Unit 11: Kinetics And Equilibrium

Skills

1. Understand theory of reaction kinetics
2. Understand factors affecting reaction rate
3. Using Table I
4. Drawing and interpreting PE diagrams
5. Distinguishing between endo/exo PE diagrams
6. Defining and identifying changes in entropy
7. Understanding and defining equilibrium
8. Understanding equilibrium through Le Chatelier's Principle
9. Applying Le Chatelier with types of stressors

Unit 11: Vocabulary:

| Word |  |
| :---: | :--- |
| Kinetics |  |
| Rate Determining Step |  |
| Effective Collision |  |
| Exothermic |  |
| Potential Energy |  |
| Enthalpy |  |
| Activation Energy |  |
| Activated Complex |  |
| Forward Reaction |  |
| Ehemical Equilibrium |  |
| Shatelier's Principle |  |
| Stress |  |

Kinetics = study of the $\qquad$ or $\qquad$ at which reactions occur

- A REACTION is the BREAKING and REFORMING of $\qquad$ to make entirely new compounds as products
$\square$ Reaction Mechanism = STEP BY STEP PROCESS needed to make a product; how you get from "a" to "b" (like a recipe)

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                                    REACTANTS }->\mathrm{ PRODUCTS
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Just like when we bake a cake we must follow directions o CAN'T OMIT any STEPS!

- CAN'T CHANGE THE ORDER of the steps!
- CAN'T OMIT any REACTANTS (ingredients)

Determine whether each of the following chemical reactions is an example of a slow or fast reaction. Explain why knowing this relative rate of $r \times n$ is significant.

## Rusting

## Alka seltzer in water

Styrofoam decomposing

## Weathering of rocks

Bleach removing color
$\qquad$

## WHAT DETERMINES THE RATE OF A REACTION? Time to Race!!

1. NUMBER OF STEPS = more steps can mean a slower reaction
2. RATE DETERMINING STEP = the $\qquad$ of the reaction; most important factor influencing reaction rate

## Rate Determining Step

$$
\begin{array}{ll}
\mathrm{A}+\mathrm{B}+\mathrm{C} \rightarrow \mathrm{ABC} \quad \text { (2 steps) } \\
\mathrm{A}+\mathrm{B} \rightarrow \mathrm{AB} & \text { (slow) } \\
\mathrm{AB}+\mathrm{C} \rightarrow \mathrm{ABC} & \text { (fast) }
\end{array}
$$



Collision Theory: In order for are action to occur, reactant PARTICLES MUST $\qquad$ and have the following when doing so:

1. Proper amount of energy
2. Proper

ALIGNMENT/DIRECTION/ORIENTATION


Only when particles collide with these two conditions are met will there be an COLLISION, resulting in a reaction

Skill 2: Identify and understand 6 Factors Affecting Rate of Reaction:

| Factor | How Rate Affected | Why does it increase the rate? |
| :---: | :---: | :---: |
| 1. Nature of Reactants | $\qquad$ substances react <br> FASTER $\qquad$ substances reac $\dagger$ <br> SLOWER | $\square$ Ionic = smaller $\rightarrow$ <br> (Fewer bonds to break; Fewer Steps!) <br> Covalent = larger (MORE bonds to break; MORE steps): |
| 2. Concentration <br> MOLECULES ON UPPER LAYER HAVE A LOWER CHANCE OF COMBINING | INCREASE concentration, $\qquad$ rxn rate | The MORE PARTICLES in a given space, the LESS SPACE b/w particles $\rightarrow$ MORE COLLISIONS |
|  | Why does concentration affec | the rate of reaction? |
| 3. Pressure | INCREASE pressure, INCREASE rxn rate (affects GASES ONLY!) | Increasing pressure $\qquad$ VOLUME which DECREASES SPACE b/w particles $\rightarrow$ MORE COLLISIONS |
| 4. Temperature | INCREASE temperature, INCREASE rxn rate | Greater SPEED $\rightarrow$ $\qquad$ total COLLISIONS Greater AVERAGE KE $\rightarrow$ collisions take place with MORE energy |
| 5. Catalyst | SPEEDS UP THE RXN WITHOUT CHANGING THE NATURE OF THE REACTANTS/PRODUCT | Provides a SHORTCUT or $\qquad$ $\qquad$ for the mechanism Lowers the ACTIVATION ENERGY for the reaction |
|  | Define Catalysis: |  |
| 6. Surface Area | INCREASE the surface area (by making PIECES SMALLER) INCREASES the rxn rate (How many surfaces are there?) | Increasing surface area EXPOSES MORE REACTANT PARTICLES to possible collisions |

