

GCE

Biology B

H422/03: Practical skills in biology

A Level

Mark Scheme for June 2022

OCR (Oxford Cambridge and RSA) is a leading UK awarding body, providing a wide range of qualifications to meet the needs of candidates of all ages and abilities. OCR qualifications include AS/A Levels, Diplomas, GCSEs, Cambridge Nationals, Cambridge Technicals, Functional Skills, Key Skills, Entry Level qualifications, NVQs and vocational qualifications in areas such as IT, business, languages, teaching/training, administration and secretarial skills.

It is also responsible for developing new specifications to meet national requirements and the needs of students and teachers. OCR is a not-for-profit organisation; any surplus made is invested back into the establishment to help towards the development of qualifications and support, which keep pace with the changing needs of today's society.

This mark scheme is published as an aid to teachers and students, to indicate the requirements of the examination. It shows the basis on which marks were awarded by examiners. It does not indicate the details of the discussions which took place at an examiners' meeting before marking commenced.

All examiners are instructed that alternative correct answers and unexpected approaches in candidates' scripts must be given marks that fairly reflect the relevant knowledge and skills demonstrated.

Mark schemes should be read in conjunction with the published question papers and the report on the examination.

© OCR 2022

June 2022

MARKING INSTRUCTIONS

PREPARATION FOR MARKING

RM ASSESSOR

- 1. Make sure that you have accessed and completed the relevant training packages for on-screen marking: *RM Assessor Online Training*; *OCR Essential Guide to Marking*.
- 2. Make sure that you have read and understood the mark scheme and the question paper for this unit. These are available in RM Assessor.
- 3. Log-in to RM Assessor and mark the **required number** of practice responses ("scripts") and the **required number** of standardisation responses.

MARKING

- 1. Mark strictly to the mark scheme.
- 2. Marks awarded must relate directly to the marking criteria.
- 3. The schedule of dates is very important. It is essential that you meet the RM Assessor 50% and 100% (traditional 50% Batch 1 and 100% Batch 2) deadlines. If you experience problems, you must contact your Team Leader (Supervisor) without delay.
- 4. If you are in any doubt about applying the mark scheme, consult your Team Leader by telephone, email or via the RM Assessor messaging system.

5. Work crossed out:

Where a candidate has crossed out a response and provided a clear alternative then the crossed-out response is not marked. Where no alternative response has been provided, examiners may give candidates the benefit of the doubt and mark the crossed-out response where legible.

Rubric Error Responses – Optional Questions

Where candidates have a choice of question across a whole paper or a whole section and have provided more answers than required, then all responses are marked and the highest mark allowable within the rubric is given. Enter a mark for each question answered into RM assessor, which will select the highest mark from those awarded. (The underlying assumption is that the candidate has penalised themselves by attempting more questions than necessary in the time allowed.)

Multiple Choice Question Responses

When a multiple choice question has only a single, correct response and a candidate provides two responses (even if one of these responses is correct), then no mark should be awarded (as it is not possible to determine which was the first response selected by the candidate).

When a question requires candidates to select more than one option/multiple options, then local marking arrangements need to ensure consistency of approach.

Contradictory Responses

When a candidate provides contradictory responses, then no mark should be awarded, even if one of the answers is correct.

Short Answer Questions (requiring only a list by way of a response, usually worth only one mark per response) Where candidates are required to provide a set number of short answer responses then only the set number of responses should be marked. The response space should be marked from left to right on each line and then line by line until the required number of responses have been considered. The remaining responses should not then be marked. Examiners will have to apply judgement as to whether a 'second response' on a line is a development of the 'first response', rather than a separate, discrete response. (The underlying assumption is that the candidate is attempting to hedge their bets and therefore getting undue benefit rather than engaging with the question and giving the most relevant/correct responses.)

Short Answer Questions (requiring a more developed response, worth two or more marks)

If the candidates are required to provide a description of, say, three items or factors and four items or factors are provided, then mark on a similar basis – that is downwards (as it is unlikely in this situation that a candidate will provide more than one response in each section of the response space.)

