

Questions

1. CHEMISTRY, M5 2019 HSC 16 MC

At equilibrium, a 1.00 L vessel contains 0.0430 mol of H_2 , 0.0620 mol of I_2 , and 0.358 mole of HI. The system is represented by the following equation:

$$H_{2}\left(g\right)+I_{2}\left(g\right)\
ightleftharpoons 2\,HI\left(g\right)$$

Which of the following is closest to the value of the equilibrium constant, K_{eq} , for this reaction?

- **A.** 0.0208
- **B.** 48.1
- **C.** 134
- **D.** 269

2. CHEMISTRY, M5 EQ-Bank 3 MC

Consider the following reaction.

$$2 \operatorname{NOCl}(g) \rightleftharpoons 2 \operatorname{NO}(g) + \operatorname{Cl}_2(g)$$

What is the equilibrium expression for this reaction?

- A. $\frac{2[NO][Cl_2]}{2[NOCl]}$
- B. $\frac{[\text{NO}]^2[\text{Cl}_2]}{[\text{NOCl}]^2}$
- c. $\frac{2[\text{NOCl}]}{2[\text{NO}][\text{Cl}_2]}$
- D. $\frac{\mathrm{[NOCl]}^2}{2\mathrm{[NO]}^2\mathrm{[Cl_2]}}$

3. CHEMISTRY, M5 2022 HSC 13 MC

Nitrosyl bromide decomposes according to the following equation.

$$2 \operatorname{NOBr}(g) \rightleftharpoons 2 \operatorname{NO}(g) + \operatorname{Br}_2(g)$$

A 0.64 mol sample of NOBr is placed in an evacuated 1.00 L flask. After the system comes to equilibrium, the flask contains 0.46 mol NOBr.

What are the concentrations of NO and Br_2 in the flask at equilibrium?

	$[\mathrm{NO}] \left(\mathrm{mol} \ \mathrm{L}^{-1} \right)$	$\left[\mathrm{Br}_2\right]\left(\mathrm{mol}\ \mathrm{L}^{-1}\right)$
A.	0.18	0.09
в.	0.18	0.18
С.	0.36	0.18
D.	0.92	0.46

4. CHEMISTRY, M5 2022 HSC 8 MC

A system is described as follows.

$$2 \operatorname{NaHCO}_3(s) \rightleftharpoons \operatorname{Na}_2 \operatorname{CO}_3(s) + \operatorname{CO}_2(g) + \operatorname{H}_2 \operatorname{O}(g)$$

What is the equilibrium expression for this system?

A.
$$K_{eq} = [\mathrm{CO}_2]$$

B.
$$K_{eq} = [CO_2][H_2O]$$

C.
$$K_{eq} = rac{1}{[\mathrm{CO_2}][\mathrm{H_2O}]}$$

$$\textbf{D.} \quad K_{eq} = \frac{\left[\text{Na}_2\text{CO}_3\right]\left[\text{CO}_2\right]\left[\text{H}_2\text{O}\right]}{\left[\text{NaHCO}_3\right]^2}$$

5. CHEMISTRY, M5 2023 HSC 7 MC

A mixture of 0.8 mol of CO(g) and 0.8 mol of $H_2(g)$ was placed in a sealed 1.0 L container. The following reaction occurred.

$$\mathrm{CO}(g) + 2\mathrm{H}_2(g) \rightleftharpoons \mathrm{CH}_3\mathrm{OH}(g)$$

When equilibrium was established, the mixture contained 0.5 mol of $\mathrm{CO}(\mathrm{g})$

What amount of $H_2(g)$ was present at equilibrium?

- **A.** 0.2 mol
- **B.** 0.4 mol
- C. 0.6 mol
- **D.** 1.0 mol

6. CHEMISTRY, M5 EQ-Bank 10 MC

At a certain temperature, the K_{eq} for the following reaction is 75.

$$2 O_3(g) \rightleftharpoons 3 O_2(g)$$

0.4 mol of O_3 and 1.2 mol of O_2 were introduced to a 5 L reaction vessel.

Which row of the table correctly identifies the direction of the equilibrium shift and the reason for the shift?

	$Direction\ favoured$	Reason
A.	Left	$Q>K_{eq}$
В.	Left	$Q < K_{eq}$
C.	Right	$Q>K_{eq}$
D.	Right	$Q < K_{eq}$

7. CHEMISTRY, M6 2020 HSC 14 MC

The equation for the autoionisation of water is shown.

$$2 \,\mathrm{H}_2\mathrm{O}\left(\mathrm{l}\right) \;
ightleftharpoons \;\mathrm{H}_3\mathrm{O}^+\left(\mathrm{aq}\right) + \mathrm{OH}^-\left(\mathrm{aq}\right)$$

At 50°C the water ionisation constant, K_w is 5.5×10^{-14} .

What is the pH of water at 50°C?

- **A.** 5.50
- **B.** 6.63
- **C.** 6.93
- **D.** 7.00

8. CHEMISTRY, M5 2019 HSC 17 MC

A student makes a solution with a final volume of 200 mL by mixing 100 mL of 0.0500 mol L $^{-1}$ barium nitrate solution with 100 mL of 0.100 mol L $^{-1}$ sodium hydroxide solution.

Which row of the table correctly identifies if a precipitate will form under these conditions and the reason?

	Will a precipitate form?	Reason
A.	Yes	$Q > K_{sp}$
B.	Yes	$Q < K_{sp}$
C.	No	$Q > K_{sp}$
D.	No	$Q < K_{sp}$

9. CHEMISTRY, M5 2021 HSC 19 MC

A quantity of silver nitrate is added to 250.0 mL of 0.100 mol L $^{-1}$ potassium sulfate at 298 K in order to produce a precipitate. Silver nitrate has a molar mass of 169.9 g mol $^{-1}$.

What mass of silver nitrate will cause precipitation to start?

