**1.** This question is about covalent compounds of nitrogen.

Ammonia, NH<sub>3</sub>, is manufactured by reacting nitrogen and hydrogen gases. This is a reversible reaction and the equilibrium is shown below.

 $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ 

 $\Delta H = -92 \text{ kJ mol}^{-1}$ 

i. This is an example of a dynamic equilibrium.

State 2 features of a dynamic equilibrium.

1			

2

\_\_\_\_\_\_[2] State and explain the conditions of temperature and pressure that would produce a large equilibrium yield ii.

2. The reversible reaction below is at equilibrium.

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

What is the expression for  $K_c$ ?

- $[SO_2(g)]^2 [O_2(g)]$ Α  $[SO_3(g)]^2$
- $[SO_3(g)]^2$ В  $[SO_2(g)]^2 [O_2(g)]$  $2[SO_2(g)] + [O_2(g)]$ C
- 2[SO<sub>3</sub>(g)] D  $2[SO_2(g)] + [O_2(g)]$

2[SO<sub>3</sub>(g)]

[3]

3. This question is about two oxides of sulfur: sulfur dioxide, SO<sub>2</sub>, and sulfur trioxide, SO<sub>3</sub>.

SO<sub>3</sub> decomposes to form SO<sub>2</sub> and O<sub>2</sub>, as shown in **Equilibrium 18.1**.

$$SO_3(g) \rightleftharpoons SO_2(g) + \frac{1}{2}O_2(g)$$

 $\Delta H = +99 \text{ kJ mol}^{-1}$ 

**Equilibrium 18.1** 

i. 2.25 moles of SO<sub>3</sub> is heated to 550 °C in the presence of a catalyst and the resulting mixture allowed to reach equilibrium.

The equilibrium mixture contains 0.900 mol of SO<sub>2</sub> and the total pressure is 2.80 atm.

Calculate the numerical value for  $K_p$  for **Equilibrium 18.1** under these conditions and state the units of  $K_p$ .

Give your answer to 3 significant figures.

$K_p =$	 	 	
units	 	 	[5]

ii. The numerical values of  $K_p$  for **Equilibrium 18.1** at temperatures  $T_1$  and  $T_2$  are shown below.

Temperature	<b>K</b> <sub>p</sub>
<i>T</i> <sub>1</sub>	3.3 × 10⁻⁵
<i>T</i> <sub>2</sub>	7.7 × 10 <sup>-2</sup>

Explain why  $T_2$  is a higher temperature than  $T_1$ .

[2]

[21
[4]

iii. Suggest how the value of  $K_p$  would change if the reaction was repeated with no catalyst added and the pressure of the system increased.

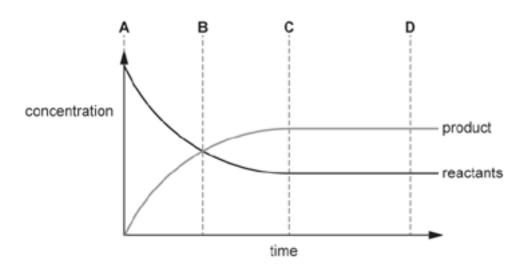
Tick  $(\checkmark)$  one box in each row.

Change	Decrease	No change	Increase
No catalyst			
Increased pressure			

**4.** The reversible reaction between hydrogen and iodine to form hydrogen iodide is  $H_2(g) + I_2(g) = 2HI(g)$ 

The graph shows how the concentrations of the reactants and product change as the reaction reaches a dynamic equilibrium.

At which point on the graph is the equilibrium reached?



Your answer [1]

**5(a).** This question is about oxides of nitrogen.

An investigation is carried out on the equilibrium system shown below.

$$2NO_2(g) \rightleftharpoons N_2O_4(g)$$
  $\Delta H = -57.4 \text{ kJ mol}^{-1}$ 

i. A sealed flask containing 6.00 moles of  $NO_2(g)$  is heated to a constant temperature and allowed to reach equilibrium.

The equilibrium mixture contains 5.40 mol of NO<sub>2</sub>(g), and the total pressure is 5.00 atm.

Determine the value of  $K_p$  and give your answer to **3** significant figures.

Include an expression for  $K_p$  and the units of  $K_p$  in your answer.

<i>K</i> <sub>p</sub> =	units	[5]
higher temperature at ar	n increased pressure. T	he system is

ii. The sealed flask in **(a)(i)** is then heated to a higher temperature at an increased pressure. The system is allowed to reach equilibrium again.