## Answer the following based on this lab and your knowledge of reaction rates:

1. Which event must always occur for a chemical reaction to take place?
(1) Formation of a precipitate
(2) Effective collisions between reaction particles
(3) Formation of a gas
(3) addition of a catalyst to the reaction system
2. Increasing the temperature increases the rate of a reaction by
(1) lowering the activation energy
(2) increasing the activation energy
(3) lowering the frequency of a effective collisions between reacting molecules
(4) Increasing the frequency of effective collisions between reacting molecules
3. A 5.0 gram sample of zinc and a 50. Millimeter sample of hydrochloric acid are using in a chemical reaction. Which combination of these samples has the FASTEST reaction rate?
(1) Zinc strip and 1.0 M HCl
(2) Zinc powder and 1.0 M HCl
(3) Zinc strip and 3.0 M HCl
(3) Zinc powder and 3.0 M HCl
4. What is required for a chemical reaction to occur?
A) standard temperature and pressure
B) a catalyst added to the reaction system
C) effective collisions between reactant particles
D) an equal number of moles of reactants and products
5. Given the balanced equation representing a reaction:
$2 \mathrm{HCl}+\mathrm{Na}_{2} \mathrm{SO}_{3}(\mathrm{aq}) \rightarrow \mathrm{S}(\mathrm{s})+\mathrm{H}_{2} \mathrm{SO}_{3}+2 \mathrm{NaCl}(\mathrm{aq})$
(1) Activation energy decreases
(2) Frequency of effective collision decreases
(3) Activation energy increases
(4) Frequency of effective collision increases
6. A student conducts an experiment to determine how the temperature of water affects the rate at which an antacid tablet dissolves in the water. The student has three antacid tablets of the same size and compositions.

Dissolving Data for Theses Antaeld Tablots

| Buker | Original Temparature at Water ('C) | Time for Tatlet to Disestive <br> (s) |
| :---: | :---: | :---: |
| 1 | 20. | 40. |
| 2 | 36. | 25 |
| 3 | 45. | 10. |

(A) Describe the effect of water temp on the rate of dissolving
(B) Explain in terms of collision theory, how water temperature influence the rate of dissolving

Table I (of the Reference Tables) tells us if particular reactions are exothermic or endothermic based on sign of the $\Delta \mathrm{H}$ value!
$\Delta \mathbf{H}$ : Enthalpy of a Reaction, how much $\qquad$ is gained or $\qquad$ during a reaction. Equal to the PE OF THE PRODUCTS - PE OF THE REACTANTS.

## RECALL; Two types of chemical reactions:

1. Exothermic: Reactions that $\qquad$ energy.

- $\Delta \mathrm{H}=$ negative value (-) energy released (on right)
- $A+B \rightarrow C+D+$ energy

Example: Sodium in water - lots of heat (and fire!) produced as product; heat felt on a test tube during a reaction
2. Endothermic: Reactions that $\qquad$ energy.

- $\Delta \mathrm{H}=$ positive value (+) energy absorbed (on left)
- $\mathrm{A}+\mathrm{B}+\mathrm{ENERGY} \rightarrow \mathrm{C}+\mathrm{D}$

Example: baking (need oven to supply heat)

## Determine if the following reactions are endo or exothermic:

1. $\mathrm{N}_{2}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NO}_{2}(\mathrm{~g})$ $\qquad$
2. $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$ $\qquad$

The * at the bottom of TABLE I tells you all you need to remember...
3. $2 \mathrm{NH}_{3}(\mathrm{~g}) \rightarrow \mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g})$ $\qquad$

## Practice:

| Reaction | $\Delta \mathbf{H}$ <br> $(k J)$ | Endothermic or <br> Exothermic |
| :--- | :--- | :--- |
| $\mathrm{CH}_{4}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ |  |  |
| $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ |  |  |
| $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{l})+3 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ |  |  |
| ${ }^{*} \mathrm{CO}(\mathrm{g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{CO}_{2}(\mathrm{~g})$ |  |  |
| ${ }^{*} 4 \mathrm{NO}(\mathrm{g}) \rightarrow 2 \mathrm{~N}_{2}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g})$ |  |  |

1. If you reverse a reaction, what happens to the magnitude of $\Delta H$ ? What happens to the sign?
2. If you double the concentration of the reactants and the products, what happens to the magnitude of $\Delta \mathrm{H}$ ? What happens to the sign?