- **A.** 0.00510 g
- **B.** 0.186 g
- **C.** 0.465 g
- **D.** 0.854 g

10. CHEMISTRY, M5 2023 HSC 20 MC

Nitrogen monoxide and oxygen combine to form nitrogen dioxide, according to the following equation.

$$2\,\mathrm{NO}\left(\mathrm{g}
ight) + \mathrm{O}_{2}\left(\mathrm{g}
ight) \
ightleftharpoons 2\,\mathrm{NO}_{2}\left(\mathrm{g}
ight) \quad K_{eq} = 2.47\, imes 10^{12}$$

A 2.00 L vessel is filled with 1.80 mol of $NO_2(g)$ and the system is allowed to reach equilibrium.

What is the equilibrium concentration of NO(g)?

- **A.** 0.00 mol L^{-1}
- **B.** $4.34 \times 10^{-5} \text{ mol } L^{-1}$
- **C.** $6.90 \times 10^{-5} \text{ mol } L^{-1}$
- **D.** $8.69 \times 10^{-5} \ mol \ L^{-1}$

11. CHEMISTRY, M5 2024 HSC 15 MC

The thermal decomposition of lithium peroxide $(\mathrm{Li}_2\mathrm{O}_2)$ is given by the equation shown.

$$2\,\mathrm{Li_2O_2(s)} \rightleftharpoons 2\,\mathrm{Li_2\,O\,(s)} + \mathrm{O_2\,(g)}$$

Mixtures of Li_2O_2 , Li_2O and O_2 were allowed to reach equilibrium in two identical, closed containers, P and Q, at the same temperature. The amount of Li_2O_2 (s) in container P is double that in container Q. The amount of Li_2O (s) is the same in each container.

What is the ratio of $\left[O_{2}\left(g\right)\right]$ in container P to $\left[O_{2}\left(g\right)\right]$ in container Q?

- **A**. 1:1
- **B.** 2:1
- **C.** 3:2
- **D.** 5:4

12. CHEMISTRY, M5 2024 HSC 18 MC

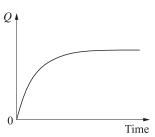
A reaction mixture, not at equilibrium, is composed of both $N_2O_4(g)$ and $NO_2(g)$ in a closed container. The reaction quotient for the system, Q, is given.

$$Q = \frac{\left[\text{NO}_2\right]^2}{\left[\text{N}_2\text{O}_4\right]}$$

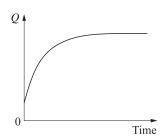
The rate of the forward reaction is initially greater than the rate of the reverse reaction.

Which diagram shows how Q changes over time for this mixture?

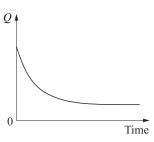
A.



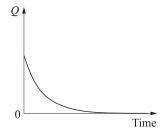
B.



C.



D.



13. CHEMISTRY, M5 EQ-Bank 13 MC

0.20 moles of phosphorus pentachloride were heated to 200°C in a 2 L container in the presence of a vanadium catalyst according to the following reaction.

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

At equilibrium, the mixture was found to contain 0.16 moles of chlorine.

Which of the following is the equilibrium constant for this reaction at this temperature?

- **A.** 0.32
- **B.** 0.64
- **C.** 1.56
- **D.** 3.13

14. CHEMISTRY, M5 EQ-Bank 22

Hydrogen gas reacts with iodine gas to form hydrogen iodide according to the following equation.

$$H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$$
 at 700 k

At equilibrium, the concentrations for H_2 , I_2 and HI are as follows: 0.326 mol L^{-1} , 0.326 mol L^{-1} and 2.39 mol L^{-1} respectively.

What is the value of the equilibrium constant for this reaction? (2 marks)

15. CHEMISTRY, M5 2019 HSC 31

GTEINIGTKT, INIG 2019 FIGG 31
The following reaction occurs in an aqueous solution.
${ m HgCl_4}^{2-} \left({ m aq} ight) + { m Cu}^{2+} \left({ m aq} ight) \ ightleftharpoons = { m CuCl_4}^{2-} \left({ m aq} ight) + { m Hg}^{2+} \hspace{0.5cm} K_{eq} = 4.55 imes 10^{-11}$
A solution containing a mixture of ${ m HgCl_4}^{2-}$ (aq) and ${ m Cu^{2+}}$ (aq) ions is prepared. The initial concentration of each ion is 0.100 mol L $^{-1}$ and there are no other ions present.
Calculate the concentration of ${ m Hg^{2+}}$ ${ m (aq)}$ ions once the system has reached equilibrium. <i>(4 marks)</i>

16. CHEMISTRY, M5 2024 HSC 23

Consider the following equilibrium system.

$$[\text{Co} (\text{H}_2\text{O})_6]^{2+} (\text{aq}) + 4 \, \text{Cl}^- (\text{aq}) \ \rightleftharpoons [\text{CoCl}_4]^{2-} (\text{aq}) + 6 \, \text{H}_2\text{O} (\text{l})$$

 $[Co(H_2O)_6]^{2+}$ (aq) is pink and $[CoCl_4]^{2-}$ (aq) is blue. When a solution of these ions and chloride ions is heated, the mixture becomes more blue.

Relate the observed colour change to the change in K_{eq} . (3 marks)

Butanoic acid is a natural product and a component of human sweat.	
Calculate the value of K_a for butanoic acid if a 0.10 mol L $^{-1}$ solution has a pH of 2.9 at 298 K. (3 mark	rs)
18. CHEMISTRY, M5 EQ-Bank 24	
When a sample of solid silver chloride is added to a 1.00×10^{-2} mol L $^{-1}$ sodium chloride solution, or some of the silver chloride dissolves.	ıly
When a sample of solid silver chloride is added to a 1.00×10^{-2} mol L ⁻¹ sodium chloride solution, or some of the silver chloride dissolves. Calculate the equilibrium concentration of silver ions in the resulting solution, given that the K_{sp} of sil chloride is 1.8×10^{-10} . (3 marks)	
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19. CHEMISTRY, M5 2023 HSC 31

Copper(II) ions (Cu^{2+}) form a complex with lactic acid ($C_3H_6O_3$), as shown in the equation.

$$\mathrm{Cu}^{2+}\left(\mathrm{aq}\right)+2\,\mathrm{C}_{3}\mathrm{H}_{6}\mathrm{O}_{3}\left(\mathrm{aq}\right) \rightleftharpoons \left[\mathrm{Cu}(\mathrm{C}_{3}\mathrm{H}_{6}\mathrm{O}_{3})_{2}\right]^{2+}\left(\mathrm{aq}\right)$$

This complex can be detected by measuring its absorbance at 730 nm. A series of solutions containing known concentrations of $\left[Cu(C_3H_6O_3)_2 \right]^{2+}$ were prepared, and their absorbances measured.

