaction in which doubling the concentration of A and halving the concentration of B (keeping all other variables constant) will lead to a two fold increase in the rate.

The following 3 questions refer to the choices below:
(A) Reaction Mechanism
(B) Catalyst
(C) Rate-Determining step
(D) Enthalpy Change
(E) Collision Theory
40. Describes the series of steps that a chemical reaction is broken in to
41. Is present at the start of a reaction and is also present at the end of a reaction
42. The specific part of a chemical reaction that can be used to determine the rate equation
43. A sealed rigid container holds two gases, A and B . If additional moles of gas B are added to the container while keeping the temperature constant, what can be said of the reaction under these conditions.
a. The rate of the reaction will increase, as will the rate constant, k .
b. the rate of the reaction will remain unchanged as will the rate constant, k .
c. the rate of the reaction will increase but the rate constant, k , will remain unchanged.
d. the rate of the reaction will remain unchanged but the rate constant, k , will increase.
e. the rate of reaction will decrease as will the rate constant, k .
44. A reaction occurs in the gas phase according to the chemical equation shown below. which statement correctly reflects the relative changes in concentrations of reactant and products in the forward reaction?

$$
\mathrm{N}_{2}+3 \mathrm{H}_{2} \rightarrow 2 \mathrm{NH}_{3}
$$

a. $\left[\mathrm{H}_{2}\right]$ decreases at the same rate $\left[\mathrm{N}_{2}\right]$ decreases.
b. $\left[\mathrm{H}_{2}\right]$ decreases at $2 / 3$ the rate $\left[\mathrm{NH}_{3}\right]$ increases
c. $\left[\mathrm{N}_{2}\right]$ increases at the same rate as $\left[\mathrm{NH}_{3}\right]$ decreases
d. $\left[\mathrm{N}_{2}\right]$ increases at $1 / 2$ the rate $\left[\mathrm{NH}_{3}\right]$ decreases
e. $\left[\mathrm{N}_{2}\right]$ decreases at $1 / 2$ the rate $\left[\mathrm{NH}_{3}\right]$ increases
45. For a reaction that is first order with respect to [D] and second order with respect to [E], which of the following will result in no change to the overall reaction rate.
a. Doubling [D] and halving [E]
b. Doubling [E] and halving [D]
c. Doubling [D] and Doubling [E]
d. Increasing [D] by a factor of four and halving [E]
e. Increasing [D] by a factor of four and dividing [E] by a factor of four.
46. The rate of a chemical reaction will be increased by all of the following factors EXCEPT
a. the presence of a high activation energy
b. the presence of a catalyst
c. increasing the surface area of a reactant
d. an increase in temperature
e. increasing the concentration of the reactants

47. In the Maxwell-Boltzman distribution plot for a given reaction at two different temperatures, what can be said about the total area underneath the two curves when comparing the curve for the relatively hot temperature to the curve for the relatively low temperature.
a. the total area under the hot curve is greater since more particles possess the minimum activation energy required.
b. the total area under the cold curve is greater since more particles possess the minimum activation energy required
c. the total areas are the same since the number of moles of gas does not change as a result of heating or cooling.
d. the total areas are the same since the reaction has a fixed activation energy
e. the total areas are the same since gases occupy the same space regardless of temperature.
48. Which of the following statements is/are true in relation to rate laws?
I. A rate law must be determined experimentally and cannot be deduced from a chemical equation alone.
II. Zero-order have rate constants equal to rates
III. A "straight line" graph can be obtained for a first-order reaction by plotting the reciprocal of the concentration of reactant ( $y$-axis) against time ( x -axis)
a. I only
b. II only
c. III only
d. I and II only
e. I,II, and III
49. If the units in the rate in a given experiment are $\mathrm{mol}^{-1}$ $\min ^{-1}$ and the units of all concentrations are $\mathrm{mol} \mathrm{L}^{-1}$, then the units of the rate constant associated with the rate law

