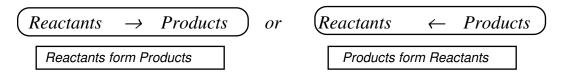
<u>Chemistry 12</u> <u>Unit 2- Equilibrium</u> <u>Notes</u>

It's important to know that many chemical reactions are *reversible*. That is:



For example, under certain conditions, one mole of the colourless gas N_2O_4 will *decompose* to form two moles of **brown NO₂ gas**:

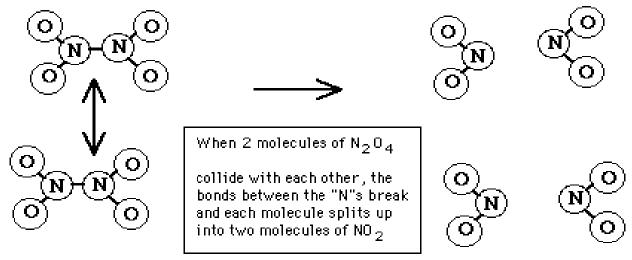


Under other conditions, you can take 2 moles of brown NO_2 gas and change it into one mole of N_2O_4 gas:



In other words, this reaction, as written may go *forward* or in *reverse*, depending on the conditions.

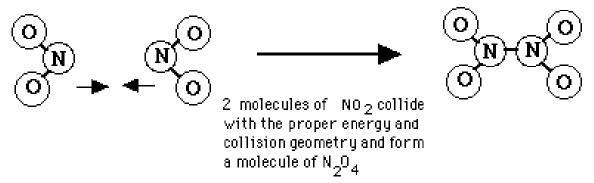
If we were to put some N_2O_4 in a flask, the N_2O_4 molecules would collide with each other and some of them would break apart to form NO_2 .



This process is indicated by the *forward reaction*:

$$N_2O_4 \rightarrow 2NO_2$$

Once this has happened for awhile, there is a build up of NO_2 molecules in the same flask Once in awhile, two NO_2 molecules will collide with *each other* and **join** to form a molecule of N_2O_4 !



This process, as you might have guessed is indicated by the *reverse reaction*:

 $N_2O_4 \leftarrow 2 NO_2$

Two things you'll have to realize is that as long as there is N_2O_4 present, the <u>forward</u> reaction will keep on happening and as long as there is NO_2 present, the <u>reverse</u> reaction will keep on happening!

Also, you must keep in mind that all these molecules are mixed in the same container!

At one particular time a molecule of N_2O_4 might be breaking up, and *at the same time* two molecules of NO_2 might be joining to form another molecule of N_2O_4 ! So here's an important thing to understand:

In any reversible reaction, the forward reaction and the reverse reaction are going on **at the same time**!

This is sometimes shown with a **double arrow**:

$$N_2O_4 \rightleftharpoons 2 NO_2$$

 \uparrow
The double arrow means that
both the forward and reverse
reaction are happening at the
same time.

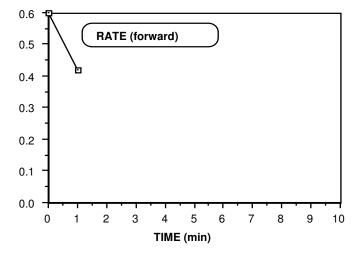
Just a little comment here. The word "*happening*" has a similar meaning to the word " *dynamic* ". Just remember that!

Now what we're going to do is look at how the *rate* of the forward reaction changes if we put some pure N_2O_4 in a flask:

If we put some pure N_2O_4 in a flask (No NO₂ yet!), there will be a high *concentration* of N_2O_4 . That is, there will be *lots* of N_2O_4 molecules to collide with each other. So at the beginning of our little experiment, (which we will call " time " 0 ") the *rate* of the *forward* reaction is quite *fast*.

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N_2O_4 \rightarrow 2 NO_2
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So if we were to make a *graph* of the *rate of the forward reaction* vs. *time*, the graph might start out something like this:



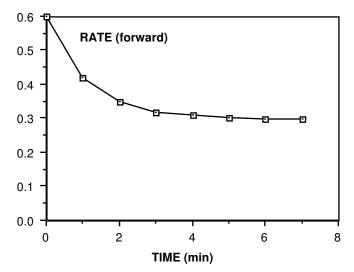
OK, now you might ask: "Why does the rate of the forward reaction go down ?

Well, if you recall Unit 1, as the forward reaction proceeds:

$$N_2O_4 \rightarrow 2NO_2$$

the N_2O_4 is used up and so it's *concentration goes down*. Also, you must remember that *if the concentration of a reactant goes down, there is less chances of collisions and the rate of the reaction decreases*.

