

Question	Answers	Extra information	Mark	AO Spec reference
01.1	circle drawn around CH <sub>3</sub> ;	Accept a clear annotation that indicates $\mathrm{CH}_3$	1	AO1 3.1.4.1
01.2	(R <sub>f</sub> =) 0.38;	The correct answer of 0.38 should be awarded 2 marks even when no working is shown Accept any value in the range 0.36 to 0.40. If the final answer is incorrect, 1 mark should be awarded for dividing any value by 5 (cm).	2	AO2 3.1.4.1 AT g
01.3	R <sub>f</sub> = 0.13; glutamine;		2	AO2 3.1.4.1 AT g
01.4	similar chemical properties; similar R groups; similar solubility;	Accept all other suitable answers	2 max	AO2 3.1.4.1 AT g
01.5	leave a larger gap between the pencil line and the solvent; add cap to jar (to prevent solvent evaporation); monitor solvent rather than leaving it for a set time;		2 max	AO3 3.1.4.1 AT g
02.1	sodium hydroxide; volumes; purple / violet;		3	AO1 3.1.4.1 AT f
02.2	amine of one amino acid reacts with carboxyl group of another amino acid; condensation reaction <b>OR</b> water released / produced;	Accept 'amino' for 'amine' and 'carboxylic acid' for 'carboxyl group'.	2	AO1 3.1.4.1

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Question	Answers	Extra information	Mark	AO Spec reference
02.3	peptide bond shown between the amine group of aspartic acid and the carboxyl group of lysine; peptide bond shown between the amine group of lysine and the carboxyl group of aspartic acid;;	Award one mark for a bond using the NH2 in lysine's R group and/ or the COOH in aspartic acid's R group.	3	AO2 3.1.4.1
03.1	C, H, O, N; S;	Accept names or chemical symbols for each element	2	AO2 3.1.4.1
03.2	(34350 × 3) = 103050 (RNA nucleotides)	Accept 103 053 (if a stop codon is referenced) or 103 056 (if a stop codon and a start codon are referenced)	1	AO2 3.4.2
03.3	<pre>primary structure is a polypeptide containing 348 amino acids; secondary structure has alpha helices; formed by hydrogen bonds; tertiary structure has disulphide bonds; (rhodopsin is) a glycoprotein / conjugated; retinal is a cofactor / prosthetic group; <i>idea that</i> (rhodopsin must have) hydrophobic regions (within the membrane) and hydrophilic regions (either side of the membrane);</pre>	The descriptions should be linked to the correct level of protein structure for marks 1–4 to be awarded.	5 max	AO2 3.1.4.1
03.4	active site <b>or</b> specific 3D shape (which can be inferred from the complementary binding of ADP); <i>idea of</i> subunits changing shape for induced fit; hydrophilic regions on the exterior (i.e. exposed to water and dissolved ADP, phosphate and H <sup>+</sup> ions);		2 max	AO2 3.1.4.2

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Question	Answers	Extra information	Mark	AO Spec reference
03.5	tertiary;		1	AO2 3.1.4.1 3.1.4.2
03.6	hydrogen bonds break; enzyme denatures; tertiary structure changed; active site no longer complementary to substrate / ADP;		3 max	AO1 3.1.4.2
04.1	<i>idea of</i> dilute 1 part protein solution to 4 parts water;		1	AO2 3.1.4.1 AT b and c
04.2	use a clean pipette each time; remember or write down the order of the solutions (because cuvettes cannot be labelled);		1 max	AO3 3.1.4.1 AT b and c
04.3	the biuret test produces a purple colour; (which means) green light is absorbed; (using the filter) increases the resolution of the results;		2 max	AO3 3.1.4.1 AT b and c
04.4	<i>x</i> -axis labelled 'percentage (concentration of) protein (solution)' <b>AND</b> <i>y</i> -axis labelled 'absorbance / AU'; straight line from origin showing a positive correlation;		2	AO2 3.1.4.1 AT b and c
04.5	protein solution concentration / volume; temperature; volume of protease solution; biuret solution concentration / volume; pH;		2 max	AO3 3.1.4.1 AT b and c
05.1	$1.58 imes 10^{-7}$ (mol dm <sup>-3</sup> );	Allow one mark for 0.000 000 158 (not in standard form)	2	AO2 3.1.4.2

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Question	Answers	Extra information	Mark	AO Spec reference
05.2	<b>temperature</b> more kinetic energy; hydrogen bonds break;		4 max	AO1 3.1.4.2
	<ul> <li>pH</li> <li>change in charges;</li> <li>hydrogen and ionic bonds break;</li> <li>general</li> <li>active site/tertiary structure changes shape;</li> </ul>			
05.3	competitive inhibitor binds to active site and non-competitive inhibitor binds to allosteric site; non-competitive inhibitor changes tertiary structure; non-competitive inhibitor lowers V <sub>max</sub> ;	Allow 'site other than active site' for 'allosteric site' Or reverse argument Or reverse argument Do not allow reversible/ irreversible because these terms can apply to either inhibitor	3	AO1 3.1.4.2
06.1	curved line below the original line; new line labelled as 'lower activation energy with enzyme';	Allow alternative wording	2	AO2 3.1.4.2
06.2	(original model was) lock and key; active site and substrate(s) have specific and complementary shapes; (new model is) induced fit;		4 max	AO1 3.1.4.2
	providing a better fit;			

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Question	Ans	wers		Extra information	Extra information Mark
07	The following are suitable topic areas fro to describe the formation of polymers. A monomers. Lipids are therefore not cons Please note that to obtain full credit, stud the formation of polymers, not just write These topics can include the enzymes an formation, when relevant. In order to full highest mark bands students must also in to demonstrate a synoptic approach to th		25		
	Specification reference	Topic area			
	3.1.1	Monomers and polymers			
	3.1.2	Carbohydrates			
	3.1.4.1	General properties of proteins			
	3.1.4.2	Many proteins are enzymes			
	3.1.5.1	Structure of DNA and RNA			
	3.1.5.2	DNA replication			
	3.4.2	DNA and protein synthesis			
	Students may be able to show the releva <b>Note:</b> other topics from beyond the spec relate to the title and contain factually co standard. Credit should not be given for t below A-level standard.	nce of other topics from the specification. ification can be used, providing they prrect material of at least an A-level topics beyond the specification which are			

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#### **Skills box answers**





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