## Equilibria

This topic illustrates that many chemical reactions are reversible and involve an equilibrium process. The consideration of the many factors that can affect an equilibrium is an important aspect of physical chemistry.
a explain, in terms of rates of the forward and reverse reactions, what is meant by a reversible reaction and dynamic equilibrium
b state Le Chatelier's principle and apply it to deduce qualitatively (from appropriate information) the effects of changes in temperature, concentration or pressure on a system at equilibrium
c state whether changes in temperature, concentration or pressure or the presence of a catalyst affect the value of the equilibrium constant for a reaction
d deduce expressions for equilibrium constants in terms of concentrations, $\mathrm{K}_{\mathrm{c}}$, and partial pressures, $K_{p}$ (treatment of the relationship between $K_{p}$ and $K_{c}$ is not required)
e calculate the values of equilibrium constants in terms of concentrations or partial pressures from appropriate data
f calculate the quantities present at equilibrium, given appropriate data (such calculations will not require the solving of quadratic equations)
describe and explain the conditions used in the Haber process and the Contact process, as examples of the importance of an understanding of chemical equilibrium in the chemical industry

## EQUILIBRIA

7 Equilibria
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## Learning outcomes

Candidates should be able to:
7.1 Chemical equilibria: reversible reactions; dynamic equilibrium
a) explain, in terms of rates of the forward and reverse reactions, what is meant by a reversible reaction and dynamic equilibrium
b) state Le Chatelier's principle and apply it to deduce qualitatively (from appropriate information) the effects of changes in temperature, concentration or pressure on a system at equilibrium
c) state whether changes in temperature, concentration or pressure or the presence of a catalyst affect the value of the equilibrium constant for a reaction
d) deduce expressions for equilibrium constants in terms of concentrations, $K_{c}$, and partial pressures, $K_{p}$ (treatment of the relationship between $K_{p}$ and $K_{c}$ is not required)
e) calculate the values of equilibrium constants in terms of concentrations or partial pressures from appropriate data
f) calculate the quantities present at equilibrium, given appropriate data (such calculations will not require the solving of quadratic equations)
g) describe and explain the conditions used in the Haber process and the Contact process, as examples of the importance of an understanding of chemical equilibrium in the chemical industry

## EQUILIBRIA

## REVERSIBLE REACTIONS

In most reactions the reactants are completely converted to products i.e. the reaction proceeds in one direction only. Such reactions are referred to as irreversible reactions.

There are some reactions, which can be reversed by altering the conditions of the reactions.

For example when gaseous ammonia and hydrogen chloride are brought together at room temperature white dense fumes of ammonium chloride is formed.

However, on heating, ammonium chloride decomposes to give ammonia and hydrogen chloride. Such reactions are described as reversible reactions.

## REVERSIBLE REACTIONS

If in a reversible reaction the conditions required for the forward reaction and the backward reaction are the same, a stage will be reached when the rate of the forward reaction becomes equal to the rate of the backward reaction.

Then the system is said to be in a state of chemical equilibrium.
The equilibrium is dynamic. i.e. the forward and the backward reactions keep on occurring continually without affecting the concentration of the components.

Thus, at equilibrium the concentration of the constituents do not change with time.

## CHARACTERISTICS OF THE EQUILIBRIUM STATE

A system is said to be in a state of dynamic equilibrium when in a reversible reaction, the rate of the forward reaction is equal to the rate of the backward reaction.

1. The rate of the forward reaction is equal to the rate of the backward reaction.
2. The concentration of the constituents at equilibrium does not change with time.
3. Equilibrium is a dynamic state - opposing changes at molecular level are continually taking place.
4. The position of equilibrium is sensitive to changes of temperature and pressure or concentration.
5. The equilibrium may be attained from either direction.

## EQUILIBRIUM LAW

For an equilibrium reaction of the form:

$$
w A+x B \rightleftharpoons y C+z D
$$

Then (at a constant temperature) $=\frac{[C]^{y} \cdot[\mathrm{D}]^{\mathrm{z}}}{[\mathrm{A}]^{\mathrm{w}} \cdot[\mathrm{B}]^{\mathrm{x}}} \quad=$ a constant, $\left(\mathrm{K}_{\mathrm{c}}\right)$
Where [] denotes the equilibrium concentration in $\mathrm{mol} \mathrm{dm}^{-3} . \mathbf{K}_{\mathbf{c}}$ is known as the Equilibrium Constant.

Note: Solids are NOT a part of equilibrium expressions because the concentration of solids is ALWAYS constant.

## EXAMPLE 1

Consider the reaction $2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g})$

If the equilibrium concentrations are:
$\mathrm{SO}_{2}: 0.016 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{O}_{2}: 0.0083 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{SO}_{3}: 0.15 \mathrm{~mol} \mathrm{dm}^{-3}$,
Calculate the value of the equilibrium constant $\mathrm{K}_{\mathrm{c}}$, stating its units.

$$
\mathrm{K}_{\mathrm{c}}=\frac{\left[\mathrm{SO}_{3}\right]^{2}}{\left[\mathrm{SO}_{2}\right]^{2} \times\left[\mathrm{O}_{2}\right]}=\frac{(0.15)^{2}}{(0.016)^{2} \times 0.0083}=1.1 \times 10^{4} \mathrm{~mol}^{-1} \mathrm{dm}^{3}
$$

The units can be calculated by realizing:

$$
\frac{\text { conc }^{2}}{\operatorname{conc}^{2} \times \text { conc }}=\frac{1}{\operatorname{conc}} \quad \text { which is }=\frac{1}{\mathrm{~mol} \mathrm{dm}^{-3}}=\mathrm{mol}^{-1} \mathrm{dm}^{3}
$$

## EXAMPLE 2

What are the units of the equilibrium constant $K_{c}$ for the following reactions?
(a) $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})$
(b) $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{g})$
a) Units are $\frac{\text { conc }^{2}}{\text { conc } x \operatorname{conc}^{3}}=\frac{1}{\operatorname{conc}^{2}}=\mathrm{mol}^{-2} \mathrm{dm}^{6}$
b) Units are $\frac{\text { conc }^{2}}{\text { conc } \times \operatorname{conc}} \quad=$ no units as all the concentration terms cancel

## CALCULATING Kc - EXAMPLE 1

Consider the equilibrium $\mathrm{P}+2 \mathrm{Q} \rightleftharpoons \mathrm{R}+\mathrm{S}$ (all species are aqueous)
One mole of $P$ and one mole of $Q$ are mixed in $1 \mathrm{dm}^{3}$ water. Once equilibrium has been achieved 0.6 moles of $P$ are present. How many moles of $Q, R$ and $S$ are present at equilibrium? What is the $\mathrm{K}_{\mathrm{c}}$ ?
moles that reacted
moles @ equilibrium
[conc] @ equilibrium

|  | $P$ | $Q$ | $R$ | $S$ |
| :---: | :---: | :---: | :---: | :---: |
| initial moles | 1 | 1 | 0 | 0 |

$\qquad$
0.6
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

## CALCULATING Kc - EXAMPLE 1

The method is:

1. Construct the balanced equation, including state symbols (aq), (g) etc.
2. Write down the expression for $\mathrm{K}_{\mathrm{c}}$.
3. Calculate the moles of each substance that reacted.
4. Calculate the moles of each substance at equilibrium.
5. Divide these by the volume to get the concentration of each substance, use V if no volume given.
6. Put the equilibrium values into the expression for Kc and calculate the answer and work out the units.

| CALCULATING Kc - EXAMPLE 1 <br> Consider the equilibrium $\mathrm{P}+2 \mathrm{Q} \rightleftharpoons \mathrm{R}+\mathrm{S}$ (all species are aqueous) |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |
| One mole of $P$ and one mole of $Q$ are mixed in $1 \mathrm{dm}^{3}$ water achieved 0.6 moles of P are present. |  |  |  |  |  |  |
|  |  |  |  |  |  | If 0.6 mol of P remain of the original $1 \mathrm{~mol}, 0.4 \mathrm{~mol}$ have reacted. |
| initial moles | P | Q | R 0 |  |  | The equation states that 2 moles of $Q$ react with every 1 mole of $P$. |
| moles that reacted | 0.4 | 0.8 | 0.4 | 0.4 |  | This means that $0.8(2 \times 0.4)$ moles of $Q$ have reacted, leaving 0.2 |
| moles @ equilibrium | 0.6 | 0.2 | 0.4 | 0.4 |  | moles. |
| [conc] @ equilibrium | 0.6 | 0.2 | 0.4 | 0.4 |  | One mole of R and S are produced from every mole of $P$ that reacts. |
| $\mathrm{K}_{\mathrm{c}}=$ |  |  |  |  |  | This means 0.4 moles of R and 0.4 moles of $S$ are present at equilibrium. |

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## CALCULATING Kc - EXAMPLE 2

One mole of ethanoic acid reacts with one mole of ethanol at 298K. When equilibrium is reached it is found that two thirds of the acid has reacted. Calculate the value of $\mathrm{K}_{\mathrm{c}}$.

| $\mathrm{CH}_{3} \mathrm{COOH}(\mathrm{I})$ | $+\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}(\mathrm{I})$ | $\rightleftharpoons$ | $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{CH}_{2} \mathrm{CH}_{3}(\mathrm{I})$ | + |
| ---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{CH}_{3} \mathrm{COOH}$ | $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}$ | $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{CH}_{2} \mathrm{CH}_{3}$ | $\mathrm{H}_{2} \mathrm{O}$ |
| initial moles | 1 | 1 | 0 | 0 |
| moles that reacted | $2 / 3$ | $2 / 3$ | $2 / 3$ | $2 / 3$ |
| moles @ equilibrium | $1-2 / 3=1 / 3$ | $1-2 / 3=1 / 3$ | $2 / 3$ | $2 / 3$ |
| [conc] @ equilibrium | $1 / 3 / \mathrm{V}$ | $1 / 3 / \mathrm{V}$ | $2 / 3 / \mathrm{V}$ | $2 / 3 / \mathrm{V}$ |

$$
\mathrm{K}_{\mathrm{c}}=\frac{\left[\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{CH}_{2} \mathrm{CH}_{3}\right]\left[\mathrm{H}_{2} \mathrm{O}\right]}{\left[\mathrm{CH}_{3} \mathrm{COOH}\right]\left[\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}\right]}=\frac{2 / 3 / \mathrm{V} \cdot 2 / 3 / \mathrm{V}}{1 / 3 / \mathrm{V} .1 / 3 / \mathrm{V}}=4
$$

## CALCULATING Kc - EXAMPLE 3

2.6 mol of ethanoic acid, $\mathrm{CH}_{3} \mathrm{COOH}$, was mixed with 1.6 mol of ethan $-1,2-$ diol and allowed to reach equilibrium when 1.1 mol of the ester $\mathrm{CH}_{3} \mathrm{COOCH}_{2} \mathrm{CH}_{2} \mathrm{OCOCH}_{3}$, was present. The total volume was $250 \mathrm{~cm}^{3}$. Calculate the value of the equilibrium constant, $\mathrm{K}_{\mathrm{c}}$.
$2 \mathrm{CH}_{3} \mathrm{COOH}(\mathrm{I})+\mathrm{CH}_{2}(\mathrm{OH}) \mathrm{CH}_{2} \mathrm{OH}(\mathrm{I}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{COOCH}_{2} \mathrm{CH}_{2} \mathrm{OCOCH}_{3}(\mathrm{I})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$

## CALCULATING Kc - EXAMPLE 3

|  | $\mathrm{CH}_{3} \mathrm{COOH}$ | $\mathrm{CH}_{2}(\mathrm{OH}) \mathrm{CH}_{2} \mathrm{OH}$ | ester |
| :---: | :---: | :---: | :---: | $\mathrm{H}_{2} \mathrm{O}$

## CALCULATIONS INVOLVING Kc

Calculate the amount of hydrogen iodide at equilibrium when 1.0 mol of hydrogen and 1.0 mol of iodine are heated at a temperature T until equilibrium is reached in a container of volume $10 \mathrm{dm}^{3}$.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \quad \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g}) \quad \mathrm{K}_{\mathrm{c}}=49 \text { (no units) }
$$

## CALCULATIONS INVOLVING Kc

This can be done in some cases where the number of molecules on the two sides are the same:

1. Write down the expression for $\mathrm{K}_{\mathrm{c}}$ (there will always be a mark for this).
2. Assume that $x$ moles of the substance(s) on the left have reacted.
3. Calculate the equilibrium moles of each substance and their concentrations in terms on $x$.
4. Substitute these values into the expression for $\mathrm{K}_{\mathrm{c}}$ and solve for x by taking the square root of both sides.

## CALCULATIONS INVOLVING Kc

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g}) \quad \mathrm{K}_{\mathrm{c}}=\frac{[\mathrm{HI}]^{2}}{\left[\mathrm{H}_{2}\right]\left[\mathrm{I}_{2}\right]}=49
$$

Let $\times$ mol of hydrogen that react.

|  | $\mathrm{H}_{2}$ | $\mathrm{I}_{2}$ | HI |
| ---: | :---: | :---: | :---: |
| initial moles | 1 | 1 | 0 |
| moles that reacted | x | x | $2 x$ |
| moles @ equilibrium | $1-\mathrm{x}$ | $1-\mathrm{x}$ | $2 x$ |
| [conc] @ equilibrium | $(1-\mathrm{x}) / 10$ | $(1-\mathrm{x}) / 10$ | $2 x / 10$ |

## CALCULATIONS INVOLVING Kc

$$
\begin{gathered}
\mathrm{K}_{\mathrm{c}}=\frac{[H 1]^{2}}{\left[\mathrm{H}_{2}\right]\left[1_{2}\right]}=\frac{(2 \mathrm{x} / 10)}{(1-\mathrm{x}) / 10 \cdot(1-\mathrm{x}) / 10}=\frac{4 x^{2}}{(1-\mathrm{x})^{2}}=49 \\
\frac{2 \mathrm{x}}{1-\mathrm{x}}= \pm 7
\end{gathered}
$$

$$
\begin{array}{cc}
\frac{2 x}{1-x}=+7 & 2 x=7-7 x \\
9 x=7 \\
x=7 / 9=0.78
\end{array}
$$

As the moles of HI at equilibrium is $2 x$, there are 1.56 moles of HI at equilibrium.

$$
\frac{2 x}{1-x}=-7
$$

$$
\begin{gathered}
2 x=-7+7 x \\
-5 x=-7
\end{gathered}
$$

$$
x=-7 / 5=-1.4
$$

As the moles of $\mathrm{H}_{2}$ at equilibrium is $1-$ $x$, that would result in -0.4 moles of $\mathrm{H}_{2}$ at equilibrium. NOT POSSIBLE

## EQUILIBRIUM CONSTANT Kp

$K_{p}$ is the equilibrium constant in terms of the partial pressures of the gases in the equilibrium reaction.

For the reaction: $\quad \mathrm{nA}(\mathrm{g})+\mathrm{mB}(\mathrm{g}) \rightleftharpoons \mathrm{xC}(\mathrm{g})+\mathrm{yD}(\mathrm{g})$

$$
K_{p}=\frac{p C^{x} \times p D^{y}}{p A^{n} \times p B^{m}}
$$

Partial pressure of gas $A$ :
$p(A)=$ mole fraction of $A \times$ total pressure
Mole fraction of $A=\frac{\text { moles of } A}{\text { total moles of gas }}$

1. Use the value of the partial pressure of any gas in the equilibrium mixture.
2. As with $K_{c}$, the units of $K_{p}$ depend on the stoichiometry of the reaction.

## EXAMPLE 1

Calculate the partial pressure of sulfur dioxide in a mixture containing $0.016 \mathrm{~mol} \mathrm{SO}_{2}, 0.0083 \mathrm{~mol} \mathrm{O}_{2}$ and $0.15 \mathrm{~mol}_{3}$ in a vessel at a pressure of 7.0 atm . and hence the $\mathrm{K}_{\mathrm{p}}$ of the following reaction:

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

Total number of moles $=0.016+0.0083+0.15=0.1743 \mathrm{~mol}$

## EXAMPLE 1

$\mathrm{SO}_{2}: \quad$ Mole fraction of $\mathrm{SO}_{2}=0.016 / 0.1743=0.0918$
Partial Pressure of $\mathrm{SO}_{2}=0.0918 \times 7.0=0.643 \mathrm{~atm}$
$\mathrm{O}_{2}$ : Mole fraction of $\mathrm{O}_{2}=0.0083 / 0.1743=0.0476$
Partial Pressure of $\mathrm{O}_{2}=0.0476 \times 7.0=0.333 \mathrm{~atm}$
$\mathrm{SO}_{3}: \quad$ Partial Pressure of $\mathrm{SO}_{3}=7-(0.643+0.333)=6.024$

$$
K_{p}=\frac{\mathrm{p}\left(\mathrm{SO}_{3}\right)^{2}}{\mathrm{p}\left(\mathrm{SO}_{2}\right)^{2} \times \mathrm{p}\left(\mathrm{O}_{2}\right)}=\frac{(6.024)^{2}}{(0.643)^{2} \times 0.333}=264 \mathrm{~atm}^{-1}
$$

## EXAMPLE 2

When some phosphorous pentachloride $\mathrm{PCl}_{5}$, was heated to a temperature T in a container, $78 \%$ of it decomposed and the pressure in the container was 2.2 atm .

