

AQA Qualifications

## A-level **Chemistry**

Paper 2 (7405/2): Organic and Physical Chemistry Mark scheme

7405 Specimen paper

Version 0.5

Question	Marking guidance	Mark	AO	Comments
01.1	Consider experiments 1 and 2: [B constant]			
	[A] increases × 3: rate increases by 3 <sup>2</sup> therefore 2nd order with respect to A	1	AO3 1a	
	Consider experiments 2 and 3:			
	[A] increases × 2: rate should increase × 2 <sup>2</sup> but only increases × 2			
	Therefore, halving [B] halves rate and so 1st order with respect to B	1	AO3 1a	
	Rate equation: $rate = k[A]^2[B]$	1	AO3 1b	
01.2	rate = $k [C]^2 [D]$ therefore $k = \text{rate} / [C]^2 [D]$	1	AO2h	
	$k = \frac{7.2 \times 10^{-4}}{(1.9 \times 10^{-2})^2 \times (3.5 \times 10^{-2})} = 57.0$	1	AO2h	Allow consequential marking on incorrect transcription
	$mol^{-2} dm^{+6} s^{-1}$	1	AO2h	Any order
01.3	rate = $57.0 \times (3.6 \times 10^{-2})^2 \times 5.4 \times 10^{-2} = 3.99 \times 10^{-3} \text{ (mol dm}^{-3} \text{ s}^{-1})$ <b>OR</b>	1	AO2h	
	Their $k \times (3.6 \times 10^{-2})^2 \times 5.4 \times 10^{-2}$			

01.4	Reaction occurs when molecules have $E \ge E_a$ Doubling T causes many more molecules to have this $E$ Whereas doubling [E] only doubles the number with this $E$	1 1 1	AO1a AO1a AO1a	
01.5	$E_a = RT(\ln A - \ln k)/1000$	1	AO1b	Mark is for rearrangement of equation and factor of 1000 used correctly to convert J into kJ
	$E_{\rm a} = 8.31 \times 300 (23.97 - (-5.03))/1000 = 72.3 (kJ mol^{-1})$	1	AO1b	

Question	Marking guidance	Mark	AO	Comments
02.1	Marking guidance  Gradient drawn on graph  0.020 0.018 0.016 0.014 0.012  [Butadiene] / mol dm <sup>-3</sup> 0.010 0.008 0.006 0.004 0.002 0.000 0 1000 2000 3000 4000 5000 6000 7000 8000 9000 Time / s	Mark 1	AO3 1a	Line must touch the curve at 0.012 but must not cross the curve.
	Timers			

02.2				Extended response
02.2	Stage 1: Rate of reaction when concentration = 0.0120 mol dm <sup>-3</sup> From the tangent			Extended response
	Change in [butadiene] = $-0.0160 - 0$ and change in time = $7800 - 0$ Gradient = $-(0.0160 - 0)/(7800 - 0) = -2.05 \times 10^{-6}$	1	AO3 1a	
	Rate = $2.05 \times 10^{-6}$ (mol dm <sup>-3</sup> s <sup>-1</sup> )	1	AO3 1a	
	Stage 2: Comparison of rates and concentrations Initial rate/rate at $0.0120 = (4.57 \times 10^{-6})/(2.05 \times 10^{-6}) = 2.23$ Inital concentration/concentration at point where tangent drawn = $0.018/0.012 = 1.5$	1	AO3 1a AO3 1a	Marking points in stage 2 can be in either order
	Stage 3: Deduction of order  If order is 2, rate should increase by factor of $(1.5)^2 = 2.25$ this is approximately equal to 2.23 therefore order is 2nd with respect to butadiene	1	AO3 1b	

Question	Marking guidance	Mark	AO	Comments
03.1	2,2,4-trimethylpentane	1	AO1a	
03.2	5	1	AO2b	
03.3	$C_{20}H_{42} \longrightarrow C_8H_{18} + 2C_3H_6 + 3C_2H_4$	1	AO2b	
03.4	Mainly alkenes formed	1	AO1b	
03.5	4 (monochloro isomers)	1	AO2b	
	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	1	AO2a	
03.6	CI	1	AO2a	

03.7	$C_8H_{17}^{35}CI = 96.0 + 17.0 + 35.0 = 148.0$ and $C_8H_{17}^{37}CI = 96.0 + 17.0 + 37.0 = 150.0$	1	AO1b	Both required
	$M_{\rm r}$ of this C <sub>8</sub> H <sub>17</sub> Cl $(1.5 \times 148.0) + (1.0 \times 150.0) = 148.8$ 2.5 2.5	1	AO1b	
03.8	<u>24.6</u> <u>2.56</u> <u>72.8</u> = 2.05 : 2.56 : 2.05 12 1 35.5			
	Simplest ratio = $\frac{2.05}{2.05}$ : $\frac{2.56}{2.05}$ : $\frac{2.05}{2.05}$			
	= 1 : 1.25 : 1	1	AO2b	
	Whole number ratio ( $\times$ 4) = 4 : 5 : 4	1	AO2b	
	$MF = C_8 H_{10} CI_8$	1	AO2b	