Potential Energy Diagrams: Recall, we have talked about chemical bonds having stored energy (AKA potential energy). For that reason, chemists use diagrams called Potential Energy Diagrams to illustrate the potential (or stored) energy changes that occur during specific chemical reactions


## Vocabulary of a Potential Energy Diagram:

Potential Energy of the Reactants:

- The amount of energy that a system $\qquad$ with (the starting point in the graph.

Potential Energy of the Products:

- The amount of energy that the system $\qquad$ with.
$\square$ ACTIVATED COMPLEX: (The PEAK)
- $\qquad$ energy point of the reaction; this is where full of the reactants occurs. Remember, this must happen for the reaction to be successful.

ㅁ Activation Energy:

- Amount of ENERGY NEEDED TO GET A $\qquad$ or to FORM THE ACTIVATED COMPLEX of a reaction (you must get "over the hump" in order for a reaction to occur)
$\square$ Activation energy of the Forward reaction:
- Energy needed to get "over the hump" going forward (left to right).Activation Energy of the Reverse reaction:
- Energy needed to get "over the hump" going backward ( right to left).

ㅁ Heat of the Reaction

- ( $\left.\Delta \mathrm{H}=\mathrm{H}_{\mathrm{p}}-\mathrm{H}_{\mathrm{r}}\right)$

Skill 5: Distinguish between Endo and Exothermic potential energy diagrams
ENDOTHERMIC Potential Energy Diagrams $\rightarrow$ POSITIVE $\Delta H$

- Product side (to the R) always HIGHER than the reactant side (to the L) meaning that ENERGY is $\qquad$ _.


Reaction Coordinate

EXOTHERMIC Potential Energy Diagrams $\rightarrow$ NEGATIVE $\Delta H$

- Product side (to the R) always LOWER than the reactant side (to the L) meaning that ENERGY is $\qquad$ _.
- Most common types of rxns because less energy has to be put in to get the rxn started (LOWER activation energy)

$$
\begin{aligned}
& \text { Label the following: } \\
& \text { A = Potential Energy of the Reactants } \\
& B=\text { Potential Energy of the Products } \\
& C=\text { Potential Energy of the Activated } \\
& \text { Complex } \\
& D=\text { Activation energy of the Forward rxn } \\
& E=\text { Activation Energy of the Reverse rxn } \\
& \text { F=Heat of the Reaction }\left(\Delta H=H_{p}-H_{r}\right)
\end{aligned}
$$

## HOW EXACTLY DOES A CATALYST SHORTEN THE REACTION TIME NEEDED FOR A REACTION TO COMPLETE and how do we show it?

The $\qquad$ is lowered ..
OR
The ACTIVATION ENERGY is decreased ...

> OR

The REACTION $\qquad$ is shortened.

Draw the catalyzed reaction pathway on the diagram to the right $\rightarrow$



Determine the values of the following from the graph above:
$\qquad$
PE
Reactants:
Energy:
$A+B \quad \rightarrow$
$C+D$

PE
Products: $\qquad$
Activated complex: $\qquad$
Enthalpy of Rxn: $\qquad$
Activation Energy forward: $\qquad$
Activation Energy Reverse: $\qquad$
Endo or Exo?: $\qquad$

## Part 2

Using "Increase" or "Decrease" or "Remains the Same" to describe how adding a catalyst would affect the following:

PE Reactants: $\qquad$
PE Products: $\qquad$
Activated complex: $\qquad$
Enthalpy of Rxn: $\qquad$
Activation Energy forward: $\qquad$
Activation Energy Reverse: $\qquad$
$\square$ Entropy ( $\Delta \mathbf{S}$ ): Degree of $\qquad$ , CHAOS, DISORDER or "MESSINESS" in a system; nature tends to proceed to a state of GREATER entropy, or disorder.

- The MORE ORDER you have, the $\qquad$ ENTROPY in your system.
- The LESS ORDER you have, the $\qquad$ ENTROPY in your system.
$\qquad$ is the most significant factor in determining $\Delta \mathrm{S}$ :

$$
\text { Changing from }(\mathbf{s}) \rightarrow(\mathrm{l}) \rightarrow(\mathrm{aq}) \rightarrow(\mathrm{g})=\text { INCREASED ENTROPY }
$$

Draw particle diagrams to illustrate each of the following phases:

(s)



(g)
*Entropy $\qquad$ when a compound is broken down.
*Entropy $\qquad$ when a compound is created and bonds are formed.