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

Calculate the value of the equilibrium constant Kp .

$$
K_{p}=\frac{p\left(\mathrm{PCl}_{3}\right) \times p\left(\mathrm{Cl}_{2}\right)}{p\left(\mathrm{PCl}_{5}\right)}
$$

## EXAMPLE 2

The method is:

1. Write down the expression for $K_{p}$.
2. Calculate the moles of each substance at equilibrium - starting amount is given, assume there was 1 mol of the reactant on the left.
3. Add up all the equilibrium moles to get the total number of moles. Work out the mole fraction of each gas (moles/total moles). Work out the partial pressure of each gas (mole fraction $x$ total pressure).
4. Put these partial pressures into the expression for $K_{p}$ and calculate its value with units.

## EXAMPLE 2

Assume that there was 1 mol of $\mathrm{PCl}_{5}$ to start with. As $78 \%$ reacted, 0.22 mol was present at equilibrium. 0.78 mol of $\mathrm{PCl}_{3}$ and $\mathrm{Cl}_{2}$ were formed.

| initial moles | $\mathrm{PCl}_{5}$ | $\mathrm{PCl}_{3}$ | $\mathrm{Cl}_{2}$ |
| ---: | :---: | :---: | :---: |
| moles that reacted | 0.78 | 0 | 0 |
| moles @ equilibrium | 0.22 | 0.78 | 0.78 |
| mole fraction | $0.22 / 1.78=0.124$ | 0.78 | 0.78 |
| partial pressure | $0.124 \times 2.2=0.273$ | $0.438 \times 2.2=0.964$ | $0.78=0.438$ |
|  | $K_{p}=\frac{p\left(\mathrm{PCl}_{3}\right) \times p\left(\mathrm{Cl}_{2}\right)}{p\left(\mathrm{PCl}_{5}\right)}$ | $=\frac{0.964 \times 0.964}{0.272}$ | $=3.48=0.438$ |
|  |  |  |  |

## CALCULATING Kp

When nitrogen ( 1 mole) and hydrogen ( 3 moles) react at constant temperature at a pressure of $8 \times 10^{6} \mathrm{~Pa}$, the equilibrium mixture was found to contain 0.7 moles of ammonia. Calculate $K_{p}$.

|  | $\mathrm{N}_{2}(\mathrm{~g})$ | $+\mathrm{H}_{2}(\mathrm{~g})$ | $\rightleftharpoons$ | $2 \mathrm{NH}_{3}(\mathrm{~g})$ |
| ---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{N}_{2}$ | $\mathrm{H}_{2}$ | $\mathrm{NH}_{3}$ |  |
| initial moles | 1 | 3 | 0 |  |
| moles that reacted | 0.35 | 1.05 | 0.7 |  |
| moles @ equilibrium | 0.65 | 1.95 | 0.7 |  |
| partial pressure | $0.65 / 3.3 \times 8 \times 10^{6}$ | $1.95 / 3.3 \times 8 \times 10^{6}$ | $0.7 / 3.3 \times 8 \times 10^{6}$ |  |

## CALCULATING Kp

$$
\begin{aligned}
K_{p}=\frac{p\left(\mathrm{NH}_{3}\right)^{2}}{p\left(\mathrm{~N}_{2}\right) \times p\left(\mathrm{H}_{2}\right)^{3}} & =\frac{\left(1.7 \times 10^{6}\right)^{2}}{\left(1.58 \times 10^{6}\right) \times\left(4.77 \times 10^{6}\right)^{3}} \\
& =1.73 \times 10^{-14} \mathrm{~Pa}^{-2}
\end{aligned}
$$

Note: The ratio of partial pressures is the same as the mole ratio at equilibrium.

## SKILL CHECK

When nitrogen ( 1 mole) and hydrogen ( 3 moles) react at constant temperature at a pressure of $8 \times{ }^{6} \mathrm{~Pa}$, the equilibrium mixture was found to contain 0.7 moles of ammonia. Calculate $\mathrm{K}_{\mathrm{p}}$.

## SKILL CHECK

Given that $\mathrm{PCl}_{5}$ at 700 K and 1.1 atm total pressure dissociated $33 \%$, find K.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

## SKILL CHECK

A chemist carries out an investigation on the equilibrium system shown below.

$$
2 \mathrm{CO}(\mathrm{~g})+2 \mathrm{NO}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{CO}_{2}(\mathrm{~g})+\mathrm{N}_{2}(\mathrm{~g})
$$

The chemist mixes 0.46 mol of CO with 0.45 mol of NO . The mixture is left to reach equilibrium at constant temperature.

The chemist analyses the equilibrium mixture and finds that 0.25 mol NO remains and the total pressure was 5 atm. What is the value of $K_{p}$.

## SKILL CHECK

1.0 mol of $\mathrm{N}_{2} \mathrm{O}_{4}$ was allowed to reach equilibrium at 400 K . At equilibrium the partial pressure of $\mathrm{N}_{2} \mathrm{O}_{4}$ was found to be 0.15 atm . Given that the $\mathrm{K}_{\mathrm{p}}$ for the reaction is 48 atm , calculate the partial pressure of $\mathrm{NO}_{2}$ in the equilibrium mixture.

$$
\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \rightleftharpoons \mathbf{2} \mathrm{NO}_{2}(\mathrm{~g})
$$

## VALUE OF K AND THE EXTENT OF A REACTION

If the value of $\mathrm{K}_{\mathrm{c}}$ is large it means that the position of the equilibrium is well to the right (towards products).

If the value of $K_{c}$ is close to 1 it means that the position of the equilibrium is close to being halfway.

If the value of $\mathrm{K}_{c}$ is very small ( $<1$ ), the position of equilibrium is well to the left (towards reactants).

Reminder: the concentration of pure solids and pure liquids is a constant and so is not included in the expression for $\mathrm{K}_{\mathrm{c}}$.

$$
\text { e.g. } \mathrm{C}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g})
$$

$$
\mathrm{K}_{\mathrm{c}}=\frac{[\mathrm{CO}]\left[\mathrm{H}_{2}\right]}{\left[\mathrm{H}_{2} \mathrm{O}\right]}
$$

## VALUE OF K AND THE EXTENT OF A REACTION

Only the partial pressures of gases are included in the expression for Kp ; the other substances are ignored.

Example State the expression for Kp for the reaction

$$
\mathrm{CaCO}_{3}(\mathrm{~s}) \rightleftharpoons \mathrm{CaO}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g})
$$

and calculate the pressure of carbon dioxide when calcium carbonate is heated to $800^{\circ} \mathrm{C}$ given that $\mathrm{K}_{p}=0.80 \mathrm{~atm}$.

$$
\mathrm{K}_{p}=p\left(\mathrm{CO}_{2}\right)=0.80 \mathrm{~atm}
$$



## LE CHATELIER'S PRINCIPLE

If a change in condition is made to a closed system in equilibrium, the system responds to counteract the change as much as possible.

Conditions include concentration, pressure and temperature
The system counteracts the change by making the direction of the reaction move against the change.

Note: Catalysts do not affect the position of equilibrium! They only speed up the rate of reaching equilibrium - getting there faster!

## EFFECT OF CONCENTRATION ON EQUILIBRIUM

TWO GENERAL RULES:

1. If the concentration of a substance is increased, the system will counteract the change by forcing the reaction to go in the direction that uses up the 'extra' substance.
2. If the concentration of a substance is decreased, the system will counteract the change by forcing the reaction to go in the direction that makes more of the substance.

## Example

$$
\mathrm{A}(\mathrm{~g})+2 \mathrm{~B}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{C}(\mathrm{~g}) \quad \Delta \mathrm{H}=-200 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

If concentration of $C$ is increased, the equilibrium will move in a way so as to oppose the change, that is, decrease C by favoring the backward reaction.

## EFFECT OF CONCENTRATION ON EQUILIBRIUM



## EFFECT OF PRESSURE ON EQUILIBRIUM



## EFFECT OF TEMPERATURE ON EQUILIBRIUM

Le Chatelier's principle also works for temperature changes.
If the temperature is increased, the reaction will go in the direction that reduces the temperature by taking in energy (converting heat energy to chemical energy), i.e. the endothermic direction.

If the temperature is decreased, the reaction will go in the direction that increases the temperature by giving out energy (converting chemical energy to heat energy), i.e. the exothermic direction.

## EFFECT OF TEMPERATURE ON EQUILIBRIUM

## Example

$$
\mathrm{A}(\mathrm{~g})+2 \mathrm{~B}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{C}(\mathrm{~g}) \quad \Delta \mathrm{H}=-200 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

If temperature increases, the equilibrium shifts in a way so as to decrease the temperature of the system. Hence, the backward reaction is favored as it is the endothermic reaction and results in a temperature decrease.

## Note:

- Low temperatures favor exothermic reactions.
- High temperatures favor endothermic reactions.


## EXOTHERMIC CHANGES

$\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \quad \boldsymbol{\Delta H}=\mathbf{- 9 2} \mathbf{~ k J ~ m o l}{ }^{-1}$


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## EXOTHERMIC CHANGES

$\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2}(\mathrm{~g}) \quad \Delta \mathrm{H}=+57 \mathrm{~kJ} \mathrm{~mol}^{-1}$


## CONDITIONS AFFECTING EQUILIBRIUM - EXAMPLE 1

Ethanol is made on an industrial scale by reacting ethene with steam at $300^{\circ} \mathrm{C}$.

$$
\mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{~g})
$$

Explain the effect that each of the following changes has on the yield of ethanol.
A. Increasing the pressure.
B. Increasing the concentration of ethene.
C. Using a catalyst.

## CONDITIONS AFFECTING EQUILIBRIUM - EXAMPLE 1

A. Increasing the pressure would increase the yield of ethanol. There are two molecules on the left of the equation and only one on the right. The system would act to reduce the increased pressure by forming fewer gas molecules, moving the equilibrium to the right (forward reaction).
B. Increasing the concentration of ethene would increase the yield of ethanol. The system would act to reduce the increased concentration by moving to the right (forward reaction) and forming more products.
C. Catalyst have no effect on yield - they only affect the rate.

## CHOOSING THE CONDITIONS

For industrial processes, the best conditions give high yields (more products) at a fast rate.

Changing conditions affects both rate and position of equilibrium.
‘Compromise conditions’ of pressure and temperature are used to give reasonable yields at fast rates.

## CHOOSING THE CONDITIONS

| CHANGES | RATE | EQUILIBRIUM POSITION (YIELD) |
| :---: | :---: | :---: |
| Increased <br> concentration of <br> reactants | Faster | Moves to the right <br> More products = higher yield |
| Increased pressure <br> (gases) | Faster | Moves to the side of the reaction with fewer gas molecules <br> For some reactions this decreases yield |
| Increased temperature | Faster | Moves in the endothermic direction <br> For exothermic reactions this decreases yield |
| Use of a catalyst | Faster | No effect |

## CONDITIONS AFFECTING EQUILIBRIUM - EXAMPLE 2

Explain whether the yield of the following reactions would increase or decrease if the temperature was increased.
A. $\mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightleftharpoons \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{g})$
B. $\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{g})$
$\Delta \mathrm{H}=-46 \mathrm{~kJ} \mathrm{~mol}^{-3}$
$\Delta \mathrm{H}=+90 \mathrm{~kJ} \mathrm{~mol}^{-3}$
C. $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})+\mathrm{C}(\mathrm{s}) \rightleftharpoons \mathrm{CO}(\mathrm{g})+\mathrm{H}_{2}(\mathrm{~g})$
$\Delta \mathrm{H}=+13 \mathrm{~kJ} \mathrm{~mol}^{-3}$

## CONDITIONS AFFECTING EQUILIBRIUM - EXAMPLE 2

A. The reaction is exothermic in the forward direction. An increase in the temperature moves the reaction in the endothermic direction, so it reverses and less yield of $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$ is formed.
B. The reaction is endothermic in the forward direction. An increase in the temperature moves the reaction in the endothermic direction, so it goes forward and more yield of NO is formed.
C. The reaction is endothermic in the forward direction. An increase in the temperature moves the reaction in the endothermic direction, so it goes forward and more yield of CO and $\mathrm{H}_{2}$ is formed.

## EFFECT OF TEMPERATURE ON K

In the following case:

$$
\mathrm{A}(\mathrm{~g})+2 \mathrm{~B}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{C}(\mathrm{~g}) \Delta \mathrm{H}=-200 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Only temperature affects $K_{c}$ and $K_{p}$. If temperature increases, $K_{c}$ decreases. This is because increasing temperature will make the reaction move backwards, increasing the value of the denominator and lowering the value of numerator.

Note: Always write the effect on equilibrium and rate. For example, for an exothermic reaction, a low temperature would slow down the rate and increase the yield.

## EFFECT OF TEMPERATURE ON K




## MEGA LECTURE

## ACID AND BASES

## ACIDS AND BASES

## ARRHENIUS THEORY

Acids are substances that produce $\mathbf{H}^{+}$ions when they dissolve in water.

Bases are substances that react with $\mathrm{H}^{+}+$ions to produce water.
Alkalis are bases which dissolve in water to produce $\mathrm{OH}^{-}$ions.
Hence, the stronger the dissociation, the stronger the acid.
However, $\mathrm{H}^{+}$ions exist as Hydronium $\mathrm{H}_{3} \mathrm{O}^{+}$ions, associating with water molecules by coordinate bonds.

## ARRHENIUS THEORY

Arrhenius theory limits itself to the concentration of $\mathrm{H}^{+}$and $\mathrm{OH}^{-}$ions determining the strength of acids and bases.

Hence, this theory is restricted to polar solvents like water.
However, this is also why Hydrochloric and Ethanoic acid show acidity.

$$
\begin{aligned}
\text { strong acid: } \mathrm{HCl} & \rightarrow \mathrm{H}^{+}+\mathrm{Cl}- \\
\text { weak acid: } \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H} & \rightleftharpoons \mathrm{H}^{+}+\mathrm{CH}_{3} \mathrm{CO}_{2}-
\end{aligned}
$$

## BRONSTED-LOWRY THEORY

According to this, an acid is a proton $(\mathrm{H}+)$ donor, and bases are proton acceptors.

In the following reaction HCl is an acid because it donates a proton to water. This means that water is acting as a Brønsted- Lowry base. The water is accepting a proton.

$$
\mathrm{H}^{+} \text {donated }
$$



```
\(\mathrm{HCl}(\mathrm{g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightarrow \mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})+\mathrm{Cl}^{-}(\mathrm{aq})\)
acid base
```


## BRONSTED-LOWRY THEORY

Also, a base has to be a substance with a lone pair, to coordinate with the proton.

$$
\mathrm{HCl}(\mathrm{~g}) \quad+\quad \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \quad \longrightarrow \quad \mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq}) \quad+\quad \mathrm{Cl}^{-}(\mathrm{aq})
$$



BRONSTED-LOWRY THEORY


## BRONSTED-LOWRY THEORY

$$
\mathrm{NH}_{3}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{NH}_{4}^{+}+\mathrm{OH}^{-}
$$

A conjugate acid is a base that has accepted a proton.
A conjugate base is an acid that has donated a proton.
Hence, the $\mathrm{NH}_{4}{ }^{+}$ion is said to be the conjugate acid of $\mathrm{NH}_{3}$, and likewise, $\mathrm{NH}_{3}$ is the conjugate base of $\mathrm{NH}_{4}{ }^{+}$ion.
$\mathrm{H}_{2} \mathrm{O}$ is the conjugate acid of the $\mathrm{OH}^{-}$ion, and the $\mathrm{OH}^{-}$ion is the conjugate base of $\mathrm{H}_{2} \mathrm{O}$.

## BRONSTED-LOWRY THEORY

This suggests that a substance can both be an acid and a base depending on the situation.


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## STRENGTH OF ACIDS

The strength of acids is judged by their tendency to donate protons.
The strength of bases is judged by their tendency to accept protons.

$$
\mathrm{HCl}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{Cl}-
$$

The above reaction proceeds to completion, therefore, HCl is a stronger acid than $\mathrm{H}_{3} \mathrm{O}^{+}$, and $\mathrm{H}_{2} \mathrm{O}$ is a stronger base than Cl .

## STRENGTH OF ACIDS <br> $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{H}_{3} \mathrm{O}^{+}+\mathrm{CH}_{3} \mathrm{CO}_{2}{ }^{-}$

Th above reaction however is a reversible reaction, with the equilibrium position well towards the left.
$\mathrm{H}_{3} \mathrm{O}^{+}$is a stronger acid than $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}$, and $\mathrm{CH}_{3} \mathrm{CO}_{2}-$ is a stronger base than $\mathrm{H}_{2} \mathrm{O}$.

Hence, strong acids have weak conjugate bases, and vice versa.