Explain why it is difficult to predict how these changes in reaction conditions affect the amount of N<sub>2</sub>O<sub>4</sub>(g) formed at equilibrium.

[3]

/ 1	_ \	$\sim$	1 - C.	11	<b>1</b>	- <b>c</b>	-I:CC 1	:	- <b>f</b> '1	: -I - A	, as the only	4
,	<b>~</b> 1	1/1/1	, raacte ti	1 1111\/ \A/ITM	OVVICED TO	n torm s	a aittarant	AVIAD A	n nitragan		ac tha Aniv	nradiiat
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Oxide A is collected and cooled to 75.0 °C at a pressure of 101 kPa.

Under these conditions, oxide **A** is a gas that occupies a volume of 74.0 cm<sup>3</sup> and has a mass of 0.280 g.

Calculate the molar mass of oxide **A** and suggest its molecular formula.

6. Chlorine trifluoride can be decomposed into its elements forming the equilibrium mixture below.

$$2C/F_3(g) \rightleftharpoons CI_2(g) + 3F_2(g)$$
  $\Delta H = +318 \text{ kJ mol}^{-1}$  colourless green yellow gas gas

Which statement(s) is/are correct?

- 1 The decomposition is a redox reaction.
- 2 When the equilibrium mixture is cooled, the colour fades.
- **3** The decomposition has a negative entropy change.
- **A** 1, 2 and 3
- B Only 1 and 2
- C Only 2 and 3
- **D** Only 1

Your answer [1]

**7.** Bromine, Br<sub>2</sub>, can be produced by the reaction:

$$5Br^{\scriptscriptstyle -}(aq) + BrO_3\,{}^{\scriptscriptstyle -}(aq) + 6H^{\scriptscriptstyle +}(aq) \rightarrow 3Br_2(aq) + 3H_2O(I)$$

A student investigates the rate of this reaction by carrying out four experiments at the same temperature. The student's results are shown below.

Experiment	[Br <sup>-</sup> ] / mol dm <sup>-3</sup>	[BrO <sub>3</sub> <sup>-</sup> ] / mol dm <sup>-3</sup>	[H+] / mol dm <sup>-3</sup>	Initial rate / mol dm <sup>-3</sup> s <sup>-1</sup>
1	2.00 × 10 <sup>-2</sup>	1.20 × 10 <sup>-1</sup>	8.00 × 10 <sup>-2</sup>	2.52 × 10 <sup>-4</sup>
2	6.00 × 10 <sup>-2</sup>	1.20 × 10 <sup>-1</sup>	8.00 × 10 <sup>-2</sup>	7.56 × 10 <sup>-4</sup>
3	4.00 × 10 <sup>-2</sup>	6.00 × 10 <sup>-2</sup>	8.00 × 10 <sup>-2</sup>	2.52 × 10 <sup>-4</sup>
4	2.00 × 10 <sup>-2</sup>	6.00 × 10 <sup>-2</sup>	$4.00 \times 10^{-1}$	3.15 × 10 <sup>−3</sup>

Explain how the reaction orders can be determined from the student's results, and determine the rate equation and rate constant for this reaction.
[6

[5]

8(a). The reaction between sulfur dioxide, SO <sub>2</sub> (g) and oxygen,	O <sub>2</sub> (g), to form sulfur trioxide,	SO <sub>3</sub> (g), is a key step
in the industrial manufacture of sulfuric acid.		

This is a reversible reaction, shown in **Equilibrium 24.1**:

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

$$\Delta H = -197 \text{ kJ mol}^{-1}$$

**Equilibrium 24.1** 

Why is <b>Equilibrium 24.1</b> a homogeneous equilibrium?	
	[1]
<b>(b).</b> Le Chatelier's principle can be used to predict how different conditions affect the equilibrium position in <b>Equilibrium 24.1</b> .	
Explain how changing pressure, temperature and using a catalyst affect the equilibrium yield of SO <sub>3</sub> .	
In your answer, use le Chatelier's principle and other chemical concepts, where appropriate.	

(c). A mixture of  $SO_2(g)$  and  $O_2(g)$  is allowed to reach equilibrium at a constant temperature.

The equilibrium concentrations are shown in the table.