NOTE: If there is no phase change, count up the \# molecules on each side

$$
\text { RULE: \# moles } \downarrow=\text { ENTROPY } \downarrow=-\Delta \mathrm{S}, \quad \text { AND } \quad \text { If } \# \text { moles } \boldsymbol{\uparrow}=\mathrm{ENTROPY} \boldsymbol{\uparrow}=+\Delta \mathrm{S})
$$

For the following determine if there is an increase, decrease, or no change in entropy

1. $2 \mathrm{KClO} 3(\mathrm{~s}) \rightarrow 2 \mathrm{KCl}(\mathrm{s})+3 \mathrm{O}_{2}(\mathrm{~g})$ $\qquad$
2. $\mathrm{H}_{2} \mathrm{O}_{(\mathrm{I})} \rightarrow \mathrm{H}_{2} \mathrm{O}_{(\mathrm{s})}$ $\qquad$
3. $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$ $\qquad$
4. $\mathrm{NaCl}(\mathrm{s}) \rightarrow \mathrm{Na}^{+}(\mathrm{aq})+\mathrm{Cl}^{-}(\mathrm{aq})$ $\qquad$
5. $\mathrm{KCl}_{(\mathrm{s})} \rightarrow \mathrm{KCl}_{(\mathrm{I})}$ $\qquad$
6. $\mathrm{CO}_{2}(\mathrm{~s}) \rightarrow \mathrm{CO}_{2}(\mathrm{~g})$ $\qquad$
$\square$ Equilibrium occurs WHEN THE $\qquad$ OF THE FORWARD REACTION EQUALS THE OF THE REVERSE REACTION in a closed system.
$\square$ When equilibrium is reached, IT DOES NOT MEAN that the reactants and products are of equal QUANTITIES. So...

- DOUBLE ARROWS () instead of a single arrow. This allows us to illustrate that the reactions are proceeding in both directions (forward and reverse).
- Equilibrium is $\qquad$ which means that it is constantly CHANGING or FLUCTUATING
- Equilibrium means that reactant and product are CONSTANT. *Equilibrium does NOT mean that reactant and product concentrations are equal.*


TIME FOR BAILING BEAKERS......WHO WILL BE VICTORIOUS?

## TYPES OF EQUILIBRIUM (all occur in

$\qquad$ ) *IT'S ALL ABOUT THE EQUAL RATES*

1. Physical Equilibrium: Equilibrium that involves physical changes

- Phase Equilibrium - occurs during a PHASE CHANGE

RATE of MELTING = RATE of FREEZING (sealed container @ $0^{\circ} \mathrm{C}$ )
RATE of EVAPORATION = RATE of CONDENSATION (sealed container @ $100^{\circ} \mathrm{C}$ )

- Solution Equilibrium - occurs at a solution's $\qquad$ POINT RATE of DISSOLVING $=$ RATE of CRYSTALLIZATION example: $\mathrm{NaCl}_{(\mathrm{s})} \leftarrow \rightarrow \mathrm{NaCl}_{(\mathrm{aq})}$

2. Chemical Equilibrium:

- $\qquad$ of the FORWARD RXN = $\qquad$ of the REVERSE RXN
OR
- $\qquad$ of BREAKING BONDS = $\qquad$ of FORMING BONDS

1. Which statement describes a chemical reaction at equilibrium?
A) The products are completely consumed in the reaction.
B) The reactants are completely consumed in the reaction.
C) The reaction rates of the products and reactants are equal.
D) The reaction rates of the products and reactants are constant.
2. Which two processes are at equilibrium in a saturated sugar solution?
a. evaporation and condensation
c. decomposition and synthesis
b. dissolving and crystallization
d. ionization and recombination

Le Chatelier's principle explains HOW A SYSTEM AT EQUILIBRIUM WILL RESPOND TO $\qquad$ .