## SKILL CHECK 1

Identify which reactants are acids and which are bases:
(a) $\mathrm{NH}_{4}{ }^{+}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{+}$
(b) $\mathrm{HCOOH}+\mathrm{HClO}_{2} \rightleftharpoons \mathrm{HCOOH}_{2}{ }^{+}+\mathrm{ClO}_{2}{ }^{-}$
(c) $\mathrm{HClO}_{4}+\mathrm{CH}_{3} \mathrm{COOH} \rightleftharpoons \mathrm{ClO}_{4}^{-}+\mathrm{CH}_{3} \mathrm{COOH}_{2}{ }^{+}$
(d) $\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{HNO}_{3} \rightleftharpoons \mathrm{HSO}_{4}^{-}+\mathrm{H}_{2} \mathrm{NO}_{3}{ }^{+}$
(e) $\mathrm{H}_{2} \mathrm{~S}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HS}^{-}+\mathrm{H}_{3} \mathrm{O}^{+}$

## MEGA LECTURE

## EQUILIBRIA WS 1

## SECTION A

1 In an experiment, 2.00 mol of hydrogen and 3.00 mol of iodine were heated together in a sealed container and allowed to reach equilibrium at a fixed temperature. The container had a fixed volume of $1.00 \mathrm{dm}^{3}$. At equilibrium, there were 2.40 mol of iodine present in the mixture.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g})
$$

What is the value of the equilibrium constant, $K_{c}$ ?
A 0.107
B 0.357
C 0.429
D 2.33

2 An aqueous solution was prepared containing a mixture of 1.0 mol of $\mathrm{AgNO}_{3}$ and 1.0 mol of $\mathrm{FeSO}_{4}$ in $1.00 \mathrm{dm}^{3}$ of water. When equilibrium was established, there was 0.44 mol of $\mathrm{Ag}^{+}(\mathrm{aq})$ in the mixture.

$$
\mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{Fe}^{2+}(\mathrm{aq}) \rightleftharpoons \mathrm{Ag}(\mathrm{~s})+\mathrm{Fe}^{3+}(\mathrm{aq})
$$

What is the numerical value of $K_{\mathrm{c}}$ ?
A 0.62
B 1.40
C 1.62
D 2.89

3 The equation for the reaction between carbon monoxide and hydrogen is shown.

$$
\mathrm{CO}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
$$

What are the units of $K_{\mathrm{p}}$ for this reaction?
A kPa
B $\mathrm{kPa}^{-1}$
C $\mathrm{kPa}^{2}$
D $\mathrm{kPa}^{-2}$

4 When solid ammonium chloride dissociates at a certain temperature in a $0.500 \mathrm{dm}^{3}$ container, ammonia and hydrogen chloride are formed.

$$
\mathrm{NH}_{4} \mathrm{Cl}(\mathrm{~s}) \rightleftharpoons \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{HCl}(\mathrm{~g})
$$

The initial amount of ammonium chloride was 1.00 mol , and when the system had reached equilibrium there was 0.300 mol of ammonium chloride.

What is the numerical value of $K_{\mathrm{c}}$ for this reaction under these conditions?
A 0.490
B 1.63
C $\quad 1.96$
D 3.27

5 The equilibrium constant, $K_{c}$, for the reaction shown is $2 \mathrm{~mol}^{-2} \mathrm{dm}^{6}$, at 600 K .

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

What is the concentration of $\mathrm{NH}_{3}$ at equilibrium, at 600 K , when the equilibrium concentrations of $\mathrm{N}_{2}$ and $\mathrm{H}_{2}$ are both $2 \mathrm{~mol} \mathrm{dm}^{-3}$ ?
A $\sqrt{8} \mathrm{moldm}^{-3}$
B $\sqrt{16} \mathrm{moldm}^{-3}$
C $\sqrt{32} \mathrm{moldm}^{-3}$
D $32 \mathrm{moldm}^{-3}$

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6 Catalysts are an important feature of many industrial processes and biochemical reactions.
Which row correctly describes the effect of a catalyst on a reversible chemical reaction?

|  | position of equilibrium | effect on value of $\Delta H$ |
| :---: | :---: | :---: |
| A | moved to right | decreased |
| B | unaffected | decreased |
| C | unaffected | increased |
| D | unaffected | unaffected |

7 The table shows the partial pressures in an equilibrium mixture formed by the Haber process.

| substance | partial pressure/kPa |
| :--- | :---: |
| nitrogen | 7000 |
| hydrogen | 8000 |
| ammonia | 4000 |

$$
3 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{N}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

What is the numerical value of the equilibrium constant, $K_{\mathrm{p}}$, for this reaction?
A $4.46 \times 10^{-9}$
B $4.76 \times 10^{-5}$
C $7.14 \times 10^{-5}$
D $2.24 \times 10^{8}$
8 Nitrogen reacts with hydrogen to produce ammonia.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

A mixture of 2.00 mol of nitrogen, 6.00 mol of hydrogen and 2.40 mol of ammonia is allowed to reach equilibrium in a sealed vessel of volume $1 \mathrm{dm}^{3}$. It was found that 2.32 mol of nitrogen were present in the equilibrium mixture.

What is the value of $K_{\mathrm{c}}$ ?
A $\frac{(1.76)^{2}}{(2.32)(6.96)^{3}}$
B $\frac{(1.76)^{2}}{(2.32)(6.32)^{3}}$
C $\frac{(2.08)^{2}}{(2.32)(6.32)^{3}}$
D $\frac{(2.40)^{2}}{(2.32)(6.00)^{3}}$

9 Lithium reacts with nitrogen at room temperature to form solid $\mathrm{Li}_{3} \mathrm{~N}$.
Three vessels of equal volume are connected by taps, $A$ and $B$, as shown.

10


At the start, $A$ and $B$ are closed, the left-hand vessel is evacuated, the middle vessel has the indicated reaction at equilibrium and the right-hand vessel contains lithium only.

Which action would allow the equilibrium mixture to contain the most ammonia?
A Keep both A and B closed.
B Open both A and B .
C Open A only.
D Open B only.
11 The table shows the partial pressures in an equilibrium mixture formed by the Haber process.

| substance | partial pressure/kPa |
| :--- | :---: |
| nitrogen | 7000 |
| hydrogen | 8000 |
| ammonia | 4000 |

$$
3 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{N}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

What is the numerical value of the equilibrium constant, $K_{p}$, for this reaction?
A $4.46 \times 10^{-9}$
B $4.76 \times 10^{-5}$
C $7.14 \times 10^{-5}$
D $2.24 \times 10^{8}$
12 Hydrogen iodide dissociates into hydrogen and iodine.

$$
2 \mathrm{HI}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})
$$

In an experiment, $b$ mol of hydrogen iodide were put into a sealed vessel at pressure p. At equilibrium, $x$ mol of the hydrogen iodide had dissociated.

Which expression for $K_{\mathrm{p}}$ is correct?
A $\frac{x^{2}}{(b-x)^{2}}$
B $\frac{x^{2} p^{2}}{(b-x)^{2}}$
C $\frac{x^{2} p^{2}}{4 b(b-x)}$
D $\frac{x^{2}}{4(b-x)^{2}}$

13 For the reaction shown, an equilibrium is established at a temperature of 700 K .
The equilibrium constant, $K_{p}$, for the reaction is 9.80 kPa . The partial pressure of $\mathrm{N}_{2} \mathrm{O}_{4}$ at equilibrium is 80.0 kPa .

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

What is the partial pressure of $\mathrm{NO}_{2}$ at equilibrium?
A 8.16 kPa
B $\quad 28.0 \mathrm{kPa}$
C $\quad 66.6 \mathrm{kPa}$
D $\quad 784 \mathrm{kPa}$

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14 Ammonia is produced by the Haber process.

$$
\mathrm{N}_{2}+3 \mathrm{H}_{2} \rightleftharpoons 2 \mathrm{NH}_{3} \quad \Delta H^{\ominus}=-92 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

A fault in the temperature control during the process resulted in the temperature changing to $600^{\circ} \mathrm{C}$ for two hours.

What effect did this have on the ammonia production during this time?
A Ammonia was formed faster. The equilibrium yield decreased.
B Ammonia was formed faster. The equilibrium yield increased.
C Ammonia was formed slower. The equilibrium yield decreased.
D Ammonia was formed slower. The equilibrium yield increased.
15 Ammonia is manufactured from nitrogen and hydrogen using the Haber process.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

What is the expression for $K_{\mathrm{c}}$ for this equilibrium?
A $\frac{2\left[\mathrm{NH}_{3}(\mathrm{~g})\right]}{\left[\mathrm{N}_{2}(\mathrm{~g})\right]+3\left[\mathrm{H}_{2}(\mathrm{~g})\right]}$
B $\frac{2\left[\mathrm{NH}_{3}(\mathrm{~g})\right]}{\left[\mathrm{N}_{2}(\mathrm{~g})\right] \times 3\left[\mathrm{H}_{2}(\mathrm{~g})\right]}$
C $\frac{\left[\mathrm{NH}_{3}(\mathrm{~g})\right]^{2}}{\left[\mathrm{~N}_{2}(\mathrm{~g})\right]+\left[\mathrm{H}_{2}(\mathrm{~g})\right]^{3}}$
D $\frac{\left[\mathrm{NH}_{3}(\mathrm{~g})\right]^{2}}{\left[\mathrm{~N}_{2}(\mathrm{~g})\right] \times\left[\mathrm{H}_{2}(\mathrm{~g})\right]^{3}}$
16 Two moles of compound $P$ were placed in a sealed container. The container was heated and $P$ was partially decomposed to produce $Q$ and $R$ only. A dynamic equilibrium between $P, Q$ and $R$ was established.

At equilibrium $x$ moles of $R$ were present and the total number of moles present was $\left(2+\frac{x}{2}\right)$.
What is the equation for this reversible reaction?
A $\mathrm{P} \rightleftharpoons 2 \mathrm{Q}+\mathrm{R}$
B $\quad 2 \mathrm{P} \rightleftharpoons 2 \mathrm{Q}+\mathrm{R}$
C $2 \mathrm{P} \rightleftharpoons \mathrm{Q}+\mathrm{R}$
D $2 P \rightleftharpoons Q+2 R$

17 A reaction involved in the Contact process is shown.

$$
2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g}) \quad \Delta H^{\circ}=-197 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The reaction is investigated at 200 kPa and 700 K and the value of the equilibrium constant, $K_{\mathrm{p}}$, is found to be Y . The reaction is then investigated at 1000 kPa and 700 K and the value of $K_{\mathrm{p}}$ is found to be $Z$.

Which statement comparing Y and Z is correct?
A $Y$ and $Z$ are the same.
B Y is greater than Z .
C Z is 2.2 times greater than Y .
D Z is 5.0 times greater than Y .
18 Nitrogen dioxide, $\mathrm{NO}_{2}$, is a brown gas.
Dinitrogen tetroxide, $\mathrm{N}_{2} \mathrm{O}_{4}$, is a colourless gas.
An equilibrium is established between $\mathrm{NO}_{2}$ and $\mathrm{N}_{2} \mathrm{O}_{4}$ in a closed vessel.

$$
\underset{\text { brown }}{2 \mathrm{NO}_{2}(\mathrm{~g})} \rightleftharpoons \underset{\text { colourless }}{\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})} \quad \Delta H=-57 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which row describes the effects of changing conditions on the colour of an equilibrium mixture of $\mathrm{NO}_{2}$ and $\mathrm{N}_{2} \mathrm{O}_{4}$ ?

|  | increasing the pressure | increasing the temperature |
| :---: | :---: | :---: |
| A | colour becomes darker | colour becomes darker |
| B | colour becomes darker | colour becomes lighter |
| C | colour becomes lighter | colour becomes darker |
| D | colour becomes lighter | colour becomes lighter |

[W'17 2 Q9]
19 The reaction between sulfur dioxide and oxygen is reversible.

$$
2 \mathrm{SO}_{2}+\mathrm{O}_{2} \rightleftharpoons 2 \mathrm{SO}_{3}
$$

$$
\Delta H^{\ominus}=-196 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which conditions of pressure and temperature favour the reverse reaction?

|  | pressure | temperature |
| :---: | :---: | :---: |
| A | high | high |
| B | high | low |
| C | low | high |
| D | low | low |

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20 In a particular reversible reaction the yield of product is increased

- if the temperature is increased;
- if the pressure is decreased.

Which equation could describe this reversible reaction?
A $\mathrm{CH}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightleftharpoons 3 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{CO}(\mathrm{g}) \quad \Delta H=+206 \mathrm{~kJ} \mathrm{~mol}^{-1}$
B $4 \mathrm{NH}_{3}(\mathrm{~g})+3 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{~N}_{2}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \quad \Delta H=-227 \mathrm{~kJ} \mathrm{~mol}^{-1}$
C $2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \quad \Delta H=-58 \mathrm{~kJ} \mathrm{~mol}^{-1}$
D $3 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{O}_{3}(\mathrm{~g})$
$\Delta H=+143 \mathrm{~kJ} \mathrm{~mol}^{-1}$

21 Hydrogen is produced industrially from methane as shown in the equation.

$$
\mathrm{CH}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \quad \Delta H^{\ominus}=+205 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which conditions would most favour the formation of hydrogen?

|  | pressure | temperature |
| :---: | :---: | :---: |
| A | high | high |
| B | high | low |
| C | low | high |
| D | low | low |

[J'18 P13 Q9]
22 The chemical equilibrium shown is formed when ammonia is produced in the Haber process.

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

The following concentrations are found to be present at equilibrium under certain conditions.

| $\mathrm{N}_{2}$ | $\mathrm{H}_{2}$ | $\mathrm{NH}_{3}$ |
| :---: | :---: | :---: |
| $0.200 \mathrm{~mol} \mathrm{dm}^{-3}$ | $0.300 \mathrm{~mol} \mathrm{dm}^{-3}$ | $0.600 \mathrm{~mol} \mathrm{dm}^{-3}$ |

What is the numerical value of $K_{c}$ under these conditions?
A 0.0150
B 6.0
C 10.0
D 66.7

23 Silver chloride and silver iodide form equilibria when added to water.

$$
\begin{array}{rll}
\mathrm{AgCl}(\mathrm{~s}) & \rightleftharpoons \mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{Cl}^{-}(\mathrm{aq}) & K_{\mathrm{c}}=K_{1} \\
\mathrm{AgI}(\mathrm{~s}) \rightleftharpoons \mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{I}^{-}(\mathrm{aq}) & K_{\mathrm{c}}=K_{2}
\end{array}
$$

Each equilibrium position lies well to the left.
Silver iodide will not dissolve in aqueous ammonia. Silver chloride will dissolve in aqueous ammonia. Another equilibrium is formed.

$$
\mathrm{Ag}^{+}(\mathrm{aq})+2 \mathrm{NH}_{3}(\mathrm{aq}) \rightleftharpoons \mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}^{+}(\mathrm{aq}) \quad K_{\mathrm{c}}=K_{3}
$$

The position of this equilibrium lies to the right.
What is the order of magnitude for these three equilibrium constants?
A $K_{1}>K_{2}>K_{3}$
B $K_{2}>K_{1}>K_{3}$
C $K_{3}>K_{1}>K_{2}$
D $K_{3}>K_{2}>K_{1}$

24 In this question you should assume that all gases behave ideally.
Hydrogen and iodine react reversibly in the following reaction. The system reaches dynamic equilibrium.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g}) \quad \Delta H=-9.5 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which statement must be true for the $K_{p}$ of this equilibrium to be constant?
A The partial pressures of $\mathrm{H}_{2}, \mathrm{I}_{2}$ and HI are equal.
B The external pressure is constant.
C The forward and reverse reactions have stopped.
D The temperature is constant.
[ N '18 P11 Q9]
250.200 mol of sulfur dioxide and 0.200 mol of oxygen are placed in a $1.00 \mathrm{dm}^{3}$ sealed container. The gases are allowed to react until equilibrium is reached.

$$
2 \mathrm{SO}_{2}+\mathrm{O}_{2} \rightleftharpoons 2 \mathrm{SO}_{3}
$$

At equilibrium there is 0.100 mol of $\mathrm{SO}_{3}$ in the container.
What is the value of $K_{c}$ ?
A $\quad 0.150 \mathrm{moldm}^{-3}$
B $\quad 0.800 \mathrm{~mol} \mathrm{dm}^{-3}$
C $\quad 1.25 \mathrm{~mol}^{-1} \mathrm{dm}^{3}$
D $\quad 6.67 \mathrm{~mol}^{-1} \mathrm{dm}^{3}$

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## SECTION B

The responses $\mathbf{A}$ to $\mathbf{D}$ should be selected on the basis of

| A | B | C | D |
| :---: | :---: | :---: | :---: |
| $\mathbf{1}, \mathbf{2}$ and $\mathbf{3}$ <br> are <br> correct | $\mathbf{1}$ and $\mathbf{2}$ <br> only are <br> correct | $\mathbf{2}$ and $\mathbf{3}$ <br> only are <br> correct | $\mathbf{1}$ only <br> is <br> correct |

1 Methanoic acid molecules, $\mathrm{HCO}_{2} \mathrm{H}$, and hydrogen carbonate ions, $\mathrm{HCO}_{3}{ }^{-}$, can both behave as acids.

Why does a solution of methanoic acid have a lower pH than a solution of sodium hydrogen carbonate of the same concentration?
$1 \mathrm{HCO}_{2} \mathrm{H}$ molecules dissociate more fully than $\mathrm{HCO}_{3}{ }^{-}$ions do.
2 Each $\mathrm{HCO}_{2} \mathrm{H}$ molecule has two hydrogen atoms; each $\mathrm{HCO}_{3}{ }^{-}$ion only has one.
3 Methanoic acid is a weaker acid than sodium hydrogen carbonate.