As the reaction continues, the *slower rate* will use up N_2O_4 more slowly, so the $[N_2O_4]$ will not decrease so quickly and therefore the rate will not decrease quite as quickly. (Read the last sentence over a couple of times and make sure it makes sense to you!) For those "graph wise" people, you will probably guess that this means that the *slope of the line* on the graph gets more gradual. The rest of the graph might look something like this:

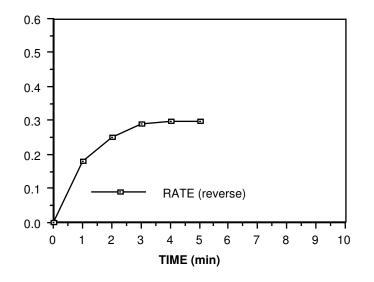


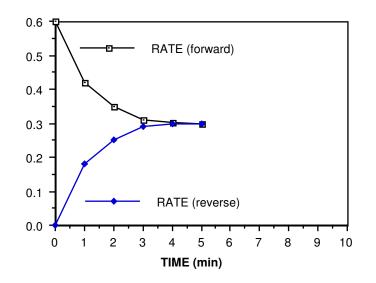
NOW, it's time we consider the rate of the reverse reaction.

As you might recall, when we have a container full of pure N_2O_4 , initially there is **no** NO_2 in the container. Since there is **no** NO_2 , there are no NO_2 molecules to collide with each other, and the **rate of the reverse reaction is zero**.

But of course, as time goes on, NO₂ is formed from the *forward reaction* ($N_2O4 \rightarrow 2NO_2$) so in a short time, some NO₂ molecules can start colliding and the *reverse reaction* will begin:

As MORE NO₂ is formed by the forward reaction, *the rate of the reverse reaction gradually increases.* Now, for you "graph buffs", the graph of the *Rate of the reverse reaction* vs. *Time* might look like this:





OK. Now, lets look at the graph for the forward rate *and* the reverse rate together:

NOW, focus your attention on the graph at " Time = 4 minutes " . You will notice that at this point:

the rate of the forward reaction = the rate of the reverse reaction

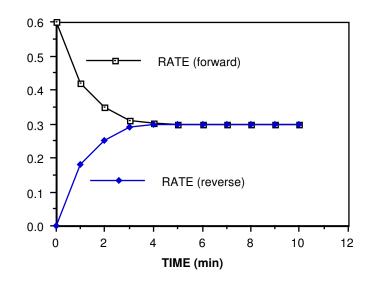
At this point, NO₂ is being used up *at the same rate* that it is being formed:

$$N_2O_4 \rightleftharpoons 2NO_2$$

Because this is so, you should be able to convince yourself that the [NO₂] is no longer changing!

Because the reverse rate is equal to the forward rate, N_2O_4 is being formed *at the same rate* it is being used up. So, also *the* [N_2O_4] *is no longer changing either!* Can you predict what will happen to the graph after 4.0 minutes?

YOU GUESSED IT! The rates of the forward reaction and the reverse reaction, no longer change because the $[N_2O_4]$ is constant and the $[NO_2]$ is also constant. The graph will look like this:



The situation happening from 4.0 minutes on in this graph has a special name and a special significance. At this point, the system (meaning the container, the N_2O_4 and the NO_2) is said to be *at <u>equilibrium</u>*. To describe it even more precisely, we can say that *we have reached a state of dynamic equilibrium*.

Here are some things that you must *understand* about *dynamic equilibrium*:

- 1. The reaction *has <u>not</u> stopped!*
- 2. The forward and the reverse reaction *continue* to take place, but their rates are *equal* so there are no changes in concentrations of reactants or products. (The forward and reverse reactions are said to be " balanced ") eg. for the reaction:

$$N_2O_4 \rightleftharpoons 2NO_2$$

for each N₂O₄ molecule that breaks up to form two NO₂ molecules, two *other* NO₂ *microscopic level*, so we don't see individual molecules reacting.

- 3. As far as we can see from the "outside", there *appears to be* nothing happening. All *observable* properties are constant. These include the concentrations of all reactants and products, the total pressure, colour, temperature etc.
- 4. If no changes were made in conditions and nothing is added or taken away, a system at equilibrium would remain that way forever, the forward and reverse reactions "ticking away", but balanced so that no observable changes happen.