Question	Marking guidance	Mark	AO	Comments
04.1	3-methylbutan-2-ol	1	AO1a	
04.2	CH <sub>3</sub> H <sub>3</sub> C—C—C—CH <sub>3</sub> H O	1	AO2g	Allow (CH <sub>3</sub> ) <sub>2</sub> CHCOCH <sub>3</sub>
04.3	Elimination	1	AO1a	
04.4	CH <sub>3</sub>   H <sub>3</sub> C—C=C-CH <sub>3</sub>   H	1	AO2g	Allow (CH <sub>3</sub> ) <sub>2</sub> C=CHCH <sub>3</sub>
	CH <sub>3</sub>   H <sub>3</sub> C—C—C—CH <sub>2</sub>   H H	1	AO2g	Allow (CH <sub>3</sub> ) <sub>2</sub> CHCH=CH <sub>2</sub>

04.5	Position	1	AO1a	
04.6	СВА	1	AO3 1b	
04.7	H <sub>3</sub> C——CH <sub>2</sub> CH <sub>3</sub> OH	1	AO2g	Allow (CH <sub>3</sub> ) <sub>2</sub> C(OH)CH <sub>2</sub> CH <sub>3</sub>
04.8	$H_3C$ — $C$ — $CH_2OH$ $CH_3$	1	AO2e	Allow (CH <sub>3</sub> ) <sub>3</sub> CCH <sub>2</sub> OH

Question	Marking guidance	Mark	AO	Comments
05.1	Secondary	1	AO1a	
05.2	Nitrogen and oxygen are very electronegative Therefore, C=O and N-H are polar Which results in the formation of a hydrogen bond between O and H In which a lone pair of electrons on an oxygen atom is strongly attracted to the $\delta\text{+H}$	1 1 1 1	AO1a AO1a AO1a AO1a	

Question	Marking guidance	Mark	AO	Comments
06.1	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	1	AO2a	
06.2	+	1	AO2a	
06.3	CH <sub>3</sub> H +	1	AO2a	Allow $(CH_3)_3$ $\stackrel{+}{N}$ $-CH_2$ $-COOH$ $(Br)$
06.4	2-amino-3-hydroxybutanoic acid	1	AO2a	

06.5	+ ŅH <sub>3</sub>	1	AO2a	
	 (CH <sub>2</sub> ) <sub>4</sub>			
	H <sub>3</sub> N—C—COOH			
	н			

Question	Marking guidance	Mark	AO	Comments
07.1	H CH <sub>3</sub>	1	AO1a	
	Ċ──Ċ I I CH <sub>3</sub> CI			
	Addition	1	AO1a	
07.2	ų ų	1	AO2e	
	H H   HO-C-C-OH   CH <sub>3</sub> CH <sub>3</sub>			
	O CH <sub>3</sub> H O O CH <sub>3</sub>	1	AO2e	
07.3	<b>Q</b> is biodegradable	1	AO2g	
	Polar C=O group or $\delta$ + C in <b>Q</b> (but not in <b>P</b> )	1	AO2c	
	Therefore, can be attacked by nucleophiles (leading to breakdown)	1	AO2c	

Question	Marking guidance	Mark	AO	Comments
08.1	2-deoxyribose	1	AO1a	
08.2	Base A Top N–H forms hydrogen bonds to lone pair on O of guanine The lone pair of electrons on N bonds to H–N of guanine A lone pair of electrons on O bonds to lower H–N of guanine	1 1 1 1	AO3 1b AO2a AO2a AO2a	If Base B stated, allow 1 mark only for response including hydrogen bonding  Allow all 4 marks for a correct diagram showing the hydrogen bonding  Students could also answer this question using labels on the diagram
08.3	Allow either of the nitrogen atoms with a lone pair NOT involved in bonding to cytosine	1	AO2a	
08.4	Use in very small amounts / target the application to the tumour	1	AO2e	

Question	Marking guidance	Mark	AO	Comments
09.1	(nucleophilic) addition-elimination	1	AO1a	Not electrophilic addition-elimination  Allow C <sub>6</sub> H <sub>5</sub> or benzene ring
	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	4	AO2a	Allow attack by :NH <sub>2</sub> C <sub>6</sub> H <sub>5</sub> M2 not allowed independent of M1, but allow M1 for correct attack on C+  M3 for correct structure with charges but lone pair on O is part of M4  M4 (for three arrows and lone pair) can be shown in more than one structure

