## UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS GCE Advanced Subsidiary Level and GCE Advanced Level

## MARK SCHEME for the May/June 2009 question paper for the guidance of teachers

## 9700 BIOLOGY

9700/31

Paper 31 (Advanced Practical Skills 1), maximum raw mark 40

This mark scheme is published as an aid to teachers and candidates, to indicate the requirements of the examination. It shows the basis on which Examiners were instructed to award marks. It does not indicate the details of the discussions that took place at an Examiners' meeting before marking began, which would have considered the acceptability of alternative answers.

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

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Question	Expected	d Answe	rs						Additional Guidance	Mark
1 (a) (i) Re	cord colou	r of pap	er and pl	H for gluco	se and	ethanol.				·
MMO collection 2	with refer	ence to g	lucose ar	nd ethanol	record a	a colour for	each;			[1]
0011001101112	one colou	ır matche	s one pH	or betwee	n two pł	d values:			Credit ONLY the given pH values	[1]
	pH	5.2	5.5	5.8	6.1	6.4	6.7		Credit less than/<5.2 with yellow or	1.,
	colours	brown/ orange/		pink/ brown	pink/ purple	<b>I</b>	c/magenta/		more than/>6.7 with purple/AW	
	OR (if no from scale				to gluce	ose and et	hanol two pH va	lues		
(ii) De	cide which	other co	oncentra	tions to m	ake and	d complete	the table.			·
MMO decisions 3	(%) 0 and 50 or 0, 1	•	•	which are	evenly/s	serially spa	aced e.g. 0, 20,	30, 40,		[1]
	correct vo	orrect volumes used to dilute up to 10 cm <sup>3</sup> AND correct %;							[1]	
	(tubes list	,			% to mo	ost concent	trated/highest %		Ignore where 0 is listed	[1]

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(iii) Pre	epare space to record colour of each piece of paper and pH.		
PDO recording 2	single table AND all cells drawn AND %/percent(age) (top or left of data heading only);	heading heading heading heading heading Do not credit if % in body of table	[1]
	(headings) colour AND pH;	Do not credit if pH/colour in body of table	[1]
MMO collection 2	two different colours for two tubes recorded;	Collection of colour for two tubes. Credit colour differences such as light orange vs orange	[1]
	(collected data/colours or pH) clearly for 1 min(ute)/start and 10 min(utes)/end;	Credit colours only or pH values only – looking for clear collection of 1 minute and 10 minutes Credit for one tube of data for 1 min and 10 min as minimum	[1]
(b) (i) Ide	entify a significant error – read complete answer for any correct one.		
ACE interpretation 1	judging colour/matching exact colours/colours very close together/pH paper narrow scale/colours not on scale/between colours/identification of colours; idea of timing not the same/different/described; loss of CO <sub>2</sub> /gas/AW;	Do not credit timing unqualified	[max 1]
(ii) Sta	ate degree of uncertainty (of syringes).	•	1
ACE interpretation 1	+/– AND half total division AND cm³;	Error with one reading is +/– half the smallest division with correct units as use syringe to measure single volume and release al contents	[1]

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(c) (i) Plo	0	x-axis temp(erature) (/)°C AND y-axis (number/no. of) bubbles/min or	Do NOT credit bubbles min <sup>1</sup>	[1]
,		per min or min <sup>-1</sup> ;		
	S	scale as 10°C to 2 cm and 5 to 2 cm; Credit origin other than 0 e.g. 5/10/15 if labelled Credit unlabelled origin only if should be 0	Do not credit S if awkward scale or if less than half grid on y or x axis	[1]
	Р	plotting correct points using crosses/dots in circles only; Do not credit if an extra point plotted at 25°C.  No cross larger than x or more than one blob larger than o.  All plots must be on horizontal lines except for the 4 to 2 cm scale points within a square of the intersection/centre of dot must not touch horizontal lines.	Do not credit P plotting if awkward scale or if only blobs/dots/blobs in circles Do not credit dot with cross  Credit x in circles	[1]
	L	line of best fit (no more than 2 points on one side)/points joined with straight line; Quality – line no thicker than 1 mm thick max Complete line should be smooth/not feathery.	Credit line of best fit – no extrapolation Joins point to point no extrapolation beyond first and last points	[1]
(ii) Est	timate	e enzyme activity at 25°C.		
ACE interpretation 1	corre min	ect reading using candidate's graph at 25°C AND bubbles per minute or -1;	Credit whole number of bubbles only Credit 0.5 up or down	[1]