$$
\text { Rate }=k[A]^{2}[B]:
$$

a. $\mathrm{mol} \mathrm{L}^{-1} \min ^{-1}$
b. $\quad \mathrm{L}^{2} \mathrm{~mol}^{-2} \mathrm{~min}^{-1}$
c. $\mathrm{mol}^{2} \mathrm{~L}^{-2} \mathrm{~min}^{-1}$
d. $\mathrm{mol}^{-3} \mathrm{~L}^{3} \mathrm{~min}^{-1}$
e. $\mathrm{mol}^{-2} \mathrm{~L}^{-2} \mathrm{~min}^{-1}$
50. For the rate law shown below, where the rate is measured in units of $\mathrm{M} \mathrm{s}^{-1}$ and the concentrations of X and Y are measured in M , which of the following statements is true?

$$
\text { Rate }=\mathrm{k}[\mathrm{X}]^{2}[\mathrm{Y}]
$$

I. the units of $\mathrm{k}=\mathrm{M}^{2} \mathrm{~s}^{-1}$
II. doubling the concentration of $y$ while keeping all other conditions constant will not affect the rate.
III. The reaction can be described as third-order.
a. I only
b. III only
c. I and II only
d. II and III only
e. I, II, and III
51. For the rate law, Rate $=k[Q]^{2}[R]^{2}$, which statement is correct?
a. The reaction is fourth order overall, and doubling both the $[Q]$ and $[R]$ while keeping the temperature constant will lead to a bigger k and a 16 -fold increase in the rate
b. The reaction is fourth order overall, and doubling both the $[\mathrm{Q}]$ and $[\mathrm{R}]$ while keeping the temperature constant will lead to a 16 -fold increase in the rate
c. The reaction is fourth order overall, and doubling both the $[\mathrm{Q}]$ and $[\mathrm{R}]$ while keeping the temperature constant will lead to a bigger k and an 8 -fold increase in the rate
d. The reaction is second order overall, and doubling both the $[\mathrm{Q}]$ and $[\mathrm{R}]$ while keeping the temperature constant will lead to a bigger k and a 16 -fold increase in the rate
e. The reaction is second order overall, and doubling both the $[\mathrm{Q}]$ and $[\mathrm{R}]$ while keeping the temperature constant will lead to a 16 -fold increase in the rate
52. The reaction $\mathrm{X}+\mathrm{Y} \rightarrow \mathrm{Z}$ has the rate law, Rate $=\mathrm{k}[\mathrm{X}]^{2}[\mathrm{Y}]$. If the concentration of X is doubled and the concentration of Y is tripled, the rate would be increased by a factor of
a. 2
b. 4
c. 5
d. 6
e. 12

For the next 2 questions, the proposed mechanism for a given chemical reaction is shown below.

$$
\begin{array}{lll}
\text { Step 1: } & \mathrm{A}+\mathrm{B} \leftrightharpoons \mathrm{~F} & \text { (fast) } \\
\text { Step 2: } & \mathrm{F}+\mathrm{A} \rightarrow \mathrm{C}+\mathrm{G} & \text { (slow) } \\
\text { Step 3: } & \mathrm{G}+\mathrm{A} \leftrightharpoons \mathrm{C}+\mathrm{D} & \text { (fast) }
\end{array}
$$

53. Which is the correct rate law?
a. $\quad$ rate $=\mathrm{k}[\mathrm{A}]^{2}[\mathrm{~B}]$
b. $\quad$ rate $=k[A][B][F][G]$
c. $\quad$ rate $=k[A][B]$
d. $\quad$ rate $=k[A]^{3}[\mathrm{~B}]$
e. $\quad$ rate $=k[\mathrm{~A}]^{2}[\mathrm{~B}][\mathrm{F}]$
54. Which reaction represents the overall equation?
a. $\mathrm{A}+\mathrm{B} \leftrightharpoons \mathrm{F}$
b. $\quad 3 \mathrm{~A}+\mathrm{B} \rightarrow 2 \mathrm{C}+\mathrm{D}$
c. $3 \mathrm{~A}+\mathrm{B}+\mathrm{F}+\mathrm{G} \rightarrow 2 \mathrm{C}+\mathrm{D}+\mathrm{G}+\mathrm{F}$
d. $\mathrm{A}+\mathrm{B} \rightarrow \mathrm{C}+\mathrm{D}$
e. $\mathrm{A}_{3}+\mathrm{B} \rightarrow \mathrm{C}_{2}+\mathrm{D}$
55. Which answer choice correctly matches the integrated rate law shown:

$$
\frac{1}{[A]_{t}}=k t+\frac{1}{[A]_{o}}
$$

a. A zero order reaction where a plot of
$\frac{1}{[A]_{t}}$ versus $t$ will be a straight line with slope $=\mathrm{k}$
b. A first order reaction where a plot of
$\frac{1}{[A]_{t}}$ versus $t$ will be a straight line with slope $=\mathrm{k}$
c. A first order reaction where a plot of
$\frac{1}{[A]_{t}}$ versus $t$ will be a straight line with slope $=-\mathrm{k}$
d. A second order reaction where a plot of
$\frac{1}{[A]_{t}}$ versus $t$ will be a straight line with slope $=\mathrm{k}$
e. A second order reaction where a plot of $\frac{1}{[A]_{t}}$ versus $t$ will be a straight line with slope $=-\mathrm{k}$
56. Which of the following plots will yield a straight line for a first order reaction where the slope $=-k$ ?
a. $\quad \ln [\mathrm{A}]$ versus time
b. [A] versus rate
c. $\frac{1}{[A]}$ versus time
d. time versus $\frac{1}{[A]}$
e. $\quad \ln k$ versus time
57. The reaction of a tertiary haloalkane (represented by RX below) with hydroxide ions is found to follow the mechanism:

$$
\begin{array}{ll}
\text { Step 1 } & \mathrm{RX} \rightarrow \mathrm{R}^{+}+\mathrm{X}^{-} \\
\text {Step } 2 & \mathrm{R}^{+}+\mathrm{OH}^{-} \rightarrow \mathrm{ROH}
\end{array}
$$

If Step 1 is found to be the slow step in the mechanism, which of the following statements is true?
I. The reaction will be found to be first-order with respect to RX
II. The reaction has an intermediate that is represented by $\mathrm{R}^{+}$.
III. The reaction rate is independent of the concentration of hydroxide ions.
a. I only
b. II only
c. III only
d. I and II only
e. I, II, and III
58. For a mechanism to be feasible, which of the following criteria must be filled?
I. The sum of the overall steps must add up to the overall stoichiometry of the chemical equation
II. The rate determining (slow step) must be the last step in the mechanism.
III. The rate determining (slow step) must include all of the reactants that appear in the rate equation where the stoichiometric coefficients match the powers.
a. I only
b. III only
c. I and III only
d. II and III only
e. I, II, and III
59. Which of the following is true of all catalysts?
a. They are used up in chemical reactions
b. They are always transition metals
c. They do not take part in chemical reactions.
d. They work by increasing the activation energy of a reaction.
e. They are present at the beginning of a a reaction and are unchanged at the end.

The following 2 questions refer to the following equation and table below:

$$
2 \mathrm{~W}+3 \mathrm{X}+\mathrm{Y} \rightarrow \mathrm{Z}
$$

| Trial | $[\mathbf{W}]$ <br> $\mathbf{m o l ~ L}^{-\mathbf{1}}$ | $[\mathbf{X}]$ <br> $\mathbf{m o l ~ L}^{-\mathbf{1}}$ | $[\mathbf{Y}]$ <br> $\mathbf{m o l ~ L}^{-\mathbf{1}}$ | Rate of Production of Z <br> $\mathbf{m o l ~ L}^{-\mathbf{1}}$ |
| :---: | :---: | :---: | :---: | :---: |
| 1 | 0.001 | 0.001 | 0.001 | $2.1 \times 10^{-4}$ |
| 2 | 0.002 | 0.001 | 0.001 | $2.1 \times 10^{-4}$ |
| 3 | 0.001 | 0.003 | 0.001 | $6.3 \times 10^{-4}$ |
| 4 | 0.002 | 0.001 | 0.0005 | $5.25 \times 10^{-5}$ |