- STRESS = Any change in TEMPERATURE, CONCENTRATION, or PRESSURE put upon an system at equilibrium
- When a STRESS is added to a system at equilibrium, the system will $\qquad$ in order to relieve that stress and reach a new equilibrium.
- SHIFT = an increase in the $\qquad$ of EITHER the forward OR the reverse rxn
- SHIFT TO RIGHT (TOWARD PRODUCTS):
- Rate of FORWARD reaction INCREASES $(\rightarrow)$
- Reactants $\rightarrow_{\leftarrow}$ Products *Favors PRODUCTS
- SHIFT TO LEFT (TOWARD REACTANTS):
- Rate of REVERSE reaction INCREASES ( $\leftarrow)$
- Reactants $\leftarrow_{\rightarrow}$ Products *Favors REACTANTS


## Skill 7a: Applying Le Chatelier with Types of Stressors

Initial Stress: Concentration
$\square$ When the concentration of a reactant or product is INCREASED, the

ADD, AWAY reaction will SHIFT $\qquad$ from the increase (use up the excess.

Example 1: $\quad 4 \mathrm{NH}_{3}(\mathrm{~g})+5 \mathrm{O}_{2}(\mathrm{~g}) \leftarrow \rightarrow 4 \mathrm{NO}(\mathrm{g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})+$ HEAT

1. If we add $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$, the system would shift to the $\qquad$ and the $\left[\mathrm{NH}_{3}\right]$ would $\qquad$ .
2. If we add $\mathrm{O}_{2}(\mathrm{~g})$, the system would shift to the $\qquad$ and the [NO] would $\qquad$ -.
3. If we add $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$, the system would shift to the $\qquad$ and the [ NO ] would $\qquad$ .
$\square$ When the concentration of a reactant or product is DECREASED: The reaction will SHIFT $\qquad$ the side that has experienced the

## Take, Towards

 decrease in concentration (replaces what was taken)Example 2: $\quad 4 \mathrm{NH}_{3}(\mathrm{~g})+5 \mathrm{O}_{2}(\mathrm{~g}) \leftarrow \rightarrow 4 \mathrm{NO}(\mathrm{g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})+\mathrm{HEAT}$

1. If we remove oxygen, the system will shift to the $\qquad$ and the $\left[\mathrm{NH}_{3}\right]$ will $\qquad$ .
2. If we remove water, the system will shift to the $\qquad$ and the [NO] will $\qquad$ .
3. If we remove ammonia, which concentration(s) will decrease? $\qquad$
4. If we remove $\mathrm{NO}(\mathrm{g})$, which concentration(s) would increase? $\qquad$

Initial Stress: Temperature (involves increasing or decreasing the "HEAT" component of a reaction)
$\square$ When temperature (or HEAT) is decreased: The reaction will SHIFT $\qquad$ the rxn side containing "HEAT" (in the EXOTHERMIC direction)

## Take, Towards

$$
\mathrm{A}+\mathrm{B} \leftarrow \mathrm{C}+\mathrm{D}+\text { HEAT }
$$

Example 1: $\quad 4 \mathrm{NH}_{3}(\mathrm{~g})+5 \mathrm{O}_{2}(\mathrm{~g}) \leftrightarrow 4 \mathrm{NO}(\mathrm{g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})+$ HEAT

1. If we remove heat, which concentration(s) will decrease? $\qquad$
2. If we remove heat, which concentration(s) will increase? $\qquad$
$\square$ When temperature (or HEAT) is increased: When temperature (or HEAT) is increased: The reaction will SHIFT $\qquad$ from the rxn side containing "HEAT" (in the ENDOTHERMIC direction)
$A+B+$ energy $\leftarrow \rightarrow C+D$

$$
\text { Example 2: } \quad 4 \mathrm{NH}_{3}(\mathrm{~g})+5 \mathrm{O}_{2}(\mathrm{~g}) \leftrightarrow 4 \mathrm{NO}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g})+\mathrm{HEAT}
$$

3. If we added heat, which concentration(s) will decrease? $\qquad$
4. If we added heat, which concentration(s) will increase? $\qquad$

Initial Stress: Pressure.....EFFECTS GASES ONLY!!!!
$\square$ INCREASE PRESSURE: rxn shifts to side with $\qquad$ \# GAS MOLECULES (or least \# moles of gas)
$\square$ DECREASE PRESSURE: rxn shifts to side with GREATEST \# GAS MOLECULES (or greatest \# moles of gas)

NOTE: If the rxn contains NO GAS MOLECULES or if the rxn has the SAME \# GAS MOLECULES on each side, there is NO EFFECT and NO SHIFT results from an increase or decrease in pressure

## Example 1: $\quad \mathrm{CO}_{2}(\mathrm{~g}) \leftrightarrow \rightarrow \mathrm{CO}_{2}(\mathrm{aq})$

4. If we increase the pressure, the concentrations of which species will increase? $\qquad$

## Example 2: <br> $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \longleftrightarrow \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$

