



Applied Science

Advanced GCE G623

Cells and Molecules

Mark Scheme for June 2010

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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.

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Planning Exercise

Investigate the effects of incubation temperature on the yield of juice after enzyme treatment from one variety of English apple.

Marking of the plan:

- 1 Read the material presented.
- 2 Then *award 1 mark* if *scientific terminology* has been used appropriately. Record using the letter Y.
- 3 Then re-read, this time point marking up to 24, by placing letters A to X in the margin where you see evidence of the marking criteria.
- 4 The same piece of evidence can be used to award one criterion only.

Marking Point	Marking Criteria	Mark	Additional notes	
A	easily recognised safety procedures highlighted; At least 3 from: glassware; enzyme allergy; sharps (knives); electrical (water baths/blenders); burns (boiling water)	1	Evidence of something that is going to make doing the investigation safer – an active document, a working document related to the plan: ref to allergic reactions & enzymes	
В	prediction made; R = ref to 'body temperature' R = ref to tinned apples	1	A statement related to effect of temperature on juice yield.	
С	with justification; Accept ref to enzyme activity & temp if linked to research or insert; R ref to 55°C as optimum.	1	Statement related to enzyme activity: inactivity/denaturation of enzyme; possible link to molecular structure of protein; molecular movement; lock & key model; kinetic energy;	
D	description of preliminary work;	1	e.g. how to prepare tissue (do not credit twice) / mass of tissue to be used / dilution of enzyme(s)/ volume of enzyme/ type of enzyme/ source of enzyme/ range of temperatures to use / incubation time/ source of tissue /age of tissue:	
E	clear and in detail;	1	Explain how to do it.	
F	reason (for doing it) explained;	1	Explain why it's necessary for completion of the whole investigation.	
G	clear and in detail;	1	Extra information.	
н	at least two secondary sources of information identified;	Source & preparation of ti pulp; treatment pectinase; conti	State at least 2 references (allow OCR t as one source). website address needed. description of named text	
I	relevance explained;	emperature; ncubation; time -iltration:	explanation as to how ences helped in the planning.	
J	basic practical skills and accuracy;	clarification; Veasurement o	ble method / list of instructions. c. 'Is it a feasible approach?'	
к	sound practical skills and accuracy;	1	unaided? Are quantities shown? Is it repeatable to appropriate degree of accuracy?	

Marking Point	Marking Criteria	Mark	Additional notes
L	range of appropriate equipment listed; R = tinned apples;	1	List of names of main items of equipment and materials needed for the investigation. Generic terms: beakers, flasks etc are OK here.
Μ	full range of appropriate equipment listed; List <u>must</u> include apples	1	Qualifications noted. Indication of number of each, specific sizes, e.g. 250 cm ³ beaker, 1dm ³ flask. If any major item missing do not award (i.e. apples; pectinase; method of temperature control)
Ν	appropriate number of measurements stated;	1	Mentions at least 2 replicates / repeats
0	need for range of measurements stated;	1	Statement: e.g. 'To enable comparison to be made (to identify optimum temperature which gives maximum yield of juice)
Р	appropriate range stated;	1	Related to prediction made – accept at least 4 different temperatures (5 ideal)
Q	relevant variables are identified (stated); Need to identify controlled variables;	1	At least 2 from: Control variables age of tissue / mass of tissue / source of fruit / variety of fruit / source of extract / volume of extract / dilution of extract / cold stored or not / time in cold storage/ incubation time/ type of enzyme/ concentration of enzyme/; Independent/dependent: incubation temperature/ volume of juice;
R	how variables to be controlled explained;	1	How for at least 2 of the variables.
S	one suitable method to display data;	1	One display of results e.g. Table. (Clear headings & units in headers)
т	additional method to display data;	1	Any <u>different</u> display e.g. graph (with appropriate units).
U	simple data handling;	1	mean / colour comparison / use of graph data
V	possible conclusions; Accept use of graph & reading off (i.e. 'optimum temperature of max yield')	erature ations of water residue of n glassware; ment % error	Statements of expectations or bservations to confirm or reject rediction made in B . What would your results need to show to onfirm or reject your prediction?'
W	recognises sources of erfor	1	At least two examples: equipment / materials / specific human error (limit to 1 human error)
x	suggests methods for improving accuracy and or validity;	A = Repeat using narrov temp range around optimum;	 uracy: relate to 'W' or use of rnative technique(s). D / OR R = ref to body temp for improvement dity: state aspect of collected data to compared with secondary sources.
Marks	Maximum for plan = 25	using increat enzyme cor	ased ific terminology)