Substance	Equilibrium concentration / mol dm <sup>-3</sup>	
SO <sub>2</sub> (g)	3.0 × 10 <sup>-3</sup>	
O <sub>2</sub> (g)	3.5 × 10 <sup>-3</sup>	
SO <sub>3</sub> (g)	5.0 × 10 <sup>-2</sup>	

	·	
i.	Write the expression for $K_c$ and calculate the numerical value for $K_c$ in <b>Equilibrium 24.1</b> at this constemperature.	tant
	Give your answer to an appropriate number of significant figures and in standard form.	
	$K_c = \dots dm^3 mc$	ol <sup>-1</sup> <b>[2]</b>
ii.	In the industrial production of $SO_3$ , an excess of $O_2(g)$ is used rather than a 2:1 proportion of $SO_2(g)$ $O_2(g)$ which would match the stochiometry in <b>Equilibrium 24.1</b> .	to
	Suggest, in terms of equilibrium, why an excess of $O_2(g)$ is used industrially.	
		[1]
). Wh	hich prediction can be made using le Chatelier's principle?	
	The effect of a catalyst on the reaction rate.  The effect of a catalyst on the equilibrium position.	
C	The effect of temperature on the reaction rate.	
D	The effect of concentration on the equilibrium position.	
Your	answer [1]	
<b>0.</b> Fo	our equilibrium reactions are set up	

The concentration of each gas in the equilibrium mixtures is  $0.1 \text{ mol dm}^{-3}$ .

Which equilibrium has a numerical  $K_{\text{\tiny c}}$  value of 0.01?

- $CH_4(g) + 2H_2O(g) \leftrightharpoons CO_2(g) + 4H_2(g)$ Α
- В  $N_2(g) + 3H_2(g) \leftrightharpoons 2NH_3(g)$
- С  $H_2(g) + I_2(g) \leftrightarrows 2HI(g)$
- $2\mathsf{NO}_2(g) \leftrightarrows \mathsf{N}_2\mathsf{O}_4(g)$ D

[1] Your answer

**11(a).** This question is about equilibria involving hydrogen.

\* Hydrogen is used industrially to manufacture ammonia.

The equilibrium is shown below.

$$N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$$

$$\Delta H = -92 \text{kJ mol}^{-1}$$

**Equilibrium 20.1** 

1.20 mol  $N_2(g)$  is mixed with 3.60 mol  $H_2(g)$  in a 8.00 dm<sup>3</sup> container.

The mixture is heated to 550 °C with an iron catalyst and allowed to reach equilibrium.

The equilibrium mixture contains 0.160 mol of NH<sub>3</sub>.

Determine the equilibrium constant $K_c$ for <b>Equilibrium 20.1</b> , and explain why the operational conditions used by industry may be different from those required for a maximum equilibrium yield of ammonia.		
	_	
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(b). In industry, hydrogen is used to reduce the ir	ron oxide Fe <sub>3</sub> O <sub>4</sub> as shown in <b>Equilibrium 20.2</b> .
---	--

The reaction is carried out at 500 °C.

 $Fe_3O_4(s) + 4H_2(g) \rightleftharpoons 3Fe(s) + 4H_2O(g)$  Equilibrium 20.2

i.	When the temperature is decreased, the value of $\textit{K}_p$ decreases. Determine whether the forward reaction is exothermic or endothermic. Explain your answer.		
			[1]
ii.	Two students are discussing the eff	ect of pressure on the equilibrium posit	ion of <b>Equilibrium 20.2</b> .
	Student 1 says:		
	"There are more moles of products the left hand side."	than reactants, so increasing the press	ure will shift the equilibrium to
	Student 2 disagrees.		
	Determine which student is correct.	Justify your answer.	
			[1]
<b>12</b> . Th	is question is about the manufacture	of hydrogen, H <sub>2</sub> .	
In indu	ustry, hydrogen is manufactured from	n methane, as shown in <b>Equilibrium 4</b> .	1.
CH₄(g	$O(g) + H_2O(g) \rightleftharpoons CO(g) + 3H_2(g)$	$\Delta H = +206 \text{ kJ mol}^{-1}$	Equilibrium 4.1
	dustrial process is carried out at 15 am. A nickel catalyst is used.	atmospheres pressure and at a tempera	ature of 800°C using an excess
i.	* Explain why these conditions are	used industrially.	