1. If we increase the pressure, in which direction will the equilibrium shift? (Count Moles) $\qquad$
2. If we increase the pressure, the concentration of which species will increase initially? $\qquad$
3. If we decrease the pressure, the concentration of which species will decrease initially? $\qquad$
4. If we decrease the pressure, the concentration of which species will increase initially? $\qquad$

## Chemical System: $\quad 2 \mathrm{SO}_{3}+$ heat $\longleftrightarrow 2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})$

a) Stress: Increasing Temperature:
a. Reaction will shift $\qquad$ Concentration of $\mathrm{SO}_{3}$ $\qquad$
b. Rate of forward reaction $\qquad$ Concentration of $\mathrm{SO}_{2}$ $\qquad$
c. Rate of reverse reaction $\qquad$ Concentration of $\mathrm{O}_{2}$ $\qquad$
b) Stress: Increasing Pressure:
a. Reaction will shift $\qquad$ Concentration of $\mathrm{SO}_{3}$ $\qquad$
b. Rate of forward reaction $\qquad$
c. Rate of reverse reaction $\qquad$
Concentration of $\mathrm{SO}_{2}$ $\qquad$ Concentration of $\mathrm{O}_{2}$ $\qquad$
b) Stress: Decreasing Pressure:
a. Reaction will shift $\qquad$ Concentration of $\mathrm{SO}_{3}$ $\qquad$
b. Rate of forward reaction $\qquad$
c. Rate of reverse reaction $\qquad$
Concentration of $\mathrm{SO}_{2}$ $\qquad$
Concentration of $\mathrm{O}_{2}$ $\qquad$
Chemical System: $\quad 3 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{N}_{2} \leftarrow \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})+$ heat
a) Stress: Increasing Temperature:
a. Reaction will shift $\qquad$
b. Rate of forward reaction $\qquad$
Concentration of $\mathrm{H}_{2}$ $\qquad$
c. Rate of reverse reaction $\qquad$
Concentration of $\mathrm{N}_{2}$ $\qquad$
Concentration of $\mathrm{NH}_{3}$ $\qquad$
b) Stress: Decreasing temperature
a. Reaction will shift $\qquad$ Concentration of $\mathrm{H}_{2}$ $\qquad$
b. Rate of forward reaction $\qquad$ Concentration of $\mathrm{N}_{2}$ $\qquad$
c. Rate of reverse reaction $\qquad$ Concentration of $\mathrm{NH}_{3}$ $\qquad$
c) Stress: Adding $\mathrm{NH}_{3}$ :
a. Reaction will shift $\qquad$ Concentration of $\mathrm{H}_{2}$ $\qquad$
b. Rate of forward reaction $\qquad$ Concentration of $\mathrm{H}_{2}$ $\qquad$
c. Rate of reverse reaction $\qquad$ Concentration of $\mathrm{NH}_{3}$ $\qquad$
Chemical System:
$\mathrm{BaSO}_{4}(\mathrm{~s}) \longleftrightarrow \mathrm{Ba}^{2+}(\mathrm{aq})+\mathrm{SO}_{4}{ }^{2-}(\mathrm{aq})$
a) Stress: Adding $\mathrm{Ba}\left(\mathrm{NO}_{3}\right)_{2}(\mathrm{~s})$
a. Reaction will shift $\qquad$
b. Rate of forward reaction $\qquad$
Concentration of $\mathrm{Ba}^{2+}$
Concentration of $\mathrm{SO}_{4}{ }^{2-}$ $\qquad$
c. Rate of reverse reaction $\qquad$
b) Stress: Removing $\mathrm{Ba}\left(\mathrm{NO}_{3}\right)_{2}$ (s)
a. Reaction will shift $\qquad$
b. Rate of forward reaction $\qquad$
c. Rate of reverse reaction $\qquad$
c) Stress: Removing $\mathrm{SO}_{4}{ }^{2-}$
a. Reaction will shift $\qquad$ Concentration of $\mathrm{Ba}^{2+}$ $\qquad$
b. Rate of forward reaction $\qquad$
c. Rate of reverse reaction $\qquad$

Concentration of $\mathrm{SO}_{4}{ }^{2-}$ $\qquad$

Match the change to the equilibrium system below with the letter of the appropriate response. Each letter can be used once, more than once, or not at all.

$$
2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \leftrightarrow 2 \mathrm{SO}_{3}(\mathrm{~g})
$$