$Concentration\ of \Big[\mathrm{Cu}(\mathrm{C_3H_6O_3})_2\Big]^{2+}\ (\mathrm{mol}\ \mathrm{L}^{-1})$	Absorbance
0.000	0.00
0.010	0.13
0.020	0.28
0.030	0.43
0.040	0.57
0.050	0.72

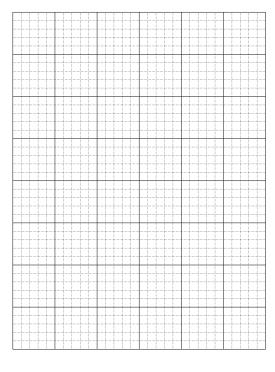
Two solutions containing Cu^{2+} and $C_3H_6O_3$ were mixed. The initial concentrations of each in the resulting solution are shown in the table.

Species	$\begin{array}{c} Initial\ Concentration \\ (\bmod\ L^{-1}) \end{array}$
Cu^{2+}	0.056
$\mathrm{C_3H_6O_3}$	0.111

When the solution reached equilibrium, its absorbance at 730 nm was 0.66.

You may assume that under the conditions of this experiment, the only species present in the solution are those present in the equation above, and that $\left[Cu(C_3H_6O_3)_2\right]^{2+}$ is the only species that absorbs at 730 nm.

With the support of a line graph, calculate the equilibrium constant for the reaction. (7 marks)



20. CHEMISTRY, M6 2019 HSC 27

	reflationship between the acid dissociation constant, K_a , and the corresponding conjugate base association constant, K_b , is given by:
٨٠	$K_a imes K_b = K_w$ ssume that the temperature for part (a) and part (b) is 25°C.
	Figure that the temperature for part (a) and part (b) is 2.5 C. Figure 4. The K_a of hypochlorous acid (HOCl) is 3.0×10^{-8} .
S	Show that the K_b of the hypochlorite ion, OCl^- , is $~3.3 imes10^{-7}$. (1 mark)
b. 7	The conjugate base dissociation constant, K_{b_t} is the equilibrium constant for the following equation:
	$\mathrm{OCl}^-(aq) + \mathrm{H}_2\mathrm{O}(l) ightleftharpoons \mathrm{HOCl}(aq) + \mathrm{OH}^-(aq)$
(Calculate the pH of a 0.20 mol L^{-1} solution of sodium hypochlorite (NaOCl). (4 mark)
•	
·	

21. CHEMISTRY, M5 2020 HSC 35

In aqueous solution, iodide ions (I^-) react rapidly with iodine (I_2) to form triiodide ions I_3^- , making the equilibrium system shown in the chemical equation:

$$\mathrm{I}^-(aq) + \mathrm{I}_2(aq)
ightleftharpoons \mathrm{I}_3^-(aq)$$

The following relationships can be derived from the reaction mechanism:

$$\begin{split} \left[\mathbf{I}^{-}\right]_{\mathit{eq}} &= 2\left[\mathbf{I}_{2}\right]_{\mathit{eq}} \\ \left[\mathbf{I}^{-}\right]_{\mathit{initial}} &= 4\left[\mathbf{I}_{2}\right]_{\mathit{eq}} + 3\left[\mathbf{I}_{3}^{\;-}\right]_{\mathit{eq}} \end{split}$$

where 'initial' designates the initial concentration and 'eq' designates the equilibrium concentration.

The absorbance of the solution in the UV-Vis spectrum is given by:

$$A = \left[\operatorname{I}_3^{\,-}
ight] imes 2.76 imes 10^4$$

Determine the value of the equilibrium constant, given that A=0.745 at equilibrium and $\left[\mathrm{I}^{-}\right]_{initial}=7.00\times10^{-4}~\mathrm{mol}~\mathrm{L}^{-1}$. (4 marks)

22. CHEMISTRY, M5 2021 HSC 31

Ammonia is produced according to the following equilibrium equation.

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

There are 4.50 moles of nitrogen gas, 1.00 mole of hydrogen gas and 5.80 moles of ammonia in a 10.0 L vessel. The system is at equilibrium at 298 K. The value of K_{eq} at this temperature is 748 .

How many moles of nitrogen gas need to be added to the vessel to increase the amount of ammonia by 0.050 moles? (4 marks)

23. CHEMISTRY, M5 2024 HSC 30

An equilibrium mixture of hydrogen, carbon dioxide, water and carbon monoxide is in a closed, 1 L container at a fixed temperature as shown:

$$H_2(g) + CO_2(g) \implies H_2O(g) + CO(g)$$
 $K_{eq} = 1.600$

The initial concentrations are

$$[H_2]=1.000~\text{mol}~L^{-1},~[CO_2]=0.500~\text{mol}~L^{-1},~[H_2O]=0.400~\text{mol}~L^{-1}$$
 and $[CO]=2.000~\text{mol}~L^{-1}.$

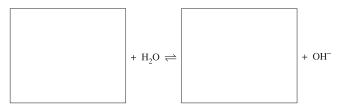
An unknown amount of $CO\left(g\right)$ was added to the same container, and the temperature was kept constant. After the new equilibrium had been established, the concentration of $H_2O\left(g\right)$ was found to be 0.200 mol L^{-1} .