60. The rate law is:
a. rate $=k[\mathrm{~W}]^{2}[\mathrm{X}]^{3}[\mathrm{Y}]$
b. rate $=k[\mathrm{~W}]^{2}[\mathrm{X}][\mathrm{Y}]$
c. $\quad$ rate $=k[\mathrm{~W}][\mathrm{X}][\mathrm{Y}]$
d. $\quad$ rate $=k[\mathrm{X}]^{3}[\mathrm{Y}]^{2}$
e. $\quad$ rate $=k[\mathrm{X}][\mathrm{Y}]^{2}$
61. The rate of consumption of X in Experiment 1 is:
a. $\quad 2.1 \times 10^{-4} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{~min}^{-1}$
b. $-2.1 \times 10^{-4} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{~min}^{-1}$
c. $\quad 6.3 \times 10^{-4} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{~min}^{-1}$
d. $-6.3 \times 10^{-4} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{~min}^{-1}$
e. $\quad-4.2 \times 10^{-4} \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{~min}^{-1}$
62. If tripling the concentration of a single reactant R in a multi-reactant reaction (while leaving all other conditions unchanged) leads to a nine-fold increase in rate, it can be deduced
a. that the complete rate law is Rate $=k[R]$
b. that the complete rate law is Rate $=k[R]^{2}$
c. that the complete rate law is Rate $=k[R]^{3}$
d. that the complete rate law is Rate $=k[R]^{9}$
e. only that the order with respect to $[R]$ is 2

For the next 2 questions consider the proposed mechanism shown below for the reaction between nitrogen dioxide and carbon monoxide.

$$
\begin{array}{lll}
\text { Step 1: } & \mathrm{NO}_{2}+\mathrm{NO}_{2} \rightarrow \mathrm{NO}_{3}+\mathrm{NO} & \text { (slow) } \\
\text { Step 2: } & \mathrm{NO}_{3}+\mathrm{CO} \rightarrow \mathrm{NO}_{2}+\mathrm{CO}_{2} & \text { (fast) }
\end{array}
$$

63. What is the Rate law for the reaction?
a. Rate $=k\left[\mathrm{NO}_{2}\right]$
b. Rate $=k\left[\mathrm{NO}_{2}\right]^{2}$
c. Rate $=k\left[\mathrm{NO}_{2}\right]^{2}\left[\mathrm{NO}_{3}\right][\mathrm{NO}]$
d. Rate $=k\left[\mathrm{NO}_{2}\right]\left[\mathrm{NO}_{3}\right][\mathrm{NO}]$
e. Rate $=k\left[\mathrm{NO}_{3}\right][\mathrm{CO}]$
64. Identify any intermediates in the Reaction:
a. $\mathrm{NO}_{2}$ only
b. $\mathrm{NO}_{3}$ only
c. $\mathrm{NO}_{2}$ and $\mathrm{NO}_{3}$
d. $\mathrm{NO}, \mathrm{NO}_{2}$, and $\mathrm{NO}_{3}$
e. $\mathrm{NO}_{2}, \mathrm{NO}_{3}$, and NO

The following 3 questions refer to the popular demonstration called "Elephants Toothpaste" in which the mechanism is believed to be:

$$
\begin{array}{ll}
\text { Step 1: } & \mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{I}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{OI}^{-} \\
\text {Step 2: } & \mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{OI}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2}+\mathrm{I}^{-}
\end{array}
$$

