

<u>Home</u> <u>Gameboard</u> Chemistry Physical Kinetics Competing Reactions

# **Competing Reactions**



A compound X can produce two possible products, Y and Z, under the same reaction conditions. An energy profile for this reaction can be represented as shown below.

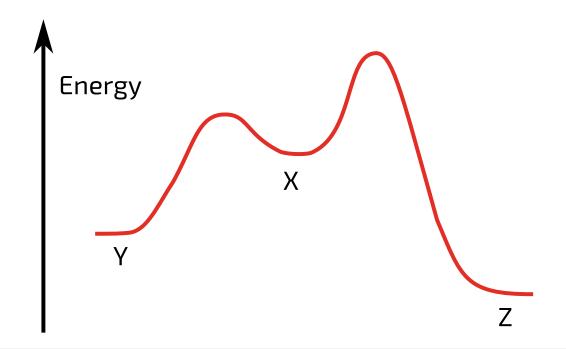


Figure 1: A reaction profile for the reaction described.

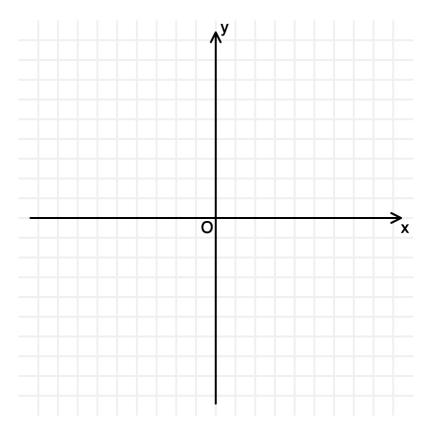
### Part A Activation energy

The activation energy is the	energy required for a rea	action to take place. It is given by the	
difference in energy between the	and the	state. In the set-up above, the reaction	
producing product Y has a	activation energy than the	tivation energy than the process producing product ${f Z}$ due to the	
state being	in energy for the former.		
Items:   minimum maximum optima	al starting materials products	transition higher lower	

#### Part B Molecular speed distribution

What do we call the distribution of molecular speeds?

What does this distribution look like (assume particle speed is on the x-axis, and number of particles on the y-axis)?



#### Part C Temperature dependence

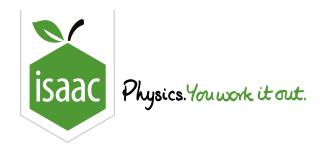
Which product (Y or Z) will be favoured at lower temperatures and which will be favoured at higher temperatures? Explain your reasoning. At lower temperatures, the reaction is under control, and so the reaction with the lower is favoured. This means product is favoured at low temperatures. At higher control, and so the reaction with the lower temperatures, the reaction is under is favoured. This means product is favoured at high temperatures. Items:  $\mathbf{Z}$  $\mathbf{Y}$ kinetic thermodynamic energy of products energy of reactants activation energy

### Part D Gold catalyst

ature at which the reaction is carried out. Which of the following statements help(s) to explain this tion?
Gold increases the activation energy for the formation of Y
Gold decreases the activation energy for the formation of Y
Gold increases the activation energy for the formation of Z
Gold decreases the activation energy for the formation of Z
Gold lowers the energy of X
Gold lowers the energy of Y
Gold lowers the energy of Z

The addition of a gold catalyst to the above reaction results in the formation only of Z, regardless of the

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<u>Home</u> <u>Gameboard</u>

Chemistry

Physical

Kinetics

Ammonia Decomposition

# **Ammonia Decomposition**



In an experiment, the decomposition of gaseous ammonia to  $N_2$  and  $H_2$  on a hot tungsten wire was studied by monitoring the change in pressure with time. The rate equation for this reaction may be written in terms of either the concentration or the partial pressure of ammonia. In terms of the partial pressure of ammonia,  $p_{\rm NH_3}$ , it is:

$$\mathrm{rate} = -k(p_{\mathrm{NH_3}})^{lpha},$$

where  $\alpha$  is the order of the reaction and k is the rate constant.

At the start of the reaction, the only gas present was ammonia at an initial pressure of  $200 \,\mathrm{mmHg}$ . The table below shows how the total pressure varied with time.