	d H₂O(g) and leaves the mixture to reach e	
A chemist mixes CH <sub>4</sub> (g) and	d $H_2O(g)$ and leaves the mixture to reach $\epsilon$	equilibrium.
A chemist mixes CH <sub>4</sub> (g) and	d $H_2O(g)$ and leaves the mixture to reach $\epsilon$	
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$	d $H_2O(g)$ and leaves the mixture to reach $\epsilon$	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$	d $H_2O(g)$ and leaves the mixture to reach $e^{-1}$ $H_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$	d $H_2O(g)$ and leaves the mixture to reach $e^{-1}$ $H_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$ The equilibrium mixture contains	the H <sub>2</sub> O(g) and leaves the mixture to reach $\Theta$ $H_2(g) \qquad \qquad \Delta H = +206 \text{ kJ mol}^{-1}$ $\text{Ins the following concentrations.}$	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$ The equilibrium mixture contains Substance	$d_1H_2O(g)$ and leaves the mixture to reach each $d_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$ ins the following concentrations.	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$ The equilibrium mixture contains	$H_2O(g)$ and leaves the mixture to reach $H_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$ ins the following concentrations.  Concentration/mol dm <sup>-3</sup> 0.111	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$ The equilibrium mixture contains  Substance $CH_4(g)$ $H_2O(g)$	$H_2O(g)$ and leaves the mixture to reach $H_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$ ins the following concentrations.	equilibrium.
A chemist mixes $CH_4(g)$ and $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3$ The equilibrium mixture contains the equilibrium mixture $CH_4(g) = CH_4(g) = CO(g) = CO(g)$	$H_2O(g)$ and leaves the mixture to reach $e^{\frac{1}{2}}H_2(g)$ $\Delta H = +206 \text{ kJ mol}^{-1}$ ins the following concentrations. $\frac{\text{Concentration/mol dm}^{-3}}{0.111}$ 0.682 0.510	equilibrium.

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3.2.3 Chemical Equilibrium

**13(a).** Two students plan to investigate **Equilibrium 4.1**, shown below.

$$CoCl_4^{2-}(aq) + 6H_2O(I)$$
  $\Rightarrow$   $[Co(H_2O)_6]^{2+}(aq) + 4Cl^{-}(aq)$  Equilibrium 4.1

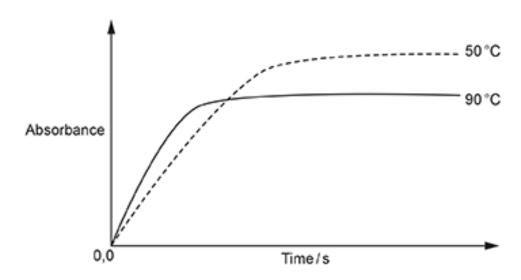
The students are supplied with the equilibrium mixture in **Equilibrium 4.1** at room temperature.

- One student heats 20 cm<sup>3</sup> of the mixture to 50°C.
- The other student heats 20 cm<sup>3</sup> of the mixture to 90°C.

The students use colorimetry to observe how the colour of the equilibrium mixture changes over time.

- The colorimeter is set up so that the greater the absorbance, the greater the concentration of  $[Co(H_2O)_6]^{2+}$ .
- The initial absorbance is set to zero.
- The absorbance is recorded every 30 seconds.

The students plot the graph below from the results of the experiment.



Use the graph and relevant chemical theory to answer the following. Include all reasoning:

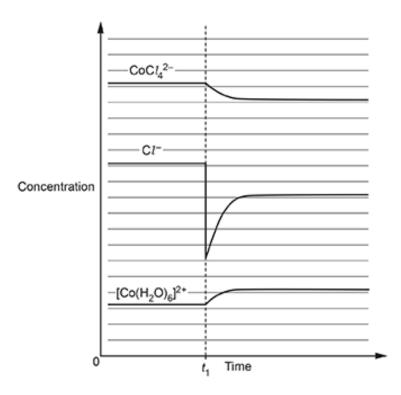
- Explain the different initial rates at 50°C and 90°C.
- Predict the sign of  $\Delta H$  for the forward reaction in **Equilibrium 4.1**.

[3]

**(b).** The students investigate how addition of aqueous silver nitrate, AgNO<sub>3</sub>(aq), affects the equilibrium position in **Equilibrium 4.1**.

The graph shows the changes in the equilibrium concentrations of  $CoCI_4$  <sup>2-</sup>, CI and  $[Co(H_2O)_6]^{2+}$  after addition of the  $AgNO_3(aq)$ .

The AgNO<sub>3</sub>(aq) is added at time =  $t_1$ 



i.	Explain why the C $I$ concentration drops sharply at time = $t_1$ .		
		[1]	
ii.	Explain the changes in concentration of $CoCl_4$ <sup>2-</sup> , $Cl$ and $[Co(H_2O)_6]^{2+}$ after time = $t_1$ . Refer to <b>Equilibrium 4.1</b> in your answer.		