PMT

June 2022

Longer Answer Questions (requiring a developed response)

Where candidates have provided two (or more) responses to a medium or high tariff question which only required a single (developed) response and not crossed out the first response, then only the first response should be marked. Examiners will need to apply professional judgement as to whether the second (or a subsequent) response is a 'new start' or simply a poorly expressed continuation of the first response.

- 6. Always check the pages (and additional objects if present) at the end of the response in case any answers have been continued there. If the candidate has continued an answer there then add a tick to confirm that the work has been seen.
- 7. There is a NR (No Response) option. Award NR (No Response)
 - if there is nothing written at all in the answer space
 - OR if there is a comment which does not in any way relate to the question (e.g. 'can't do', 'don't know')
 - OR if there is a mark (e.g. a dash, a question mark) which isn't an attempt at the question.

Note: Award 0 marks – for an attempt that earns no credit (including copying out the question).

8. The RM Assessor **comments box** is used by your Team Leader to explain the marking of the practice responses. Please refer to these comments when checking your practice responses. **Do not use the comments box for any other reason.**

If you have any questions or comments for your Team Leader, use the phone, the RM Assessor messaging system, or email.

9. Assistant Examiners will send a brief report on the performance of candidates to their Team Leader (Supervisor) via email by the end of the marking period. The report should contain notes on particular strengths displayed as well as common errors or weaknesses. Constructive criticism of the question paper/mark scheme is also appreciated.

10. For answers marked by levels of response:

Read through the whole answer from start to finish, using the Level descriptors to help you decide whether it is a strong or weak answer. The indicative scientific content in the Guidance column indicates the expected parameters for candidates' answers, but be prepared to recognise and credit unexpected approaches where they show relevance. Using a 'best-fit' approach based on the skills and science content evidenced within the answer, first decide which set of level descriptors, Level 1, Level 2 or Level 3, best describes the overall quality of the answer.

Once the level is located, award the higher or lower mark:

The higher mark should be awarded where the level descriptor has been evidenced and all aspects of the communication statement (in italics) have been met.

The lower mark should be awarded where the level descriptor has been evidenced but aspects of the communication statement (in italics) are missing.

In summary:

The skills and science content determines the level.

The communication statement determines the mark within a level.

Level of response questions on this paper are 2(c) and 4(b)(ii).