Here are a couple of other things to consider before we summarize everything:

- 1. Changing the temperature can alter the rates of the reactions at equilibrium. This could "throw off" the balance. So, *for a system at equilibrium, the <u>temperature</u> must remain constant and uniform throughout the system.*
- 2. Letting material into or out of the system will affect rates so *a system at equilibrium is a <u>closed system</u>.*
- 3. Again, consider the equilibrium reaction: $N_2O_4 \rightleftharpoons 2NO_2$ In the example that we did to construct the graphs, we had started with *pure* N_2O_4 and no NO₂. The forward reaction rate was high at the start, but the reverse reaction rate eventually "caught up", the rates became equal and *equilibrium* was established. Can you guess what would happen if we had started with *pure* NO_2 instead (no N₂O₄)? The reverse rate would start out high and the forward rate, zero. In time, the forward rate would " catch up ". When the rates became *equal*, again *equilibrium* would be established.

We can summarize all this by saying that *the equilibrium can be approached from the* <u>*left*</u> (*starting with reactants*) *or from the* <u>*right*</u> (*starting with products*)

Just a little term before we summarize: The word *macroscopic* means *large scale or visible or observable*. (The opposite is *microscopic*, which means too small to see eg. molecular level). Some *macroscopic* properties are total pressure, colour, concentrations, temperature, density etc. Alright, let's summarize:

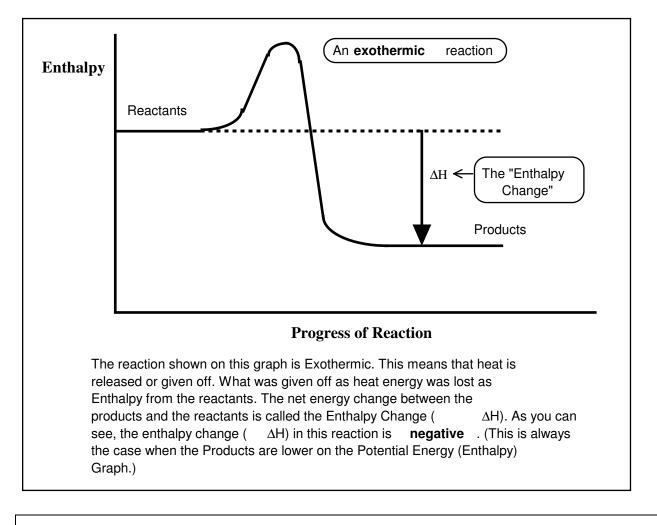
Characteristics of a System at Dynamic Equilibrium

- 1. The *rate* of the *forward* reaction = The *rate* of the *reverse* reaction
- 2. *Microscopic* processes (the forward and reverse reaction) continue in a *balance* which yields *no macroscopic changes*. (so nothing *appears* to be happening.)
- 3. The system is *closed* and the *temperature* is *constant* and *uniform* throughout.
- 4. The equilibrium can be approached from the *left* (starting with *reactants*) or from the *right* (starting with *products*).

<u>Enthalpy</u>

Enthalpy is "*The heat content of a system*. " Another way to think of *enthalpy* is as "*Chemical Potential Energy*".

Any change in the Potential Energy of a system means the same thing as the "Enthalpy Change ". The symbol for Enthalpy is " H ". Therefore the "change in Enthalpy" of a chemical reaction is called " ΔH ". In Chemistry 12, a Potential Energy Diagram is the same thing as an "Enthalpy Diagram



In an <u>Exothermic</u> Reaction (ΔH is negative), the <u>Enthalpy</u> is decreasing.

In an <u>Endothermic</u> Reaction (*AH* is positive), the <u>Enthalpy</u> is increasing.

If the "Heat Term" is written right in the equation. (a "thermochemical equation".)

If the heat term is on the *left* side, it means heat is being *used up* and it's *endothermic*.

If the heat term is on the *right* side, heat is being *released* and it's *exothermic*.

Look at the following examples:

1.	$A + B \rightleftharpoons C + D$	$\Delta H = -24 \text{ kJ}$	is <i>exothermic</i> so <u>enthalpy is <i>decreasing</i></u> .
2.	$X + Y \rightleftharpoons Z \qquad \Delta H =$	87 kJ	is endothermic so enthalpy is increasing.
3.	$E + D \rightleftharpoons F + 45 \text{ kJ}$		is <i>exothermic</i> so <u>enthalpy is <i>decreasing</i></u> .
4.	G + J + 36 kJ ₹ L +	М	is endothermic so enthalpy is increasing.