Using this information, calculate the unknown amount (in mol) of CO(g) that was added to the container. (4 marks)

24. CHEMISTRY, M5 2020 HSC 27

A student makes up a solution of propan-2-amine in water with a concentration of 1.00 mol L^{-1} .

a. Using structural formulae, complete the equation for the reaction of propan-2-amine with water. (2 marks)



b. The equilibrium constant for the reaction of propan-2-amine with water is 4.37×10^{-4} . Calculate the concentration of hydroxide ions in this solution. (3 marks)

25. CHEMISTRY, M6 2020 HSC 33

Excess solid calcium hydroxide is added to a beaker containing 0.100 L of $2.00 mol L^{-1}$ hydrochloric acid and the mixture is allowed to come to equilibrium.

a.	Show that the amount (in mol) of calcium hydroxide that reacts with the hydrochloric acid is 0.100 mol. (2 marks)
	It is valid in this instance to make the simplifying assumption that the amount of calcium ions present a equilibrium is equal to the amount generated in the reaction in part (a).
	Calculate the pH of the resulting solution. (4 marks)

26. CHEMISTRY, M5 2023 HSC 37

When performing industrial reductions with ${ m CO(g)}$, the following equilibrium is of great importance. $2{ m CO}({ m g}) \ ightharpoons { m CO_2}({ m g}) + { m C}({ m s}) \qquad K_{eq} = 10.00 \ { m at } 1095 \ { m K}$	
A 1.00 L sealed vessel at a temperature of 1095 K contains $CO(g)$ at a concentration of 1.10 × 10 ⁻² m L ⁻¹ , $CO_2(g)$ at a concentration of 1.21 × 10 ⁻³ mol L ⁻¹ , and excess solid carbon.	nol
. Is the system at equilibrium? Support your answer with calculations. (2 marks)	
. Carbon dioxide gas is added to the system above and the mixture comes to equilibrium. The equilibrium concentrations of $CO\left(g\right)$ and $CO_{2}\left(g\right)$ are equal. Excess solid carbon is present and the temperature remains at 1095 K.	um e
Calculate the amount (in mol) of carbon dioxide added to the system. (3 marks)	

$$BrCH_2COOH(aq) \rightleftharpoons BrCH_2COOH(octan-l-ol)$$

The equilibrium constant expression for this system is

$$K_{eq} = rac{[ext{BrCH}_2 ext{COOH}(octan-l-ol)]}{[ext{BrCH}_2 ext{COOH}(aq)]}$$

An aqueous solution of bromoacetic acid with an initial concentration of 0.1000 mol L $^{-1}$ is shaken with an equal volume of octan-1-ol. Bromoacetic acid does not dissociate in octan-1-ol but does dissociate in water, with $K_a=1.29\times 10^{-3}$. When the system has reached equilibrium, the $[\mathrm{H}^+]$ is 9.18×10^{-3} mol L $^{-1}$.

Calculate the equilibrium concentration of aqueous bromoacetic acid and hence, or otherwise, calculate the K_{eq} for the octan-1-ol and water system. (4 marks)

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Worked Solutions

1. CHEMISTRY, M5 2019 HSC 16 MC

$$egin{aligned} K_{eq} &= rac{[ext{HI}]^2}{[ext{H}_2][ext{I}_2]} \ &= rac{0.358^2}{(0.0430)(0.0620)} \ &= 48.1 \end{aligned}$$

$$\Rightarrow B$$

2. CHEMISTRY, M5 EQ-Bank 3 MC

ightarrow All elements in the reaction are included in the K_{eq} expression (i.e. no solids or pure liquids are present which would be omitted)

$$K_{eq} = rac{[ext{NO}]^2 [ext{Cl}_2]}{[ext{NOCl}]^2}$$

$$\Rightarrow B$$

3. CHEMISTRY, M5 2022 HSC 13 MC

	NOBr	NO	Br_2
Initial	0.64	0	0
Change	-0.18	+0.18	+0.09
Equilibrium	0.46	0.18	0.09

$$[\mathrm{NO}] = 0.18 \; \mathrm{mol} \; \mathrm{L}^{-1}$$
 $[\mathrm{Br_2}] = 0.09 \; \mathrm{mol} \; \mathrm{L}^{-1}$
 $\Rightarrow A$

4. CHEMISTRY, M5 2022 HSC 8 MC

- \rightarrow The concentrations of solids and pure liquids are omitted from the equilibrium expression because they have a constant concentration.
- \rightarrow Thus, the equilibrium expression is:

$$K_{eq} = \mathrm{[CO_2][H_2O]}$$

$$\Rightarrow B$$

5. CHEMISTRY, M5 2023 HSC 7 MC

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

	CO(g)	$2\mathrm{H}_2\left(\mathrm{g} ight)$	$\mathrm{CH_{3}OH}$
Initial	0.8	0.8	0
Change	-x	-2x	+x
Equilibrium	0.5	0.2	0.03

$$x = 0.3$$

 $\Rightarrow A$

6. CHEMISTRY, M5 EQ-Bank 10 MC

$$O_2 \colon c = \frac{n}{V} = \frac{1.2}{5} = 0.24 \, \text{mol} \, L^{-1}$$

$$O_3 \colon c = rac{n}{V} = rac{0.4}{5} = 0.08 \, \mathrm{mol} \, L^{-1}$$

$$K_{eq} = rac{[{
m O}_2]^3}{[{
m O}_2]^2} = 75 \, {
m (given)}$$

$$Q = \frac{0.24^3}{0.08^2} = 2.16$$

Since $Q < K_{eq}$, the reaction will shift right to favour the products until $Q = K_{eq}$ $\Rightarrow D$

7. CHEMISTRY, M6 2020 HSC 14 MC

$$K_w = [\mathrm{H_3O^+}][\mathrm{OH^-}]$$

Since $[H_3O^+] = [OH^-]$:

$$\left[{\rm H_3O^+} \right]^2 = 5.5 imes 10^{-14}$$

$$\left[{
m H_{3}O^{+}}
ight]\,=\,2.3 imes10^{-7}\,\,{
m mol}\,{
m L^{-1}}$$

$$\begin{aligned} \mathrm{pH} &= -\mathrm{log}_{10} \left(2.3 \times 10^{-7} \right) \, = \, 6.63 \\ &\Rightarrow B \end{aligned}$$