$\qquad$

1) $\mathrm{O}_{2}$ is added to the reaction
a) The equilibrium shifts to the right
2) $\mathrm{SO}_{3}$ is removed from the reaction
b) The equilibrium shifts to the left
3) $\mathrm{SO}_{3}$ is added to the reaction
c) there is no change in the equilibrium
$\qquad$
$\qquad$ 4) The pressure is increased
4) $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})+53 \mathrm{~kJ} \longleftrightarrow \rightarrow \quad 2 \mathrm{HI}(\mathrm{g})$

Using directional arrows, describe what would happens if:
a. Increase Temperature: Shifts to the $\qquad$ , $\qquad$ [HI], $\qquad$ $\left[\mathrm{H}_{2}\right]$, $\qquad$ [ ${ }_{2}$ ]
b. Increase $\left[\mathrm{H}_{2}(\mathrm{~g})\right]$ : Shifts to the $\qquad$
$\qquad$ [HI], $\qquad$ [ 12 ]
c. Increase $[\mathrm{HI}(\mathrm{g})]$ : Shifts to the $\qquad$ , $\qquad$ [ $\mathrm{H}_{2}$ ], $\qquad$ $[12]$
d. Decrease Pressure:
6) $\mathrm{PCl}_{5}(\mathrm{~g})+$ heat $\leftrightarrow \rightarrow \quad \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})$

What happens if:
a. Adding $\mathrm{Cl}_{2}$ : Shifts to the $\qquad$ , $\qquad$ [ $\mathrm{PCl}_{5}$ ], $\qquad$ $\left[\mathrm{PCl}_{3}\right]$
b. Increasing Pressure: Shifts to the $\qquad$
$\qquad$ [ $\mathrm{PCl}_{5}$ ], $\qquad$ [ $\mathrm{PCl}_{3}$ ], $\qquad$ $\left[\mathrm{Cl}_{2}\right]$
c. Lowering Temperature: Shifts to the $\qquad$ , $\qquad$ [ $\mathrm{PCl}_{5}$ ], $\qquad$ [ $\mathrm{PCl}_{3}$ ], $\qquad$ $\left[\mathrm{Cl}_{2}\right]$
d. Removing $\mathrm{PCl}_{3}$ : Shifts to the $\qquad$ [ $\mathrm{PCl}_{5}$ ], $\qquad$ $\left[\mathrm{PCl}_{3}\right]$
7) For the following reaction, what will occur if pressure is increased? Why?

$$
2 \mathrm{NO}_{2}(\mathrm{~g}) \leftarrow \rightarrow \mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})
$$

8) Given the reaction at equilibrium : $\mathrm{A}(\mathrm{g})+\mathrm{B}_{(\mathrm{g})} \leftrightarrow \mathrm{C}_{(\mathrm{g})}+\mathrm{D}_{(\mathrm{g})}$ The addition of a catalyst will:
a) shift equilibrium to the right
b) shift equilibrium to the left
c) increase the rate of the forward and reverse reactions
d) have no effect on the rate of the forward and reverse reactions
9) Consider the equation of the following reaction at equilibrium:

$$
X+Y \leftarrow \rightarrow 2 Z+\text { heat }
$$

The concentration of the product can be increased by
a) adding a catalys $\dagger$
b) adding more heat to the system
c) increasing the concentration of $Y$
d) decreasing the concentration of $Z$
10) Consider the following equation: $\mathrm{H}_{2(\mathrm{~g})}+\mathrm{Cl}_{2(\mathrm{~g})} \leftarrow \rightarrow 2 \mathrm{HCl}_{(\mathrm{g})}$

Which change will result in an increase in the concentration of chloride gas?
a) decreasing the pressure on the system
b) decreasing the concentration of HCl
c) increasing the concentration of $\mathrm{H}_{2}$
d) increasing the concentration of HCl
11) Consider the following equation:
$\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \leftarrow \mathrm{gNO}_{(\mathrm{g})}$
As the concentration of $\mathrm{N}_{2(\mathrm{~g})}$ increases, the concentration of $\mathrm{O}_{2(\mathrm{~g})}$ will
a) decrease
b) increase
c) remain the same
d) vary directly
12) Referencing the equation in question 11 , why does increasing the pressure increase the rate of the forward reaction?

