

## INTERNATIONAL AS BIOLOGY (9610) BL01

Unit 1 The Diversity of Living Organisms

Mark scheme

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Question	Marking guidance	Mark	Comments
01.1	(Glucose) 6;	2	
	(Ribose) 5;		

Question	Marking guidance	Mark	Comments
01.2	Nitrogen;	1	Ignore N

Question	Marking guidance	Mark	Comments
01.3	R-groups/side chains are <b>different</b> / contain <b>different</b> numbers of carbon atoms;	1	

Question	Marking guidance	Mark	Comments
01.4	Heat with Benedict's solution <b>to</b> confirm a negative result / sucrose is not a reducing sugar;	3	For 'confirm a negative result' accept stays blue or no colour change
	2. Boil/heat (sucrose) with an acid then <b>neutralise</b> with alkali;		2. Accept named examples of acids/alkalis
	3. Heat with Benedict's solution and red/orange colour indicates		3. Award only if mp2 has been attempted
	sucrose is a non-reducing sugar;		3. For 'heat' ignore 'warm'/'heat gently'/'put in a water bath' but accept stated temperatures ≥60 °C
			Heat must be stated again, do not accept using residual heat from mp2

Question	Marking guidance	Mark	Comments
02.1	1. <b>A</b> = Phosphate (group);		1. Reject phosphorus
			1. Accept P <sub>i</sub> /PO <sub>4</sub> <sup>3-</sup>
	2. <b>B</b> = Deoxyribose;		2. Ignore pentose/sugar
		3	2. Reject ribose
	3. <b>C</b> = Nitrogen-containing/nitrogenous/organic base;		Accept Adenine and Thymine and Cytosine and Guanine
			3. Reject uracil

Question	Marking guidance	Mark	Comments
02.2	1088 / 1.088 x 10 <sup>3</sup> (mm);	2	Allow 1.09 x 10 <sup>3</sup> (mm);
			One mark for correct answer given in µm
			One mark for correct answer given in the incorrect order of magnitude
			One mark for correct working but incorrect answer eg
			(0.34÷1000) x 3.2x10 <sup>9</sup> x 10 <sup>-3</sup>

Question	Marking guidance	Mark	Comments
02.3	1. DNA in mitochondria is circular <b>and</b> DNA in the nucleus is linear;	2	
	2. DNA in mitochondria is not associated with proteins		
	Or		
	DNA in the nucleus is associated with proteins/histones;		
			Allow DNA in mitochondria has no/fewer introns;

Question	Marking guidance	Mark	Comments
02.4	1. (DNA) helicase breaks hydrogen bonds (between DNA strands);	5	Accept H bonds for hydrogen bonds
	Or		Reject 'hydrolyses hydrogen bonds'
	(DNA) helicase separates the DNA strands (between 2 DNA strands);		
	2. Both strands act as templates		Allow description of both DNA strands being copied
	Or		and the second and th
	Each strand acts as a template;		
	3. (Free) nucleotides attach by complementary / specific base pairing / AT and GC;		
	4. DNA polymerase joins nucleotides (to the new DNA strand);		4. Reject if DNA polymerase catalyses complementary
	5. Reference to condensation reactions / formation of phosphodiester bonds (between nucleotides);		base pairing or if DNA polymerase catalyses nucleotides joining to template strand

Question	Marking guidance	Mark	Comments
03.1	(Number of chromosomes per cell) 2, 2;	2	Mark per column
	(Mass of DNA per cell) 400, 200;		

Question	Marking guidance	Mark	Comments
03.2	(Meiosis 1)	2	
	<ol> <li>Homologous chromosomes separate/are pulled/move apart (to opposite poles);</li> </ol>		Allow chromosomes of each pair for homologous chromosomes
	(Meiosis 2)		
	<ol><li>(Sister) chromatids separate/are pulled/move apart (to opposite poles);</li></ol>		

Question	Marking guidance	Mark	Comments
03.3	(Random) fertilisation/fusion of gametes;	1	Allow mutations
			Ignore random mating/interbreeding

