

3(a)	heterogeneous AND MnO ₂ is in a different state / phase to the reactants / H ₂ O ₂
3(b)(i)	M1: (expt 2 and 3) [I⁻] × 2 initial rate × 2 so 1st order wrt [I⁻] M2: (expt 1 and 2) [I⁻] × 2, [H₂O₂] × 0.5 initial rate × 1 OR (expt 1 and 4) [I⁻] × 4 initial rate × 4 AND [H⁺] × 4 OR (expt 1 and 3) [I⁻] × 4, [H₂O₂] × 0.5 initial rate × 2 so 1st order wrt [H₂O₂] M3: (expt 1 and 4) [I⁻] × 4, [H⁺] × 4 initial rate × 4 so zero order wrt [H⁺] OR (expt 3 and 4) [H₂O₂] × 2 initial rate × 2 AND [H⁺] × 4 so zero order wrt [H⁺] OR (expt 3 and 4) [H₂O₂] × 2, [H⁺] × 4 initial rate × 2 so 1st order wrt [H₂O₂] M4: rate = k [H₂O₂] [I⁻] OR rate = k [H₂O₂] [I⁻] [H⁺]⁰
3(b)(ii)	M1: $k = 2.42 \times 10^{-3} \div (0.045 \times 0.030) = 1.79 \text{ min 2sf}$ M2: units = dm³ mol⁻¹ s⁻¹
3(c)(i)	M1: calculation of one $t_{1/2} = 60 \text{ s} \pm 5 \text{ s}$ M2: two $t_{1/2}$ calculated that are constant
3(c)(ii)	$k = \ln 2 \div 60 = 0.0116$ OR $k = 0.693 \div 60 = 0.0116 \text{ min 2sf}$
3(d)	increases k and increases the rate of reaction

2(a)(i)	homolytic fission
2(a)(ii)	$\cdot\ddot{\text{O}}:\ddot{\text{N}}:\ddot{\text{O}} \quad \text{H}:\ddot{\text{O}}:\ddot{\text{N}}:\ddot{\text{O}}\cdot$
2(a)(iii)	115–120°
2(b)	M1: $3.825 \times 10^{-3} / 0.003825 / 0.00383 / 0.0038 / 3.8 \times 10^{-3}$ M2: units = mol⁻¹ dm³ s⁻¹
2(c)	not constant half-life because overall second order
2(d)	8.17 / 8.2
2(e)	- reactants are adsorbed onto the catalyst surface - bonds within reactant molecules are weakened - products are desorbed from catalyst surface Any two [1], all three [2]

4(a)(i)	<table border="1"> <tr> <td>the order of reaction with respect to [H₂]</td><td>1</td></tr> <tr> <td>the order of reaction with respect to [NO]</td><td>2</td></tr> <tr> <td>overall order of the reaction</td><td>3</td></tr> </table>	the order of reaction with respect to [H ₂]	1	the order of reaction with respect to [NO]	2	overall order of the reaction	3
the order of reaction with respect to [H ₂]	1						
the order of reaction with respect to [NO]	2						
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4(a)(ii)	rate $\times \frac{1}{4}$						
4(a)(iii)	rate $\times 27$						
4(a)(iv)	a four-particle collision is unlikely						
4(a)(v)	step 1 2NO → N₂O₂ step 2 N₂O₂ + H₂ → N₂O + H₂O step 3 N₂O + H₂ → N₂ + H₂O						
4(a)(vi)	intermediate formed (step 2) then used up later (step 3)						
4(b)	$k = 0.693 / 720 = 9.625 \times 10^{-4} \text{ min 2sf}$ AND units: s ⁻¹						
4(c)(i)	homogeneous AND Co ³⁺ / catalyst / it is in the same phase / same state as the reactants						
4(c)(ii)	M1: (equation 1) 2Co³⁺ + 2I⁻ → 2Co²⁺ + I₂ M2: (equation 2) 2Co²⁺ + S₂O₈²⁻ → 2Co³⁺ + 2SO₄²⁻						
4(c)(iii)	repulsion of two negative / same charge ions slows the reaction / raises E_{A}						

2(a)(i)	to increase the rate of drying / removal of (distilled) water														
2(a)(ii)	iodine is not (very) soluble (in water)														
2(a)(iii)	mass change is (too / very) small														
2(a)(iv)	repeat without any iodine														
2(b)	<table border="1"> <tr> <th>time / s</th><th>mass of tin reacted / g</th></tr> <tr> <td>0</td><td>0.000</td></tr> <tr> <td>100</td><td>0.069</td></tr> <tr> <td>200</td><td>0.120</td></tr> <tr> <td>300</td><td>0.218</td></tr> <tr> <td>400</td><td>0.291</td></tr> <tr> <td>500</td><td>0.359</td></tr> </table>	time / s	mass of tin reacted / g	0	0.000	100	0.069	200	0.120	300	0.218	400	0.291	500	0.359
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2(c)(i)	M1 points plotted correctly <i>centre of all crosses should be on line for $\times$ value.</i> <i>y value should be on the line for point 1, 3</i> <i>point 2, 6 – just below line / on line</i> <i>4 – midway between two lines</i> <i>5 – just above line / on line</i>
	M2 straight line of best fit drawn
2(c)(ii)	most anomalous point circled (expected to be at time 200 s, point 3, below line of best fit) AND the balance/mass reading was taken before 200 s / too early
2(d)	M1 two acceptable coordinates from line of best fit expressed in the form (x,y). M2 gradient correctly calculated from points listed for M1. answer correctly rounded to 3 significant figures.

2(e)(i)	<table> <tr> <th>volume of 0.400 mol dm⁻³ I₂ solution used / cm³</th><th>volume of methylbenzene used / cm³</th></tr> <tr> <td>100.0</td><td>0.0</td></tr> <tr> <td>75.0</td><td>25.0</td></tr> <tr> <td>50.0</td><td>50.0</td></tr> <tr> <td>25.0</td><td>75.0</td></tr> </table>	volume of 0.400 mol dm ⁻³ I ₂ solution used / cm ³	volume of methylbenzene used / cm ³	100.0	0.0	75.0	25.0	50.0	50.0	25.0	75.0
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2(e)(ii)	[I ₂]										
2(e)(iii)	M1 no AND the rate of reaction is proportional to the concentration of iodine / a conclusion based on calculations done for M2 M2 explained using actual values from columns 3 and 4 in Table 2.2										