

Mark schemes

1 A

[1]

2 (a) M1 $(K_c =) \frac{[\text{CH}_3\text{CH}_2\text{OH}]}{[\text{CH}_2 = \text{CH}_2][\text{H}_2\text{O}]}$

Penalise missing brackets or use of (); allow correct molecular formulae in correct expression (and allow CH_2CH_2); ignore powers shown as 1

1

M2 $\text{mol}^{-1} \text{dm}^3$

Units must be in simplest form on one line (or $\text{dm}^3 \text{mol}^{-1}$)

Units are consequential on expression in M1 ($\text{mol}^{-1} \text{dm}^3$ only scores if it is the units for the expression in M1)

1

(b) M1 $\frac{[\frac{4.40}{2.00}]}{[\frac{0.70}{2.00}] \times [\frac{1.20}{2.00}]}$ or $\frac{2.20}{0.35 \times 0.60}$ or $\frac{4.40}{0.70 \times 1.20} \times 2.20$

10.5 (3sf) scores both marks;

Correct value to 2sf (10) or 4sf or more (10.476...) scores 1 mark

Volume not used is CE=0

If use incorrect expression for K_c in part (b) then no marks in part (b)

1

M2 10.5 (must be 3sf)

If a value from the question is copied incorrectly into the expression, could still score M2 if then used correctly in calculation (AE -1)

Ignore units

1

[4]

3 C

[1]

4 C

[1]

5 (a) (i) Curve drawn from origin with peak clearly lower and to right.

*New curve crosses original once only, finishes above original and does **not** clearly curve up*

IGNORE relative areas

1

- (ii) (Relative areas under curves indicate) many (owtte) more molecules with E greater than or equal to E_a (at higher T) or reverse argument

ALLOW 'particles'

IGNORE 'atoms'

1

(Large) increase in (number of) successful (owtte) collisions per unit time

OR 'frequency of successful collisions'

1

- (b) (i) Yield increases

Yield decreases/stays the same $CE = 0$

If not answered mark on

1

More moles/molecules (of gas) on left/fewer on right/3 on left 1 on right

1

Equilibrium shifts/moves (to right) to reduce pressure/oppose higher pressure

No M3 if 'more moles on right' in M2

IGNORE 'favours'

NOT just 'oppose the change'

QoL means that M3 is only awarded if these ideas are clearly linked in one statement

1

- (ii) Higher T would increase rate but decrease yield/make less methanol

OR

Lower T decreases rate but increases yield;

*If no mention of both rate **AND** (idea of) yield max 1*

1

Chosen T is a compromise/balance (between rate and yield) owtte

1

[8]

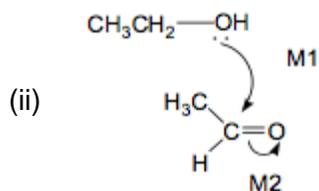
6

- (a) (i) Nucleophilic addition

Any extra loses the mark

Allow minor spelling errors e.g. nucleophyllic

1



M1 for arrow from lone pair on oxygen in ethanol to C of C=O (or to space half way between O and C)

M2 for arrow from C=O bond to oxygen in ethanal

Do not allow M2 as first step without nucleophilic attack, but can allow M1 for attack on C+ produced

+ rather than δ^+ on C=O loses M2

Ignore any further steps

Mark independently

1
1

(b) (i) Equal mixture of enantiomers/optical isomers OWTTE

1

(ii) (Non-superimposable) mirror images

Ignore rotates light in opposite directions

Ignore stereoisomers

1

(c) (i) Ethanal 0.33

1

Ethanol 4.16

Allow 4.2 for ethanol

1

$$(ii) \quad K_c = \frac{[acetal][H_2O]}{[CH_3CHO][CH_3CH_2OH]^2} \text{ or with names}$$

$$\frac{(0.37/0.31)(0.65/0.31)}{(0.58/0.31)(3.76/0.31)^2} \text{ OR } \frac{(0.37)(0.65)}{(0.58)(3.76)^2} \times 0.31$$