8. CHEMISTRY, M5 2019 HSC 17 MC

$$Ba(OH)_2(s) \rightleftharpoons Ba^{2+}(aq) + 2OH^-(aq)$$

$$K_{sp} = 2.55 imes 10^{-4}$$

$${
m [Ba^{2+}]} = rac{(0.05)(0.1)}{(0.2)} = 0.025\,{
m M}$$

$${
m [OH^-]}=rac{(0.1)(0.1)}{(0.2)}=0.05\,{
m M}$$

$$Q = [{\rm Ba^{2+}}][{\rm OH^-}]^2$$

$$=(0.025)(0.05)^2$$

$$=6.25 imes 10^{-6}$$

Since $Q < K_{sp}$, no precipitate forms.

$$\Rightarrow D$$

9. CHEMISTRY, M5 2021 HSC 19 MC

The reaction when silver nitrate is added to potassium sulfate is:

$$2 \mathrm{AgNO}_3(aq) + \mathrm{K}_2 \mathrm{SO}_4(aq) \ o \ \mathrm{Ag}_2 \mathrm{SO}_4(s) + 2 \mathrm{KNO}_3(aq)$$

♦ Mean mark 45%

Since each K_2SO_4 molecule has 1 sulfate ion

$$\left[{{{
m SO}_4^{2 - }}} \right] = {{
m K}_2}{{
m SO}_4} = 0.100\,\,{
m mol}\,{{
m L}^{ - 1}}$$

$$Ag_2SO_4(s) \leftrightharpoons 2Ag^+(aq) + SO_4^{2-}(aq)$$

$$\mathrm{K}_{sp} = \left[\mathrm{Ag}^+
ight]^2 \left[\mathrm{SO}_4^{2-}
ight]$$

From the data sheet:

$${
m K}_{sp}=1.20~{
m x}~10^{-5}$$

$$1.20 \times 10^{-5} = \left\lceil \mathrm{Ag}^+ \right\rceil^2 \times \left\lceil \mathrm{SO_4^{\,2-}} \right\rceil$$

$$1.20 \times 10^{-5} = \left[Ag^{+} \right]^{2} \times \left[0.100 \right]$$

$$\left\lceil Ag^+\right\rceil = 0.01095\dots \, \text{mol} \, L^{-1}$$

$$[Ag^+] = [AgNO_3]$$

$$n(AgNO_3) = c \times V = 0.01095... \times 0.250 = 0.00273... mol$$

$$m(AgNO_3) = n \times MM = 0.00273... \times 169.9 = 0.465~g$$

$$\Rightarrow \mathit{C}$$

♦♦♦ Mean mark 36%

10. CHEMISTRY, M5 2023 HSC 20 MC

 \rightarrow As 1.80 mol of $NO_2\left(g\right)$ is added to the solution, the reverse reaction can be used to determine the equilibrium concentration of $NO\left(g\right)$.

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

$$ightarrow$$
 Reverse reaction $\ K_{eq} = rac{[{
m O}_2][{
m NO}]^2}{[{
m NO}_2]^2}$

 \rightarrow Forward reaction K_{eq} is the inverse of K_{eq} of the reverse reaction:

$$K_{eq} = rac{1}{2.47 imes 10^{12}} = 4.0486 imes 10^{-13}$$

	$2\mathrm{NO}_2\left(\mathrm{g} ight)$	2 NO (g)	$O_2(g)$
Initial	0.9	0	0
Change	-2x	+2x	+x
Equilibrium	0.9 - 2x	2x	x

- ightarrow -2x is very small as the K_{eq} for the reaction is very small, thus 0.9-2xpprox 0.9.
- \rightarrow By substituting the values into the K_{eq} for the reverse reaction:

$$4.0486 imes 10^{-13} = rac{(x)(2x)^2}{(0.9)^2} \ = rac{4x^3}{(0.9)^2} \ 4x^3 = 3.279 imes 10^{-13} \ x = 4.344 imes 10^{-5}$$

$$\begin{array}{l} \rightarrow \left[\mathrm{NO_2}\right] = 2\,\times 4.344\,\times 10^{-5} = 8.69\,\times 10^{-5}~\mathrm{mol}~\mathrm{L}^{-1} \\ \Rightarrow D \end{array}$$

11. CHEMISTRY, M5 2024 HSC 15 MC

- ightarrow When calculating the K_{eq} of a system, substances in solid states are all given a value of 1.
- ightarrow The equilibrium constant of the above reaction is $K_{eq}=\left[\mathrm{O}_{2}\left(\mathrm{g}
 ight)
 ight]$
- ightarrow As both mixtures reached equilibrium, the K_{eq} values for each mixture is the same, hence the ratio of $\left[O_{2}\left(\mathbf{g}\right)\right]$ in each container is 1:1.
- $\Rightarrow A$

♦♦ Mean mark 34%.

♦♦ Mean mark 39%

 \rightarrow When calculating the reaction quotient of a chemical reaction, the formula is $\frac{products}{max + to t^{-1}}$.

♦ Mean mark 43%

→ The equation for the reaction taking place is:

12. CHEMISTRY, M5 2024 HSC 18 MC

$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$

- \rightarrow As the rate of the forward reaction is greater than the rate of the reverse reaction, $\left\lceil NO_2 \right\rceil^2$ will increase and $\left\lceil N_2O_4 \right\rceil$ will decrease.
- \rightarrow Hence the value for the reaction quotient, $\frac{\left[NO_{2}\right]^{2}}{\left[N_{2}O_{4}\right]}$, will increase.
- ightarrow As it states both $m N_2O_4\left(g
 ight)$ and $m NO_2\left(g
 ight)$ are present in the intial system, the value for Q will not be zero.