2 The following equilibrium is an exothermic reaction in the forward direction.

$$
2 \mathrm{CrO}_{4}{ }^{2-}(\mathrm{aq})+2 \mathrm{H}^{+}(\mathrm{aq}) \rightleftharpoons \mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I})
$$

What happens when the concentration of $\mathrm{CrO}_{4}{ }^{2-}$ ions increases and the temperature decreases?

1 The concentration of $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$ ions increases.
2 The equilibrium constant increases.
3 The activation energy decreases.

3 R and S react together.

$$
R+S \rightleftharpoons T
$$

Which factors affect the rate of the forward reaction?
1 the activation energy of the reaction
2 the enthalpy change of the reaction
3 the equilibrium constant of the reaction
4 The Brønsted-Lowry theory describes acid and base character.
When concentrated sulfuric acid and concentrated nitric acid are mixed, the following reactions occur.

$$
\begin{gathered}
\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{HNO}_{3} \rightleftharpoons \mathrm{HSO}_{4}^{-}+\mathrm{H}_{2} \mathrm{NO}_{3}^{+} \\
\mathrm{H}_{2} \mathrm{NO}_{3}^{+} \rightleftharpoons \mathrm{H}_{2} \mathrm{O}+\mathrm{NO}_{2}^{+} \\
\mathrm{H}_{2} \mathrm{O}+\mathrm{H}_{2} \mathrm{SO}_{4} \rightleftharpoons \mathrm{HSO}_{4}^{-}+\mathrm{H}_{3} \mathrm{O}^{+}
\end{gathered}
$$

Which species are bases in these reactions?
1 H.SO.

5 Which statements about reversible reactions are correct?
1 An increase in concentration of a reactant always increases the concentration of the product.
2 An increase in temperature always increases the rate at which the equilibrium is established.
3 An increase in temperature always increases the concentration of the product at equilibrium.

6 In this question, all gases can be assumed to behave ideally.
A chemist heats a mixture of nitrogen and oxygen gases in a sealed container at a constant temperature until the mixture reaches a dynamic equilibrium containing $\mathrm{N}_{2}(\mathrm{~g}), \mathrm{O}_{2}(\mathrm{~g})$ and $\mathrm{NO}(\mathrm{g})$.

$$
\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})
$$

The chemist repeats the experiment at the same temperature using the same initial amounts of $\mathrm{N}_{2}(\mathrm{~g})$ and $\mathrm{O}_{2}(\mathrm{~g})$, but at a much higher pressure.

Which statements about the second experiment at higher pressure are correct?
1 At higher pressure, there are more particles per unit volume.
2 The composition of the equilibrium mixture does not change.
3 There are more collisions per second so equilibrium is reached faster.
7 In the manufacture of sulfuric acid, the following exothermic reaction occurs.

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

Which changes will move the position of the equilibrium to the right?
1 increasing the pressure
2 increasing the temperature
3 using twice as much catalyst
8 Methanol, $\mathrm{CH}_{3} \mathrm{OH}$, can be produced industrially by reacting CO with $\mathrm{H}_{2}$.

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

$$
\Delta H=-91 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The process can be carried out at $4 \times 10^{3} \mathrm{kPa}$ and 1150 K .
Which statements about this reaction are correct?
1 Increasing the temperature will increase the rate of reaction because more effective collisions will occur.

2 Lowering the temperature will reduce the rate of reaction because the forward reaction is exothermic.

3 Increasing the pressure will reduce the rate of reaction because there are a larger number of moles on the left-hand side of the equation.

9 In the manufacture of sulfuric acid, the following exothermic reaction occurs.

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

Which changes will move the position of the equilibrium to the right?
1 increasing the pressure

10 Ammonia and chlorine react together in the gas phase.

$$
8 \mathrm{NH}_{3}+3 \mathrm{Cl}_{2} \rightarrow \mathrm{~N}_{2}+6 \mathrm{NH}_{4} \mathrm{Cl}
$$

Which statements are correct?
1 Ammonia behaves as a reducing agent.
2 Ammonia behaves as a base.
3 The oxidation number of hydrogen changes.

11 In which reactions does $\mathrm{NH}_{3}$ behave as a Brønsted-Lowry acid?
$12 \mathrm{NH}_{3} \rightarrow \mathrm{NH}_{2}^{-}+\mathrm{NH}_{4}^{+}$
$2 \mathrm{HSO}_{4}^{-}+\mathrm{NH}_{3} \rightarrow \mathrm{SO}_{4}{ }^{2-}+\mathrm{NH}_{4}{ }^{+}$
$3 \mathrm{Ag}^{+}+2 \mathrm{NH}_{3} \rightarrow\left[\mathrm{Ag}\left(\mathrm{NH}_{3}\right)_{2}\right]^{+}$

12 Which statements are correct when a reversible reaction is at equilibrium?
1 All species are at equal concentration.
2 The concentrations of all species remain constant.
3 The rate of the forward reaction equals the rate of the reverse reaction.

13 Ammonia and chlorine react as shown.

$$
8 \mathrm{NH}_{3}+3 \mathrm{Cl}_{2} \rightarrow \mathrm{~N}_{2}+6 \mathrm{NH}_{4} \mathrm{Cl}
$$

Which statements are correct?
1 Each nitrogen atom is oxidised.
2 Each chlorine atom is reduced.
3 Ammonia behaves as a base.

## EQUILIBRIA WS 2

(c) At temperatures above $1500 \mathrm{~K}, \mathrm{HCl}$ will decompose.

A sample of 0.300 mol of HCl decomposed in a sealed container.
The resulting equilibrium mixture was found to contain $1.50 \times 10^{-2} \mathrm{~mol}$ of $\mathrm{Cl}_{2}$.
(i) Calculate the amounts, in mol, of $\mathrm{H}_{2}$ and HCl present in the equilibrium mixture.

$$
\begin{aligned}
& \mathrm{H}_{2}= \\
& \text { mol } \\
& \mathrm{HCl}=
\end{aligned}
$$

(ii) Calculate the mole fraction of each gas in the equilibrium mixture.

$$
\begin{aligned}
& \text { mole fraction of } \mathrm{HCl}= \\
& \text { mole fraction of } \mathrm{H}_{2}= \\
& \text { mole fraction of } \mathrm{Cl}_{2}=
\end{aligned}
$$

(d) In another experiment under different conditions, an equilibrium mixture was produced with mole fractions for each species as shown.

| species | mole fraction |
| :---: | :---: |
| HCl | 0.88 |
| $\mathrm{H}_{2}$ | 0.06 |
| $\mathrm{Cl}_{2}$ | 0.06 |

(i) Write the expression for the equilibrium constant, $K_{p}$, for the decomposition of HCl .

$$
2 \mathrm{HCl}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

$$
K_{p}=
$$

## $\xrightarrow[\mathbf{9 8}]{\text { Online Classes : Megalecture@gmail.com }}$

(ii) Explain why the total pressure of the system does not need to be known for $K_{p}$ to be calculated for this experiment.
$\qquad$
$\qquad$
(iii) Calculate the value of $K_{p}$ for this experiment.

$$
K_{p}=
$$

20 (b) The equation for the preparation of hydrogen chloride using concentrated sulfuric acid is shown.

$$
\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{NaCl} \rightarrow \mathrm{NaHSO}_{4}+\mathrm{HCl}
$$

(i) Use the Brønsted-Lowry theory of acids and bases to identify the base and its conjugate acid in this reaction. Explain your answer.

Brønsted-Lowry base (base-I) $=$ $\qquad$
conjugate acid (acid-II) = $\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) Explain why the reaction of concentrated sulfuric acid and sodium iodide is not suitable for the preparation of hydrogen iodide.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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2 (c) Hydrogen chloride undergoes a reversible reaction with oxygen.

$$
4 \mathrm{HCl}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{Cl}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
$$

The reaction is carried out at $400^{\circ} \mathrm{C}$ in the presence of a copper(II) chloride catalyst.
(i) Use the data in the table to calculate the overall enthalpy change of reaction.

| compound | enthalpy change of <br> formation $/ \mathrm{kJ} \mathrm{mol}^{-1}$ |
| :---: | :---: |
| $\mathrm{HCl}(\mathrm{g})$ | -92 |
| $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ | -242 |

enthalpy change of reaction =
(ii) State the type of catalyst used in this reaction. Explain how a catalyst is able to increase the rate of a chemical reaction.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(iii) The reaction exists in dynamic equilibrium.

The reaction was repeated at $1000^{\circ} \mathrm{C}$ and the same pressure.
State and explain the effect on the composition of the equilibrium mixture of the change in temperature.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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(iv) When 1.60 mol of HCl are mixed in a sealed container with 0.500 mol of $\mathrm{O}_{2}$ at $400^{\circ} \mathrm{C}$, 0.600 mol of $\mathrm{Cl}_{2}$ and 0.600 mol of $\mathrm{H}_{2} \mathrm{O}$ are formed.

The total pressure inside the container is $1.50 \times 10^{5} \mathrm{~Pa}$.

- Calculate the amounts, in mol, of HCl and $\mathrm{O}_{2}$ in the equilibrium mixture

$$
\begin{aligned}
& \mathrm{HCl}=\ldots . . . . . . . . . . . . . . . . . . . . . . . . . . . ~ m o l ~ \\
& \mathrm{O}_{2}=\ldots . . . . . . . . . . . . . . . . . . . . . . . . . . . . ~
\end{aligned} \mathrm{~mol}
$$

- Calculate the mole fraction of $\mathrm{Cl}_{2}$ and hence the partial pressure of $\mathrm{Cl}_{2}$ in the equilibrium mixture.
(v) In a separate experiment, an equilibrium reaction mixture was found to contain the four gases at the partial pressures shown in the table.

| gas | HCl | $\mathrm{O}_{2}$ | $\mathrm{Cl}_{2}$ | $\mathrm{H}_{2} \mathrm{O}$ |
| :---: | :---: | :---: | :---: | :---: |
| partial pressure $/ \mathrm{Pa}$ | $4.8 \times 10^{4}$ | $3.0 \times 10^{4}$ | $3.6 \times 10^{4}$ | $3.6 \times 10^{4}$ |

$$
K_{\mathrm{p}}=\frac{\left(p_{\mathrm{Cl}_{2}}\right)^{2} \times\left(p_{\mathrm{H}_{2}}\right)^{2}}{\left(p_{\mathrm{HCl},}\right)^{4} \times p_{\mathrm{O}_{2}}}
$$

Use this information and the expression given for $K_{\mathrm{p}}$ to calculate a value for $K_{\mathrm{p}}$. State the units of $K_{p}$.

$$
\begin{aligned}
K_{\mathrm{p}} & =\ldots . . . . . . . . . . . . . . . . . . . . . . . . . . ~
\end{aligned} .
$$

(vi) The reaction is repeated without a catalyst.

State the effect of this on $K_{p}$.

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3 Ammonia, $\mathrm{NH}_{3}$, is manufactured from nitrogen and hydrogen by the Haber process.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \quad \Delta H=-92 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(c) At a pressure of $2.00 \times 10^{7} \mathrm{~Pa}, 1.00 \mathrm{~mol}$ of nitrogen, $\mathrm{N}_{2}(\mathrm{~g})$, was mixed with 3.00 mol of hydrogen, $\mathrm{H}_{2}(\mathrm{~g})$. The final equilibrium mixture formed contained 0.300 mol of ammonia, $\mathrm{NH}_{3}(\mathrm{~g})$.
(i) Calculate the amounts, in mol, of $\mathrm{N}_{2}(\mathrm{~g})$ and $\mathrm{H}_{2}(\mathrm{~g})$ in the equilibrium mixture.

$$
\begin{aligned}
& \mathrm{N}_{2}(\mathrm{~g})= \\
& \text { mol } \\
& \mathrm{H}_{2}(\mathrm{~g})= \\
& \text { mol }
\end{aligned}
$$

(ii) Calculate the partial pressure of ammonia, $\mathrm{pNH}_{3}$, in the equilibrium mixture.

Give your answer to three significant figures.

$$
p \mathrm{NH}_{3}=
$$

Pa [3]

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(d) In another equilibrium mixture the partial pressures are as shown.

| substance | partial pressure $/ \mathrm{Pa}$ |
| :---: | :---: |
| $\mathrm{N}_{2}(\mathrm{~g})$ | $2.20 \times 10^{6}$ |
| $\mathrm{H}_{2}(\mathrm{~g})$ | $9.62 \times 10^{5}$ |
| $\mathrm{NH}_{3}(\mathrm{~g})$ | $1.40 \times 10^{4}$ |

(i) Write the expression for the equilibrium constant, $K_{\mathrm{p}}$, for the production of ammonia from nitrogen and hydrogen.
$K_{\mathrm{p}}=$
(ii) Calculate the value of $K_{\mathrm{p}}$ for this reaction.

State the units.

$$
\begin{array}{r}
K_{\mathrm{p}}= \\
\text { units }=
\end{array}
$$

$\qquad$
$\qquad$
(iii) This reaction is repeated with the same starting amounts of nitrogen and hydrogen. The same temperature is used but the container has a smaller volume.

State the effects, if any, of this change on the yield of ammonia and on the value of $K_{p}$. effect on yield of ammonia $\qquad$
effect on value of $K_{p}$ $\qquad$

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4 At 450 K phosphorus $(\mathrm{V})$ chloride, $\mathrm{PCl}_{5}(\mathrm{~g})$, decomposes to form phosphorus(III) chloride, $\mathrm{PC} l_{3}(\mathrm{~g})$, and chlorine, $\mathrm{Cl}_{2}(\mathrm{~g})$. A dynamic equilibrium is established as shown.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \quad \Delta H=+124 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(a) The enthalpy change of formation of $\mathrm{PCl}_{3}(\mathrm{~g})$ under these conditions is given.

$$
\Delta H_{\mathrm{f}} \mathrm{PCl}_{3}(\mathrm{~g})=-320 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Calculate the enthalpy change of formation of $\mathrm{PCl}_{5}(\mathrm{~g})$ under these conditions.
Include a sign with your answer.

> enthalpy change =
$\mathrm{kJmol}^{-1}$
(b) (i) State and explain the effect of increasing temperature on the rate of decomposition of $\mathrm{PCl}_{5}(\mathrm{~g})$.
$\qquad$
$\qquad$
$\qquad$
(ii) State and explain the effect of increasing temperature on the percentage of $\mathrm{PCl}_{5}(\mathrm{~g})$ that decomposes.
$\qquad$
$\qquad$
$\qquad$
(c) Explain the meaning of the term dynamic equilibrium and the conditions necessary for it to become established.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(d) When 2.00 mol of $\mathrm{PCl}_{5}(\mathrm{~g})$ are decomposed at 450 K and $1.00 \times 10^{5} \mathrm{~Pa}$ the resulting equilibrium mixture contains 0.800 mol of $\mathrm{Cl}_{2}(\mathrm{~g})$.
(i) Calculate the partial pressure of phosphorus(V) chloride, $\mathrm{pPC} l_{5}$, in this equilibrium mixture.
$p \mathrm{PCl}{ }_{5}=$ $\qquad$
(ii) Write the expression for the equilibrium constant, $K_{\mathrm{p}}$, for the decomposition of $\mathrm{PC} l_{5}(\mathrm{~g})$.
$K_{\mathrm{p}}=$
(iii) The partial pressures of $\mathrm{PCl}_{3}(\mathrm{~g})$ and of $\mathrm{Cl}_{2}(\mathrm{~g})$ in this equilibrium mixture are both $2.86 \times 10^{4} \mathrm{~Pa}$.

Calculate the value of $K_{\mathrm{p}}$ and state its units.

[2]

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5 (c) Hydrogen chloride undergoes a reversible reaction with oxygen.

$$
4 \mathrm{HCl}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{Cl}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
$$

The reaction is carried out at $400^{\circ} \mathrm{C}$ in the presence of a copper(II) chloride catalyst.
(ii) State the type of catalyst used in this reaction. Explain how a catalyst is able to increase the rate of a chemical reaction.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(iii) The reaction exists in dynamic equilibrium.

The reaction was repeated at $1000^{\circ} \mathrm{C}$ and the same pressure.
State and explain the effect on the composition of the equilibrium mixture of the change in temperature.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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The equation for this stage of the Contact Process is shown.

$$
2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g}) \quad \Delta H=-196 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(d) (i) State and explain the effect of increasing temperature on the rate of production of $\mathrm{SO}_{3}$.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) State and explain the effect of increasing temperature on the yield of $\mathrm{SO}_{3}$.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(e) The $\mathrm{SO}_{3}$ produced is converted to sulfuric acid in two stages. In the first stage the $\mathrm{SO}_{3}$ is reacted with concentrated sulfuric acid to produce oleum, $\mathrm{H}_{2} \mathrm{~S}_{2} \mathrm{O}_{7}$. The oleum is then reacted with water to form sulfuric acid.

Suggest an equation for the reaction of oleum, $\mathrm{H}_{2} \mathrm{~S}_{2} \mathrm{O}_{7}$, with water to form sulfuric acid.

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6 (d) 2.00 moles of $\mathrm{SO}_{2}(\mathrm{~g})$ and 2.00 moles of $\mathrm{O}_{2}(\mathrm{~g})$ are sealed in a container with a suitable catalyst, at constant temperature and pressure. The resulting equilibrium mixture contains 1.98 moles of $\mathrm{SO}_{3}(\mathrm{~g})$.
The total volume of the equilibrium mixture is $40.0 \mathrm{dm}^{3}$.