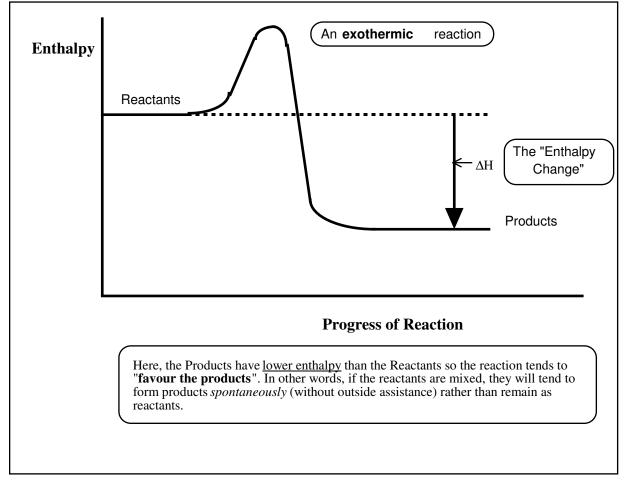
Systems will tend toward a state of *lower potential energy* if nothing else is acting upon them.

In Chemistry, we are interested in is chemical potential energy, otherwise known as enthalpy!

Chemical systems will tend toward a state of <u>minimum enthalpy</u> if sufficient activation energy is available and <u>no other factors are considered</u>.

Another way of stating this might be:

A chemical reaction will favour the side (reactants or products) with <u>minimum enthalpy</u> if no other factors are considered.



Thus for an *exothermic reaction*, if no other factors are considered:

The *products will be favoured* because the products have *minimum enthalpy*. In other words, there is a natural tendency here for reactants to **spontaneously** form products.

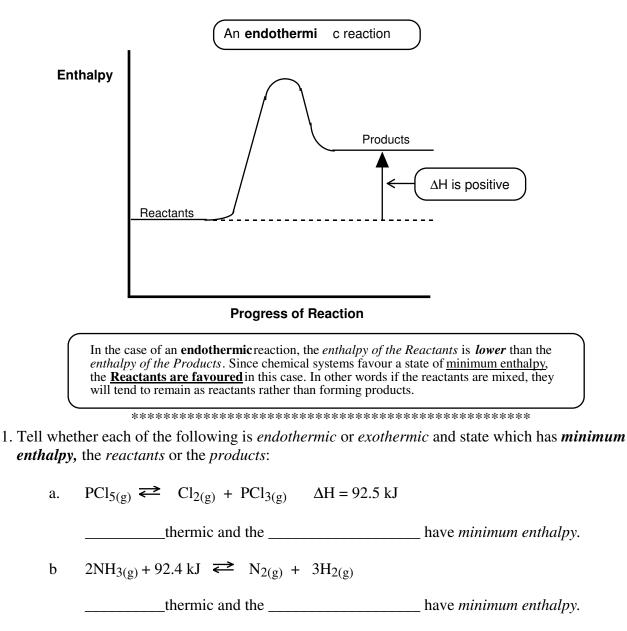
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In an <i>endothermic reaction</i> , the _	have minimum
· _	

enthalpy, so the ______ will be favoured. In other words, if

the reactants are mixed they will (tend to remain as reactants / spontaneously form products)

Let's look at a diagram for an *endothermic reaction*:



 $c \qquad CO_{(g)} + 3H_{2(g)} \rightleftarrows CH_{4(g)} + H_2O_{(g)} + 49.3 \text{ kJ}$

______thermic and the ______ have *minimum enthalpy*.

d. $Cl_{2(g)} \rightleftharpoons Cl_{2(aq)} \Delta H = -25 \text{ kJ}$

______thermic and the _______have *minimum enthalpy*.

Chemistry 12

2.	When no other factors are considered, a reaction will move in such a way (left or right) in			
	order to achieve a state of	enthalpy.		
3	Given the equation: $2NH_{3(g)} + 92.4 \text{ kJ} \rightleftharpoons N_{2(g)} + 3H_{2(g)}$			
	If only the <i>enthalpy</i> is considered, the (reactant / products)			
4.	Given the equation: $Cl_{2(g)} \rightleftharpoons Cl_{2(aq)} \Delta H = -25 \text{ kJ}$			
	If only the <i>enthalpy</i> is considered, the (reactant / products)			
5	If the reaction : $CO_{(g)} + 3H_{2(g)} \rightleftharpoons CH_{4(g)} + H_2O_{(g)} + 49.3 \text{ kJ}$			
	was proceeding to the <i>right</i> , the enthalpy would be	ing. Is this a		
	favourable change?			
6.	If the reaction: $PCl_{5(g)} \rightleftharpoons Cl_{2(g)} + PCl_{3(g)} \Delta H = 92.5 \text{ kJ}$			
	was proceeding to the <i>right</i> , the enthalpy would be	ing. Is this a		
	<i>favourable</i> change?			
7.	If the reaction: $Cl_{2(g)} \rightleftharpoons Cl_{2(aq)} \Delta H = -25 \text{ kJ}$			
	was proceeding to the <i>right</i> , the enthalpy would be	ing. Is this a		
	favourable change?			
8	If the reaction: $2NH_{3(g)} + 92.4 \text{ kJ} \iff N_{2(g)} + 3H_{2(g)}$			
	was proceeding to the <i>right</i> , the enthalpy would be	ing. Is this a		
	favourable change?			