Question	Marking guidance	Mark	Comments
04.1	1. The rate is the same up to 20–30 minutes/eq;	3 max	Accept initial rates are the same
	2. Faster uptake for <b>P</b> than <b>Q</b> ;		For full marks must have marking point 4 plus any other 2
	3. (Uptake of) <b>P</b> is linear/increases throughout/does not level off <b>and</b> uptake of <b>Q</b> levels off/stops;		marking points
	4. Correct manipulation of comparative figures eg <b>P</b> 9.5 μg cm <sup>-3</sup> higher than <b>Q</b> at 10 hours/max concentration eg <b>P</b> 14 μg cm <sup>-3</sup> and <b>Q</b> 4.5 μg cm <sup>-3</sup>		
	OR		
	Other relevant figures eg uptake of <b>Q</b> levels off in the range of 3.8 to 4.2 hours / at 4.5 µg cm <sup>-3</sup> ;		

Question	Marking guidance	Mark	Comments
04.2	Active transport is against a concentration gradient <b>and</b> diffusion is down a concentration gradient;	2 max	
	Active transport requires (energy in the form of) ATP (diffusion does not);		2. Ignore active transport requires energy unqualified
	Active transport requires membrane/carrier proteins (simple diffusion does not involve proteins);		

Question	Marking guidance	Mark	Comments
04.3	Substance P rate of uptake does not decrease/level off;	2	
	<ol> <li>Substance P rate of uptake is not affected by concentration gradient/does not stop when concentration inside cell equals that outside;</li> </ol>		

Question	Marking guidance	Mark	Comments
04.4	<ol> <li>Temperature affects the rate of enzyme reactions/metabolism Or 37°C is optimal temperature for enzymes Or (So) temperature is not a limiting factor/enough kinetic energy for enzymes to work efficiently Or Too high a temperature would denature enzymes/transport proteins;</li> <li>Enzymes are needed for respiration/to generate ATP Or Enzyme/ATPase needed to hydrolyse ATP (to release energy);</li> </ol>	2	Ignore idea that 37 °C is human body temperature  Ignore too high a temperature would kill the cells

Question	Marking guidance	Mark	Comments
05.1	32;;	2	One mark for correct answer but with incorrect number of significant figures (eg 32.0 / 32.3 / 32.26) or with incorrect rounding (eg 32.2)

Question	Marking guidance	Mark	Comments
05.2	Bacteria have a large surface area to volume ratio;	2	Accept converse
	Idea that bacteria can just use diffusion (through cell membranes to exchange gases)     Or     Idea of bacteria having a short diffusion distance;		

Question	Marking guidance	Mark	Comments
05.3	1. Many alveoli/capillaries <b>so</b> provide a large surface area;	3	Both feature and explanation needed for each mark point
	2. Thin epithelium/wall/surface of alveoli/capillaries <b>so</b> short <u>diffusion</u> distance/pathway;		2. Ignore reference to thin cell membrane
	<ol> <li>Constant ventilation/breathing/circulation so maintains concentration gradient;</li> </ol>		

Question	Marking guidance	Mark	Comments
06.1	<ol> <li>So that sequence/order of movement (of amino acids/polypeptide/proteins) through organelles can be tracked/recorded         Or         To identify organelles with (radioactively labelled) amino acids;</li> <li>(So) a comparison could be made between organelles/to control for differences in radioactivity take up between cells;</li> </ol>	2	Accept converse eg 'non-labelled amino acids cannot be tracked' etc

Question	Marking guidance	Mark	Comments
06.2	Translation;	1	

Question	Marking guidance	Mark	Comments
06.3	Named bonds form eg hydrogen/ionic/disulfide/hydrophobic;	3	Reject peptide bond
	2. Chaperone proteins assist in the folding (of polypeptides);		2. Allow forms alpha haliv or bata placted about for
	3. The polypeptide folds into the secondary/tertiary structure;		Allow forms alpha-helix or beta-pleated sheets for secondary structure;

Question	Marking guidance	Mark	Comments
06.4	Organelle <b>X</b> Rough endoplasmic reticulum/RER/ <u>transport</u> vesicle/ribosome;	1	Both need to be correct
	Organelle <b>Z</b> (Secretory) vesicle/lysosome;		

Question	Marking guidance	Mark	Comments
06.5	Vesicles containing radioactive amino acids/polypeptides fuse to create the Golgi	1	
	Or		
	(Radioactive) polypeptides/proteins enter the Golgi apparatus to be modified/packaged/transported;		