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

(i) Write the expression for the equilibrium constant, $K_{\mathrm{c}}$, for the reaction between $\mathrm{SO}_{2}(\mathrm{~g})$ and $\mathrm{O}_{2}(\mathrm{~g})$ to produce $\mathrm{SO}_{3}(\mathrm{~g})$.
$K_{\mathrm{c}}=$
(ii) Calculate the amount, in moles, of $\mathrm{SO}_{2}(\mathrm{~g})$ and $\mathrm{O}_{2}(\mathrm{~g})$ in the equilibrium mixture.

$$
\begin{aligned}
& \mathrm{SO}_{2}(\mathrm{~g})= \\
& \text { mol } \\
& \mathrm{O}_{2}(\mathrm{~g})= \\
& \text { mol } \\
& \text { [2] }
\end{aligned}
$$

(iii) Use your answers to (d)(i) and (d)(ii) to calculate the value of $K_{\mathrm{c}}$ for this equilibrium mixture. Give the units of $K_{c}$.

$$
\begin{array}{r}
K_{\mathrm{c}}=\ldots . . . . . . . . . . . . . . . . . . . . . . . . . . ~ \\
\text { units }=\ldots . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . . ~ \\
{[3]} \\
{\left[J^{\prime} 18 \mathrm{P} 23 \mathrm{Q} 1\right]}
\end{array}
$$

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7 (c) At a pressure of $2.00 \times 10^{7} \mathrm{~Pa}, 1.00 \mathrm{~mol}$ of nitrogen, $\mathrm{N}_{2}(\mathrm{~g})$, was mixed with 3.00 mol of hydrogen, $\mathrm{H}_{2}(\mathrm{~g})$. The final equilibrium mixture formed contained 0.300 mol of ammonia, $\mathrm{NH}_{3}(\mathrm{~g})$.
(i) Calculate the amounts, in mol , of $\mathrm{N}_{2}(\mathrm{~g})$ and $\mathrm{H}_{2}(\mathrm{~g})$ in the equilibrium mixture.

$$
\begin{aligned}
& \mathrm{N}_{2}(\mathrm{~g})=\ldots . . . . . . . . . . . . . . . . . . . . . . . . . ~ \\
& \mathrm{H}_{2}(\mathrm{~g})=\ldots \mathrm{l} \\
&
\end{aligned}
$$

(ii) Calculate the partial pressure of ammonia, $\mathrm{pNH}_{3}$, in the equilibrium mixture.

Give your answer to three significant figures.

$$
p \mathrm{NH}_{3}=
$$

Pa [3]

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(d) In another equilibrium mixture the partial pressures are as shown.

| substance | partial pressure/Pa |
| :---: | :---: |
| $\mathrm{N}_{2}(\mathrm{~g})$ | $2.20 \times 10^{6}$ |
| $\mathrm{H}_{2}(\mathrm{~g})$ | $9.62 \times 10^{5}$ |
| $\mathrm{NH}_{3}(\mathrm{~g})$ | $1.40 \times 10^{4}$ |

(i) Write the expression for the equilibrium constant, $K_{p}$, for the production of ammonia from nitrogen and hydrogen.
$K_{\mathrm{p}}=$
(ii) Calculate the value of $K_{\mathrm{p}}$ for this reaction.

State the units.

$$
\begin{aligned}
& K_{\mathrm{p}}= \\
& \text { units }=
\end{aligned}
$$

(iii) This reaction is repeated with the same starting amounts of nitrogen and hydrogen. The same temperature is used but the container has a smaller volume.

State the effects, if any, of this change on the yield of ammonia and on the value of $K_{p}$.
effect on yield of ammonia $\qquad$
effect on value of $K_{\mathrm{p}}$

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8 At 450 K phosphorus $(\mathrm{V})$ chloride, $\mathrm{PCl}_{5}(\mathrm{~g})$, decomposes to form phosphorus(III) chloride, $\mathrm{PC} l_{3}(\mathrm{~g})$, and chlorine, $\mathrm{Cl}_{2}(\mathrm{~g})$. A dynamic equilibrium is established as shown.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \quad \Delta H=+124 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(a) The enthalpy change of formation of $\mathrm{PCl}_{3}(\mathrm{~g})$ under these conditions is given.

$$
\Delta H_{\mathrm{f}} \mathrm{PCl}_{3}(\mathrm{~g})=-320 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Calculate the enthalpy change of formation of $\mathrm{PCl}_{5}(\mathrm{~g})$ under these conditions.
Include a sign with your answer.
enthalpy change =
$\qquad$ $\mathrm{kJmol}^{-1}$
(b) (i) State and explain the effect of increasing temperature on the rate of decomposition of $\mathrm{PCl}_{5}(\mathrm{~g})$.
$\qquad$
$\qquad$
$\qquad$
(ii) State and explain the effect of increasing temperature on the percentage of $\mathrm{PCl}_{5}(\mathrm{~g})$ that decomposes.
$\qquad$
$\qquad$
$\qquad$
(c) Explain the meaning of the term dynamic equilibrium and the conditions necessary for it to become established.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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(d) When 2.00 mol of $\mathrm{PCl}_{5}(\mathrm{~g})$ are decomposed at 450 K and $1.00 \times 10^{5} \mathrm{~Pa}$ the resulting equilibrium mixture contains 0.800 mol of $\mathrm{Cl}_{2}(\mathrm{~g})$.
(i) Calculate the partial pressure of phosphorus(V) chloride, $\mathrm{pPC} l_{5}$, in this equilibrium mixture.
$\mathrm{pPCl}_{5}=$ Pa [2]
(ii) Write the expression for the equilibrium constant, $K_{\mathrm{p}}$, for the decomposition of $\mathrm{PC} l_{5}(\mathrm{~g})$.

$$
K_{\mathrm{p}}=
$$

(iii) The partial pressures of $\mathrm{PCl}_{3}(\mathrm{~g})$ and of $\mathrm{Cl}_{2}(\mathrm{~g})$ in this equilibrium mixture are both $2.86 \times 10^{4} \mathrm{~Pa}$.

Calculate the value of $K_{\mathrm{p}}$ and state its units.


## EQUILIBRIA WS 3

## SECTION A

1 For which equilibrium does $K_{c}$ have no units?
A $\mathrm{C}(\mathrm{s})+\mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightleftharpoons \mathrm{CO}(\mathrm{g})+\mathrm{H}_{2}(\mathrm{~g})$
B $\mathrm{CH}_{3} \mathrm{OH}(\mathrm{I})+\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}(\mathrm{I}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{CH}_{3}(\mathrm{I})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I})$
C $\mathrm{Cu}^{2+}(\mathrm{aq})+4 \mathrm{NH}_{3}(\mathrm{aq}) \rightleftharpoons \mathrm{Cu}\left(\mathrm{NH}_{3}\right)_{4}^{2+}(\mathrm{aq})$
D $\quad \mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2}(\mathrm{~g})$

2 When 0.20 mol of hydrogen gas and 0.15 mol of iodine gas are heated at 723 K until equilibrium is established, the equilibrium mixture is found to contain 0.26 mol of hydrogen iodide.

The equation for the reaction is as follows.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g})
$$

What is the correct expression for the equilibrium constant $K_{c}$ ?
A $\frac{2 \times 0.26}{0.20 \times 0.15}$
B $\frac{(2 \times 0.26)^{2}}{0.20 \times 0.15}$
C $\frac{(0.26)^{2}}{0.07 \times 0.02}$
D $\frac{(0.26)^{2}}{0.13 \times 0.13}$
3 Two equilibria are shown below.

| reaction I | $2 X_{2}(\mathrm{~g})+Y_{2}(\mathrm{~g}) \rightleftharpoons 2 X_{2} Y(\mathrm{~g})$ |
| :--- | :--- |
| reaction II | $X_{2} Y(\mathrm{~g}) \rightleftharpoons X_{2}(\mathrm{~g})+\frac{1}{2} Y_{2}(\mathrm{~g})$ |

The numerical value of $K_{c}$ for reaction $I$ is 2 .
Under the same conditions, what is the numerical value of $K_{c}$ for reaction II?
A $\frac{1}{\sqrt{2}}$
B $\quad \frac{1}{2}$
C $\frac{1}{4}$
D -2

4 For the reaction

$$
\mathrm{W}(\mathrm{aq})+2 X(\mathrm{aq}) \rightleftharpoons 2 Y(\mathrm{aq})+3 Z(\mathrm{aq})
$$

what are the correct units for the equilibrium constant $K_{c}$ ?
A $\mathrm{moldm}^{-3}$
B $\mathrm{mol}^{2} \mathrm{dm}^{-6}$
C $\mathrm{mol}^{-1} \mathrm{dm}^{3}$
D $\mathrm{mol}^{-2} \mathrm{dm}^{6}$

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5 A dimer, $X$, is stable when solid but a dynamic equilibrium is set up in solution.

$$
X(a q) \rightleftharpoons 2 Y(a q)
$$

A solution of $X$ has an initial concentration of $0.50 \mathrm{~mol} \mathrm{dm}^{-3}$. When equilibrium has been reached $[\mathrm{X}(\mathrm{aq})]$ has fallen to $0.25 \mathrm{~mol} \mathrm{dm}^{-3}$.

The changes in $[\mathrm{X}(\mathrm{aq})]$ and $[\mathrm{Y}(\mathrm{aq})]$ are plotted against time until equilibrium is reached. The value of $K_{\mathrm{c}}$ is then calculated.

Which graph and value for $K_{c}$ are correct?

|  | graph | $\mathrm{K}_{\mathrm{c}} / \mathrm{mol} \mathrm{dm}^{-3}$ |
| :---: | :---: | :---: |
| A |  | 1 |
| B |  | $2$ |
| C |  | 1 |
| D |  | 2 |

6 In an experiment, $b$ mol of hydrogen iodide were put into a sealed vessel under pressure $p$. At equilibrium, $x \mathrm{~mol}$ of the hydrogen iodide had dissociated, the reaction being represented by the following equation.

$$
2 \mathrm{HI}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})
$$

Which expression for $K_{p}$ is correct?
A $\frac{x^{2}}{(b-x)^{2}}$
B $\frac{x^{2} \mathrm{p}^{2}}{(b-x)^{2}}$
C $\frac{x^{2} \mathrm{p}^{2}}{4 b(b-x)}$
D $\frac{x^{2}}{4(b-x)^{2}}$
7 The value of the equilibrium constant, $K_{c}$, for the reaction to form ethyl ethanoate from ethanol and ethanoic acid is 4.0 at $60^{\circ} \mathrm{C}$.

$$
\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}+\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}+\mathrm{H}_{2} \mathrm{O}
$$

When 1.0 mol of ethanol and 1.0 mol of ethanoic acid are allowed to reach equilibrium at $60^{\circ} \mathrm{C}$, what is the number of moles of ethyl ethanoate formed?
A $\frac{1}{3}$
B $\frac{2}{3}$
C $\frac{1}{4}$
D $\frac{3}{4}$

8 Nitrogen dioxide decomposes on heating according to the following equation.

$$
2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})
$$

When 4 mol of nitrogen dioxide were put into a $1 \mathrm{dm}^{3}$ container and heated to a constant temperature, the equilibrium mixture contained 0.8 mol of oxygen.

What is the value of the equilibrium constant, $K_{c}$, at the temperature of the experiment?
A $\frac{0.8^{2} \times 0.8}{4^{2}}$
B $\frac{1.6 \times 0.8}{2.4^{2}}$
C $\frac{1.6^{2} \times 0.8}{4^{2}}$
D $\frac{1.6^{2} \times 0.8}{2.4^{2}}$

9 Dinitrogen tetroxide dissociates into nitrogen dioxide on heating.

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

In an experiment the partial pressures of the gases at equilibrium were found to be $\mathrm{NO}_{2}, 0.33 \mathrm{~atm}$; $\mathrm{N}_{2} \mathrm{O}_{4}, 0.67$ atm.

What is the numerical value of $K_{p}$ at the temperature of the experiment?
A 0.16
B 0.49
C 0.65
D 2.03

10 At a total pressure of 1.0 atm , dinitrogen tetraoxide is $50 \%$ dissociated at a temperature of $60^{\circ} \mathrm{C}$, according to the following equation.

$$
\mathrm{N}_{2} \mathrm{O}_{4} \rightleftharpoons 2 \mathrm{NO}_{2}
$$

D 2 atm

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11 At high temperatures, steam decomposes into its elements according to the following equation.

$$
2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})
$$

In one experiment at 1 atm pressure, it was found that $20 \%$ of the steam had been converted into hydrogen and oxygen.

What are the values of the equilibrium partial pressures, in atm, of the components of this equilibrium?

|  | partial pressure of steam | partial pressure of hydrogen | partial pressure of oxygen |
| :---: | :---: | :---: | :---: |
| A | $\frac{0.80 \times 1}{1.0}$ | $\frac{0.10 \times 1}{1.0}$ | $\frac{0.10 \times 1}{1.0}$ |
| B | $\frac{0.80 \times 1}{1.0}$ | $\frac{0.133 \times 1}{1.0}$ | $\frac{0.067 \times 1}{1.0}$ |
| C | $\frac{0.80 \times 1}{1.0}$ | $\frac{0.20 \times 1}{1.0}$ | $\frac{0.10 \times 1}{1.0}$ |
| D | $\frac{0.80 \times 1}{1.1}$ | $\frac{0.20 \times 1}{1.1}$ | $\frac{0.10 \times 1}{1.1}$ |

12 When vanadium(II) compounds are dissolved in water, the following equilibrium is established.

$$
\mathrm{V}^{2+}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{~V}^{3+}+\frac{1}{2} \mathrm{H}_{2}+\mathrm{OH}^{-}
$$

What would alter the composition of the equilibrium mixture in favour of the $\mathrm{V}^{2+}$ ions?
A adding an acid
B adding a reagent that selectively precipitates $\mathrm{V}^{3+}$ ions
C allowing the hydrogen to escape as it forms
D making the solution more alkaline
13 Which statement concerning the equilibrium reaction below is true?

$$
2 \mathrm{CrO}_{4}^{2-}(\mathrm{aq})+2 \mathrm{H}^{+}(\mathrm{aq}) \rightleftharpoons \mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I})
$$

A An increase in acid concentration will result in an increase in the concentration of $\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})$.

B A redox reaction is taking place.
C The addition of a catalyst will result in an increase in the concentration of $\mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}(\mathrm{aq})$.
D The equilibrium constant, $K_{\mathrm{c}}$, has no units.
14 For the equilibrium $2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g})$, what will change the value of $K_{p}$ ?
A adding a catalyst
B adding more $\mathrm{O}_{2}$
C increasing the pressure
D increasing the temperature

15 An experiment is set up to measure the rate of hydrolysis of methyl ethanoate.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{CH}_{3}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{CH}_{3} \mathrm{OH}
$$

The hydrolysis is found to be slow in neutral aqueous solution but it proceeds at a measurable rate when the solution is acidified with hydrochloric acid.

What is the function of the hydrochloric acid?
A to dissolve the methyl ethanoate
B to ensure that the reaction reaches equilibrium
C to increase the reaction rate by catalytic action
D to suppress ionisation of the ethanoic acid formed
16 The equilibrium

$$
\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g}) \quad \Delta H=+180 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

contributes to a series of reactions producing photochemical smog.
Which factors would affect the value of $K_{p}$ of the above equilibrium?

|  | change in <br> pressure | change in <br> temperature | presence or absence <br> of a catalyst |
| :---: | :---: | :---: | :---: |
| A | $\checkmark$ | $\checkmark$ | $x$ |
| B | $\checkmark$ | $x$ | $\checkmark$ |
| C | $x$ | $\checkmark$ | $\checkmark$ |
| D | $x$ | $\checkmark$ | $x$ |

17 When sulfur trioxide is manufactured from sulfur dioxide and oxygen, using the Contact process, which condition affects the value of the equilibrium constant, $K_{c}$ ?

A adjusting the temperature
B adjusting the pressure
C using a catalyst
D removing $\mathrm{SO}_{3}$ from the equilibrium mixture
18 The Haber process for the manufacture of ammonia is represented by the following equation.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \quad \Delta H=-92 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which statement is correct about this reaction when the temperature is increased?
A Both forward and backward rates increase.
B The backward rate only increases.
C The forward rate only increases.
D There is no effect on the backward or forward rate.

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19 Swimming pool water can be kept free of harmful bacteria by adding aqueous sodium chlorate(I), NaOCl . This reacts with water to produce HOCl molecules which kill bacteria.

$$
\mathrm{OCl}^{-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{OH}^{-}(\mathrm{aq})+\mathrm{HOCl}(\mathrm{aq})
$$

In bright sunshine, the $\mathrm{OCl}^{-}$ion is broken down by ultra-violet light.

$$
\mathrm{OCl}^{-}(\mathrm{aq})+\text { uv light } \rightarrow \mathrm{Cl}^{-}(\mathrm{aq})+1 / 2 \mathrm{O}_{2}(\mathrm{~g})
$$

Which method would maintain the highest concentration of $\mathrm{HOCl}(\mathrm{aq})$ ?
A acidify the pool water
B add a solution of chloride ions
C add a solution of hydroxide ions
D bubble air through the water
20 The dissociation of dinitrogen tetraoxide into nitrogen dioxide is represented by the equation below.

$$
\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2}(\mathrm{~g}) ; \quad \Delta H^{\ominus}=+57 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

If the temperature of an equilibrium mixture of the gases is increased at constant pressure, will the volume of the mixture increase or decrease and why?