09.2	The minimum quantity of hot water was used:			
	To ensure the hot solution would be saturated / crystals would form on cooling	1	AO1b	
	The flask was left to cool before crystals were filtered off:			
	Yield lower if warm / solubility higher if warm	1	AO1b	
	The crystals were compressed in the funnel:			
	Air passes through the sample not just round it	1	AO1b	Allow better drying but not water squeezed out
	A little cold water was poured through the crystals:			
	To wash away soluble impurities	1	AO1b	
09.3	Water	1	AO3 1b	Do not allow unreacted reagents
	Press the sample of crystals between filter papers	1	AO3 2b	Allow give the sample time to dry in air
09.4	$M_{\rm r}$ product = 135.0	1	AO2h	
	Expected mass = $5.05 \times \frac{135.0}{93.0} = 7.33 \text{ g}$	1	AO2h	
	Percentage yield = $\frac{4.82}{7.33} \times 100 = 65.75 = 65.8(\%)$	1	AO1b	Answer must be given to this precision

09.5	NHCOCH <sub>3</sub> $+ NO_2^+ \longrightarrow NHCOCH_3$ $+ NO_2^+ \longrightarrow NO_2$ OR $C_6H_5NHCOCH_3 + NO_2^+ \rightarrow C_6H_4(NHCOCH_3)NO_2 + H^+$	1	AO2c	
09.6	Electrophilic substitution	1	AO1a	
09.7	Hydrolysis	1	AO3 1a	
09.8	Sn/HCI	1	AO1b	Ignore acid concentration; allow Fe/HCI

Question		Marking guidance	Mark	AO	Comments
10	IR				Extended response
	M1	Absorption at 3360 cm <sup>-1</sup> shows OH alcohol present	1	AO3 1a	Deduction of correct structure without explanation scores maximum of 4 marks as this does not show a
	NMR				clear, coherent line of reasoning.
	M2	There are 4 peaks which indicates 4 different environments of hydrogen	1	AO3 1a	Maximum of 6 marks if no structure given OR if coherent logic not displayed in the explanations of
	МЗ	The integration ratio = 1.6 : 0.4 : 1.2 : 2.4	1	AO3 1a	how two of OH, CH <sub>3</sub> and CH <sub>2</sub> CH <sub>3</sub> are identified.
		The simplest whole number ratio is 4 : 1 : 3 : 6			
	M4	The singlet (integ 1) must be caused by H in OH alcohol	1	AO3 1a	
	M5	The singlet (integ 3) must be due to a $\text{CH}_3$ group with no adjacent H	1	AO3 1b	
	M6	Quartet + triplet suggest CH <sub>2</sub> CH <sub>3</sub> group	1	AO3 1b	
	M7	Integration 4 and integration 6 indicates two equivalent CH <sub>2</sub> CH <sub>3</sub> groups	1	AO3 1b	
		ÇH₂CH₃			
	M8	H <sub>3</sub> C—C—OH	1	AO3 1b	
		CH <sub>2</sub> CH <sub>3</sub>			

Question		Marking guidance	Mark	AO	Comments
11.1	$CH_3CH_2COCH_3 + 2[H] \longrightarrow CH_3CH_2CH(OH)CH_3$		1	AO1b	
11.2		All stages are covered and the explanation of each stage is generally correct and virtually complete.  Answer is communicated coherently and shows a logical progression from stage 1 to stage 2 then stage 3.  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 from stage 1 to stage 3.  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.  Insufficient correct chemistry to gain a mark.	6	1 AO1a 5 AO2a	Indicative Chemistry content  Stage 1: Formation of product  Nucleophilic attack Planar carbonyl group Hattacks from either side (stated or drawn)  Stage 2: Nature of product Product of step 1 shown This exists in two chiral forms (stated or drawn) Equal amounts of each enantiomer/racemic mixture formed  Stage 3: Optical activity Optical isomers/enantiomers rotate the plane of polarised light equally in opposite directions With a racemic/equal mixture the effects cancel

Question	Marking guidance	Mark	AO	Comments
12.1	HBr OR HCI OR H <sub>2</sub> SO <sub>4</sub>	1	AO1b	Allow HI or HY
12.2	Electrophilic addition	1	AO1a	
	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	4	AO2a	Allow consequential marking on acid in 12.1 and allow use of HY
12.3	The major product exists as a pair of enantiomers	1	AO2a	
	The third isomer is 1-bromobutane (minor product)	1	AO2a	
	Because it is obtained via primary carbocation	1	AO2a	