Ignore slips in acetal structure or formula $C_6H_{14}O_2$

If K_c wrong, allow M4 only for units conseq to their K_c

If volume omitted (gives 2.93×10^{-2}) may only score M1 and M4

If volume used = 310 cm^3 allow M2 then award M3 for $9.08 - 9.23$ only and M4 for $\text{mol}^{-1} \text{ cm}^3$ only

Treat error in converting 310 cm^3 to dm^3 as AE

M1
M2

$$9.1 \times 10^{-3}$$

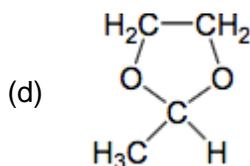
Allow range $9.08 \times 10^{-3} - 9.23 \times 10^{-3}$

M3

$$\text{mol}^{-1}\text{dm}^3$$

Not $\text{moles}^{-1}\text{dm}^3$

M4



1

[12]

7



State symbols essential

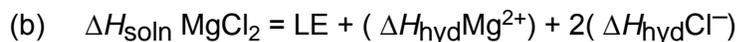
Do not allow this equation with H_2O on the LHS

Ignore + aq on the LHS

Allow H_2O written over the arrow / allow equation written as an equilibrium

Allow correct equations to form $[\text{Mg}(\text{H}_2\text{O})_6]^{2+}$ ions

1



$$\Delta H_{\text{soln}} \text{MgCl}_2 = 2493 - 1920 + (2 \times -364)$$

$$= -155 \text{ (kJ mol}^{-1}\text{)}$$

M1 for expression in words or with correct numbers

Ignore units, but penalise incorrect units

1
1

(c) M1: Solubility decreases (as temp increases)

M2: the enthalpy of solution is exothermic / reaction is exothermic / backwards reaction is endothermic

M3: (According to Le Chatelier) the equilibrium moves to absorb heat/reduce temperature/oppose the increase in temperature (in the endothermic direction)

If M1 is incorrect then CE=0/3

If answer to (b) is a +ve value, allow:

M1: Solubility increases (as temp increases)

M2: Enthalpy of solution is endothermic etc

M3: (According to Le Chatelier) the equilibrium moves to absorb heat/reduce the temperature/oppose the increase in temperature (in the endothermic direction)

1
1
1

[6]

8

(a) amount of X = 0.50 – 0.20 = 0.30 (mol)

1

amount of Y = 0.50 – 2 × 0.20 = 0.10 (mol)

1

(b) Axes labelled with values, units and scales that use over half of each axis

All three of values, units and scales are required for the mark

1

Curve starts at origin

1

Then flattens at 30 seconds at 0.20 mol

1

(c) Expression = $K_c = \frac{[Z]}{[X][Y]^2}$

1

$[Y]^2 = \frac{[Z]}{[X] K_c}$

1

$[Y] = (0.35 / 0.40 \times 2.9)^{0.5} = 0.5493 = 0.55 \text{ (mol dm}^{-3}\text{)}$

Answer must be to 2 significant figures

1

(d) Darkened / went more orange

1

The equilibrium moved to the right

1

To oppose the increased concentration of Y

1

(e) The orange colour would fade

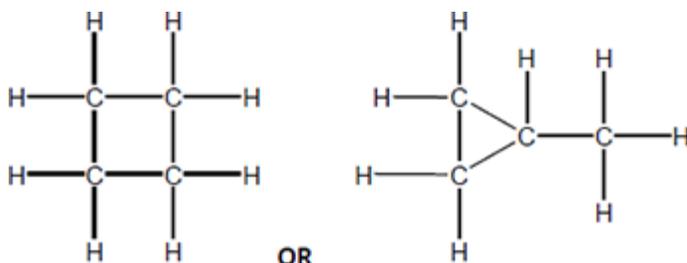
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[12]

9

(a) Alkenes

1



OR
*Correctly drawn molecule of cyclobutane or methyl cyclopropane,
need not be displayed formula*

1

(b) C_6H_{14} (or correct alkane structure with 6 carbons)

Allow hexane or any other correctly named alkane with 6 carbons

1

(c) Poly(but-2-ene)

1

(d) High pressure

Allow pressure \geq MPa

Mention of catalyst loses the mark

1

(e) This question is marked using levels of response. Refer to the Mark Scheme Instructions for Examiners for guidance on how to mark this question.