A The volume will increase, but only because of a shift of equilibrium towards the right.
B The volume will increase, both because of a shift of equilibrium towards the right and also because of thermal expansion.

C The volume will stay the same, because any thermal expansion could be exactly counteracted by a shift of equilibrium towards the left.

D The volume will decrease, because a shift of equilibrium towards the left would more than counteract any thermal expansion.
$21 \mathrm{PCl}_{5}$ dissociates as follows.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightarrow \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

The extent of dissociation is $13 \%$ at $160^{\circ} \mathrm{C}$ and $100 \%$ at $300^{\circ} \mathrm{C}$.
Which pair of statements about this formation of $\mathrm{PCl}_{3}$ is correct?

|  | shape of $\mathrm{PC}_{3}$ molecule | the reaction is |
| :---: | :---: | :---: |
| A | pyramidal | endothermic |
| B | pyramidal | exothermic |
| C | trigonal | endothermic |
| D | trigonal | exothermic |

22 In an industrial process, two gases $X$ and $Y$ react together to form a single gaseous product $Z$.

$$
X(\mathrm{~g})+Y(\mathrm{~g}) \rightleftharpoons Z(\mathrm{~g})
$$

The percentage yield of product $Z$ varies according to the pressure and the temperature as shown in the graphs.


Which statement about this equilibrium reaction is correct?
A Decreasing the temperature decreases the value of the equilibrium constant.
B Decreasing the temperature increases the rate of this reaction.
C Increasing the pressure increases the value of the equilibrium constant.
D The reaction is exothermic in the forward direction.

23 The percentage of ammonia obtainable, if equilibrium were established during the Haber process, is plotted against the operating pressure for two temperatures, $400^{\circ} \mathrm{C}$ and $500^{\circ} \mathrm{C}$.

Which diagram correctly represents the two graphs?


C



D


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24 The table gives the concentrations and pH values of the aqueous solutions of two compounds, X and $Y$. Either compound could be an acid or a base.

|  | $X$ | $Y$ |
| :--- | :---: | :---: |
| concentration | $2 \mathrm{moldm}^{-3}$ | $2 \mathrm{moldm}^{-3}$ |
| pH | 6 | 9 |

Student $P$ concluded that $X$ is a strong acid.
Student $Q$ concluded that the extent of dissociation is lower in $X(a q)$ than in $Y(a q)$.
Which of the students are correct?
A both P and Q
B neither P nor Q
C P only
D Q only
25 Which substance, in $1 \mathrm{~mol} \mathrm{dm}^{-3}$ aqueous solution, would have the same hydrogen ion concentration as $1 \mathrm{~mol} \mathrm{dm}^{-3}$ of hydrochloric acid?

A ethanoic acid
B nitric acid
C sodium hydroxide
D sulphuric acid

26 Why is ethanoic acid a stronger acid in liquid ammonia than in aqueous solution?
A Ammonia is a stronger base than water.
B Ammonium ethanoate is completely ionised in aqueous solution.
C Ammonium ethanoate is strongly acidic in aqueous solution.
D Liquid ammonia is a more polar solvent than water.
27 The following equilibrium is set up in a mixture of concentrated nitric and sulfuric acids.

$$
\mathrm{HNO}_{3}+\mathrm{H}_{2} \mathrm{SO}_{4} \rightleftharpoons \mathrm{H}_{2} \mathrm{NO}_{3}^{+}+\mathrm{HSO}_{4}^{-}
$$

Which row correctly describes the behaviour of each substance in the equilibrium mixture?

|  | $\mathrm{HNO}_{3}$ | $\mathrm{H}_{2} \mathrm{SO}_{4}$ | $\mathrm{H}_{2} \mathrm{NO}_{3}{ }^{+}$ | $\mathrm{HSO}_{4}{ }^{-}$ |
| :---: | :---: | :---: | :---: | :---: |
| A | acid | acid | base | base |
| B | acid | base | base | acid |
| C | base | acid | acid | base |
| D | base | acid | base | acid |

28 The table shows the enthalpy change of neutralisation per mole of water formed, $\Delta H$, for various acids and bases.

| acid | base | $\Delta H / \mathrm{kJ} \mathrm{mol}^{-1}$ |
| :---: | :---: | :---: |
| hydrochloric acid | sodium hydroxide | -57.0 |
| $\mathbf{P}$ | sodium hydroxide | -54.0 |
| hydrochloric acid | $\mathbf{Q}$ | -52.0 |
| nitric acid | $\mathbf{R}$ | -57.0 |

What are $\mathbf{P}, \mathbf{Q}$ and $\mathbf{R}$ ?

|  | P | $\mathbf{Q}$ | $\mathbf{R}$ |
| :---: | :---: | :---: | :---: |
| A | ethanoic acid | ammonia | potassium hydroxide |
| B | ethanoic acid | sodium hydroxide | ammonia |
| C | sulphuric acid | ammonia | potassium hydroxide |
| D | sulphuric acid | sodium hydroxide | ammonia |

29 The esterification reaction

$$
\text { ethanol + ethanoic acid } \rightleftharpoons \text { ethyl ethanoate + water }
$$

is an equilibrium. The forward reaction is exothermic.
How can the value of the equilibrium constant $K_{\mathrm{c}}$ be increased?
A by adding a little concentrated sulfuric acid as a catalyst
B by increasing the initial concentration of ethanol
C by lowering the temperature
D by raising the temperature

30 An aqueous solution was prepared containing $1.0 \mathrm{~mol}^{\mathrm{m}} \mathrm{AgNO}_{3}$ and $1.0 \mathrm{~mol}^{2}$ of $\mathrm{FeSO}_{4}$ in $1.00 \mathrm{dm}^{3}$ of water. When equilibrium was established, there was 0.44 mol of $\mathrm{Ag}^{+}(\mathrm{aq})$ in the mixture.

$$
\mathrm{Ag}^{+}(\mathrm{aq})+\mathrm{Fe}^{2+}(\mathrm{aq}) \rightleftharpoons \mathrm{Ag}(\mathrm{~s})+\mathrm{Fe}^{3+}(\mathrm{aq})
$$

What is the numerical value of $K_{\mathrm{c}}$ ?
A 0.35
B 0.62
C 1.62
D 2.89

31 For the equilibrium $2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g})$, what will change the value of $K_{\mathrm{p}}$ ?
A adding a catalyst
B adding more $\mathrm{O}_{2}$
C increasing the pressure
D increasing the temperature

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32 When gaseous iodine is heated with hydrogen at $450^{\circ} \mathrm{C}$, an equilibrium is established.

$$
\underset{\text { colourless }}{\mathrm{H}_{2}(\mathrm{~g})}+\underset{\text { purple }}{\mathrm{I}_{2}(\mathrm{~g})} \rightleftharpoons \underset{\text { colourless }}{2 \mathrm{HI}(\mathrm{~g})} \quad \Delta H=+53 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which change of conditions will cause the purple colour of the equilibrium mixture to become paler?

A decrease in pressure
B decrease in temperature
C increase in pressure
D increase in temperature
33 For the reaction

$$
W(a q)+2 X(a q) \rightleftharpoons 2 Y(a q)+3 Z(a q)
$$

what are the correct units for the equilibrium constant $K_{c}$ ?
A $\mathrm{moldm}^{-3}$
B $\mathrm{mol}^{2} \mathrm{dm}^{-6}$
C $\mathrm{mol}^{-1} \mathrm{dm}^{3}$
D $\mathrm{mol}^{-2} \mathrm{dm}^{6}$

34 Two moles of compound $P$ were placed in a vessel. The compound $P$ was partly decomposed by heating. A dynamic equilibrium between chemicals $P, Q$ and $R$ was established.

At equilibrium, $x$ mol of $R$ were present and the total number of moles present was $(2+x)$.
What is the equation for this equilibrium?
A $\quad \mathrm{P} \rightleftharpoons 2 \mathrm{Q}+\mathrm{R}$
B $\quad 2 \mathrm{P} \rightleftharpoons 2 \mathrm{Q}+\mathrm{R}$
C $\quad 2 P \rightleftharpoons Q+R$
D $2 P \rightleftharpoons Q+2 R$

35 Methanol is manufactured by reacting carbon dioxide and hydrogen.

$$
\mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad \Delta H=-49 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

What would increase the equilibrium yield of methanol in this process?
A adding a catalyst
B adding an excess of steam
C increasing the pressure
D increasing the temperature

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36 An experiment is set up to measure the rate of hydrolysis of ethyl ethanoate.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}
$$

The hydrolysis is found to be slow in neutral aqueous solution but it proceeds at a measurable rate when the solution is acidified with hydrochloric acid.

What is the function of the hydrochloric acid?
A to dissolve the ethyl ethanoate
B to ensure that the reaction reaches equilibrium
C to increase the reaction rate by catalytic action
D to suppress ionisation of the ethanoic acid formed
37 The reaction between sulfur dioxide and oxygen is a dynamic equilibrium.

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

What happens when the pressure of the system is increased?
A The rate of reaction will decrease and the position of the equilibrium will move to the left.
B The rate of reaction will decrease and the position of the equilibrium will move to the right.
C The rate of reaction will increase and the position of the equilibrium will move to the left.
D The rate of reaction will increase and the position of the equilibrium will move to the right.
38 Nitrogen reacts with hydrogen to produce ammonia.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

A mixture of 2.00 mol of nitrogen, 6.00 mol of hydrogen, and 2.40 mol of ammonia is allowed to reach equilibrium in a sealed vessel of volume $1 \mathrm{dm}^{3}$ under certain conditions. It was found that 2.32 mol of nitrogen were present in the equilibrium mixture.

What is the value of $K_{\mathrm{c}}$ under these conditions?
A $\frac{(1.76)^{2}}{(2.32)(6.96)^{3}}$
B $\frac{(1.76)^{2}}{(2.32)(6.32)^{3}}$
C $\frac{(2.08)^{2}}{(2.32)(6.32)^{3}}$
D $\frac{(2.40)^{2}}{(2.32)(6.00)^{3}}$

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39 Nitrogen reacts with hydrogen to produce ammonia.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

A mixture of 1.00 mol of nitrogen, 3.00 mol of hydrogen and 1.98 mol of ammonia is allowed to reach equilibrium in a sealed vessel under certain conditions. It was found that 1.64 mol of nitrogen were present in the equilibrium mixture.

What is the value of $K_{\mathrm{c}}$ under these conditions?
A $\frac{(0.70)^{2}}{(1.64)(4.92)^{3}}$
B $\frac{(1.34)^{2}}{(1.64)(3.64)^{3}}$
C $\frac{(1.64)(4.92)^{3}}{(0.70)^{2}}$
D $\frac{(1.64)(3.64)^{3}}{(1.34)^{2}}$
40 The equilibrium constant, $K_{\mathrm{c}}$, for the reaction $\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{g})$, is 60 at $450^{\circ} \mathrm{C}$.
What is the number of moles of hydrogen iodide in equilibrium with 2 mol of hydrogen and 0.3 mol of iodine at $450^{\circ} \mathrm{C}$ ?
A $\frac{1}{100}$
B $\frac{1}{10}$
C 6
D 36

41 Hydrogen and carbon dioxide gases are mixed in equal molar amounts at 800 K . A reversible reaction takes place.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{CO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2} \mathrm{O}(\mathrm{~g})+\mathrm{CO}(\mathrm{~g})
$$

At equilibrium, the partial pressures of $\mathrm{H}_{2}$ and $\mathrm{CO}_{2}$ are both 10.0 kPa . $K_{\mathrm{p}}$ is 0.288 at 800 K .
What is the partial pressure of CO in the equilibrium mixture?
A 5.37 kPa
B $\quad 18.6 \mathrm{kPa}$
C 28.8 kPa
D 347 kPa

42 The formation of hydrogen and ethyne, $\mathrm{C}_{2} \mathrm{H}_{2}$, from methane reaches dynamic equilibrium.

$$
2 \mathrm{CH}_{4}(\mathrm{~g}) \rightleftharpoons 3 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{C}_{2} \mathrm{H}_{2}(\mathrm{~g})
$$

What are the units of $K_{c}$ ?
A $\mathrm{moldm}^{-3}$
B $\mathrm{mol}^{2} \mathrm{dm}^{-6}$
C $\mathrm{mol}^{3} \mathrm{dm}^{-9}$
D $\mathrm{mol}^{4} \mathrm{dm}^{-12}$

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43 One mole of phosphorus $(\mathrm{V})$ chloride, $\mathrm{PCl}_{5}$, is heated to 600 K in a sealed flask of volume $1 \mathrm{dm}^{3}$. Equilibrium is established and measurements are taken.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

The experiment is repeated with one mole of phosphorus $(\mathrm{V})$ chloride heated to 600 K in a sealed flask of volume $2 \mathrm{dm}^{3}$.

How will the measurements vary?
A The equilibrium concentrations of $\mathrm{PCl}_{3}(\mathrm{~g})$ and $\mathrm{Cl}_{2}(\mathrm{~g})$ are higher in the second experiment.
B The equilibrium concentration of $\mathrm{PCl}_{5}(\mathrm{~g})$ is lower in the second experiment.
C The equilibrium concentrations of all three gases are the same in both experiments.
D The value of the equilibrium constant is higher in the second experiment.
44 In which reaction is the underlined substance acting as a base?
A $\mathrm{HNO}_{3}+\underline{\mathrm{H}_{2} \mathrm{SO}_{4}} \rightarrow \mathrm{H}_{2} \mathrm{NO}_{3}{ }^{+}+\mathrm{HSO}_{4}^{-}$
B $\mathrm{HSiO}_{3}^{-}+\underline{\mathrm{HCN}} \rightarrow \mathrm{CN}^{-}+\mathrm{H}_{2} \mathrm{O}+\mathrm{SiO}_{2}$
C $\mathrm{HNO}_{2}+\underline{\mathrm{HCO}_{3}^{-}} \rightarrow \mathrm{H}_{2} \mathrm{O}+\mathrm{CO}_{2}+\mathrm{NO}_{2}^{-}$
D $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}+\mathrm{CH}_{2} \mathrm{ClCO} 2 \mathrm{H} \rightarrow \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{OH}+\mathrm{CH}_{2} \mathrm{ClCO}_{2}^{-}$
45 One molecule of haemoglobin, Hb , can bind with four molecules of oxygen according to the following equation.

$$
\mathrm{Hb}(\mathrm{aq})+4 \mathrm{O}_{2}(\mathrm{aq}) \rightleftharpoons \mathrm{Hb}\left(\mathrm{O}_{2}\right)_{4}(\mathrm{aq})
$$

When the equilibrium concentration of $\mathrm{O}_{2}$ is $7.6 \times 10^{-6} \mathrm{~mol} \mathrm{dm}^{-3}$, the equilibrium concentrations of Hb and $\mathrm{Hb}\left(\mathrm{O}_{2}\right)_{4}$ are equal.

What is the value of $K_{\mathrm{c}}$ for this equilibrium?
A $3.0 \times 10^{20}$
B $\quad 1.3 \times 10^{5}$
C $7.6 \times 10^{-6}$
D $\quad 3.3 \times 10^{-21}$

46 Nitrogen dioxide, $\mathrm{NO}_{2}$, exists in equilibrium with dinitrogen tetroxide, $\mathrm{N}_{2} \mathrm{O}_{4}$.

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

$$
\Delta H=-57 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Which conditions give the greatest percentage of $\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})$ at equilibrium?

|  | pressure | temperature |
| :---: | :---: | :---: |
| A | high | high |
| B | high | low |
| C | low | high |
| D | low | low |

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47 When a sample of HI is warmed to a particular temperature the equilibrium below is established.

$$
2 \mathrm{HI}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})
$$

At this temperature, it is found that the partial pressure of $\mathrm{HI}(\mathrm{g})$ is 28 times the partial pressure of $\mathrm{H}_{2}(\mathrm{~g})$.

What is the value of $K_{p}$ at this temperature?
A $1.28 \times 10^{-3}$
B 0.035
C 28
D 784

48 A mixture of nitrogen and hydrogen gases, at a temperature of 500 K , was put into an evacuated vessel of volume $6.0 \mathrm{dm}^{3}$. The vessel was then sealed.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

The mixture was allowed to reach equilibrium. It was found that 7.2 mol of $\mathrm{N}_{2}$ and 12.0 mol of $\mathrm{H}_{2}$ were present in the equilibrium mixture. The value of the equilibrium constant, $K_{\mathrm{c}}$, for this equilibrium is $6.0 \times 10^{-2}$ at 500 K .