Question	Marking guidance	Mark	Comments
06.6	1.5;	1	Accept –1.5

Question	Marking guidance	Mark	Comments
06.7	(Secretory) vesicles (containing radioactive amino acids) fused with the cell membrane/carried out exocytosis/leave the cell;	2	
	(Radioactive) amino acids/polypeptides move into other organelles/form lysosomes;		Accept decreased due to radioactive decay

Question	Marking guidance	Mark	Comments
07.1	30.4 / 30 (range 29–31);;	2	Allow for one mark for:
			<u>(6.9 – 4.8);</u> 6.9
			Or Or
			0.304;
			Allow one mark for correct answer from use of 6.8 or 7.0 instead of 6.9

Question	Marking guidance	Mark	Comments
07.2	H+ increases / products are acidic / (fatty) acids produced;	1	Accept fatty acid/FFAs concentration increases

Question	Marking guidance	Mark	Comments
07.3	pH probe/meter/sensor;	1	Ignore litmus paper/universal indicator

Question	Marking guidance	Mark	Comments
07.4	One from the following:  1. Temperature;	1	Ignore amount/volume of milk Ignore concentration of enzymes (in the milk)
	2. Type/source of milk Or Concentration of milk/concentration of triglycerides/fat in the milk Or Initial pH of milk Or Age of the milk;		

Question	Marking guidance	Mark	Comments
07.5	Substrate/triglycerides used up/equilibrium reached;	2	1. Ignore milk used up
	2. (pH too low so) enzyme denatured;		

Question	Marking guidance	Mark	Comments
07.6	(In pasteurised) lower concentration of enzymes/lipase     Or	2	Accept converse
	Enzymes/lipase are denatured;		
	(So) fewer successful collisions     Or     Fewer enzyme-substrate complexes formed;		
	OR		
	3. Heating kills (most of) the bacteria;		
	<ol> <li>(So in pasteurised) less/no lipase (is released into the milk)</li> <li>Or</li> </ol>		
	Only the lipase naturally in the milk is present (so lower concentration);		

Question	Marking guidance	Mark	Comments
07.7	Reference to using at least 5 enzyme concentrations;	5 max	
	Description of how to make enzyme concentration eg serial dilutions etc;		
	Equilibrate/eq enzyme and substrate/triglycerides separately at correct temperature;		
	4. Add enzyme/lipase to the substrate/triglycerides;		
	5. Measure the pH at intervals;		
	Idea of repeating experiment without enzyme/with denatured enzyme;		Do not award if lipase concentration of 0 is stated as one of the enzyme concentrations
	7. Idea of controlling triglyceride concentration/concentration of milk/temperature/volume of solutions;		7. Only accept references to controlling pH if clear that this isn't the dependent variable
	8. Repeat for each enzyme concentration and calculate a mean;		7. Reject if reference to control of pH / use of buffer solution

Question	Marking guidance	Mark	Comments
08.1	1. Same genus;	2	
	2. Share a <b>recent</b> common ancestor;		

Question	Marking guidance	Mark	Comments
08.2	(Comparisons of:)	3	
	1. The (base) sequence of DNA or of mRNA;		
	2. The frequency of specific base sequences or alleles;		
	3. The amino acid sequence of a particular protein eg haemoglobin;		Allow the following or correct descriptions of:
			4. Immunological comparison
			5. DNA hybridisation
			6. Genetic fingerprinting

Question	Marking guidance	Mark	Comments
08.3	Any <b>two</b> from:	2 max	
	Described difficulty of measuring penguins in the wild eg difficult to catch or possible damage caused whilst taking measurements / less accurate measurements from living specimens;		Ignore easier to take measurements unqualified
	Difficulty of reaching/locating sampling areas (in extreme conditions);		2. Allow reference to more expensive to sample penguins in the wild
	3. Described problems associated with scientist disturbing nesting sites, breeding etc;		

Question	Marking guidance	Mark	Comments
08.4	Idea of:	1	
	All four species would more likely be protected (as the population sizes would be smaller) rather that treating them as one large population		
	Or		
	Specific conservation plans could be put into place for each population		
	Or		
	Some populations may be more at risk than others so can monitor certain populations more carefully;		

Question	Marking guidance	Mark	Comments
08.5	Breed individuals from one colony with another;	2	
	2. Failure to produce fertile offspring indicates they are different species;		