 $\Rightarrow B$

13. CHEMISTRY, M5 EQ-Bank 13 MC

$$PCl_{5}\left(g\right) \; \rightleftharpoons PCl_{3}\left(g\right) + Cl_{2}\left(g\right)$$

$$[PCl_5]_{init} = \frac{n}{V} = \frac{0.20}{2} = 0.10\, mol\, L^{-1}$$

	$[PCl_5]$	$[PCl_3]$	$[Cl_2]$
Initial	0.10	0	0
Change	-0.08	+0.08	+0.08
Equilibrium	0.02	0.08	0.08

$$egin{aligned} K_{eq} &= rac{[ext{PCl}_3][ext{Cl}_2]}{[ext{PCl}_5]} &= rac{0.08^2}{0.02} = 0.32 \ \Rightarrow A \end{aligned}$$

14. CHEMISTRY, M5 EQ-Bank 22

$$K_{eq} = rac{[ext{HI}]^2}{[ext{H}_2][ext{I}_2]} = rac{2.39^2}{[0.326][0.326]} = 53.7$$

15. CHEMISTRY, M5 2019 HSC 31

$$K_{eq} = rac{[{
m CuCl_4}^{2-}][{
m Hg}^{2+}]}{[{
m HgCl_4}^{2-}][{
m Cu}^{2+}]}$$

	$[\mathrm{HgCl_4}^{2-}]$	$[\mathrm{Cu}^{2+}]$	$[\operatorname{CuCl}_4{}^{2-}]$	$[\mathrm{Hg}^{2+}]$
Initial	0.100	0.100	0	0
Change	-x	-x	+x	+x
Equilibrium	0.100 - x	0.100 - x	x	x

Since x is small $\Rightarrow 0.100 - x \approx 0.100$

$$4.55 imes 10^{-11} = rac{x imes x}{(0.100 - x)(0.100 - x)}$$

$$4.55 imes 10^{-11} = rac{x^2}{(0.100)^2} \ x^2 = 4.55 imes 10^{-11} imes (0.100)^2 \ x = \sqrt{4.55 imes 10^{-11} imes (0.100)^2} \ = 6.75 imes 10^{-7} ext{ mol L}^{-1}$$

$$\therefore [{\rm Hg^{2+}}] = 6.75\,\times 10^{-7}\,{\rm mol}\,{\rm L^{-1}}$$

16. CHEMISTRY, M5 2024 HSC 23

- \rightarrow When the solution is heated and the mixture becomes more blue, it suggests that the concentration of $[CoCl_4]^{2-}$ (aq) is increasing.
- \rightarrow The increase in temperature favoured the forward endothermic reaction and shifts the equilibrium position to the products.
- \rightarrow Therefore the concentrations of $[Co(H_2O)_6]^{2+}$ (aq) and Cl^- (aq) will decrease.

$$ightarrow$$
 As $K_{eq}=rac{\left[\left[\mathrm{CoCl_4}
ight]^2-
ight]}{\left[\left[\mathrm{Co}\left(\mathrm{H_2O}
ight)_6
ight]^{2+}
ight]\left[\mathrm{Cl}^-
ight]^4}$, K_{eq} will increase.

17. CHEMISTRY, M5 EQ-Bank 23

	$[\mathrm{C_3H_7COOH}]$	$[\mathrm{C_3H_7COO^-}]$	$[\mathrm{H}^+]$
Initial	0.10	0	0
Change	$-1.26 imes10^{-3}$	$+1.26 imes10^{-3}$	$+1.26 imes10^{-3}$
Equilibrium	0.0987	$1.26 imes10^{-3}$	$1.26 imes10^{-3}$

$$K_a = rac{\left(1.26 imes 10^{-3}
ight)^2}{0.0987} \ = 1.6 imes 10^{-5}$$

18. CHEMISTRY, M5 EQ-Bank 24

$$AgCl(s) \rightleftharpoons Ag^{+}(aq) + Cl^{-}(aq)$$

	[AgCl(s)]	$\left[\mathrm{Ag^{+}}\left(\mathrm{aq} ight) ight]$	$\left[\mathrm{Cl^{-}\left(aq ight)} ight]$
Initial		0	$1.00 imes10^{-2}$
Change		+x	+x
Equilibrium		x	$1.00 imes 10^{-2} + x$

Let
$$x = [Ag^+]$$

$$K_{sp}=[\mathrm{Ag}^+][\mathrm{Cl}^-]$$

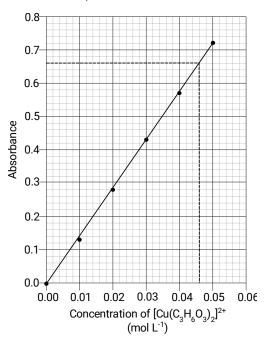
$$K_{sp} = x(1.00 \, imes 10^{-2} + x) = 1.80 \, imes 10^{-10}$$

Since x is small, $1.00 \times 10^{-2} + x \approx 1.00 \times 10^{-2}$

$$x (1.00 \times 10^{-2}) = 1.80 \times 10^{-10}$$

 $x = 1.80 \times 10^{-8}$
 $\therefore [\mathrm{Ag^+}] = 1.80 \times 10^{-8} \,\mathrm{mol}\,\mathrm{L}^{-1}$

19. CHEMISTRY, M5 2023 HSC 31



From graph:

$$0.66 \ absorbance \ \Rightarrow \ \left[\left[Cu(C_3H_6O_3)_2 \right]^{2+} \right] = 0.046 \ mol \ L^{-1}$$

	Cu^{2+}	$2\mathrm{C_3H_6O_3(aq)}$	$\left[\mathrm{Cu}(\mathrm{C_3H_6O_3})_2 ight]^{2+} \mathrm{(aq)}$
Initial	0.056	0.111	0
Change	-0.046	-0.092	+0.046
Equilibrium	0.010	0.019	0.046

$$egin{aligned} K_{eq} &= rac{\left[\left[\mathrm{Cu}(\mathrm{C_3H_6O_3})_2
ight]^{2+}
ight]}{\left[\mathrm{Cu}^{2+}
ight]\left[\mathrm{C_3H_6O_3}
ight]^2} \ &= rac{0.046}{0.010 imes 0.019^2} \ &= 1.3 imes 10^4 \end{aligned}$$

20. CHEMISTRY, M6 2019 HSC 27

a.
$$K_a imes K_b = K_w \ \Rightarrow \ K_b = \frac{K_w}{K_a}$$
 $K_b = \frac{1.0 imes 10^{-14}}{3.0 imes 10^{-8}}$ $= 3.3 imes 10^{-7}$

$$\textbf{b.} \ \ \mathrm{OCl^{-}}\left(\mathrm{aq}\right) + \mathrm{H}_{2}\mathrm{O}\left(l\right) \ \rightleftharpoons \mathrm{HOCl}\left(\mathrm{aq}\right) + \mathrm{OH^{-}}\left(\mathrm{aq}\right)$$

♦ Mean mark (b) 45%.