Time / s	Total pressure / $\mathrm{mmHg}$
100	214
200	227
300	238
400	249
500	259
600	270
800	292

#### Part A Equation

Write a balanced equation, including state symbols, for the decomposition of one mole of ammonia.

#### Part B Pressure vs concentration

Using the ideal gas law, we can show that pressure is directly proportional to concentration at constant temperature. What is the proportionality constant we need to multiply concentration by to get the pressure?

The following symbols may be useful: R, T, V, n, p

#### Part C Total pressure

The initial partial pressure of ammonia is  $p_0$  mmHg. After a time t it is  $(p_0 - x)$  mmHg. Find an expression for the *total* pressure of reactants and products,  $p_T$ , at time t, expressed in mmHg (don't include the unit in your expression).

The following symbols may be useful:  $p_0$ , t, x

#### Part D Partial pressure

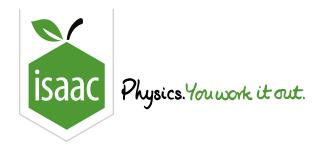
Find an expression for the *partial pressure of ammonia* at time t as a function of  $p_0$  and  $p_T$ , expressed in mmHg (don't include the unit in your expression).

The following symbols may be useful:  $p_0$ ,  $p_T$ 

Part E	Graph
Which o	f the following statements are true about the graph of the partial pressure of ammonia against time?
	It appears to be approximately a straight line.
	It appears to be approximately a parabola
	It appears to be approximately S-shaped
	It has a positive gradient throughout
	It has zero gradient throughout
	It has a negative gradient throughout
	It has both a positive and a negative gradient over different intervals.
	The graph is steepest at around $t=400\mathrm{s}$
	The graph gets steeper at higher times
Part F	Order
What is	the order of the reaction, $\alpha$ ?
Part G	Rate constant
What is	the rate constant of the reaction? Give your answer to 2 s.f.
Adapted with Gameboard:	permission from OCR, STEP Chemistry, Jun 1999, Question 2
Janichudiu.	

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**STEM SMART Chemistry Week 49** 



<u>Home</u> <u>Gameboard</u> Chemistry Physical Kinetics A Rusty Carillon

## A Rusty Carillon



An excavated Chinese ancient bronze musical instrument, carillon, was covered entirely by rust. Chemical analysis showed that the rust contains CuCl,  $Cu_2O$  and  $Cu_2(OH)_3Cl$ . Simulation experiments showed that CuCl was formed first under the action of both air and Cl containing aqueous solution and then  $Cu_2(OH)_3Cl$  produced in the following way:

$$Cu \longrightarrow CuCl \longrightarrow Cu_2(OH)_3Cl$$

Rate constants  $k_c$  for this reaction were measured at various temperatures in a simulation experiment in order to obtain its kinetic parameters. The results of the experiment are given below.

Temperature / $^{\circ}\mathrm{C}$	$k_c$ / $ m moldm^{-3}s^{-1}$
25	$1.29  imes 10^{-4}$
40	$2.50 imes10^{-4}$

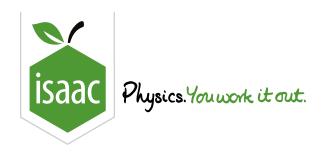
#### Part A Activation energy

Find the value of the activation energy of this reaction.

Adapted from the International Chemistry Olympiad, Beijing 1995, Problem 1.2

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meboard Chemistry

Physical

Entropy

Essential Pre-Uni Chemistry G2.2

# Essential Pre-Uni Chemistry G2.2



Use the following standard molar entropy values in  $J\,K^{-1}\,\mathrm{mol}^{-1}$  to help answer the questions in this section.