Qu	Question		Expected Answers	Marks	Additional Guidance
1	а		advantages: magnifies objects (over 500 000 times) / higher magnification / can see cell ultra-structure ✓ has a higher, resolving power/resolution / possible to investigate greater depth of field ✓	2	accept 'shows up more organelles than a light microscope'
			disadvantages - any two from: cost ✓ special accommodation ✓ needs skilled operative / difficult to operate ✓ preparation of specimens lengthy/complex /tissue sample thin/ ref to dehydration process/ complex staining ✓ material may be distorted / produces artefacts or distorts image ✓ high vacuum required ✓ living material cannot be viewed / ORA ✓	2	ignore ref to B & W images or does not show true colour image;
	b	i	65mm (accept range 64 to 66) √	1	allow correct measurement of maximum diameter in mm
		ii	correct conversion to $\mu m \checkmark$ division by magnification value \checkmark	2	accept answer range $9.4 - 9.7 (\mu m)$ award two marks for correct answer allow ecf from b(i) ; Limit to 1 mark if calculation method correct but answer incorrect; $64 = 9.4$; $65 = 9.6$; $67 = 9.7 (\mu m)$
	С	i	X = Golgi (body/apparatus) ✓ Y = mitochondrion / matrix ✓	2	ignore crista
		ii	 X = makes secretory vesicles / lysosome formation / produces, glycoproteins/mucin, / transports lipids / stores lipids / modifies glycolipids ✓ Y = aerobic respiration / production of ATP/ Krebs Cycle / TCA cycle / oxidative phosphorylation / link reaction ✓ 	2	accept 'receives proteins and modifies them'. for Y accept reference to electron transport chain
			Total	11	

Que	esti	on	Expected Answers	Marks	Additional Guidance
2	а		<pre>starch: iodine/iodine KI, solution ✓ protein: Biuret (reagent) / sodium hydroxide (solution) and dilute copper sulfate (solution) ✓ purple/lilac ✓</pre>	а	1 mark for each correct box, test reagent (s) must be complete for the mark. Accept ref to 'iodine in potassium iodine' owtte
			lipid/fat: ethanol and water ✓		reject ethanol test/ emulsion test / ethanol;
			dilute HCl and sodium bicarbonate and Benedict's (reagent) √		accept sodium hydrogen carbonate instead of bicarbonate; accept ref to any alkali
			Green / yellow / Orange / red (precipitate) ✓ H→N-c-c <o h→n-c-c<o="" h→n-c<="" th=""><th></th><th>accept any appropriate colour change i.e brown / brick red)</th></o>		accept any appropriate colour change i.e brown / brick red)
	b	i	NH₂ group ✓ –COOH group ✓	2	accept diagram without bonds shown within NH_2 and COOH groups
		ii	peptide (bond) √	1	
		iii	hydrolysis √	1	

Q	Question		Expected Answers	Marks	Additional Guidance
2	b	iv	primary structure: sequence / order, of amino acids (in a polypeptide) ✓ secondary structure: Coiling/folding, of the, polypeptide/chain of amino acids/peptide chain/primary structure, / (α) helix / (β) pleated sheet / hydrogen bonds / between amino acids in (same) chain / (between) –NH and –CO / AVP e.g. random coiling ✓	2	reject ref to protein chains/structures joining together
			Total	12	