11. Annotations available in RM Assessor

Marking Annotations

Annotation	Use				
BOD	Benefit of Doubt				
CON	Contradiction				
×	Cross				
ECF	Error Carried Forward				
GM	Given Mark				
~~~·	Extendable horizontal wavy line (to indicate errors / incorrect science terminology)				
I	Ignore				
•	Large dot (various uses as defined in mark scheme)				
	Highlight (various uses as defined in mark scheme)				
NBOD	Benefit of the doubt not given				
<b>~</b>	Tick				
<b>^</b>	Omission Mark				
BP	Blank Page				
Ц	Level 1 answer in Level of Response question				
L2	Level 2 answer in Level of Response question				
L3	Level 3 answer in Level of Response question				

12. Abbreviations, annotations and conventions used in the detailed Mark Scheme (to include abbreviations and subject-specific conventions).

Annotation	Meaning
I	alternative and acceptable answers for the same marking point
✓	Separates marking points
DO NOT ALLOW	Answers which are not worthy of credit
IGNORE	Statements which are irrelevant
ALLOW	Answers that can be accepted
()	Words which are not essential to gain credit
	Underlined words must be present in answer to score a mark
ECF	Error carried forward
AW	Alternative wording
ORA	Or reverse argument

## 13. Subject-specific Marking Instructions

### INTRODUCTION

Your first task as an Examiner is to become thoroughly familiar with the material on which the examination depends. This material includes:

- the specification, especially the assessment objectives
- the question paper
- the mark scheme.

You should ensure that you have copies of these materials.

You should ensure also that you are familiar with the administrative procedures related to the marking process. These are set out in the OCR booklet **Instructions for Examiners**. If you are examining for the first time, please read carefully **Appendix 5 Introduction to Script Marking: Notes for New Examiners**.

Please ask for help or guidance whenever you need it. Your first point of contact is your Team Leader.

June 2	022
--------	-----

C	Questi	on	Answer	Mark	AO	Guidance
1	(a)	(i)	secretes (digestive) enzymes ✓	1	2.3	ALLOW named example of digestive enzyme
1	(a)	(ii)	alpha / α <b>AND</b> beta / β ✓	1	2.3	IGNORE endocrine IGNORE islets of Langerhans IGNORE 'a' and 'b' cells
1	(a)	(iiii)	FIRST CHECK ON ANSWER LINE If answer = 55.6 award 2 marks 20mm x1000 = 20 000μm ✓ 20 000 ÷ 360 = 55.6 (μm) ✓	2	2.4	<ul> <li>ALLOW measurement of 20mm +/- 0.5mm i.e. 54.2 (using 19.5mm) or 56.9 (using 20.5mm)</li> <li>Apply ECF e.g. 19.0 → 19000 / 360 = 52.8 would gain one mark (for ECF 2nd MP for working)</li> <li>ALLOW any correct rounding for this working mark e.g. 55.56, 55.556 but only 1 mark maximum if answer less or more than 3 sig figs.</li> <li>ALLOW 2cm / 360 = 0.005555cm (as alternative for 2nd working mark)</li> </ul>

) (iv)	<ul> <li><i>correct reference to</i> calibration of the eye piece graticule using the stage micrometer ✓</li> <li>replace stage micrometer with specimen slide AND use, same / specified/ AW, magnification ✓</li> <li><u>count the number of graticule divisions that cover the (linear) dimension of the cells AND</u></li> </ul>	max 2	1.2	<b>ALLOW</b> 'eyepiece units' as alternative wording for 'graticule divisions'
	using the stage micrometer ✓ replace stage micrometer with specimen slide AND use, same / specified/ AW, magnification ✓ count the number of graticule divisions that cover the (linear) dimension of the cells			'graticule divisions'
	AND use, same / specified/ AW, magnification ✓ count the number of graticule divisions that cover the (linear) dimension of the cells			
	(linear) dimension of the cells			
	multiply number of graticule divisions by known length of (stage) micrometer / AW ✓			
	AVP ✓			e.g. micrometer has (scale with) divisions of , known length / 1mm / 0.1mm / 100µm
)	glucose tolerance test, indicates the person has impaired glucose tolerance / AW AND fasting test, indicates the person (probably) has diabetes / AW ✓	max 2	3.1	
	<i>idea that</i> the person's glucose tolerance may have worsened (over the two months) ✓			
	<i>dea that</i> different test conclusions could have been caused by fault in the person's preparation for the tests ✓			e.g. the person might not have fasted for the correct amount of time