65. Identify the catalyst.
a. $\mathrm{H}_{2} \mathrm{O}_{2}$
b. $\mathrm{I}^{-}$
c. $\mathrm{H}_{2} \mathrm{O}$
d. $\mathrm{OI}^{-}$
e. $\mathrm{O}_{2}$
66. Identify the Intermediate:
a. $\mathrm{H}_{2} \mathrm{O}_{2}$
b. $\mathrm{I}^{-}$
c. $\mathrm{H}_{2} \mathrm{O}$
d. $\mathrm{OI}^{-}$
e. $\mathrm{O}_{2}$
67. Which chemical equation is consistent with the mechanism provided?
a. $2 \mathrm{H}_{2} \mathrm{O}_{2} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2}$
b. $2 \mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{I}^{-} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2}$
c. $\quad 1 / 2 \mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{I}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2}$
d. $\quad 1 / 2 \mathrm{H}_{2} \mathrm{O}_{2}+\mathrm{I}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{OI}^{-}+\mathrm{O}_{2}$
e. $2 \mathrm{H}_{2} \mathrm{O}_{2} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}+\mathrm{OI}^{-}+\mathrm{O}_{2}$
68. The nuclear decay of an isotope of this element is used to measure the age of archaeological artifacts.
a. C
b. O
c. N
d. $F$
e. Ne
69. If $75 \%$ of a sample of pure ${ }^{3}{ }_{1} \mathrm{H}$ decays in 24.6 years, what is the half- life of ${ }^{3}{ }_{1} \mathrm{H}$
a. 24.6 years
b. $\quad 18.4$ years
c. 12.3 years
d. 6.15 years
e. 3.07 years
70. Strontium-90 decays through the emission of beta particles. It has a half- life of 29 years. How long does it take for 80 percent of a sample of strontium-90 to decay?
a. 9.3 years
b. 21 years
c. 38 years
d. 67 years
e. 96 years
71. After 44 minutes, a sample of ${ }^{44}{ }_{19} \mathrm{~K}$ is found to have decayed to 25 percent of the original amount present. What is the half-life of ${ }^{44}{ }_{19} \mathrm{~K}$ ?
a. 11 minutes
b. 22 minutes Answer Key said (e)
c. 44 minutes
d. 66 minutes
e. 88 minutes
72. A sample of radioactive cadmium has a half life of 15 days. If you have a sample that originally weighed 40 g and later weighs 2.5 g . How much time elapsed between the initial and final masses?
a. 600 days
b. 100 days
c. 60 days
d. 45 days
e. $\quad 37.5$ days
73. The half-life of ${ }^{14} \mathrm{C}$ is 5570 years. How many years will it take for $90 \%$ of a sample to decompose?
a. 5,013 years
b. 11,000 years
c. 18,600 years
d. 23,000 years
e. 50,130 years
74. Tritium decays by a first-order process that has half-life of 12.5 years. How many years will it take to reduce the radioactivity of a tritium sample to $15 \%$ of it original value?
a. 64 yrs
b. 54 yrs
c. 34 yrs
d. 24 yrs
e. 14 yrs
75. For a first order reaction that has a half-life of 69 s at $80^{\circ} \mathrm{C}$, the value of the rate constant, k , is closest to?
a. $\quad 0.01 \mathrm{~s}^{-1}$
b. $0.1 \mathrm{~s}^{-1}$
c. $1 \mathrm{~s}^{-1}$
d. $10 \mathrm{~s}^{-1}$
e. $100 \mathrm{~s}^{-1}$
76. If the half-life of a reaction is independent of concentration, what is the order of the reaction?
a. zero
b. first
c. second
d. half-life is unrelated to the order of the reaction
e. unable to be determined without knowing starting concentrations
77. A particular nuclear decay has a rate constant of 0.00346 $\mathrm{min}^{-1}$ What is the half-life?
a. 3.3 hours
b. 1.6 hours
c. $\quad 0.0012 \mathrm{~min}$
d. 0.0024 min
e. $\quad 0.0050 \mathrm{~min}$
78. The half-life of ${ }^{99} \mathrm{Tc}$ is 6.00 hours. If it takes exactly 12.00 hours for the manufacturer to deliver a ${ }^{99} \mathrm{Tc}$ sample to a hospital, how much must be shipped in order for the hospital to receive 10.0 mg ?
a. $\quad 40.0 \mathrm{mg}$
b. $\quad 30.0 \mathrm{mg}$
c. $\quad 20.0 \mathrm{mg}$
d. $\quad 15.0 \mathrm{mg}$
e. $\quad 4.0 \mathrm{mg}$
79. b Use the table to check the relationship between concentration changes and rate changes. $\frac{\operatorname{rate}_{2}}{\text { rate }_{1}}=\frac{k_{2}}{k_{1}}\left(\frac{\operatorname{concA} A_{2}}{\operatorname{conc} A_{1}}\right)^{x}\left(\frac{\operatorname{conc} B_{2}}{\operatorname{conc} B_{1}}\right)^{y}$ and substitute to find the order of A $\frac{5 \times 10^{-4}}{2.5 \times 10^{-4}}=\frac{k_{2}}{k_{1}}\left(\frac{0.20}{0.10}\right)^{x}\left(\frac{0.10}{0.10}\right)^{y}$ thus $2=(2)^{x}$ and $\mathrm{x}=1$, thus first order with respect to A. To solve for order of B use trials $3 / 2 \frac{125 \times 10^{-4}}{5 \times 10^{-4}}=\frac{k_{2}}{k_{1}}\left(\frac{0.20}{0.20}\right)^{1}\left(\frac{0.50}{0.10}\right)^{y}$ thus $25=(5)^{y}$ which means $\mathrm{y}=2$, thus second order with respect to reactant B
80. a The order of the reaction indicates which plot will give a straight line. Since you are told second order, it is the plot of $\frac{1}{[A]}$ vs time that will be a straight line.
81. e Second order: $[\text { conc } \times 2]^{2}=$ rate $\times 4$
82. d A catalyst is a chemical that gets used up during an early step, but returns as a product during a later step.