Mark Scheme

Question		on	Expected Answers	Marks	Additional Guidance
3	а	i	correct plots $\checkmark \checkmark$	2	5 correct plots = 2 marks
					3 - 4 correct plots = 1 mark
					(Accept accurate dots; tolerance +/- ½ square)
			appropriate line of heat fit .	1	smooth curve, no bairu/tram lines (teleranes $\pm 1/2$ equare)
				I	
					accept graph lines which follow IoB guidelines:
					Points may be joined with a curve of best fit if values are likely to
					fall on such a curve.
					Alternatively points may be joined with straight lines if the position
					or intermediate points can not be predicted reliably.
	b	i	correct data quote from graph = 5.6 (cm ³) \checkmark	1	accept 5.7 (cm ³)
		ii	5.6 ÷ 0.5 √	2	allow ecf from b(i);
			11.2 (cm³ min⁻¹) ✓		1 mark for conversion of time to minutes in rate formula
					1 mark for correct answer
			If 5.7 used from b(i):		
			5.7 ÷ 0.5;		Alternative: 1 mark for volume per sec (volume ÷ 30) =
			11.4 (cm ³ min ⁻¹)		0.187 (cm ³ sec ⁻¹)
					1 mark for conversion to rate per min i.e. $0.187 \times 60 = 11.2$
					(cm [°] min ⁻)
					accept tangents drawn on graph for calculation of rate

Question		on	Expected Answers	Marks	Additional Guidance
Q1 3	d d	on	Expected Answers [Level 1] Candidate shows a high level of understanding & includes a detailed description & explanation of two effects of temperature on enzyme activity, including at least four valid points, expressed clearly and logically. (4 - 5 marks) [Level 2] Candidate shows some understanding & includes a description & limited explanation of two effects of temperature on enzyme activity, expressed clearly and logically. (2 - 3 marks) [Level 3] Candidate shows some understanding & includes a description & limited explanation written in a sentence. (1 mark)	Marks 5	Additional Guidance valid points may include: ref to enzyme binding to active site/lock & key/active site specificity√ reference to 'optimum temperature' √ slow activity/reaction at low temperatures √ suitable ref movement/energy, of, molecules/enzyme/substrate, related to low temperature √ few collisions (between enzyme & substrate) / enzyme- substrate complex formed at low temperature√ more frequent collisions (between enzyme & substrate) / enzyme-substrate complex formed as temp increases √ (in warm temp) collisions (occur with more energy) to break bonds√ at high temperatures enzymes are denatured √ ref to enzyme/substrate complex can not form √ molecule vibrates breaking bonds (within enzyme molecule)
			Total	11	AVP e.g. suitable sketched graph ✓
			Total	11	

Qu	esti	on	Expected Answers	Marks	Additional Guidance
4	а	i	P = (eye piece) graticule √ Q = (stage) micrometer √	2	
		ii	any three from: line up scales of P with Q ✓ use Q to calibrate P before use / ref to calibration ✓ Scale Q is in mm/ cm / detail on scale (at specific magnification) ✓ convert divisions on P to actual measurements ✓ stage micrometers are usually calibrated for set magnification ✓	3	accept reference to calculations from Fig 4.1 <i>any three from:</i> 100 epg units = 25 x 0.1 mm / 2.5 mm ✓ 1 epg unit = 0.025 mm or 25 µm ✓ diameter of cell = number of epg units ✓ actual diameter of cell = diameter in epg units x 2.5µm ✓ ignore ref to 'measure cell' since this uses epg.
	b	i	diagnosis: anaemia ✓ explanation: fewer red blood cells observed / very few / paler red blood cells / less haemoglobin / irregular shaped red blood cells ✓	2	award diagnosis & explanation independently; explanation must link to photographs. reject 'few'
		ii	diagnosis: (lymphocytic) leukaemia ✓ explanation: large(r) numbers of lymphocytes/white blood cells / irregular shaped white blood cells / enlarged nuclei of, lymphocytes/white blood cells ✓	2	

Mark Scheme

(Question		Expected Answers	Marks	Additional Guidance
	L C		 any two from: reliability of test / possibility of error arising during testing ✓ human rights issues including, employment/insurance/mortgage facilities ✓ whether or not to pursue abortion ✓ how serious a defect has to be before abortion might be considered ✓ cost effectiveness of screening ✓ whether or not to risk starting a family ✓ AVP ✓ ✓ 	2	AVP e.g. religious issues / cultural issues AVP 'right to life of unborn foetus'
			Total	11	

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