$ m H_2O\left(l ight)$	69.9	$\mathrm{HCl}(\mathrm{g})$	186.8	$\mathrm{NaCl}\left(\mathrm{s}\right)$	72.1
$ m H_2O\left(g ight)$	188.7	$\mathrm{Cl}_{2}\left( \mathrm{g} ight)$	223.1	$\rm ZnCl_2\left(s\right)$	111.5
$\mathrm{H}_{2}\left( \mathrm{g}\right)$	130.7	$\mathrm{H}_{2}\mathrm{SO}_{4}\left(\mathrm{l} ight)$	156.9	$\mathrm{Zn}\left( \mathrm{s}\right)$	41.6
Na(s)	51.2	${ m Zn}({ m g})$	150.0	$\mathrm{NaHSO_4}(\mathrm{s})$	113.0
$O_2(g)$	205.2	$\mathrm{CO}_{2}\left( \mathrm{g} ight)$	213.6	$\mathrm{C}\left(\mathrm{s}\right)$ graphite	5.7

Calculate the standard entropy change when...

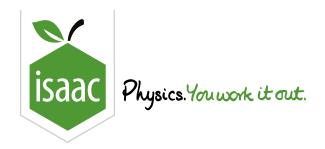
### Part A (a)

 $2.50\,\mathrm{mol}$  of solid zinc chloride decomposes into its gaseous elements.

### Part B (b)

 $2.0\,\mathrm{g}$  of sodium reacts fully with chlorine gas.

Part C (c)
$10.0\mathrm{mg}$ of zinc vapour deposits onto a surface.
Part D (d)
$40.0\mathrm{m}^3$ (at RTP) of steam condenses. Give your answer to 3 significant figures.
Part E (e)
$200\mathrm{cm^3}$ of water $(1.00\mathrm{gcm^{-3}})$ is electrolysed.
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Home Gameboard Chemistry Physical Entropy Essential Pre-Uni Chemistry H1.3

## Essential Pre-Uni Chemistry H1.3



Pa	rt	Α	(a)
			<b>( -</b> . <i>)</i>

A phase change at constant pressure has an enthalpy change of  $58\,\mathrm{kJ\,mol^{-1}}$  and an entropy change of  $85\,\mathrm{J\,K^{-1}\,mol^{-1}}$ . Find the temperature at which the entropy change of the surroundings is equal in magnitude and opposite in sign to that of the system (i.e. the temperature at which the phase change occurs at this pressure.)

#### Part B (b)

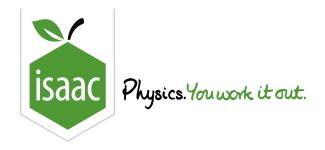
The neutralisation of one mole of acid at  $305\,\mathrm{K}$  increases the entropy of the surroundings by  $180\,\mathrm{J}\,\mathrm{K}^{-1}$  at constant pressure. Find the enthalpy of neutralisation.

#### Part C (c)

The reaction of  $3.5\,\mathrm{g}$  of lithium raises the entropy of the surroundings by  $220\,\mathrm{J\,K^{-1}}$  at constant pressure and  $294\,\mathrm{K}$ . Find the molar enthalpy change for the reaction.

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Home Gameboard Chemistry Physical Entropy Essential Pre-Uni Chemistry H2.1

# Essential Pre-Uni Chemistry H2.1



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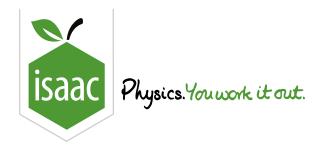
At  $298 \, \mathrm{K}$  the decomposition of hydrogen peroxide has an enthalpy change of  $-98 \, \mathrm{kJ \, mol^{-1}}$  and an entropy change of  $62.9 \, \mathrm{J \, K^{-1} \, mol^{-1}}$ . Calculate the free energy change at constant pressure.

#### Part B (b)

The free energy change for a reaction at constant pressure is  $-4.5 \, \mathrm{kJ \, mol^{-1}}$ . If the temperature is  $320 \, \mathrm{K}$  and the entropy change is  $-12 \, \mathrm{J \, K^{-1} \, mol^{-1}}$ , find the enthalpy change.