As you can see by looking at the exercises above, there are two ways of looking at what happens to the *enthalpy*:

If the reaction is <u>exothermic</u>, the <u>products</u> have minimum enthalpy and the formation of products (move toward the <u>right</u>) is favourable.

If the reaction is <u>endothermic</u>, the <u>reactants</u> have minimum enthalpy and the formation of products (move toward the right) is <u>un</u>favourable. In this case the formation of <u>reactants</u> (move toward the <u>left</u>) <u>is favourable</u>.

Now, consider the simple *melting* of water:

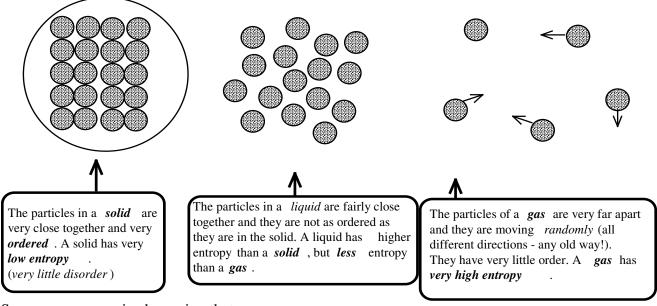
 $H_2O_{(s)}$ + heat \rightleftharpoons $H_2O_{(l)}$ (the subscript (s) stands for solid) (the subscript (l) stands for liquid)

If we were to look at only the enthalpy in this process, you can see that the reactant $(H_2O_{(s)})$ would have minimum enthalpy and would be favoured. So *all of the water in the universe should exist only as a <u>solid</u>! (It would not be favourable for water to exist as a liquid!) We would all be frozen solid!!!!!*

The answer to this problem lies in looking at *another factor* that governs equilibrium. That factor is called *entropy* (or *randomness* or *disorder*)

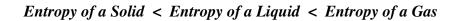
<u>Entropy</u>

Entropy simply means <u>disorder</u>, or lack of order.



In Grade 8, you probably learned about the arrangement of molecules in solids, liquids and gases.

So we can summarize by saying that:



We can look at a chemical equation with subscripts showing the phases and tell which has *maximum entropy*, the *reactants* or the *products*.

In other words, they can look at an equation and tell whether *entropy* is *increasing* or *decreasing* as the reaction *proceeds to the right*.

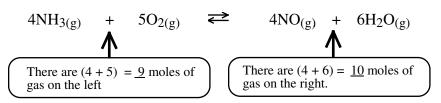
In the following examples, the *entropy is increasing* (or the *products* have *greater entropy*):

1. There is a *gas* (or gases) on the *right*, when there are *no gases* on the *left* of the equation:

 $CaCO_{3(s)} + 2 HCl_{(aq)} \rightleftharpoons CaCl_{2(aq)} + CO_{2(g)} + H_2O_{(l)}$

a gas is formed on the right.

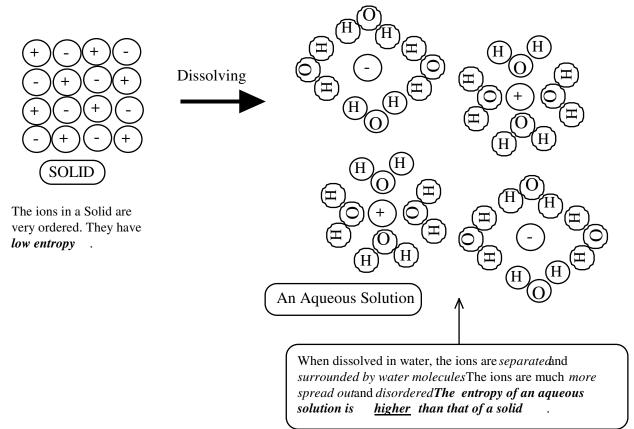
2. When there are *gases on both sides*, the *products* have *greater entropy* when there are *more moles of gas on the right* (add up coefficients of gases on left and right.):



Another way to look at the last example is to say that:

" The side with the greater number of moles of gas has the greatest entropy. "

3. When a *solid dissolves in water*, the *products* (the aqueous solution of ions) have *greater entropy*. This makes sense because:

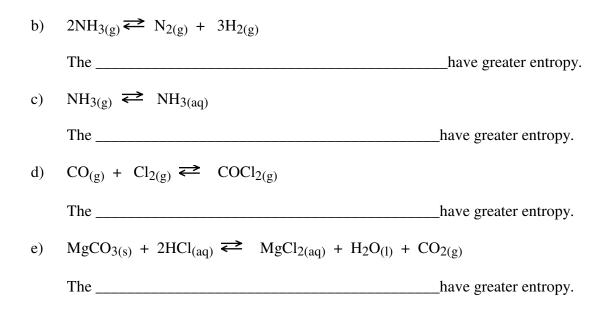


So in order of lowest to highest **entropy**:

Solids < Liquids < Aqueous solutions < Gases < More moles of Gas

Here are few exercises for you:

- 9. For each of the following, decide whether the *reactants* or the *products* have *greater entropy*:
 - a) $I_{2(s)} \rightleftharpoons I_{2(aq)}$ The ______ have greater entropy.



If you have any questions about these, check with your teacher!

Remember: $H_2O_{(s)}$ + heat \rightleftharpoons $H_2O_{(l)}$

We decided that all the H₂O in the universe should remain as a *solid* because $H_2O_{(s)}$ has *lower enthalpy* than $H_2O_{(l)}$ and nature *favours a state of* <u>minimum enthalpy</u>.

Well, now we can explain why there is some liquid water in the universe (lots of it):

 $H_2O_{(l)}$ has <u>higher entropy</u> than $H_2O_{(s)}$

There is a natural tendency in nature toward maximum disorder or maximum entropy!

Chemical systems will tend toward a state of <u>maximum entropy</u> if no other factors are considered.

Another was of stating this might be:

A chemical reaction will favour the side (reactants or products) with <u>maximum entropy</u> if no other factors are considered.

Remember, the other factor which controlled reactions was <u>*enthalpy*</u>. (chemical potential energy). Also remember that:

Chemical systems will tend toward a state of <u>minimum enthalpy</u> if sufficient activation energy is available and <u>no other factors are considered</u>.

or

A chemical reaction will favour the side (reactants or products) with <u>minimum enthalpy</u> if no other factors are considered.

You figure out which has the most *enthalpy* (reactants or products) by looking at the ΔH or the *heat term*.

Also, remember that you can figure out which has the more *entropy* (reactants or products) by looking at the subscripts which represent the phases.

Also, we can combine the rules about "natural tendencies" to come up with this:

In nature, there is a tendency toward *minimum enthalpy* and *maximum entropy*.

Now, let's consider this process again:

$$H_2O_{(s)}$$
 + heat \rightleftharpoons $H_2O_{(l)}$

The two tendencies are said to "oppose each other" in this case:

The tendency toward *minimum enthalpy* would *favour the reactant* !(since you have to add heat energy to $H_2O_{(s)}$ to get $H_2O_{(l)}$, $H_2O_{(s)}$ has *minimum enthalpy*)

In this case the tendency toward *maximum entropy* would tend to *favour the product*. (A *liquid* has more *entropy* (disorder) than a *solid*)

We say that:

When the two tendencies *oppose each other* (one favours reactants, the other favours products), the reaction will *reach a state of equilibrium*.

That is, there will be some reactants and some products present. The relative amounts of each depends on conditions like temperature, pressure, concentration etc.

Since this is the case with : $H_2O_{(s)}$ + heat $\rightleftharpoons H_2O_{(l)}$, there is some solid water and some liquid water in the universe. (In other words, there is a state of equilibrium) Which one is present in the greater amount is determined largely by the *temperature*.

Now, lets consider another simple process: A glass bottle is knocked down from a high shelf onto a concrete floor and the glass shatters:

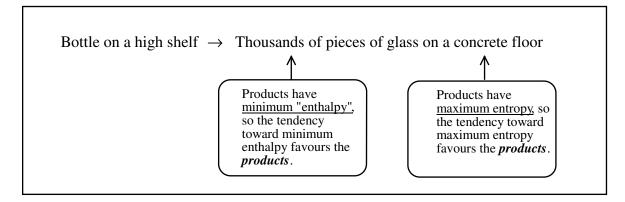
Bottle on a high shelf \rightarrow Thousands of pieces of glass on a concrete floor

The bottle falls *down* and not up! This happens because there is a natural tendency toward *minimum gravitational potential energy* (like *minimum enthalpy* in chemistry)

In other words the tendency toward *minimum gravitational potential energy* favours the *products* (the low bottle rather than the high)

(The person who knocked the bottle off of the shelf was simply supplying the " activation energy ")

Remember that the bottle broke into thousands of pieces when it hit the concrete. The broken pieces of glass have *more disorder* (*entropy*) than the bottle, so in this process, the tendency toward *maximum entropy* also *favours the products*!