		<ul> <li>AND fasting test, indicates the person (probably) has diabetes / AW ✓</li> <li><i>idea that</i> the person's glucose tolerance may have worsened (over the two months)</li> <li>✓</li> <li><i>dea that</i> different test conclusions could have been caused by fault in the person's preparation for the tests</li> </ul>	AND         fasting test, indicates the person (probably) has         diabetes / AW ✓         idea that the person's glucose tolerance may have         worsened (over the two months)         ✓         dea that different test conclusions could have been         caused         by fault in the person's preparation for the tests	AND         fasting test, indicates the person (probably) has         diabetes / AW ✓         idea that the person's glucose tolerance may have         worsened (over the two months)         ✓         dea that different test conclusions could have been         caused         by fault in the person's preparation for the tests

H42	2/03			Ma	rk So	cheme		June 2022
1	(c)	H CH2OH H OH H OH H OH H OH OH	н	✓		1	1.1	<b>DO NOT ALLOW</b> lower bonds joining to H rather than O in the OH groups.
1	(d)	carbohydrate molecule	reagent used in tests for identification	number of glycosidic bonds per molecule		2	1.1 1.2	Award one mark per correct column
		glucose	Benedict's (test)	0 / none				
		lactose	Benedict's (test)	1 / one				
		amylose	iodine <u>solution</u> / potassium iodide <u>solution</u>	many				IGNORE "iodine K-I"
			$\checkmark$	√				

12

PMT

June	2022
------	------

2	(a)		agree with / supports the conclusion heroin is more harmful in 3 of the 4 categories ✓ disagree with / undermines the conclusion alcohol has a greater economic cost ✓ general points correct use of the combined harm scores to justify the data, supporting / undermining, the conclusion ✓ other (named) categories of harm have not been considered (in the graph) ✓ AVP ✓	max 4	3.2	Allow a maximum of one mark from MP1 and MP2 if no reference to the data supporting or undermining the conclusion e.g. the total harm score for heroin is higher than for alcohol, so supports the conclusion <b>OR</b> the total harm scores are (approximately) the same, so undermines the conclusion e.g. effects on relationships and mental / psychological effects are not shown e.g. many more people (may) use alcohol than heroin (due to availability) idea that one drug may be more harmful than the other to, <u>society</u> / the individual idea that subjective as to which categories are most important no statistical analysis of the data has been carried out / no error bars plotted (to show variation in data )
2	(b)	(i)	intrinsic / integral (protein) ✓	1	2.1	ALLOW <u>ligand-gated</u> ion (channel protein) IGNORE 'channel protein' as given in figure 2.2

June	2022
------	------

	(1)	()				
2	(b)	(ii)		3	2.5	ALLOW the use of the term 'neurotransmitter' in place of
					3.1	'GABA' (provided no other neurotransmitter is referred
						to)
			comparison			
			heroin, stops / reduces / AW, release of GABA / AW, from			
			presynaptic neurone / into synaptic cleft			
			AND			
			alcohol, mimics / AW, GABA, so binds to GABA receptors			
			$\checkmark$			
			explanation			
			heroin, prevents / reduces, GABA from reaching / AW, the			
			postsynaptic neurone			
			AND			
			prevents / reduces, the generation / AW, of an action			
			potential (in the inhibitory postsynaptic neurone) $\checkmark$			
			alcohol (binds to GABA receptor so) stimulates, the			
			generation / AW, of an action potential (in the inhibitory			
			postsynaptic neurone) ✓			
			(alcohol causes channel to remain open for longer			
			causing) more CI ions to enter, causing hyperpolarisation			ALLOW idea of 'more negative / more polarised, as a
			AND			result of channels remaining open for longer' in place of
			reduces chance of / inhibits, the generation of an action			'hyperpolarisation'
			potential 🗸			
		I	1			

2 (c)*	Please refer to the marking instructions on page 4 of thi	s mark s	scheme	for guidance on how to mark this question.
	<ul> <li>In summary: Read through the whole answer. (Be prepared to recognise Using a 'best-fit' approach based on the science content of t Level 3, best describes the overall quality of the answer. Then, award the higher or lower mark within the level, accor o award the higher mark where the Communication Stat o award the lower mark where aspects of the Communic • The science content determines the level.</li> <li>• The Communication Statement determines the mark w</li> </ul>	the answe ding to th tement ha cation Sta	er, first c ne <b>Comı</b> as been atement	decide which of the level descriptors, <b>Level 1</b> , <b>Level 2</b> or <b>munication Statement</b> (shown in italics): met.