	OCl-	HOCl	$\mathrm{OH^-}$
Initial	0.20	0	0
Change	-x	+x	+x
Equilibrium	0.20 - x	x	x

$$K_b = rac{ ext{[HOCl][OH^-]}}{ ext{[OCl^-]}} = rac{x^2}{ ext{(0.20-}x)}$$

Assume $0.20 - x \approx 0.20$ because x is negligible:

$$3.3 \times 10^{-7} = rac{x^2}{0.20 - x}$$
 $x = \sqrt{3.3 \times 10^{-7} \times 0.20}$ $= 2.5690 \times 10^{-4} \text{ mol L}^{-1}$

$$\begin{split} [\text{OH}^-] &= 2.5690 \, \times 10^{-4} \; \text{mol} \; L^{-1} \\ \text{pOH} &= -\log_{10}[\text{OH}^-] = -\log_{10}(2.5690 \, \times 10^{-4}) = 3.59 \\ \therefore \, \text{pH} &= 14 - 3.59 = 10.41 \end{split}$$

21. CHEMISTRY, M5 2020 HSC 35

$$\begin{split} A &= [I_3{}^-]_{eq} \, \times 2.76 \! \times 10^4 \\ 0.745 &= [I_3{}^-]_{eq} \, \times 2.76 \! \times 10^4 \\ [I_3{}^-]_{eq} &= 2.70 \times 10^{-5} \; \text{mol L}^{-1} \end{split}$$

$$egin{aligned} [\mathrm{I}^-\]_{initial} &= 4\ [\mathrm{I}_2\]_{\mathrm{eq}} + 3\ [\mathrm{I}_3^-\]_{\mathrm{eq}} \ & \ 7.00 imes 10^{-4} = 4[\mathrm{I}_2]_{\mathrm{eq}} + \left(3 imes 2.70 imes 10^{-5}
ight) \ & \ [\mathrm{I}_2\]_{\mathrm{eq}} = rac{7.00 imes 10^{-4} - \left(3 imes 2.70 imes 10^{-5}
ight)}{4} \ & = 1.55 imes 10^{-4}\ \mathrm{mol}\ \mathrm{L}^{-1} \end{aligned}$$

$${
m [I^-]}_{
m eq} = 2 {
m [I_2]}_{
m eq} = 2 imes \left(1.55 imes 10^{-4}
ight) = 3.10 imes 10^{-4} \; {
m mol} \; {
m L}^{-1}$$

$$egin{aligned} K_{ ext{eq}} &= rac{igl[ext{I}_3^{-} igr]_{ ext{eq}}}{igl[ext{I}^{-} igr]_{ ext{eq}} imes igl[ext{I}_2 igr]_{ ext{eq}}} \ &= rac{2.70 imes 10^{-5}}{3.10 imes 10^{-4} imes 1.55 imes 10^{-4}} \ &= 564 \end{aligned}$$

22. CHEMISTRY, M5 2021 HSC 31

concentration

Mean mark 55%.

 N_2

 $3\mathrm{H}_2$

 $2NH_3$

♦ Mean mark 44%

$$K_{eq} = rac{\left[NH_3
ight]^2}{\left[N_2
ight]\left[H_2
ight]^3} \ 748 = rac{0.585^2}{rac{4.475 + x}{10} imes 0.0925^3} \ 748 imes rac{4.475 + x}{10} imes 0.0925^3 = 0.585^2 \ rac{4.475 + x}{10} = rac{0.585^2}{748 imes 0.0925^3} \ 4.475 + x = rac{10 imes 0.585^2}{748 imes 0.0925^3} \ x = rac{10 imes 0.585^2}{748 imes 0.0925^3} - 4.475 \ = 1.3 ext{ moles (1 d.p.)}$$

: 1.3 moles of nitrogen must be added to the equilibrium mixture.

23. CHEMISTRY, M5 2024 HSC 30

$$\rightarrow$$
 n_{intial}(CO(g)) = 0.400 and n_{final}(CO(g)) = 0.200.

♦ Mean mark 55%.

_

Change in the number of moles in CO (g) = 0.400 - 0.200 = 0.200 mol in 1 L

	$\mathrm{H}_{2}\left(\mathrm{g}\right)$	$\mathrm{CO}_{2}\left(\mathrm{g} ight)$	$\mathrm{H_{2}O}\left(\mathrm{g}\right)$	$\mathrm{CO}_{2}\left(\mathrm{g} ight)$
Initial	1	0.5	0.4	2 + x
Change	+0.2	+0.2	-0.2	-0.2
Equilibrium	1.2	0.7	0.2	1.8 + x

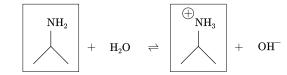
→ Since all substances are present in a 1 L container, the concentrations of each substance is equal to the number of moles of that substance present at equilibrium

$$K_{eq} = rac{[ext{H}_2 ext{O}\,(ext{g})][ext{CO}\,(ext{g})]}{[ext{H}_2\,(ext{g})][ext{CO}_2\,(ext{g})]} \ 1.600 = rac{0.2 imes(1.8+x)}{1.2 imes0.7} \ 1.8+x = 1.6 imesrac{1.2 imes0.7}{0.2} \ x = 6.72-1.8 \ = 4.92 ext{ mol}$$

 \rightarrow 4.92 mol of CO (g) were added to the container.

24. CHEMISTRY, M5 2020 HSC 27

a.



♦ Mean mark (a) 48%.

b.