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ACE improvements 1	control of any variable/use a water bath/same type of yeast/same volume of yeast/keep time the same/stagger the start/have separate experiments/(keep pH same) using buffer;	Credit in either accuracy or reliability  Do not credit ref to enzymes or amount of yeast	[1]
	Accuracy: collect volume using measuring cylinder/gas syringe/video to count bubbles/AW;	Accuracy: (change method of measuring to obtain results as close as possible to the true value)	[1]
	Reliability 1: increase number or range of temperatures/2 extra named examples;	Reliable: (to have results which are as repeatable as possible)	[1]
	Reliability 2: repeat more/several times/twice/obtain three readings/(at each temp);		[1]
	Reliability 3: calculate mean;	Credit only two reliability marks	[1]
			[max 3]
(d) State w	hether you think the hypothesis is supported by the student's results. Expl	ain your answer.	
ACE conclusion 2	hypothesis true/yes/OR re-states the hypothesis OR partly true/true but only; Either	Needs clear statement Do not credit idea that totally wrong	[1]
	(true for) 15 to 40°C as increases from 5 to 18/ or any two correct temps within 15 and 40/41 with two correct numbers of bubbles/may have two temps and difference in number of bubbles/calculated rate of increase/gradient;	Credit temp as long as units are present once	[max 1]
	OR ONLY TRUE: 15 to 40°C as increases from 5 to 18/ or any two correct temps with two correct numbers of bubbles/may have two		

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Question	Expected Answers		Additional Guidance	Mark
2 (a) Draw a	large, low power, plan diagram of qua	rter. Label phloem and xylem in a vas	cular bundle.	1
PDO layout 1	clear, sharp, unbroken lines AND no sh cannot fit totally within the 6 cm by 6 cm			[1]
MMO collection 3	no cells AND outline irregular;			[1]
	clear layer/layers below cortex AND corpoint of layer below; Ignore epidermis (			[1]
	at least 3 complete vascular bundles AN twice length of smallest vascular bundle Do not credit if whole section drawn			[1]
MMO decision	phloem and xylem are labelled on the la as shown by candidate;			[1]
(b) (ii) Cal	culate mean width of the cells in micro	ometres. Mark clearly the cells you use	ed. Show your working.	
MMO collection 1	cell(s) marked on Fig 2.2 AND Credit separate or	selected cell(s) measured with mm or cm shown;  Do not credit if line is not on the fig.	Check recorded cell measurements are at or between 0.5 to 5.5 cm or 5 mm to 55 mm Ignore more sig. figs.  OR if single line is drawn	[1]
	one line across cells selected	Do not credit measurements in metres or micrometres.	check measurement of line is less than 170 mm/	
MMO decision 1	5 or more complete cells measured;			[1]
	Credit even if not shown on Fig. 2.2. Do not credit if partial cells included.			
PDO display 2	shows any number of measurements addivided by number of measurements/ce		Ignore the answer	[1]

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	divides by 400 and multiplies by (cm) 10 000/10 <sup>4</sup> or (mm) 1000/10 <sup>3</sup> ;	E.g. cell 1 = 20 mm cell 2 = 21 mm cell 3 = 17 mm cell 4 = 16 mm cell 5 = 21 mm	[1]
		Mean = $\frac{20 + 21 + 17 + 16 + 21}{5}$ = 19 mm 19 mm = 19 × 1000 µm = 19 000 Mag. = ×400 Actual = $\frac{19000}{400}$ = 475 Must show division by 400	
(ii) Lak	pel with a line and the letter X, the area from which section may have been		
ACE conclusion 1	central vascular tissue root cap	Credit anywhere between bracket for central vascular tissue and end of label line for root cap  Do not credit if longitudinal section shown	[1]

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PDO layout 1	clear, sharp, AND unbroken lines	no shading	AND	cannot fit totally into the 6 cm by 6 cm grid;
	Ignore cell walls crossing over as cells appear to overlap.	Do not credi cytoplasm or walls.		
MMO collection 1	3 complete cells marked o			nplete cells touching; onal incomplete cells.
PDO recording 1	DO recording  2 nuclei each drawn near Nucleus not in centre of c Do not credit if additional plasmodesmata have bee		f cell; elles or cell w	vall details e.g.
MMO decisions 2	space between outer wall	and cytoplasr	n shown in pa	ert of two cells;
	Ignore labelling.  nucleus and cell wall corre	ectly labelled;		
	Do not credit nucleolus.			

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PDO recording 1	organise as a table/ Venn diagram/ ruled connected boxes	headed	comparative stateme opposite each other;		[1]
ACE			T	_	[max
interpretation 3	feature	Fig. 2.2	Fig. 2.4		
	nuclei Ignore ref. to nuclear membrane/size	few/less distinct/less stained	more/dense/obvious/ more stained;		
	chromosomes/dividing cells/mitosis/anaphase	present	not visible		
	interphase	fewer in interphase	all in interphase;		
	cytoplasm/ plasma membrane	more/fills cells not visible/pulled away	less/does not fill cells/ visible/pulled away;		
	size/ number of cells Credit more (tightly packed)	smaller/calculated mean/ecf/more	larger/calculated more than Fig 2.2/fewer;		
	shape of cells	have oval/corners/angular/h exagonal	rounded/AW;		
	(Intercellular) spaces/gaps	fewer/more/larger; Credit either way			

[Total: 19]