	<ul> <li>Level 3 (5-6 marks)         Comprehensive experimental and safe method producing valid data and detailed statistical analysis for the investigation.     </li> <li>There is a well-developed line of reasoning which is clear and logically structured. The information presented is relevant and substantiated.</li> <li>Level 2 (3-4 marks)         Detailed experimental method and simple statistical analysis for the investigation.     </li> <li>There is a line of reasoning with some structure. The information presented is relevant and supported by some evidence.     </li> <li>Level 1 (1-2 marks)         Simple detail of experimental method assessment and some numerical processing of data     </li> </ul>	6	3.3 3.4	<ul> <li>Indicative scientific points may include (but are not limited to):</li> <li><i>Experimental method</i> <ul> <li>Details of sample sizes (repeated with at least 10 subjects)</li> <li>Details of the independent variable (e.g. 'caffeine vs no caffeine consumed', or 'volume of caffeinated drink consumed', or 'concentration of caffeine consumed' (if the original drink is diluted for different groups, for example))</li> <li>Details of group design (e.g. two groups: one consuming caffeine, or several groups receiving different caffeine concentrations, plus a control group)</li> <li>Details of standardisation of method (i.e. same procedure every time)</li> <li>Details of the measurement of the dependent variable (e.g. distance to reaction time conversion table or online software)</li> </ul> </li> </ul>
	The information is basic and communicated in an unstructured way. The information is supported by limited evidence and the relationship to the evidence may not be clear.			<ul> <li>Health &amp; safety</li> <li>Gain consent from participants</li> <li>Risk assessment: allergies, heart conditions etc</li> </ul>

<b>0 marks</b> No response or no response worthy of credit.	<ul> <li>Consideration of screen exposure, flicker rate etc (for those methods using online reaction time method)</li> <li>Possible contraindications with caffeine intake</li> </ul>
	<ul> <li>Statistical analysis</li> <li>Idea of identifying and excluding (or replacing) anomalies</li> <li>Mean calculations</li> <li>Standard deviation</li> <li>Appropriate statistical testing (e.g. unpaired t-test if two separate groups are used, paired t-test if same subjects used and tested before/after intake of caffeine, or Spearman's Rank Correlation Coefficient test if a range of caffeine concentrations are used)</li> </ul>
	<ul> <li>Validity <ul> <li>Minimum of 10 in each group for t-test</li> <li>Use of dominant hand to catch ruler Stated time period between ingestion of caffeine and testing (to allow for absorption and effect to take place)</li> <li>Details of control variables (e.g. age of participants, biological sex, typical caffeine intake/ diet, volume of caffeine consumed) or unbiased assignment to groups</li> </ul> </li> </ul>

3	(a)	(i)		2	2.5	ALLOW measurements to be +/- 0.5mm
			FIRST CHECK ON ANSWER LINE If answer = 0.59 award 2 marks			ALLOW 1 mark for alternative sig figs e.g. 0.6, 0.588
			47 (mm) / 80 (mm) = 0.5875 ✓			
			$R_{f}$ (to 2 significant figures) = 0.59 $\checkmark$			<b>Apply ECF</b> for MP2 for incorrect measurements e.g. 48mm / 80mm = 0.60
3	(a)	(ii)	pigments have, different, solubilities in / affinity for, the (new / different) solvent / mobile phase OR (new) solvent has, different, affinity for stationary phase ✓	1	2.7	
3	(a)	(iii)	ninhydrin	1	1.2	
3	(b)	(i)	centrifuge (sample) <b>AND</b> <i>idea of</i> samples spun at (high) speed (in tubes)	max 2	1.2 2.7	
			separation by density / AW ✓			e.g. pellets are denser so they settle at the bottom
			decanting supernatant to leave sediment which is the 'pellet' ✓			
3	(b)	(ii)	reduce / AW, <u>enzyme</u> activity (in chloroplasts) ✓	1	2.7	<b>IGNORE</b> references to 'enzymes being denatured' or 'enzyme activity prevented'
3	(b)	(iii)	statement red (filter) ✓ explanation (as it) absorbs green wavelengths OR	2	2.7	ALLOW orange
			idea that the decrease in the, intensity / AW, of the			

	blue colour of the DCPIP is observed		
	$\checkmark$		

3	(b)	(iv)	cuvette ✓ reduced ✓ 1 / one <b>AND</b> 3 / three ✓			3	1.2 2.7	ACCEPT in either order