83. d Since the first step is the slow step, the stoichiometry of that step results in the order of the reaction, thus it s first order for both $\mathrm{NO}_{2}$ and $\mathrm{F}_{2}$. Also, F is an intermediate because it is a product of one step, but gets used up in a subsequent step.
84. c "Same phase" meaning liquid vs solid vs gas, heterogeneous catalyst are in a different phase than the reactants they are catalyzing.
85. a It doesn't get much easier than this!
86. e No effect on the rate since the reaction is zero order with respect to $\mathrm{H}^{+}$
87. b While the rate changes, the rate constant will remain constant. Rate constants only change when temperature changes or in the presence of a catalyst. Rate constants do not change with concentration or pressure changes.
88. d The pressure decreases because two molecules are converted into one. The rate of all reactions decrease over time. The entropy decreases because fewer molecules result from the reaction. Of course the number of C molecules will increase during the reaction. Since the concentration of A and B will decrease over the course of the reaction, their collisions will decrease.
89. $d$ The formation of the product is half of the rate of the loss of $Y$. It is correct to change the sign to positive, because we are representing the formation of product, and the negative sign on the reactant to represent loss.
90. c Since it is stated that equilibrium is not established, there is no significant reverse reaction, thus adding product does not cause the formation of more B.
91. e Half life is an indication of reaction rate. Shorter half life indicates faster reactions.
92. e Given the rate law, insert the values given and solve for $\mathrm{k} ; \quad 0.048=k[4]^{2}[1]$
93. c Since the reaction is third order overall, with a molarity cubed, $\mathrm{M}^{3}$, the rate constant must have $\mathrm{M}^{-2}$ as well as the sec ${ }^{-1}$. Remember that $\mathrm{M}^{-2}$ is the same as $\mathrm{L}^{2} \mathrm{~mol}^{-2}$. Time is always in the denominator of every rate constant, and molarity will be in the denominator raised to one power less than the overall order of the reaction.
94. c You could make a RICE table for this reaction. since B becomes 0.4 and it started at 1 , it must have reacted 0.6 to get there. The $2: 1$ stoichiometry means that A will have reacted 1.2 , resulting in a final quantity of $2.8(4-1.2=2.8)$.
95. d Concentration of reactants always decrease over the course of all reactions.
96. c As the concentrations of the reactants decrease over the course of a reaction, the frequency of collisions will decrease, but the effectiveness of the collisions will remain the same, as that is a function of the molecules themselves, the temperature, and the presence of any catalyst.
97. c The reaction will proceed at a faster rate, so the loss of A will be at a greater rate. The rate constant changes at different temperatures. It increases at higher temps and decreases at lower temps.
98. a Yup, it's a list of steps.
99. a 12 days is 4 half lives, the amount decayed will be $100>50>25>12.5>6.25 \%$ remains, thus $94 \%$ of the original material will have decayed.
100. c Add the individual orders of each reactant to get the overall order.
101. c The coefficients of mechanism steps can be used to write the rate law of the individual steps but the high coefficients in the overall reaction, are certainly not likely to correspond to the order of the reaction with respect to each reactant. IntermediateA must be a reactant in a later step, just as $\mathrm{BrO}_{3}{ }^{-}$must also show up in a later step in or before the slow step.
102. d Increasing concentration always increases collision frequency, but may not always increase rate if a reaction is zero order with respect to the increasing reactant.
103. e Addition of $\mathrm{OH}^{-}$will react with $\mathrm{H}^{+}$reducing its concentration. Removal of $\mathrm{H}^{+}$will decrease the rate. Addition of $\mathrm{H}_{2} \mathrm{O}$ will dilute the reactants causing the reaction rate to decrease.
104. a $\mathrm{Br}_{2}$ is a liquid so it will have no effect on the reaction rate since the concentration will remain constant for a pure liquid regardless of how much is in the container.
105. c Consider the graph to the right and values of Ea and $\Delta \mathrm{H}$ to answer the question.
106. e The addition of a catalyst provides an alternative reaction pathway, with a modified or different activated complex that has lower potential energy. This lowers the activation energy, $E_{a}$, thus increasing the reaction rate.
107. b Changing the temperature will change the rate constant. Changing concentration and size of container will change the rate, but not the rate constant.
108. b An alternative pathway.
109. e Higher activation energy is a higher hump, and is harder to get over, making for fewer successful collisions.