Level 3

All stages are covered and the explanation of each stage is generally correct and virtually complete.

Answer communicates the whole process coherently and shows a logical progression from stage 1 and stage 2 (in either order) to stage 3.

5–6 marks

Level 2

All stages are covered but the explanation of each stage may be incomplete or may contain inaccuracies OR two stages are covered and the explanations are generally correct and virtually complete.

Answer is mainly coherent and shows progression. Some steps in each stage may be out of order and incomplete.

3–4 marks

Level 1

Two stages are covered but the explanation of each stage may be incomplete or may contain inaccuracies, OR only one stage is covered but the explanation is generally correct and virtually complete.

Answer includes isolated statements but these are not presented in a logical order or show confused reasoning.

1–2 marks

Level 0

Insufficient correct chemistry to gain a mark.

0 marks

Indicative chemistry content

Stage 1: consider effect of higher temperature on yield

(Or vice versa for lower temperature)

- Le Chatelier's principle predicts that equilibrium shifts to oppose any increase in temperature
- Exothermic reaction, so equilibrium shifts in endothermic direction / to the left
- So a Higher T will reduce yield

Stage 2: consider effect of higher temperature on rate

(Or vice versa for lower temperature)

- At higher temperature, more high energy molecules
- more collisions have $E > E_a$
- So rate of reaction increases / time to reach equilibrium decreases

Stage 3: conclusion

Industrial conditions chosen to achieve (cost-effective) balance of suitable yield at reasonable rate

[11]

10

(a) Bonds broken = $2(\text{C}=\text{O}) + 3(\text{H}-\text{H}) = 2 \times 743 + 3 \times \text{H}-\text{H}$

Bonds formed = $3(\text{C}-\text{H}) + (\text{C}-\text{O}) + 3(\text{O}-\text{H}) = 3 \times 412 + 360 + 3 \times 463$

Both required

1

$$-49 = [2 \times 743 + 3 \times (\text{H}-\text{H})] - [3 \times 412 + 360 + 3 \times 463]$$

$$3(\text{H}-\text{H}) = -49 - 2 \times 743 + [3 \times 412 + 360 + 3 \times 463] = 1450$$

Both required

1

$$\text{H}-\text{H} = 483 \text{ (kJ mol}^{-1}\text{)}$$

Allow 483.3(3)

1

(b) Mean bond enthalpies are not the same as the actual bond enthalpies in CO₂ (and / or methanol and / or water)

1

(c) The carbon dioxide (produced on burning methanol) is used up in this reaction

1

(d) 4 mol of gas form 2 mol

1

At high pressure the position of equilibrium moves to the right to lower the pressure / oppose the high pressure

1

This increases the yield of methanol

1

(e) Impurities (or sulfur compounds) block the active sites

Allow catalyst poisoned

1

(f) Stage 1: moles of components in the equilibrium mixture

Extended response question



Initial moles	1.0	3.0	0	0
Eqm moles	(1-0.86) = 0.14	(3-3×0.86) = 0.42	0.86	0.86

1

Stage 2: Partial pressure calculations

Total moles of gas = 2.28

Partial pressures = mol fraction × p_{total}

1

$$p_{\text{CO}_2} = \text{mol fraction} \times p_{\text{total}} = 0.14 \times 500 / 2.28 = 30.7 \text{ kPa}$$

$$p_{\text{H}_2} = \text{mol fraction} \times p_{\text{total}} = 0.42 \times 500 / 2.28 = 92.1 \text{ kPa}$$