There is **no** "equilibrium" here when the process is finished. That bottle has completely fallen down and it is all broken. (This bottle is no longer on the shelf and it is no longer an "unbroken bottle")

We can summarize what happened here:

Processes in which <u>both</u> the tendency toward *minimum enthalpy* and toward *maximum entropy* favour the *products*, will go to completion.

(ie. All reactants will be converted into products. There will be no reactants left once the process is finished!)

Here's an example of a chemical reaction in which this happens:

$$2K_{(s)} + 2H_2O_{(l)} \rightleftharpoons 2KOH_{(aq)} + H_{2(g)} + heat$$

This process is *exothermic* (the heat term is on the right) so the *products have lower enthalpy*.

The tendency toward *minimum enthalpy* favours the *products*.

There is a mole of gas on the right $(H_{2(g)})$ and no gases in the reactants Therefore, the *products* have greater entropy.

The tendency toward *maximum entropy* favours the *products*.

Since *both tendencies favour the products*, this reaction *will go to completion*.

That is, all of the reactants (assuming you have the correct mole ratios eg. 2 moles of K to 2 moles of H_2O) will be converted to products.

If one reactant is *in excess*, the *limiting reactant* will be *completely consumed*.

So, if you put a little bit of potassium in a beaker of water, the reaction will keep going until all of the potassium is used up. There will be *no* potassium left once the reaction is complete.

In other words, the reverse reaction does not occur!

Let's consider one more process:

$$2KOH_{(aq)} + H_{2(g)} + heat \rightleftharpoons 2K_{(s)} + 2H_2O_{(l)}$$

In this case, the tendency toward <u>minimum enthalpy</u> favours the <u>reactants</u>, and the

tendency toward *maximum entropy* also favours the *reactants*.

Processes in which <u>both</u> the tendency toward *minimum enthalpy* and toward *maximum entropy* favour the *reactants*, will <u>not occur at all!</u>.

(ie. None of the reactants will be converted into products. There will be no products formed!)

NOTE: This would be like thousands of pieces of glass spontaneously sticking together, forming a bottle and jumping up onto a high shelf! This does not occur at all. (At least I've never seen it happen!)

To summarize:

When the two tendencies *oppose each other* (one favours reactants, the other favours products), the reaction will *reach a state of equilibrium*.

Processes in which <u>both</u> the tendency toward *minimum enthalpy* and toward *maximum entropy* favour the <u>products</u>, will <u>go to completion</u>.

Processes in which <u>both</u> the tendency toward *minimum enthalpy* and toward *maximum entropy* favour the <u>reactants</u>, will <u>not occur at all!</u>.

10. For each of the following reactions decide which has *minimum enthalpy* (reactants or products), which has *maximum entropy* (reactants or products), and if the reactants are mixed, what will happen? (go to completion/ reach a state of equilibrium/not occur at all).

a) $PCl_{3(g)} + Cl_{2(g)} \rightleftharpoons PCl_{5(g)}$; $\Delta H = -92.5 \text{ kJ}$			
	The	has/have minimum enthalpy.	
	The	has/have maximum entropy.	
	If PCl_3 and Cl_2 are put together, what should h equilibrium/not occur at all)	appen?(go to completion/ reach a state of	
b)	$2NO_{2(g)} \rightleftharpoons N_2O_{4(g)} + energy$		
	The	has/have minimum enthalpy.	
	The	has/have maximum entropy.	
	If NO ₂ was put in a flask, what should happen?(go to completion/ reach a state of equilibrium/not occur at all)		
c)	$P_{4(s)} + 6H_{2(g)} + 37 \text{ kJ} \rightleftharpoons 4PH_{3(g)}$		
	The	has/have minimum enthalpy.	