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<u>Home</u> <u>Gameboard</u> Chemistry Physical Equilibrium Essential Pre-Uni Chemistry I1.1

# Essential Pre-Uni Chemistry I1.1



Write expressions for  $K_p$  for the following gas phase equilibria, using the notation p(A) for the partial pressure of A:

### Part A (a)

 $CH_3CHCH_2 \mathop{\Longrightarrow}\limits_{} CH_2CH_2CH_2$ 

The following symbols may be useful: C, H, p

#### Part B (b)

 $3\,O_2 \mathop{\Longrightarrow}\nolimits 2\,O_3$ 

The following symbols may be useful: 0, p

### Part C (c)

$$H_2 + I_2 \mathop{\Longrightarrow}\nolimits 2\,HI$$

The following symbols may be useful: H,  $\,$  I,  $\,$  p

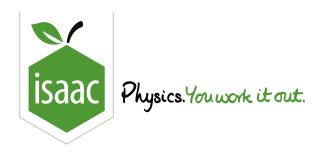
## Part D (d)

$$N_2 + 3\,H_2 \mathop{\Longrightarrow}\nolimits 2\,NH_3$$

The following symbols may be useful: H, N, p

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Chemistry

Physical Equilibri

**Equilibrium** Essential Pre-Uni Chemistry I2.9

# Essential Pre-Uni Chemistry I2.9



The hydration of chloroethanal has an equilibrium constant,  $K_{\rm c}$  , of 37.0~[2].

$$ClCH_{2}CHO\left(aq\right)+H_{2}O\left(l\right) \Longrightarrow ClCH_{2}CH(OH)_{2}\left(aq\right)$$

Complete the following table by providing any missing initial or equilibrium concentrations.

Initial [ClCH <sub>2</sub> CHO]	$\begin{array}{c} \textbf{Initial} \\ [\text{ClCH}_2\text{CH}(\text{OH})_2] \end{array}$	Equilibrium $[ClCH_2CHO]$	$\begin{array}{c} \textbf{Equilibrium} \\ [\text{ClCH}_2\text{CH}(\text{OH})_2] \end{array}$
$1.20\mathrm{moldm^{-3}}$	$0.00\mathrm{moldm^{-3}}$	(a)	(b)
$0.00\mathrm{moldm^{-3}}$	(c)	(d)	$0.292\mathrm{moldm^{-3}}$
(e)	$0.100\mathrm{moldm^{-3}}$	$0.0184\mathrm{moldm^{-3}}$	(f)
$14.0\mathrm{mmoldm^{-3}}$	$0.800\mathrm{mmoldm^{-3}}$	(g)	(h)
$161\mathrm{mgdm^{-3}}$	$0.00\mathrm{mgdm^{-3}}$	(i)	(j)

### Part A (a)

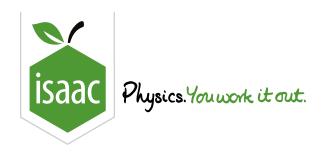
(a) in  $\rm mol\,dm^{-3}$ 

### Part B (b)

(b) in  $m mol\,dm^{-3}$ 

Part C (c)
(c) in $ m moldm^{-3}$
Part D (d)
(d) in $ m moldm^{-3}$
Part E (e)
(e) in $ m moldm^{-3}$
Part F (f)
(f) in $ m moldm^{-3}$
Part G (g)
(g) in $ m mmoldm^{-3}$

Part H (h)
(h) in ${ m mmoldm^{-3}}$
Part I (i)
(i) in ${ m mg}{ m dm}^{-3}$
Part J (j)
(j) in ${ m mg}{ m dm}^{-3}$
Part K (k)
(k) Trichloroethanal is almost fully hydrated in aqueous solution, with a $K_{\rm c}$ value of around $10^4$ [3], to give "chloral hydrate". Give the approximate concentration in ${ m moldm^{-3}}$ of chloral hydrate, (to 2 significant figures), required to maintain an equilibrium concentration of $1.0{ m nmolcm^{-3}}$ of the free, unhydrated form.
[ <b>2</b> ] Tadashi Okuyama, Howard Maskill, 'Organic Chemistry: A Mechanistic Approach' OUP Oxford, $2013;$ ISBN $0199693277, 9780199693276$
${f [3]} Ibid.$
STEM SMART Chemistry Week 49



Home Gameboard Chemistry Physical Acids & Bases Essential Pre-Uni Chemistry J5.10

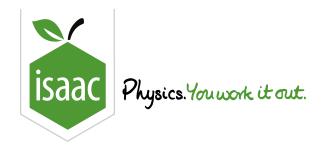
# Essential Pre-Uni Chemistry J5.10



A buffer of pH 7.8 is prepared by taking  $200\,\mathrm{cm^3}$  of  $0.020\,\mathrm{mol\,dm^{-3}}$  "tris" solution and adding dilute hydrochloric acid from a burette until the pH is correct. If this requires  $1.35\,\mathrm{cm^3}$  of  $2.0\,\mathrm{mol\,dm^{-3}}$  HCl(aq), calculate the  $pK_a$  of the conjugate acid of "tris".