M3 is for partial pressures of both reactants

Alternative M3 =

$$pp_{\text{CO}_2} = 0.0614 \times 500$$

$$pp_{\text{H}_2} = 0.1842 \times 500$$

1

$$p_{\text{CH}_3\text{OH}} = \text{mol fraction} \times p_{\text{total}} = 0.86 \times 500 / 2.28 = 188.6 \text{ kPa}$$

$$p_{\text{H}_2\text{O}} = \text{mol fraction} \times p_{\text{total}} = 0.86 \times 500 / 2.28 = 188.6 \text{ kPa}$$

M4 is for partial pressures of both products

Alternative M4 =

$$pp_{\text{CH}_3\text{OH}} = 0.3772 \times 500$$

$$pp_{\text{H}_2\text{O}} = 0.3772 \times 500$$

1

Stage 3: Equilibrium constant calculation

$$K_p = p_{\text{CH}_3\text{OH}} \times p_{\text{H}_2\text{O}} / p_{\text{CO}_2} \times (p_{\text{H}_2})^3$$

1

$$\text{Hence } K_p = 188.6 \times 188.6 / 30.7 \times (92.1)^3 = 1.483 \times 10^{-3} = 1.5 \times 10^{-3}$$

Answer must be to 2 significant figures

1

$$\text{Units} = \text{kPa}^{-2}$$

1

[16]

11

(a) Stage 1: Moles of acid at equilibrium

Moles of sodium hydroxide in each titration

$$= (3.20 \times 2.00 \times 10^{-1}) / 1000 = 6.40 \times 10^{-4}$$

Extended response

1

Sample = 10 cm³ so moles of acid in 250 cm³ of equilibrium mixture

$$= 25 \times 6.40 \times 10^{-4} = 1.60 \times 10^{-2}$$

M2 can only be scored if = answer to M1 x 25

1

Stage 2: Moles of ester and water formed

$$\text{Moles of acid reacted} = 8.00 \times 10^{-2} - 1.60 \times 10^{-2} = 6.40 \times 10^{-2}$$

= moles ester and water formed

M3 is $8.00 \times 10^{-2} - M2$

1

Stage 3: Moles of ethanol at equilibrium

$$\text{Moles of ethanol remaining} = 1.20 \times 10^{-1} - 6.40 \times 10^{-2} = 5.60 \times 10^{-2}$$

M4 is $1.20 \times 10^{-1} - M3$

1

Stage 4: Calculation of equilibrium constant

$$K_c = [\text{CH}_3\text{COOCH}_2\text{CH}_3] [\text{H}_2\text{O}] / [\text{CH}_3\text{COOH}] [\text{CH}_3\text{CH}_2\text{OH}]$$

1

$$= (6.40 \times 10^{-2})^2 / (1.60 \times 10^{-2})(5.60 \times 10^{-2})$$

$$= 4.5714 = 4.57$$

M6 is M3² / M2 × M4

Answer must be given to 3 significant figures

1

(b)

	Rough	1	2	3
Final burette reading / cm³	4.60	8.65	12.85	16.80
Initial burette reading / cm³	0.10	4.65	8.65	12.85
Titre / cm³	4.50	4.00	4.20	3.95

1

(c) Mean = $4.00 + 3.95 / 2 = 3.98$ (cm³)

Allow 3.975 (cm³)

1

Titres 1 and 3 are concordant

Allow titre 2 is not concordant

1

(d) Thymol blue

1

(e) Percentage uncertainty: $0.15 / 3.98 \times 100 = 3.77\%$

Allow consequential marking on mean titre from 2.3

1

(f) Use a lower concentration of NaOH

1

So that a larger titre is required (reduces percentage uncertainty in titre)

1

[13]

12

A

[1]