

Cambridge AS & A Level

CHEMISTRY Paper 2

Topical Past Paper Questions

+ Answer Scheme

2015 - 2021







Chapter 7

Equilibria





7.1 Chemical equilibria: reversible reactions, dynamic equilibrium

 $26.\ 9701_s21_qp_23\ Q:\ 2$

Methanol, CH₃OH, is soluble in water because it forms hydrogen bonds with water molecules.

(a) Draw a fully labelled diagram to show how a hydrogen bond forms between a water molecule and a methanol molecule.

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[3]

(b) Methanol has a melting point of -97.6 °C and a boiling point of 64.7 °C.

A sample of pure liquid methanol is added to a flask and then sealed. The sealed flask is left for several days at constant temperature. The vapour pressure is then measured as 17 kPa.

(1)	Describe what is meant by the term <i>vapour pressure of methanol</i> .
	[2]
(ii)	Explain why some of the liquid becomes a vapour.
	[1]
(iii)	Suggest and explain why the vapour pressure of water at room temperature is lower than the vapour pressure of methanol at room temperature. Refer to the correct intermolecular forces in your answer.
	[O



(c) Methanol is made by reacting carbon monoxide with hydrogen.

$$CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$$

Carbon monoxide and hydrogen react at $1.0 \times 10^7 \, \text{Pa}$ and $200 \, ^{\circ}\text{C}$. Eventually the reaction mixture reaches dynamic equilibrium.

The table shows the amounts of each species present in the mixture.

	CO(g)	H ₂ (g)	CH ₃ OH(g)
initial amount/mol	1.0	2.0	0
equilibrium amount/mol	0.030	0.060	0.97

(i)	Explain what is meant by dynamic equilibrium	m.
		[2]
(ii)	Calculate the partial pressure of methanol v Show your working.	apour at equilibrium under these conditions.
		an
	-0	Pa [2]
(iii)	Write an expression for the equilibrium con your answer.	stant, K_p , for this reaction. State the units in
	K _p =	
	***	units = [2]
		[Total: 14]





 $27.\ 9701_s18_qp_23\ Q:\ 1$

The elements sodium to chlorine, in the third period, all form oxides.

(a) Draw a diagram to show the shape of the molecule of each of the oxides, SO_3 and Cl_2O . Name each shape.

In SO₃ each oxygen atom forms a double bond with the sulfur atom.

SO ₃	C <i>l</i> ₂ O

(b) (i) Explain why the melting point of MgO is higher than that of Na₂O.

[2]

(ii) Explain why the melting point of SiO₂ is much higher than that of SO₃.



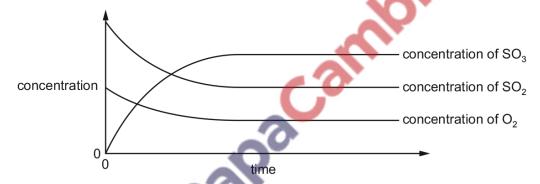


$$2SO_2(g) + O_2(g) \implies 2SO_3(g)$$
 $\Delta H = -196 \text{ kJ mol}^{-1}$

(i)	Explain why increasing the total pressure, at constant temperature, increases the rate of
	production of SO ₂ and increases the yield of SO ₂ .

rate	 	 			
	 	 	_	0	[4]

The graph shows how the concentrations of all three species in the system change with time for a typical reaction mixture. The gradients of all three lines decrease with time and then level off in this dynamic equilibrium.



(ii) Explain why the gradients of the SO ₂ and O ₃ lines decrease with ti	(ii)	Explain why the	gradients of the	SO, and O, li	nes decrease with tim
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(iii) Explain why all three lines become horizontal.

		[1]

(iv) Suggest a reason why the initial gradient of the SO_2 line is steeper than that of the O_2 line.

[4]





(d) 2.00 moles of SO₂(g) and 2.00 moles of O₂(g) are sealed in a container with a suitable catalyst, at constant temperature and pressure. The resulting equilibrium mixture contains 1.98 moles of $SO_3(g)$.

The total volume of the equilibrium mixture is 40.0 dm³.

$$2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$$

(i) Write the expression for the equilibrium constant, K_c , for the reaction between $SO_2(g)$ and $O_2(g)$ to produce $SO_3(g)$.

$$K_{c} =$$

[1]

[3]

[Total: 22]

(ii) Calculate the amount, in moles, of SO₂(g) and O₂(g) in the equilibrium mixture.

$$SO_2(g) =$$
 mol $O_2(g) =$ mol [2]

(iii) Use your answers to (d)(i) and (d)(ii) to calculate the value of K_c for this equilibrium mixture. Give the units of K_c .





$$28.9701_{w17}_{qp}_{2}$$
 Q: 2

At 450 K phosphorus(V) chloride, $PCl_5(g)$, decomposes to form phosphorus(III) chloride, $PCl_3(g)$, and chlorine, $Cl_2(g)$. A dynamic equilibrium is established as shown.

$$PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$$
 $\Delta H = +124 \text{ kJ mol}^{-1}$

(a) The enthalpy change of formation of $PCl_3(g)$ under these conditions is given.

$$\Delta H_{\rm f} \, PC \, l_3(g) = -320 \, kJ \, mol^{-1}$$

Calculate the enthalpy change of formation of $PCl_5(g)$ under these conditions.