	$\mathrm{C_{3}H_{7}NH_{2}}$	$C_3H_7NH_3$	OH-
Initial	1.00	0	0
Change	-x	+x	+x
Equilibrium	1.00 - x	x	x

$$K_b = rac{[{
m C_3H_7NH_3}^+][{
m OH}^-\;]}{[{
m C_3H_7NH_2}\;]} = rac{x^2}{(1.00-x)}$$

Assume 1.00 - x = 1.00 because x is negligible:

$$4.37 imes 10^{-4} = rac{x^2}{1.00}$$
 $x = \sqrt{4.37 imes 10^{-4}}$ $= 0.0209 \; ext{mol L}^{-1}$

$$\Rightarrow [OH^-] = 0.0209 \; mol \; L^{-1}$$

Mean mark (b) 51%.

25. CHEMISTRY, M6 2020 HSC 33

$$\begin{array}{l} \textbf{a.} & \operatorname{Ca(OH)_2(s)} + 2\operatorname{HCl}\left(aq\right) \longrightarrow \operatorname{CaCl_2}\left(aq\right) + 2\operatorname{H_2O}\left(l\right) \\ \\ & \operatorname{n(HCl)} = c \times V = 2.00 \times 0.100 \ = 0.200 \ \text{mol} \\ \\ & \operatorname{n(Ca(OH)_2)} = \frac{\operatorname{n(HCL)}}{2} = \frac{0.200}{2} = \ 0.100 \, \text{mol} \end{array}$$

b.
$$Ca(OH)_2(s) \rightleftharpoons Ca^{2+}(aq) + 2OH^-(aq)$$

$$\left[Ca^{2+}\right] = \frac{n}{V} = \frac{0.100}{0.100} = 1.00 \text{ mol } L^{-1}$$

$$\begin{split} K_{sp} &= [Ca^{2+}][OH^-\,]^2 \\ 5.02 \times 10^{-6} &= 1.00 \times [OH^-]^2 \\ [OH^-] &= \sqrt{5.02 \times 10^{-6}} \\ &= 2.24 \times 10^{-3} \; mol \, L^{-1} \\ pOH &= -\log_{10} \left(2.24 \times 10^{-3}\right) \\ &= 2.650 \end{split}$$

$$\therefore$$
 pH = 14 - 2.650 = 11.35

26. CHEMISTRY, M5 2023 HSC 37

a.
$$Q = \frac{[\text{CO}_2]}{[\text{CO}]^2}$$
$$= \frac{1.21 \times 10^{-3}}{(1.10 \times 10^{-2})^2}$$
$$= 10.0$$

Since $Q = K_{eq}$, system is in equilibrium.

b. Given $[CO] = [CO_2]$,

♦♦♦ Mean mark (b) 20%.

$$egin{aligned} K_{eq} &= rac{[ext{CO}_2]}{[ext{CO}]^2} = rac{1}{[ext{CO}]} = 10.00 \ \ &\Rightarrow [ext{CO}] = rac{1}{10.00} = 0.1000 \, ext{mol L}^{-1} \ \ &\Rightarrow [ext{CO}_2] = 0.1000 \, ext{mol L}^{-1} \end{aligned}$$

From this point, the change in ${\rm CO}$ and ${\rm CO}_2$ concentrations can be calculated...

	2 CO (g)	$\mathrm{CO}_{2}\left(\mathrm{g} ight)$	C(s)
Initial	$1.10 imes10^{-2}$	$1.21 imes 10^{-3}$	
Change	+0.0890	+0.0988	
Equilibrium	0.1000	0.1000	

However, the change in moles of ${\rm CO_2}$ in the system consists of:

- Change in CO₂ concentration
- Change in CO concentration (as some of the added CO₂ was converted into CO)

n(CO₂) required to increase [CO] by 0.0988 mol (1 litre vessel)

Formula ratio shows $CO_2: CO = 1 \text{ mol}: 2 \text{ mol}$

 $n(CO_2)$ to add to increase $[CO_2] = 0.0988$ mol (1 litre vessel)

 $n(\mathrm{CO_2})_{total\ to\ add} = 0.0988\ mol + n(\mathrm{CO_2}\ to\ make\ CO)$

$$n(\mathrm{CO_2})$$
 to add to increase $[\mathrm{CO}] = \frac{0.0890}{2} = 0.0445 \; mol$

$$n({
m CO_2})_{
m total\ to\ add} = 0.0988 + 0.0445 = 0.143\ mol$$

♦♦♦ Mean mark (b) 24%.

27. CHEMISTRY, M5 2024 HSC 39

→ The ionisation of bromoacetic acid in water is:

$$BrCH_2COOH(aq) \implies BrCH_2COO^-(aq) + H^+(aq)$$

→ At equilibrium

 $[BrCH_{2}COO^{-}\,(aq)]=[H^{+}\,(aq)]=9.18\times 10^{-3}\ mol\ L^{-1}$ as the are formed in a 1 : 1 .

$$egin{aligned} K_a &= rac{[ext{H}^+][ext{BrCH}_2 ext{COO}^-]}{[ext{BrCH}_2 ext{COOH}]_{ ext{eq}}} \ &= rac{[ext{H}^+][ext{BrCH}_2 ext{COO}^-]}{K_a} \ &= rac{(9.18 imes 10^{-3})^2}{1.29 imes 10^{-3}} \ &= 0.06533 ext{ mol L}^{-1} \end{aligned}$$

◆◆ Mean mark 27%.

COMMENT: Students
who identified the acid
conc in the organic
solvent often succeeded
in this question.

$$\begin{split} [BrCH_{2}COOH]_{total} &= [BrCH_{2}COOH\,(aq)]_{eq} + [BrCH_{2}COO^{-}\,(aq)] + [BrCH_{2}COOH(octan-1-ol)]_{eq} \\ [BrCH_{2}COOH(octan-1-ol)]_{eq} &= 0.1000 - 0.06533 - 9.18 \times 10^{-3} = 0.02549 \, \text{mol} \, L^{-1} \end{split}$$

→ Since the volume of the aqueous solution of bromoacetic acid and octane is the same, the concentration values between the water and octane solutions can be added/subtracted in one equation and mole calculations are not required.

$$K_{eq} = rac{
m [BrCH_2COOH(octan-1-ol)]_{eq}}{
m [BrCH_2COOH\,(aq)]_{eq}} = rac{0.02549}{0.06533} = 0.390~(3~{
m sig.})$$

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