Gameboard:

**STEM SMART Chemistry Week 49** 



Home Gameboard Chemistry Physical Acids & Bases pH in Ester Hydrolysis

# pH in Ester Hydrolysis



The hydrolysis of an ester can be carried out in both alkaline and acidic conditions.

### Part A Equations

Fill in the gaps to complete the reaction equations for hydrolysis of ethyl ethanoate under acidic and alkaline conditions respectively.

$$CH_{3}C(=O)OCH_{2}CH_{3} + H_{2}O \xrightarrow{acid} +$$

$$CH_{3}C(=O)OCH_{2}CH_{3} + NaOH \xrightarrow{alkali} +$$

Items:

### Part B pH rate profile

A study of the rate of hydrolysis of ethyl ethanoate in buffered solutions gave the following pH rate profile:

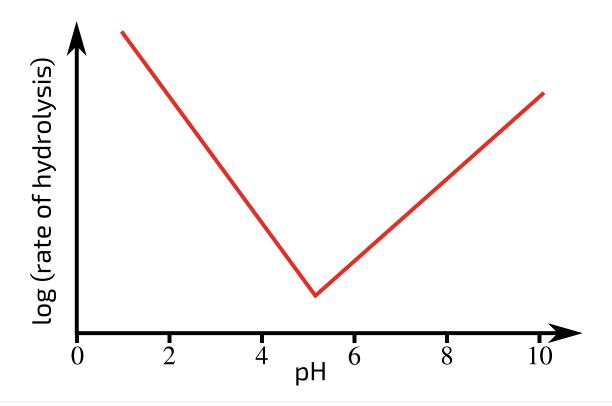


Figure 1: pH rate profile for ester hydrolysis

Which of the following statements help explain the trends seen in this profile?

- at higher pH, the  $\mathrm{H}^+$  concentration is higher, speeding up acid-catalysed hydrolysis
- at higher pH, the OH<sup>-</sup> concentration is higher, speeding up base-catalysed hydrolysis
- at lower pH, the  $\mathrm{H}^+$  concentration is higher, speeding up acid-catalysed hydrolysis
- at lower pH the OH<sup>-</sup> concentration is higher, speeding up base-catalysed hydrolysis

Part C pH 5.5
At pH $5.5$ , the rate for the acid hydrolysis reaction is the same as the rate for the alkaline hydrolysis reaction. What are the concentrations of $\mathrm{H^{+}}$ and $\mathrm{OH^{-}}$ at this pH?
Give your answers to 2 sf.
Concentration of $\mathrm{H}^+$ :
Concentration of $\mathrm{OH}^-$ :
Part D Rate constant comparison
Which of the following statements about the rate constants for the two hydrolysis processes is/are correct?
The rate constants for the acid and alkaline hydrolysis are equal, as the rate for both processes can be made equal under certain conditions.
The rate constant for the acid hydrolysis is higher than the rate constant for the alkaline hydrolysis, as the rate for both processes is equal when the ${\rm H^+}$ concentration is higher than the ${\rm OH^-}$ concentration.
The rate constant for the alkaline hydrolysis is higher than the rate constant for the acid hydrolysis, as the rate for both processes is equal when the $\mathrm{H}^+$ concentration is higher than the $\mathrm{OH}^-$ concentration.
The rate constant for the acid hydrolysis is higher than the rate constant for the alkaline hydrolysis, as the acid hydrolysis dominates at neutral pH when the $\mathrm{H}^+$ and $\mathrm{OH}^-$ concentrations are equal.
The rate constant for the alkaline hydrolysis is higher than the rate constant for the acid hydrolysis, as the alkaline hydrolysis dominates at neutral pH when the $\mathrm{H}^+$ and $\mathrm{OH}^-$ concentrations are equal.

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