Fill in the blanks using the word bank. Only one word in the bank is used TWICE.

| activated complex | Le Chatelier's Principle | entropy |
| :--- | :--- | :--- |
| heterogeneous reaction | double arrow | reaction mechanism |
| activation energy | potential energy | exothermic reaction |
| homogeneous reaction | endothermic reaction | shift |
| catalyst | rate | heat of reaction |
| kinetics | enthalpy | stress |
| chemical equilibrium | rate-determining step |  |

The branch of chemistry concerned with the rates of chemical changes is called
$\qquad$ . A chemical change in which all the reactants are in the same phase is called $a(n)$ $\qquad$ . One in which the reactants are in different phases is called $a(n)$ $\qquad$ . A substance that speeds up a chemical change without being permanently altered or affecting the nature of the reaction is called $a(n)$ $\qquad$ .

The series of steps by which reacting particles rearrange themselves to form products is called the $\qquad$ The slowest step in such a series is the
$\qquad$ . A short-lived, high-energy arrangement of particles that is formed when reacting particles collide at the proper angle with the proper amount of energy is $a(n)$ $\qquad$ . The minimum amount of energy needed to form this arrangement is called the $\qquad$ . Because this energy is stored inside the particles, it is an example of $\qquad$ . The reactants and the products of any reaction have different amounts of this kind of stored energy. The difference between these two amounts of energy is the $\qquad$ .

The heat content of a substance is called its $\qquad$ . The change in this quantity that occurs during a chemical reaction is called the $\qquad$ $\Delta H$. The sign of the quantity $\Delta H$ is positive in the case of $a(n)$ $\qquad$ . It is negative in the case of $a(n)$ $\qquad$ .

When forward and backward reactions occur at the same $\qquad$ , a state of
$\qquad$ exists. A(n) $\qquad$ is used in an equation to
symbolize this state.
When conditions such as temperature are changed, a chemical reaction is said to be placed under $\mathrm{a}(\mathrm{n})$ $\qquad$ . Under such changing conditions, equilibrium can undergo $a(n)$ ___ in direction that tends to counteract the imposed changes. This generalization is known as $\qquad$ . The measure of the randomness of a system is its $\qquad$ .

1) If you were given the $\Delta \mathrm{H}$ value of a reaction, you could determine whether the reaction was exothermic or endothermic. Explain how you could do so.
$\qquad$
$\qquad$
2) In an exothermic reaction, Hproducts will always be (larger/smaller) than Hreactants

## Enthalpy and Table I

Look in your Reference Tables (Table I) and state whether the following reactions are exothermic or endothermic.
3. Sodium hydroxide dissociating into a positive sodium ion and a negative hydroxide ion

## Exo or Endo

1. Methane $\left(\mathrm{CH}_{4}\right)$ combining with oxygen to produce carbon dioxide and water
2. Potassium nitrate dissociating into a positive potassium ion and a negative nitrate ion
3. Carbon monoxide combining with oxygen to form carbon dioxide
4. **A positive lithium ion combining with a negative bromine ion to form lithium bromide

| Process | $\Delta H$ | Exo or Endo | Entropy change |
| :---: | :---: | :---: | :---: |
| 1. $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})}$ |  |  |  |
| 2. $\mathrm{C}_{(\mathrm{s})}+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{CO}_{2}(\mathrm{~g})$ |  |  |  |
| 3. $\mathrm{CO}_{2}(\mathrm{~g}) \rightarrow \mathrm{C}_{(\mathrm{s})}+\mathrm{O}_{2}(\mathrm{~g})$ |  |  |  |
| 4. ${ }^{\text {C }}$ (s) $+\mathrm{H}_{2}(\mathrm{~g}) \rightarrow \mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g})$ |  |  |  |
| 5. $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ |  |  |  |
| 6. $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}(\mathrm{~s})+6 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 6 \mathrm{CO}_{2}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ |  |  |  |
| 7. $\mathrm{Br}-(\mathrm{aq})+\mathrm{Li+}(\mathrm{aq}) \rightarrow \mathrm{LiBr}$ (s) |  |  |  |
| 8. $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{HI}(\mathrm{g})$ |  |  |  |
| 9. $\mathrm{NaOH}_{(\mathrm{s})} \rightarrow \mathrm{Na}^{+}(\mathrm{aq})+\mathrm{OH}_{-}(\mathrm{aq})$ |  |  |  |
| 10. $2 \mathrm{CO}_{2}(\mathrm{~g}) \rightarrow \mathrm{O}_{2}(\mathrm{~g})+2 \mathrm{CO}(\mathrm{g})$ |  |  |  |