Include a sign with your answer.



(b)	(i)	State and explain the effect of increasing temperature on the rate of decomposition of $PCl_5(g)$.
		[2]
	(ii)	State and explain the effect of increasing temperature on the percentage of $PCl_5(g)$ that decomposes.
		[2]
(c)		plain the meaning of the term <i>dynamic equilibrium</i> and the conditions necessary for it to come established.





(d)	When 2.00 mol of PC $l_5(g)$ are decomposed at 450 K and 1.00×10^5 Pa the resulting equilibrium
	mixture contains 0.800 mol of $Cl_2(g)$.

(i)	Calculate the partial	pressure of pho	sphorus(V) chloride	pPC1	in this ed	nuilibrium	mixture
111	Calculate the partial	pressure of pric	JOHN GOLV	, ci iloride.	$DI \cup \iota_{E}$	111 11113 60	adilibrialir	HINKUIC

$$pPCl_5 =Pa$$
 [2]

(ii) Write the expression for the equilibrium constant, K_p , for the decomposition of $PCl_5(g)$.

[1]

(iii) The partial pressures of $PCl_3(g)$ and of $Cl_2(g)$ in this equilibrium mixture are both $2.86 \times 10^4 Pa$.

Calculate the value of $K_{\scriptscriptstyle D}$ and state its units.



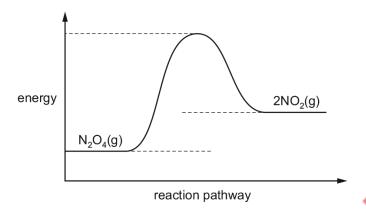
[Total: 12]



Dinitrogen tetraoxide, N₂O₄, and nitrogen dioxide, NO₂, exist in dynamic equilibrium with each other.

$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$
 $\Delta H = +54 \text{ kJ mol}^{-1}$

The energy profile for this reaction is shown.



- (a) Add labelled arrows to the energy profile to indicate
 - the enthalpy change of the reaction, ∆H,
 - the activation energy of the forward reaction, E_a.

[2]

- (b) $0.0500 \, \text{mol}$ of N_2O_4 was placed in a sealed vessel of volume $1.00 \, \text{dm}^3$, at a temperature of $50 \, ^{\circ}\text{C}$ and a pressure of $1.68 \times 10^5 \, \text{Pa}$. The mass of the resulting equilibrium mixture was $4.606 \, \text{g}$.
 - (i) Calculate the average molecular mass, M_r , of the resulting equilibrium mixture. Give your answer to **three** significant figures.



$$M_{r} = \dots [2]$$

(ii) The number of moles of N_2O_4 that dissociated can be represented by n.

State, in terms of n, the amount, in moles, of NO_2 in the equilibrium mixture.

moles of
$$NO_2$$
 =[1]





The number of moles of N_2O_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)	[1] State, in terms of n , the mole fraction of NO_2 in the equilibrium mixture.
	[1]
In t	his equilibrium mixture, the mole fraction of 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 N ₂ O ₄ = mol
	amount of NO_2 = mol [2]
(vi)	Write the expression for the equilibrium constant, K_{p} , for this equilibrium.
	$K_{p} =$ [1]
vii)	Use the total pressure of the mixture, 1.68×10^5 Pa, to calculate the value of the equilibrium constant, K_p , and give its units.
	$\mathcal{K}_{_{\mathrm{p}}}$ =
	units =
	[3]
	[Total: 13]





7.2 Ionic equilibria

$$30.\ 9701_s18_qp_21\ \ Q{:}\ 1$$

Sulfuric acid is manufactured by the Contact process.

One stage in this process is the conversion of sulfur dioxide into sulfur trioxide in the presence of a heterogeneous catalyst of vanadium(V) oxide, V_2O_5 .

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2O=S=O(g) + O=O(g)
$$\implies$$
 2O=S=O(g) $\Delta H = -196 \text{ kJ mol}^{-1}$

(a)	(i)	State the effect of a catalyst on a reaction. Explain how a catalyst causes this effect.
		[2]
	(ii)	State the meaning of the term <i>heterogeneous</i> as applied to catalysts.
		[1 ¹]

(b) Some bond energies are given.

bond	bond energy/kJ mol ⁻¹		
S=O (in SO ₂)	534		
0=0	496		

Use the data, and the enthalpy change for the conversion of sulfur dioxide into sulfur trioxide, to calculate a value for the S=O bond energy in SO₃.