3	(b)	(v)	<ul> <li>(A = tube) 4 AND (B = tube) 2 ✓</li> <li>any two from:</li> <li><i>idea that</i> DCPIP is decolorised, when it is reduced / when it accepts electrons (from light-dependent reactions) ✓</li> <li>(tube) B / 2, has a high<u>er</u>, concentration / number, of, chloroplasts / photosynthetic pigments ora</li> <li>✓</li> <li>(rate of) light absorption / light-dependent reactions, is great<u>er</u> / fast<u>er</u> / AW , in , (tube) B / 2 ora</li> </ul>				3.1	ACCEPT 'supernatant' for '4' and 'pellet in <u>light'</u> for '2' ACCEPT 'ETC' or 'photosystem I' for 'light-dependent reactions'
4	(a)		Feature         peptidoglycan cell         wall	Gram- positive bacteria ✓	Gram- negative bacteria √	3	1.1 1.2	2 correct rows = 1 mark 3 correct rows = 2 marks 4 correct rows = 3 marks IGNORE crosses in empty cells

4 (b)

		Mark S	cheme		June 2022
lipopolysacchario outer envelope		✓ 			
plasma membrar	ne 🗸	$\checkmark$			
is stained with crys violet during the Gram staining procedure		✓			
has a final colour pink after Gram staining		~			
$\checkmark\checkmark\checkmark$					
(i) Any two from			max 2	2.8	
	extrapolation of control group data points to the y axis at the first solid line above $10^2$ which equates to 200 $\checkmark$				ALLOW ecf for incorrect use of log scale but evidence
x 10 (takes into acc	x 10 (takes into account the final volume) = 2000 $\checkmark$				of dilution factor being taken into account
2.00 x 10 ³ ✓	2.00 x 10 ³ ✓				

4	(b)	(ii)*	Please refer to the marking instructions on page 4 of this mark scheme for guidance on how to mark this question.
			<i>In summary:</i> Read through the whole answer. (Be prepared to recognise and credit unexpected approaches where they show relevance.) Using a 'best-fit' approach based on the science content of the answer, first decide which of the level descriptors, <b>Level 1</b> , <b>Level 2</b> or <b>Level 3</b> , best describes the overall quality of the answer. Then, award the higher or lower mark within the level, according to the <b>Communication Statement</b> (shown in italics): • award the higher mark where the Communication Statement has been met. • award the lower mark where aspects of the Communication Statement have been missed.

June 2	2022
--------	------

Level 3 (5-6 marks)Description and detailed explanation for all 3 groups.There is a well-developed line of reasoning which is clear and logically structured. The information presented is	6	2.7 3.3 3.4	Indicative scientific points may include (but are not limited to): Description Inear increase for all three groups between 2 and
Image: and logically structured. The information presented is relevant and substantiated.         Level 2 (3-4 marks)         Description of all 3 groups and brief explanation for at least 2 groups         OR         Description for at least 1 group and detailed explanation for at least one group.         There is a line of reasoning with some structure. The information presented is relevant and supported by some evidence.			<ul> <li>Intear increase for all three groups between 2 and 4 hours</li> <li>linear increase/ exponential growth, for control group between 2 and 8 hours</li> <li>plateau, after 8 hours/ between 8 to 14h, for control group</li> <li>plateau, after 4 hours/ between 4 to 14h, for P</li> <li>peak at 4h for Q</li> <li>(gradual/inconsistent) decrease, after 4 hours / between 4 to 14h, for Q</li> <li>antibiotic Q is most effective, as after 14h, there are the least number of (bacterial) cells</li> </ul>
<b>Level 1 (1-2 marks)</b> Description for at least 1 group and an attempt at explanation for at least one group.The information is basic and communicated in an unstructured way. The information is supported by limited evidence and the relationship to the evidence may not be clear.			<ul> <li>Explanation</li> <li>Control group: <ul> <li>exponential growth before carrying capacity is reached</li> <li>(stationary phase occurs) when waste products have accumulated</li> <li>nutrients are (becoming) limited (in stationary phase)</li> </ul> </li> </ul>
<b>0 marks</b> No response or no response worthy of credit.			<ul> <li>Antibiotic P:</li> <li>the antibiotic is bacteriostatic</li> <li>replication is prevented (but bacteria are not killed)</li> <li>binary fission is prevented</li> <li>named reason e.g. protein synthesis is stopped, tRNA is prevented from binding to ribosomes</li> </ul>