What is the concentration of ammonia present in the equilibrium mixture at 500 K ?
A $0.58 \mathrm{moldm}^{-3}$
B $\quad 0.76 \mathrm{~mol} \mathrm{dm}^{-3}$
C $3.5 \mathrm{moldm}^{-3}$
D $27 \mathrm{moldm}^{-3}$
49 Ammonia is made by the Haber process. The reactants are nitrogen and hydrogen.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \quad \Delta H-\mathrm{ve}
$$

What will increase the rate of the forward reaction?
A adding argon to the mixture but keeping the total volume constant
B decreasing the temperature
C increasing the total pressure by reducing the total volume at constant temperature
D removing ammonia as it is made but keeping the total volume of the mixture the same
50 Nitrogen monoxide reacts with oxygen in a reversible reaction according to the equation shown below.

$$
2 \mathrm{NO}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2}(\mathrm{~g})
$$

The partial pressures of each of the components in an equilibrium mixture are shown in the table.

| partial pressure $\mathrm{NO} / \mathrm{kPa}$ | partial pressure $\mathrm{O}_{2} / \mathrm{kPa}$ | partial pressure $\mathrm{NO}_{2} / \mathrm{kPa}$ |
| :---: | :---: | :---: |
| 10 | 30 | 20 |

What is the numerical value of the equilibrium constant, $K_{p}$, for this equilibrium?
A $6.67 \times 10^{-2}$
B $1.33 \times 10^{-1}$
C 7.50
D 15.0

## SECTION B

The responses $\mathbf{A}$ to $\mathbf{D}$ should be selected on the basis of

| A | B | C | D |
| :---: | :---: | :---: | :---: |
| $\mathbf{1 , 2}$ and 3 |  |  |  |
| are |  |  |  |
| correct |  |  |  |$\quad$| $\mathbf{1}$ and $\mathbf{2}$ |
| :---: |
| only are |
| correct |$\quad$| $\mathbf{2}$ and 3 |
| :---: |
| only are |
| correct |$\quad$| $\mathbf{1}$ only |
| :---: |
| is |
| correct |

1 A reversible reaction is catalysed.
Which statements about the effects of the catalyst on this system are correct?
1 The catalyst alters the mechanism of the reaction.
2 The catalyst reduces the energy of activation (the energy barrier) for both the forward and the backward reaction.

3 The catalyst alters the composition of the equilibrium mixture.
2 Two bulbs $\mathbf{R}$ and $\mathbf{S}$, connected by a mercury manometer, are held in a thermostat, as shown. The volume of $\mathbf{R}$ is twice that of $\mathbf{S}$. $\mathbf{R}$ contains gas, $X$, at the same pressure as the nitrogen in $\mathbf{S}$.


When the temperature is increased, which gases in bulb $\mathbf{R}$ would cause the mercury level in the right-hand limb of the manometer to rise?

1 an equilibrium mixture $\mathrm{N}_{2} \mathrm{~F}_{4}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NF}_{2}(\mathrm{~g}) ; \Delta H$ positive
2 an equilibrium mixture $\mathrm{CH}_{3} \mathrm{NC}(\mathrm{g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{CN}(\mathrm{g}) ; \Delta H$ negative
3 nitrogen
3 Hydroxyapatite, $\mathrm{Ca}_{5}\left(\mathrm{PO}_{4}\right)_{3} \mathrm{OH}$, is the main constituent of tooth enamel. In the presence of saliva, the following equilibria exist.

$$
\begin{aligned}
& \mathrm{Ca}_{5}\left(\mathrm{PO}_{4}\right)_{3} \mathrm{OH}(\mathrm{~s}) \rightleftharpoons 5 \mathrm{Ca}^{2+}(\mathrm{aq})+3 \mathrm{PO}_{4}^{3-}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \\
& \mathrm{HPO}_{4}^{2-}(\mathrm{aq}) \rightleftharpoons \mathrm{H}^{+}(\mathrm{aq})+\mathrm{PO}_{4}^{3-}(\mathrm{aq})
\end{aligned}
$$

Which of the following statements help to explain why tooth enamel is dissolved more readily when saliva is acidic?

1 The hydroxide ions are neutralised by the acid.
2 The phosphate ion $\mathrm{PO}_{4}^{3-}(\mathrm{aq})$ accepts $\mathrm{H}^{+}(\mathrm{aq})$
3 Calcium ions react with acids.

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4 Under given conditions, what governs the rate of a forward reaction?
1 the activation energy of the reaction
2 the enthalpy change of the reaction
3 the equilibrium constant of the reaction

5 Carbon monoxide burns readily in oxygen to form carbon dioxide.
What can be deduced from this information?
1 The +4 oxidation state of carbon is more stable than the +2 state.
2 The standard enthalpy change of formation of carbon dioxide is more negative than that of carbon monoxide.

3 The value of the equilibrium constant for the reaction, $2 \mathrm{CO}(\mathrm{g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{CO}_{2}(\mathrm{~g})$, is likely to be high.

6 Phosphorus pentachloride is introduced into an empty gas syringe which has a movable, tightlyfitting plunger. The gas is allowed to expand until equilibrium is reached at a temperature at which the phosphorus pentachloride partially dissociates.

$$
\mathrm{PCl}_{5}(\mathrm{~g}) \rightleftharpoons \mathrm{PCl}_{3}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$



Which statements are correct?
1 The equilibrium pressure inside the syringe will be greater than atmospheric pressure.
2 When the plunger is pushed in the equilibrium adjusts to produce more $\mathrm{PCl}_{5}(\mathrm{~g})$.
3 The volume of gas in the syringe at equilibrium will be greater than if no dissociation had occurred.

7 Catalysts are used in many reversible reactions in the chemical industry. Vanadium(V) oxide is used in this way in the Contact process for the formation of $\mathrm{SO}_{3}$.

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

What effect does vanadium $(\mathrm{V})$ oxide have on this equilibrium?
1 It speeds up the forward reaction.
2 It increases the value of $K_{p}$.
3 It increases the value of $E_{\mathrm{a}}$ for the reverse reaction.

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8 Which statements about the Haber process for the industrial production of ammonia are correct?
1 The equilibrium constant $K_{p}$ increases with pressure.
2 As the temperature increases, the equilibrium constant for the forward reaction becomes smaller.

3 The process is usually carried out at between $450^{\circ} \mathrm{C}$ and $550^{\circ} \mathrm{C}$ at a pressure of at least 150 atm.

9 Which equilibria, in which all species are gaseous, would have equilibrium constants, $K_{p}$, with no units?

1 sulfur dioxide and oxygen in equilibrium with sulfur trioxide
2 hydrogen and iodine in equilibrium with hydrogen iodide
3 carbon monoxide and steam in equilibrium with carbon dioxide and hydrogen
10 Which of the following can act as a Bronsted-Lowry acid?
$1 \mathrm{H}_{3} \mathrm{O}^{+}$
$2 \mathrm{NH}_{4}^{+}$
$3 \mathrm{H}_{2} \mathrm{O}$

11 Which statements are correct in terms of the Brønsted-Lowry theory of acids and bases?
1 Water can act as either an acid or a base.
2 Sulfuric acid, $\mathrm{H}_{2} \mathrm{SO}_{4}$, does not behave as an acid when dissolved in ethanol, $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$.
3 The ammonium ion acts as a base when dissolved in liquid ammonia.
12 Concentrated sulphuric acid behaves as a strong acid when it reacts with water.

$$
\mathrm{H}_{2} \mathrm{SO}_{4}(\mathrm{I})+\mathrm{aq} \rightarrow \mathrm{H}^{+}(\mathrm{aq})+\mathrm{HSO}_{4}^{-}(\mathrm{aq})
$$

The $\mathrm{HSO}_{4}^{-}$ion formed behaves as a weak acid.

$$
\mathrm{HSO}_{4}^{-}(\mathrm{aq}) \rightleftharpoons \mathrm{H}^{+}(\mathrm{aq})+\mathrm{SO}_{4}^{2-}(\mathrm{aq})
$$

Which statements are true for $1.0 \mathrm{moldm}^{-3}$ sulphuric acid?
$1\left[\mathrm{H}^{+}(\mathrm{aq})\right]$ is high
2 [ $\left.\mathrm{SO}_{4}^{2-}(\mathrm{aq})\right]$ is high
$3 \quad\left[\mathrm{HSO}_{4}^{-}(\mathrm{aq})\right]=\left[\mathrm{SO}_{4}^{2-}(\mathrm{aq})\right]$

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13
The equilibrium constant $K_{p}$ for the reaction

$$
X(\mathrm{~g})+Y(\mathrm{~g}) \rightleftharpoons Z(\mathrm{~g})
$$

varies with temperature as shown in the diagram below.


Which conclusions can be drawn from this information?
1 The reaction is exothermic in the forward direction.
2 The equilibrium mixture contains a greater proportion of $Z$ at higher pressures.
3 The equilibrium mixture contains a greater proportion of $Z$ at higher temperatures.
14 What are necessary properties of a dynamic equilibrium?
1 Equal amounts of reactants and products are present.
2 Concentrations of reactants and products remain constant.
3 The rate of the forward reaction is the same as the rate of the reverse reaction.
15 If $\mathrm{N}_{2} \mathrm{O}_{4}$ gas is placed in a sealed vessel the following equilibrium is established.

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

The forward reaction is endothermic.
What happens when the temperature is increased?
1 The equilibrium constant increases.
2 The partial pressure of $\mathrm{NO}_{2}$ increases.
3 The activation energy is unchanged.

## EQUILIBRIA WS 4

1 Hydrogen iodide dissociates into its elements according to the equation below.

$$
2 \mathrm{HI}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g})
$$

(a) Write the expression for the equilibrium constant, $K_{c}$.
(b) At $120^{\circ} \mathrm{C}$ the equilibrium mixture contains $1.47 \mathrm{~mol} \mathrm{dm}^{-3}$ of $\mathrm{HI}(\mathrm{g}), 0.274 \mathrm{~mol} \mathrm{dm}^{-3}$ each of $\mathrm{H}_{2}(\mathrm{~g})$ and $\mathrm{I}_{2}(\mathrm{~g})$.

Calculate the value of $K_{c}$ for the equilibrium at $120^{\circ} \mathrm{C}$.
(c) Suggest and explain why it would be more difficult to determine $K_{c}$ for this equilibrium at room temperature.
$\qquad$
$\qquad$
$\qquad$
$\qquad$

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2 NO is also formed when nitrosyl chloride, NOCl, dissociates according to the following equation.

$$
2 \mathrm{NOCl}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

Different amounts of the three gases were placed in a closed container and allowed to come to equilibrium at $230^{\circ} \mathrm{C}$. The experiment was repeated at $465^{\circ} \mathrm{C}$.

The equilibrium concentrations of the three gases at each temperature are given in the table below.

|  | concentration $/ \mathrm{mol} \mathrm{dm}^{-3}$ |  |  |
| :---: | :---: | :---: | :---: |
| temperature $/{ }^{\circ} \mathrm{C}$ | NOCl | NO | $\mathrm{Cl}_{2}$ |
| 230 | $2.33 \times 10^{-3}$ | $1.46 \times 10^{-3}$ | $1.15 \times 10^{-2}$ |
| 465 | $3.68 \times 10^{-4}$ | $7.63 \times 10^{-3}$ | $2.14 \times 10^{-4}$ |

(c) (i) Write the expression for the equilibrium constant, $K_{c}$, for this reaction. Give the units.
(ii) Calculate the value of $K_{c}$ at each of the temperatures given. $230^{\circ} \mathrm{C}$
$465^{\circ} \mathrm{C}$
(iii) Is the forward reaction endothermic or exothermic? Explain your answer.
$\qquad$
$\qquad$
(b) Ethanoic acid, $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}$, reacts with ethanol, $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}$, to produce ethyl ethanoate and water. The reaction is an example of dynamic equilibrium.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}+\mathrm{H}_{2} \mathrm{O}
$$

(i) Explain what is meant by dynamic equilibrium.
$\qquad$
$\qquad$
(ii) Write an expression for the equilibrium constant, $K_{\mathrm{C}}$, for this reaction.
(c) A mixture of 6.0 g of ethanoic acid and 6.0 g of ethanol was added to 4.4 g of ethyl ethanoate and the overall mixture allowed to reach equilibrium. It was found that 0.040 mol of ethanoic acid was present in the equilibrium mixture.
(i) Calculate the number of moles of each compound, both initially and at equilibrium. Place the results in the spaces provided.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}+\mathrm{H}_{2} \mathrm{O}
$$

initially $\qquad$
$\qquad$0.00
at equilibrium 0.040

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(ii) Calculate the equilibrium constant, $K_{\mathrm{c}}$, for the reaction.
(iii) Explain why $K_{\mathrm{c}}$ in this reaction has no units.

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4 The reaction of ethanoic acid with ethanol to form ethyl ethanoate and water is an example of dynamic equilibrium. It is catalysed by the presence of $\mathrm{H}^{+}$ions.

$$
\begin{gathered}
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}(\mathrm{I})+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{I}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}(\mathrm{I})+\mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \\
\Delta H=+14 \mathrm{~kJ} \mathrm{~mol}^{-1}
\end{gathered}
$$

(a) (i) Explain what is meant by dynamic equilibrium.
(ii) On the axes below, draw and label a reaction pathway/energy diagram for both the catalysed and uncatalysed reaction.

Also label your diagram with the $\Delta H$ value given.

(b) (i) State the expression for the equilibrium constant, $K_{\mathrm{c}}$, for this reaction.
(ii) State why, in determining the value of $K_{\mathrm{c}}$ for this reaction, it is only necessary to know the number of moles rather than the concentrations of each substance.
$\qquad$
$\qquad$
(c) In an experiment to determine $K_{c}, 1.00 \mathrm{~mol}$ of ethanoic acid was allowed to reach equilibrium at $60^{\circ} \mathrm{C}$ with 1.00 mol of ethanol in the presence of 0.100 mol of $\mathrm{H}^{+}$ dissolved in 1.00 mol of water. It forms an homogeneous mixture.
(i) At equilibrium, it was found that 0.43 mol of ethanoic acid was present.

How many moles of each of the following are present at equilibrium?
ethanol $\qquad$
ethyl ethanoate $\qquad$
water
(ii) Use these values to calculate the equilibrium constant, $K_{\mathrm{c}}$.

5 Alcohols and esters are important organic compounds which are widely used as solvents.
Esters such as ethyl ethanoate can be formed by reacting carboxylic acids with alcohols.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{C}_{2} \mathrm{H}_{5}+\mathrm{H}_{2} \mathrm{O}
$$

This reaction is an example of a dynamic equilibrium.
(a) Explain what is meant by the term dynamic equilibrium.
$\qquad$
$\qquad$
(b) Write the expression for the equilibrium constant for this reaction, $K_{c}$.
(c) For this equilibrium, the value of $K_{c}$ is 4.0 at 298 K .

A mixture containing 0.5 mol of ethanoic acid, 0.5 mol ethanol, 0.1 mol ethyl ethanoate and 0.1 mol water was set up and allowed to come to equilibrium at 298 K . The final volume of solution was $\mathrm{Vdm}{ }^{3}$.

Calculate the amount, in moles, of each substance present at equilibrium.

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6 Ethanoic acid can be reacted with alcohols to form esters, an equilibrium mixture being formed.

$$
\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}+\mathrm{ROH} \rightleftharpoons \mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{R}+\mathrm{H}_{2} \mathrm{O}
$$

The reaction is usually carried out in the presence of an acid catalyst.
(a) Write an expression for the equilibrium constant, $K_{\mathrm{c}}$, for this reaction, clearly stating the units.
$K_{\mathrm{c}}=$
units
In an experiment to determine $K_{\mathrm{c}}$ a student placed together in a conical flask 0.10 mol of ethanoic acid, 0.10 mol of an alcohol ROH , and 0.005 mol of hydrogen chloride catalyst.
The flask was sealed and kept at $25^{\circ} \mathrm{C}$ for seven days.
After this time, the student titrated all of the contents of the flask with $2.00 \mathrm{moldm}^{-3} \mathrm{NaOH}$ using phenolphthalein indicator.
At the end-point, $22.5 \mathrm{~cm}^{3}$ of NaOH had been used.
(b) (i) Calculate the amount, in moles, of NaOH used in the titration.
(ii) What amount, in moles, of this NaOH reacted with the hydrogen chloride?
(iii) Write a balanced equation for the reaction between ethanoic acid and NaOH .
(iv) Hence calculate the amount, in moles, of NaOH that reacted with the ethanoic acid.

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(c) (i) Use your results from (b) to calculate the amount, in moles, of ethanoic acid present at equilibrium. Hence complete the table below.

|  | $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{H}$ | ROH | $\mathrm{CH}_{3} \mathrm{CO}_{2} \mathrm{R}$ | $\mathrm{H}_{2} \mathrm{O}$ |
| :--- | :---: | :---: | :---: | :---: |
| initial <br> amount $/ \mathrm{mol}$ | 0.10 | 0.10 | 0 | 0 |
| equilibrium <br> amount $/ \mathrm{mol}$ |  |  |  |  |

(ii) Use your results to calculate a value for $K_{\mathrm{c}}$ for this reaction.
(d) Esters are hydrolysed by sodium hydroxide. During the titration, sodium hydroxide reacts with ethanoic acid and the hydrogen chloride, but not with the ester.

Suggest a reason for this.
$\qquad$
$\qquad$
(e) What would be the effect, if any, on the amount of ester present if all of the water were removed from the flask and the flask kept for a further week at $25^{\circ} \mathrm{C}$ ?

Explain your answer.
$\qquad$
$\qquad$
$\qquad$

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7 Under certain conditions the equilibrium pressures of the three gases are

| nitrogen | 44.8 atm, |
| :--- | ---: |
| hydrogen | 105.6 atm, |
| ammonia | 37.2 atm. |

(i) Write an expression for the equilibrium constant, $K_{\mathrm{p}}$, for the Haber Process.
(ii) Calculate $K_{\mathrm{p}}$ from these data, giving the units.