110. $d$ The stoichiometry of the overall reaction will give us answers to the questions about comparative rates of formation of products compared to loss of reactants.
111. $d$ The curve shifts to the right and flattens out, making for more molecules under the curve that will have the minimum energy required for successful reaction collisions.
112. b Same $\Delta \mathrm{H}$, but a lower activation energy.
113. b Reactant and products at same level on both graphs, but a lower hump to represent a lower activation energy.
114. a
115. e
116. b
117. d
118. a
119. b
120. c
121. c
122. e
123. d
124. a
125. c
126. d
127. b
128. b
129. b
130. e
131. a
132. b
133. d
134. a
135. e
136. a
137. e
138. e
139. d
140. e
141. b
142. b
143. b
144. d
145. a
146. a
147. c
148. d
149. e
150. c
151. consider the decay $100 \gg 50 \gg 25 \gg 12.5$, thus just over 3 half lives occur. Looking over the answer options, c should be an easy pick.
152. c
153. a
154. b
155. a Use the equation $t_{\text {halfife }}=\frac{0.693}{k}$ and substitute: $t_{\text {halfife }}=\frac{0.693}{0.00346}$. Look for the easy math and solve for $\sim 200$ which is just over 3 hours.
156. a For this problem you must work backwards. If delivery takes 12 hours, that's two half lives. To have 10 mg remaining, work backwards $10 \ll 20 \ll 40 \mathrm{mg}$ must be sent.