H42	2/03		Mark S	cheme		June 2022
						<ul> <li>Antibiotic Q:</li> <li>the antibiotic is bactericidal</li> <li>bacteria are killed</li> <li>named reason e.g. peptidoglycan synthesis is stopped, plasma membrane is damaged, plasma membrane ruptures</li> </ul>
4	(c)	(i)	<i>idea that</i> <u>only</u> , glycolysis / substrate level phosphorylation, occurs ✓	1	2.1	ACCEPT no Krebs cycle / oxidative phosphorylation, occurs ACCEPT substrate-linked phosphorylation
4	(c)	(ii)	plasma / cell <u>surface</u> , membrane ✓	1	2.1	ACCEPT on mesosome

PMT

June	2022
------	------

E	(a)	(:)		mey 2	2.2	Only award mark if abanga and avalanction are bath
Э	(a)	(i)		max 3	2.3	Only award mark if change and explanation are both
					2.7	given. Correct explanation must be paired with the
			,			change matched.
			change:			
			increase temperature (by using a water bath) / stir / mix			ACCEPT heat in a water bath
			AND			
			explanation:			
			(water) molecules have more kinetic energy ✓			ACCEPT '(water) molecules gain energy and move
						faster'
			change:			
			increase sucrose concentration			
			AND			
			explanation:			
			<i>idea of</i> osmotic / water potential, gradient increases $\checkmark$			
			change:			
			increase, number of foldings / (surface) area, of, bladder /			
			semipermeable membrane			
			AND			
						ALLOW 'size' in place of 'surface area'
			explanation:			
			increased, surface area (for osmosis / diffusion) $\checkmark$			
			<i>change</i> : reduce thickness of bladder / use a (named)			
			membrane with reduced thickness			
			AND			
			<i>explanation</i> : reduced diffusion, pathway / distance $\checkmark$			
			abangai			
			change:			
			AVP			
			AND			e.g. increase height of thistle funnel <b>AND</b> reduces
			explanation:			pressure acting on goat bladder membrane
			allow AVP ✓			

# Mark Scheme

5	(a)	(ii)	mm / cm ✓	1	2.4	<b>IGNORE</b> conversion of heights to volumes as this is the processed data not the DV (raw data) e.g. mm min ⁻¹ / cm min ⁻¹ / ml min ⁻¹ / mm ³ min ⁻¹ / cm ³ min ⁻¹
5	(a)	(iii)	<u>Visking</u> tubing / <u>dialysis</u> membrane ✓	1	2.3	
5	(b)	(i)	proximal convoluted tubule ✓	1	1.1	ACCEPT PCT
5	(b)	(ii)	collecting duct ✓	1	1.1	ACCEPT distal convoluted tubule / DCT

#### Need to get in touch?

If you ever have any questions about OCR qualifications or services (including administration, logistics and teaching) please feel free to get in touch with our customer support centre.

Call us on

#### 01223 553998

Alternatively, you can email us on

support@ocr.org.uk

For more information visit



ocr.org.uk

Twitter/ocrexams

/ocrexams

/company/ocr

/ocrexams



OCR is part of Cambridge University Press & Assessment, a department of the University of Cambridge.

For staff training purposes and as part of our quality assurance programme your call may be recorded or monitored. © OCR 2022 Oxford Cambridge and RSA Examinations is a Company Limited by Guarantee. Registered in England. Registered office The Triangle Building, Shaftesbury Road, Cambridge, CB2 8EA.

Registered company number 3484466. OCR is an exempt charity.

OCR operates academic and vocational qualifications regulated by Ofqual, Qualifications Wales and CCEA as listed in their qualifications registers including A Levels, GCSEs, Cambridge Technicals and Cambridge Nationals.

OCR provides resources to help you deliver our qualifications. These resources do not represent any particular teaching method we expect you to use. We update our resources regularly and aim to make sure content is accurate but please check the OCR website so that you have the most up-to-date version. OCR cannot be held responsible for any errors or omissions in these resources.

Though we make every effort to check our resources, there may be contradictions between published support and the specification, so it is important that you always use information in the latest specification. We indicate any specification changes within the document itself, change the version number and provide a summary of the changes. If you do notice a discrepancy between the specification and a resource, please <u>contact us</u>.

Whether you already offer OCR qualifications, are new to OCR or are thinking about switching, you can request more information using our Expression of Interest form.

Please get in touch if you want to discuss the accessibility of resources we offer to support you in delivering our qualifications.