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8 Concern over the ever-increasing use of fossil fuels has led to many suggestions for alternative sources of energy. One of these, suggested by Professor George Olah, winner of a Nobel Prize in chemistry, is to use methanol, $\mathrm{CH}_{3} \mathrm{OH}$, which can be obtained in a number of different ways.

Methanol could be used instead of petrol in a conventional internal combustion engine or used to produce electricity in a fuel cell.
(a) Construct a balanced equation for the complete combustion of methanol.

When hydrocarbon fuels are completely burned in an internal combustion engine, several toxic pollutants may be formed.
(b) State two toxic pollutants that can be produced after complete combustion of a hydrocarbon fuel in an internal combustion engine.
$\qquad$
$\qquad$
Methanol may be manufactured catalytically from synthesis gas, a mixture of $\mathrm{CO}, \mathrm{CO}_{2}$ and $\mathrm{H}_{2}$. The CO is reacted with $\mathrm{H}_{2}$ to form methanol, $\mathrm{CH}_{3} \mathrm{OH}$.

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g}) \quad \Delta H=-91 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(c) From your understanding of Le Chatelier's principle, state two conditions that could be used in order to produce a high yield of methanol.

In each case, explain why the yield would increase.
condition 1
explanation $\qquad$
$\qquad$
condition 2 $\qquad$
explanation $\qquad$

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Carbon monoxide, which can be used to make methanol, may be formed by reacting carbon dioxide with hydrogen.

$$
\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad K_{c}=1.44 \text { at } 1200 \mathrm{~K}
$$

(d) (i) It has been suggested that, on a large scale, this reaction could be helpful to the environment.

Explain, with reasons, why this would be the case.
$\qquad$
$\qquad$
(ii) A mixture containing 0.50 mol of $\mathrm{CO}_{2}, 0.50 \mathrm{~mol}$ of $\mathrm{H}_{2}, 0.20 \mathrm{~mol}$ of CO and 0.20 mol of $\mathrm{H}_{2} \mathrm{O}$ was placed in a $1.0 \mathrm{dm}^{3}$ flask and allowed to come to equilibrium at 1200 K .

Calculate the amount, in moles, of each substance present in the equilibrium mixture at 1200 K .

|  | $\mathrm{CO}_{2}$ | + | $\mathrm{H}_{2}$ | $\rightleftharpoons$ | CO | + |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| $\mathrm{H}_{2} \mathrm{O}$ <br> initial <br> moles | 0.50 |  | 0.50 |  | 0.20 |  |
| 0.20 |  |  |  |  |  |  |

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9 Methanol, $\mathrm{CH}_{3} \mathrm{OH}$, is considered to be a possible alternative to fossil fuels, particularly for use in vehicles.

Methanol can be produced from fossil fuels and from agricultural waste. It can also be synthesised from carbon dioxide and hydrogen.

$$
\mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
$$

The synthesis of methanol is carried out at about 500 K with a pressure of between 40 and 100 atmospheres (between $4 \times 10^{6} \mathrm{~Pa}$ and $10 \times 10^{7} \mathrm{~Pa}$ ) and using a catalyst. The use of such conditions will affect both the rate of reaction and the equilibrium yield.

In the spaces below, explain the effects of higher temperature, higher pressure, and the use of a catalyst on the equilibrium yield of methanol.
higher temperature
effect
explanation $\qquad$
$\qquad$
higher pressure
effect
explanation $\qquad$
$\qquad$
use of catalyst
effect $\qquad$
explanation $\qquad$
$\qquad$

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10
Methanol may be synthesised from carbon monoxide and hydrogen.

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

The operating conditions for this reaction are as follows.

| pressure | 200 atmospheres $\left(2 \times 10^{7} \mathrm{~Pa}\right)$ |
| :--- | :--- |
| temperature | 600 K |
| catalyst | oxides of $\mathrm{Cr}, \mathrm{Cu}$, and Zn |

In the spaces below, explain how each of these conditions affects the rate of formation of methanol.
pressure
$\qquad$
$\qquad$
$\qquad$
temperature
$\qquad$
$\qquad$
$\qquad$
catalyst
$\qquad$
$\qquad$
$\qquad$

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11 Hydrogen iodide can be made by heating together hydrogen gas and iodine vapour. The reaction is incomplete.

$$
\mathrm{H}_{2}(\mathrm{~g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{HI}(\mathrm{~g})
$$

(b) Write an expression for $K_{\mathrm{c}}$ and state the units.
$K_{\mathrm{c}}=$ $\qquad$ units
(c) For this equilibrium, the numerical value of the equilibrium constant $K_{c}$ is 140 at 500 K and 59 at 650 K .

Use this information to state and explain the effect of the following changes on the equilibrium position.
(i) increasing the pressure applied to the equilibrium
$\qquad$
(ii) decreasing the temperature of the equilibrium
$\qquad$
$\qquad$
(d) A mixture of 0.02 mol of hydrogen and 0.02 mol of iodine was placed in a $1 \mathrm{dm}^{3}$ flask and allowed to come to equilibrium at 650 K

Calculate the amount, in moles, of each substance present in the equilibrium mixture at 650 K.

|  | $\mathrm{H}_{2}(\mathrm{~g})$ | + | $\mathrm{I}_{2}(\mathrm{~g})$ |
| ---: | :---: | :---: | :---: |
| initial moles | 0.02 | 0.02 |  |
|  |  |  | $2 \mathrm{HI}(\mathrm{g})$ |
|  |  |  |  |

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12 Hydrogen is the most abundant element in the Universe, although on Earth only very small quantities of molecular hydrogen have been found to occur naturally.

Hydrogen is manufactured on a large scale for use in the chemical industry and is also regarded as a possible fuel to replace fossil fuels in internal combustion engines.
(a) State one large scale use of hydrogen in the chemical industry.

One common way of producing hydrogen on a large scale for use in the chemical industry is by the steam 'reforming' of methane (natural gas), in which steam and methane are passed over a catalyst at $1000-1400 \mathrm{~K}$ to produce carbon monoxide and hydrogen.

$$
\mathrm{CH}_{4}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \quad \Delta H=+206 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(b) Use the information above to state and explain the effect on the equilibrium position of the following changes.
(i) increasing the pressure applied to the equilibrium
$\qquad$
$\qquad$
(ii) decreasing the temperature of the equilibrium
$\qquad$
$\qquad$
(c) What will be the effect on the rate of the reaction of increasing the pressure at which it is carried out? Explain your answer.
$\qquad$
$\qquad$
$\qquad$
(d) Further hydrogen can be obtained by the 'water-gas shift' reaction in which the carbon monoxide produced is reacted with steam.

$$
\mathrm{CO}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g}) \quad K_{\mathrm{c}}=6.40 \times 10^{-1} \text { at } 1100 \mathrm{~K}
$$

A mixture containing 0.40 mol of $\mathrm{CO}, 0.40 \mathrm{~mol}$ of $\mathrm{H}_{2} \mathrm{O}, 0.20 \mathrm{~mol}^{2} \mathrm{CO}_{2}$ and 0.20 mol of $\mathrm{H}_{2}$ was placed in a $1 \mathrm{dm}^{3}$ flask and allowed to come to equilibrium at 1100 K
(i) Give an expression for $K_{\mathrm{c}}$ for this reaction.
(ii) Calculate the amount, in moles, of each substance present in the equilibrium mixture at 1100 K .

|  | $\mathrm{CO}(\mathrm{g})$ | + | $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$ | $\rightleftharpoons$ | $\mathrm{CO}_{2}(\mathrm{~g})$ |
| :--- | :---: | :---: | :---: | :---: | :---: |
| initial moles | 0.40 | 0.40 |  | $\mathrm{H}_{2}(\mathrm{~g})$ |  |
| in |  | 0.20 |  | 0.20 |  |

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13 Ammonium nitrate fertiliser is manufactured from ammonia. The first reaction in the manufacture of the fertiliser is the catalytic oxidation of ammonia to form nitrogen monoxide, NO. This is carried out at about $1 \times 10^{3} \mathrm{kPa}$ (10 atmospheres) pressure and a temperature of 700 to $850^{\circ} \mathrm{C}$.

$$
4 \mathrm{NH}_{3}(\mathrm{~g})+5 \mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 4 \mathrm{NO}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad \Delta H^{\ominus}=-906 \mathrm{kJmol}^{-1}
$$

(a) Write the expression for the equilibrium constant, $K_{\mathrm{p}}$, stating the units.
$K_{\mathrm{p}}=$
units
(b) What will be the effect on the yield of NO of each of the following? In each case, explain your answer.
(i) increasing the temperature
$\qquad$
$\qquad$
$\qquad$
(ii) decreasing the applied pressure
$\qquad$
$\qquad$
$\qquad$

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14 Methanol, $\mathrm{CH}_{3} \mathrm{OH}$, can be produced industrially by reacting carbon monoxide, CO , with hydrogen, $\mathrm{H}_{2}$.

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g}) \quad \Delta H=-91 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The process is carried out at $4 \times 10^{3} \mathrm{kPa}$ ( 40 atmospheres) and 1150 K .
(a) (i) State Le Chatelier's Principle.
$\qquad$
$\qquad$
$\qquad$
(ii) From your understanding of Le Chatelier's Principle, state the conditions of temperature and pressure that could be used in order to produce an increased yield of methanol in this process.
In each case, explain why the yield would increase.
temperature $\qquad$
explanation $\qquad$
$\qquad$
pressure
explanation $\qquad$

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(b) The carbon monoxide for use in the production of methanol may be formed by reacting carbon dioxide with hydrogen.

$$
\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \quad K_{\mathrm{c}}=1.44 \text { at } 1200 \mathrm{~K}
$$

A mixture containing 0.70 mol of $\mathrm{CO}_{2}, 0.70 \mathrm{~mol}$ of $\mathrm{H}_{2}, 0.30 \mathrm{~mol}$ of CO and 0.30 mol of $\mathrm{H}_{2} \mathrm{O}$ was placed in a $1 \mathrm{dm}^{3}$ flask and allowed to come to equilibrium at 1200 K .

Calculate the amount, in moles, of each substance present in the equilibrium mixture at 1200 K.

|  | $\mathrm{CO}_{2}+\mathrm{H}_{2} \rightleftharpoons \mathrm{CO}+\mathrm{H}_{2} \mathrm{O}$ |  |  |  |
| :--- | :--- | :--- | :--- | :--- | :--- |
| initial <br> moles | 0.70 | 0.70 | 0.30 | 0.30 |

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15 Nitrogen dioxide, $\mathrm{NO}_{2}$, can enter the atmosphere in a variety of ways.
(a) (i) State one natural and one man-made source of atmospheric $\mathrm{NO}_{2}$. natural $\qquad$ man-made $\qquad$
(ii) Write an equation to show how $\mathrm{NO}_{2}$ leads to the formation of nitric acid in acid rain.
(iii) Use equations to illustrate the catalytic role of $\mathrm{NO}_{2}$ in the formation of sulfuric acid in acid rain.
$\qquad$
$\qquad$
$\qquad$
(b) Nitrogen dioxide exists in equilibrium with dinitrogen tetroxide, $\mathrm{N}_{2} \mathrm{O}_{4}$.

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

2.00 mol of dinitrogen tetroxide was sealed in a container at 350 K . After equilibrium had been established the total pressure was 140 kPa and the mixture of gases contained 1.84 mol of dinitrogen tetroxide.
(i) Give the expression for the equilibrium constant, $K_{\mathrm{p}}$, for this equilibrium.

$$
K_{p}=
$$

(ii) Calculate the number of moles of $\mathrm{NO}_{2}$ present at equilibrium.
(iii) Calculate the total number of moles of gas present at equilibrium and hence the mole fraction of each gas present at equilibrium.

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(iv) Calculate the partial pressure of each gas present at equilibrium.
(v) Calculate the value of the equilibrium constant, $K_{\mathrm{p}}$, at 350 K .

Give your answer to three significant figures and include the units.

$$
\begin{aligned}
& K_{p}= \\
& \text { units }=
\end{aligned}
$$

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16 The Contact process for the manufacture of sulfuric acid was originally patented in the 19th century and is still in use today.

The key step in the overall process is the reversible conversion of sulfur dioxide to sulfur trioxide in the presence of a vanadium $(\mathrm{V})$ oxide catalyst.

$$
2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g}) \quad \Delta H=-196 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(a) One way in which the sulfur dioxide for this reaction is produced is by heating the sulfide ore iron pyrites, $\mathrm{FeS}_{2}$, in air. Iron(III) oxide is also produced. Write an equation for this reaction.
(d) The conversion of sulfur dioxide into sulfur trioxide is carried out at a temperature of $400^{\circ} \mathrm{C}$.
(i) With reference to Le Chatelier's Principle and reaction kinetics, state and explain one advantage and one disadvantage of using a higher temperature.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) State the expression for the equilibrium constant, $K_{\mathrm{p}}$, for the formation of sulfur trioxide from sulfur dioxide.
$K_{\mathrm{p}}=$
(iii) 2.00 moles of sulfur dioxide and 2.00 moles of oxygen were put in a flask and left to reach equilibrium.
At equilibrium, the pressure in the flask was $2.00 \times 10^{5} \mathrm{~Pa}$ and the mixture contained 1.80 moles of sulfur trioxide.

Calculate $K_{p}$. Include the units.

$$
K_{\mathrm{p}}=
$$

$\qquad$
$\qquad$

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17 The Haber process for the manufacture of ammonia, $\mathrm{NH}_{3}$, was originally devised at the start of the 20th century and was developed into a full-scale industrial process by Carl Bosch in 1913.

The key step in the process is the reversible reaction of nitrogen and hydrogen in the presence of an iron catalyst.

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g}) \quad \Delta H=-92 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

(a) The hydrogen for this reaction can be formed by reacting methane with steam, during which carbon monoxide is also produced. Write an equation for this reaction.
$\qquad$
(b) Use the Boltzmann distribution shown to explain why a catalyst increases the rate of this reaction.

$\qquad$
$\qquad$
$\qquad$
$\qquad$
(c) Draw a three-dimensional diagram to show the shape of an ammonia molecule. Name this shape and state the bond angle.
shape $\qquad$ bond angle

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(d) The Haber process is typically carried out at a temperature of $400^{\circ} \mathrm{C}$.
(i) With reference to Le Chatelier's Principle and reaction kinetics, state and explain one advantage and one disadvantage of using a higher temperature.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
(ii) State the expression for the equilibrium constant, $K_{p}$, for the formation of ammonia from nitrogen and hydrogen in the Haber process.
$K_{\mathrm{p}}=$
(iii) 2.00 moles of nitrogen and 3.00 moles of hydrogen were put in a vessel and left to reach equilibrium.

At equilibrium, the pressure was $2.00 \times 10^{7} \mathrm{~Pa}$ and the mixture contained 1.60 moles of ammonia.

Calculate $K_{p}$. Include the units.



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18 Dinitrogen tetraoxide, $\mathrm{N}_{2} \mathrm{O}_{4}$, and nitrogen dioxide, $\mathrm{NO}_{2}$, exist in dynamic equilibrium with each other.

$$
\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}_{2}(\mathrm{~g}) \quad \Delta H=+54 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The energy profile for this reaction is shown.

(a) Add labelled arrows to the energy profile to indicate

- the enthalpy change of the reaction, $\Delta H$,
- the activation energy of the forward reaction, $E_{\mathrm{a}}$.
(b) 0.0500 mol of $\mathrm{N}_{2} \mathrm{O}_{4}$ was placed in a sealed vessel of volume $1.00 \mathrm{dm}^{3}$, at a temperature of $50^{\circ} \mathrm{C}$ and a pressure of $1.68 \times 10^{5} \mathrm{~Pa}$. The mass of the resulting equilibrium mixture was 4.606 g .
(i) Calculate the average molecular mass, $M_{r}$, of the resulting equilibrium mixture. Give your answer to three significant figures.

[2]
(ii) The number of moles of $\mathrm{N}_{2} \mathrm{O}_{4}$ that dissociated can be represented by $n$.

State, in terms of $n$, the amount, in moles, of $\mathrm{NO}_{2}$ in the equilibrium mixture.

The number of moles of $\mathrm{N}_{2} \mathrm{O}_{4}$ remaining at equilibrium is $(0.05-n)$.
(iii) State, in terms of $n$, the total amount, in moles, of gas in the equilibrium mixture.
(iv) State, in terms of $n$, the mole fraction of $\mathrm{NO}_{2}$ in the equilibrium mixture.

In this equilibrium mixture, the mole fraction of $\mathrm{NO}_{2}$ is 0.400 .
(v) Use your answers to (ii) and (iv) to calculate the amount in moles of each gas in the equilibrium mixture. Give your answers to three significant figures.
amount of $\mathrm{N}_{2} \mathrm{O}_{4}=$
mol
amount of $\mathrm{NO}_{2}=$
mol
(vi) Write the expression for the equilibrium constant, $K_{\mathrm{p}}$, for this equilibrium.
$K_{\mathrm{p}}=$
(vii) Use the total pressure of the mixture, $1.68 \times 10^{5} \mathrm{~Pa}$, to calculate the value of the equilibrium constant, $K_{\mathrm{p}}$, and give its units.

$$
\begin{aligned}
K_{p} & =\ldots . \ldots . . . . . . . . . . . . . . . . . . . . . . . ~
\end{aligned} .
$$