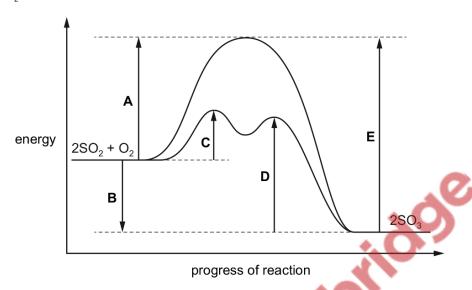
S=O bond energy in
$$SO_3 = \dots kJ mol^{-1}$$
 [2]





The Contact process is usually carried out at a temperature of about 400 °C and a pressure just above atmospheric pressure. Using a higher or lower temperature and pressure would affect both the rate of production of sulfur trioxide and the yield of sulfur trioxide.

(c) A reaction pathway diagram for both the catalysed and uncatalysed reactions between SO_2 and O_2 is shown.



The letters A-E represent energy changes.

Complete the table by stating which letter, A–E, represents the energy change described.

energy change	letter
the energy change for the production of SO ₃	
the activation energy for the production of SO_3 in the absence of a catalyst	
the activation energy for the first step in the decomposition of SO ₃ in the presence of a catalyst	

[3]





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

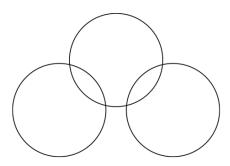
		$2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$ $\Delta H = -196 \text{ kJ mol}^{-1}$
(d)	(i)	State and explain the effect of increasing temperature on the rate of production of SO ₃ .
		[3]
((ii)	State and explain the effect of increasing temperature on the yield of SO_3 .
		29
		[3]
(e)	The	SO ₃ produced is converted to sulfuric acid in two stages. In the first stage the SO ₃ is
		cted with concentrated sulfuric acid to produce oleum, $H_2S_2O_7$. oleum is then reacted with water to form sulfuric acid.
	Sug	gest an equation for the reaction of oleum, $H_2S_2O_7$, with water to form sulfuric acid.
		[1]



[1]



- (f) SO_2 reacts with water to form sulfurous acid. Sulfurous acid is a weak Brønsted-Lowry acid, while sulfuric acid is a strong Brønsted-Lowry acid.
 - (i) Complete the 'dot-and-cross' diagram to show the bonding in a molecule of SO₂. Show outer electrons only.



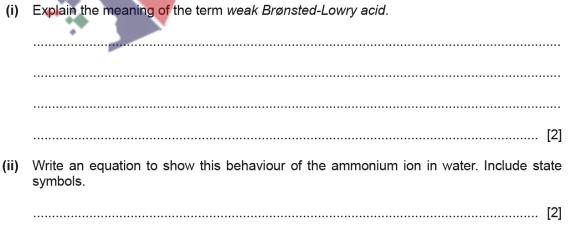
(ii)	State the meaning of the term strong Brønsted-Lowry acid.
	[2]
(iii)	Write an equation to show the acid-base behaviour of sulfuric acid with water. Include state symbols.
	[2]
	[Total: 20]





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		$_{\rm s18_qp_22~Q:~2}$ iium iron(II) sulfate, (NH ₄) ₂ Fe(SO ₄) ₂ , has a relative formula mass, $M_{\rm r}$, of 284.					
(a)	Dei	fine the term <i>relative formula mass</i> .					
			[2]				
(b)	On	e of the cations in ammonium iron(II) sulfate is the ammonium ion, NH_4^+ .					
	(i)	Draw a 'dot-and-cross' diagram of an ammonium ion. Show outer shell electrons only.					
		Use × to show electrons from nitrogen. Use ● to show electrons from hydrogen.					
			[2]				
	(ii)	Suggest the shape of an ammonium ion and predict the bond angle.					
		shape					
		bond angle					
			[2]				
(c)	In a	aqueous solution the ammonium ion acts as a weak Brønsted-Lowry acid.					







(d) Mohr's salt, $(NH_4)_2Fe(SO_4)_2 \cdot xH_2O$, is the hydrated form of ammonium iron(II) sulfate.

x represents the number of moles of water in 1 mole of the salt.

A student wanted to determine the value of x. 0.784g of the hydrated salt was dissolved in water and this solution was acidified.

All of the solution was titrated with $0.0200\,\text{mol\,dm}^{-3}$ potassium manganate(VII). $20.0\,\text{cm}^3$ of this potassium manganate(VII) solution was required for complete reaction with the Fe²⁺ ions.

(i) Use changes in oxidation numbers to balance the equation for the reaction taking place.

$$\mathsf{MnO_4^-(aq)} +\mathsf{Fe^{2^+}(aq)} +\mathsf{H^+(aq)} \to\mathsf{Mn^{2^+}(aq)} +\mathsf{Fe^{3^+}(aq)} +\mathsf{H_2^-O(l)}$$

(ii) State the role of the Fe2+ ions in this reaction.

Explain your answer.	O	
	7	
	 	[2]

(iii) Calculate the amount, in moles, of manganate(VII) ions that reacted.

(iv) Calculate the amount, in moles, of Fe²⁺ ions in the sample of the salt.



amount = mol [1]





(v) Calculate the relative formula mass of $(NH_4)_2Fe(SO_4)_2 \cdot xH_2O$.

relative formula mass = [1]

(vi) Calculate the value of x.

Ralpa Callina