		The	has/have maxi	mum entropy.
		If $P_{4(s)}$ and $6H_{2(g)}$ was put in a flask, what should happen?(go to completion/reach a state of equilibrium/not occur at all)		
	d)	$2PbO_{(s)} + 4NO_{2(s)}$	$_{g)} + O_{2(g)} \rightleftharpoons 2Pb(NO_3)_{2(s)}; \Delta H = -59$	97 kJ
		The	has/have minin	num enthalpy.
		The	has/have max	timum entropy.
		If PbO _(s) and NO _{2(g)} of equilibrium/not occur at al	were put in a flask, what should happen?(go 1)	to completion/ reach a state
M	ore Qu	estions:		
1.	Wh	at is meant by <i>enthalp</i>	y?	
2.	Wh	at is meant by <i>entropy</i>	?	
3.		endothermic reaction, num enthalpy.	the	have
4.		n exothermic reaction imum enthalpy.	, the	have
5.	a) li	quids b) gases c) aqu	order from <i>least entropy</i> to <i>greatest entropy</i> eous solutions d) solids	
6.			< < y toward	
	and		entropy.	
7.	-	rocess in which <i>entrop</i> o completion/ reach a state of	y <i>increases</i> and <i>enthalpy decreases</i> will equilibrium/not occur at all)	
8.	-	-	y <i>increases</i> and <i>enthalpy increases</i> will equilibrium/not occur at all)	

9.	A process in which <i>entropy decreases</i> and <i>enthalpy decreases</i> will (go to completion/ reach a state of equilibrium/not occur at all)		
10.	A process in which <i>entropy decreases</i> and <i>enthalpy increases</i> will (go to completion/ reach a state of equilibrium/not occur at all)		
11.	A process in which both the enthalpy and entropy trends favour reactants will		
	(go to completion/ reach a state of equilibrium/not occur at all)		
12.	A process in which both the enthalpy and entropy trends favour products will		
	(go to completion/ reach a state of equilibrium/not occur at all)		
13.	A process in which the enthalpy and entropy trends oppose each other will		
	(go to completion/ reach a state of equilibrium/not occur at all)		
14.	In each of the following, state which has the maximum entropy, (reactants or products)		
	a) $C_{(s).} + O_{2(g)} \rightleftharpoons CO_{2(g)}$		
	b) $2Al(s) + 6HCl_{(aq)} \rightleftharpoons 3H_{2(g)} + 2AlCl_{3(aq)}$		
	c) $2SO_{3(g)} \rightleftharpoons 2SO_{2(g)} + O_{2(g)}$		
	d) $HCl_{(g)} \rightleftharpoons H^+_{(aq)} + Cl^{(aq)}$		
	e) $KOH_{(s)} \rightleftharpoons K^+_{(aq)} + OH^{(aq)}$		
15.	For each of the following reactions decide which has <i>minimum enthalpy</i> (reactants or products) which has <i>maximum entropy</i> (reactants or products), and if the reactants are mixed, what will happen? (go to completion/ reach a state of equilibrium/not occur at all). Assume there is sufficient activation energy to initiate any spontaneous reaction.		
	a) $PCl_{5(g)} \rightleftharpoons PCl_{3(g)} + Cl_{2(g)}$; $\Delta H = +92.5 \text{ kJ}$		
	The has/have minimum enthalpy.		
	The has/havemaximum entropy.		

If PCl₅ is put in a flask what should happen?(go to completion/ reach a state of equilibrium/not occur at all)

b)	$2NO_{(g)} + O_{2(g)} \rightleftharpoons 2NO_{2(g)} + energy$		
	The	has/have minimum enthalpy.	
	The	has/have maximum entropy.	
	If NO and O ₂ were put in a flask, what should happen?(go to completion/ reach a state of equilibrium/not occur at all)		
c)	$\overline{\text{Na}_2\text{CO}_{3(s)}}$ + $2\text{HCl}_{(aq)}$	$2NaCl_{(aq)} + CO_{2(g)} + H_2O_{(l)} + 27.7 kJ$	
	The	has/have minimum enthalpy.	
	The	has/have maximum entropy.	
	If $Na_2CO_{3(s)}$ + 2HCl _(aq) were put in a flask, what should happen?(go to completion/ reach a state of equilibrium/not occur at all)		
d)	$2Pb(NO_3)_{2(s)} + 597 \text{ kJ} \rightleftharpoons 2PbO_{(s)} + 4NO_{2(g)} + O_{2(g)};$		
	The	has/have minimum enthalpy.	
	The	has/have maximum entropy.	
	If Pb(NO ₃) ₂ was put in a flask, what should happen?(go to completion/ reach a state of equilibrium/not occur at all)		
Rea	ctions which result in a/an	in enthalpy and a/an	
		in entropy will <i>always</i> be spontaneous .	
Rea	ctions which result in a/an	in enthalpy and a/an	
		_ in entropy will <i>always</i> be non-spontaneous .	

Do Worksheet 2-1